Electronic Transport in Metallic Systems and Generalized Kinetic Equations*

A. L. Kuzemsky
Bogoliubov Laboratory of Theoretical Physics,
Joint Institute for Nuclear Research,
141980 Dubna, Moscow Region, Russia.
E-mail:kuzemsky@theor.jinr.ru
http://theor.jinr.ru/~kuzemsky

Abstract

This paper reviews some selected approaches to the description of transport properties, mainly electroconductivity, in crystalline and disordered metallic systems. A detailed qualitative theoretical formulation of the electron transport processes in metallic systems within a model approach is given. Generalized kinetic equations which were derived by the method of the nonequilibrium statistical operator are used. Tight-binding picture and modified tight-binding approximation (MTBA) were used for describing the electron subsystem and the electron-lattice interaction correspondingly. The low- and high-temperature behavior of the resistivity was discussed in detail. The main objects of discussion are nonmagnetic (or paramagnetic) transition metals and their disordered alloys. The choice of topics and the emphasis on concepts and model approach makes it a good method for a better understanding of the electrical conductivity of the transition metals and their disordered binary substitutional alloys, but the formalism developed can be applied (with suitable modification), in principle, to other systems. The approach we used and the results obtained complements the existent theories of the electrical conductivity in metallic systems. The present study extends the standard theoretical format and calculation procedures in the theories of electron transport in solids.

Keywords: Transport phenomena in solids; electrical conductivity in metals and alloys; transition metals and their disordered alloys; tight-binding and modified tight-binding approximation; method of the nonequilibrium statistical operator; generalized kinetic equations.

*International Journal of Modern Physics B (IJMPB), Volume: 25, Issue: 23-24 (2011) p.3071-3183
## Contents

1 Introduction  \hspace{1cm} 2

2 Metals and Nonmetals. Band Structure  \hspace{1cm} 3

3 Many-Particle Interacting Systems and Current operator  \hspace{1cm} 8

4 Tight-Binding and Modified Tight-Binding Approximation  \hspace{1cm} 11
   4.1 Tight-binding approximation  \hspace{1cm} 11
   4.2 Interacting electrons on a lattice and the Hubbard model  \hspace{1cm} 13
   4.3 Current operator for the tight-binding electrons  \hspace{1cm} 14
   4.4 Electron-lattice interaction in metals  \hspace{1cm} 17
   4.5 Modified tight-binding approximation  \hspace{1cm} 21

5 Charge and Heat Transport  \hspace{1cm} 23
   5.1 Electrical resistivity and Ohm law  \hspace{1cm} 23
   5.2 Drude-Lorentz model  \hspace{1cm} 24
   5.3 The low- and high-temperature dependence of conductivity  \hspace{1cm} 27
   5.4 Conductivity of alloys  \hspace{1cm} 30
   5.5 Magnetoresistance and the Hall effect  \hspace{1cm} 32
   5.6 Thermal conduction in solids  \hspace{1cm} 34

6 Linear Macroscopic Transport Equations  \hspace{1cm} 35

7 Statistical Mechanics and Transport Coefficients  \hspace{1cm} 37
   7.1 Variational principles for transport coefficients  \hspace{1cm} 38
   7.2 Transport theory and electrical conductivity  \hspace{1cm} 38

8 The Method of Time Correlation Functions  \hspace{1cm} 39
   8.1 Linear response theory  \hspace{1cm} 40
   8.2 Green functions in the theory of irreversible processes  \hspace{1cm} 45
   8.3 The electrical conductivity tensor  \hspace{1cm} 49
   8.4 Linear response theory: pro et contra  \hspace{1cm} 51

9 The Nonequilibrium Statistical Operator Method and Kinetic Equations  \hspace{1cm} 56

10 Generalized Kinetic Equations and Electroconductivity  \hspace{1cm} 57
   10.1 Basic formulas  \hspace{1cm} 57
   10.2 Electrical conductivity  \hspace{1cm} 59

11 Resistivity of Transition Metal with Non-spherical Fermi Surface  \hspace{1cm} 61
   11.1 Generalized kinetic equations  \hspace{1cm} 62
   11.2 Temperature dependence of R  \hspace{1cm} 65
   11.3 Equivalence of NSO approach and Kubo formalism  \hspace{1cm} 67
   11.4 High-temperature resistivity and MTBA  \hspace{1cm} 69

12 Resistivity of Disordered Alloys  \hspace{1cm} 77

13 Discussion  \hspace{1cm} 85
1 Introduction

Transport properties of matter constitute the transport of charge, mass, spin, energy and momentum. It has not been our aim to discuss all the aspects of the charge and thermal transport in metals. We are concerned in the present work mainly with some selected approaches to the problem of electric charge transport (mainly electroconductivity) in crystalline and disordered metallic systems. Only the fundamentals of the subject are treated. In the present work we aim to obtain a better understanding of the electrical conductivity of the transition metals and their disordered binary substitutional alloys both by themselves and in relationship to each other within the statistical mechanical approach. Thus our consideration will concentrate on the derivation of generalized kinetic equations suited for the relevant models of metallic systems. The problem of the electronic transport in solids is an interesting and actual part of the physics of condensed matter. It includes the transport of charge and heat in crystalline and disordered metallic conductors of various nature. Transport of charge is connected with an electric current. Transport of heat has many aspects, main of which is the heat conduction. Other important aspects are the thermoelectric effects. The effect, termed Seebeck effect, consists of the occurrence of a potential difference in a circuit composed of two distinct metals at different temperatures. Since the earlier seminal attempts to construct the quantum theory of the electrical, thermal and thermoelectric and thermomagnetic transport phenomena there is a great interest in the calculation of transport coefficients in solids in order to explain the experimental results as well as to get information on the microscopic structure of materials.

A number of physical effects enter the theory of quantum transport processes in solids at various density of carriers and temperature regions. A variety of theoretical models has been proposed to describe these effects. Theory of the electrical and heat conductivities of crystalline and disordered metals and semiconductors have been developed by many authors during last decades. There exist a lot of theoretical methods for the calculation of transport coefficients as a rule having a fairly restricted range of validity and applicability. In the present work the description of the electronic and some aspects of heat transport in metallic systems are briefly reviewed, and the theoretical approaches to the calculation of the resistance at low and high temperature are surveyed. As a basic tool we use the method of the nonequilibrium statistical operator (NSO). It provides a useful and compact description of the transport processes. Calculation of transport coefficients within NSO approach was presented and discussed in the author’s work. The present paper can be considered as the second part of the review article. The close related works on the study of electronic transport in metals are briefly summarized in the present work. It should be emphasized that the choice of generalized kinetic equations among all other methods of the theory of transport in metals is related with its efficiency and compact form. They are an alternative (or complementary) tool for studying of transport processes, which complement other existing methods.

Due to the lack of space many interesting and actual topics must be omitted. An important and extensive problem of thermoelectricity was mentioned very briefly; thus it has not been possible to do justice to all the available theoretical and experimental results of great interest. The thermoelectric and transport properties of the layered high-$T_c$ cuprates were reviewed by us already in the extended review article. Another interesting aspect of transport in solids which we did not touched is the spin transport. The spin degree of freedom of charged carriers in metals and semiconductors has attracted in last decades big attention and continues to play a key role in the development of many applications, establishing a field that is now known as spintronics. Spin transport and manipulation in not only ferromagnets but also nonmagnetic materials are currently being studied actively in a variety of artificial structures and designed new materials. This enables the fabrication of spintronic...
properties on intention. A study on spintronic device structures was reported as early as in late sixties. Studies of spin-polarized internal field emission using the magnetic semiconductor EuS sandwiched between two metal electrodes opened a new epoch in electronics. Since then, many discoveries have been made using spintronic structures. Among them is giant magnetoresistance in magnetic multilayers. Giant magnetoresistance has enabled the realization of sensitive sensors for hard-disk drives, which has facilitated successful use of spintronic devices in everyday life. There is big literature on this subject and any reasonable discussion of the spin transport deserves a separate extended review. We should mention here that some aspects of the spin transport in solids were discussed by us in Refs. In the present study a qualitative theory for conductivity in metallic systems is developed and applied to systems like transition metals and their disordered alloys. The nature of transition metals is discussed in details and the tight-binding approximation and method of model Hamiltonians are described. For the interaction of the electron with the lattice vibrations we use the modified tight-binding approximation (MTBA). Thus this approach can not be considered as the first-principle method and has the same shortcomings and limitations as describing a transition metal within the Hubbard model. In the following pages, we shall present a formulation of the theory of the electrical transport in the approach of the nonequilibrium statistical operator. Because several other sections in this review require a certain background in the use of statistical-mechanical methods, physics of metals, etc., it was felt that some space should be devoted to this background. Sections 2 to 8 serves as an extended introduction to the core sections 9-12 of the present paper. Thus those sections are intended as a brief summary and short survey of the most important notions and concepts of charge transport (mainly electroconductivity) for the sake of a self-contained formulation. We wish to describe those concepts which have proven to be of value, and those notions which will be of use in clarifying subtle points.

First, in order to fix the domain of study, we must briefly consider the various formulation of the subject and introduce the basic notions of the physics of metals and alloys.

### 2 Metals and Nonmetals. Band Structure

The problem of the fundamental nature of the metallic state is of long standing. It is well known that materials are conveniently divided into two broad classes: insulators (nonconducting) and metals (conducting). More specific classification divided materials into three classes: metals, insulators, and semiconductors. The most characteristic property of a metal is its ability to conduct electricity. If we classify crystals in terms of the type of bonding between atoms, they may be divided into the following five categories (see Table 1).

Ultimately we are interested in studying all of the properties of metals. At the outset it is natural to approach this problem through studies of the electrical conductivity and closely related

---

**Table 1: Five Categories of Crystals**

| Type of Crystal                  | substances                              |
|---------------------------------|-----------------------------------------|
| ionic                           | alkali halides, alkaline oxides, etc.   |
| homopolar bounded (covalent)    | diamond, silicon, etc.                  |
| metallic                         | various metals and alloys               |
| molecular                        | Ar, He, O2, H2, CH4, etc.               |
| hydrogen bonded                  | ice, KH2PO4, fluorides, etc.            |
problem of the energy band structure.\textsuperscript{32,35}

The energy bands in solids represent the fundamental electronic structure of a crystal just as the atomic term values represent the fundamental electronic structure of the free atom. The behavior of an electron in one-dimensional periodic lattice is described by Schrödinger equation

\begin{equation}
\frac{d^2 \psi}{dx^2} + \frac{2m}{\hbar^2} (E - V) \psi = 0,\end{equation}

(2.1)

where $V$ is periodic with the period of the lattice $a$. The variation of energy $E(k)$ as a function of quasi-momentum within the Brillouin zones, and the variation of the density of states $D(E)dE$ with energy, are of considerable importance for the understanding of real metals. The assumption that the potential $V$ is small compared with the total kinetic energy of the electrons (approximation of nearly free electrons) is not necessarily true for all metals. The theory may also be applied to cases where the atoms are well separated, so that the interaction between them is small. This treatment is usually known as the approximation of "tight binding"\textsuperscript{13} In this approximation the behavior of an electron in the region of any one atom being only slightly influenced by the field of the other atoms\textsuperscript{33,52} Considering a simple cubic structure, it is found that the energy of an electron may be written as

\begin{equation}
E(k) = E_a - t_\alpha - 2t_\beta (\cos(k_xa) + \cos(k_ya) + \cos(k_za)),\end{equation}

(2.2)

where $t_\alpha$ is an integral depending on the difference between the potentials in which the electron moves in the lattice and in the free atom, and $t_\beta$ has a similar significance\textsuperscript{33,52} (details will be given below). Thus in the tight-binding limit, when electrons remain to be tightly bound to their original atoms, the valence electron moves mainly about individual ion core, with rare hopping from ion to ion. This is the case for the $d$-electrons of transition metals. In the typical transition metal the radius of the outermost $d$-shell is less than half the separation between the atoms. As a result, in the transition metals the $d$-bands are relatively narrow. In the nearly free-electron limit the bands are derived from the $s$- and $p$-shells which radii are significantly larger than half the separation between the atoms. Thus, according this simplified picture simple metals have nearly-free-electron energy bands (see Fig.1). Fortunately in the case of simple metals the combined results of the energy band calculation and experiment have indicated that the effects of the interaction between the electrons and ions which make up the metallic lattice is extremely weak. It is not the case for transition metals and their disordered alloys\textsuperscript{53,54}

An obvious characterization of a metal is that it is a good electrical and thermal conductor.\textsuperscript{1,2,13,55,56} Without considering details it is possible to see how the simple Bloch picture outlined above accounts for the existence of metallic properties, insulators, and semiconductors. When an electric
current is carried, electrons are accelerated, that is promoted to higher energy levels. In order that this may occur, there must be vacant energy levels, above that occupied by the most energetic electron in the absence of an electric field, into which the electron may be excited. At some conditions there exist many vacant levels within the first zone into which electrons may be excited. Conduction is therefore possible. This case corresponds with the noble metals. It may happen that the lowest energy in the second zone is lower than the highest energy in the first zone. It is then possible for electrons to begin to occupy energy contained within the second zone, as well as to continue to fill up the vacant levels in the first zone and a certain number of levels in the second zone will be occupied. In this case the metallic conduction is possible as well. The polyvalent metals are materials of this class.

If, however, all the available energy levels within the first Brillouin zone are full and the lowest possible electronic energy at the bottom of the second zone is higher than the highest energy in the first zone by an amount $\Delta E$, there exist no vacant levels into which electrons may be excited. Under these conditions no current can be carried by the material and an insulating crystal results. For another class of crystals, the zone structure is analogous to that of insulators but with a very small value of $\Delta E$. In such cases, at low temperatures the material behaves as an insulator with a higher specific resistance. When the temperature increases a small number of electrons will be thermally excited across the small gap and enter the second zone, where they may produce metallic conduction. These substances are termed semiconductors [13, 55, 56], and their resistance decreases with rise in temperature in marked contrast to the behavior of real metals (for a detailed review of semiconductors see Refs. [57, 58]).

The differentiation between metal and insulator can be made by measurement of the low frequency electrical conductivity near $T = 0$ K. For the substance which we can refer as an ideal insulator the electrical conductivity should be zero, and for metal it remains finite or even becomes infinite. Typical values for the conductivity of metals and insulators differ by a factor of the order $10^{10} - 10^{15}$. So big difference in the electrical conductivity is related directly to a basic difference in the structural and quantum chemical organization of the electron and ion subsystems of solids. In an insulator the position of all the electrons are highly connected with each other and with the crystal lattice and a weak direct current field cannot move them. In a metal this connection is not so effective and the electrons can be easily displaced by the applied electric field. Semiconductors occupy an intermediate position due to the presence of the gap in the electronic spectra.

An attempt to give a comprehensive empirical classification of solids types was carried out by Zeitz [55] and Kittel [56]. Zeitz reanalyzed the generally accepted classification of materials into three broad classes: insulators, metals and semiconductors and divided materials into five categories: metals, ionic crystals, valence or covalent crystals, molecular crystals, and semiconductors. Kittel added one more category: hydrogen-bonded crystals. Zeitz also divided metals further into two major classes, namely, monoatomic metals and alloys.

Alloys constitute an important class of the metallic systems. [25, 49, 55, 56, 59, 61] A metal alloy is a mixed material that has metal properties and is made by melting at least one pure metal along with another pure chemical or metal. Examples of metal alloys $Cu - Zn$, $Au - Cu$ and an alloy of carbon and iron, or copper, antimony and lead. Brass is an alloy of copper and zinc, and bronze is an alloy of copper and tin. Alloys of titanium, vanadium, chromium and other metals are used in many applications. The titanium alloys (interstitial solid solutions) form a big variety of equilibrium phases. Alloy metals are usually formed to combine properties of metals and the exact proportion of metals in an alloy will change the characteristic properties of the alloy. We confine ourselves to those alloys which may be regarded essentially as very close to pure metal with the properties intermediate to those of the constituents.

There are different types of monoatomic metals within the Bloch model for the electronic struc-
ture of a crystal: simple metals, alkali metals, noble metals, transition metals, rare-earth metals, divalent metals, trivalent metals, tetravalent metals, pentavalent semimetals, lantanides, actinides and their alloys. The classes of metals according to crude Bloch model provide us with a simple qualitative picture of variety of metals. This simplified classification takes into account the state of valence atomic electrons when we decrease the interatomic separation towards its bulk metallic value. Transition metals have narrow $d$-bands in addition to the nearly-free-electron energy bands of the simple metals.$^{53, 54}$ In addition, the correlation of electrons plays an essential role.$^{53, 54, 62}$ The Fermi energy lies within the $d$-band so that the $d$-band is only partially occupied. Moreover the Fermi surface have much more complicated form and topology. The concrete calculations of the band structure of many transition metals ($Nb$, $V$, $W$, $Ta$, $Mo$, etc.) can be found in Refs.$^{13, 32–35, 53, 63–66}$ and in Landolt-Bornstein reference books.$^{60, 67}$

The noble metal atoms have one $s$-electron outside of a just completed $d$-shell. The $d$-bands of the noble metals lie below the Fermi energy but not too deeply. Thus they influence many of the physical properties of these metals. It is, in principle, possible to test the predictions of the single-electron band structure picture by comparison with experiment. In semiconductors it has been performed with the measurements of the optical absorption, which gives the values of various energy differences within the semiconductor bands. In metals the most direct approach is related to the experiments which studied the shape and size of the Fermi surfaces. In spite of their value, these data represent only a rather limited scope in comparison to the many properties of metals which are not so directly related to the energy band structure. Moreover, in such a picture there are many weak points: there is no sharp boundary between insulator and semiconductor, the theoretical values of $\Delta E$ have discrepancies with experiment, the metal-insulator transition$^{68}$ cannot be described correctly, and the notion "simple" metal have no single meaning.$^{69}$ The crude Bloch model even met more serious difficulties when it was applied to insulators. The improved theory of insulating state was developed by Kohn$^{70}$ within a many-body approach. He proposed a new and more comprehensive characterization of the insulating state of matter. This line of reasoning was continued further in Refs.$^{68, 71, 72}$ on a more precise and firm theoretical and experimental basis.

Anderson$^{50}$ gave a critical analysis of the Zeitz and Kittel classification schemes. He concluded that "in every real sense the distinction between semiconductors and metals or valence crystals as to type of binding, and between semiconductor and any other type of insulator as to conductivity, is entirely artificial; semiconductors do not represent in any real sense a distinct class of crystal"$^{60}$ (see, however Refs.$^{13, 23, 38, 55, 56}$). Anderson has pointed also the extent to which the standard classification falls. His conclusions were confirmed by further development of solid state physics. During the last decades a lot of new substances and materials were synthesized and tested. Their conduction properties and temperature behavior of the resistivity are differed substantially and constitute a difficult task for consistent classification.$^{73}$ (see Fig.1). Bokij$^{74}$ carried out an interesting analysis of notions "metals" and "nonmetals" for chemical elements. According to him, there are typical metals ($Cu$, $Au$, $Fe$) and typical nonmetals ($O$, $S$, halogens), but the boundary between them and properties determined by them are still an open question. The notion "metal" is defined by a number of specific properties of the corresponding elemental substances, e.g. by high electrical conductivity and thermal capacity, the ability to reflect light waves(luster), plasticity, and ductility. Bokij emphasizes,$^{74}$ that when defining the notion of a metal, one has also to take into account the crystal structure. As a rule, the structure of metals under normal conditions are characterized by rather high symmetries and high coordination numbers (c.n.) of atoms equal to or higher than eight, whereas the structures of crystalline nonmetals under normal conditions are characterized by lower symmetries and coordination numbers of atoms (2-4).

It is worth noting that such topics like studies of the strongly correlated electronic systems$^{52}$ high-$T_c$ superconductivity$^{52}$ colossal magnetoresistance$^5$ and multiferroicity$^5$ have led to a new
development of solid state physics during the last decades. Many transition-metal oxides show very large ("colossal") magnitudes of the dielectric constant and thus have immense potential for applications in modern microelectronics and for the development of new capacitance-based energy-storage devices. These and other interesting phenomena to a large extent first have been revealed and intensely investigated in transition-metal oxides. The complexity of the ground states of these materials arises from strong electronic correlations, enhanced by the interplay of spin, orbital, charge and lattice degrees of freedom. These phenomena are a challenge for basic research and also bear big potentials for future applications as the related ground states are often accompanied by so-called "colossal" effects, which are possible building blocks for tomorrow’s correlated electronics. The measurement of the response of transition-metal oxides to electric fields is one of the most powerful techniques to provide detailed insight into the underlying physics that may comprise very different phenomena, e.g., charge order, molecular or polaronic relaxations, magnetocapacitance, hopping charge transport, ferroelectricity or density-wave formation. In the recent work, authors thoroughly discussed the mechanisms that can lead to colossal values of the dielectric constant, especially emphasizing effects generated by external and internal interfaces, including electronic phase separation. The authors of the work studied the materials showing so-called colossal dielectric constants (CDC), i.e. values of the real part of the permittivity exceeding 1000. Since long, materials with high dielectric constants are in the focus of interest, not only for purely academic reasons but also because new high-ε′ materials are urgently sought after for the further development of modern electronics. In addition, authors of the work provided a detailed overview and discussion of the dielectric properties of CaCu₃Ti₄O₁₂ and related systems, which is today’s most investigated material with colossal dielectric constant. Also a variety of
Table 2. Metallic and Semimetallic Elements

| item                        | number | elements                        |
|-----------------------------|--------|---------------------------------|
| alkali metals               | 5      | Li, Na, K, Rb, Cs               |
| noble metals                | 3      | Cu, Ag, Au                      |
| polyvalent simple metals    | 11     | Be, Mg, Zn, Hg, Al, Ga, In, Tl, Sn, Pb |
| alkali-earth metals         | 4      | Ca, Sr, Ba, Ra                  |
| semi-metals                 | 4      | As, Sb, Bi, graphite            |
| transition metals           | 23     | Fe, Ni, Co, etc.                |
| rare earths                 | 14     |                                |
| actinides                   | 4      |                                |

Further transition-metal oxides with large dielectric constants were treated in detail, among them the system \(La_{2-x}Sr_xNiO_4\) where electronic phase separation may play a role in the generation of a colossal dielectric constant. In general, for the miniaturization of capacitive electronic elements materials with high-\(\varepsilon'\) are prerequisite. This is true not only for the common silicon-based integrated-circuit technique but also for stand-alone capacitors. Nevertheless, as regards to metals, the workable practical definition of Kittel can be adopted: metals are characterized by high electrical conductivity, so that a portion of electrons in metal must be free to move about. The electrons available to participate in the conductivity are called conduction electrons. Our picture of a metal, therefore, must be that it contains electrons which are free to move, and which may, when under the influence of an electric field, carry a current through the material.

In summary, the 68 naturally occurring metallic and semimetallic elements\(^\text{[19]}\) can be classified as it is shown in Table 2.

### 3 Many-Particle Interacting Systems and Current operator

Let us now consider a general system of \(N\) interacting electrons in a volume \(\Omega\) described by the Hamiltonian

\[
H = \left( \sum_{i=1}^{N} \frac{\vec{p}_i^2}{2m} + \sum_{i=1}^{N} U(\vec{r}_i) \right) + \frac{1}{2} \sum_{i \neq j} v(\vec{r}_i - \vec{r}_j) = H_0 + H_1. \tag{3.1}
\]

Here \(U(\vec{r})\) is a one-body potential, e.g., an externally applied potential like that due to the field of the ions in a solid, and \(v(\vec{r}_i - \vec{r}_j)\) is a two-body potential like the Coulomb potential between electrons. It is essential that \(U(\vec{r})\) and \(v(\vec{r}_i - \vec{r}_j)\) do not depend on the velocities of the particles. It is convenient to introduce a ”quantization” in a continuous space\(^\text{[27,49]}\) via the operators \(\Psi^\dagger(\vec{r})\) and \(\Psi(\vec{r})\) which create and destroy a particle at \(\vec{r}\). In terms of \(\Psi^\dagger\) and \(\Psi\) we have

\[
H = \int d^3r \Psi^\dagger(\vec{r}) \left( -\frac{\nabla^2}{2m} + U(\vec{r}) \right) \Psi(\vec{r}) \tag{3.2}
\]

\[
+ \frac{1}{2} \int d^3r d^3r' \Psi^\dagger(\vec{r}) \Psi^\dagger(\vec{r}') v(\vec{r} - \vec{r}') \Psi(\vec{r}') \Psi(\vec{r})
\]

Studies of flow problems lead to the continuity equation\(^\text{[28,42]}\)

\[
\frac{\partial n(\vec{r}, t)}{\partial t} + \nabla \cdot \vec{j} = 0. \tag{3.3}
\]
This equation based on the concept of conservation of certain extensive variable. In nonequilibrium thermodynamics\textsuperscript{12} the fundamental flow equations are obtained using successively mass, momentum, and energy as the relevant extensive variables. The analogous equation are known from electromagnetism. The central role plays a global conservation law of charge, \( \dot{q}(t) = 0 \), for it refers to the total charge in a system. Charge is also conserved locally.\textsuperscript{80} This is described by Eq. (3.3), where \( n(\vec{r}, t) \) and \( \vec{j} \) are the charge and current densities, respectively.

In quantum mechanics there is the connection of the wavefunction \( \psi(\vec{r}, t) \) to the particle mass-probability current distribution \( \vec{J} \)

\[ \vec{J}(\vec{r}, t) = \frac{\hbar}{2m} (\psi^* \nabla \psi - \psi \nabla \psi^*), \tag{3.4} \]

where \( \psi(\vec{r}, t) \) satisfy the time-dependent Schrödinger equation\textsuperscript{79,81}

\[ i\hbar \frac{\partial}{\partial t} \psi(\vec{r}, t) = H \psi(\vec{r}, t). \tag{3.5} \]

Consider the motion of a particle under the action of a time-independent force determined by a real potential \( V(\vec{r}) \). Equation (3.5) becomes

\[ \left( \frac{\vec{p}^2}{2m} + V \right) \psi = \frac{\hbar}{2m} \nabla^2 \psi + V \psi = i\hbar \frac{\partial}{\partial t} \psi. \tag{3.6} \]

It can be shown that for the probability density \( n(\vec{r}, t) = \psi^* \psi \) we have

\[ \frac{\partial n}{\partial t} + \nabla \vec{J} = 0. \tag{3.7} \]

This is the equation of continuity and it is quite general for real potentials. The equation of continuity mathematically states the local conservation of particle mass probability in space.

A thorough consideration of a current carried by a quasi-particle for a uniform gas of fermions, containing \( N \) particles in a volume \( \Omega \), which was assumed to be very large, was performed within a semi-phenomenological theory of Fermi liquid\textsuperscript{82} This theory describes the macroscopic properties of a system at zero temperature and requires knowledge of the ground state and the low-lying excited states. The current carried by the quasi-particle \( \vec{k} \) is the sum of two terms: the current which is equal to the velocity \( v_k \) of the quasi-particle and the backflow of the medium.\textsuperscript{82} The precise definition of the current \( \vec{J} \) in an arbitrary state \( |\varphi \rangle \) within the Fermi liquid theory is given by

\[ J = \langle \varphi \sum_i \frac{p_i}{m} |\varphi \rangle, \tag{3.8} \]

where \( p_i \) is the momentum of the \( i \)th particle and \( m \) its bare mass. To measure \( J \) it is necessary to use a reference frame moving with respect to the system with the uniform velocity \( \hbar q/m \). The Hamiltonian in the rest frame can be written

\[ H = \sum_i \frac{p_i^2}{2m} + V. \tag{3.9} \]

It was assumed that \( V \) depends only on the positions and the relative velocities of the particles; it is not modified by a translation. In the moving system only the kinetic energy changes; the apparent Hamiltonian becomes

\[ H_q = \sum_i \frac{(p_i - \hbar q)^2}{2m} + V = H - \hbar q \sum_i \frac{p_i}{m} + N \frac{(\hbar q)^2}{2m}. \tag{3.10} \]
Taking the average value of $H_q$ in the state $|\varphi\rangle$, and let $E_q$ be the energy of the system as seen from the moving reference frame, one find in the $\lim q \to 0$

$$\frac{\partial E_q}{\partial q_\alpha} = -\hbar \langle \varphi | \sum_i \frac{p_i m}{m} | \varphi \rangle = -\hbar J_{\alpha},$$

(3.11)

where $\alpha$ refers to one of the three coordinates. This expression gives the definition of current in the framework of the Fermi liquid theory. For the particular case of a translationally invariant system the total current is a constant of the motion, which commutes with the interaction $V$ and which, as a consequence, does not change when $V$ is switched on adiabatically. For the particular state containing one quasi-particle $\vec{k}$ the total current $J_k$ is the same as for the ideal system

$$J_k = \frac{(hk)}{m}.$$  

(3.12)

This result is a direct consequence of Galilean invariance.

Let us consider now the many-particle Hamiltonian (3.2)

$$H = H_1 + H_2.$$  

(3.13)

It will also be convenient to consider density of the particles in the following form

$$n(\vec{r}) = \sum_i \delta(\vec{r} - \vec{r}_i).$$

The Fourier transform of the particle density operator becomes

$$n(\vec{q}) = \int d^3r \exp(-i\vec{q}\vec{r}) \sum_i \delta(\vec{r} - \vec{r}_i) = \sum_i \exp(-i\vec{q}\vec{r}_i).$$

(3.14)

The particle mass-probability current distribution $\vec{J}$ in this "lattice" representation will take the form

$$\vec{J}(\vec{r}) = n(\vec{r})\vec{v} = \frac{1}{2} \sum_i \left\{ \frac{\vec{p}_i}{m} \delta(\vec{r} - \vec{r}_i) + \delta(\vec{r} - \vec{r}_i) \frac{\vec{p}_i}{m} \right\} =$$

$$\frac{1}{2} \sum_i \left\{ \frac{\vec{p}_i}{m} \exp(-i\vec{q}\vec{r}_i) + \exp(-i\vec{q}\vec{r}_i) \frac{\vec{p}_i}{m} \right\},$$

$$[\vec{r}_i, \vec{p}_k] = i\hbar \delta_{ik}.$$  

Here $\vec{v}$ is the velocity operator. The direct calculation shows that

$$[n(\vec{q}), H] = \frac{1}{2} \sum_i \left\{ \frac{\vec{q}\vec{p}_i}{m} \exp(-i\vec{q}\vec{r}_i) + \exp(-i\vec{q}\vec{r}_i) \frac{\vec{q}\vec{p}_i}{m} \right\} = \vec{q}\vec{J}(\vec{q}).$$

(3.16)

Thus the equation of motion for the particle density operator becomes

$$\frac{dn(\vec{q})}{dt} = \frac{i}{\hbar} [H, n(\vec{q})] = -\frac{i}{\hbar} \vec{q}\vec{J}(\vec{q}),$$

(3.17)

or in another form

$$\frac{dn(\vec{r})}{dt} = \text{div}\vec{J}(\vec{r}),$$

(3.18)

which is the continuity equation considered above. Note, that

$$[n(\vec{q}), H_1]_- = [n(\vec{q}), H_2]_- = 0.$$
These relations hold in general for any periodic potential and interaction potential of the electrons which depend only on the coordinates of the electrons.

It is easy to check the validity of the following relation

\[
\left[ n(\vec{q}), H \right], n^\dagger(\vec{q}) \right] = \frac{N q^2}{m}.
\]  

(3.19)

This formula is the known f-sum rule\textsuperscript{[82]} which is a consequence from the continuity equation (for a more general point of view see Ref.\textsuperscript{[83]}).

Now consider the second-quantized Hamiltonian \textsuperscript{[3.2]}. The particle density operator has the form\textsuperscript{[77,84,85]}

\[
n(\vec{r}) = e\Psi^\dagger(\vec{r})\Psi(\vec{r}), \quad n(\vec{q}) = \int d^3r \exp(-i\vec{q}\vec{r})n(\vec{r}).
\]

(3.20)

Then we define

\[
\vec{j}(\vec{r}) = \frac{eh}{2mi}(\Psi^\dagger\nabla\Psi - \Psi\nabla\Psi^\dagger).
\]

(3.21)

Here \(\vec{j}\) is the probability current density, i.e. the probability flow per unit time per unit area perpendicular to \(\vec{j}\). The continuity equation will persist for this case too. Let us consider the equation motion

\[
\frac{dn(\vec{r})}{dt} = -\frac{i}{\hbar}[n(\vec{r}), H_1] - \frac{i}{\hbar}[n(\vec{r}), H_2] = \frac{eh}{2mi}(\Psi^\dagger(\vec{r})\nabla^2\Psi(\vec{r}) - \nabla^2\Psi^\dagger(\vec{r})\Psi(\vec{r})).
\]

(3.22)

Note, that \([n(\vec{r}), H_2] \equiv 0\).

We find

\[
\frac{dn(\vec{r})}{dt} = -\nabla\vec{j}(\vec{r}).
\]

(3.23)

Thus the continuity equation have the same form in both the "particle" and "field" versions.

4 Tight-Binding and Modified Tight-Binding Approximation

Electrons and phonons are the basic elementary excitations of a metallic solid. Their mutual interactions manifest themselves in such observations as the temperature dependent resistivity and low-temperature superconductivity. In the quasiparticle picture, at the basis of this interaction is the individual electron-phonon scattering event, in which an electron is deflected in the dynamically distorted lattice. We consider here the scheme which is called the modified tight-binding interaction (MTBA). But firstly, we remind shortly the essence of the tight-binding approximation. The main purpose in using the tight-binding method is to simplify the theory sufficiently to make workable. The tight-binding approximation considers solid as a giant molecule.

4.1 Tight-binding approximation

The main problem of the electron theory of solids is to calculate the energy level spectrum of electrons moving in an ion lattice.\textsuperscript{[52]} The tight binding method for energy band calculations has generally been regarded as suitable primarily for obtaining a simple first approximation to a complex band structure. It was shown that the method should also be quite powerful in quantitative calculations from first principles for a wide variety of materials. An approximate treatment requires to obtain energy levels and electron wave functions for some suitable chosen one-particle potential (or pseudopotential), which is usually local. The standard molecular orbital
theories of band structure are founded on an independent particle model. As atoms are brought together to form a crystal lattice the sharp atomic levels broaden into bands. Provided there is no overlap between the bands, one expects to describe the crystal state by a Bloch function of the type,

\[ \psi_{\vec{k}}(\vec{r}) = \sum_n e^{i\vec{k}\cdot\vec{R}_n} \phi(\vec{r} - \vec{R}_n), \]  

(4.1)

where \( \phi(\vec{r}) \) is a free atom single electron wave function, for example such as \( 1s \) and \( \vec{R}_n \) is the position of the atom in a rigid lattice. If the bands overlap or approach each other one should use instead of \( \phi(\vec{r}) \) a combination of the wave functions corresponding to the levels in question, e.g. \( (a\phi(1s) + b\phi(2p)) \), etc. In the other words, this approach, first introduced to crystal calculation by F.Bloch, expresses the eigenstates of an electron in a perfect crystal in a linear combination of atomic orbitals and termed LCAO method.\(^{52,91–94}\)

Atomic orbitals are not the most suitable basis set due to the nonorthogonality problem. It was shown by many authors\(^{52,95–97}\) that the very efficient basis set for the expansion (4.1) is the atomic-like Wannier functions \( \{w(\vec{r} - \vec{R}_n)\}\).\(^{52,95–97}\) These are the Fourier transforms of the extended Bloch functions and are defined as

\[ w(\vec{r} - \vec{R}_n) = N^{-1/2} \sum_{\vec{k}} e^{-i\vec{k}\cdot\vec{R}_n} \psi_{\vec{k}}(\vec{r}). \]  

(4.2)

Wannier functions \( w(\vec{r} - \vec{R}_n) \) form a complete set of mutually orthogonal functions localized around each lattice site \( \vec{R}_n \) within any band or group of bands. They permit one to formulate an effective Hamiltonian for electrons in periodic potentials and span the space of a singly energy band. However, the real computation of Wannier functions in terms of sums over Bloch states is a complicated task.\(^{33,97}\)

To define the Wannier functions more precisely let us consider the eigenfunctions \( \psi_{\vec{k}}(\vec{r}) \) belonging to a particular simple band in a lattice with the one type of atom at a center of inversion. Let it satisfy the following equations with one-electron Hamiltonian \( H \)

\[ H\psi_{\vec{k}}(\vec{r}) = E(\vec{k})\psi_{\vec{k}}(\vec{r}), \quad \psi_{\vec{k}}(\vec{r} + \vec{R}_n) = e^{-i\vec{k}\cdot\vec{R}_n} \psi_{\vec{k}}(\vec{r}), \]  

(4.3)

and the orthonormality relation \( \langle \psi_{\vec{k}} | \psi_{\vec{k}'} \rangle = \delta_{\vec{k}\vec{k}'} \) where the integration is performed over the \( N \) unit cells in the crystal. The property of periodicity together with the property of the orthonormality lead to the orthonormality condition of the Wannier functions

\[ \int d^3r w^*(\vec{r} - \vec{R}_n)w(\vec{r} - \vec{R}_m) = \delta_{nm}. \]  

(4.4)

The set of the Wannier functions is complete, i.e.

\[ \sum_i w^*(\vec{r} - \vec{R}_i)w(\vec{r} - \vec{R}_i) = \delta(\vec{r} - \vec{r}). \]  

(4.5)

Thus it is possible to find the inversion of the Eq.(4.2) which has the form

\[ \psi_{\vec{k}}(\vec{r}) = N^{-1/2} \sum_{\vec{k}} e^{i\vec{k}\cdot\vec{R}_n} w(\vec{r} - \vec{R}_n). \]  

(4.6)

These conditions are not sufficient to define the functions uniquely since the Bloch states \( \psi_{\vec{k}}(\vec{r}) \) are determined only within a multiplicative phase factor \( \varphi(\vec{k}) \) according to

\[ w(\vec{r}) = N^{-1/2} \sum_{\vec{k}} e^{i\varphi(\vec{k})} w_{\vec{k}}(\vec{r}), \]  

(4.7)
where \( \varphi(\vec{k}) \) is any real function of \( \vec{k} \), and \( u_{\vec{k}}(\vec{r}) \) are Bloch functions. The phases \( \varphi(\vec{k}) \) are usually chosen so as to localize \( w(\vec{r}) \) about the origin. The usual choice of phase makes \( \psi_{\vec{k}}(\vec{0}) \) real and positive. This leads to the maximum possible value in \( w(\vec{0}) \) and \( w(\vec{r}) \) decaying exponentially away from \( \vec{r} = 0 \). In addition, function \( \psi_{\vec{k}}(\vec{r}) \) with this choice will satisfy the symmetry properties
\[
\psi_{-\vec{k}}(\vec{r}) = (\psi_{\vec{k}}(\vec{r}))^* = \psi_{\vec{k}}(-\vec{r}).
\]

It follows from the above consideration that the Wannier functions are real and symmetric,
\[
w(\vec{r}) = (w(\vec{r}))^* = w(-\vec{r}).
\]

Analytical, three dimensional Wannier functions have been constructed from Bloch states formed from a lattice Gaussians. Thus, in the condensed matter theory, the Wannier functions play an important role in the theoretical description of transition metals, their compounds and disordered alloys, impurities and imperfections, surfaces, etc.

### 4.2 Interacting electrons on a lattice and the Hubbard model

There are big difficulties in description of the complicated problems of electronic and magnetic properties of a metal with the \( d \) band electrons which are really neither "local" nor "itinerant" in a full sense. A better understanding of the electronic correlation effects in metallic systems can be achieved by the formulation of the suitable flexible model that could be used to analyze major aspects of both the insulating and metallic states of solids in which electronic correlations are important.

The Hamiltonian of the interacting electrons with pair interaction in the second-quantized form is given by Eq. (4.23). Consider this Hamiltonian in the Bloch representation. We have
\[
\Psi_\sigma(\vec{r}) = \sum_k \varphi_{\vec{k}\sigma}(\vec{r})a_{k\sigma}, \quad \Psi_\sigma^\dagger(\vec{r}) = \sum_k \varphi_{\vec{k}\sigma}^*(\vec{r})a_{k\sigma}^\dagger.
\]  

Here \( \varphi_\sigma(\vec{k}) \) is the Bloch function satisfying the equation
\[
H_1(r)\varphi_{k\sigma}(\vec{r}) = E_\sigma(\vec{k})\varphi_{k\sigma}(\vec{r}), \quad E_\sigma(\vec{k}) = E_\sigma(-\vec{k}),
\]
\[
\varphi_{\vec{k}}^-(\vec{r}) = \exp(i\vec{k}\vec{r})u_k(r), \quad u_k(r + l) = u_k(r);
\]
\[
\varphi_{k\sigma}(\vec{r}) = \varphi_{-k\sigma}(-\vec{r}), \quad \varphi_{k\sigma}^*(\vec{r}) = \varphi_{-k\sigma}(\vec{r}).
\]

The functions \( \{\varphi_{k\sigma}(\vec{r})\} \) form a complete orthonormal set of functions
\[
\int d^3 r \varphi_{k\sigma}^*(\vec{r})\varphi_{l\sigma}(\vec{r}) = \delta_{kl},
\]
\[
\sum_k \varphi_{k\sigma}(\vec{r}')\varphi_{k\sigma}^*(\vec{r}) = \delta(r - r').
\]

We find
\[
H = \sum_{mn} \langle m|H_1|n\rangle a_{m\sigma}^\dagger a_{n\sigma} + \frac{1}{2} \sum_{klmn} \langle kl|H_2|mn\rangle a_{k\sigma}^\dagger a_{l\sigma}^\dagger a_{m\sigma} a_{n\sigma} =
\]
\[
\sum_{k\sigma} \langle \varphi_{k\sigma}^*|H_1|\varphi_{k\sigma}\rangle a_{k\sigma}^\dagger a_{k\sigma} +
\]
\[
\frac{1}{2} \sum_{k_1k_2k_3k_4} \sum_{\alpha\beta\mu\nu} \langle \varphi_{k_4\alpha|\nu k_3\beta\mu k_2|\nu k_1\alpha\beta} |H_2|\varphi_{k_2\beta\mu k_1\alpha\beta}\rangle a_{k_4\mu\nu}^\dagger a_{k_3\mu\nu} a_{k_2\beta}\alpha a_{k_1\alpha}\beta.
\]
Since the method of second quantization is based on the choice of suitable complete set of orthogonal normalized wave functions, we take now the set \{w_\lambda(\vec{r} - \vec{R}_n)\} of the Wannier functions. Here \( \lambda \) is the band index. The field operators in the Wannier-function representation are given by

\[
\Psi_\sigma(\vec{r}) = \sum_n w_\lambda(\vec{r} - \vec{R}_n) a_{n\lambda\sigma}, \quad \Psi^\dagger_\sigma(\vec{r}) = \sum_n w^*_\lambda(\vec{r} - \vec{R}_n) a^\dagger_{n\lambda\sigma}.
\]  

(4.12)

Thus we have

\[
a^\dagger_{n\lambda\sigma} = N^{-1/2} \sum_\vec{k} e^{-i\vec{k}\cdot\vec{R}_n} a^\dagger_{\vec{k}\lambda\sigma}, \quad a_{n\lambda\sigma} = N^{-1/2} \sum_\vec{k} e^{i\vec{k}\cdot\vec{R}_n} a_{\vec{k}\lambda\sigma},
\]  

(4.13)

Many of treatment of the correlation effects are effectively restricted to a non-degenerate band. The Wannier functions basis set is the background of the widely used Hubbard model. The Hubbard model\(^{99,100}\) is, in a certain sense, an intermediate model (the narrow-band model) and takes into account the specific features of transition metals and their compounds by assuming that the \( d \) electrons form a band, but are subject to a strong Coulomb repulsion at one lattice site. The single-band Hubbard Hamiltonian is of the form\(^{62,99}\)

\[
H = \sum_{ij\sigma} t_{ij} a^\dagger_{i\sigma} a_{j\sigma} + U/2 \sum_{\sigma} n_{i\sigma} n_{i,-\sigma}.
\]  

(4.14)

Here \( a^\dagger_{i\sigma} \) and \( a_{i\sigma} \) are the second-quantized operators of the creation and annihilation of the electrons in the lattice state \( w(\vec{r} - \vec{R}_i) \) with spin \( \sigma \). The Hamiltonian includes the intra-atomic Coulomb repulsion \( U \) and the one-electron hopping energy \( t_{ij} \). The corresponding parameters of the Hubbard Hamiltonian are given by

\[
t_{ij} = \int d^3r w^*(\vec{r} - \vec{R}_i) H_1(r) w(\vec{r} - \vec{R}_j),
\]  

(4.15)

\[
U = \int \int d^3r d^3r' w^*(\vec{r} - \vec{R}_i) w^*(\vec{r'} - \vec{R}_i) \frac{e^2}{|\vec{r} - \vec{r'}|} w(\vec{r'} - \vec{R}_i) w(\vec{r} - \vec{R}_i).
\]  

(4.16)

The electron correlation forces electrons to localize in the atomic-like orbitals which are modelled here by a complete and orthogonal set of the Wannier wave functions \{w(\vec{r} - \vec{R}_j)\}. On the other hand, the kinetic energy is increased when electrons are delocalized. The band energy of Bloch electrons \( E(k) \) is defined as follows:

\[
t_{ij} = N^{-1} \sum_{\vec{k}} E(k) \exp[i\vec{k}\cdot(\vec{R}_i - \vec{R}_j)],
\]  

(4.17)

where \( N \) is the number of lattice sites. The Pauli exclusion principle which does not allow two electrons of common spin to be at the same site, \( n^2_{i\sigma} = n_{i\sigma} \), plays a crucial role. Note, that the standard derivation of the Hubbard model presumes the rigid ion lattice with the rigidly fixed ion positions. We note that \( s \)-electrons are not explicitly taken into account in our model Hamiltonian. They can be, however, implicitly taken into account by screening effects and effective \( d \)-band occupation.

### 4.3 Current operator for the tight-binding electrons

Let us consider again a many-particle interacting systems on a lattice with the Hamiltonian \( \text{(4.11)} \). At this point, it is important to realize the fundamental difference between many-particle system which is uniform in space and many-particle system on a lattice. For the many-particle systems on a lattice the proper definition of current operator is a subtle problem. It was shown above that
a physically satisfactory definition of the current operator in the quantum many-body theory is given based upon the continuity equation. However, this point should be reconsidered carefully for the lattice fermions which are described by the Wannier functions.

Let us remind once again that the Bloch and Wannier wave functions are related to each other by the unitary transformation of the form

\[ \varphi_k(\vec{r}) = N^{-1/2} \sum_{\vec{R}_n} w(\vec{r} - \vec{R}_n) \exp[i \vec{k} \cdot \vec{R}_n], \]

\[ w(\vec{r} - \vec{R}_n) = N^{-1/2} \sum_{\vec{k}} \varphi_k(\vec{r}) \exp[-i \vec{k} \cdot \vec{R}_n]. \]

The number occupation representation for a single-band case lead to

\[ \Psi_\sigma(\vec{r}) = \sum_n w(\vec{r} - \vec{R}_n) a_{n\sigma}, \quad \Psi_\sigma^\dagger(\vec{r}) = \sum_n w^*(\vec{r} - \vec{R}_n) a_{n\sigma}^\dagger. \]

In this representation the particle density operator and current density take the form

\[ n(\vec{r}) = \sum_{ij} \sum_\sigma w^*(\vec{r} - \vec{R}_i) w(\vec{r} - \vec{R}_j) a_{i\sigma}^\dagger a_{j\sigma}, \]

\[ \vec{j}(\vec{r}) = \frac{e\hbar}{2mi} \sum_{ij} \sum_\sigma [w^*(\vec{r} - \vec{R}_i) \nabla w(\vec{r} - \vec{R}_j) - \nabla w^*(\vec{r} - \vec{R}_i) w(\vec{r} - \vec{R}_j)] a_{i\sigma}^\dagger a_{j\sigma}. \]

The equation of the motion for the particle density operator will consists of two contributions

\[ \frac{dn(\vec{r})}{dt} = -\frac{i}{\hbar} [n(\vec{r}), H_1] - \frac{i}{\hbar} [n(\vec{r}), H_2]. \]

The first contribution is

\[ [n(\vec{r}), H_1] = \sum_{mn} \sum_\sigma F_{nm}(\vec{r})(t_{mi} a_{n\sigma}^\dagger a_{i\sigma} - t_{in} a_{i\sigma}^\dagger a_{n\sigma}). \]

Here the notation was introduced

\[ F_{nm}(\vec{r}) = w^*(\vec{r} - \vec{R}_n) w(\vec{r} - \vec{R}_m). \]

In the Bloch representation for the particle density operator one finds

\[ [n(\vec{k}), H_1] = \sum_{mn} \sum_\sigma F_{nm}(\vec{k})(t_{mi} a_{n\sigma}^\dagger a_{i\sigma} - t_{in} a_{i\sigma}^\dagger a_{n\sigma}), \]

where

\[ F_{nm}(\vec{k}) = \int d^3r \exp[-i \vec{k} \cdot \vec{r}] F_{nm}(\vec{r}) = \int d^3r \exp[-i \vec{k} \cdot \vec{r}] w^*(\vec{r} - \vec{R}_n) w(\vec{r} - \vec{R}_m). \]

For the second contribution \([n(\vec{r}), H_2]\) we find

\[ [n(\vec{r}), H_2] = \frac{1}{2} \sum_{mn} \sum_{fst \sigma \sigma'} F_{nm}(\vec{r}) : \]

\[ \left( \langle mf|H_2|st\rangle a_{n\sigma}^\dagger a_{f\sigma'}^\dagger a_{t\sigma'} a_{s\sigma} - \langle fm|H_2|st\rangle a_{m\sigma}^\dagger a_{f\sigma'}^\dagger a_{t\sigma'} a_{s\sigma} \right. \]

\[ + \left. \langle fs|H_2|tn\rangle a_{f\sigma}^\dagger a_{s\sigma'}^\dagger a_{t\sigma'} a_{n\sigma} - \langle fs|H_2|tn\rangle a_{f\sigma}^\dagger a_{s\sigma'}^\dagger a_{t\sigma'} a_{n\sigma} \right). \]
For the single-band Hubbard Hamiltonian the last equation will take the form

$$[n(\vec{r}), H_2] = U \sum_{mn} \sum_{\sigma} F_{nm}(\vec{r}) a_{n\sigma}^\dagger a_{m\sigma}(n_{m-\sigma} - n_{n-\sigma}).$$

(4.27)

The direct calculations give for the case of electrons on a lattice \( e \) is a charge of an electron

$$\frac{dn(\vec{r})}{dt} =$$

$$\frac{eh}{2mi} \sum_{ij} \sum_{\sigma} [w^*(\vec{r} - \vec{R}_i) \nabla^2 w(\vec{r} - \vec{R}_j) - \nabla^2 w^*(\vec{r} - \vec{R}_i) w(\vec{r} - \vec{R}_j)] a_{i\sigma}^\dagger a_{j\sigma} -$$

$$-ieU \sum_{ij} \sum_{\sigma} F_{ij}(\vec{r}) a_{i\sigma}^\dagger a_{j\sigma} (n_{j-\sigma} - n_{i-\sigma}).$$

(4.28)

Taking into account that

$$\text{div} \vec{j}(\vec{r}) =$$

$$\frac{eh}{2mi} \sum_{ij} \sum_{\sigma} [w^*(\vec{r} - \vec{R}_i) \nabla^2 w(\vec{r} - \vec{R}_j) - \nabla^2 w^*(\vec{r} - \vec{R}_i) w(\vec{r} - \vec{R}_j)] a_{i\sigma}^\dagger a_{j\sigma},$$

we find

$$\frac{dn(\vec{r})}{dt} = -\text{div} \vec{j}(\vec{r}) - ieU \sum_{ij} \sum_{\sigma} F_{ij}(\vec{r}) a_{i\sigma}^\dagger a_{j\sigma} (n_{j-\sigma} - n_{i-\sigma}).$$

(4.30)

This unusual result was analyzed critically by many authors. The proper definition of the current operator for the Hubbard model has been the subject of intensive discussions\(^{101-111}\) To clarify the situation let us consider the ”total position operator” for our system of the electrons on a lattice

$$\vec{R} = \sum_{j=1}^{N} \vec{R}_j.$$ (4.31)

In the ”quantized” picture it has the form

$$\vec{R} = \sum_{j} \int d^3r \Psi^\dagger(\vec{r}) \vec{R}_j \Psi(\vec{r})$$

$$= \sum_{j} \sum_{mn} \sum_{\mu} \int d^3r \vec{R}_j w^*(\vec{r} - \vec{R}_m) w(\vec{r} - \vec{R}_n) a_{m\mu}^\dagger a_{n\mu}$$

$$= \sum_{j} \sum_{m} \sum_{\mu} \vec{R}_j a_{m\mu}^\dagger a_{m\mu},$$

where we took into account the relation

$$\int d^3r w^*(\vec{r} - \vec{R}_m) w(\vec{r} - \vec{R}_n) = \delta_{mn}.$$ (4.33)

We find that

$$[\vec{R}, a_{i\sigma}^\dagger]_- = \sum_{m} \vec{R}_m a_{i\sigma}^\dagger,$$ (4.34)

$$[\vec{R}, a_{i\sigma}]_- = -\sum_{m} \vec{R}_m a_{i\sigma},$$

$$[\vec{R}, a_{i\sigma}^\dagger a_{i\sigma}]_- = 0.$$
Let us consider the local particle density operator \( n_{i\sigma} = a_{i\sigma}^\dagger a_{i\sigma} \).
\[
\frac{dn_{i\sigma}}{dt} = -\frac{i}{\hbar}[n_{i\sigma}, H] = \sum_j t_{ij} (a_{i\sigma}^\dagger a_{j\sigma} - a_{j\sigma}^\dagger a_{i\sigma}).
\] (4.35)

It is clear that the current operator should be defined on the basis of the equation
\[
\vec{j} = e\left(\frac{-i}{\hbar}\right)[\vec{R}, H]_-. 
\] (4.36)

Defining the so-called polarization operator\(^{101, 103, 106, 107}\)
\[
P = e \sum_m \sum_{\sigma} \vec{R}_m n_{m\sigma},
\] (4.37)
we find the current operator in the form
\[
\vec{j} = \dot{P} = e\left(\frac{-i}{\hbar}\right) \sum_{mn} \sum_{\sigma} (\vec{R}_m - \vec{R}_n) t_{mn} a_{m\sigma}^\dagger a_{n\sigma}.
\] (4.38)

This expression of the current operator is a suitable formulae for studying of the transport properties of the systems of correlated electron on a lattice\(^{112, 113}\). The consideration carried out in this section demonstrate explicitly the specific features of the many-particle interacting systems on a lattice.

### 4.4 Electron-lattice interaction in metals

In order to understand quantitatively the electrical, thermal and superconducting properties of metals and their alloys one needs a proper description of an electron-lattice interaction.\(^86\) In the physics of molecules\(^{115}\) the concept of an intermolecular force requires that an effective separation of the nuclear and electronic motion can be made. This separation is achieved in the Born-Oppenheimer approximation.\(^{115, 116}\) Closely related to the validity of the Born-Oppenheimer approximation is the notion of adiabaticity. The adiabatic approximation is applicable if the nuclei is much slower than the electrons. The Born-Oppenheimer approximation consists of separating the nuclear motion and in computing only the electronic wave functions and energies for fixed position of the nuclei. In the mathematical formulation of this approximation, the total wave function is assumed in the form of a product both of whose factors can be computed as solutions of two separate Schrödinger equations. In most applications the separation is valid with sufficient accuracy, and the adiabatic approach is reasonable, especially if the electronic properties of molecules are concerned.

The conventional physical picture of a metal adopts these ideas\(^{86, 87}\) and assumes that the electrons and ions are essentially decoupled from one another with an error which involves the small parameter \(m/M\), the ratio between the masses of the electron and the ion. The qualitative arguments for this statement are the following estimations. The maximum lattice frequency is of the order \(10^{13}\) sec\(^{-1}\) and is quite small compared with a typical atomic frequency. This latter frequency is of order of \(10^{15}\) sec\(^{-1}\). If the electrons are able to respond in times of the order of atomic times then they will effectively be following the motion of the lattice instantaneously at all frequencies of vibration. In other words the electronic motion will be essentially adiabatic. This means that the wave functions of the electrons adjusting instantaneously to the motion of the ions. It is intuitively clear that the electrons would try to follow the motion of the ions in such a way as to keep the system locally electrically neutral. In other words, it is expected that the electrons will try to respond to the motion of the ions in such a way as to screen out the local
charge fluctuations. The construction of an electron-phonon interaction requires the separation of the Hamiltonian describing mutually interacting electrons and ions into terms representing electronic quasiparticles, phonons, and a residual interaction. For the simple metals the interaction between the electrons and the ions can be described within the pseudopotential method or the muffin-tin approximation. These methods could not handled well the d bands in the transition metals. They are too narrow to be approximated as free-electron-like bands but too broad to be described as core ion states. The electron-phonon interaction in solid is usually described by the Fröhlich Hamiltonian. We consider below the main ideas and approximations concerning to the derivation of the explicit form of the electron-phonon interaction operator.

Consider the total Hamiltonian for the electrons with coordinates \( \vec{r}_i \) and the ions with coordinates \( \vec{R}_m \), with the electron cores which can be regarded as tightly bound to the nuclei. The Hamiltonian of the \( N \) ions is

\[
H = -\frac{\hbar^2}{2M} \sum_{m=1}^{N} \nabla^2 \vec{R}_m - \frac{\hbar^2}{2m} \sum_{i=1}^{ZN} \nabla^2 \vec{r}_i + \frac{1}{2} \sum_{i,j=1}^{ZN} \frac{e^2}{|\vec{r}_i - \vec{r}_j|} + \sum_{n>m}^{N} V_i(\vec{R}_m - \vec{R}_n) + \sum_{m=1}^{N} U_{ie}(\vec{r}_i; \vec{R}_m). \tag{4.39}
\]

Each ion is assumed to contribute \( Z \) conduction electrons with coordinates \( \vec{r}_i \) \( (i = 1, \ldots, ZN) \). The first two terms in Eq. (4.39) are the kinetic energies of the electrons and the ions. The third term is the direct electron-electron Coulomb interaction between the conduction electrons. The next two terms are short for the potential energy for direct ion-ion interaction and the potential energy of the \( ZN \) conduction electrons moving in the field from the nuclei and the ion core electrons, when the ions take instantaneous position \( \vec{R}_m \) \( (m = 1, \ldots, N) \). The term \( V_i(\vec{R}_m - \vec{R}_n) \) is the interaction potential of the ions with each other, while \( U_{ie}(\vec{r}_i; \vec{R}_m) \) represents the interaction between an electron at \( \vec{r}_i \) and an ion at \( \vec{R}_m \). Thus the total Hamiltonian of the system can be represented as the sum of an electronic and ionic part.

\[
H = H_e + H_i, \tag{4.40}
\]

where

\[
H_e = -\frac{\hbar^2}{2m} \sum_{i=1}^{ZN} \nabla^2 \vec{r}_i + \frac{1}{2} \sum_{i,j=1}^{ZN} \frac{e^2}{|\vec{r}_i - \vec{r}_j|} + \sum_{m=1}^{N} U_{ie}(\vec{r}_i; \vec{R}_m), \tag{4.41}
\]

and

\[
H_i = -\frac{\hbar^2}{2M} \sum_{m=1}^{N} \nabla^2 \vec{R}_m + \sum_{n>m}^{N} V_i(\vec{R}_m - \vec{R}_n). \tag{4.42}
\]

The Schrödinger equation for the electrons in the presence of fixed ions is

\[
H_e \Psi(\vec{K}, \vec{R}, \vec{r}) = E(\vec{K}, \vec{R}) \Psi(\vec{K}, \vec{R}, \vec{r}), \tag{4.43}
\]

in which \( \vec{K} \) is the total wave vector of the system, \( \vec{R} \) and \( \vec{r} \) denote the set of all electronic and ionic coordinates. It is seen that the energy of the electronic system and the wave function of the electronic state depend on the ionic positions. The total wave function for the entire system of electrons plus ions \( \Phi(\vec{Q}, \vec{R}, \vec{r}) \) can be expanded, in principle, with respect to the \( \Psi \) as basis functions

\[
\Phi(\vec{Q}, \vec{R}, \vec{r}) = \sum_{\vec{K}} L(\vec{Q}, \vec{K}, \vec{R}) \Psi(\vec{K}, \vec{R}, \vec{r}). \tag{4.44}
\]
We start with the approach which uses a fixed set of basis states. Let us suppose that the ions of the crystal lattice vibrate around their equilibrium positions $\vec{R}^0_m$ with a small amplitude, namely $\vec{R}_m = \vec{R}^0_m + \vec{u}_m$, where $\vec{u}_m$ is the deviation from the equilibrium position $\vec{R}^0_m$. Let us consider an idealized system in which the ions are fixed in these positions. Suppose that the energy bands $E_n(\vec{k})$ and wave functions $\psi_n(\vec{k}, \vec{r})$ are known. As a result of the oscillations of the ions, the actual crystal potential differs from that of the rigid lattice. This difference is possible to treat as a perturbation. This is the Bloch formulation of the electron-phonon interaction.

To proceed we must expand the potential energy $V(\vec{r} - \vec{R})$ of an electron at $\vec{r}$ in the field of an ion at $\vec{R}_m$ in the atomic displacement $\vec{u}_m$

$$V(\vec{r} - \vec{R}_m) \simeq V(\vec{r} - \vec{R}^0_m) - \vec{u}_m \nabla V(\vec{r} - \vec{R}^0_m) + \ldots$$  \hspace{1cm} (4.45)

The perturbation potential, including all atoms in the crystal, is

$$\tilde{V} = - \sum_m \vec{u}_m \nabla V(\vec{r} - \vec{R}^0_m).$$  \hspace{1cm} (4.46)

This perturbation will produce transitions between one-electron states with the corresponding matrix element of the form

$$M_{mk,nq} = \int \psi^*_{m}(\vec{k}, \vec{r}) \tilde{V} \psi_n(\vec{q}, \vec{r}) d^3 r.$$  \hspace{1cm} (4.47)

To describe properly the lattice subsystem let us remind that the normal coordinate $Q_{\vec{q},\nu}$ is defined by the relation \cite{52,56}

$$(\vec{R}_m - \vec{R}^0_m) = \vec{u}_m = (\hbar/2N^0 M)^{1/2} \sum_{\vec{q},\nu} Q_{\vec{q},\nu} \vec{e}_\nu(\vec{q}) \exp(i\vec{q}\vec{R}^0_m),$$  \hspace{1cm} (4.48)

where $N$ is the number of unit cells per unit volume and $\vec{e}_\nu(\vec{q})$ is the polarization vector of the phonon. The Hamiltonian of the phonon subsystem in terms of normal coordinates is written as \cite{52,56}

$$H_1 = \sum_{\mu,\vec{q}} \left( \frac{1}{2} P_{\vec{q},\mu}^{\dagger} P_{\vec{q},\mu} + \frac{1}{2} \Omega_{\vec{q},\mu}^2 Q_{\vec{q},\mu}^{\dagger} Q_{\vec{q},\mu} \right),$$  \hspace{1cm} (4.49)

where $\mu$ denote polarization direction and the sum $\sum_{\vec{q},\nu}$ is restricted to the Brillouin zone denoted as $BZ$. It is convenient to express $u_m$ in terms of the second quantized phonon operators

$$\vec{u}_m = (\hbar/2N^0 M)^{1/2} \sum_{\vec{q},\nu} [\omega_{\nu}^{1/2}(\vec{q})]^{-1} \vec{e}_\nu(\vec{q})[\exp(i\vec{q}\vec{R}^0_m) b_{\vec{q},\nu} + \exp(-i\vec{q}\vec{R}^0_m) b_{\vec{q},\nu}^{\dagger}],$$  \hspace{1cm} (4.50)

in which $\nu$ denotes a branch of the phonon spectrum, $\vec{e}_\nu(\vec{q})$ is the eigenvector for a vibrational state of wave vector $\vec{q}$ and branch $\nu$, and $b_{\vec{q},\nu}^{\dagger}(b_{\vec{q},\nu})$ is a phonon creation (annihilation) operator. The matrix element $M_{mk,nq}$ becomes

$$M_{mk,nq} = -(\hbar/2N^0 M)^{1/2} \sum_{\vec{q},\nu} (\vec{e}_\nu(\vec{k} - \vec{q}) A_{mn}(\vec{k}, \vec{q})[\omega_{\nu}(\vec{k} - \vec{q})]^{-1/2}(b_{\vec{q}-\vec{k},\nu} + b_{\vec{q}-\vec{k},\nu}^{\dagger})).$$  \hspace{1cm} (4.51)

Here the quantity $A_{mn}$ is given by

$$A_{mn}(\vec{k}, \vec{q}) = N \int \psi^*_{m}(\vec{k}, \vec{r}) \nabla V(\vec{r}) \psi_n(\vec{q}, \vec{r}) d^3 r.$$  \hspace{1cm} (4.52)

It is well known \cite{52,56} that there is the distinction between normal processes in which vector $(\vec{k} - \vec{q})$ is inside the Brillouin zone and Umklapp processes in which vector $(\vec{k} - \vec{q})$ must be brought back
into the zone by addition of a reciprocal lattice vector $\vec{G}$.

The standard simplification in the theory of metals consists of replacement of the Bloch functions $\psi_n(\vec{q}, \vec{r})$ by the plane waves

$$\psi_n(\vec{q}, \vec{r}) = V^{-1/2} \exp(i\vec{q}\vec{r}),$$

in which $V$ is the volume of the system. With this simplification we get

$$A_{mn}(\vec{k}, \vec{q}) = i(\vec{k} - \vec{q}) V((\vec{k} - \vec{q})),$$

(4.53)

Introducing the field operators $\psi(\vec{r}), \psi(\vec{r})^\dagger$ and the fermion second quantized creation and annihilation operators $a_{nk}^\dagger, a_{nk}$ for an electron of wave vector $\vec{k}$ in band $n$ in the plane wave basis

$$\psi(\vec{r}) = \sum_{\vec{q}n} \psi_n(\vec{q}, \vec{r}) a_{nk}$$

and the set of quantities

$$\Gamma_{mn,\nu}(\vec{k}, \vec{q}) = -\left(\hbar/2M\omega_\nu(\vec{k} - \vec{q})\right)^{1/2} \hat{c}_\nu(\vec{k} - \vec{q}) A_{mn}(\vec{k}, \vec{q}),$$

we can write an interaction Hamiltonian for the electron-phonon system in the form

$$H_{ei} = \mathcal{N}^{1/2} \sum_{nl\nu} \sum_{\vec{k}\vec{q}} \Gamma_{mn,\nu}(\vec{k}, \vec{q})\left(a_{nk}^\dagger a_{lq} b_{\vec{k} - \vec{q},\nu} + a_{nk}^\dagger a_{lq} b_{\vec{q} - \vec{k},\nu})\right).$$

(4.54)

This Hamiltonian describes the processes of phonon absorption or emission by an electron in the lattice, which were first considered by Bloch. Thus the electron-phonon interaction is essentially dynamical and affects the physical properties of metals in a characteristic way.

It is possible to show\(^8\) that in the Bloch momentum representation the Hamiltonian of a system of conduction electrons in metal interacting with phonons will have the form

$$H = H_e + H_i + H_{ei},$$

(4.55)

where

$$H_e = \sum_{\vec{p}} E(\vec{p}) a_{\vec{p}}^\dagger a_{\vec{p}},$$

(4.56)

$$H_i = \frac{1}{2} \sum_{\vec{q},\nu} \omega_\nu(\vec{q})(b_{\vec{q}}^\dagger b_{\vec{q}} + b_{-\vec{q}}^\dagger b_{-\vec{q}}),$$

(4.57)

$$H_{ei} = \sum_{\nu} \sum_{\vec{p}'=\vec{p}+\vec{q}+\vec{G}} \Gamma_{\nu}(\vec{p} - \vec{p}') a_{\vec{p}'}^\dagger a_{\vec{p}}(b_{\vec{q} + \vec{G},\nu} + b_{-\vec{q} - \vec{G},\nu}).$$

(4.58)

The Fröhlich model ignores the Umklapp processes ($\vec{G} \neq 0$) and transverse phonons and takes the unperturbed electron and phonon energies as

$$E(\vec{p}) = \frac{\hbar^2 \vec{p}^2}{2m} - E_F, \quad \omega(\vec{q}) = v_s q, \quad (q < q_m).$$

Here $v_s$ is the sound velocity of the free phonon. The other notation are:

$$|\vec{q}| = q, \quad |\vec{p}| = p, \quad q_m = (6\pi^2 n_i)^{1/3}.$$
Thus we obtain

$$H_{ei} = \sum_{p,q} v(q)a_{p+q}^\dagger a_p(b_{q} + b_{-q}^\dagger),$$

(4.59)

where $v(q)$ is the Fourier component of the interaction potential

$$v(q) = g(\omega(q)/2)^{1/2}, \quad g = [2E_F/3Mn_iv_s^2]^{1/2}. $$

Here $n_i$ is the ionic density. The point we should like to emphasize in the present context is that the derivation of this Hamiltonian is based essentially on the plane wave representation for the electron wave function.

### 4.5 Modified tight-binding approximation

Particular properties of the transition metals, their alloys and compounds follow, to a great extent, from the dominant role of $d$-electrons. The natural approach to description of electron-lattice effects in such type of materials is the modified tight-binding approximation (MTBA). The electron-phonon matrix element in the Bloch picture is taken between electronic states of the undeformed lattice. For transition metals it is not easy task to estimate the electron-lattice interaction matrix element due to the anisotropy and other factors. There is an alternative description, introduced by Fröhlich and which was termed the modified tight-binding approximation (MTBA). In this approach the electrons are moving adiabatically with the ions. Moreover, the coupling of the electron to the displacement of the ion nearest to it, is already included in zero order of approximation. This is the basis of modified tight-binding calculations of the electron-phonon interaction which purports to remove certain difficulties of the conventional Bloch tight-binding approximation for electrons in narrow band. The standard Hubbard Hamiltonian should be rederived in this approach in terms of the new basis wave functions for the vibrating lattice. This was carried out by Barisic, Labbe and Friedel. They derived a model Hamiltonian which is a generalization of the single-band Hubbard model including the lattice vibrations. The hopping integral $t_{ij}$ of the single-band Hubbard model (4.14) is given by

$$t_{ij} = \int d^3r w^*(\vec{r} - \vec{R}_j) \left( \frac{1}{2m} \frac{\hbar^2 p^2}{2} + \sum_l V_{sf}(\vec{r} - \vec{R}_l) \right) w(\vec{r} - \vec{R}_i).$$

Here we assumed that $V_{sf}$ is a short-range, self-consistent potential of the lattice suitable screened by outer electrons. Considering small vibrations of ions we replace in Eq.(4.60) the ion position $\vec{R}_i$ by $(\vec{R}_0^i + \vec{u}_i)$, i.e. its equilibrium position plus displacement. The unperturbed electronic wave functions must be written as a Bloch sum of displaced and suitable (approximately) orthonormalized atomic-like functions

$$\int d^3r w^*(\vec{r} - \vec{R}_j^0 - \vec{u}_j)w(\vec{r} - \vec{R}_i^0 - \vec{u}_i) \approx \delta_{ij}. $$

(4.61)

As it follows from Eq.(4.61), the creation and annihilation operators $a_{k\sigma}^\dagger, a_{k\sigma}$ may be introduced in the deformed lattice so as to take partly into account the adiabatic follow up of the electron upon the vibration of the lattice. The Hubbard Hamiltonian Eq.(4.14) can be rewritten in the form

$$H = t_0 \sum_{i\sigma} h_{i\sigma} + \sum_{i\neq j} t(\vec{R}_j^0 + \vec{u}_j - \vec{R}_i^0 - \vec{u}_i)a_{i\sigma}^\dagger a_{j\sigma} + U/2 \sum_{i\sigma} n_{i\sigma} n_{i-\sigma}. $$

(4.62)

For small displacements $\vec{u}_i$, we may expand $t(\vec{R})$ as

$$t(\vec{R}_j^0 + \vec{u}_j - \vec{R}_i^0 - \vec{u}_i) \approx t(\vec{R}_j^0 - \vec{R}_i^0) + \frac{\partial t(\vec{R})}{\partial \vec{R}}|_{\vec{R}=\vec{R}_j^0-\vec{R}_i^0}(\vec{u}_j - \vec{u}_i) + \ldots $$

(4.63)
Table 3: Slater coefficients

| \( q_0 \) (\( A^{-1} \)) | element | element | element | element | element | element |
|--------------------------|--------|--------|--------|--------|--------|--------|
| 0.93                     | Ti     | V      | Cr     | Mn     | Fe     | Co     |
| 0.91                     | Zr     | Nb     | Mo     | Tc     | Ru     | Rh     |
| 0.87                     | Hf     | Ta     | W      | Re     | Os     | Ir     |

Using the character of the exponential decrease of the Slater and Wannier functions the following approximation may be used:\textsuperscript{125–127}

\[
\frac{\partial t(R)}{\partial R} \simeq -q_0 \frac{R}{|R|} t(R). \tag{4.64}
\]

Here \( q_0 \) is the Slater coefficient\textsuperscript{125} originated in the exponential decrease of the wave functions of \( d \)-electrons; \( q_0^{-1} \) related to the range of the \( d \) function and is of the order of the interatomic distance. The Slater coefficients for various metals are tabulated\textsuperscript{125} The typical values are given in Table 3.

It is of use to rewrite the total model Hamiltonian of transition metal \( H = H_e + H_i + H_{ei} \) in the quasi-momentum representation. We have

\[
H_e = \sum_{k\sigma} E(k)a_{k\sigma}^\dagger a_{k\sigma} + U/2N \sum_{k_1k_2k_3k_4G} a_{k_1\uparrow}^\dagger a_{k_2\uparrow} a_{k_3\uparrow}^\dagger a_{k_4\uparrow} \delta(\vec{k}_1 - \vec{k}_2 + \vec{k}_3 - \vec{k}_4 + \vec{G}). \tag{4.65}
\]

For the tight-binding electrons in crystals we use \( E(k) = 2 \sum_{q} t(a_{\alpha}) \cos(k_\alpha a_{\alpha}) \), where \( t(\vec{a}) \) is the hopping integral between nearest neighbours, and \( a_{\alpha}(\alpha = x, y, z) \) denotes the lattice vectors in a simple lattice with an inversion center.

The electron-phonon interaction is rewritten as

\[
H_{ei} = \sum_{kk_1qG} \sum_{\nu} g_{kk_1}^\nu a_{k\sigma}^\dagger a_{k\sigma}(b_{q\nu}^\dagger + b_{-q\nu}) \delta(\vec{k}_1 - \vec{k} + \vec{q} + \vec{G}), \tag{4.66}
\]

where

\[
g_{kk_1}^\nu = \left( \frac{1}{(N M \omega_\nu(k))} \right)^{1/2} I_{kk_1}^\nu, \tag{4.67}
\]

\[
I_{kk_1}^\nu = 2i q_0 \sum_{\alpha} t(\vec{a}_{\alpha}) \frac{\vec{a}_{\alpha} \vec{e}_{\nu}(\vec{q})}{|\vec{a}_{\alpha}|} \left( \sin(\vec{a}_{\alpha} \vec{k}) - \sin(\vec{a}_{\alpha} \vec{k}_1) \right), \tag{4.68}
\]

where \( N \) is the number of unit cells in the crystal and \( M \) is the ion mass. The \( \vec{e}_{\nu}(q) \) are the polarization vectors of the phonon modes. Operators \( b_{q\nu}^\dagger \) and \( b_{q\nu} \) are the creation and annihilation phonon operators and \( \omega_\nu(k) \) are the acoustical phonon frequencies. Thus we can describe the transition metal by the one-band model which takes into consideration the electron-electron and electron-lattice interaction in the framework of the MTBA. It is possible to rewrite (4.66) in the following form\textsuperscript{126, 127, 129, 130}

\[
H_{ei} = \sum_{\nu} \sum_{kq} V_{\nu}(\vec{k}, \vec{k} + \vec{q}) Q_{\vec{q} \nu} a_{k+q\sigma}^\dagger a_{k\sigma}, \tag{4.69}
\]

where

\[
V_{\nu}(\vec{k}, \vec{k} + \vec{q}) = \frac{2i q_0}{(N M)^{1/2}} \sum_{\alpha} t(\vec{a}_{\alpha}) e_{\nu}(q) \left( \sin \vec{a}_{\alpha} \vec{k} - \sin \vec{a}_{\alpha} \vec{k} - \vec{q} \right). \tag{4.70}
\]
The one-electron hopping $t(\vec{a}_\alpha)$ is the overlap integral between a given site $\vec{R}_m$ and one of the two nearby sites lying on the lattice axis $\vec{a}_\alpha$. For the ion subsystem we have

$$H_i = \frac{1}{2} \sum_{q\nu} \left( P_{q\nu}^+ P_{q\nu} + \omega_{\nu}^2(q) \right) Q_{q\nu}^+ Q_{q\nu} = \sum_{q\nu} \omega_{\nu}(q)(b_{q\nu}^\dagger b_{q\nu} + 1/2),$$

where $P_{q\nu}$ and $Q_{q\nu}$ are the normal coordinates. Thus, as in the Hubbard model, the $d$- and $s(p)$-bands are replaced by one effective band in our model. However, the $s$-electrons give rise to screening effects and are taken into effects by choosing proper value of $U$ and the acoustical phonon frequencies. It was shown by Ashkenazi, Dacorogna and Peter that the MTBA approach for calculating electron-phonon coupling constant based on wave functions moving with the vibrating atoms lead to same physical results as the Bloch approach within the harmonic approximation. For transition metals and narrow band compounds the MTBA approach seems to be yielding more accurate results, especially in predicting anisotropic properties.

5 Charge and Heat Transport

We now tackle the transport problem in a qualitative fashion. This crude picture has many obvious shortcomings. Nevertheless, the qualitative description of conductivity is instructive. Guided by this instruction the results of the more advanced and careful calculations of the transport coefficients will be reviewed below in the next sections.

5.1 Electrical resistivity and Ohm law

Ohm law is one of the equations used in the analysis of electrical circuits. When a steady current flow through a metallic wire, Ohm law tell us that an electric field exists in the circuit, that like the current this field is directed along the uniform wire, and that its magnitude is $J/\sigma$, where $J$ is the current density and $\sigma$ the conductivity of the conducting material. Ohm law states that, in an electrical circuit, the current passing through most materials is directly proportional to the potential difference applied across them. A voltage source, $V$, drives an electric current, $I$, through resistor, $R$, the three quantities obeying Ohm law: $V = IR$. In other terms, this is written often as: $I = V/R$, where $I$ is the current, $V$ is the potential difference, and $R$ is a proportionality constant called the resistance. The potential difference is also known as the voltage drop, and is sometimes denoted by $E$ or $U$ instead of $V$. The SI unit of current is the ampere; that of potential difference is the volt; and that of resistance is the ohm, equal to one volt per ampere. The law is named after the physicist Georg Ohm, who formulated it in 1826. The continuum form of Ohm’s law is often of use

$$\mathbf{J} = \sigma \cdot \mathbf{E},$$

where $\mathbf{J}$ is the current density (current per unit area), $\sigma$ is the conductivity (which can be a tensor in anisotropic materials) and $\mathbf{E}$ is the electric field. The common form $V = I \cdot R$ used in circuit design is the macroscopic, averaged-out version. The continuum form of the equation is only valid in the reference frame of the conducting material.

A conductor may be defined as a material within which there are free charges, that is, charges that are free to move when a force is exerted on them by an electric field. Many conducting materials, mainly the metals, show a linear dependence of $I$ on $V$. The essence of Ohm law is this linear relationship. The important problem is the applicability of Ohm law. The relation $R \cdot I = W$ is the generalized form of Ohm law for the current flowing through the system from terminal $A$ to terminal $B$. Here $I$ is a steady dc current, which is zero if the work $W$ done per unit charge is zero,
while $I \neq 0$ or $W \neq 0$. If the current in not too large, the current $I$ must be simply proportional to $W$. Hence one can write $R \cdot I = W$, where the proportionality constant is called the resistance of the two-terminal system. The basic equations are:

$$\nabla \times E = 4\pi n,$$  \hspace{1cm} (5.2)

Gauss law, and

$$\frac{\partial n}{\partial t} + \nabla \cdot J = 0,$$  \hspace{1cm} (5.3)

charge conservation law. Here $n$ is the number density of charge carriers in the system. Equations (5.2) and (5.3) are fundamental. The Ohm law is not. However, in the absence of nonlocal effects, Eq.(5.1) is still valid. In an electric conductor with finite cross section it must be possible a surface conditions on the current density $J$. Ohm law does not permit this and cannot, therefore, be quite correct. It has to be supplemented by terms describing a viscous flow. Ohm law is a statement of the behavior of many, but not all conducting bodies, and in this sense should be looked upon as describing a special property of certain materials and not a general property of all matter.

### 5.2 Drude-Lorentz model

The phenomenological picture described above requires the microscopic justification. We are concerned in this paper with the transport of electric charge and heat by the electrons in a solid. When our sample is in uniform thermal equilibrium the distribution of electrons over the eigenstates available to them in each region of the sample is described by the Fermi-Dirac distribution function and the electric and heat current densities both vanish everywhere. Non-vanishing macroscopic current densities arise whenever the equilibrium is made non-uniform by varying either the electrochemical potential or the temperature from point to point in the sample. The electron distribution in each region of the crystal is then perturbed because electrons move from filled states to adjacent empty states.

The electrical conductivity of a material is determined by the mobile carriers and is proportional to the number density of charge carriers in the system, denoted by $n$, and their mobility, $\mu$, according to

$$\sigma \simeq ne\mu.$$  \hspace{1cm} (5.4)

Only in metallic systems the number density of charge carriers is large enough to make the electrical conductivity sufficiently large. The precise conditions under which one substance has a large conductivity and another substance has low ones are determined by the microscopic physical properties of the system such as energy band structure, carrier effective mass, carrier mobility, lattice properties, and the presence of impurities and imperfections.

Theoretical considerations of the electric conductivity were started by P. Drude within the classical picture about hundred years ago. He put forward a free electron model that assumes a relaxation of the independent charge carriers due to driving forces (frictional force and the electric field). The current density was written as

$$J = \frac{ne^2}{m} E\tau.$$  \hspace{1cm} (5.5)

Here $\tau$ is the average time between collisions, $E$ is the electric field, $m$ and $e$ are the mass and the charge of the electron. The electric conductivity in the Drude model is given by

$$\sigma = \frac{ne^2\tau}{m}.$$  \hspace{1cm} (5.6)
The time $\tau$ is called the mean lifetime or electron relaxation time. Then the Ohm’s law can be expressed as the linear relation between current density $\vec{J}$ and electric field $\vec{E}$

$$\vec{J} = \sigma \vec{E}. \quad (5.7)$$

The electrical resistivity $R$ of the material is equal to

$$R = \frac{E}{J}. \quad (5.8)$$

The free-electron model of Drude is the limiting case of the total delocalization of the outer atomic electrons in a metal. The former valence electrons became conduction electrons. They move independently through the entire body of the metal; the ion cores are totally ignored. The theory of Drude was refined by Lorentz. Drude-Lorentz theory assumed that the free conduction electrons formed an electron gas and were impeded in their motion through crystal by collisions with the ions of the lattice. In this approach, the number of free electrons $n$ and the collision time $\tau$, related to the mean free path $r_l = 2\tau v$ and the mean velocity $v$, are still adjustable parameters. Contrary to this, in the Bloch model for the electronic structure of a crystal, though each valence electron is treated as an independent particle, it is recognized that the presence of the ion cores and the other valence electrons modifies the motion of that valence electron. In spite of its simplicity, Drude model contains some delicate points. Each electron changes its direction of propagation with an average period of $2\tau$. This change of propagation direction is mainly due to a collision of an electron with an impurity or defect and the interaction of electron with a lattice vibration. In an essence, $\tau$ is the average time of the electron motion to the first collision. Moreover, it is assumed that the electron forgets its history on each collision, etc. To clarify these points let us consider the notion of the electron drift velocity. The electrons which contribute to the conductivity have large velocities, that is large compared to the drift velocity which is due to the electric field, because they are at the top of the Fermi surface and very energetic. The drift velocity of the carriers $v_d$ is intimately connected with the collision time $\tau$

$$v_d = \alpha \tau,$$

where $\alpha$ is a constant acceleration between collision of the charge carriers. In general, the mean drift velocity of a particle over $N$ free path is

$$v_d \sim \frac{1}{2} \alpha \left[ \tau + (\Delta t)^2 / \tau \right].$$

This expression shows that the drift velocity depends not only on the average value $\tau$ but also on the standard deviation $(\Delta t)$ of the distribution of times between collisions. An analysis shows that the times between collisions have an exponential probability distribution. For such a distribution, $\Delta t = \tau$ and one obtains $v_d = \alpha \tau$ and $J = ne^2/mE\tau$. Assuming that the time between collisions always has the same value $\tau$ we find that $(\Delta t) = 0$ and $v_d = \frac{1}{2} \alpha \tau$ and $J = ne^2/2mE\tau$. The equations (5.7) and (5.8) are the most fundamental formulas in the physics of electron conduction. Note, that resistivity is not zero even at absolute zero, but is equal to the so called "residual resistivity". For most typical cases it reasonably to assume that scattering by impurities or defects and scattering by lattice vibrations are independent events. As a result, the relation (5.6) will take place. There is a big variety (and irregularity) of the resistivity values for the elements not speaking on the huge variety of substances and materials.$^{[134][137]}$

In a metal with spherical Fermi surface in the presence of an electric field $\vec{E}$, the Fermi surface would affect a $\Delta \vec{k}$ displacement, $\Delta \vec{k} = \vec{k} - \vec{k}_0$. The simplest approximation is to suppose a rigid displacement of the Fermi sphere with a single relaxation time $\tau$,

$$h \frac{d\vec{k}}{dt} + h \frac{(\vec{k} - \vec{k}_0)}{\tau} = e\vec{E}. \quad (5.9)$$
Thus we will have at equilibrium

$$\Delta \vec{k} = \frac{e\tau}{\hbar} \vec{E}. \quad (5.10)$$

The corresponding current density will take the form

$$\vec{J} = \frac{2}{(2\pi)^3} \int_{\Omega_k} e\vec{v}d\Omega_k = \frac{2}{(2\pi)^3} \int_{S_k} e\vec{v}\Delta \vec{k}\delta \vec{S}_{k0}. \quad (5.11)$$

We get from Eq.(5.7)

$$\sigma = \frac{2}{(2\pi)^3} \frac{e^2\tau}{\hbar} \int_{S_k} \vec{v}d\vec{S}_{k0}. \quad (5.12)$$

Let us consider briefly the frequency dependence of $\sigma$. Consider a gas of noninteracting electrons of number density $n$ and collision time $\tau$. At low frequencies collisions occur so frequently that the charge carriers are moving as if within a viscous medium, whereas at high frequencies the charge carriers behave as if they were free. These two frequency regimes are well-known in the transverse electromagnetic response of metals. The electromagnetic energy given to the electrons is lost in collisions with the lattice, which is the "viscous medium". The relevant frequencies in this case satisfy the condition $\omega\tau \ll 1$. Thus in a phenomenological description one should introduce a conductivity $\sigma$ and viscosity $\eta$ by

$$\sigma = \frac{e^2\tau n}{m}, \quad \eta = \frac{1}{2}mv^2n\tau. \quad (5.13)$$

On the other hand, for $\omega\tau \gg 1$ viscous effects are negligible, and the electrons behave as the nearly free particles. For optical frequencies they can move quickly enough to screen out the applied field. Thus, two different physical mechanisms are suitable in the different regimes defined by $\omega\tau \ll 1$ and $\omega\tau \gg 1$.

In a metal impurity atoms and phonons determine the scattering processes of the conduction electrons. The electrical force on the electrons is $e\vec{E}$. The "viscous" drag force is given by $-m\vec{v}/\tau$. Then one can write the equation

$$m\ddot{\vec{v}} = e\vec{E} - \frac{m\vec{v}}{\tau}. \quad (5.14)$$

For $\vec{E} \sim \exp(-i\omega t)$, the oscillating component of the current is given by

$$\vec{J}(\omega) = ne\vec{v}(\omega) = \sigma(\omega)\vec{E}(\omega), \quad (5.15)$$

where

$$\sigma(\omega) = \frac{\sigma_0}{1 - i\omega\tau}, \quad \sigma_0 = \frac{e^2n\tau}{m}. \quad (5.16)$$

For low frequencies we may approximate Eq.(5.14) as $\vec{v} \sim (e\tau/m)\vec{E}$. For high frequencies we may neglect the collision term, so $\vec{v} \sim (e/m)\vec{E}$. Thus the behavior of the conductivity as a function of frequency can be described on the basis of the formula Eq.(5.16).

Let us remark on a residual resistivity, i.e. the resistivity at absolute zero. Since real crystals always contain impurities and defects the resistivity is not equal zero even at absolute zero. If one assume that the scattering of a wave caused by impurities (or defects) and by lattice vibrations are independent events, then the total probability for scattering will be the sum of the two individual probabilities. The scattering probability is proportional to $1/\tau$, where $\tau$ is the mean lifetime or relaxation time of the electron motion. Denoting by $1/\tau_1$ the scattering probability due to impurities and defects and by $1/\tau_2$ the scattering probability due to lattice vibrations we obtain for total probability the equality

$$\frac{1}{\tau} = 1/\tau_1 + 1/\tau_2; \quad 1/\sigma = 1/\sigma_1 + 1/\sigma_2. \quad (5.17)$$
This relation is called Matthiessen rule. In practice, this relation is not fulfilled well (see Refs. 139, 140). The main reason for the violation of the Matthiessen rule are the interference effects between phonon and impurity contributions to the resistivity. Refs. 139, 140 give a comprehensive review of the subject of deviation from Matthiessen rule and detailed critical evaluation of both theory and experimental data.

5.3 The low- and high-temperature dependence of conductivity

One of the most informative and fundamental properties of a metal is the behavior of its electrical resistivity as a function of temperature. The temperature dependence of the resistivity is a good indicator of important scattering mechanisms for the conduction electrons. It can also suggests in a general way what the solid-state electronic structure is like. There are two limiting cases, namely, the low temperature dependence of the resistivity for the case when \( T \leq \theta_D \), where \( \theta_D \) is effective Debye temperature, and the high temperature dependence of the resistivity, when \( T \geq \theta_D \).

The electrical resistivity of metals is due to two mechanisms, namely, (i) scattering of electrons on impurities (static imperfections in the lattice), and (ii) scattering of electrons by phonons. Simplified treatment assumes that one scattering process is not influenced by the other (Matthiessen rule). The first process is usually temperature independent. For a typical metal the electrical resistivity \( R(T) \), as a function of the absolute temperature \( T \), can be written as

\[
R(T) = R_0 + R_i(T),
\]

where \( R_0 \) is the residual electrical resistivity independent of \( T \), and \( R_i(T) \) is the temperature-dependent intrinsic resistivity. The quantity \( R_0 \) is due to the scattering of electrons from chemical and structural imperfections. The term \( R_i(T) \) is assumed to result from the interaction of electrons with other degrees of freedom of a crystal. In general, for the temperature dependence of the resistivity three scattering mechanisms are essential, (i) electron-phonon scattering, (ii) electron-magnon scattering and (iii) electron-electron scattering. The first one gives \( T^5 \) or \( T^3 \) dependence at low temperatures. The second one, the magnon scattering is essential for the transition metals because some of them show ferromagnetic and antiferromagnetic properties. This mechanism can give different temperature dependence due to the complicated (anisotropic) dispersion of the magnons in various structures. The third mechanism, the electron-electron scattering is responsible for the \( R \sim T^2 \) dependence of resistivity.

Usually, the temperature-dependent electrical resistivity is tried to fit to an expression of the form

\[
R(T) = R_0 + R_i(T) = R_0 + AT^5 + BT^2 + (CT^3) + \ldots
\]

This dependence corresponds to Matthiessen rule, where the different terms are produced by different scattering mechanisms. The early approach for studying of the temperature variation of the conductivity was carried out by Sommerfeld, Bloch and Houston. Houston explained the temperature variation of conductivity applying the wave mechanics and assuming that the wave-lengths of the electrons were in most cases long compared with the interatomic distance. He then solves the Boltzmann equation, using for the collision term an expression taken from the work of Debye and Waller on the thermal scattering of x-rays. He obtained an expression for the conductivity as a function of a mean free path, which can be determined in terms of the scattering of the electrons by the thermal vibrations of the lattice. Houston found a resistance proportional to the temperature at high temperatures and to the square of temperature at low temperatures. The model used by Houston for the electrons in a metal was that of Sommerfeld - an ideal gas in a structureless potential well. Bloch improved this approach by taking the periodic structure of the lattice into account. For the resistance law at low temperatures both Houston and Bloch...
results were incorrect. Houston realized that the various treatments of the mean free path would give different variations of resistance with temperature. In his later work\textsuperscript{111} he also realized that the Debye theory of scattering was inadequate at low temperatures. He applied the Brillouin theory of scattering and arrived at $T^5$ law for the resistivity at low temperatures and $T$ at high temperatures. Later on, it was shown by many authors\textsuperscript{22} that the distribution function obtained in the steady state under the action of an electric field and the phonon collisions does indeed lead to $R \sim T^5$. The calculations of the electron-phonon scattering contribution to the resistivity by Bloch\textsuperscript{122} and Gruneisen\textsuperscript{113} lead to the following expression

$$R(T) \sim \frac{T^5}{\theta^5} \int_0^{\theta/T} dz \frac{z^5}{(e^z - 1)(1 - e^{-z})},$$

(5.20)

which is known as the Bloch-Gruneisen law.

A lot of efforts has been devoted to the theory of transport processes in simple metals\textsuperscript{111,114,115} such as the alkali metals. The Fermi surface of these metals is nearly spherical, so that band-structure effects can be either neglected or treated in some simple approximation. The effect of the electron-electron interaction in these systems is not very substantial. Most of the scattering is due to impurities and phonons. It is expected that the characteristic $T^2$ dependence of electron-electron interaction effects can only be seen at very low temperatures $T$, where phonon scattering contributes a negligible $T^5$ term. In the non-simple metallic conductors, and in transition metals, the Fermi surfaces are usually far from being isotropic. Moreover, it can be viewed as the two-component system\textsuperscript{140} where one carrier is an electron and the other is an inequivalent electron (as in $s - d$ scattering) or a hole. It was shown that anisotropy such as that arising from a nonspherical Fermi surface or from anisotropic scattering can yield a $T^2$ term in the resistivity at low temperatures, due to the deviations from Matthiessen rule. This term disappears at sufficiently high $T$. The electron-electron Umklapp scattering contributes a $T^2$ term even at high $T$. It was conjectured (see Ref.\textsuperscript{144}) that the effective electron-electron interaction due to the exchange of phonons should contribute to the electrical resistivity in exactly the same way as the direct Coulomb interaction, namely, giving rise to a $T^2$ term in the resistivity at low temperatures. The estimations of this contribution show\textsuperscript{145} that it can alter substantially the coefficient of the $T^2$ term in the resistivity of simple and polyvalent metals. The role of electron-electron scattering in transition metals was discussed in Refs.\textsuperscript{146,147} A calculation of the electrical and thermal resistivity of $Nb$ and $Pd$ due to electron-phonon scattering was discussed in Ref.\textsuperscript{122} A detailed investigation\textsuperscript{153} of the temperature dependence of the resistivity of $Nb$ and $Pd$ showed that a simple power law fit cannot reconcile the experimentally observed behavior of the transition metals. Matthiessen rule breaks down and simple Bloch-Gruneisen theory is inadequate to account for the experimental data. In particular, in Ref.\textsuperscript{154} it has been shown that the resistivity of $Pd$ can be expressed by a $T^2$ function where, on the other hand, the temperature dependence of the resistivity of $Nb$ should be represented by a function of $T$ more complicated than the $T^3$. It seems to be plausible that the low-temperature behavior of the resistivity of transition metals may be described by a rational function of $(AT^5 + BT^2)$. This conjecture will be justified in section \ref{sec:results}.

For real metallic systems the precise measurements show a quite complicated picture in which the term $R_s(T)$ will not necessarily be proportional to $T^5$ for every metal (for detailed review see Refs.\textsuperscript{111,115,125}). The purity of the samples and size-effect contributions and other experimental limitations can lead to the deviations from the $T^5$ law. There are a lot of other reasons for such a deviation. First, the electronic structures of various pure metals differ very considerably. For example, the Fermi surface of sodium is nearly close to the spherical one, but those of transition and rare-earth metals are much more complicated, having groups of electrons of very different velocities. The phonon spectra are also different for different metals. It is possible to formulate that the $T^5$ law can be justified for a metal of a spherical Fermi surface and for a Debye phonon
spectrum. Moreover, the additional assumptions are an assumption that the electron and phonon systems are separately in equilibrium so that only one phonon is annihilated or created in an electron-phonon collision, that the Umklapp processes can be neglected, and an assumption of a constant volume at any temperature. Whenever these conditions are not satisfied in principle, deviations from $T^5$ law can be expected. This takes place, for example, in transition metals as a result of the $s-d$ transitions due to the scattering of $s$ electrons by phonons. This process can be approximately described as being proportional to $T^3$ with $\gamma$ somewhere between 5 and 3. The $s-d$ model of electronic transport in transition metals was developed by Mott In this model, the motion of the electrons is assumed to take place in the nearly-free-electron-like $s$-band conduction states. These electrons are then assumed to be scattered into the localized $d$ states. Owing to the large differences in the effective masses of the $s$ and $d$ bands, large resistivity result. In Ref the temperature of the normal-state electrical resistivity of very pure niobium was reported. The measurements were carried out in the temperature range from the superconducting transition ($T_c = 9.25K$) to 300K in zero magnetic field. The resistance-versus-temperature data were analyzed in terms of the possible scattering mechanisms likely to occur in niobium. To fit the data a single-band model was assumed. The best fit can be expressed as

\[ R(T) = (4.98 \pm 0.7) \times 10^{-5} + (0.077 \pm 3.0) \times 10^{-7} T^2 + (3.10 \pm 0.23) \times 10^{-7} [T^3 J_3(\theta_D/T)/7.212] + (1.84 \pm 0.26) \times 10^{-10} [T^5 J_5(\theta_D/T)/124.4], \]

where $J_3$ and $J_5$ are integrals occurring in the Wilson and Bloch theories and the best value for $\theta_D$, the effective Debye temperature, is $(270 \pm 10)K$. Over most of the temperature range below 300 K, the $T^3$ Wilson term dominates. Thus it was concluded that inter-band scattering is quite important in niobium. Because of the large magnitude of inter-band scattering, it was difficult to determine the precise amount of $T^2$ dependence in the resistivity. Measurements of the electrical resistivity of the high purity specimens of niobium were carried out in Refs. It was shown that Mott theory is obeyed at high temperature in niobium. In particular, the resistivity curve reflects the variation of the density of states at the Fermi surface when the temperature is raised, thus demonstrating the predominance of $s-d$ transitions. In addition, it was found impossible to fit a Bloch-Gruneisen or Wilson relation to the experimental curve. Several arguments were presented to indicate that even a rough approximation of the Debye temperature has no physical significance and that it is necessary to take the Umklapp processes into account. Measurements of low-temperature electrical and thermal resistivity of tungsten and vanadium showed the effects of the electron-electron scattering between different branches of the Fermi surface in tungsten and vanadium, thus concluding that electron-electron scattering does contribute measurable to electrical resistivity of these substances at low temperature. In transition metal compounds, e.g. $MnP$ the electron-electron scattering is attributed to be dominant at low temperatures, and furthermore the $3d$ electrons are thought to carry electric current. It is remarkable that the coefficient of the $T^2$ resistivity is very large, about a hundred times those of $Ni$ and $Pd$ in which $s$ electrons coexist with $d$ electrons and electric current is mostly carried by the $s$ electrons. This fact suggests strongly that in $MnP$ $s$ electrons do not exist at the Fermi level and current is carried by the $3d$ electrons. This is consistent with the picture that in transition metal compounds the $s$ electrons are shifted up by the effect of antibonding with the valence electrons due to a larger mixing matrix, compared with the $3d$ electrons, caused by their larger orbital extension. It should be noted that the temperature coefficients of resistance can be positive and negative in different materials. A semiconductor material exhibits the temperature dependence of the resistivity quite different than in metal. A qualitative explanation of this different behavior follows from considering the number of free charge carriers per unit volume, $n$, and their mobility, $\mu$. In metals $n$ is essentially constant, but $\mu$ decreases with increasing temperature, owing to increased
lattice vibrations which lead to a reduction in the mean free path of the charge carriers. This
decrease in mobility also occurs in semiconductors, but the effect is usually masked by a rapid
increase in $n$ as more charge carriers are set free and made available for conduction. Thus, intrinsic
semiconductors exhibit a negative temperature coefficient of resistivity. The situation is different
in the case of extrinsic semiconductors in which ionization of impurities in the crystal lattice is
responsible for the increase in $n$. In these, there may exist a range of temperatures over which
essentially all the impurities are ionized, that is, a range over which $n$ remains approximately
constant. The change in resistivity is then almost entirely due to the change in $\mu$, leading to a
positive temperature coefficient.

It is believed that the electrical resistivity of a solid at high temperatures is primarily due to the
scattering of electrons by phonons and by impurities. It is usually assumed, in accordance with
Matthiessen rule that the effect of these two contributions to the resistance are simply additive. At
high temperature (not lower than Debye temperature) lattice vibrations can be well represented by
the Einstein model. In this case, $1/\tau_2 \sim T$, so that $1/\sigma_2 \sim T$. If the properties and concentration
of the lattice defects are independent of temperature, then $1/\sigma_1$ is also independent of temperature
and we obtain

$$1/\sigma \simeq a + bT,$$

where $a$ and $b$ are constants. However, this additivity is true only if the effect of both impurity
and phonon scattering can be represented by means of single relaxation times whose ratio is in-
dependent of velocity. It was shown that the addition of impurities will always decrease the
conductivity. Investigations of the deviations from Matthiessen rule at high temperatures in
relation to the electron-phonon interaction were carried out in Refs. It was shown, in
particular, that changes in the electron-phonon interaction parameter $\lambda$, due to dilute impurities
were caused predominantly by interference between electron-phonon and electron-impurity scatter-
ing.

The electronic band structures of transition metals are extremely complicated and make calcula-
tions of the electrical resistivity due to structural disorder and phonon scattering very difficult. In
addition the nature of the electron-phonon matrix elements is not well understood. The analysis
of the matrix elements for scattering between states was performed in Ref. It was concluded
that even in those metals where a fairly spherical Fermi surface exists, it is more appropriate to
think of the electrons as tightly bound in character rather than free electron-like. In addition, the
"single site" approximations are not likely to be appropriate for the calculation of the transport
properties of structurally-disordered transition metals.

### 5.4 Conductivity of alloys

The theory of metallic conduction can be applied for explaining the conductivity of alloys. According to the Bloch-Gruneisen theory, the contribution of the electron-phonon interaction to the $dc$ electrical resistivity of a metal at high temperatures is essentially governed by two factors, the absolute square of the electron-phonon coupling constant, and the thermally excited mean square lattice displacement. Since the thermally excited mean lattice displacement is proportional to the number of phonons, the high temperature resistivity $R$ is linearly proportional to the absolute temperature $T$, and the slope $dR/dT$ reflects the magnitude of the electron-phonon coupling constant. However, in many high resistivity metallic alloys, the resistivity variation $dR/dT$ is found to be far smaller than that of the constituent materials. In some cases $dR/dT$ is not even always positive. There are two types of alloy, one of which the atoms of the different metals are distributed at random over the lattice points, another in which the atoms of the components are regularly arranged. Anomalous behavior in electrical resistivity was observed in many amorphous and disordered substances. At low temperatures, the resistivity increases
in $T^2$ instead of the usual $T^5$ dependence. Since $T^2$ dependence is usually observed in alloys which include a large fraction of transition metals, it has been considered to be due to spins. In some metals, $T^2$ dependence might be caused by spins. However, it can be caused by disorder itself. The calculation of transport coefficients in disordered transition metal alloys become a complicated task if the random fluctuations of the potential are too large. It can be shown that strong potential fluctuations force the electrons into localized states. Another anomalous behavior occurs in highly resistive metallic systems which is characterized by small temperature coefficient of the electrical resistivity, or by even negative temperature coefficient.

According to Matthiessen rule, the electrical resistance of a dilute alloy is separable into a temperature-dependent part, which is characteristic of the pure metal, and a residual part due to impurities. The variation with temperature of the impurity resistance was calculated by Taylor. The total resistance is composed of two parts, one due to elastic scattering processes, the other to inelastic ones. At the zero of temperature the resistance is entirely due to elastic scattering, and is smaller by an amount $\gamma_0$ than the resistance that would be found if the impurity atom were infinitely massive. The factor $\gamma_0$ is typically of the order of $10^{-2}$. As the temperature is raised the amount of inelastic scattering increases, while the amount of elastic scattering decreases. However, as this happens the ordinary lattice resistance, which varies as $T^5$, starts to become appreciable. For a highly impure specimen for which the lattice resistance at room temperature, $R_\theta$, is equal to the residual resistance, $R_0$, the total resistance at low temperatures will have the form

$$R(T) \approx 10^{-2}(\frac{T}{\theta})^2 + 500(\frac{T}{\theta})^5 + R_0.$$  \hspace{1cm} (5.23)

The first term arises from incoherent scattering and the second from coherent scattering, according to the usual Bloch-Gruneisen theory. It is possible to see from this expression that $T^2$ term would be hidden by the lattice resistance except at temperatures below $\theta/40$. This represents a resistance change of less than $10^{-5}R_0$, and is not generally really observable.

In disordered metals the Debye-Waller factor in electron scattering by phonons may be an origin for negative temperature coefficient of the resistivity. The residual resistivity may decreases as $T^2$ with increasing temperature because of the influence of the Debye-Waller factor. But resulting resistivity increases as $T^2$ with increasing temperature at low temperatures even if the Debye-Waller factor is taken into account. It is worthy to note that the deviation from Matthiessen rule in electrical resistivity is large in the transition metal alloys and dilute alloys. In certain cases the temperature dependence of the electrical resistivity of transition metal alloys at high temperatures can be connected with change electronic density of states. The electronic density of states for $V-Cr$, $Nb-Mo$ and $Ta-W$ alloys have been calculated in the coherent potential approximation. From these calculated results, temperature dependence of the electrical resistivity $R$ at high temperature have been estimated. It was shown that the concentration variation of the temperature dependence in $R/T$ is strongly dependent on the shape of the density of states near the Fermi level.

Many amorphous metals and disordered alloys exhibit a constant or negative temperature coefficient of the electrical resistivity in contrast to the positive temperature coefficient of the electrical resistivity of normal metals. Any theoretical models of this phenomenon must include both the scattering (or collision) caused by the topological or compositional disorder, and also the modifications to this collision induced by the temperature or by electron-phonon scattering. If one assumes that the contributions to the resistivity from scattering mechanisms other than the electron-phonon interaction are either independent of $T$, like impurity scattering, or are saturated at high $T$, like magnetic scattering, the correlation between the quenched temperature dependence and high resistivity leads one to ask whether the electron-phonon coupling constant is affected by the collisions of the electrons.
The effect of collisions on charge redistributions is the principal contributor to the electron-phonon interaction in metals. It is studied as a mechanism which could explain the observed lack of temperature dependence of the electrical resistivity of many concentrated alloys. The collision time-dependent free electron deformation potential can be derived from a self-consistent linearized Boltzmann equation. The results indicate that the collision effects are not very important for real systems. It can be understood assuming that the charge redistribution produces only a negligible correction to the transverse phonon-electron interaction. In addition, although the charge shift is the dominant contribution to the longitudinal phonon-electron interaction, this deformation potential is not affected by collisions until the root mean square electron diffusion distance in a phonon period is less than the Thomas-Fermi screening length. This longitudinal phonon-electron interaction reduction requires collision times of the order of $10^{-19}\text{sec}$ in typical metals before it is effective. Thus, it is highly probable that it is never important in real metals. Hence, this collision effect does not account for the observed, quenched temperature-dependence of the resistivity of these alloys. However, these circumstances suggest that the validity of the adiabatic approximation, i.e., the Born-Oppenheimer approximation, should be relaxed far beyond the previously suggested criteria. All these factors make the proper microscopic formulation of the theory of the electron-phonon interaction in strongly disordered alloys a very complicated problem. A consistent microscopic theory of the electron-phonon interaction in substitutionally disordered crystalline transition metal alloys was formulated by Wysokinski and Kuzemsky within the MTBA. This approach combines the Barisic, Labbe and Friedel model with the more complex details of the CPA (coherent potential approximation).

The low-temperature resistivity of many disordered paramagnetic materials often shows a $T^{3/2}$ rather than a $T^2$ dependence due to spin-fluctuation-scattering resistivity. The coefficient of the $T^{3/2}$ term often correlates with the magnitude of the residual resistivity as the amount of disorder is varied. A model calculations that exhibits such behavior were carried out in Ref. The absence of disorder the spin-fluctuations drag suppresses the spin-fluctuation $T^2$ term in the resistivity. Disorder produces a finite residual resistivity and also produces a finite spin-fluctuation-scattering rate.

### 5.5 Magnetoresistance and the Hall effect

The Hall effect and the magnetoresistance are the manifestations of the Lorentz force on a subsystem of charge carries in a conductor constrained to move in a given direction and subjected to a transverse magnetic field. Let us consider a confined stream of carriers, each having a charge $e$ and a steady-state velocity $v_x$ due to the applied electric field $E_x$. A magnetic field $H$ in the $z$ direction produces a force $F_y$ which has the following form

$$
\vec{F} = e \left( \vec{E} + \frac{1}{c} \vec{\sigma} \times \vec{H} \right).
$$

The boundary conditions lead to the equalities

$$
F_y = 0 = E_y - \frac{1}{c} v_x H_z.
$$

The transverse field $E_y$ is termed the Hall field $E_y \equiv E^H$ and is given by

$$
E^H = \frac{1}{c} v_x H_z = \frac{J_x H_z}{nec}; \quad J_x = ne v_x,
$$

where $J_x$ is the current density and $n$ is the charge carrier concentration. The Hall field can be related to the current density by means of the Hall coefficient $R_H$

$$
E^H = R_H J_x H_z; \quad R_H = \frac{1}{nec}.
$$
The essence of the Hall effect is that Hall constant is inversely proportional to the charge carrier density \( n \), and that it is negative for electron conduction and positive for hole conduction. A useful notion is the so-called Hall angle which is defined by the relation

\[
\theta = \tan^{-1}(E_y / E_x) .
\]  

(5.28)

Thus the Hall effect may be regarded as the rotation of the electric field vector in the sample as a result of the applied magnetic field. The Hall effect is an effective practical tool for studying the electronic characteristics of solids. The above consideration helps to understand how thermomagnetic effects can arise in the framework of simple free-electron model. The Lorentz force acts as a velocity selector. In other words, due to this force the slow electrons will be deflected less than the more energetic ones. This effect will lead to a temperature gradient in the transverse direction. This temperature difference will result in a transverse potential difference due to the Seebeck coefficient of the material. This phenomenon is called the Nernst-Ettingshausen effect. It should be noted that the simple expression for the Hall coefficient \( R_H \) is the starting point only for the studies of the Hall effect in metals and alloys. It implies \( R_H \) is temperature independent and that \( E^H \) varies linearly with applied field strength. Experimentally the dependence \( R_H = 1/nee \) do not fit well the situation in any solid metal. Thus there are necessity to explain these discrepancies. One way is to consider an effective carrier density \( n^*(n) \) which depends on \( n \), where \( n \) is now the mean density of electrons calculated from the valency. This interrelation is much more complicated for the alloys where \( n^*(n) \) is the function of the concentration of solute too. It was shown that the high-field Hall effect reflects global properties of the Fermi surface such as its connectivity, the volume of occupied phase space, etc. The low-field Hall effect depends instead on microscopic details of the dominant scattering process. A quantum-mechanical theory of transport of charge for an electron gas in a magnetic field which takes account of the quantization of the electron orbits has been given by Argyres. Magnetoresistance is an important galvanomagnetic effect which is observed in a wide range of substances and under a variety of experimental conditions. The transverse magnetoresistance is defined by

\[
\rho_{MR}(H) = \frac{R(H) - R}{R} \equiv \frac{\Delta R}{R},
\]  

(5.29)

where \( R(H) \) is the electrical resistivity measured in the direction perpendicular to the magnetic field \( H \), and \( R \) is the resistivity corresponding to the zero magnetic field. The zero-field resistivity \( R \) is the inverse of the zero-field conductivity and is given approximately by

\[
R \sim \frac{m^* \langle v \rangle}{ne^l},
\]  

(5.30)

according to the simple kinetic theory applied to a single-carrier system. Here \( e \), \( m^* \), \( n \), \( \langle v \rangle \) and \( l \) are respectively, charge, effective mass, density, average speed, and mean free path of the carrier. In this simplified picture the four characteristics, \( e \), \( m^* \), \( n \), and \( \langle v \rangle \), are unlikely to change substantially when a weak magnetic field is applied. The change in the mean free path \( l \) should then approximately determine the behavior of the magnetoresistance \( \Delta R/R \) at low fields.

The magnetoresistance practically of all conducting pure single-crystals has been experimentally found to be positive and a strong argument for this were given on the basis of nonequilibrium statistical mechanics. In some substances, e.g. carbon, \( CdSe \), \( Eu_2 CuSi_3 \), etc., magnetoresistance is negative while in \( CdMnSe \) is positive and much stronger than in \( CdSe \). A qualitative interpretation of the magnetoresistance suggests that those physical processes which make the mean free path larger for greater values of \( H \) should contribute to the negative magnetoresistance. Magnetic scattering leads to negative magnetoresistance, characteristic for ferro- or paramagnetic case, which comes from the suppression of fluctuation of the localized spins by the magnetic field.
A comprehensive derivation of the quantum transport equation for electric and magnetic fields was carried out by Mahan. More detailed discussion of the various aspects of theoretical calculation of the magnetoresistance in concrete substances are given in Refs. 188, 199, 200, 202.

5.6 Thermal conduction in solids

Electric and thermal conductivities are intimately connected since the thermal energy also is mainly transported by the conduction electrons. The thermal conductivity of a variety of substances, metals and nonmetals, depends on temperature region and varies with temperature substantially (see Fig. 3). Despite a rough similarity in the form of the curves for metallic and nonmetallic materials there is a fundamental difference in the mechanism whereby heat is transported in these two types of materials. In metals, heat is conducted by electrons; in nonmetals, it is conducted through coupled vibrations of the atoms. The empirical data show that the better the electrical conduction of a metal, the better its thermal conduction. Let us consider a sample with a temperature gradient $dT/dx$ along the $x$ direction. Suppose that the electron located at each point $x$ has thermal energy $E(T)$ corresponding to the temperature $T$ at the point $x$. It is possible to estimate the net thermal energy carried by each electron as

$$E(T) - E(T + \frac{dT}{dx} \tau v \cos \theta) = -\frac{dE(T)}{dT} \frac{dT}{dx} \tau v \cos \theta.$$  \hspace{1cm} (5.31)

Here we denote by $\theta$ the angle between the propagation direction of an electron and the $x$ direction and by $v$ the average speed of the electron. Then the average distance travelled in the $x$ direction by an electron until it scatters is $\tau v \cos \theta$. The thermal current density $J_q$ can be estimated as

$$J_q = -n \frac{dE(T)}{dT} \frac{dT}{dx} \tau v^2 \langle \cos^2 \theta \rangle,$$  \hspace{1cm} (5.32)

where $n$ is the number of electrons per unit volume. If the propagation direction of the electron is random, then $\langle \cos^2 \theta \rangle = 1/3$ and the thermal current density is given by

$$J_q = -\frac{1}{3} n \frac{dE(T)}{dT} \tau v^2 \frac{dT}{dx}.$$  \hspace{1cm} (5.33)
Here \( ndE(T)/dT \) is the electronic heat capacity \( C_e \) per unit volume. We obtain for the thermal conductivity \( \kappa \) the following expression

\[
\kappa = \frac{1}{3} C_e v^2; \quad Q = -\kappa \frac{dT}{dx}.
\]  

(5.34)

The estimation of \( \kappa \) for a degenerate Fermi distribution can be given by

\[
\kappa = \frac{1}{3} \frac{\pi^2 k_B^2 T}{2 \zeta_0} \frac{6 \zeta_0}{5 m} = \frac{\pi^2 k_B^2 T}{5 m} n,
\]

(5.35)

where \( 1/2mv^2 = 3/5\zeta_0 \) and \( k_B \) is the Boltzmann constant. It is possible to eliminate \( n\tau \) with the aid of equality \( \tau = m\sigma/ne^2 \). Thus we obtain

\[
\frac{\kappa}{\sigma} \approx \frac{\pi^2 k_B^2}{5 e^2} T.
\]

(5.36)

This relation is called by the Wiedemann-Franz law. The more precise calculation gives a more accurate factor value for the quantity \( \pi^2/5 \) as \( \pi^2/3 \). The most essential conclusion to be drawn from the Wiedemann-Franz law is that \( \kappa/\sigma \) is proportional to \( T \) and the proportionality constant is independent of the type of metal. In other words, a metal having high electrical conductivity has a high thermal conductivity at a given temperature. The coefficient \( \kappa/\sigma T \) is called the Lorentz number. At sufficiently high temperatures, where \( \sigma \) is proportional to \( 1/T \), \( \kappa \) is independent of temperature. Qualitatively, the Wiedemann-Franz law is based upon the fact that the heat and electrical transport both involve the free electrons in the metal. The thermal conductivity increases with the average particle velocity since that increases the forward transport of energy. However, the electrical conductivity decreases with particle velocity increases because the collisions divert the electrons from forward transport of charge. This means that the ratio of thermal to electrical conductivity depends upon the average velocity squared, which is proportional to the kinetic temperature.

Thus, there are relationships between the transport coefficients of a metal in a strong magnetic field and a very low temperatures. Examples of such relations are the Wiedemann-Franz law for the heat conductivity \( \kappa \), which we rewrite in a more general form

\[
\kappa = LT\sigma,
\]

(5.37)

and the Mott rule\(^{207}\) for the thermopower \( S \)

\[
S = eLT\sigma^{-1} \frac{d\sigma}{d\mu}.
\]

(5.38)

Here \( T \) is the temperature, \( \mu \) denotes the chemical potential. The Lorentz number \( L = 1/3(\pi k_B/e)^2 \), where \( k_B \) is the Boltzmann constant, is universal for all metals. Useful analysis of the Wiedemann-Franz law and the Mott rule was carried out by Nagaev\(^{208}\).

### 6 Linear Macroscopic Transport Equations

We give here a brief refresher of the standard formulation of the macroscopic transport equations from the most general point of view\(^{209}\). One of the main problems of electron transport theory is the finding of the perturbed electron distribution which determines the magnitudes of the macroscopic current densities. Under the standard conditions it is reasonably to assume that the gradients of the electrochemical potential and the temperature are both very small. The macroscopic current densities are then linearly related to those gradients and the ultimate objective of
### Table 4: Fluxes and Generalized Forces

| Process          | Flux  | Generalized force | Tensor character |
|------------------|-------|-------------------|------------------|
| Electrical conduction | \( \vec{J}_e \) | \( \nabla \phi \)   | vector           |
| Heat conduction  | \( \vec{J}_q \) | \( \nabla (1/T) \) | vector           |
| Diffusion        | Diffusion flux \( \vec{J}_p \) | \( -(1/T) \nabla n \) | vector           |
| Viscous flow     | Pressure tensor \( \mathbf{P} \) | \( -(1/T) \nabla \vec{u} \) | Second-rank tensor |
| Chemical reaction| Reaction rate \( W_r \) | Affinity \( a_r/T \) | Scalar           |

the theory of transport processes in solids (see Table 4). Let \( \eta \) and \( T \) denote respectively the electrochemical potential and temperature of the electrons. We suppose that both the quantities vary from point to point with small gradients \( \nabla \eta \) and \( \nabla T \). Then, at each point in the crystal, electric and heat current densities \( \vec{J}_e \) and \( \vec{J}_q \) will exist which are linearly related to the electromotive force \( \vec{E} = 1/e \nabla \eta \) and \( \nabla T \) by the basic transport equations

\[
\vec{J}_e = L_{11} \vec{E} + L_{12} \nabla T, \tag{6.1}
\]

\[
\vec{J}_q = L_{21} \vec{E} + L_{22} \nabla T. \tag{6.2}
\]

The coefficients \( L_{11}, L_{12}, L_{21} \) and \( L_{22} \) in these equations are the transport coefficients which describe irreversible processes in linear approximation. We note that in a homogeneous isothermal crystal, \( \vec{E} \) is equal to the applied electric field \( \vec{E} \). The basic transport equations in the form (6.1) and (6.2) describe responses \( \vec{J}_e \) and \( \vec{J}_q \) under the influence of \( \vec{E} \) and \( \nabla T \). The coefficient \( L_{11} = \sigma \) is the electrical conductivity. The other three coefficients, \( L_{12}, L_{21} \) and \( L_{22} \) have no generally accepted nomenclature because these quantities are hardly ever measured directly. From the experimental point of view it is usually more convenient to fix \( \vec{J}_e \) and \( \nabla T \) and then measure \( \vec{E} \) and \( \vec{J}_q \). To fit the experimental situation the equations (6.1) and (6.2) must be rewritten in the form

\[
\vec{E} = R \vec{J}_e + S \nabla T, \tag{6.3}
\]

\[
\vec{J}_q = \Pi \vec{J}_e - \kappa \nabla T, \tag{6.4}
\]

where

\[
R = \sigma^{-1}, \tag{6.5}
\]

\[
S = -\sigma^{-1} L_{12}, \tag{6.6}
\]

\[
\Pi = L_{21} \sigma^{-1}, \tag{6.7}
\]

\[
\kappa = L_{21} \sigma^{-1} L_{12} = L_{22}, \tag{6.8}
\]

which are known respectively as the resistivity, thermoelectric power, Peltier coefficient and thermal conductivity. These are the quantities which are measured directly in experiments.

All the coefficients in the above equations are tensors of rank 2 and they depend on the magnetic induction field \( \vec{B} \) applied to the crystal. By considering crystals with full cubic symmetry, when \( \vec{B} = 0 \), one reduces to a minimum the geometrical complications associated with the tensor character of the coefficients. In this case all the transport coefficients must be invariant under all the operations in the point group \( m3m \).

This high degree of symmetry implies that the coefficients must reduce to scalar multiples of the unit tensor and must therefore be replaced by scalars. When
$\vec{B} \neq 0$ the general form of transport tensors is complicated even in cubic crystals. In the case, when an expansion to second order in $\vec{B}$ is sufficient the conductivity tensor takes the form

$$\sigma_{\alpha\beta}(\vec{B}) = \sigma_{\alpha\beta}(0) + \sum_{\mu} \frac{\partial \sigma_{\alpha\beta}(0)}{\partial B_\mu} B_\mu + \frac{1}{2} \sum_{\mu\nu} \frac{\partial^2 \sigma_{\alpha\beta}(0)}{\partial B_\mu B_\nu} B_\mu B_\nu + \ldots$$

(6.9)

Here the $(\alpha\beta)$-th element of $\sigma$ referred to the cubic axes $(0xyz)$. For the case when it is possible to confine ourselves by the proper rotations in $m3m$ only, we obtain that

$$\sigma_{\alpha\beta}(0) = \sigma_0 \delta_{\alpha\beta},$$

(6.10)

where $\sigma_0$ is the scalar conductivity when $\vec{B} = 0$. We also find that

$$\frac{\partial \sigma_{\alpha\beta}(0)}{\partial B_\mu} = \varsigma \epsilon_{\alpha\beta\gamma},$$

$$\frac{\partial^2 \sigma_{\alpha\beta}(0)}{\partial B_\mu B_\nu} = 2\xi \delta_{\alpha\beta} \delta_{\mu\nu} + \eta[\delta_{\alpha\mu} \delta_{\beta\nu} + \delta_{\alpha\nu} \delta_{\beta\mu}] + 2\zeta \delta_{\alpha\beta} \delta_{\alpha\mu} \delta_{\alpha\nu},$$

where $\varsigma, \xi, \eta, \zeta$ are all scalar and $\epsilon_{\alpha\beta\gamma}$ is the three-dimensional alternating symbol. Thus we obtain a relation between $\vec{j}$ and $\vec{E}$ (with $\nabla T = 0$)

$$\vec{j} = \sigma_0 \vec{E} + \varsigma \vec{E} \times \vec{B} + \xi \vec{B}^2 \vec{E} + \eta \vec{B} (\vec{B} \vec{E}) + \zeta \Phi \vec{E},$$

(6.11)

where $\Phi$ is a diagonal tensor with $\Phi_{\alpha\alpha} = B_\alpha^2$ (see Refs.12, 14). The most interesting transport phenomena is the electrical conductivity under homogeneous isothermal conditions. In general, the calculation of the scalar conductivity coefficients $\sigma_0, \varsigma, \xi, \eta, \zeta$ is complicated task. As was mentioned above, these coefficients are not usually measured directly. In practice, one measures the corresponding terms in the expression for $\vec{E}$ in terms of $\vec{j}$ up to terms of second order in $\vec{B}$. To show this clearly, let us iterate the equation (6.11). We then find that

$$\vec{E} = R_0 \vec{j} - R_H \vec{j} \times \vec{B} + R_0 (b \vec{B}^2 \vec{j} + c \vec{B} (\vec{B} \vec{j}) + d \Phi \vec{j}),$$

(6.12)

where

$$R_0 = \sigma_0^{-1}, \quad R_H = \sigma_0^{-2} \varsigma,$$

(6.13)

are respectively the low-field resistivity and Hall constant, and

$$b = -R_0 (\xi + \varsigma \zeta); \quad c = -R_0 (\eta - \varsigma \zeta); \quad d = -\eta \zeta$$

(6.14)

are the magnetoresistance coefficients. These are the quantities which are directly measured.

7 Statistical Mechanics and Transport Coefficients

The central problem of nonequilibrium statistical mechanics is to derive a set of equations which describe irreversible processes from the reversible equations of motion. The consistent calculation of transport coefficients is of particular interest because one can get information on the microscopic structure of the condensed matter. There exist a lot of theoretical methods for calculation of transport coefficients as a rule having a fairly restricted range of validity and applicability. The most extensively developed theory of transport processes is that based on the Boltzmann equation. However, this approach has strong restrictions and can reasonably be applied to a strongly rarefied gas of point particles. For systems in the state of
statistical equilibrium, there is the Gibbs distribution by means of which it is possible to calculate an average value of any dynamical quantity. No such universal distribution has been formulated for irreversible processes. Thus, to proceed to the solution of problems of statistical mechanics of nonequilibrium systems, it is necessary to resort to various approximate methods. During the last decades, a number of schemes have been concerned with a more general and consistent approach to transport theory.\cite{42,43,45,211,217} This field is very active and there are many aspects to the problem.\cite{45,220,221,223,224}

### 7.1 Variational principles for transport coefficients

The variational principles for transport coefficients are the special techniques for bounding transport coefficients, originally developed by Kohler, Sondheimer, Ziman, etc. (see Refs.\cite{2,216}). This approach is equally applicable for both the electronic and thermal transport. It starts from a Boltzmann-like transport equation for the space-and-time-dependent distribution function $f_{\vec{q}}$ or the occupation number $n_q(\vec{r},t)$ of a single quasiparticle state specified by indices $q$ (e.g., wave vector for electrons or wave vector and polarization, for phonons). Then it is necessary to find or fit a functional $F[f_{\vec{q}}, n_q(\vec{r},t)]$ which has a stationary point at the distribution $f_{\vec{q}}, n_q(\vec{r},t)$ satisfying the transport equation, and whose stationary value is the suitable transport coefficient. By evaluating $F$ for a distribution only approximately satisfying the transport equation, one then obtains an upper or lower bound on the transport coefficients. Let us mention briefly the phonon-limited electrical resistivity in metals.\cite{2,49} With the neglect of phonon drag, the electrical resistivity can be written

$$R \leq \frac{1}{2k_B T} \int \int (\Phi_\vec{k} - \Phi_\vec{k}')^2 W(\vec{k},\vec{k}') d\vec{k} d\vec{k}' \left| \int e\vec{v}_\vec{k} \Phi_\vec{k} \left( \frac{\partial f^0}{\partial \epsilon(\vec{k})} \right) d\vec{k} \right|^2.$$  \hspace{1cm} (7.1)

Here $W(\vec{k},\vec{k}')$ is the transition probability from an electron state $\vec{k}$ to a state $\vec{k}'$, $\vec{v}_\vec{k}$ is the electron velocity and $f^0$ is the equilibrium Fermi-Dirac statistical factor. The variational principle\cite{9} tells us that the smallest possible value of the right hand side obtained for any function $\Phi_\vec{k}$ is also the actual resistivity. In general we do not know the form of the function $\Phi_\vec{k}$ that will give the right hand side its minimal value. For an isotropic system the correct choice is $\Phi_\vec{k} = \vec{u}\vec{k}$, where $\vec{u}$ is a unit vector along the direction of the applied field. Because of its simple form, this function is in general used also in calculations for real systems. The resistivity will be overestimated but it can still be a reasonable approximation. This line of reasoning leads to the Ziman formula for the electrical resistivity

$$R \leq \frac{3\pi}{e^2 k_B T S^2 k_F^2} \sum_\nu \int \int \frac{d^2 \vec{k} d^2 \vec{k}'}{v v'} \frac{(\vec{k} - \vec{k}')^2 \omega_\nu(\vec{q}) A^2_\nu(\vec{k},\vec{k}')}{(\exp(h\nu(\vec{q})/k_B T) - 1) (1 - \exp(-h\nu(\vec{q})/k_B T))}.$$  \hspace{1cm} (7.2)

Here $k_F$ is the average magnitude of the Fermi wave vector and $S$ is the free area of the Fermi surface. It was shown in Refs.\cite{224} that the formula for the electrical resistivity should not contains any electron-phonon enhancements in the electron density of states. The electron velocities in Eq.(7.2) are therefore the same as those in Eq.(7.1).

### 7.2 Transport theory and electrical conductivity

Let us summarize the results of the preceding sections. It was shown above that in zero magnetic field, the quantities of main interest are the conductivity $\sigma$ (or the electrical resistivity, $R = 1/\sigma$) and the thermopower $S$. When a magnetic field $\vec{B}$ is applied, the quantity of interest is a magnetoresistance

$$\varrho_{MR} = \frac{R(\vec{B}) - R(\vec{B} = 0)}{R(\vec{B} = 0)}$$  \hspace{1cm} (7.3)
and the Hall coefficient $R_H$. The third of the (generally) independent transport coefficients is the thermal conductivity $\kappa$. The important relation which relates $\kappa$ to $R$ at low and high temperatures is the Wiedemann-Franz law. In simple metals and similar metallic systems, which have well-defined Fermi surface it is possible to interpret all the transport coefficients mentioned above, the conductivity (or resistivity), thermopower, magnetoresistance, and Hall coefficient in terms of the rate of scattering of conduction electrons from initial to final states on the Fermi surface. Useful tool to describe this in an approximate way is the Boltzmann transport equation, which, moreover usually simplified further by introducing a concept of a relaxation time. In the approach of this kind we are interested in low-rank velocity moments of the distribution function such as the current

$$j = e \int \tilde{v} f(\tilde{v}) \, d^3v.$$  \hspace{1cm} (7.4)

In the limit of weak fields one expects to find Ohm law $j = \sigma E$. The validity of such a formulation of the Ohm law was analyzed by Bakshi and Gross. This approach was generalized and developed by many authors. The most popular kind of consideration starts from the linearized Boltzmann equation which can be derived assuming weak scattering processes. For example, for the scattering of electrons by "defect" (substituted atom or vacancy) with the scattering potential $V^d(\vec{r})$ the perturbation theory gives

$$R \sim \frac{3\pi^2 m^2 \Omega_0 N}{4\hbar^3 e^2 k_F^2} \int_0^{2k_F} |\langle k + q | V^d(\vec{r}) | k \rangle|^2 q^3 dq,$$  \hspace{1cm} (7.5)

where $N$ and $\Omega_0$ are the number and volume of unit cells, $k_F$ the magnitude of the Fermi wavevector, the integrand $\langle k + q | V^d(\vec{r}) | k \rangle$ represents the matrix elements of the total scattering potential $V^d(\vec{r})$, and the integration is over the magnitude of the scattering wavevector $q$ defined by $\vec{q} = \vec{k} - \vec{k}'$.

Lax have analyzed in detail the general theory of carriers mobility in solids. Luttinger and Kohn, Greenwood, Chester and Thellung, and Fujiita developed approaches to the calculation of the electrical conductivity on the basis of the generalized quantum kinetic equations. The basic theory of transport for the case of scattering by static impurities has been given in the works of Kohn and Luttinger and Greenwood. In these works the usual Boltzmann transport equation and its generalizations were used to write down the equations for the occupation probability in the case of a weak, uniform and static electric field. It was shown that in the case of static impurities the exclusion principle for the electrons has no effect at all on the scattering term of the transport equation. In the case of scattering by phonons, where the electrons scatter inelastically, the exclusion principle plays a very important role and the transport problem is more involved. On the other hand transport coefficients can be calculated by means of theory of the linear response such as the Kubo formulae for the electrical conductivity. New consideration of the transport processes in solids which involve weak assumptions and easily generalizable methods are of interest because they increase our understanding of the validity of the equations and approximations used. Moreover, it permits one to consider more general situations and apply the equations derived to a variety of physical systems.

8 The Method of Time Correlation Functions

The method of time correlation functions is an attempt to base a linear macroscopic transport equation theory directly on the Liouville equation. In this approach one starts with complete $N$-particle distribution function which contains all the information about the system. In the method of time correlation functions it is assumed that the $N$-particle distribution
function can be written as a local equilibrium \( N \)-particle distribution function plus correction terms. The local equilibrium function depends upon the local macroscopic variables, temperature, density and mean velocity and upon the position and momenta of the \( N \) particles in the system. The corrections to this distribution functions is determined on the basis of the Liouville equation. The main assumption is that at some initial time the system was in local equilibrium (quasi-equilibrium) but at later time is tending towards complete equilibrium. It was shown by many authors (for comprehensive review see Refs.\[42,43,45\]) that the suitable solutions to the Liouville equation can be constructed and an expression for the corrections to local equilibrium in powers of the gradients of the local variables can be found as well. The generalized linear macroscopic transport equations can be derived by retaining the first term in the gradient expansion only. In principle, the expressions obtained in this way should depend upon the dynamics of all \( N \) particles in the system and apply to any system, regardless of its density.

### 8.1 Linear response theory

The linear response theory was anticipated in many works (see Refs.\[42,43,45,237,238\] for details) on the theory of transport phenomena and nonequilibrium statistical mechanics. The important contributions have been made by many authors. By solving the Liouville equation to the first order in the external electric field, Kubo\[239,240\] formulated an expression for the electric conductivity in microscopic terms. He used linear response theory to give exact expressions for transport coefficients in terms of correlation functions for the equilibrium system. To evaluate such correlation functions for any particular system, approximations have to be made.

In this section we shall formulate briefly some general expressions for the conductivity tensor within the linear response theory. Consider a many-particle system with the Hamiltonian of a system denoted by \( H \). This includes everything in the absence of the field; the interaction of the system with the applied electric field is denoted by \( H_{\text{ext}} \). The total Hamiltonian is

\[
H = H + H_{\text{ext}}. \tag{8.1}
\]

The conductivity tensor for an oscillating electric field will be expressed in the form\[239\]

\[
\sigma_{\mu\nu} = \int_{0}^{\beta} \int_{0}^{\infty} \text{Tr} \rho_{0} j_{\nu}(0) j_{\mu}(t + i\hbar \lambda) e^{-i\omega t} dt d\lambda, \tag{8.2}
\]

where \( \rho_{0} \) is the density matrix representing the equilibrium distribution of the system in absence of the electric field

\[
\rho_{0} = e^{-\beta H} / \text{Tr} e^{-\beta H}, \tag{8.3}
\]

\( \beta \) being equal to \( 1/k_{B}T \). Here \( j_{\mu}, j_{\nu} \) are the current operators of the whole system in the \( \mu, \nu \) directions respectively, and \( j_{\mu} \) represents the evolution of the current as determined by the Hamiltonian \( H \)

\[
j_{\mu}(t) = e^{iHt/\hbar} j_{\mu} e^{-iHt/\hbar}. \tag{8.4}
\]

Kubo derived his expression Eq.(8.2) by a simple perturbation calculation. He assumed that at \( t = -\infty \) the system was in the equilibrium represented by \( \rho_{0} \). A sinusoidal electric field was switched on at \( t = -\infty \), which however was assumed to be sufficiently weak. Then he considered the equation of motion of the form

\[
i\hbar \frac{\partial}{\partial t} \rho = [H + H_{\text{ext}}(t), \rho]. \tag{8.5}
\]
The change of $\rho$ to the first order of $H_{\text{ext}}$ is given by

$$\rho - \rho_0 = \frac{1}{i\hbar} \int_{-\infty}^{t} e^{-H't'/\hbar} [H_{\text{ext}}(t'), \rho_0] e^{(H't'/\hbar)} + O(H_{\text{ext}}). \quad (8.6)$$

Therefore the averaged current will be written as

$$\langle j_\mu(t) \rangle = \frac{1}{i\hbar} \int_{-\infty}^{t} \text{Tr}[H_{\text{ext}}(t'), \rho_0] j_\mu(-t') dt', \quad (8.7)$$

where $H_{\text{ext}}(t')$ will be replaced by $-e_d E(t')$, $e_d$ being the total dipole moment of the system. Using the relation

$$[A, e^{-\beta H}] = \hbar ie^{-\beta H} \int_{0}^{\beta} e^{\lambda H} [A, \rho] e^{-\lambda H} d\lambda, \quad (8.8)$$

the expression for the current can be transformed into Eq. (8.7). The conductivity can be also written in terms of the correlation function $\langle j_\nu(0)j_\mu(t) \rangle_0$. The average sign $\langle \ldots \rangle_0$ means the average over the density matrix $\rho_0$.

The correlation of the spontaneous currents may be described by the correlation function

$$\Xi_{\mu\nu}(t) = \langle j_\nu(0)j_\mu(t) \rangle_0 = \langle j_\nu(t)j_\mu(t+\tau) \rangle_0. \quad (8.9)$$

The conductivity can be also written in terms of these correlation functions. For the symmetric ("s") part of the conductivity tensor Kubo\cite{239} derived a relation of the form

$$\text{Re} \sigma_{\mu\nu}^s(\omega) = \frac{1}{\varepsilon_\beta(\omega)} \int_{0}^{\infty} \Xi_{\mu\nu}(t) \cos\omega t dt, \quad (8.10)$$

where $\varepsilon_\beta(\omega)$ is the average energy of an oscillator with the frequency $\omega$ at the temperature $T = 1/k_B \beta$. This equation represents the so-called fluctuation-dissipation theorem, a particular case of which is the Nyquist theorem for the thermal noise in a resistive circuit. The fluctuation-dissipation theorems were established\cite{244, 245} for systems in thermal equilibrium. It relates the conventionally defined noise power spectrum of the dynamical variables of a system to the corresponding admittances which describe the linear response of the system to external perturbations. The linear response theory is very general and effective tool for the calculation of transport coefficients of the systems which are rather close to a thermal equilibrium. Therefore, the two approaches, the linear response theory and the traditional kinetic equation theory share a domain in which they give identical results. A general formulation of the linear response theory was given by Kubo\cite{240, 243} for the case of mechanical disturbances of the system with an external source in terms of an additional Hamiltonian.

A mechanical disturbance is represented by a force $F(t)$ acting on the system which may be given function of time. The interaction energy of the system may then be written as

$$H_{\text{ext}}(t) = -AF(t), \quad (8.11)$$

where $A$ is the quantity conjugate to the force $F$. The deviation of the system from equilibrium is observed through measurements of certain physical quantities. If $\Delta \bar{B}(t)$ is the observed deviation of a physical quantity $B$ at the time $t$, we may assume, if only the force $F$ is weak enough, a linear relationship between $\Delta \bar{B}(t)$ and the force $F(t)$, namely

$$\Delta \bar{B}(t) = \int_{-\infty}^{t} \phi_{BA}(t, t') F(t') dt', \quad (8.12)$$
where the assumption that the system was in equilibrium at $t = -\infty$, when the force had been switched on, was introduced. This assumption was formulated mathematically by the asymptotic condition,

$$F(t) \sim e^{\varepsilon t} \quad \text{as} \quad t \to -\infty \quad (\varepsilon > 0).$$

Eq. (8.13) assumes the causality and linearity. Within this limitation it is quite general. Kubo called the function $\phi_{BA}$ of response function of $B$ to $F$, because it represents the effect of a delta-type disturbance of $F$ at the time $t'$ shown up in the quantity $B$ at a later time $t$. Moreover, as it was claimed by Kubo, the linear relationship (8.12) itself not in fact restricted by the assumption of small deviations from equilibrium. In principle, it should be true even if the system is far from equilibrium as far as only differentials of the forces and responses are considered. For instance, a system may be driven by some time-dependent force and superposed on it a small disturbance may be exerted; the response function then will depend both on $t$ and $t'$ separately. If, however, we confine ourselves only to small deviations from equilibrium, the system is basically stationary and so the response functions depend only on the difference of the time of pulse and measurement, $t$ and $t'$, namely

$$\phi_{BA}(t, t') = \phi_{BA}(t - t').$$

(8.14)

In particular, when the force is periodic in time

$$F(t) = \Re F e^{i\omega t},$$

(8.15)

the response of $B$ will have the form

$$\Delta \bar{B}(t) = \Re \chi_{BA}(\omega) F e^{i\omega t},$$

(8.16)

where $\chi_{BA}(\omega)$ is the admittance

$$\chi_{BA}(\omega) = \int_{-\infty}^{t} \phi_{BA}(t)e^{-i\omega t}dt.$$  

(8.17)

More precisely, the response $\Delta \bar{B}(t)$ to an external periodic force $F(t) = F \cos(\omega t)$ conjugate to a physical quantity $A$ is given by Eq. (8.16), where the admittance $\chi_{BA}(\omega)$ is defined as

$$\chi_{BA}(\omega) = \lim_{\varepsilon \to 0} \int_{0}^{\infty} \phi_{BA}(t)e^{-(i\omega + \varepsilon)t}dt.$$  

(8.18)

The response function $\phi_{BA}(\omega)$ is expressed as

$$\phi_{BA}(t) = i\text{Tr}[A, \rho]B(t) = -i\text{Tr}[A, B(t)]$$

$$= \int_{0}^{\beta} \text{Tr}\rho \dot{A}(-i\lambda)B(t)dt = -\int_{0}^{\beta} \text{Tr}\rho A(-i\lambda)\dot{B}d\lambda,$$

(8.19)

where $\rho$ is the canonical density matrix

$$\rho = \exp(-\beta(H - \Omega)), \quad \exp(-\beta\Omega) = \text{Tr} \exp(-\beta H).$$

(8.20)
In certain problems it is convenient to use the relaxation function defined by

\[
\Phi_{BA}(t) = \lim_{\varepsilon \to +0} \int_t^\infty \phi_{BA}(t') e^{-it'} dt' = i \int_t^\infty \langle [B(t'), A] \rangle dt' = -\int_t^\infty dt' \int_0^\beta d\lambda \langle A(i\lambda) \dot{B}(t') \rangle = \int_0^\beta d\lambda \left( \langle A(-i\lambda) B(t) \rangle - \lim_{t \to \infty} \langle A(-i\lambda) B(t) \rangle \right) = \int_0^\beta d\lambda \langle A(-i\lambda) B(t) \rangle - \beta \lim_{t \to \infty} \langle AB(t) \rangle = \int_0^\beta d\lambda \langle A(-i\lambda) B(t) \rangle - \beta \langle A^0 B^0 \rangle.
\]

It is of use to represent the last term in terms of the matrix elements

\[
\int_0^\beta d\lambda \langle A(-i\lambda) B(t) \rangle - \beta \langle A^0 B^0 \rangle = \left( 1/ \sum_i \exp(-\beta E_i) \right) \sum_{n,m} \langle n|A|m\rangle \langle m|B|n\rangle e^{-it(E_n-E_m)} \frac{e^{-\beta E_n} - e^{-\beta E_m}}{E_m - E_n}.
\]

Here \(|m\rangle\) denotes an eigenstate of the Hamiltonian with an eigenvalue \(E_m\) and \(A^0\) and \(B^0\) are the diagonal parts of \(A\) and \(B\) with respect to \(H\). The response function \(\chi_{BA}(\omega)\) can be rewritten in terms of the relaxation function. We have

\[
\chi_{BA}(\omega) = -\lim_{\varepsilon \to +0} \int_0^\infty \phi_{BA}(t) e^{-(i\omega + \varepsilon)t} dt = \phi_{BA}(0) - i\omega \lim_{\varepsilon \to +0} \int_0^\infty \phi_{BA}(t) e^{-(i\omega + \varepsilon)t} dt = -\lim_{\varepsilon \to +0} \int_0^\infty dt \int_0^\beta d\lambda e^{-(i\omega + \varepsilon)t} \frac{d}{dt} \langle AB(t + i\lambda) \rangle = i \lim_{\varepsilon \to +0} \int_0^\infty dt e^{-(i\omega + \varepsilon)t} \left( \langle AB(t + i\beta) \rangle - \langle AB(t) \rangle \right) = i \lim_{\varepsilon \to +0} \int_0^\infty dt e^{-(i\omega + \varepsilon)t} \langle [B(t), A] \rangle = \sum_{n,m} \frac{A_{mn} B_{nm}}{\omega + \omega_{mn} + i\varepsilon} (e^{-\beta E_n} - e^{-\beta E_m}),
\]

where

\[
A_{mn} = \frac{\langle m|A|n\rangle}{(\sum_n e^{-\beta E_n})^{1/2}}.
\]
In particular, the static response $\chi_{BA}(0)$ is given by

$$\chi_{BA}(0) = \phi_{BA}(0)$$

(8.24)

$$= \int_0^\beta d\lambda \left( \langle AB(i\lambda) \rangle - \lim_{t \to \infty} \langle AB(t + i\lambda) \rangle \right)$$

$$= i \lim_{\varepsilon \to +0} \int_0^\infty e^{-\varepsilon t} dt \left( \langle AB(t + i\beta) \rangle - \langle AB(t) \rangle \right)$$

$$= i \lim_{\varepsilon \to +0} \int_0^\infty e^{-\varepsilon t} dt [\langle B(t), A \rangle].$$

This expression can be compared with the isothermal response defined by

$$\chi_{BA}^T = \int_0^\beta \left( \langle AB(i\lambda) \rangle - \langle A \rangle \langle B \rangle \right) d\lambda.$$ 

(8.25)

The difference of the two response functions is given by

$$\chi_{BA}^T - \chi_{BA}(0) = \lim_{t \to \infty} \int_0^\beta d\lambda \langle AB(t + i\lambda) \rangle - \beta \langle A \rangle \langle B \rangle$$

(8.26)

$$= \beta \left( \lim_{t \to \infty} \langle AB(t) \rangle - \langle A \rangle \langle B \rangle \right).$$

The last expression suggests that it is possible to think that the two response functions are equivalent for the systems which satisfy the condition

$$\lim_{t \to \infty} \langle AB(t) \rangle = \langle A \rangle \langle B \rangle.$$ 

(8.27)

It is possible to speak about these systems in terms of ergodic (or quasi-ergodic) behavior, however, with a certain reservation (for a recent analysis of the ergodic behavior of many-body systems see Refs. [237, 256]).

It may be of use to remind a few useful properties of the relaxation function.

If $A$ and $B$ are both Hermitian, then

$$\Phi_{BA}(t) = \text{real}, \quad \int_0^\infty \Phi_{AA}(t) \geq 0.$$ 

(8.28)

The matrix-element representation of the relaxation function have the form

$$\Phi_{BA}(t) = \sum_{m,n} \left( R_{mn}^1 \cos(\omega_{mn}) - R_{mn}^2 \sin(\omega_{mn}) \right)$$

(8.29)

$$+ i \sum_{m,n} \left( R_{mn}^1 \sin(\omega_{mn}) - R_{mn}^2 \cos(\omega_{mn}) \right),$$

where

$$R_{mn}^1 = \frac{1}{2} (A_{nm} B_{mn} + A_{mn} B_{nm}) R_{mn}^3,$$

$$R_{mn}^2 = \frac{1}{2i} (A_{nm} B_{mn} - A_{mn} B_{nm}) R_{mn}^3,$$

$$A_{mn} = \frac{\langle m | A | n \rangle}{(\sum_n e^{-\beta E_n})^{1/2}},$$

$$R_{mn}^3 = \frac{e^{-\beta E_m} - e^{-\beta E_n}}{\omega_{mn}}, \quad \omega_{mn} = E_m - E_n.$$
This matrix-element representation is very useful and informative. It can be shown that the relaxation function has the property

\[ \text{Im} \Phi_{BA}(t) = 0, \] (8.30)

which follows from the odd symmetry of the matrix-element representation. The time integral of the relaxation function is given by

\[ \int_0^\infty \Phi_{BA}(t)dt = \frac{\pi \beta}{2} \sum_{m,n} \left( A_{nm}B_{mn} + B_{nm}A_{mn} \right) e^{-\beta E_n} \delta(\omega_{mn}). \] (8.31)

In particular, for \( A = B \), we obtain

\[ \int_0^\infty \Phi_{AA}(t)dt = \pi \beta \sum_{m,n} |A_{nm}|^2 e^{-\beta E_n} \delta(\omega_{mn}) \geq 0. \] (8.32)

It can be shown also \(^{246}\) that if \( A \) and \( B \) are both bounded, then we obtain

\[ \int_0^\infty \Phi_{BA}(t)dt = i \int_0^{\infty} dt \int_t^{\infty} \langle [\dot{B}(t'), \dot{A}] \rangle dt' \] (8.33)
\[ = -i \int_0^{\infty} dt \langle [B(t), \dot{A}] \rangle = i \langle [A, B] \rangle. \]

For \( A = B \), we have

\[ \int_0^\infty \Phi_{AA}(t)dt = 0, \quad \int_0^\infty dt \int_0^\beta d\lambda \langle \dot{A}(-i\lambda), \dot{A}(t) \rangle = 0 \] (8.34)

and

\[ \lim_{t \to \infty} \langle \dot{A}\dot{B}(t) \rangle = 0, \] (8.35)

if \( A \) and \( B \) are both bounded.

Application of this analysis may not be limited to admittance functions. For example, if one write a frequency dependent mobility function \( \mu(\omega) \) as

\[ \mu(\omega) = [i\omega + \gamma(\omega)]^{-1}, \] (8.36)

the frequency-dependent friction \( \gamma(\omega) \) is also related to a function \( \phi(t) \), which is in fact the correlation function of a random force. An advanced analysis and generalization of the Kubo linear response theory was carried out in series of papers by Van Vliet and co-authors. Fluctuations and response in nonequilibrium steady state were considered within the nonlinear Langevin equation approach by Ohta and Ohkuma. It was shown that the steady probability current plays an important role for the response and time-correlation relation and violation of the time reversal symmetry.

### 8.2 Green functions in the theory of irreversible processes

Green functions are not only applied to the case of statistical equilibrium. They are a convenient means of studying processes where the deviation from the state of statistical equilibrium is small. The use of the Green functions permits one to evaluate the transport coefficients of these processes. Moreover, the transport coefficients are written in terms of Green functions evaluated for the unperturbed equilibrium state without explicitly having recourse to setting up a transport equation. The linear response theory can be reformulated in terms of double-time temperature-dependent (retarded and advanced) Green functions. We shall give a brief
account of this reformulation and its simplest applications to the theory of irreversible processes.

The retarded two-time thermal Green functions arise naturally within the linear response formalism, as it was shown by Zubarev. To show this we consider the reaction of a quantum-mechanical system with a time-independent Hamiltonian $H$ when an external perturbation

$$H_{\text{ext}}(t) = -AF(t), \quad (8.37)$$

is switched on. The total Hamiltonian is equal to

$$\mathcal{H} = H + H_{\text{ext}}, \quad (8.38)$$

where we assume that there is no external perturbation at $\lim t \to -\infty$

$$H_{\text{ext}}(t) |_{\lim t \to -\infty} = 0. \quad (8.39)$$

The last condition means that

$$\lim_{t \to -\infty} \rho(t) = \rho_0 = e^{-\beta H} / [\text{Tr} e^{-\beta H}], \quad (8.40)$$

where $\rho(t)$ is a statistical operator which satisfies the equation of motion

$$i\hbar \frac{\partial}{\partial t} \rho(t) = [H + H_{\text{ext}}(t), \rho], \quad (8.41)$$

This equation of motion together with the initial condition (8.40) suggests to look for a solution of Eq.(8.41) of the form

$$\rho(t) = \rho + \Delta \rho(t). \quad (8.42)$$

Let us rewrite Eq.(8.42), taking into account that $[H, \rho] = 0$, in the following form

$$i\hbar \frac{\partial}{\partial t} (\rho + \Delta \rho(t)) = i\hbar \frac{\partial}{\partial t} \Delta \rho(t) = [H + H_{\text{ext}}(t), \rho + \Delta \rho(t)] = [H, \Delta \rho(t)] + [H_{\text{ext}}(t), \rho] + [H_{\text{ext}}(t), \Delta \rho(t)]. \quad (8.43)$$

Neglecting terms $H_{\text{ext}}(t) \Delta \rho$, since we have assumed that the system is only little removed from a state of statistical equilibrium, we get then

$$i\hbar \frac{\partial}{\partial t} \Delta \rho(t) = [H, \Delta \rho(t)] + [H_{\text{ext}}(t), \rho], \quad (8.44)$$

where

$$\Delta \rho(t) |_{\lim t \to -\infty} = 0. \quad (8.45)$$

Processes for which we can restrict ourselves in Eq.(8.44) to terms linear in the perturbation are called linear dissipative processes. For a discussion of higher-order terms it is convenient to introduce a transformation

$$\Delta \rho(t) = e^{-iHt/\hbar} \varrho(t) e^{iHt/\hbar}, \quad (8.46)$$

Then we have

$$i\hbar \frac{\partial}{\partial t} \Delta \rho(t) = [H, \Delta \rho(t)] + e^{-iHt/\hbar} \left( i\hbar \frac{\partial}{\partial t} \varrho(t) \right) e^{iHt/\hbar}. \quad (8.47)$$
This equation can be transformed to the following form

\[ i\hbar \frac{\partial}{\partial t} \rho(t) = [e^{iHt/\hbar} H_{\text{ext}}(t)e^{-iHt/\hbar}, \rho] + [e^{iHt/\hbar} H_{\text{ext}}(t)e^{-iHt/\hbar}, \rho(t)], \tag{8.48} \]

where

\[ \rho(t)|_{t \to -\infty} = 0. \tag{8.49} \]

In the equivalent integral form the above equation reads

\[ \rho(t) = \frac{1}{\hbar} \int_{-\infty}^{t} d\lambda [e^{iH\lambda/\hbar} H_{\text{ext}}(\lambda)e^{-iH\lambda/\hbar}, \rho] \tag{8.50} \]

\[ + \frac{1}{\hbar} \int_{-\infty}^{t} d\lambda [e^{iH\lambda/\hbar} H_{\text{ext}}(\lambda)e^{-iH\lambda/\hbar}, \rho(\lambda)]. \]

This integral form is convenient for the iteration procedure which can be written as

\[ \rho(t) = \frac{1}{\hbar} \int_{-\infty}^{t} d\lambda [e^{iH\lambda/\hbar} H_{\text{ext}}(\lambda)e^{-iH\lambda/\hbar}, \rho] \tag{8.51} \]

\[ + \left( \frac{1}{\hbar} \right)^2 \int_{-\infty}^{t} d\lambda \int_{-\infty}^{\lambda} d\lambda' \left[ e^{iH\lambda/\hbar} H_{\text{ext}}(\lambda)e^{-iH\lambda/\hbar}, [e^{iH\lambda'/\hbar} H_{\text{ext}}(\lambda')e^{-iH\lambda'/\hbar}, \rho] \right] + \ldots \]

In the theory of the linear reaction of the system on the external perturbation usually the only first term is retained

\[ \Delta \rho(t) = \frac{1}{\hbar} \int_{-\infty}^{t} d\tau e^{-iH(t-\tau)/\hbar}[H_{\text{ext}}(\tau), \rho]e^{iH(t-\tau)/\hbar}. \tag{8.52} \]

The average value of observable \( A \) is

\[ \langle A \rangle_t = \text{Tr}(A\rho(t)) = \text{Tr}(A\rho_0) + \text{Tr}(A\Delta \rho(t)) = \langle A \rangle + \Delta \langle A \rangle_t. \tag{8.53} \]

From this we find

\[ \Delta \langle A \rangle_t = \frac{1}{\hbar} \int_{-\infty}^{t} d\tau \cdot \text{Tr} \left( e^{iH(t-\tau)/\hbar} A e^{-iH(t-\tau)/\hbar} H_{\text{ext}}(\tau) \rho - H_{\text{ext}}(\tau) e^{iH(t-\tau)/\hbar} A e^{-iH(t-\tau)/\hbar} \rho \right) + \ldots \]

\[ = \frac{1}{\hbar} \int_{-\infty}^{t} d\tau \langle [A(\tau - t), H_{\text{ext}}(\tau)]_-, \rho \rangle + \ldots \tag{8.54} \]

Introducing under the integral the sign function

\[ \theta(t - \tau) = \begin{cases} 1 & \text{if } \tau < t, \\ 0 & \text{if } \tau > t, \end{cases} \tag{8.55} \]

and extending the limit of integration to \(-\infty < \tau < +\infty\) we finally find

\[ \Delta \langle A \rangle_t = \int_{-\infty}^{\infty} d\tau \frac{1}{\hbar} \theta(\tau) \langle [A(\tau), H_{\text{ext}}(t - \tau)]_-, \rho \rangle + \ldots \tag{8.56} \]

Let us consider an adiabatic switching on a periodic perturbation of the form

\[ H_{\text{ext}}(t) = B \exp \frac{1}{\hbar} (E + i\varepsilon)t. \]
The presence in the exponential function of the infinitesimal factor \( \varepsilon > 0, \varepsilon \to 0 \) make for the adiabatic switching of the perturbation. Then we obtain
\[
\Delta \langle A \rangle_t = \exp \frac{1}{i\hbar} (E + i\varepsilon) t \int_{-\infty}^{\infty} d\tau \exp \frac{-1}{i\hbar} (E + i\varepsilon) \tau \frac{1}{i\hbar} \theta(\tau) ([A(\tau), B]_-). \tag{8.57}
\]

It is clear that the last expression can be rewritten as
\[
\Delta \langle A \rangle_t = \exp \frac{1}{i\hbar} (E + i\varepsilon) t \int_{-\infty}^{\infty} d\tau \exp \frac{-1}{i\hbar} (E + i\varepsilon) \tau G^{ret}(A, B; \tau) = \tag{8.58}
\exp \frac{1}{i\hbar} (E + i\varepsilon) t G^{ret}(A, B; E) = \exp \frac{1}{i\hbar} (E + i\varepsilon) t \langle [A|B] \rangle_{E+i\varepsilon}. \]

Here \( E = \hbar \omega \) and \( \langle [A|B] \rangle_{E+i\varepsilon} \) is the Fourier component of the retarded Green function \( \langle [A(t)|B(\tau)] \rangle \).

The change in the average value of an operator when a periodic perturbation is switched on adiabatically can thus be expressed in terms of the Fourier components of the retarded Green functions which connect the perturbation operator and the observed quantity.

In the case of an instantaneous switching on of the interaction
\[
H_{ext}(t) = \begin{cases} 
0 & \text{if } t < t_0, \\
\sum_{\Omega} \exp \left( \Omega t / i\hbar \right) V_{\Omega} & \text{if } t > t_0, 
\end{cases} \tag{8.59}
\]

where \( V_{\Omega} \) is an operator which does not explicitly depend on the time, we get
\[
\Delta \langle A \rangle_t = \sum_{\Omega} \int_{t_0}^{\infty} d\tau \langle [A(t)|V_{\Omega}(\tau)] \rangle \exp \frac{1}{i\hbar} (\Omega + i\varepsilon) \tau, \tag{8.60}
\]
i.e., the reaction of the system can also be expressed in terms of the retarded Green functions.

Now we can define the generalized susceptibility of a system on a perturbation \( H_{ext}(t) \) as
\[
\chi(A, B; E) = \chi(A, zB; E) = \lim_{z \to 0} \frac{1}{z} \Delta \langle A \rangle_t \exp \frac{1}{i\hbar} (E + i\varepsilon) t = \langle [A|B] \rangle_{E+i\varepsilon}. \tag{8.61}
\]

In the time representation the above expression reads
\[
\chi(A, B; E) = \frac{1}{i\hbar} \int_{-\infty}^{\infty} dt \exp \frac{-1}{i\hbar} (E + i\varepsilon) t \theta(t) ([A(t), B]_-). \tag{8.62}
\]

This expression is an alternative form of the fluctuation-dissipation theorem, which show explicitly the connection of the relaxation processes in the system with the dispersion of the physical quantities.

The particular case where the external perturbation is periodic in time and contains only one harmonic frequency \( \omega \) is of interest. Putting in that case \( \Omega = \pm i\hbar \omega \) in Eq.\( 8.60 \), since
\[
H_{ext}(t) = -h_0 \cos \omega t e^{\varepsilon t} B, \tag{8.63}
\]
where \( h_0 \), the amplitude of the periodic force, is a \( c- \)number and where \( B \) is the operator part of the perturbation, we get
\[
\Delta \langle A \rangle_t = -h_0 \exp \left( \frac{1}{i\hbar} \omega t + \varepsilon t \right) \langle [A|B] \rangle_{E=\hbar \omega} \tag{8.64}
- h_0 \exp \left( -\frac{1}{i\hbar} \omega t + \varepsilon t \right) \langle [A|B] \rangle_{E=-\hbar \omega}.
\]
Taking into account that $\langle A \rangle_t$ is a real quantity we can write it as follows
\begin{equation}
\Delta \langle A \rangle_t = \text{Re} \left( \chi(E) \hbar_0 e^{\frac{1}{\hbar} \omega t + i \varepsilon t} \right). \tag{8.65}
\end{equation}

Here $\chi(E)$ is the complex admittance, equal to
\begin{equation}
\chi(E) = -2\pi \langle \langle A | B \rangle \rangle_{E=\hbar \omega}. \tag{8.66}
\end{equation}

These equations elucidate the physical meaning of the Fourier components of the Green function $\langle \langle A | B \rangle \rangle_{E=\hbar \omega}$ as being the complex admittance that describes the influence of the periodic perturbation on the average value of the quantity $A$.

### 8.3 The electrical conductivity tensor

When a uniform electric field of strength $\vec{E}$ is switched on then the perturbation acting upon the system of charged particles assumes the form $H_{\text{ext}} = -\vec{E} \cdot \vec{d}(t)$, where $\vec{d}(t)$ is the total dipole moment of the system. In this case the average operator $A(t)$ is the current density operator $\vec{j}$ and the function $\chi$ is the complex electrical conductivity tensor denoted by $\sigma_{\alpha\beta}(\omega)$. If the volume of the system is taken to be equal to unity, then we have
\begin{equation}
\frac{d}{dt} \alpha(t) = j_{\alpha}(t). \tag{8.67}
\end{equation}

The Kubo formula (8.65) relates the linear response of a system to its equilibrium correlation functions. Here we consider the connection between the electrical conductivity tensor and Green functions. Let us start with a simplified treatment when there be switched on adiabatically an electrical field $\vec{E}(t)$, uniform in space and changing periodically in time with a frequency $\omega$
\begin{equation}
\vec{E}(t) = \vec{E} \cos \omega t. \tag{8.68}
\end{equation}

The corresponding perturbation operator is equal to
\begin{equation}
H_{\text{ext}}(t) = -e \sum_j (\vec{E} \vec{r}_j) \cos \omega t e^{\varepsilon t}. \tag{8.69}
\end{equation}

Here $e$ is the charge of an electron, and the summation is over all particle coordinates $\vec{r}_j$. Under the influence of the perturbation there arises in the system an electrical current
\begin{equation}
\alpha(t) = \int_{-\infty}^{\infty} d\tau \langle \langle j_{\alpha}(t); H_{\text{ext}}(\tau) \rangle \rangle, \tag{8.70}
\end{equation}

where
\begin{equation}
H_{\text{ext}}(\tau) = H^{\prime}_1(\tau) \cos \omega \tau e^{\varepsilon \tau}, \tag{8.71}
\end{equation}
\begin{equation}
H^{\prime}_1(\tau) = -e \sum_{\alpha} \mathcal{E}_{\alpha} \vec{r}_j^\alpha(\tau), \quad j_{\alpha}(t) = e \sum_{j} \vec{r}_j^\alpha(t). \tag{8.72}
\end{equation}

Here $\alpha(t)$ is the current density operator, if the volume of the system is taken to be unity. The equation (8.70) can be transformed to the following form
\begin{equation}
\alpha(t) = -\text{Re} \left\{ \int_{-\infty}^{\infty} d\tau \langle \langle j_{\alpha}(t); H^{\prime}_1(\tau) \rangle \rangle e^{\frac{1}{i\omega + \varepsilon} \omega t + i \varepsilon t} \right\} + \langle [j_{\alpha}(0), H^{\prime}_1(0)]_{-} \rangle e^{\frac{1}{\omega - i \varepsilon} \omega t + i \varepsilon t} \frac{1}{\omega - i \varepsilon}. \tag{8.73}
\end{equation}
Noting that
\[ \dot{H}_1^1(\tau) = -(\vec{E} \vec{j}(\tau)), \quad [r_i^\alpha, r_j^\beta] = \frac{1}{i\hbar} \delta_{\alpha\beta} \delta_{ij}, \]
we get from this equation
\[ j_\alpha(t) = \text{Re}\{\sigma_{\alpha\beta}(\omega) \mathcal{E}_\beta \exp(i\omega t + \varepsilon t)\}, \]
where
\[ \sigma_{\alpha\beta}(\omega) = -\frac{ie^2 n}{m\omega} \delta_{\alpha\beta} + \int_{-\infty}^{\infty} d\tau \langle\langle j_\alpha(0); j_\beta(\tau)\rangle\rangle \frac{\exp(i\omega t + \varepsilon t)}{i\omega + \varepsilon} \]
is the conductivity tensor, and \( n \) the number of electrons per unit volume. The first term in Eq. (8.73) corresponds to the electrical conductivity of a system of free charges and is not connected with the inter-particle interaction. As \( \omega \to \infty \) the second term decreases more strongly than the first one
\[ \lim_{\omega \to \infty} \text{Im} \omega \sigma_{\alpha\beta}(\omega) = -\frac{e^2 n}{m\omega} \delta_{\alpha\beta}, \]
and the system behaves as a collection of free charges.

Let us discuss the derivation of the conductivity tensor in general form. Consider a system of charged particles in electrical field \( \vec{E} \), which is directed along the axis \( \beta \) (\( \beta = x, y, z \)). The corresponding electrostatic potential \( \varphi^{(\beta)} \)
has the following form
\[ \varphi^{(\beta)}(\vec{r}, t) = \sum_{\Omega} \frac{1}{\Omega} \sum_{q} \varphi^{(\beta)}(q, \Omega) \exp \left( (\Omega t + i\varepsilon t)/i\hbar - i\vec{q}\vec{r} \right). \]
In the momentum representation the above expression reads
\[ \mathcal{E}_\alpha^{(\beta)}(q, \Omega) = \frac{i}{\hbar} q_\alpha \varphi^{(\beta)}(q, \Omega) = \delta_{\alpha\beta} \mathcal{E}_\beta(q, \Omega). \]
Consider the case when the perturbation \( H^{\text{ext}} \) has the form
\[ H^{\text{ext}} = \frac{1}{\Omega} e\varphi^{(\beta)}(q, \Omega) \eta_q^\dagger \exp \left( (\Omega t + i\varepsilon t)/i\hbar \right). \]
Here
\[ \eta_q = \sum_p a_{p+q}^\dagger a_p, \quad \eta_{-q}^\dagger = \eta_{-q}. \]
is the particle density operator.

A reaction of the system on the perturbation is given by
\[ I^{(\beta)}_\alpha(q, \Omega) = \Delta \langle e j_\alpha(q) \rangle = \frac{1}{V} \langle \langle e j_\alpha(q) | e \varphi^{(\beta)}(q, \Omega) \eta_q^\dagger \rangle \rangle_{\Omega+i\varepsilon} \]
\[ = e^2 \varphi^{(\beta)}(q, \Omega) \frac{1}{V} \int_{-\infty}^{\infty} d\omega J(\eta_q^\dagger, j_\alpha(q); \omega) \frac{e^{\omega/\Theta} - 1}{\Omega - \omega + i\varepsilon}, \]
where \( I^{(\beta)}_\alpha(q, \Omega) \) denotes the component of the density of the current in the \( \alpha \)-direction when external electric field directed along the \( \beta \)-axis.
8.4 Linear response theory: pro et contra

It was shown in the previous sections that the formulation of the linear response theory can be generalized so as to be applied to a rather wide class of the problems. It is worth to note that the "exact" linear expression for electrical conductivity for an arbitrary system was derived originally by Kubo\textsuperscript{240} in a slightly different form than Eq. (8.75)

\[\sigma_{\mu\nu} = \lim_{\varepsilon \to 0} \frac{1}{\varepsilon} \left( \phi_{\mu\nu}(0) + \int_{-\infty}^{\infty} dt e^{-\varepsilon t} \dot{\phi}_{\mu\nu}(t) \right),\] (8.80)

where

\[\phi_{\mu\nu}(t) = \frac{1}{i\hbar} \text{Tr} \left( n, \sum_i e_i x_{i\nu} \right) \sum_i e_i \dot{x}_{i\nu}(t)\] (8.81)

is the current response in the $\mu$--direction when a pulse of electric field $E(t)$ is applied in the $\nu$--direction at $t = 0$; $e_i$ is the charge of the $i$th particle with position vector $\vec{r}_i$ and $n$ is a density operator. In the one-electron approximation Eq. (8.80) reduces to (cf. Ref. \textsuperscript{233})

\[\sigma_{ij} = 2\pi e^2 h \sum_{nm} \langle m|v_i|n\rangle \langle n|v_j|m\rangle \left( \frac{\partial f}{\partial E} \right)_n.\] (8.82)

Here $v_i$ is the velocity operator and $f$ is the Fermi function.

To clarify the general consideration of the above sections it is of interest to consider here a simplified derivation of this formula, using only the lowest order of time-dependent perturbation theory.\textsuperscript{262} This approach is rooted in the method of derivation of Callen and Welton\textsuperscript{263} but takes explicitly into account degeneracy of the states. The linear response theory formulated by Kubo\textsuperscript{240} was motivated in part\textsuperscript{237, 238} by the prior work of Callen and Welton\textsuperscript{263} who proposed a quantum mechanical perturbative calculation with the external forces exerted on a dissipative system. They pointed out a general relationship between the power dissipation induced by the perturbation and the average of a squared fluctuation of the current of the system in thermal equilibrium.

Let us discuss first the role of dissipation. A system may be called to be dissipative if it absorbs energy when subjected to a time-periodic perturbation, and linear if the dissipation (rate of absorption of energy) is quadratic in the perturbation. For a linear system an impedance may be defined and the proportionality constant between the power and the square of the perturbation amplitude is simply related to the impedance. In the case of electrical current in a material one can write down that

\[W = \frac{1}{2} \frac{R}{(\text{Impedance})^2} = V^2,\]

where $W$ is the average power and $V$ is the voltage. If we calculate the power microscopically in some way and find it quadratic in the applied force (voltage), then comparison with this equation will give the conductivity of the substance.

Consider a situation when an electron of charge $e$ is situated a distance $x$ from the end of a resistor of length $L$ and then a voltage $V = V_0 \sin \omega t$ is applied in the $x$ direction\textsuperscript{262} The perturbation term in the Hamiltonian will be of the form

\[H_{\omega}^{\text{ext}} = V_0 \frac{x}{L} \sin \omega t.\] (8.83)

The Hamiltonian of a system in the absence of the field (but including all other interactions) is denoted by $H_0$ with corresponding wave function $\psi_n$ such that

\[H_0 \psi_n = E_n \psi_n.\] (8.84)
The total wave function may be expanded in terms of the $\psi_n$

$$\Psi_n = \sum_n a_n(t)\psi_n,$$  \hspace{1cm} (8.85)

where the coefficients $a_n(t)$ may be approximately determined by first-order perturbation theory. The rate of transition is then given by

$$\frac{dp_n}{dt} = \frac{1}{2} \frac{\pi e^2 V_0^2}{\hbar} \sum_{mn} |\langle m|x|n \rangle|^2 \left[ \delta(E_m - (E_n + \hbar \omega)) + \delta(E_m - (E_n - \hbar \omega)) \right].$$  \hspace{1cm} (8.86)

The first term corresponds to a transition to a state $E_m = E_n + \hbar \omega$ in which energy $\hbar \omega$ is absorbed, whereas the second term corresponds to a transition to a state $E_m = E_n - \hbar \omega$ in which energy is emitted. Hence, the net rate of absorption of energy is given by

$$\frac{dE_n}{dt} = \frac{1}{2} \frac{\pi e^2 V_0^2}{\hbar} \sum_{mn} |\langle m|x|n \rangle|^2 \left[ \delta(E_m - E_n - \hbar \omega) + \delta(E_m - E_n + \hbar \omega) \right],$$  \hspace{1cm} (8.87)

which is quadratic in $V_0$.

This equation gives the absorption rate for a single, isolated electrons, but in a real system we are dealing with an ensemble of these, which we shall represent by the Fermi function. One must therefore find the average absorption by averaging over all initial states $|n\rangle$ and taking the Pauli exclusion principle as well as the two spin directions into account. The result is

$$\langle \frac{dE_n}{dt} \rangle = \pi e^2 V_0^2 \omega \sum_{mn} |\langle m|x|n \rangle|^2 \left[ f(E_n) \delta(E_m - E_n - \hbar \omega)(1 - f(E_m)) - f(E_n) \delta(E_m - E_n + \hbar \omega)(1 - f(E_m)) \right] - \sum_{n'} |\langle n'|x|n + \hbar \omega \rangle|^2 \left[ f(E_{n'}) f(E_n + \hbar \omega) - f(E_{n'} + \hbar \omega) f(E_{n'}) \right],$$

where we have put $n' = n - \hbar \omega$. The above formula can be transformed to the form

$$\langle \frac{dE_n}{dt} \rangle = \pi e^2 V_0^2 \omega \sum_{mn} |\langle m|x|n \rangle|^2 \left[ f(E_n) - f(E_m) \right] \delta(E_m - E_n - \hbar \omega).$$  \hspace{1cm} (8.89)

This expression may be simplified by introducing the matrix element of the velocity operator

$$\frac{dx}{dt} = \frac{i}{\hbar} [H, x].$$  \hspace{1cm} (8.90)

In principle, on the right-hand side of Eq. (8.90) the total Hamiltonian

$$\mathcal{H} = H + H_{\text{ext}}$$  \hspace{1cm} (8.91)

should be written. In this case, however, the terms of higher than quadratic order in $V_0$ appear. For a linear system one can neglect these and use Eq. (8.90). Thus we have

$$|\langle m|x|n \rangle| = \frac{i}{\hbar} \langle n | [H, x] | m \rangle = \frac{i}{\hbar} (E_m \langle m|x|n \rangle - E_n \langle m|x|n \rangle)$$  \hspace{1cm} (8.92)
and
\[ |\langle m|\dot{x}|n\rangle|^2 = \frac{-1}{\hbar^2}(E_m - E_n)^2 |\langle m|x|n\rangle|^2 = \omega^2 |\langle m|x|n\rangle|^2. \] (8.93)

Therefore
\[ \langle \frac{dE_n}{dt} \rangle = -\frac{\pi e^2}{\omega} V_0^2 \sum_{mn} |\langle m|\dot{x}|n\rangle|^2 \left[ f(E_n) - f(E_n + \hbar \omega) \right] \delta(E_m - E_n - \hbar \omega). \] (8.94)

If we now assume the current to be in phase with the applied voltage, the average energy dissipation becomes
\[ \text{Power} = \frac{1}{2} \frac{V_0^2}{R(\omega)} \] (8.95)
as the resistance \( R(\omega) \) is now equal to the impedance \( Z(\omega) \). Referring everything to unit volume, and noting that the resistance per unit volume is the resistivity, we get for the conductivity \( \sigma \)
\[ \sigma_{xx}(\omega) = -\frac{2\pi e^2}{\omega} \sum_{mn} |\langle m|v|n\rangle \langle n|v|m\rangle| \left[ f(E_n) - f(E_n + \hbar \omega) \right] \delta(E_m - E_n - \hbar \omega). \] (8.96)

A straightforward generalization of this procedure, using a perturbation
\[ H_{\text{ext}} = \sum_i V_0 e x_i / L \sin(\omega t), \] (8.97)
leads to the definition of an impedance matrix and a conductivity tensor
\[ \sigma_{ij} = \frac{2\pi e^2}{\omega} \sum_{mn} |\langle m|v_i|n\rangle \langle n|v_j|m\rangle| \left[ f(E_n) - f(E_n + \hbar \omega) \right] \delta(E_m - E_n - \hbar \omega), \] (8.98)
which is the Kubo-Greenwood equation. The derivation presented here confirms that fact that the linear response theory is based on the fluctuation-dissipation theorem, i.e. that the responses to an external perturbation are essentially determined by fluctuations of relevant physical quantities realized in the absence of the perturbation. Thus the linear response theory has a special appeal since it deals directly with the quantum mechanical motion of a process.

The linear response theory (or its equivalent) become soon a very popular tool of the transport theory. As was expressed by Langer, the Kubo formula "probably provides the most rigorous possible point of departure for transport theory. Despite its extremely formal appearance, it has in fact proved amenable to direct evaluation for some simple models". Edwards and Chester and Thellung have used the Kubo formula to calculate the impurity resistance of a system of independent electrons, and have recovered the usual solution of the linearized Boltzmann equation. Verboven has extended this work to higher orders in the concentration of impurities and has found corrections to the conductivity not originally derived via Boltzmann techniques. It was concluded that the Kubo formula might be most fruitfully applied in the full many-body problem, where it is not clear that any Boltzmann formulation is valid. However Izuyama casted doubt on the Kubo formula for electrical conductivity. He claimed that it is not, in fact, an exact formula for electrical conductivity, but is rather a coefficient relating current to an "external" field, which coefficient is equal to the conductivity only in special case. A correlation-function formula for electrical conductivity was derived by Magan by a formalism which gives prominence to the total electric field, including fields which may arise from the charged particles which are part of the system being studied. Langer have evaluated the impurity resistance of an interacting electron gas on the basis of
the Kubo formula at low but finite temperatures. The calculations are exact to all orders in the electron-electron interactions and to lowest order in the concentration of impurities. In the previous papers,\textsuperscript{271,272} the impurity resistance of this gas was computed at absolute zero temperature. It was shown\textsuperscript{266} that the zero-temperature limit of this calculation yields the previous result. In Ref.,\textsuperscript{225} Kubo formula for thermal conductivity was evaluated for the case of an interacting electron gas and random, fixed, impurities. The heat flux was examined in some detail and it was shown that in a normal system where the many-body correlations are sufficiently weak, the Wiedemann-Franz law remains valid. The relationships between the transport coefficients of a metal in a strong magnetic field and at very low temperatures were discussed by Smrcka and Streda.\textsuperscript{273} Formulae describing the electron coefficients as functions of the conductivity were derived on the basis of the linear response theory. As was mentioned earlier, examples of such relations are the Wiedemann-Franz law for the heat conductivity $\kappa$ and the Mott rule\textsuperscript{207} for the thermopower $S$. It was shown that that the Wiedemann-Franz law and the Mott rule are obeyed even in the presence of a quantized magnetic field $\omega \tau > 1$ if the scattering of electrons is elastic and if $\hbar \omega \gg kT$.

A theoretical analysis, based on Kubo formalism, was made for the ferromagnetic Hall effect by Leribaux\textsuperscript{274} in the case of transport limited by electron-phonon scattering. The antisymmetric, off-diagonal conductivity was, to first order in magnetization, found to be of order zero in the electron-phonon interaction (assumed to be weak) and, to this order, was equivalent to Karplus and Luttinger results.\textsuperscript{275} Tanaka, Moorjani and Morita\textsuperscript{276} expressed the nonlinear transport coefficients in terms of many-time Green functions and made an attempt to calculate the higher-order transport coefficients. They applied their theory to the calculation of the nonlinear susceptibility of a Heisenberg ferromagnet and nonlinear polarizability problem. Schotte\textsuperscript{277} reconsidered the linear response theory to show the closeness of it and kinetic equations.

At the same time, several authors\textsuperscript{278–280} raised an important question as to the general validity of the correlation formulae for transport coefficients. The relation for the electrical conductivity was, in principle, not questioned as in this case (as was shown above) one may obtain it by a straightforward application of basic statistical mechanical principles and perturbation techniques. One simply calculates the response of the system to an electric field. What has been questioned, however, was the validity of the correlation relations for the transport coefficients of a fluid - the diffusion constant, the thermal conductivity, and the viscosity - where no such straightforward procedure as was used for the electrical conductivity was available.\textsuperscript{281}

However, Jackson and Mazur\textsuperscript{282} presented a derivation of the correlation formula for the viscosity which is similar in spirit, and as free of additional assumptions, as that for the electrical conductivity. The correlation formula for the viscosity was obtained by calculating statistically the first order response of a fluid, initially in equilibrium, to an external shearing force. It was shown that on the basis of this derivation, the correlation formula for the viscosity, the exactness of which had been questioned, was placed on as firm a theoretical basis as the Kubo relation for the electrical conductivity. In addition, Resibois\textsuperscript{283,284} demonstrated the complete equivalence between the kinetic approach developed by Prigogine and coworkers\textsuperscript{285} and the correlation function formalism for the calculation of linear thermal transport coefficients. It was shown that in both cases these transport coefficients are determined by the solution of an inhomogeneous integral equation for a one-particle distribution function which is the generalization to strongly coupled system of the Chapman-Enskog first approximation of the Boltzmann equation.\textsuperscript{45,286} Interesting remarks concerning the comparison of the linear response theory and Boltzmann equation approach were formulated by R. Peierls.\textsuperscript{225,287} Schofield\textsuperscript{288,289} elaborated a general derivation of the transport coefficients and thermal equilibrium correlation functions for a classical system having arbitrary number of microscopic conservation laws. This derivation gives both the structure of the correlation functions in the hydrodynamic (long wavelength) region and a generalized definition of the
transport coefficients for all wave-lengths and frequencies. A number of authors have given the formulation of nonlinear responses. It was shown that since in a nonlinear system fluctuation sources and transport coefficients may considerably depend on a nonequilibrium state of the system, nonlinear nonequilibrium thermodynamics should be a stochastic theory. From the other side the linearity of the theory itself was a source of many doubts. The most serious criticism of the Kubo linear response theory was formulated by van Kampen. He argued strenuously that the standard derivation of the response functions are incorrect. In his own words, ” the basic linearity assumption of linear response theory ... is completely unrealistic and incompatible with basic ideas of statistical mechanics of irreversible processes”. The main question raised by van Kampen concerned the logic of the linear response theory, not the results. As van Kampen expressed it, ”the task of statistical mechanics is not only to provide an expression for this (transport) coefficient in term of molecular quantities, but also to understand how such a linear dependence arises from the microscopic equations of motion”. The van Kampen’s objections to Kubo linear response theory can be reduced to the following points. In the linear response theory one solves the Liouville equation to first order in the external field (electric field). This is practically equivalent to following a perturbed trajectory in phase space in a vicinity of the order of of unperturbed trajectory. In the classical case trajectories are exponentially unstable and corresponding field should be very small. Kubo’s derivation supposed non-explicitly that macroscopic linearity (Ohm law, etc.) is the consequence of microscopic linearity, but these two notions are not identical. Macroscopic linearity is the result of averaging over many trajectories and is not the same as linear deviations from any one trajectory. In other words, van Kampen’s argument was based on the observation that due to the Lyapunov instability of phase-space trajectories, even a very small external field will rapidly drive any trajectory far away from the corresponding trajectory without field. Hence linear response theory which is based on the proportionality of the trajectory separation with the external field could only be expected to hold for extremely short times, of no physical interest.

Responses to van Kampen’s objections were given by many authors. The main arguments of these responses, were based on the deep analysis of the statistical mechanical behavior of the many-body system under the external perturbation. It was shown that in statistical mechanical calculations one deals with the probability distributions for the behavior of the many particles rather than to the behavior of an individual particle. An analysis of the structural stability of hyperbolic dynamics-averaging and some other aspects of the dynamical behavior shows that the linear separation of trajectories goes on long enough for Green-Kubo integrals to decay. Moreover, Naudts, Pule and Verbeure analyzed the long-time behavior of correlations between extensive variables for spin-lattice systems and showed that the Kubo formula, expressing the relaxation function in terms of of the linear response function, is exact in the thermodynamic limit.

It was mentioned above that in the 1950s and 1960s the fluctuation relations, the so-called Green-Kubo relations, were derived for the causal transport coefficients that are defined by causal linear constitutive relations such as Fourier law of heat flow or Newton law of viscosity. Later it was shown also that it was possible to derive an exact expression for linear transport coefficients which is valid for systems of arbitrary temperature, density. The Green-Kubo relations give exact mathematical expression for transport coefficients in terms of integrals of time correlation functions. More precisely, it was shown that linear transport coefficients are exactly related to the time dependence of equilibrium fluctuations in the conjugate flux. For a more detailed discussion of these questions see Refs.

To summarize, close to equilibrium, linear response theory and linear irreversible thermodynamics provide a relatively complete treatment. However, in systems where local thermodynamic equilibrium has broken down, and thermodynamic properties are not the same local functions of thermodynamic state variables such that they are at equilibrium, serious problems may ap-
The Nonequilibrium Statistical Operator Method and Kinetic Equations

The method of the nonequilibrium statistical operator was reviewed already in detail by us. In this section, we remind very briefly the main ideas of the nonequilibrium statistical operator (NSO) approach for the sake of a self-contained formulation. The precise definition of the nonequilibrium state is quite difficult and complicated, and is not uniquely specified. Thus the method of reducing the number of relevant variables was proposed. A large and important class of transport processes can reasonably be modelled in terms of a reduced number of macroscopic relevant variables. It was supposed that the equations of motion for the "relevant" variables (the space- and time-dependent thermodynamic "coordinates" of a many-body nonequilibrium system), can be derived directly from the Liouville equation. This can be done by defining a generalized canonical density operator depending only upon present values of the thermodynamic "coordinates". According to D. N. Zubarev the NSO method permits one to generalize the Gibbs ensemble method to the nonequilibrium case and to construct a nonequilibrium statistical operator which enables one to obtain the transport equations and calculate the kinetic coefficients in terms of correlation functions, and which, in the case of equilibrium, goes over to the Gibbs distribution. The basic hypothesis is that after small time-interval $\tau$ the nonequilibrium distribution is established. Moreover, it is supposed that it is weakly time-dependent by means of its parameter only. Then the statistical operator $\rho$ for $t \geq \tau$ can be considered as an "integral of motion" of the quantum Liouville equation

$$\frac{\partial \rho}{\partial t} + \frac{1}{i\hbar} [\rho, H] = 0. \quad (9.1)$$

Here $\partial \rho/\partial t$ denotes time differentiation with respect to the time variable on which the relevant parameters $F_m$ depend. It is important to note once again that $\rho$ depends on $t$ by means of $F_m(t)$ only. We may consider that the system is in thermal, material, and mechanical contact with a combination of thermal baths and reservoirs maintaining the given distribution of parameters $F_m$. For example, it can be the densities of energy, momentum, and particle number for the system which is macroscopically defined by given fields of temperature, chemical potential and velocity. It is assumed that the chosen set of parameters is sufficient to characterize macroscopically the state of the system. The set of the relevant parameters are dictated by the external conditions for the system under consideration and, therefore, the term $\partial \rho/\partial t$ appears as the result of the external influence upon the system; this influence causes that the system is non-stationary.

In order to describe the nonequilibrium process, it is also necessary to choose the reduced set of relevant operators $P_m$, where $m$ is the index (continuous or discrete). In the quantum case, all operators are considered to be in the Heisenberg representation

$$P_m(t) = \exp \left( \frac{iHt}{\hbar} \right) P_m \exp \left( \frac{-iHt}{\hbar} \right), \quad (9.2)$$

where $H$ does not depend on the time. The relevant operators may be scalars or vectors. The equations of motions for $P_m$ will lead to the suitable "evolution equations" In the quantum case

$$\frac{\partial P_m(t)}{\partial t} - \frac{1}{i\hbar} [P_m(t), H] = 0. \quad (9.3)$$

The time argument of the operator $P_m(t)$ denotes the Heisenberg representation with the Hamiltonian $H$ independent of time. Then we suppose that the state of the ensemble is described by a nonequilibrium statistical operator which is a functional of $P_m(t)$
\[ \rho(t) = \rho\{ \ldots P_m(t) \ldots \}. \] (9.4)

Then \( \rho(t) \) satisfies the Liouville equation \( (9.1) \). Hence the quasi-equilibrium (local-equilibrium) Gibbs-type distribution will have the form

\[ \rho_q = Q_q^{-1} \exp \left( - \sum_m F_m(t) P_m \right), \] (9.5)

where the parameters \( F_m(t) \) have the sense of time-dependent thermodynamic parameters, e.g., of temperature, chemical potential, and velocity (for the hydrodynamic stage), or the occupation numbers of one-particle states (for the kinetic stage). The statistical functional \( Q_q \) is defined by demanding that the operator \( \rho_q \) be normalized and equal to

\[ Q_q = \text{Tr} \exp \left( - \sum_m F_m(t) P_m \right). \] (9.6)

The kinetic equations are of great interest in the theory of transport processes. In the NSO approach, the main quantities involved are the following thermodynamically conjugate values:

\[ \langle P_m \rangle = - \frac{\delta \Omega}{\delta F_m(t)}; \quad F_m(t) = \frac{\delta S}{\delta \langle P_m \rangle}. \] (9.7)

The generalized transport equations which describe the time evolution of variables \( \langle P_m \rangle \) and \( F_m \) follow from the equation of motion for the \( P_m \), averaged with the nonequilibrium statistical operator \( (9.4) \). It reads

\[ \langle \dot{P}_m \rangle = - \sum_n \frac{\delta^2 \Omega}{\delta F_m(t) \delta F_n(t)} \dot{F}_n(t); \quad \dot{F}_m(t) = \sum_n \frac{\delta^2 S}{\delta \langle P_m \rangle \delta \langle P_n \rangle} \langle \dot{P}_n \rangle. \] (9.8)

### 10 Generalized Kinetic Equations and Electroconductivity

#### 10.1 Basic formulas

Let us consider a many-particle system in the quasi-equilibrium state. It is determined completely by the quasi-integrals of motion which are the internal parameters of the system. In this and following sections we will use the notation \( A_j \) for the relevant observables to distinguish it from the momentum operator \( \vec{P} \). Here, for the sake of simplicity, we shall mainly treat the simplest case of mechanical perturbations acting on the system. The total Hamiltonian of the system under the influence of homogeneous external perturbation, depending on time as \( \sim \exp(i\omega t) \) is written in the following form

\[ \mathcal{H}(t) = H + H_F(t), \quad H_F(t) = - \sum_j A_j F_j \exp(i\omega t). \] (10.1)

In the standard approach the statistical operator \( \rho \) can be considered as an "integral of motion" of the quantum Liouville equation

\[ \frac{\partial \rho(t)}{\partial t} + \frac{1}{i\hbar} [\rho(t), \mathcal{H}(t)] = 0. \] (10.2)

Using the ideas of the method of the nonequilibrium statistical operator, as it was described above, we can write

\[ \rho(t) = \varepsilon \int_{-\infty}^{t} dt_1 e^{-\varepsilon(t-t_1)} U(t, t_1) \rho(t_1) U^\dagger(t, t_1). \] (10.3)
The time-evolution operator \( U(t,t_1) \) satisfy the conditions
\[
\frac{\partial}{\partial t} U(t,t_1) = \frac{1}{i\hbar} \mathcal{H}(t) U(t,t_1),
\]
\[
\frac{\partial}{\partial t_1} U(t,t_1) = -\frac{1}{i\hbar} \mathcal{H}(t) U(t,t_1), \quad U(t,t) = 1.
\]

If we consider the special case in which \( \rho(t_1) \to \rho_0 \), where \( \rho_0 \) is an equilibrium solution of the quantum Liouville equation \([10.2]\), then we can find the nonequilibrium statistical operator from the following equation:
\[
\frac{\partial \rho^K_\varepsilon(t)}{\partial t} + \frac{1}{i\hbar} [\rho^K_\varepsilon(t), \mathcal{H}(t)] = -\varepsilon (\rho^K_\varepsilon(t) - \rho_0).
\]
(10.4)

In the \( \lim_{\varepsilon \to 0^+} \) the nonequilibrium statistical operator will corresponds to the Kubo density matrix
\[
\rho^K_\varepsilon(t) = \rho_0 = \frac{1}{i\hbar} \int_{-\infty}^{t} dt_1 e^{-\varepsilon(t-t_1)} U(t,t_1)[\rho_0, \mathcal{H}(t_1)] U^\dagger(t,t_1).
\]
(10.5)

An average of the observable \( B_j \) are defined as
\[
\lim_{\varepsilon \to 0^+} \text{Tr}(\rho^K_\varepsilon(t) B_j) = \langle B_j \rangle_t.
\]
The linear response approximation for the statistical operator is given by
\[
\rho^K_\varepsilon(t) = \rho_0 = \frac{1}{i\hbar} \sum_j A_j F_j \exp(i\omega t) \int_{-\infty}^{0} dt_1 e^{\varepsilon(t-t_1)} \exp \left( \frac{iHt_1}{\hbar} \right) [\rho_0, A_j] \exp \left( -\frac{iHt_1}{\hbar} \right).
\]
(10.6)

Using the obtained expression for the statistical operator the mean values of the relevant observables \( B_t \) can be calculated. To simplify notation only observables will be considered for which the mean value in the thermal equilibrium vanishes. In other words, in general the \( B_t \) will be replaced by \( B_t - \langle B_t \rangle_0 \), where \( \langle B_t \rangle_0 = \text{Tr}(\rho_0 B_t) \). We find
\[
\langle B_t \rangle_t \equiv \text{Tr}(\rho^K_\varepsilon(t) B_t) = \sum_j L_{ij}(\omega) F_j \exp(i\omega t),
\]
(10.7)
where the linear response coefficients (linear admittances) are given by
\[
L_{ij}(\omega) = \langle \dot{A}_j; B_i \rangle_{\omega-\varepsilon}, \quad (\varepsilon \to 0^+).
\]
(10.8)

This expression vanish for operators \( A_j \) commuting with the Hamiltonian \( \mathcal{H} \) of the system, i.e. the Kubo expressions for \( L_{ij}(\omega) \) vanish for all \( \omega \) where for \( \omega = 0 \) the correct result is given by
\[
\langle B_t \rangle = \beta \sum_j F_j \text{Tr}(\rho_0 A_j B_t) = \sum_j L_{ij}(\omega) F_j \exp(i\omega t).
\]
(10.9)

The scheme based on the NSO approach starts with the generalized Liouville equation
\[
\frac{\partial \rho_\varepsilon(t)}{\partial t} + \frac{1}{i\hbar} [\rho_\varepsilon(t), \mathcal{H}(t)] = -\varepsilon (\rho_\varepsilon(t) - \rho_\eta(t)).
\]
(10.10)

For the set of the relevant operators \( P_m \) it is follow that
\[
\frac{d}{dt} \text{Tr}(\rho_\varepsilon(t) P_m) = 0.
\]
(10.11)
Here notation are

\[ \rho_q(t) = Q^{-1} \exp[-\beta(H - \mu N - \sum m F_m(t)P_m)], \]  
\[ F_m(t) = F_m \exp(i\omega t), \quad P_m \rightarrow P_m - \langle P_m \rangle_0. \]

To find the approximate evolution equations, the \( \rho \) can be linearized with respect to the external fields \( F_j \) and parameters \( F_m \)

\[ \rho(t) = \rho_q(t) - \exp(i\omega t)\rho_0 \left( \int_{-\infty}^{0} dt_1 e^{(\varepsilon+i\omega)t_1} \int_{0}^{\beta} d\tau \right) \]
\[ \left[ -\sum_j F_j A_j(t - i\tau) + \sum_m F_m \dot{P}_m(t - i\tau) + i\omega \sum_m F_m P_m(t - i\tau) \right]. \]

As a result we find

\[ \sum_n F_n \left( \langle \dot{P}_n; P_m \rangle_{\omega-i\varepsilon} + i\omega \langle P_n; P_m \rangle_{\omega-i\varepsilon} \right) = \sum_j F_j \langle \dot{A}_j; P_m \rangle_{\omega-i\varepsilon}. \]

In other notation we obtain

\[ \sum_n F_n \left( \langle \dot{P}_n; P_m \rangle + \langle \dot{P}_n; P_m \rangle_{\omega-i\varepsilon} + i\omega \left[ \langle P_n; P_m \rangle + \langle P_n; \dot{P}_m \rangle_{\omega-i\varepsilon} \right] \right) \]
\[ \quad = \sum_j F_j \left( \langle \dot{A}_j; P_m \rangle + \langle \dot{A}_j; \dot{P}_m \rangle_{\omega-i\varepsilon} \right), \]

where

\[ \langle A; B \rangle_{\omega-i\varepsilon} = \int_{-\infty}^{\infty} dt e^{(\varepsilon-i\omega)t} \langle A(t) | B \rangle; \]
\[ \langle A|B \rangle = \int_{0}^{\beta} d\lambda \text{Tr}(\rho_0 A(-i\lambda)B), \quad \rho_0 = \frac{1}{Q} e^{-\beta H}; \]
\[ \chi_{ij} = \langle A_i | A_j \rangle, \quad \sum_k \chi_{ik} (\chi^{-1})_{kj} = \delta_{ij}. \]

Thus our generalized transport equation can be written in the following abbreviated form

\[ \sum_n F_n P_{nm} = \sum_j F_j K_{jm}. \]

### 10.2 Electrical conductivity

The general formalism of the nonequilibrium statistical operator has been the starting point of many calculations of transport coefficients in concrete physical systems. In the present section we consider some selected aspects of the theory of electron conductivity in transition metals and disordered alloys. We put \( \hbar = 1 \) for simplicity of notation. Let us consider the dc electrical conductivity

\[ \sigma = \frac{\alpha^2}{3m^2\Omega} \langle \vec{P}; \vec{P} \rangle = \frac{\alpha^2}{3m^2\Omega} \int_{-\infty}^{0} d\varepsilon e^{\varepsilon t} \int_{0}^{\beta} d\lambda \text{Tr}(\rho P(t - i\lambda)\vec{P}), \quad \varepsilon \rightarrow 0 \]
where \( \vec{P} \) is the total momentum of the electrons. Representing \( \vec{P} \) as a sum of operators \( A_i \) (relevant observables) which can be chosen properly to describe the system considered (see below), the corresponding correlation functions \( \langle A_i; A_j \rangle \) can be calculated by the set of equations

\[
\sum_k \left( i\varepsilon \delta_{ik} - i \sum_l (\hat{A}_l|A_j)(\chi^{-1})_{lk} + i \sum_l \Pi_{il}(\chi^{-1})_{lk} \right) \langle A_k; A_j \rangle = i\chi_{ij}, \quad (10.19)
\]

where

\[
\chi_{ij} = (A_i|A_j), \quad \sum_k \chi_{ik}(\chi^{-1})_{kj} = \delta_{ij}; \quad (10.20)
\]

\[
(A_i|A_j) = \int_0^\beta d\lambda \text{Tr}(\rho A_i(-i\lambda)A_j), \quad \rho = \frac{1}{Q} e^{-\beta H}; \quad (10.21)
\]

\[
\Pi_{ij} = \langle \hat{A}_i \hat{C} | C \hat{A}_j \rangle^C, \quad \hat{A}_l = i[H, A_i]. \quad (10.22)
\]

The operator \( \hat{C} = 1 - \hat{P} \) is a projection operator with \( \hat{P} = \sum_{ij} |A_i)(\chi^{-1})_{ij}A_i \) and \( \langle \ldots \rangle^C \) denotes that in the time evolution of this correlation function, operator \( L = i[H, \ldots] \) is to be replaced by \( CLC \). By solving the set of equations (10.19) the correlation functions \( \Pi_{ij} \) which we started from are replaced by the correlation functions \( \Pi_{ij} \) and with a proper choice of the relevant observables these correlation functions can be calculated in a fairly simple approximation. It is, however, difficult to go beyond this first approximation and, in particular, to take into account the projection operators. Furthermore this method is restricted to the calculation of transport coefficients where the exact linear response expressions are known; generalization to thermal transport coefficients is not trivial. In Ref. we described a kind of general formalism for the calculation of transport coefficients which includes the approaches discussed above and which can be adapted to the problem investigated.

For simplicity of notation we restrict our consideration here on the influence of a stationary external electrical field. In the linear response theory the density matrix of the system becomes

\[
\rho_{LR} = \frac{e \vec{E}}{m} \int_0^t dt' e^{\varepsilon(t'-t)} \int_0^\beta d\lambda \rho \vec{P}(t' - t - i\lambda), \quad \varepsilon \to 0, \quad (10.23)
\]

where the time dependence in \( \vec{P}(t) \) is given by the total Hamiltonian of the system without the interaction term with the electrical field. From another point of view we can say that there is a reaction of the system on the external field which can be described by relevant observables such as shift of the Fermi body or a redistribution of the single particle occupation numbers, etc. Hence, for small external fields the system can be described in a fairly good approximation by the quasi-equilibrium statistical operator of the form

\[
\rho_q = \frac{1}{Q_q} e^{-\beta(H - \sum_i \alpha_i A_i)}, \quad (10.24)
\]

where the \( A_i \) are the observables relevant for the reaction of the system and the \( \alpha_i \) are parameters proportional to the external field. Of course, the statistical operator (10.24) is not a solution of the Liouville equation, but an exact solution can be found starting from (10.24) as an initial condition:

\[
\rho_s = \rho_q - i \int_0^\omega dt' e^{\varepsilon(t'-t)} \exp(iH_s(t' - t))\rho_q \exp(-iH_s(t' - t)), \quad (10.25)
\]

where \( H_s = H - e\vec{E} \sum_i \vec{r}_i \). In order to determine the parameters \( \alpha_i \) we demand that the mean values of the relevant observables \( A_i \) are equal in the quasi-equilibrium state \( \rho_q \) and in the real state \( \rho_s \), i.e.

\[
\text{Tr}(\rho_s A_i) = \text{Tr}(\rho_q A_i). \quad (10.26)
\]
This condition is equivalent to the stationarity condition
\[
\frac{d}{dt} \text{Tr}(\rho_s A_i) = 0.
\]

For a sufficiently complete set of operators \( A_i \) the condition (10.26) ensures that \( \rho_q \) describes the system with a sufficient accuracy. Linearizing Eq. (10.26) with respect to the parameters \( \alpha_i \) and the external field \( E \), one obtains the set of equations
\[
\sum_j \alpha_j \left( -i \text{Tr}(\rho [A_j, A_i]) + \langle A_j; A_i \rangle \right) = \frac{e \vec{E}}{m} \left( \langle \vec{P} | A_i \rangle + \langle A_i | \vec{P} \rangle \right).
\]

This set of equations can be shown to be equivalent to Eq. (10.19); whereas in the higher orders of interaction the equations (10.27) are more convenient to handle because the time dependence is given here in an explicit form without any projection.

With the parameters \( \alpha_i \) the current density is given by
\[
\vec{J} = \frac{e}{m \Omega} \text{Tr}(\rho_q \vec{P}) = \frac{e}{m \Omega} \sum_j \alpha_j \langle A_j | \vec{P} \rangle.
\]

Supposing that the total momentum \( \vec{P} \) of the electrons can be built up by the operators \( A_i \), it can be shown that the Eqs. (10.27) and (10.28) include the Kubo expression for the conductivity (see below). In order to solve the system of equations (10.27) a generalized variational principle can be formulated, but the reduction of the number of parameters \( \alpha_i \) by a variational ansatz corresponds to a new restricted choice of the relevant observable \( A_i \).

11 Resistivity of Transition Metal with Non-spherical Fermi Surface

The applicability of the transport equations (10.27) and (10.28) derived above to a given problem depends strongly on the choice of the relevant operators \( A_i \). The first condition to be fulfilled is that the mean values of the occupation numbers of all quasiparticles involved in the transport process should be time independent, i.e.
\[
\text{Tr}(\rho_s(\alpha_i) n_k) = \text{Tr}(\rho_q(\alpha_i) n_k), \quad \frac{d}{dt} \text{Tr}(\rho_s(\alpha_i) n_k) = 0.
\]

Of course, this condition is fulfilled trivially for \( A_i \to n_k \), but in this case the equations (10.27) cannot be solved in practice. In the most cases, however, there can be found a reduced set of operators sufficiently complete to describe the reaction of the system on the external field. It can be shown that under certain conditions the scattering process can be described by one relaxation time only and then the set of relevant operators reduces to the total momentum of the electrons describing a homogeneous shift of the Fermi body in the Bloch (momentum) space. These conditions are fulfilled for a spherical Fermi body at temperatures small in comparison to the degeneration temperature and for an isotropic scattering mechanism where the scatterers remain in the thermal equilibrium. In this simple case the Eqs. (10.27) are reduced to the so-called resistivity formula. For non-spherical Fermi bodies the set of relevant observables has to be extended in order to take into account not only its shift in the \( k \)-space but also its deformation. Our aim is to develop a theory of electron conductivity for the one-band model of a transition metal. The model considered is the modified Hubbard model, which include the electron-electron interaction as well as the electron-lattice interaction within MTBA. The non-spherical Fermi
The studies of the electrical resistivity of many transition metals revealed some peculiarities. It is believed that these specific features are caused by the fairly complicated dispersion law of the carriers and the existence of the two subsystems of the electrons, namely the broad $s-p$ band and relatively narrow $d$-band. The Fermi surface of transition metals is far from the spherical form. In addition, the lattice dynamics and dispersion relations of the phonons is much more complicated than in simple metals. As a result, it is difficult to attribute the observed temperature dependence of the resistivity of transition metals to definite scattering mechanisms.

Here we investigate the influence of the electron dispersion on the electrical resistivity within a simplified but workable model. We consider an effective single-band model of transition metal with tight-binding dispersion relation of the electrons. We take into account the electron-electron and electron-lattice interactions within the extended Hubbard model. The electron-lattice interaction is described within the MTBA. For the calculation of the electrical conductivity the generalized kinetic equations are used which were derived by the nonequilibrium statistical operator method. In these kinetic equations the shift of the non-spherical Fermi surface and its deformation by the external electrical field are taken into account explicitly. By using the weak scattering limit the explicit expressions for the electrical resistivity are obtained and its temperature dependence is estimated.

We consider a transition metal model with one non-spherical Fermi body shifted in the $k$-space and deformed by the external electrical field. Hence, the Fermi surface equation $E(\vec{k}) = E_F$ is transformed into $\tilde{E}(\vec{k}) = E_F$, where

$$\tilde{E}(\vec{k}) = E(\vec{k}) + m\vec{v}_1 \frac{\partial E}{\partial \vec{k}} + m \sum_{i=1}^{n} \vec{v}_i \Phi^i(\vec{k}) \frac{\partial E}{\partial \vec{k}} + \ldots$$

The term proportional to $\vec{v}_1$ describes a homogeneous shift of the Fermi surface in the $k$-space and the last terms allow for deformations of the Fermi body. The polynomials $\Phi^i(\vec{k})$ have to be chosen corresponding to the symmetry of the crystal and in consequence of the equality $\tilde{E}(\vec{k} + \vec{G}) = \tilde{E}(\vec{k})$ (which is fulfilled in our tight-binding model) they should satisfy the relation

$$\Phi^i(\vec{k} + \vec{G}) = \Phi^i(\vec{k}),$$

where $\vec{G}$ is a reciprocal lattice vector. Thus the relevant operators are given by

$$A_i \rightarrow m \sum_{\vec{k}\sigma} \Phi^i(\vec{k}) \frac{\partial E}{\partial \vec{k}} \tau_{k\sigma}, \quad \Phi^1 = 1.$$  

(11.3)

For our tight-binding model we are restricting by the assumption that $\Phi^i \rightarrow 0$ for $i \geq 3$.

### 11.1 Generalized kinetic equations

The quantum-mechanical many-body system is described by the statistical operator $\rho_s$ obeying the modified Liouville equation of motion Eq.(10.10)

$$\frac{\partial \rho_s}{\partial t} + \frac{1}{i\hbar} [\rho_s, H_s] = -\varepsilon (\rho_s - \rho_q),$$

(11.4)

where $H_s = H + H_E$ is the total Hamiltonian of the system including the interaction with an external electrical field $H_E = -e\vec{E} \cdot \sum_i \vec{r}_i$ and $\rho_q$ is the quasi-equilibrium statistical operator. According to the NSO formalism the relevant operators $P_m$ should be selected. These operators
include all the relevant observables which describe the reaction of the system on the external electrical field. These relevant operators satisfy the condition

$$\text{Tr}(\rho_s(t,0)P_m) = \langle P_m \rangle = \langle P_m \rangle_q; \quad \text{Tr} \rho_s = 1. \quad (11.5)$$

This condition is equivalent to the stationarity condition

$$\frac{d}{dt} \langle P_m \rangle = 0 \rightarrow \langle [H_s, P_m] \rangle = 0. \quad (11.6)$$

In the framework of the linear response theory the operators $\rho_s$ and $\rho_q$ in Eqs. (11.5) and (11.6) should be expanded to the first order in the external electric field $\vec{E}$ and in the parameters $F_m$. Thus Eq. (11.6) becomes

$$\sum_n F_n \left( -i \text{Tr}(\rho[P_n, P_m]) + \langle \dot{P}_n; \dot{P}_m \rangle \right) = \frac{eE}{m} \left( \text{Tr}(\rho \vec{P}(-i\lambda)P_m) + \langle \vec{P}; \dot{P}_m \rangle \right), \quad (11.7)$$

where

$$\dot{P}_m = i[H_s, P_m], \quad (11.8)$$

$$\langle A; B \rangle = \int_{-\infty}^{0} dt e^{\epsilon t} \int_{0}^{\beta} d\lambda \text{Tr}(\rho A(t - i\lambda)B), \quad (11.9)$$

$$A(t) = \exp(iHt)A \exp(-iHt); \quad \rho = Q^{-1} \exp(-\beta H),$$

and $P$ is the total momentum of the electrons.

The equations (11.7) are generalized kinetic equations in which the relaxation times and particle numbers are expressed via the correlation functions. It will be shown below that these equations can be reduced to the Kubo formula for the electrical conductivity provided a relation $\vec{P}_e = \sum \vec{a}_i P_i$ can be found.

The generalized kinetic equations (11.7) can be solved and the parameters $F_m$ can be determined by using a variational principle. The current density is given by

$$\vec{j} = \frac{e}{m\Omega} \text{Tr}(\rho_s \vec{P}_e) = \frac{e}{m\Omega} \text{Tr}(\rho_q \vec{P}_e), \quad (11.10)$$

where the conditions (11.5) have been used and $\Omega$ is the volume of the system. Linearizing Eq. (11.10) in the parameter $F_m$ we find

$$\vec{j} = \frac{e}{m\Omega} \sum_m F_m \int_{0}^{\beta} d\lambda \text{Tr}(\rho P_m(t - i\lambda)\vec{P}_e) \equiv \frac{1}{R} \vec{E}, \quad (11.11)$$

where the proportionality of the $F_m$ to the external electrical field has been taken into account. For the tight-binding model Hamiltonian (4.65), (4.66) the proper set of operators $P_m$ is given by

$$P_e = P_1 = m \sum_{k\sigma} \frac{\partial E}{\partial k} \alpha_{k\sigma}^\dagger a_{k\sigma}, \quad (11.12)$$

$$P_i = m \sum_{k\sigma} \Phi_i(\vec{k}) \frac{\partial E}{\partial k} \alpha_{k\sigma}^\dagger a_{k\sigma}, \quad (i = 2 \ldots n). \quad (11.13)$$

The parameters $F_m$ are replaced by the generalized drift velocities. Then the quasi-equilibrium statistical operator $\rho_q$ take the form

$$\rho_q = \frac{1}{Q_q} \exp \left( -\beta[H + mv_1 \sum_{k\sigma} \frac{\partial E}{\partial k} \alpha_{k\sigma}^\dagger a_{k\sigma} + \sum_{i=2 \ldots n} mv_i \Phi_i(\vec{k}) \frac{\partial E}{\partial k} \alpha_{k\sigma}^\dagger a_{k\sigma}] \right), \quad (11.14)$$
It should be mentioned here that in general in $\rho_\nu$ the redistribution of the scatterers by collisions with electrons should be taken into consideration. For the electron-phonon problem, e.g., the phonon drag can be described by an additional term $\vec{v}_{ph}\vec{P}_{ph}$ in Eq. (11.2) where $\vec{v}_{ph}$ is the mean drift velocity and $\vec{P}_{ph}$ is the total momentum of the phonons. Here it will be supposed for simplicity that due to phonon-phonon Umklapp processes, etc., the phonon subsystem remains near thermal equilibrium. In the same way the above consideration can be generalized to many-band case. In the last case, the additional terms in (11.2), describing shift and deformation of other Fermi bodies should be taken into account.

For the tight-binding model Hamiltonian (4.65), (4.66) the time derivatives of the generalized momenta (generalized forces) in Eq. (11.7) are given by

$$\dot{P}_n \rightarrow \dot{P}_j = \dot{P}^{ee}_j + \dot{P}^{ei}_j,$$

$$\dot{P}^{ee}_j = i[H_{ee}, P_j] =$$

$$\frac{iU_m}{N} \sum_{k_1k_2k_3k_4G} \left( \Phi^j(k_4) \frac{\partial E}{\partial k_4} + \Phi^j(k_2) \frac{\partial E}{\partial k_2} - \Phi^j(k_3) \frac{\partial E}{\partial k_3} - \Phi^j(k_1) \frac{\partial E}{\partial k_1} \right).$$

$$\dot{P}^{ei}_j = m \sum_{k_1k_2qG} \sum_{\sigma\nu} g_{k_1k_2}^{\nu} \left( \Phi^j(k_2) \frac{\partial E}{\partial k_2} - \Phi^j(k_1) \frac{\partial E}{\partial k_1} \right).$$

We confine ourselves by the weak scattering limit. For this case the total Hamiltonian $H$ in evolution $A(t) = \exp(iHt)A\exp(-iHt)$ we replace by the Hamiltonian of the free quasiparticles $H^0 + H^0$. With this approximation the correlation functions in Eq. (11.7) can be calculated straightforwardly. We find that

$$\langle \dot{P}_j; \dot{P}_l \rangle \approx \langle \dot{P}^{ee}_j; \dot{P}^{ee}_l \rangle + \langle \dot{P}^{ei}_j; \dot{P}^{ei}_l \rangle.$$  

Restricting ourselves for simplicity to a cubic system, the correlation functions of the generalized forces are given by

$$\langle \dot{P}^{ee}_j; \dot{P}^{ee}_l \rangle =$$

$$\frac{U^2m^2\beta\pi}{N^2} \sum_{k_1k_2k_3k_4G} \sum_{k_1k_2k_3k_4G} A_j(k_1, k_2, k_3, k_4)A_l(k_1, k_2, k_3, k_4) \cdot$$

$$f_{k_1}(1 - f_{k_2})f_{k_3}(1 - f_{k_4})\delta(E(k_1) - E(k_2) + E(k_3) - E(k_4)) \cdot$$

$$\delta(k_1 - k_2 + k_3 - k_4 + \vec{G}),$$

$$\langle \dot{P}^{ei}_j; \dot{P}^{ei}_l \rangle =$$

$$2\pi m^2\beta \sum_{k_1k_2qG} \sum_{k_1k_2qG} \left( g_{k_1k_2}^{\nu} \right)^2 B_j(k_1, k_2)B_l(k_1, k_2) \cdot$$

$$f_{k_2}(1 - f_{k_1})N(q\nu)\delta(E(k_2) - E(k_1) + \omega(q\nu))\delta(k_2 - k_1 - \vec{q} + \vec{G}),$$

where

$$A_j(k_1, k_2, k_3, k_4) = \left( \Phi^j(k_4) \frac{\partial E}{\partial k_4} + \Phi^j(k_2) \frac{\partial E}{\partial k_2} - \Phi^j(k_3) \frac{\partial E}{\partial k_3} - \Phi^j(k_1) \frac{\partial E}{\partial k_1} \right),$$

$$B_j(k_1, k_2) = \left( \Phi^j(k_2) \frac{\partial E}{\partial k_2} - \Phi^j(k_1) \frac{\partial E}{\partial k_1} \right)$$

(11.22)
and

\[ f(E(k)) = f_k = [\exp \beta(E(k) - E_F) + 1]^{-1}, \]

\[ N(\omega(q\nu)) = N(q\nu) = [\exp \beta\omega(q\nu) - 1]^{-1}. \]

The correlation functions \( \langle \tilde{P}_1; \tilde{P}_1 \rangle \) vanish in the weak scattering limit. The generalized electron numbers in Eq. (11.13) become

\[ N_l = \frac{1}{m} \text{Tr} \left( \rho \tilde{P}_1(i\lambda); \tilde{P}_1 \right) = m\beta \sum_k \Phi^l(\vec{k}) \frac{\partial E}{\partial \vec{k}} f_k(1 - f_k). \]  

(11.23)

### 11.2 Temperature dependence of \( R \)

Let us consider the low-temperature dependence of the electrical resistivity obtained above. For this region we have

\[ \lim_{T \to 0} \beta f_k(1 - f_k) \to \delta(E(k) - E_F). \]

Thus the generalized electron numbers \( N_l \) in Eq. (11.23) do not depend on temperature, and the temperature dependence of \( R \) in Eq. (11.11) is given by the correlation functions (11.19) and (11.21). For the term arising from the electron-electron scattering we find

\[ \langle \tilde{P}^{ee}_j; \tilde{P}^{ee}_1 \rangle = \beta \int \int \int_{E_{max}} dE(k_1)dE(k_2)dE(k_3) F^{3}_{ll}(E(k_1), E(k_2), E(k_3)) \cdot \]

\[ f_{k_1}(1 - f_{k_2})f_{k_3}[1 - f(E(k_1) - E(k_2) + E(k_3))], \]  

(11.24)

where

\[ F^{3}_{ll}(E(k_1), E(k_2), E(k_3)) = \frac{U^2m^2\pi}{N^2} \frac{\Omega^3}{(2\pi)^6} \sum_G \int d^2S_1 \int d^2S_2 \int d^2S_3 \cdot \]

\[ \frac{A_j(k_1, k_2, k_3, k_1 - k_2 + k_3 + G)A_l(k_1, k_2, k_3, k_1 - k_2 + k_3 + G)}{\left| \frac{\partial E}{\partial \vec{k}_1} \right| \left| \frac{\partial E}{\partial \vec{k}_2} \right| \left| \frac{\partial E}{\partial \vec{k}_3} \right|} \cdot \]

\[ \delta(E(k_1) - E(k_2) + E(k_3) - E(k_1 - k_2 + k_3 + G)). \]  

(11.25)

With the substitution

\[ x = \beta(E(k_1) - E_F), \quad y = \beta(E(k_2) - E_F), \quad z = \beta(E(k_3) - E_F), \]

the expression (11.24) reads

\[ \langle \tilde{P}^{ee}_j; \tilde{P}^{ee}_1 \rangle = \beta^{-2} \int_{-\beta E_F}^{\beta E_{max} - E_F} \frac{1}{1 + \exp(x)} \frac{1}{1 + \exp(-y)} \frac{1}{1 + \exp(z)} \cdot \]

\[ \int dx dy dz \frac{dx dy dz}{1 + \exp(-x + y - z)} F^1 \left( \frac{x}{\beta} + E_F, \frac{y}{\beta} + E_F, \frac{z}{\beta} + E_F \right) \]

\[ = \beta^{-2} A^{ee}_{jl}. \]  

(11.26)

It is reasonably to conclude from this expression, that in the limits

\[ \lim \beta E_F \to \infty, \quad \lim \beta(E_{max} - E_F) \to \infty, \]

the electron-electron correlation function for low temperatures becomes proportional to \( T^2 \) for any polynomial \( \Phi^j(\vec{k}) \).

For the electron-phonon contributions to the resistivity the temperature dependence is given by
the Bose distribution function of phonons \( N(q\nu) \). Because of the quasi-momentum conservation law \( \vec{q} = \vec{k}_2 - \vec{k}_1 - \vec{G} \) the contribution of the electron-phonon Umklapp processes freezes out at low temperatures as \( \exp(\beta \omega(q_{\text{min}}) - 1) \)⁻¹, where \( q_{\text{min}} \) is the minimal distance between the closed Fermi surfaces in the extended zone scheme. For electron-phonon normal processes as well as for electron-phonon Umklapp processes in metals with an open Fermi surface the quasi-momentum conservation law can be fulfilled for phonons with \( q \to 0 \) being excited at low temperatures solely. To proceed further we use the relation \( \omega(q\nu) = q \nu \vec{v}_F(q/\vec{q}) \) and the periodicity of the quasiparticle dispersion relation with the reciprocal lattice vector \( \vec{G} \). Taking into account the relation \( \Phi^i(\vec{k} + \vec{G}) = \Phi^i(\vec{k}) \), we find to the first non-vanishing order in \( q \)

\[
g_{k,k+q+G}^\nu \approx q \left( \frac{\vec{q}}{q} \frac{\partial}{\partial k'} \right) g_{k,k'}^\nu |_{k'=k}, \tag{11.27}
\]

\[
B_j(k, k + q + G) \approx q \left( \frac{\vec{q}}{q} \frac{\partial}{\partial k'} \right) B_j(k, k') |_{k'=k}, \tag{11.28}
\]

\[
\delta(E(k + q + G) - E(k) - \omega(q\nu)) \approx \frac{1}{q} \delta \left( \frac{\vec{q}}{q} \frac{\partial E}{\partial k} + \nu_0^\nu \right). \tag{11.29}
\]

Hence, we have

\[
\langle \dot{P}_j^\nu; \dot{P}_l^\nu \rangle \approx m^2 \beta \frac{\Omega^2}{(2\pi)^3} \sum_\nu \int_0^{q_{\text{max}}} q^5 dq \int \sin(\theta_q) d\theta_q \int d\nu | \exp(\beta \nu_0 q) - 1 |^{-1} F_{jl}^2(\theta_q, \varphi_q), \tag{11.30}
\]

where

\[
F_{jl}^2 = \int dk \left( \left( \frac{\vec{q}}{q} \frac{\partial}{\partial k'} \right) g_{k,k'}^\nu |_{k'=k} \right)^2 \left( \left( \frac{\vec{q}}{q} \frac{\partial}{\partial k} \right) B_j(k, k') |_{k'=k} \right) \cdot
\]

\[
\left( \left( \frac{\vec{q}}{q} \frac{\partial}{\partial k} \right) B_j(k, k') |_{k'=k} \right) f_k(1 - f_k) \delta \left( \frac{\vec{q}}{q} \frac{\partial E}{\partial k} + \nu_0^\nu \right). \tag{11.31}
\]

In the \( k \)-integral in Eq.\((11.30)\) the integration limits have to be chosen differently for normal and Umklapp processes. With the substitution \( x = \beta \nu_0^\nu q \) we find

\[
\langle \dot{P}_j^\nu; \dot{P}_l^\nu \rangle = \beta^{-5} m^2 \Omega^2 \sum_\nu \frac{1}{(\nu_0^\nu)^5} \int_0^{\nu_{\text{max}}} dx \frac{x^5}{e^x - 1} \int \sin(\theta_q) d\theta_q \int d\varphi_q F_{jl}^2(\theta_q, \varphi_q) = A_{jl}^5 T^5. \tag{11.32}
\]

Thus we can conclude that the electron-phonon correlation function is proportional to \( T^5 \) for any polynomial \( \Phi^i \). It is worthy to note that for an open Fermi surface this proportionality follows for normal and Umklapp processes either. For a closed Fermi surface the Umklapp processes freeze out at sufficiently low temperatures, and the electron-phonon normal processes contribute to the electrical resistivity only. With the aid of the Eqs.\((11.26)\) and \((11.32)\) the generalized kinetic equations \((11.7)\) becomes

\[
\sum_{i=1}^n v_i \left( A_{ij}^e T^2 + A_{ij}^e T^5 \right) = e EN_j, \tag{11.33}
\]

For simplicity we restrict our consideration to two parameters \( v_1 \) and \( v_2 \) describing the homogeneous shift and one type of deformation of the Fermi body. Taking into consideration more parameters is straightforward but does not modify very much qualitative results. Finally, the expression for the electrical resistivity becomes

\[
R = \frac{\Omega}{3e^2 N_1^2 (A_{11}^e T^2 + A_{11}^e T^5) (A_{22}^e T^2 + A_{22}^e T^5) + N_2^2 (A_{11}^e T^2 + A_{11}^e T^5)^2 - (A_{12}^e T^2 + A_{12}^e T^5)^2 - 2N_1 N_2 (A_{12}^e T^2 + A_{12}^e T^5)^2}. \tag{11.34}
\]
In general, a simple dependence $R \sim T^2$ or $R \sim T^5$ can be expected only if one of the scattering mechanisms dominate. For example, when $A_{ij}^{ee} \approx 0$, we find

$$R = \Omega \frac{(A_{11}^{ei}A_{22}^{ei} - A_{12}^{ei})T^5}{3e^2 N_2^2 A_{22}^{ei} + N_2^2 A_{11}^{ei} - 2N_1 N_2 A_{12}^{ei}}. \quad (11.35)$$

If, on the other hand, the deformation of the Fermi body is negligible ($v_2 = 0$), then from Eq. (11.34) it follows that

$$R = \Omega \frac{A_{11}^{ei}T^2 + A_{12}^{ei}T^5}{3e^2 N_1^2}.$$ \quad (11.36)

It is interesting to note that a somewhat similar in structure to expression (11.34) have been used to describe the resistivity of the so-called strong-scattering metals. In order to improve our formula for the resistivity derived above the few bands and the interband scattering (e.g. $s - d$ scattering) as well as the phonon drag effects should be taken into account.

### 11.3 Equivalence of NSO approach and Kubo formalism

Equivalence of the generalized kinetic equations to the Kubo formula for the electrical resistivity can be outlined as following. Let us consider the generalized kinetic equations

$$\sum_n F_n \left(-i\text{Tr}(\rho[P_n, P_m]) + \langle \tilde{P}_n; \tilde{P}_m \rangle \right) = \frac{eE}{m} \left(\text{Tr}(\rho\tilde{P}_e(-i\lambda)P_m) + \langle \tilde{P}_e; \tilde{P}_m \rangle \right). \quad (11.37)$$

To establish the correspondence of these equations with the Kubo expression for the electrical resistivity, it is necessary to express the operators of the total electron momentum $\tilde{P}_e$ and the current density $\tilde{j}$ in terms of the operators $P_m$. In the other words, we suppose that there exists a suitable set of coefficients $a_i$ with the properties

$$\tilde{P}_e = \sum_i a_i P_i, \quad \tilde{j} = \frac{e}{m\Omega} \tilde{P}_e. \quad (11.38)$$

We get, by integrating Eq. (11.37) by parts, the following relation

$$\sum_n F_n \left(\int_0^\beta d\lambda \text{Tr}(\rho P_n(-i\lambda)P_m) - \varepsilon \langle P_n; P_m \rangle \right) = \frac{eE}{m} \langle \tilde{P}_e; \tilde{P}_m \rangle. \quad (11.39)$$

Supposing the correlation function $\langle P_n; P_m \rangle$ to be finite and using Eq. (11.38) we find in the limit $\varepsilon \to 0$

$$\sum_i \sum_n a_i F_n \int_0^\beta d\lambda \text{Tr}(\rho P_n(-i\lambda)P_i) = \frac{eE}{m} \langle \tilde{P}_e; \tilde{P}_e \rangle. \quad (11.40)$$

From the equality (11.40) it follows that

$$\sum_n F_n \int_0^\beta d\lambda \text{Tr}(\rho P_n(-i\lambda)P_e) = \frac{eE}{m} \langle \tilde{P}_e; \tilde{P}_e \rangle. \quad (11.41)$$

Let us emphasize again that the condition

$$\lim_{\varepsilon \to 0} \varepsilon \langle P_n; P_m \rangle = 0 \quad (11.42)$$

is an additional one for a suitable choice of the operators $P_m$. It is, in the essence, a certain boundary condition for the kinetic equations (11.37). Since the Kubo expression for the electrical conductivity

$$\sigma \sim \langle \tilde{P}_e; \tilde{P}_e \rangle \sim \langle P_n; P_m \rangle \quad (11.43)$$
should be a finite quantity, the condition (11.42) seems quite reasonable. To make a following step, we must take into account Eq.(11.10) and (11.11). It is easy to see that the r.h.s. of the Eq.(11.41) is proportional to the current density. Finally, we reproduce the Kubo expression (10.18) (for cubic systems)

$$\sigma = \frac{j}{E} = \frac{e^2}{3m^2\Omega} \langle \vec{P}_e; \vec{P}_e \rangle,$$

(11.44)

where the proportionality of the $F_m$ to the external electrical field has been taken into account.

It will be instructive to consider a concrete problem to clarify some points discussed above. There are some cases when the calculation of electrical resistivity is more convenient to be performed within the approach of the generalized transport equations (10.27) and (10.28) than within the Kubo formalism for the conductivity. We can use these two approaches as two complementary calculating scheme, depending on its convenience to treat the problem considered. To clarify this, let us start from the condition

$$\text{Tr}(\rho_{LR} B_i) = \text{Tr}(\rho_s B_i),$$

(11.45)

where the density matrices $\rho_{LR}$ and $\rho_s$ were given by equations (10.23) and (10.25). For operators $B_i$ which can be represented by linear combinations of the relevant observables $A_i$ the equation (11.45) is fulfilled exactly where for other operators the equations (11.45) seems to be plausible if the relevant observables have been chosen properly. The conditions (11.45) make it possible to determine a set of parameters which can be used in approximate expressions for the correlation functions. In simple cases the conditions (11.45) even allow to calculate the correlation functions in (10.27) without resorting to another technique. As an example we consider the one-band Hubbard model (4.14)

$$H = H_e + H_{ee}$$

(11.46)

in the strongly-correlated limit $|t|/U \ll 1$. It well known that in this limit the band splits into two sub-bands separated by the correlation energy $U$. In order to take into account the band-split we have to project the one-electron operators onto the sub-bands. The relevant operators will have the form

$$\tilde{P}^{\alpha\beta} = m \sum_{k\sigma} \frac{\partial E}{\partial k} n_{\alpha\beta}^{k\sigma},$$

(11.47)

where

$$n_{\alpha\beta}^{k\sigma} = N^1 \sum_{ij} e^{i\vec{k}(\vec{R}_i-\vec{R}_j)} a_{i\alpha}^\dagger n_{i-\sigma}^{\alpha} a_{j\sigma} n_{j-\sigma}^{\beta},$$

(11.48)

with the projection operators

$$n_{i-\sigma}^{\alpha} = \begin{cases} n_{i-\sigma} & \text{if } \alpha = +, \\ (1 - n_{i-\sigma}) & \text{if } \alpha = +. \end{cases}$$

(11.49)

Here $\tilde{P}^{\alpha\alpha}$ is the operator of the total momentum of the electrons in the sub-band $\alpha$, and $\tilde{P}^{\alpha\beta}$, $(\alpha \neq \beta)$ describes kinematical transitions between the sub-bands. It can be shown that correlation functions $\langle \tilde{P}^{\alpha\beta}, \tilde{P}^{\gamma\delta} \rangle$ and $\langle \tilde{P}^{\alpha\alpha}, \tilde{P}^{\beta\beta} \rangle$ vanish for $\alpha \neq \beta$ in the limit $|t|/U \ll 1$. To make an estimation it is necessary to decouple the higher correlation functions in $|t|/U$ and take into account nearest neighbor hopping terms only. Then the correlation functions $\langle \tilde{P}^{\alpha\alpha}, \tilde{P}^{\alpha\alpha} \rangle (\alpha = \pm)$ can be calculated directly by means of Eq.(11.45), where $B_i \rightarrow \tilde{P}^{\alpha\alpha}$. The conductivity becomes

$$\sigma = WT^{-1} \sum_{\alpha} \sum_{\alpha} \langle n_{\sigma}^{\alpha} \rangle^{-1/2} \langle n_{\sigma}^{\alpha} n_{-\sigma}^{\alpha} \rangle (\langle n_{\sigma}^{\alpha} \rangle - \langle n_{\sigma}^{\alpha} n_{-\sigma}^{\alpha} \rangle),$$

(11.50)
where
\[ W = \frac{e^2}{3m^2\Omega k} \frac{1}{\sqrt{2z}} |t| \sum_k \left( \frac{\partial E}{\partial k} \right)^2 \] (11.51)
and
\[ \langle W \rangle = \text{Tr}(\rho W), \quad \rho = \frac{1}{Q} \exp(-\beta H). \]

Here \( z \) is the number of nearest neighbors and \( t \) the nearest neighbor hopping matrix element. With the well-known expressions for the mean values \( \langle n_\alpha^n \rangle \) we find the conductivity in dependence on the electron number in the form
\[ \sigma = WT^{-1} \frac{1}{2} \frac{n(1-n)}{\sqrt{1-n/2}}, \quad (0 \leq n < 1), \] (11.52)
\[ \sigma = WT^{-1} \frac{1}{2} \exp(-U/2kT), \quad (n = 1), \] (11.53)
\[ \sigma = WT^{-1} \frac{\sqrt{2}}{\sqrt{n}} (1-n/2)(1-n)/(1-n), \quad (2 \geq n > 1). \] (11.54)

Thus, it was shown in this and in the preceding sections that the formalism of the generalized kinetic equations has certain convenient features and its own specific in comparison with Kubo formalism. The derived expressions are compact and easy to handle with.

11.4 High-temperature resistivity and MTBA

At high temperatures the temperature dependence of the electrical resistivity \( R \) of some transition metals and highly resistive metallic systems such as A15 compounds may deviate substantially from the linear dependence, which follows from the Bloch-Gruneisen law. These strong deviations from the expected behavior with a tendency to flatten to a constant resistivity value was termed resistivity saturation\(^\text{314}\) and have been studied both experimentally and theoretically by many authors\(^\text{315–327}\). The phenomenon of resistivity saturation describes a less-than-linear rise in dc electrical resistivity \( R \) when temperature \( T \) increases. It was found that this effect is common in transition metal compounds (with pronounced \( d \)-band structure) when \( R \) exceeds \( \sim 80 \mu\Omega\text{cm} \), and that \( R \) seems bounded above by a value \( R_{\text{max}} \sim 150 \mu\Omega\text{cm} \) which varies somewhat with material. In Ref.\(^\text{326}\) in particular, it was formulated that the electrical resistivity, \( R \), of a metal is usually interpreted in terms of the mean free path (the average distance, \( l \), an electron travels before it is scattered). As the temperature is raised, the resistivity increases and the apparent mean free path is correspondingly reduced. In this semi-classical picture, the mean free path cannot be much shorter than the distance, \( a \), between two atoms. This has been confirmed for many systems and was considered to be a universal behavior. Recently, some apparent exceptions were found, including alkali-doped fullerenes and high-temperature superconductors\(^\text{328}\). However, there remains the possibility that these systems are in exotic states, with only a small fraction of the conduction electrons contributing to the conductivity; the mean free path would then have to be correspondingly larger to explain the observed resistivity. The authors of Ref.\(^\text{326}\) performed a model calculation of electron conduction in alkali-doped fullerenes, in which the electrons are scattered by intramolecular vibrations. The resistivity at large temperatures implies \( l \sim a \), demonstrating that there is no fundamental principle requiring \( l > a \). At high temperatures, the semi-classical picture breaks down, and the electrons cannot be described as quasiparticles. Recent review of theoretical and experimental investigations in this field was given in Refs.\(^\text{325,349}\) (for discussion of the electronic thermal conductivity at high temperatures see Ref.\(^\text{331}\)).

The nature of saturation phenomenon of electrical resistivity is not fully understood. Resistivity
of a metallic system as a function of temperature reflects an overall electron-phonon interaction effects as well as certain contribution effects of disorder.332–334 There have been some attempts to explain the saturation phenomenon in the framework of of the Boltzmann transport theory using special assumptions concerning the band structure, etc. The influence of electron-phonon scattering on electrical resistivity at high temperatures was investigated in Refs.323, 324 in the framework of the Fröhlich Hamiltonian for the electron-phonon interaction. In Ref.323 authors calculated a temperature-dependent self-energy to the lowest non-vanishing order of the electron-phonon interaction.

However, as it was shown above, for transition metals and their disordered alloys the modified tight-binding approximation is more adequate. Moreover the anisotropic effects are described better within MTBA. Here we consider a single-band model of transition metal with the Hamiltonian

$$H = H_e + H_i + H_{ei}. \quad (11.55)$$

The electron subsystem is described by the Hubbard model4 in the Hartree-Fock approximation

$$H_e = \sum_{k\sigma} E(k\sigma) a_{k\sigma}^\dagger a_{k\sigma}, \quad E(k\sigma) = E(k) + \frac{U}{N} \sum_p \langle n_{k+\sigma} \rangle. \quad (11.56)$$

For the tight-binding electrons in crystals we use $E(k) = 2\sum_{\alpha} t_0(\vec{R}_m) \cos(\vec{k}\vec{R}_m)$, where $t_0(\vec{R}_m)$ is the hopping integral between nearest neighbours, and $\vec{R}_m(k = x, y, z)$ denotes the lattice vectors in a simple lattice in an inversion centre. For the electron-phonon interaction we use the Hamiltonian

$$H_{ei} = \sum_{\sigma} \sum_{kq} V(\vec{k}, \vec{k} + \vec{q}) Q_{\vec{q}} a_{k+q\sigma}^\dagger a_{k\sigma}, \quad Q_{\vec{q}} = \frac{1}{\sqrt{2\omega(q)}} (b_{\vec{q}} + b_{-\vec{q}}^\dagger), \quad (11.57)$$

where

$$V(\vec{k}, \vec{k} + \vec{q}) = \frac{\hbar q_0}{(NM)^{1/2}} \sum_{k\nu} \epsilon^0(\vec{R}_m) \frac{\vec{e}_\nu(\vec{q})}{|\vec{R}_m|} [\sin \vec{R}_m \vec{k} - \sin \vec{R}_m (\vec{k} + \vec{q})]. \quad (11.58)$$

The one-electron hopping $\epsilon^0(\vec{R}_m)$ is the overlap integral between a given site $\vec{R}_m$ and one of the two nearby sites lying on the lattice axis $\vec{R}_m$. Operators $b_{\vec{q}}$ and $b_{\vec{q}}^\dagger$ are creation and annihilation phonon operators and $\omega(q)$ is the acoustical phonon frequency. $\mathcal{N}$ is the number of unit cells in the crystal and $M$ is the ion mass. The $\vec{e}_\nu(\vec{q})$ are the polarization vectors of the phonon modes.

For the ion subsystem we have

$$H_i = \sum_q \omega(\vec{q}) (b_q^\dagger b_q + 1/2). \quad (11.59)$$

For the resistivity calculation we use the following formula323

$$R = \frac{\Omega}{3e^2\mathcal{N}^2} \frac{\langle \vec{F}; \vec{F} \rangle}{1 + (1/3m\mathcal{N})\langle \vec{P}; \vec{F} \rangle}. \quad (11.60)$$

Here $\mathcal{N}$ is the effective number of electrons in the band considered

$$\mathcal{N} = \frac{1}{3m} \int_0^\beta d\lambda Tr(\rho\vec{P} (-i\hbar \lambda) \vec{P}) \quad (11.61)$$

and $\vec{P}$ is the total momentum operator

$$\vec{P} = \frac{m}{\hbar} \sum_{\vec{k}} \left( \frac{\partial E(\vec{k}\sigma)}{\partial \vec{k}} \right) n_{k\sigma}, \quad n_{k\sigma} = a_{k\sigma}^\dagger a_{k\sigma}. \quad (11.62)$$
The total force $\vec{F}$ acting on the electrons is given by

$$
\vec{F} = \frac{i}{\hbar} [H, \vec{P}] = -\frac{im}{\hbar} \sum_{kq\sigma} V(\vec{k}, \vec{k} + \vec{q})(\vec{v}_{k+q,\sigma} - \vec{v}_{k,\sigma})Q_q a_{k+q,\sigma}^\dagger a_{k,\sigma}, \tag{11.63}
$$

with the velocity defined as $\vec{v}_{k,\sigma} = \partial E(\vec{k}\sigma)/\hbar \partial \vec{k}$. It is convenient to introduce a notation

$$
\frac{V(\vec{k}, \vec{k} + \vec{q})}{\sqrt{2\omega(q)}} = \frac{i\Delta F_q}{\sqrt{\Omega}}.
$$

Correlation functions in Eq. (11.60) can be expressed in terms of the double-time thermodynamic Green functions

$$
\langle \vec{F}; \vec{F} \rangle = \frac{2\pi i}{\hbar} \langle \langle \vec{F} | \vec{P} \rangle \rangle_{-i\h\epsilon}, \tag{11.64}
$$

$$
\langle \vec{P}; \vec{F} \rangle = \frac{2\pi mi}{\hbar} \langle \langle \vec{P} | \vec{F} \rangle \rangle_{-i\h\epsilon}, \tag{11.65}
$$

$$
\langle \langle \vec{F} | A \rangle \rangle_{-i\h\epsilon} = \frac{1}{2\pi} \int_{-\infty}^{\infty} dt e^{\nu t} \theta(-t) \text{Tr} (\rho[A, F(t)]), \tag{11.66}
$$

where $A$ represents either the momentum operator $\vec{P}$ or the position operator $\vec{R}$ with $\vec{P} = im[H, \vec{R}] / \hbar$.

We find the following relation

$$
\langle \vec{F}; \vec{F} \rangle = \frac{2\pi m}{\hbar^2} \sum_{kq\sigma} \frac{V(\vec{k}, \vec{k} + \vec{q})}{\sqrt{2\omega(q)}} (\vec{v}_{k+q,\sigma} - \vec{v}_{k,\sigma}) \langle \langle a_{k+q,\sigma}^\dagger a_{k,\sigma}(b_q + b_{-q}^\dagger) | \vec{P} \rangle \rangle_{-i\h\epsilon}.
$$

Thus we obtain

$$
\langle \langle \vec{F}; \vec{A} \rangle \rangle_{-i\h\epsilon} = -i \sum_{kq\sigma} \frac{V(\vec{k}, \vec{k} + \vec{q})}{\sqrt{2\omega(q)}} (\vec{v}_{k+q,\sigma} - \vec{v}_{k,\sigma}) \left( \langle \langle a_{k+q,\sigma}^\dagger a_{k,\sigma} b_q | \vec{A} \rangle \rangle_{-i\h\epsilon} - \langle \langle a_{k+q,\sigma}^\dagger a_{k,\sigma} b_{-q}^\dagger | \vec{A} \rangle \rangle_{-i\h\epsilon} \right).
$$

Calculation of the higher-order Green functions gives

$$
(E(k + q\sigma) - E(k\sigma) - \Omega_q - i\h\epsilon) \langle \langle a_{k+q,\sigma}^\dagger a_{k,\sigma} B_q | \vec{A} \rangle \rangle_{-i\h\epsilon} =
$$

$$
T_{kq}(A) + \sum_{\vec{q}'} \frac{V(\vec{k} - \vec{q}', \vec{k})}{\sqrt{2\omega(q')}} \langle \langle a_{k+q',\sigma}^\dagger a_{k-q',\sigma} (b_{q'} - b_{-q'}^\dagger) B_q | \vec{A} \rangle \rangle_{-i\h\epsilon} -
$$

$$
\frac{V(\vec{k} + \vec{q}, \vec{k} + \vec{q} + \vec{q}')}{\sqrt{2\omega(q')}} \langle \langle a_{k+q+q',\sigma}^\dagger a_{k,\sigma} (b_{q'} - b_{-q'}^\dagger) B_q | \vec{A} \rangle \rangle_{-i\h\epsilon} -
$$

$$
- \sum_{\vec{k}'} \frac{V(\vec{k}', \vec{k}' - \vec{q})}{\sqrt{2\omega(q)}} \langle \langle a_{k+q,\sigma} a_{k',-q,\sigma}^\dagger a_{k',\sigma} | \vec{A} \rangle \rangle_{-i\h\epsilon}.
$$

(11.69)
with notation

\[ \Omega_q = h\omega(q) \rightarrow B_q = b_q; \quad E_q = -h\omega(q) \rightarrow B_q = b^\dagger_{-q}, \]

\[ T_{kq}(\vec{P}) = \frac{im}{2\pi} (\vec{v}_{k+q,\sigma} - \vec{v}_{k,\sigma}) (a_{k+q,\sigma}^\dagger a_{k,\sigma} B_q), \]

\[ T_{kq}(\vec{R}) = -\frac{1}{2\pi} \sum_{\vec{q}} \frac{\partial}{\partial q^\prime} (a_{k+q+q'^{*},\sigma}^\dagger a_{k+q^{*},\sigma} B_q) \delta_{q',0}, \]

\[ \mathcal{N} = \frac{m}{3\hbar} \sum_{kq,\sigma} \delta_{q',0} \vec{v}_{k,\sigma} \frac{\partial}{\partial q^\prime} (a_{k+q^{*},\sigma}^\dagger a_{k+q,\sigma}). \]

We find also

\[ \langle \langle a_{k+q,\sigma}^\dagger a_{k-\sigma} b_q b_{q'} | A \rangle \rangle_{-i\hbar\epsilon} = (11.70) \]

\[ -\langle b_q^\dagger b_{q'}^\dagger \rangle_0 \frac{V(\vec{k} - \vec{q}, \vec{k} - \vec{q}^{'*})}{\sqrt{2\omega(\vec{q}^{'*})}} \frac{\langle \langle a_{k+q,\sigma}^\dagger a_{k,\sigma} b_q b_{q'} | A \rangle \rangle_{-i\hbar\epsilon}}{(E(k + q\sigma) - E(k - q'\sigma) - E_{q'} - \hbar\omega(q') - i\hbar\epsilon)}, \]

\[ -\langle b_q^\dagger b_{q'}^\dagger \rangle_0 \frac{V(\vec{k} + \vec{q} + \vec{q}', \vec{k} + \vec{q})}{\sqrt{2\omega(\vec{q}^{'*})}} \frac{\langle \langle a_{k+q,\sigma}^\dagger a_{k,\sigma} b_q b_{q'} | A \rangle \rangle_{-i\hbar\epsilon}}{(E(k + q + q'\sigma) - E(k\sigma) - E_{q'} - \hbar\omega(q') - i\hbar\epsilon)}. \]

Here the symmetry relations

\[ V(\vec{k} - \vec{q}, \vec{k}) = V^{*}(\vec{k}, \vec{k} - \vec{q}); \quad V(\vec{k} + \vec{q} + \vec{q}', \vec{k} + \vec{q}) = V^{*}(\vec{k} + \vec{q}, \vec{k} + \vec{q} + \vec{q}^{'*}) \]

were taken into account.

Now the Green function of interest can be determined by introducing the self-energy.

\[ \langle \langle a_{k+q,\sigma}^\dagger a_{k-\sigma} b_q b_{q'} | A \rangle \rangle_{-i\hbar\epsilon} = T_{kq}(A) \]

\[ -\langle b_q^\dagger b_{q'}^\dagger \rangle_0 \frac{M_{kq\sigma}(E_q, -i\hbar\epsilon)}{(E(k + q\sigma) - E(k - q'\sigma) - E_{q'} - M_{kq\sigma}(E_q, -i\hbar\epsilon) - i\hbar\epsilon)}. \]

The self-energy is given by

\[ \sum_{\vec{q}} \frac{1}{2\omega(\vec{q}^{'*})} \left( \langle b_q^\dagger b_{q'}^\dagger \rangle_0 \frac{|V(\vec{k} - \vec{q}^{'*})|^2}{(E(k + q\sigma) - E(k - q'\sigma) - E_{q'} - \hbar\omega(q') - i\hbar\epsilon)} + \frac{|V(\vec{k} + \vec{q} + \vec{q}^{'*})|^2}{E(k + q + q'\sigma) - E(k\sigma) - E_{q'} - \hbar\omega(q') - i\hbar\epsilon} \right) \]

\[ + \langle b_q^\dagger b_{q'}^\dagger \rangle_0 \frac{|V(\vec{k} - \vec{q}^{'*})|^2}{E(k + q\sigma) - E(k - q'\sigma) - E_{q'} + \hbar\omega(q') - i\hbar\epsilon} + \frac{|V(\vec{k} + \vec{q} + \vec{q}^{'*})|^2}{E(k + q + q'\sigma) - E(k\sigma) - E_{q'} + \hbar\omega(q') - i\hbar\epsilon} \right). \]

In equation 11.73 the energy difference \((E(k + q\sigma) - E(k - q'\sigma) - E_{q'})\) is that for the scattering process of electrons on phonons, while emission or absorption of one phonon is possible, corresponding to \(E_q\). These scattering processes are contained in the usual Boltzmann transport theory.
leading to the Bloch-Gruneisen law. The self-energy $M_{k\sigma}$ describes multiple scattering corrections to the Bloch-Gruneisen behavior to second order in $V$, which depends on the temperature via the phonon occupation numbers.

Furthermore, it is assumed that the averages of occupation numbers for phonons in the self-energy and for electrons in the effective particle number, are replaced by the Bose and Fermi distribution functions, respectively

$$\langle b_q^\dagger b_q \rangle = N_q, \quad N_q = [\exp(\beta \hbar \omega(q)) - 1]^{-1},$$  \hspace{1cm} (11.74)  

$$\langle a_{k\sigma}^\dagger a_{k\sigma} \rangle = f_k, \quad f_k = [\exp(\beta E(k\sigma) - E_F) + 1]^{-1}. \hspace{1cm} (11.75)$$

This corresponds to neglecting the influence of multiple scattering corrections on the phonon and electron distribution functions.

In order to calculate the expectation values in the inhomogeneities, Eqs. (11.72) and (11.73), the spectral theorem should be used. In the lowest non-vanishing order of the electron-phonon interaction parameter $V$ we obtain

$$\langle a_{k+q\sigma}^\dagger a_{k\sigma} B_q \rangle = \frac{V(k + q, \bar{k})}{\sqrt{2\omega(q)}} f_{k+q}(1 - f_k)\nu_q(E_q) \frac{[\exp(\beta (E(k + q\sigma) - E(k\sigma) - E_q))] - 1}{E(k + q\sigma) - E(k\sigma) - E_q}, \hspace{1cm} (11.76)$$

with

$$\nu_q(E_q) = \frac{1}{1 - \exp(-\beta E_q)} = \begin{cases} 1 + N_q & \text{if } B_q = b_q; \quad E_q = \hbar \omega(q) \\ N_q & \text{if } B_q = b_q^\dagger; \quad E_q = -\hbar \omega(q) \end{cases} \hspace{1cm} (11.77)$$

Applying the approximation scheme discussed above we have found the following expressions for the Green function with $A = P$

$$\langle \langle a_{k+q\sigma}^\dagger a_{k\sigma} B_q | \hat{P} \rangle \rangle_{-i\hbar \varepsilon} = \frac{im V(k + q, \bar{k})}{2\pi \sqrt{2\omega(q)}} (\bar{v}_{k+q,\sigma} - \bar{v}_{k,\sigma}) P_{kq\sigma}^1(E_q) \frac{1}{\Omega_{kq\sigma}(E_q) - M_{kq\sigma} - i\hbar \varepsilon}, \hspace{1cm} (11.78)$$

and for the Green function with $A = R$

$$\langle \langle a_{k+q\sigma}^\dagger a_{k\sigma} B_q | \hat{R} \rangle \rangle_{-i\hbar \varepsilon} = \frac{h V(k + q, \bar{k})}{2\pi \sqrt{2\omega(q)}} (\bar{v}_{k+q,\sigma} - \bar{v}_{k,\sigma}) P_{kq\sigma}^2(E_q) \frac{1}{\Omega_{kq\sigma}(E_q) - M_{kq\sigma} - i\hbar \varepsilon}. \hspace{1cm} (11.79)$$

We have introduced in the above equations the following notation:

$$P_{kq\sigma}^{(1)}(E_q) = f_{k+q}(1 - f_k)\nu_q(E_q)\gamma_1(\Omega_{kq\sigma}(E_q)), \hspace{1cm} (11.80)$$

$$P_{kq\sigma}^{(2)}(E_q) = (v_{k+q,\sigma} - v_{k,\sigma}) f_{k+q}(1 - f_k)\nu_q(E_q) \left( \frac{\gamma_2(\Omega_{kq\sigma}(E_q))}{\Omega_{kq\sigma}(E_q)} - \frac{\beta \exp(\beta \Omega_{kq\sigma}(E_q))}{\Omega_{kq\sigma}(E_q)} \right), \hspace{1cm} (11.81)$$

$$P_{kq\sigma}^{(3)}(E_q) = f_{k+q}(1 - f_k)\nu_q(E_q)\beta \gamma_1(\Omega_{kq\sigma}(E_q))[f_k v_{k,\sigma} - (1 - f_{k+q})v_{k+q,\sigma}], \hspace{1cm} (11.82)$$

with

$$\gamma_n(\Omega_{kq\sigma}(E_q)) = \frac{\beta \exp(\beta \Omega_{kq\sigma}(E_q)) - 1}{(\Omega_{kq\sigma}(E_q))^n}, \hspace{1cm} (11.83)$$
and

$$\Omega_{kq\sigma}(E_q) = E(k + q\sigma) - E(k\sigma) - E_q. \quad (11.84)$$

For the effective particle number we find

$$\mathcal{N} = \frac{2}{3} m\beta \sum_k (v_k)^2 f_k(1 - f_k). \quad (11.85)$$

Before starting of calculation of the resistivity it is instructive to split the self-energy into real and imaginary part ($\varepsilon \to 0$)

$$\lim_{\varepsilon \to 0} M_{kq\sigma}(\hbar\omega(q) \pm i\varepsilon) = \text{Re} M_{kq\sigma}(\hbar\omega(q)) \mp i\text{Im} M_{kq\sigma}(\hbar\omega(q)) \quad (11.86)$$

and perform an interchange of variables $k + q \to k; q \to -q$. Now for the relevant correlation functions we obtain the expressions

$$\langle \vec{P}; \vec{P}^\dagger \rangle = \frac{2m^2}{\hbar} \sum_{kq\sigma} \frac{|V(\vec{k}, \vec{k} + \vec{q})|^2}{\sqrt{2\omega(q)}} (v_{k+q\sigma} - v_{k\sigma})^2 P_{kq\sigma}^{(1)}(\hbar\omega(q)) S_{kq\sigma}(\hbar\omega(q)), \quad (11.87)$$

$$-m^2 \sum_{kq\sigma} \frac{|V(\vec{k}, \vec{k} + \vec{q})|^2}{\sqrt{2\omega(q)}} (v_{k+q\sigma} - v_{k\sigma}) \left( P_{kq\sigma}^{(2)}(\hbar\omega(q)) + P_{kq\sigma}^{(3)}(\hbar\omega(q)) \right) S_{kq\sigma}(\hbar\omega(q)), \quad (11.88)$$

where

$$S_{kq\sigma}(\hbar\omega(q)) = \frac{\text{Im} M_{kq\sigma}(\hbar\omega(q))}{(\Omega_{kq\sigma}(\hbar\omega(q)) - \text{Re} M_{kq\sigma}(\hbar\omega(q)))^2 + (\text{Im} M_{kq\sigma}(\hbar\omega(q)))^2}. \quad (11.89)$$

In order to obtain Eq. (11.89) the following symmetry relation for the self-energy was used

$$M_{kq\sigma}(\hbar\omega(q) - i\varepsilon) = -M_{kq\sigma}(\hbar\omega(q) + i\varepsilon). \quad (11.90)$$

The inspection of both the correlation functions $\langle \vec{P}; \vec{P}^\dagger \rangle$ and $\langle \vec{P}; \vec{F}^\dagger \rangle$ shows that it include two dominant parts. The first one is the scattering part $S_{kq\sigma}(\hbar\omega(q))$, which contains all the information about the scattering processes. The second part describes the occupation possibilities before and after the scattering processes ($P_{kq\sigma}^{(1)}(\hbar\omega(q))$, $P_{kq\sigma}^{(2)}(\hbar\omega(q))$, $P_{kq\sigma}^{(3)}(\hbar\omega(q))$), and includes both the Fermi and Bose distribution functions. The approximation procedure described above neglects the multiple scattering corrections in these factors for the occupation possibilities.

For further estimation of the correlation functions the quasi-elastic approximation can be used. In this case, in the energy difference $\Omega_{kq\sigma}(\hbar\omega(q))$, the phonon energy $\hbar\omega(q)$ can be neglected $\Omega_{kq\sigma}(\hbar\omega(q)) \simeq \Omega_{kq\sigma}(0)$. The phonon wave number $q$ only is taken into account via the electron dispersion relation. Furthermore, for the Bose distribution function it was assumed that

$$\langle b_q^\dagger b_q \rangle = \langle b_q b_q^\dagger \rangle \simeq (\beta\hbar\omega(q))^{-1}. \quad (11.91)$$

This approximation is reasonable at temperatures which are high in comparison to the Debye temperature $\Theta_D$.

It is well known from Bloch-Gruneisen theory that the quasi-elastic approximation does not disturb the temperature dependence of the electrical resistivity at high or low temperatures. The absolute value of the resistivity is changed, but the qualitative picture of the power law of the temperature dependence of the resistivity is not influenced.
In the framework of the quasi-elastic approximation the scattering contribution can be represented in the form
\[ S_{kq\sigma} = \frac{\text{Im}M^e_{kq\sigma}}{(\Omega_{kq\sigma}(0) - \text{Re}M^e_{kq\sigma})^2 + (\text{Im}M^e_{kq\sigma})^2}. \]  

Here the real and imaginary parts of the self-energy have the following form
\[ \text{Re}M^e_{kq\sigma} = \frac{kT}{\hbar} \sum_p \left( \frac{|V(\vec{k}, \vec{k} + \vec{p})|^2}{\omega(p)^2} P \left( \frac{1}{E(k + q\sigma) - E(k - p\sigma)} \right) \right) + \frac{|V(\vec{k} + \vec{q}, \vec{k} + \vec{q} + \vec{p})|^2}{\omega(p)^2} P \left( \frac{1}{E(k + q + p\sigma) - E(k\sigma)} \right), \]  

\[ \text{Im}M^e_{kq\sigma} = \frac{\pi kT}{\hbar} \sum_p \left( \frac{|V(\vec{k}, \vec{k} + \vec{p})|^2}{\omega(p)^2} \delta(E(k + q\sigma) - E(k - p\sigma)) \right) + \frac{|V(\vec{k} + \vec{q}, \vec{k} + \vec{q} + \vec{p})|^2}{\omega(p)^2} \delta(E(k + q + p\sigma) - E(k\sigma)). \]  

The occupation possibilities given by \( P^{(n)}_{kq\sigma} \) can be represented in the quasi-elastic approximation as
\[ P^{(1)e}_{kq\sigma} \approx \frac{kT}{\hbar\omega(q)} \delta(E_F - E(k\sigma)); \]  

\[ P^{(2)e}_{kq\sigma} = P^{(3)e}_{kq\sigma} = 0. \]

In this approximation the momentum-force correlation function disappears \( \langle \vec{F}; \vec{F} \rangle \approx 0 \). Thus we have
\[ R \approx \frac{\Omega}{3e^2 N^2} \langle \vec{F}; \vec{F} \rangle, \]  

\[ \langle \vec{F}; \vec{F} \rangle = \frac{2m^2 kT}{\hbar} \sum_{kq\sigma} \frac{|V(\vec{k}, \vec{k} + \vec{q})|^2}{\omega(q)^2} (v_{k+q\sigma} - v_{k\sigma})^2 \delta(E_F - E(k\sigma)) \cdot \]  

\[ \frac{\text{Im}M^e_{kq\sigma}}{(E(k + q\sigma) - E(k\sigma) - \text{Re}M^e_{kq\sigma})^2 + (\text{Im}M^e_{kq\sigma})^2}. \]  

The explicit expression for the electrical resistivity was calculated in Ref. The additional simplifying assumptions have been made to achieve it. For the electrons and phonons the following simple dispersion relations were taken
\[ E(k) = \frac{\hbar^2 k^2}{2m^*}; \quad \omega(q) = v_0|q|, \quad V(\vec{k}, \vec{k} + \vec{q}) \sim \sqrt{|q|}. \]  

It was shown that the estimation of \( P^{(1)e}_{kq\sigma} \) is given by
\[ P^{(1)e}_{kq\sigma} \approx \frac{Tkm^*}{v_0 \hbar^3 k_F q} \delta(k_F - k). \]  

Then, for the electrical resistivity the following result was found
\[ R \approx \frac{3V^2 m^* q_D T}{2e^2 k_F^2 \hbar} \int_0^{q_D} dq \int_{-1}^1 dz q^4 \left( \frac{\text{Im}M^e_{kq\sigma}}{(h^2 q/m^*) (zk_F + q/2) - \text{Re}M^e_{kq\sigma}} \right)^2 + \left( \left| \text{Im}M^e_{kq\sigma} \right| \right)^2. \]  

75
This result shows that the usual Bloch-Gruneisen theory of the electrical resistivity can be corrected by including the self-energy in the final expression for the resistivity. The Bloch-Gruneisen theory can be reproduced in the weak scattering limit using the relation

\[
\lim_{\mathrm{Re}(\mathrm{Im})M \to 0} \frac{\mathrm{Im}M_{kF,qz}^e}{[\hbar^2q/m^*(zk_F + q/2) - \mathrm{Re}M_{kF,qz}^e]^2 + [\mathrm{Im}M_{kF,qz}^e]^2} = \frac{\pi m^*}{\hbar^2 qk_F} \delta(z + \frac{q}{2k_F}).
\]  \hspace{1cm} (11.101)

Inserting Eq. (11.101) in the resistivity expression Eq. (11.100) gives the electrical resistivity \( R_w \) in the weak scattering limit, showing a linear temperature dependence

\[
R_w \simeq \frac{3\pi}{8} \frac{V^2(m^*)^2 q_D^5 T}{e^2 k_F^6 \hbar^3 \Theta_D}.
\]  \hspace{1cm} (11.102)

In this form, the resistivity formula contains two main parameters that influence substantially. The first is the Debye temperature \( \Theta_D \) characterizing the phonon system, and, the second, the parameter \( \alpha = (Vm^*)^2 \) describing the influence of both the electron system and the strength of the electron-phonon interaction. The numerical estimations\(^323\) were carried out for \( Nb \) and gave the magnitude of the saturation resistivity 207 \( \mu \Omega \text{cm} \).

In the theory described above the deviation from linearity in the high-temperature region of the resistivity may be caused by multiple scattering corrections. The multiple scattering processes which describe the scattering processes of electrons on the phonon system by emission or absorption of more than one phonon in terms of self-energy corrections become more and more important with increasing temperature. As was shown above, even for simple dispersion relation of electrons and phonons within one-band model the thermally induced saturation phenomenon occurs. For the anisotropic model within MTBA the extensive numerical calculations are necessary.

In subsequent paper,\(^324\) Christoph and Schiller have considered the problem of the microscopic foundation of the empirical formula\(^314\) (parallel resistor model)

\[
\frac{1}{R(T)} = \frac{1}{R_{SBT}(T)} + \frac{1}{R_{max}(T)}
\]  \hspace{1cm} (11.103)

within the framework of the transport theory of Christoph and Kuzemsky.\(^308\) The parallel resistor formula describing the saturation phenomenon of electrical resistivity in systems with strong electron-phonon interaction was derived. In Eq. (11.103) \( R_{SBT}(T) \) is the resistivity given by the semiclassical Boltzmann transport theory \( R_{SBT}(T) \sim T \) and the saturation resistivity \( R_{max} \) corresponds to the maximum metallic resistivity.\(^333\) The higher-order terms in the electron-phonon interaction were described by a self-energy which was determined self-consistently. They found for the saturation resistivity the formula\(^324\)

\[
R_{max} = \frac{3\pi^3 \hbar q_D^4}{32 e^2 k_F^6 |P(q_D/2k_F)|}.
\]  \hspace{1cm} (11.104)

Within the frame of this approach, the saturation behavior of the electrical resistivity was explained by the influence of multiple scattering processes described by a temperature-dependent damping term of one-electron energies. In the standard picture, the conventional linear temperature dependence of the resistivity \( R \sim T(T \gg \Theta_D) \) is explained by taking into account that the number of phonons is proportional to the temperature and, moreover, assuming that the electron momentum is dissipated in single-phonon scattering processes only. For an increasing number of phonons, however, the multiple scattering processes become more important and the
single scattering event becomes less effective. This argument coincides in some sense with the Yoffe-Regel criterion stating that an increase in the number of scatterers does not result in a corresponding increase of the resistivity if the mean free path of the electrons becomes comparable with the lattice distance. Indeed, the saturation resistivity coincides roughly with the inverse minimal metallic conductivity which can be derived using this criterion.

12 Resistivity of Disordered Alloys

In the present section a theory of electroconductivity in disordered transition metal alloys with the proper microscopic treatment of the non-local electron-phonon interaction is considered. It was established long ago that any deviation from perfect periodicity will lead to a resistivity contribution which will depend upon the spatial extent and lifetime of the disturbance measured in relation to the conduction electron mean free path and relaxation time. It is especially important to develop a theory for the resistivity of concentrated alloys, because of its practical significance. The electrical resistivity of disordered metal alloys and its temperature coefficient is of considerable practical and theoretical interest. The work in this field has been considerably stimulated by Mooij's paper, where it has been shown that the temperature coefficient of the resistivity of disordered alloys becomes negative if their residual resistivity exceeds a given critical value. To explain this phenomenon, one has to go beyond the weak-scattering limit and to take into account the interference effects between the static disorder scattering and the electron-phonon scattering.

In the weak-scattering limit the contributions of impurity and phonon scattering add to the total resistivity without any interference terms (Matthiessen rule). For disordered systems many physical properties can be related to the configuration-averaged Green functions. The single-site CPA (coherent potential approximation) provides a convenient and accurate approximation for it. The CPA is a self-consistent method that predicts alloy electronic properties, interpolating between those of the pure constituents over the entire range of concentrations and scattering strengths. The self-consistency condition is introduced by requiring that the coherent potential, when placed at each lattice site of the ordered lattice, reproduces all the average properties of the actual crystal. The coherent-potential approximation has been developed within the framework of the multiple-scattering description of disordered systems. A given scatterer in the alloy can be viewed as being embedded in an effective medium with a complex energy-dependent potential whose choice is open and can be made self-consistently such that the average forward scattering from the real scatterer is the same as free propagation in the effective medium. The strong scattering has been first considered by Velicky in the framework of the single-site CPA using the Kubo-Greenwood formula. These results have been extended later to a more general models. The first attempt to include the electron-phonon scattering in the CPA calculations of the resistivity was given by Chen et al.

A model was introduced in which phonons were treated phenomenologically while electrons were described in CPA. The electron-phonon interaction was described by a local operator. Chen, Weisz and Sher (CWS) have performed a model calculation on the temperature dependence of the electronic density of states and the electrical conductivity of disordered binary alloys, based on CPA solutions by introducing thermal disorder in the single-band model. They found that the effect of thermal disorder is to broaden and smear the static-alloy density of states. The electrical conductivity in weak scattering alloys always decreases with temperature. However, in the strong-scattering case, the temperature coefficient of conductivity can be negative, zero, or positive, depending on the location of the Fermi energy. Brouers and Brauwers have extended the calculation to an $s - d$ two-band model that accounts for
the general behavior of the temperature dependence of the electrical resistivity in concentrated transition metal alloys. In Ref. 358 a generalization of CWS theory 340 was made by including the effect of uniaxial strain on the temperature variation of the electronic density of states and the electrical conductivity of disordered concentrated binary alloys. The validity of the adiabatic approximation in strong-scattering alloys was analyzed by Chen, Sher and Weisz. 359 It was shown that the electron screening process in the moving lattice may be modified by lattice motion in disordered alloys. Were this modification significant, not only the effective Hamiltonian but also the whole adiabatic approximation would need to be reconsidered.

A consistent theory of the electroconductivity in disordered transition metal alloys with the proper microscopic treatment of the electron-phonon interaction was carried out by Christoph and Kuzemsky 310. They used the approach of paper 180 where a self-consistent microscopic theory for the calculation of one-particle Green functions for the electron-phonon problem in disordered transition metal alloys was developed. However, this approach cannot be simply generalized to the calculation of two-particle Green functions needed for the calculation of the conductivity by the Kubo formula. Therefore, for the sake of simplicity, in their study Christoph and Kuzemsky 310 neglected the influence of disorder on the phonons. Thus, in the model investigated here, in contrast to the GWS approach 340 the dynamics of the phonons is taken into account microscopically, but they are treated as in a virtual reference crystal.

For a given configuration of atoms the total Hamiltonian of the electron-ion system in the substitutionally disordered alloy can be written in the form 180, 310

\[ H = H_e + H_i + H_{ei}, \]  

(12.1)

where

\[ H_e = \sum_{i\sigma} \epsilon_i a^\dagger_{i\sigma} a_{i\sigma} + \sum_{ij\sigma} t_{ij} a^\dagger_{i\sigma} a_{j\sigma} \]  

(12.2)

is the one-particle Hamiltonian of the electrons. For our main interest is the description of the electron-phonon interaction, we can suppose that the electron-electron correlation in the Hubbard form has been taken here in the Hartree-Fock approximation in analogy with Eq. (11.56).

For simplicity, in this paper the vibrating ion system will be described by the usual phonon Hamiltonian

\[ H_i = \sum_{qv} \omega(q\nu)(b_{q\nu}^\dagger b_{q\nu} + 1/2). \]  

(12.3)

The electron-phonon interaction term is taken in the following form 180

\[ H_{ei} = \sum_{ij} \sum_{\alpha\sigma} T_{ij}^\alpha (u_i^\alpha - u_j^\alpha) a^\dagger_{i\sigma} a_{j\sigma}, \]  

(12.4)

where \( u_i^\alpha (\alpha = x, y, z) \) is the ion displacement from the equilibrium position \( \vec{R}_i \).

In terms of phonon operators this expression can be rewritten in the form

\[ H_{ei} = \sum_{i\neq j} \sum_{qv\sigma} A_{qv}(ij) (b_{q\nu} + b_{-q\nu}^\dagger) a^\dagger_{i\sigma} a_{j\sigma}. \]  

(12.5)

where

\[ A_{qv}(ij) = \frac{q_0}{\sqrt{2\langle M\rangle N \omega(q\nu)}} e^{i\vec{q} \cdot \vec{R}_j} e^{-i\vec{q} \cdot \vec{R}_i} \bar{e}_\nu(q) \left( e^{i\vec{q} \cdot \vec{R}_i} - e^{i\vec{q} \cdot \vec{R}_j} \right). \]  

(12.6)

Here \( \omega(q\nu) \) are the acoustic phonon frequencies, \( \langle M\rangle \) is the average ion mass, \( \bar{e}_\nu(q) \) are the polarization vectors of the phonons, and \( q_0 \) is the Slater coefficient originated in the exponential radial
decrease of the tight-binding electron wave function. It is convenient to rewrite this expression in the form

$$H_{ei} = \sum_{i \neq j} \sum_q A_q(ij)(b_q + b_{-q}^\dagger)a_i^\dagger a_j, \quad (12.7)$$

where the spin and phonon polarization indices are omitted for brevity.

The electrical conductivity will be calculated starting with the Kubo expression for the dc conductivity:

$$\sigma(i\varepsilon) = -\langle\langle \vec{J}\rangle\rangle_{i\varepsilon}, \quad (\varepsilon \to 0^+), \quad (12.8)$$

where $P = e \sum_i \vec{R}_ia_i^\dagger a_i$ and $\vec{R}_i$ is the position vector; $m/e\vec{J} = m/e\vec{P}$ is the current operator of the electrons. It has the form

$$\vec{J} = -ie \sum_{ij} (\vec{R}_i - \vec{R}_j)t_{ij}a_i^\dagger a_j, \quad (12.9)$$

Then the normalized conductivity becomes

$$\sigma^{\alpha\beta} = \frac{ie^2}{\Omega} \sum_{ij} \sum_l (\vec{R}_i - \vec{R}_j)^\alpha \vec{R}_l^\beta t_{ij} \langle\langle a_i^\dagger a_j|a_l^\dagger a_l \rangle\rangle_{i\varepsilon}, \quad (12.10)$$

where $\Omega$ is the volume of the system. It should be emphasized here that $\sigma^{\alpha\beta}$ depends on the configuration of the alloy. A realistic treatment of disordered alloys must involve a formalism to deal with one-electron Hamiltonian that include both diagonal and off-diagonal randomness. \[353–360\]

In the present study, for the sake of simplicity, we restrict ourselves to a diagonal disorder. Hence we can rewrite hopping integral $t_{ij}$ as

$$t_{ij} = \frac{1}{N} \sum_k E(k) \exp[i\vec{k}(\vec{R}_i - \vec{R}_j)]. \quad (12.11)$$

Thus to proceed it is necessary to find the Green function $G_{ij,lm} = \langle\langle a_i^\dagger a_j|a_l^\dagger a_m \rangle\rangle$. It can be calculated by the equation of motion method. Using the Hamiltonian (12.1) we find by a differentiation with respect to the left hand side

$$\sum_{nr} H_{ij,rn} G_{nr,lm}(\omega) = \langle\langle a_i^\dagger a_m \rangle\rangle_{ij} - \langle\langle a_i^\dagger a_j \rangle\rangle_{mi} + $$

$$\sum_{qn} \left( A_q(j - n)e^{i\vec{q}\vec{R}_j}\langle\langle a_i^\dagger a_n(b_q + b_{-q}^\dagger)|a_i^\dagger a_m \rangle\rangle - A_q(n - i)e^{i\vec{q}\vec{R}_n}\langle\langle a_i^\dagger a_j(b_q + b_{-q}^\dagger)|a_i^\dagger a_m \rangle\rangle \right), \quad (12.12)$$

where

$$H_{ij,rn} = (\omega - \epsilon_n + \epsilon_r)\delta_{ni}\delta_{rj} - t_{jr}\delta_{ni} + t_{mi}\delta_{rj}. \quad (12.13)$$

We define now the zeroth-order Green functions $G_{ij,lm}^0$ that obey the following equations of motion

$$\sum_{nr} H_{ij,rn} G_{nr,lm}^0 = \langle\langle a_i^\dagger a_m \rangle\rangle_{ij} - \langle\langle a_i^\dagger a_j \rangle\rangle_{mi}, \quad (12.14)$$

$$\sum_{nr} H_{rn,lm} G_{ij,nr}^0 = \langle\langle a_i^\dagger a_m \rangle\rangle_{ij} - \langle\langle a_i^\dagger a_j \rangle\rangle_{mi}, \quad (12.15)$$

79
where Eq. (12.15) has been obtained by a differentiation with respect to the right hand side of \( G_{ij,lm}^0 \). Using these definitions it can be shown that

\[
\sum_{nr} \langle \langle a_s^\dagger a_n \rangle \rangle \delta_{rt} - \langle \langle a_s^\dagger a_t \rangle \rangle \delta_{sn} \rangle G_{nr,lm}(\omega) =
\]

\[
\sum_{ij} \langle \langle a_s^\dagger a_m \rangle \rangle \delta_{ij} - \langle \langle a_s^\dagger a_j \rangle \rangle \delta_{mi} \rangle G_{st,ji}^0(\omega) +
\]

\[
\sum_{ijn} \sum_q \left( A_q(j - n)e^{iq\vec{R}_t}i\langle \langle a_s^\dagger a_n(b_q + b_{-q})|a_s^\dagger a_m \rangle \rangle -
\right.
\]

\[
A_q(n - i)e^{iq\vec{R}_n}i\langle \langle a_s^\dagger a_n(b_q + b_{-q})|a_s^\dagger a_m \rangle \rangle \right) G_{st,ji}^0(\omega) .
\] (12.16)

The r.h.s. higher order Green functions can be calculated in a similar way. To proceed we approximate the electron-phonon Green function as

\[
\langle \langle a_s^\dagger a_r b_q^\dagger b_q | B \rangle \rangle \approx N(q)\langle \langle a_s^\dagger a_r | B \rangle \rangle .
\] (12.17)

Here \( N(q) \) denotes the Bose distribution function of the phonons.

As a result we find

\[
\sum_{nr} \langle \langle a_s^\dagger a_n \rangle \rangle \delta_{rt} - \langle \langle a_s^\dagger a_t \rangle \rangle \delta_{sn} \rangle G_{nr,lm}(\omega) =
\]

\[
\omega(q) \sum_{ij} \langle \langle a_s^\dagger a_j b_q|a_s^\dagger a_m \rangle \rangle G_{st,ji}^0(\omega) -
\]

\[
\sum_{ijn} \sum_q \left( 1 + N(q) \right) A_{-q}(j - n)e^{-iq\vec{R}_t}G_{in,lm}(\omega) -
\]

\[
A_{-q}(n - i)e^{-iq\vec{R}_n}G_{nj,lm}(\omega) \right) G_{st,ji}^0(\omega) -
\]

\[
\sum_{ij} \sum_{np} \left( A_{-q}(n - p)e^{-iq\vec{R}_n} \right) \langle \langle a_p^\dagger a_j G_{nj,lm} - \langle \langle a_p^\dagger a_j G_{ip,lm} \rangle \rangle G_{st,ji}^0(\omega)
\] (12.18)

and a similar equation for \( \langle \langle a_s^\dagger a_r b_q|a_s^\dagger a_m \rangle \rangle \).

In the above equations the Green functions \( G \) and \( G^0 \) as well as the mean values \( \langle a_s^\dagger a_j \rangle \) which can be expressed by one-particle Green functions depend on the atomic configuration. For the configuration averaging (which we will denote by \( \bar{G} \)) we use the simplest approximation

\[
\bar{G} \cdot G \sim \bar{G} \cdot \bar{G} ,
\] (12.19)

i.e. in all products the configuration-dependent quantities will be averaged separately. Taking into account Eqs. (12.14) and (12.19), the averaged zeroth-order Green function \( \bar{G}_{ij,lm}^0 \) is given by the well-known CPA solution for two-particle Green function in disordered metallic alloy \( 351 \)

\[
\bar{G}_{ij,lm}^0(\omega) = \frac{1}{N^2} \sum_{k_1 k_2} e^{i\vec{k}_1(\vec{R}_m - \vec{R}_i)} e^{i\vec{k}_2(\vec{R}_j - \vec{R}_i)} F_2(\vec{k}_1, \vec{k}_2) ,
\] (12.20)

where \( F_2(\vec{k}_1, \vec{k}_2) \) is given by

\[
F_2(\vec{k}_1, \vec{k}_2) \approx i(E(\vec{k}_2) - E(\vec{k}_1)) \int d\omega \frac{\partial f}{\partial \omega} |\text{Im}(\frac{1}{\omega - \Sigma(\omega) - E(\vec{k}_1)})|^2 ,
\] (12.21)

for \( |E(\vec{k}_1) - E(\vec{k}_2)| \ll |\Sigma(E(\vec{k}_1))| \),

\[
F_2(\vec{k}_1, \vec{k}_2) \approx \frac{f(E(\vec{k}_1)) - f(E(\vec{k}_2))}{E(\vec{k}_1) - E(\vec{k}_2)} ,
\] (12.22)

for \( |E(\vec{k}_1) - E(\vec{k}_2)| \gg |\Sigma(E(\vec{k}_1))| \).
Here $\Sigma(\omega)$ denotes the coherent potential and $f(\omega)$ is the Fermi distribution function. The configurational averaged terms $\langle a_i^\dagger a_n \rangle$ are given by

$$\langle a_i^\dagger a_n \rangle = \sum_k e^{i\vec{k}(\vec{R}_n - \vec{R}_k)} F_1(\vec{k}),$$  \hspace{1cm} (12.23)$$

$$F_1(\vec{k}) = -\frac{1}{\pi} \int d\omega f(\omega) \text{Im} \left( \frac{1}{\omega - \Sigma(\omega) - E(\vec{k})} \right).$$

After the configurational averaging equations (12.19) and (12.18) can be solved by Fourier transformation and we find

$$G_{ij,lm}(\omega) = \frac{1}{N^2} \sum_{k_1k_2k_3k_4} e^{-i\vec{k} \cdot (\vec{R}_i - \vec{R}_4)} \left[ N \sum_q e^{-i\vec{q} \cdot (\vec{R}_i - \vec{R}_4)} G(\vec{k}_1, \vec{k}_2; \vec{k}_3, \vec{k}_4) \right],$$  \hspace{1cm} (12.24)$\text{where}$

$$G(\vec{k}_1, \vec{k}_2; \vec{k}_3, \vec{k}_4) \equiv G(\vec{k}_1, \vec{k}_2) = F_2(\vec{k}_1, \vec{k}_2) \delta(\vec{k}_4, \vec{k}_3) \delta(\vec{k}_3, \vec{k}_2) - \frac{F_2(\vec{k}_1, \vec{k}_2)}{F_1(\vec{k}_1) - F_1(\vec{k}_2)}.$$  \hspace{1cm} (12.25)$\text{and}$

$$A(q, k) = \frac{1}{N} \sum_k e^{-i\vec{k} \cdot (\vec{R}_i - \vec{R}_3)} A_q(i - j).$$  \hspace{1cm} (12.26)$\text{Here the following notation were introduced}$

$$X(q, \vec{k}_2) = A(q, \vec{k}_2 - \vec{q}) A(-q, \vec{k}_2) (F_1(\vec{k}_2 - \vec{q}) - 1 - N(q)),$$  \hspace{1cm} (12.27)$\text{and}$

$$Y(q, \vec{k}_1, \vec{k}_2) = A(q, \vec{k}_2 - \vec{q}) A(-q, \vec{k}_1) (F_1(\vec{k}_1) + N(q)),$$  \hspace{1cm} (12.28)$\text{Equation (12.25) is an integral equation for the Green function } G(\vec{k}_1, \vec{k}_2) \text{ to be determined.}$

The structural averaged conductivity can be obtained, in principle, by using Eq. (12.10), where the Green function $\langle \{a_i^\dagger a_j\} \rangle$ is to be replaced by $G_{ij,lm}(\omega)$ as given by Eq. (12.24). It is, however, more convenient to start with the Kubo formula in the following form

$$\sigma = \frac{ie^2}{\Omega} \lim_{\mu \to 0} \sum_k \frac{1}{p^2} \left( \frac{\partial E(k)}{\partial k} \right)^{p} \langle \langle a_k^\dagger a_{k+p} | \eta_{-p} \rangle \rangle \varepsilon,$$  \hspace{1cm} (12.31)$\text{where } \eta_{-p} = \sum_k a_k^\dagger a_{k-p} \text{ is the electron density operator. To find the Green function } \langle \langle a_k^\dagger a_{k+p} | \eta_{-p} \rangle \rangle,$

the integral equation (12.25) has to be solved. In general, this can be done only numerically, but we can discuss here two limiting cases explicitly. At first we consider the weak-scattering limit being realized for a weak disorder in the alloy, and second, we investigate the temperature coefficient of the conductivity for a strong potential scattering.
In the weak-scattering limit the CPA Green function is given by the expression

\[ F_2(\tilde{k}_1, \tilde{k}_2) \approx i(E(\tilde{k}_2) - E(\tilde{k}_1)) \frac{df}{dE(k_1)} \cdot \frac{1}{\Sigma(E(k_1))}. \quad (12.32) \]

for \( |E(\tilde{k}_2) - E(\tilde{k}_1)| \ll |\Sigma(E(\tilde{k}_1))| \).

Corresponding to this limit the following solution ansatz for the Green function \( G(k, k + p) \) can be used

\[ G(k, k + p) = \langle \langle a_k^\dagger a_{k+p}\rangle \rangle \approx i \left( \frac{\partial E(k)}{\partial k} \right) \frac{df}{dE(k)} \cdot \frac{1}{\Sigma(E(k)) + \gamma(E(k))}, \quad (12.33) \]

where \( \gamma \) describes the contribution of the electron-phonon scattering to the coherent potential. Taking into account that in the weak-scattering limit \( |\Sigma| \ll \omega(q) \), the terms \( F_2(k, k - q) \) in the right hand side denominators of Eq.\((12.25)\) can be replaced by the expression \((12.32)\), and then the integral equation \((12.25)\) becomes for \( \lim p \to 0 \)

\[
\begin{align*}
&\frac{i}{\Sigma(E(k))} \frac{1}{N} \sum_q \frac{A(\tilde{q}, \tilde{k} - \tilde{q}) A(-\tilde{q}, \tilde{k})}{E(k) - E(k - q) - \omega(q) + i\epsilon} + \\
&\frac{Z_1(k, q) + Z_2(k, q)}{E(k) - E(k - q) + \omega(q) + i\epsilon} + \\
&2 \text{ terms with } \omega(q) \to -\omega(q), \quad N(q) \to (-1 - N(q)).
\end{align*}
\]

(12.34)

Here the following notation were introduced

\[
\begin{align*}
Z_1(k, q) &= (f(E(k - q)) - 1 - N(q)) \frac{df}{dE(k)} \cdot \frac{\left( \frac{\partial E(k)}{\partial k} \right)^p}{\Sigma(E(k)) + \gamma(E(k))}, \quad (12.35) \\
Z_2(k, q) &= (f(E(k)) + N(q)) \frac{df}{dE(k - q)} \cdot \frac{\left( \frac{\partial E(k - q)}{\partial k} \right)^p}{\Sigma(E(k - q)) + \gamma(E(k - q))}, \quad (12.36) \\
Z_3(k, q) &= (1 - f(E(k)) + N(q)) \frac{df}{dE(k)} \cdot \frac{\left( \frac{\partial E(k)}{\partial k} \right)^p}{\Sigma(E(k) + \gamma(E(k))}, \quad (12.37) \\
Z_4(k, q) &= (f(E(k - q)) + N(q)) \frac{df}{dE(k)} \cdot \frac{\left( \frac{\partial E(k)}{\partial k} \right)^p}{\Sigma(E(k))}, \quad (12.38)
\end{align*}
\]

Approximating the self-energy terms \( \Sigma(E(k)) \) and \( \gamma(E(k)) \) by \( \Sigma(E_F) \equiv \Sigma \) and \( \gamma(E_F) \equiv \gamma \), respectively, the terms proportional to \( \Sigma \) cancel and \( \gamma \) can be calculated by

\[
\gamma \frac{df}{dE(k)} \cdot \left( \frac{\partial E(k)}{\partial k} \right)^p = -\frac{\pi}{N} \sum_q A(\tilde{q}, \tilde{k} - \tilde{q}) A(-\tilde{q}, \tilde{k}) \cdot (12.39)
\]

\[
\left( \left( f(E(k)) + N(q) \right) \frac{df}{dE(k - q)} \cdot \left( \frac{\partial E(k - q)}{\partial k} \right)^p \right) \delta(E(k) - E(k - q) - \omega(q)) \left( (1 - f(E(k)) + N(q)) \frac{df}{dE(k - q)} \cdot \left( \frac{\partial E(k - q)}{\partial k} \right)^p \right) \delta(E(k) - E(k - q) + \omega(q)) - \\
\left( \left( f(E(k - q)) + N(q) \right) \frac{df}{dE(k)} \cdot \left( \frac{\partial E(k)}{\partial k} \right)^p \right) \delta(E(k) - E(k - q) + \omega(q)), \quad (12.39)
\]

82
Using the approximations
\[ \frac{\partial E(k)}{\partial k} \simeq \frac{1}{m^*} k, \quad A(q, \vec{k} - \vec{q}) A(-\vec{q}, \vec{k}) \simeq A^2 q, \quad q \to 0, \]
where effective mass \( m^* = m^*(E_F) \), we find
\[ \gamma = \beta \frac{\Omega}{2 \pi N} \frac{A^2 m^*}{2(2m^* E_F)^{3/2}} \int dq q \omega(q) N(q) \left( 1 + N(q) \right) \tag{12.40} \]
and
\[ \gamma \sim \begin{cases} T^5 & \text{if } T \ll \theta_D, \\ T & \text{if } T \gg \theta_D. \end{cases} \tag{12.41} \]
For a binary alloy \( A_x B_{1-x} \) with concentrations of the constituents \( c_A \) and \( c_B \) and the corresponding atomic energies \( \epsilon_A \) and \( \epsilon_B \), in the weak-scattering limit the coherent potential is given by \( \Sigma \simeq c_A c_B(\epsilon_A - \epsilon_B)^2 D(E_F). \) \( \tag{12.42} \)
Then the conductivity becomes
\[ \sigma = \frac{e^2}{3(2\pi)^3} \int dk \left( \frac{\partial E(k)}{\partial k} \right)^2 \frac{df}{dE(k)} \cdot \tau, \tag{12.43} \]
where
\[ \tau^{-1} = \Sigma + \gamma, \tag{12.44} \]
in correspondence with the Matthiessen, Nordheim and Bloch-Gruneisen rules. Now we estimate temperature coefficient of the conductivity for a strong potential scattering. For a strongly disordered alloy the electron-phonon interaction can be considered as a small perturbation and the Green functions \( G(k, k') \) on the right hand side of equation (12.25) can be replaced by CPA Green functions \( F(k, k') \). For simplicity, on the right hand side of equation (12.25) we take into consideration only terms proportional to the Bose distribution function giving the main contribution to the temperature dependence of the conductivity. Then \( \langle a_k^\dagger a_k | \eta_{k-k_1} \rangle \) becomes \( (k_1 = k + p \approx k) \)
\[ \langle a_k^\dagger a_k | \eta_{k-k_1} \rangle = F_2(k, k_1) \left( 1 \right. \right. \]
\[ \left. \left. - \frac{2}{F_1(k) - F_1(k_1)} \sum_q A(q, \vec{k} - \vec{q}) A(-\vec{q}, \vec{k}) N(q) \cdot \right. \]
\[ \left. - \frac{F_2(k, k - q)[F_2(k - q, k_1 - q) - F_2(k, k_1)](F_1(k) - F_1(k - q))}{[F_1(k) - F_1(k - q)]^2 - \omega^2(q) F_2^2(k, k - q)} \right) \]
\[ + \frac{F_2(k - q, k)[F_2(k - q, k_1 - q) - F_2(k, k_1)](F_1(k) - F_1(k))}{[F_1(k - q) - F_1(k)]^2 - \omega^2(q) F_2^2(k - q, k)} \right). \tag{12.45} \]
Neglecting at low temperatures the terms \( \omega^2(q) F_2^2(k_1 - q, k_1) \sim q^4 \) as compared to \( [F_1(k_1) - F_1(k - q)]^2 \sim q^2 \) and using equation (12.21) for \( \omega(q) \ll |\Sigma| \), we find for small \( q \) and \( p \to 0 \)
\[ \langle a_k^\dagger a_{k+p} | \eta_{-p} \rangle \simeq F_2(k, k + p) \left[ 1 + \left( \frac{dF_1}{dE(k)} \right)^{-2} \sum_q (\Delta(q, k - q) - \Delta(q, k)) \right]. \tag{12.46} \]
Here \( \Delta(q, k) \) is the temperature-dependent correction terms to the CPA Green function are given by
\[ \Delta(q, k) = 2A^2 q N(q) \left( \int d\omega \frac{df(\omega)}{d\omega} \left| \text{Im} \left( \frac{1}{\omega - \Sigma(\omega) - E(k)} \right) \right|^2 \right)^2. \tag{12.47} \]
For temperatures $k_B T \ll E_F$ we can write
\[
\int d\omega \frac{df(\omega)}{d\omega} S(\omega, E(k)) \approx -S(E_F, E(k))
\] (12.48)
and the conductivity becomes
\[
\sigma = \sigma_{CPA} + \Delta \sigma(T),
\] (12.49)
where
\[
\sigma_{CPA} = \frac{e^2}{\Omega} \sum_k \left( \frac{\partial E(k)}{\partial k} \right)^2 \left[ \text{Im} \left( \frac{1}{E_F - \Sigma(E_F) - E(k)} \right) \right]^2
\] (12.50)
is the standard CPA expression for the conductivity and
\[
\Delta \sigma(T) = \frac{2e^2 A^2}{\Omega} \sum_k \left( \frac{\partial E(k)}{\partial k} \right)^2 \sum_q q^2 N(q) \cdot \left( \text{Im} \left( \frac{1}{E_F - \Sigma(E_F) - E(k - q)} \right) \right)^4 - \left( \text{Im} \left( \frac{1}{E_F - \Sigma(E_F) - E(k)} \right) \right)^4.
\] (12.51)
Introducing the effective mass of the electrons with $E(k) \simeq E_F$, the temperature-dependent correction to the conductivity becomes
\[
\Delta \sigma(T) \approx \frac{2e^2 A^2}{\Omega} \frac{1}{(m^*)^2} \sum_k \sum_q q^2 N(q) \left[ \text{Im} \left( \frac{1}{E_F - \Sigma(E_F) - E(k - q)} \right) \right]^4.
\] (12.52)
Here the quantity $\Delta \sigma(T)$ is positive definite and increasing with increasing temperature. Hence, in strongly disordered alloys where the electron-phonon scattering is weak as compared with the disorder scattering the temperature coefficient of the resistivity is negative. It should be mentioned, however, that the concrete temperature dependence of the correction term \[12.52\] is a crude estimation only because in the derivation of \[12.52\] the influence of the disorder on the lattice vibrations has been neglected.

One more remark is appropriate for the above consideration. For the calculation of transport coefficients in disordered $3d$ systems the classical approaches as the Boltzmann equation become useless if the random fluctuations of the potential are too large.\cite{235,337} The strong potential fluctuations force the electrons into localized states. In order to investigate the resistivity of metallic alloys near the metal-insulator transition\cite{235,337}, the corresponding formula for the resistivity can be deduced along the line described above. For a binary transition metal alloy the corresponding Hamiltonian is given by
\[
H = \sum_i \epsilon_i a_i^\dagger a_i + \sum_{ij} t_{ij} a_i^\dagger a_j
\] (12.53)
(with $\epsilon_i = \epsilon_A, \epsilon_B$ depending on the occupation of the lattice site $i$). A corresponding integral equation for the Green function $\langle \langle a_j a_i^\dagger \rangle \rangle_\omega$ can be written down. Using a simple ensemble averaging procedure and approximating the averaged Green function by the expression
\[
\langle \langle a_j a_i^\dagger \rangle \rangle_\omega \approx \frac{1}{N} \sum_k \exp[ik(R_i - R_j)] \frac{1}{\omega - \epsilon_k} \exp(- \alpha(\epsilon_k)|R_i - R_j|),
\] (12.54)
the integral equation transforms into an equation for the parameter $\alpha(\epsilon_k)^{-1}$ which is proportional to the averaged mean free path of the electrons. It can be shown then, by solving this equation for electrons at the Fermi surface $E_F$, that $\alpha(E_F)^{-1}$ and the conductivity $\sigma$ drop in a discontinued way from $\sigma_{min}$ and $\alpha_{min}$, respectively, to zero as the potential fluctuations exceed a critical value. Note that $\alpha_{min}$ is of the order $1/d$, where $d$ is the lattice parameter.
13 Discussion

In the foregoing sections we have discussed some selected statistical mechanics approaches to the calculation of the electrical conductivity in metallic systems like transition metals and their disordered alloys within a model approach. Electrons in metals are scattered by impurities and phonons. The theory of transport processes for ordinary metals was based on the consideration of various types of scattering mechanisms and, as a rule, has used the Boltzmann equation approach. The aim of the present review was to describe an alternative approach to the calculation of electroconductivity, which can be suitable for transition metals and their disordered alloys. There is an important aspect of this consideration. The approximations used here are the tight-binding and modified tight-binding, which are admittedly not ideally precise but does give (at least as the first approximation) reasonable qualitative results for paramagnetic transition metals and their disordered alloys. We studied the electronic conduction in a model of transition metals and their disordered alloys utilizing the method of generalized kinetic equations. The reasonable and workable expressions for the electrical conductivity were established and analyzed. We discussed briefly various approaches for computing electrical conductivity as well.

We hope that these considerations have been done with sufficient details to bring out their scope and workability. In this paper, we have considered the idealized Hubbard model which is the simplest (in the sense of formulation, but not solution) and most popular model of correlated lattice fermions. We believe that this technique can be applied to other model systems (e.g. multi-band Hubbard model, periodic Anderson model, etc.). As it is seen, this treatment has some advantages in comparison with the standard methods of computing electrical conductivity within the Boltzmann equation approach, namely, the very compact form. The physical picture of electron-electron and electron-phonon scattering processes in the interacting many-particle systems is clearly seen at every stage of calculations, which is not the case with the standard methods. This picture of interacting many-particle system on a lattice is far richer and gives more possibilities for the analysis of phenomena which can actually take place in real metallic systems. We believe that our approach offers a convenient way for approximate considerations of the resistivity of the correlated electron systems on a lattice. We believe that in view of the great difficulty of developing a first-principles microscopic theory of transport processes in solids, the present approach is a useful alternative for description the influence of electron-electron, electron-phonon and disorder scattering effects on the transport properties of transition metals and their disordered alloys.

In the confines of a review of this nature many topics of great practical and theoretical interest have necessarily to be omitted (see e.g. Refs. 369–371). In recent years the field of mesoscopic physics is developed rapidly. It deals with systems under experimental conditions where several quantum length scales for electrons are comparable. The physics of transport processes in such systems is rich of quantum effects, which is typically characterized by interplay of quantum interference and many-body interactions. It would be of interest to generalize the present approach to quantum transport phenomena.

In conclusion, the foregoing analysis suggests that the method of the generalized kinetic equations is an efficient and useful formalism for the studying of some selected transport processes in metallic systems.

References

[1] N. F. Mott and H. Jones, The Theory of the Properties of Metals and Alloys (Dover, New York, 1964).
[2] J. M. Ziman, *Electrons and Phonons. The Theory of Transport Phenomena in Solids* (Oxford University Press, Oxford, 1960).

[3] L. Solimar and D. Walsh, *Electrical Properties of Materials.* (Oxford University Press, Oxford, 2004).

[4] Terry M. Tritt, (ed.), *Thermal Conductivity: Theory, Properties, and Applications.* (Kluwer Academic/Plenum Publishers, New York, 2004).

[5] Feng Duan and Jin Guojun, *Introduction to Condensed Matter Physics, Vol.1,* (World Scientific, Singapore, 2005).

[6] H. J. Goldsmid, *Introduction to Thermoelectricity* (Springer-Verlag, Berlin, 2010).

[7] S. Maekawa, (ed.), *Concepts in Spin Electronics.* (Oxford University Press, Oxford, 2006).

[8] T. Dietl, D. Awschalom, M. Kaminska and H. Ohno, (eds.), *Spintronics.* (Semiconductors and Semimetals, Vol.82) (Elsevier Inc., Amsterdam, 2008).

[9] T. L. Olsen, *Electron Transport in Metals* (Pergamon Press, Oxford, 1962).

[10] A. B. Pippard, *Dynamics of Conduction Electrons* (Blackie and Son, London, 1965).

[11] G. T. Meaden, *Electrical Resistance of Metals* (Plenum Press, New York, 1965).

[12] A. C. Smith, J. F. Janak and R. B. Adler, *Electronic Conduction in Solids* (McGraw-Hill Book Co., New York, 1967).

[13] J. C. Slater, *Insulators, Semiconductors and Metals* (McGraw-Hill Book Co., New York, 1967).

[14] F. J. Blatt, *Physics of Electronic Conduction in Solids* (McGraw-Hill Book Co., New York, 1968).

[15] J. M. Ziman, (ed.), *The Physics of Metals: Volume 1, Electrons* (Cambridge University Press, Cambridge, 1969).

[16] W. B. Pearson, *The Crystal Chemistry and Physics of Metals* (John Wiley and Sons, New York, 1972).

[17] R. Berman, *Thermal Conduction in Solids* (Oxford University Press, Oxford, 1976).

[18] P. L. Rossiter, *The Electrical Resistivity of Metals and Alloys* (Cambridge University Press, Cambridge, 1987).

[19] H. Smith and H. H. Jensen, *Transport Phenomena* (Clarendon Press, Oxford, 1989).

[20] C. P. Enz, *A Course on Many-Body Theory Applied to Solid-State Physics* (World Scientific, Singapore, 1992).

[21] R. E. Hummel, *Electronic Properties of Materials* (Springer-Verlag, Berlin, 1993).

[22] J. S. Dugdale, *The Electrical Properties of Disordered Metals* (Cambridge University Press, Cambridge, 1995).

[23] M. Lundstrom, *Fundamentals of Carrier Transport* (Cambridge University Press, Cambridge, 2000).
[24] J. I. Gerstein and F. W. Smith, *The Physics and Chemistry of Materials* (John Wiley and Sons, New York, 2001).

[25] U. Mizutani, *Introduction to the Electron Theory of Metals* (Cambridge University Press, Cambridge, 2001).

[26] M. Di Ventra, *Electrical Transport in Nanoscale Systems* (Cambridge University Press, Cambridge, 2008).

[27] A. Sommerfeld and N. H. Frank, *Rev. Mod. Phys.* 3, 1 (1931).

[28] N. Mott, *Proc. Phys. Soc.* 47, 571 (1935).

[29] J. Bardeen, *Phys. Rev.* 52, 688 (1937).

[30] R. Peierls, *Ann. Physik (Leipzig)* 3, 1055 (1929).

[31] D. I. Blochinzev and L. Nordheim, *Z. Physik* 84, 168 (1933).

[32] A. P. Sutton, *Electronic Structure of Materials* (Oxford University Press, Oxford, 1993).

[33] W. A. Harrison, *Elementary Electronic Structure* (World Scientific, Singapore, 1999).

[34] J. Singleton, *Band Theory and Electronic Properties of Solids* (Oxford University Press, Oxford, 2001).

[35] R. M. Martin, *Electronic Structure: Basic Theory and Practical Methods* (Cambridge University Press, Cambridge, 2004).

[36] S. Fujita and K. Ito, *Quantum Theory of Conducting Matter* (Springer-Verlag, Berlin, 2007).

[37] X. L. Lei, *Balance Equation Approach to Electron Transport in Semiconductors* (World Scientific, Singapore, 2008).

[38] C. Jacoboni, *Theory of Electron Transport in Semiconductors. A Pathway from Elementary Physics to Nonequilibrium Green Functions* (Springer-Verlag, Berlin, 2010).

[39] M. Prasad, *J. Physics C:Solid State Phys.* C 13, 3239 (1980).

[40] D. C. Mattis, *J. Stat. Phys.* 77, 383 (1994).

[41] A. Kundu, A. Dhar and O. Narayan, *J. Stat. Mech.: Theory and Experiment*, L03001 (2009).

[42] D. N. Zubarev, *Nonequilibrium Statistical Thermodynamics* (Consultant Bureau, New York, 1974).

[43] D. N. Zubarev, *Reviews of Science and Technology: Modern Problems of Mathematics, Vol.15*, pp. 131-226, ed. by R. B. Gamkrelidze, (Izd.Nauka, Moscow, 1980)(in Russian) [English Transl.: *J. Soviet Math.* 16, 1509 (1981)].

[44] R. L. Liboff, *Kinetic Theory: Classical, Quantum, and Relativistic Description*, 3rd edn. (Springer-Verlag, Berlin, 2003).

[45] A. L. Kuzemsky, *Int. J. Mod. Phys.* B 21, 2821 (2007).

[46] C. M. Van Vliet, *Equilibrium and Non-equilibrium Statistical Mechanics* (World Scientific, Singapore, 2008).
[47] A. L. Kuzemsky and I. G. Kuzemskaya, The Structural, Superconducting and Transport Properties of the Mercurocuprates and Other Layered Systems. in Studies in High-T$_c$ Superconductors. Vol.44, ed. A. Narlikar. (Nova Science Publ., New York, 2003), P.1-80.

[48] A. L. Kuzemsky, J. Low Temp. Phys. 143, 213 (2006).

[49] N. V. Sidgwick, The Chemical Elements and their Compounds (Clarendon Press, Oxford, 1950).

[50] P. W. Anderson, Concepts in Solids (World Scientific, Singapore, 1997).

[51] A. L. Kuzemsky, Physics of Particles and Nuclei 39, 137 (2008).

[52] J. Callaway, Quantum Theory of the Solid State, 2nd ed. (Academic Press, New York, 1991).

[53] N. Mott, Adv.Phys. 13, 325 (1964).

[54] J. Friedel, J. Physics F:Metal Phys. F 3, 785 (1973).

[55] F. Zeitz, The Modern Theory of Solids (McGraw-Hill Book, New York, 1940).

[56] C. Kittel, Introduction to Solid State Physics, 7th edn. (John Wiley and Sons, New York, 1996).

[57] Semiconductors. Electronic, Transport, Optical and Other Properties (Landolt-Bornstein: Numerical Data and Functional Relationships in Science and Technology - New Series / Condensed Matter) Edited by U. Rossler, S. Adachi, R. Blachnik and R.P. Devaty, (Springer, Berlin, 2002).

[58] P. Y. Yu and M. Cardona, Fundamentals of Semiconductors: Physics and Materials Properties (Springer, Berlin, 2010).

[59] R. E. Watson and M. Weinert, Transition-Metals and Their Alloys, in Solid State Physics, eds. H. Ehrenreich and F. Spaepen, Vol.56 (Academic Press, New York, 2001) p.1.

[60] 3d, 4d, and 5d Elements, Alloys and Compounds. (Landolt-Bornstein: Numerical Data and Functional Relationships in Science and Technology - New Series / Condensed Matter, Volume 19) Edited by K. Adachi, D. Bonneberg, J.J.M. Franse and R. Gersdorf, (Springer, Berlin, 1986).

[61] Band Structures and Fermi Surfaces of Metallic Compounds and Disordered Alloys. Phonon States of Elements. Electron States and Fermi Surfaces of Alloys (Landolt-Bornstein: Numerical Data and Functional Relationships in Science and Technology - Group III Condensed Matter, Volume 13a) Edited by K.-H. Hellwege and J. L. Olsen, (Springer, Berlin, 1983).

[62] A. L. Kuzemsky, Physics of Particles and Nuclei, 40, 949 (2009).

[63] O. K. Andersen, O. Jepsen and D. Glotzel, Canonical Description of the Band Structure of Metals, in Highlights of Condensed Matter Theory (Societa Italiana di Fizica, Bologna, 1985), Vol. 89, p. 59.

[64] L. F. Mattheiss, Phys.Rev. B 1, 373 (1970).

[65] L. Hodges, R. E. Watson and H. Ehrenreich, Phys.Rev. B 5, 3993 (1972).

[66] D. G. Laurent, C. S. Wang and J. Callaway, Phys.Rev. B 17, 455 (1978).
[67] A. Goldmann, W. Gudat, and O. Rader. Electronic Structure of Solids: Photoemission Spectra and Related Data: Noble Metals, Noble Metal Halides and Nonmagnetic Transition Metals (Landolt-Bornstein: Numerical Data and Functional Relationships in Science and Technology - New Series III/ Condensed Matter, Volume 23) Edited by A. Goldmann, (Springer, Berlin, 2003).

[68] N. F. Mott, Metal-Insulator Transitions, 2nd ed. (Taylor and Francis, London, 1990).

[69] N. Wiser, Can. J. Phys. 60, 693 (1982).

[70] W. Kohn, Phys.Rev. 133, A171 (1964).

[71] P. P. Edwards, R. L. Johnston, F. Hensel, C. N. R. Rao, and D. P. Tunstall, A Perspective on the Metal-Nonmetal Transition, in Solid State Physics, eds. H. Ehrenreich and F. Spaepen, Vol.52 (Academic Press, New York, 1999) p.229.

[72] R. Resta J. Phys.:Condens. Matter 14, R625 (2002).

[73] A. B. Kaiser, in Physics of Novel Materials, ed. M. P. Das (World Scientific, Singapore, 1998) p.290.

[74] G. B. Bokij, Crystallography Reports 47, 452 (2002).

[75] C.P. Poole Jr., H. A. Farach, R. J. Creswick and R. Prozorov, Superconductivity, 2nd ed. (Academic Press, New York, 2007).

[76] P. Lunkenheimer, S. Krohns, S. Riegg, S.G. Ebbinghaus, A. Reller, and A. Loidl, Eur. Phys. J. Special Topics 180, 61 (2010).

[77] N. N. Bogoliubov and N. N. Bogoliubov, Jr., Introduction to Quantum Statistical Mechanics, 2nd ed. (World Scientific, Singapore, 2009).

[78] D. Ya. Petrina, Mathematical Foundations of Quantum Statistical Mechanics (Kluwer Academic Publ., Dordrecht, 1995).

[79] D. ter Haar, Lectures on Selected Topics in Statistical Mechanics, (Pergamon Press, Oxford, 1977).

[80] A. L. Kuzemsky, Int. J. Mod. Phys. B 24, 835 (2010).

[81] A. S. Davydov, Quantum Mechanics (Pergamon Press, Oxford, 1965).

[82] P. Nozieres, Theory of Interacting Fermi Systems (W. A. Benjamin, Inc., New York, 1964).

[83] I. M. Kim and M. H. Lee, Phys.Rev. B 24, 3961 (1981).

[84] S. Raims, Many-Electron Theory (North-Holland, Amsterdam, New York, 1972).

[85] J. W. Negele and H. Orland, Quantum Many-Particle Systems (Addison-Wesley, New York, 1988).

[86] G. Grimvall, The Electron-Phonon Interaction in Metals (North-Holland, Amsterdam, New York, 1982).

[87] G. V. Chester, Adv.Phys. 10, 357 (1961).

[88] W. A. Harrison, Physica 55, 696 (1971).
[89] C. Kittel, *Quantum Theory of Solids*, 2nd ed. (John Wiley and Sons, New York, 1987).

[90] J. Korringa, *Physica* **13**, 392 (1947).

[91] E. E. Lafon and C. C. Lin, *Phys.Rev.* **152**, 579 (1966).

[92] J. M. Tyler, T. E. Norwood and J. L. Fry, *Phys.Rev.* B **1**, 297 (1970).

[93] J. Callaway and J. L. Fry, Towards self-consistency with the tight binding approximation, in *Computational Methods in Band Theory* eds. P. M. Marcus, J. F. Janak and A. R. Williams (Plenum Press, N.Y., 1971), p. 512.

[94] C. M. Goringe, D. R. Bowler and E. Hernandez, *Rep. Prog. Phys.* **60**, 1447 (1997).

[95] G. Wannier, *Phys.Rev.* **52**, 191 (1937).

[96] E. Brown, *Phys.Rev.* **166**, 626 (1968).

[97] W. Kohn, *Phys.Rev.* B **7**, 4388 (1973).

[98] L. G. Ferreira and N. J. Parada, *Phys.Rev.* B **2**, 1614 (1970).

[99] A. L. Kuzemsky, *Rivista del Nuovo Cimento* **25**, 1 (2002).

[100] J. Hubbard, *Proc. Roy. Soc.* A **276**, 238 (1963).

[101] N. Ohata and R. Kubo, *J. Phys. Soc. Jpn.* **28**, 1402 (1970).

[102] R. A. Bari, D. Adler and R. V. Lange, *Phys.Rev.* B **2**, 2898 (1970).

[103] K. Kubo, *J. Phys. Soc. Jpn.* **31**, 30 (1971).

[104] M. L. Malwah and R. W. Bene, *Phys.Rev.* B **6**, 3305 (1972).

[105] R. A. Bari and T. A. Kaplan, *Phys.Rev.* B **6**, 4623 (1972).

[106] I. Sadakata and E. Hanamura, *J. Phys. Soc. Jpn.* **34**, 882 (1973).

[107] D. Cabib and T. A. Kaplan, *phys. stat. sol.* b **58**, 85 (1973).

[108] M. Eswaran and J. C. Kimball, *Phys.Rev.* B **11**, 3420 (1975).

[109] E. Marsch, *phys. stat. sol.* b **72**, K103 (1975).

[110] E. Marsch, *Z. Physik* B **25**, 83 (1976).

[111] W. Gasser and J. Schneider, *phys. stat. sol.* b **77**, K103 (1976).

[112] D. Cabib, *Phys.Rev.* B **12**, 2189 (1975).

[113] P. F. Maldague, *Phys.Rev.* B **16**, 2437 (1977).

[114] K. Michielsen and H. De Raedt, *Mod. Phys. Lett.* B **10**, 855 (1996).

[115] J. C. Slater, *Electronic Structure of Molecules* (McGraw-Hill Book Co., New York, 1963).

[116] M. Baer, *Beyond Born-Oppenheimer. Conical Intersections and Electronic Nonadiabatic Coupling Terms* (John Wiley and Sons, New York, 2006).
[117] H. Fröhlich, *Adv.Phys.* **3**, 325 (1954).

[118] J. A. Moriarty, *J.Physics F:Metal Phys.* **F 5**, 873 (1975).

[119] W. H. Butler, J. J. Olson, J. S. Faulkner and B. L. Gyorffy, *Phys. Rev.* B **14**, 3823 (1976).

[120] W. H. Butler, *Phys. Rev.* B **15**, 5267 (1977).

[121] A.L. Simons, C. M. Varma and W. Weber, *Phys. Rev.* B **23**, 2431 (1981).

[122] H. Fröhlich and T. K. Mitra, *J.Physics C:Solid State Phys.* C **1**, 548 (1968).

[123] T. K. Mitra, *J.Physics C:Solid State Phys.* C **2**, 52 (1969).

[124] T. K. Mitra, *J.Physics C:Solid State Phys.* C **2**, 1357 (1969).

[125] S. Barisic, J. Labbe and J. Friedel, *Phys.Rev.Lett.* **25**, 919 (1970).

[126] A. L. Kuzemsky, A. Holas and N. M. Plakida, *Physica* B **122**, 168 (1983).

[127] A. L. Kuzemsky and A. P. Zhernov, *Int. J. Mod. Phys.* B **4**, 1395 (1990).

[128] J. C. Slater, *Phys.Rev.* **36**, 57 (1930).

[129] R. A. Deegan, *Phys. Rev.* B **4**, 1389 (1971).

[130] R. A. Deegan, *Phys. Rev.* B **5**, 1183 (1972).

[131] J. Ashkenazi, M. Dacorogna and M. Peter, *Solid State Comm.* **29**, 181 (1979).

[132] J. Ashkenazi and M. Dacorogna, Phonons and Electron-phonon Coupling in Metals, in *Recent Developments in Condensed Matter Physics*, (eds.) J. Devreese et al., Vol.34 (Plenum Press, New York, 1981) p.15.

[133] For a discussion of the P. Drude works and its modern development see the memorial volume *Annalen der Physik*, **15**, Issues 7-8 (2006).

[134] D.R. Lide (ed.) *CRC Handbook of Chemistry and Physics* (CRC Press, London, 2005).

[135] W. Martienssen and H. Warlimont, *Handbook of Condensed Matter and Materials Data* (Springer, Berlin, 2005).

[136] J. Bass and K.H. Fischer, Electrical resistivity of pure metals and dilute alloys. *Metals: Electronic Transport Phenomena*. (Landolt-Bornstein: Numerical Data and Functional Relationships in Science and Technology - New Series III/ Condensed Matter, Volume 15a) Edited by K.-H. Hellwege, J. L. Olsen, (Springer, Berlin, 1982).

[137] J. Bass, J. S. Dugdale, C. L. Foiles, and A. Myers, *Metals: Electronic Transport Phenomena*. (Landolt-Bornstein: Numerical Data and Functional Relationships in Science and Technology - New Series III/ Condensed Matter, Volume 15b) Edited by K.-H. Hellwege, J. L. Olsen, (Springer, Berlin, 1985).

[138] H. Fröhlich, B. V. Paranjape, C. G. Kuper and S. Nakajima, *Proc. Phys. Soc.* A **69**, 842 (1956).

[139] J. Bass, *Adv.Phys.* **21**, 431 (1972).
[140] M. R. Cimberle, G. Bobel, and C. Rizzutto, Adv.Phys. 23, 639 (1974).
[141] W. V. Houston, Phys. Rev. 34, 279 (1929).
[142] F. Bloch, Z. Physik 59, 208 (1930).
[143] E. Gruneisen, Ann. Physik (Leipzig) 16, 530 (1933).
[144] M. Kavech and N. Wiser, Adv.Phys. 33, 257 (1984).
[145] J. Bass, W. P. Pratt and P. A. Schroeder, Rev. Mod. Phys. 62, 645 (1990).
[146] P. F. Maldague and C. A. Kukkonen, Phys. Rev. B 19, 6172 (1979).
[147] A. H. MacDonald, Phys.Rev.Lett. 44, 489 (1980).
[148] J. M. B. Lopes dos Santos and D. Sherrington, J.Physics F:Metal Phys. F 13, 1233 (1983).
[149] M. J. Rice, Phys.Rev.Lett. 20, 1439 (1968).
[150] J. E. Black, Can. J. Phys. 56, 708 (1978).
[151] C. Potter and G. J. Morgan, J.Physics F:Metal Phys. F 9, 493 (1979).
[152] W. H. Butler, Can. J. Phys. 60, 735 (1982).
[153] F. J. Pinski, P. B. Allen and W. H. Butler Phys. Rev. B 23, 5080 (1981).
[154] N. Mott, Proc. Roy. Soc. A 153, 699 (1936).
[155] G. W. Webb, Phys. Rev. 181, 1127 (1969).
[156] R. Fogelholm and O. Rapp, J. Physics F:Metal Phys. F 7, 667 (1977).
[157] S. M. Sharma, Phys.Rev. B 20, 1514 (1979).
[158] S. M. Sharma, J. Physics F:Metal Phys. F 10, L47 (1980).
[159] S. M. Sharma, J. Physics F:Metal Phys. F 11, 2367 (1981).
[160] D. K. Wagner, J. C. Garland and R. Bowers, Phys. Rev. B 3, 3141 (1971).
[161] E. L. Stone, M. D. Ewbank, J. Bass and W. P. Pratt, Phys.Lett. A 59, 168 (1976).
[162] G. J. Gautron, J. E. Zablocki, T. Y. Hsiang and H. Weinstock, J. Low Temp. Phys. 49, 185 (1982).
[163] A. Takase and T. Kasuya, J. Phys. Soc. Jpn. 48, 430 (1980).
[164] L. M. Falicov and B. Koiller, J. Solid State Chem. 12, 349 (1975).
[165] H. L. Frisch and J. L. Lebowitz, Phys. Rev. 123, 1542 (1961).
[166] D. Greig and G. J. Morgan, Phil. Mag. 27, 929 (1973).
[167] W. M. Visscher, Phys.Rev. B 17, 598 (1978).
[168] J. H. Mooij, phys. stat. sol. a 17, 521 (1973).
[169] S. P. Repetsky and T. D. Shatnii, *Theor. Math. Phys.* **131**, 832 (2002).

[170] S. P. Repetsky and T. D. Shatnii, *Ukr. J. Phys.* **48**, 853 (2003).

[171] I. G. Vyshivana, S. P. Repetsky and I. V. Shakhov, *Ukr. J. Phys.* **50**, 273 (2005).

[172] S. P. Repetsky and I. G. Vyshivana, *The Physics of Metals and Metallography* **99**, 558 (2005).

[173] F. J. Ohkawa, *J. Phys. Soc. Jpn.* **44**, 1105 (1978).

[174] P. L. Taylor, *Phys.Rev.* **135**, A1333 (1964).

[175] J. Yamashita and H. Hayakawa, *Prog. Theor. Phys.* **56**, 361 (1976).

[176] R. K. Williams and F. J. Weaver, *Phys.Rev.* B **25**, 3663 (1982).

[177] H. L. Engquist and G. Grimvall, *Phys.Rev.* B **21**, 2072 (1980).

[178] A. Bergmann, M. Kaveh, and N. Wiser, *J. Physics F:Metal Phys.* F **12**, 2985 (1982).

[179] J. Inoue and M. Shimizu, *J. Phys. Soc. Jpn.* **41**, 1211 (1976).

[180] K. I. Wysokinski and A. L. Kuzemsky, *phys. stat. sol. b* **113**, 409 (1982).

[181] P. S. Riseborough, *Phys.Rev.* B **29**, 4134 (1984).

[182] C. M. Hurd, *The Hall Effect in Metals and Alloys* (Plenum Press, New York, 1972).

[183] C. L. Chien and C. R. Westgate (eds.), *The Hall Effect and Its Applications* (Plenum Press, New York, 1980).

[184] M. Dresden, *Rev. Mod. Phys.* **33**, 265 (1961).

[185] P. N. Argyres, *Phys.Rev.* **117**, 315 (1960).

[186] S. Simons, *Proc. Phys. Soc.* **78**, 316 (1961).

[187] G. H. Wannier, *Rev. Mod. Phys.* **34**, 645 (1962).

[188] A. B. Pippard, *Proc. Roy. Soc. A* **305**, 291 (1968).

[189] J. M. Luttinger, *Phys.Rev.* **84**, 814 (1951).

[190] H. F. Budd, *Phys.Rev.* **175**, 241 (1968).

[191] G. H. Wannier, *Phys.Rev.* B **5**, 3836 (1972).

[192] F. Y. Yang, Kai Liu, Kimin Hong, D. H. Reich, P. C. Searson and C. L. Chien, *Science* **284**, 1335 (1999).

[193] M. B. Fontes, J. C. Trochez, B. Giordanengo, S. L. Budko, D. R. Sanchez, E. M. Baggio-Saitovich and M. A. Continentino, *Phys.Rev.* B **60**, 6781 (1999).

[194] N. Manyala, Y. Sidis, J. F. DiTusa, G. Aeppli, D. P. Yang and Z. Fisk, *Nature* **404**, 581 (2000).

[195] M. Sawicki, T. Dietl and J. Kossut, *Acta Phys. Pol. A* **67**, 399 (1985).
[196] J. Jaroszynski and T. Dietl, *Acta Phys. Pol.* A **69**, 1017 (1986).

[197] S. Majumdar, R. Malik, E. V. Sampathkumaran, K. Ruprecht and G. Wortmann, *Phys.Rev.* B **60**, 6770 (1999).

[198] H. Yamada and S. Takada, *Prog. Theor. Phys.* **48**, 1828 (1972).

[199] G. D. Mahan, *Phys.Rep.* **145**, 251 (1987).

[200] Y. Toyozawa, *J. Phys. Soc. Jpn.* **17**, 986 (1962).

[201] C. C. Chen and S. Fujita, *J. Phys. Chem. Solids* **28**, 607 (1967).

[202] C. A. Kukkonen and P. F. Maldague, *Phys.Rev.* B **19**, 2394 (1999).

[203] P. Carruthers, *Rev. Mod. Phys.* **33**, 92 (1961).

[204] P. G. Klemens, G. Neuer, B. Sundqvist, C. Uher, and G. K. White, *Metals: Electronic Transport Phenomena. Thermal Conductivity of Pure Metals and Alloys* (Landolt-Bornstein: Numerical Data and Functional Relationships in Science and Technology - New Series III/Condensed Matter, Volume 15c) Edited by O. Madelung and G. K. White, (Springer, Berlin, 1985).

[205] R. W. Powell, *Contemp. Phys.* **10**, 579 (1969).

[206] G. A. Slack, The Thermal Conductivity of Nonmetallic Crystals, in *Solid State Physics*, eds. F. Seitz and D. Turnbull, Vol.34 (Academic Press, New York, 1979) p.1.

[207] N. F. Mott, *Proc. Roy. Soc.* A **156**, 368 (1936).

[208] E. L. Nagaev, *Physics of Metals and Metallography* **25**, 177 (1968).

[209] S. R. de Groot and P. Mazur, *Non-Equilibrium Thermodynamics* (North-Holland, Amsterdam, New York, 1962).

[210] W. A. Wooster, *Tensors and Group Theory for the Physical Properties of Crystals* (Clarendon Press, Oxford, 1973).

[211] D. N. Zubarev, *Doklady Academii Nauk SSSR* **140**, 92 (1961).

[212] D. N. Zubarev, *Teor. Mat. Fiz.* **3**, 276 (1970).

[213] M. Toda, R. Kubo and N. Saito, *Statistical Physics: Equilibrium Statistical Mechanics*, Vol.1, (Springer, Berlin, 1992).

[214] R. Kubo, M. Toda and N. Hashitsume, *Statistical Physics: Nonequilibrium Statistical Mechanics*, Vol.2, (Springer, Berlin, 1991).

[215] J. A. McLennan, *Introduction to Nonequilibrium Statistical Mechanics*, (Prentice Hall, New Jersey, 1989).

[216] M. Ichiyama, *Phys. Rep.* **243**, 125 (1994).

[217] R. Zwanzig, *Nonequilibrium Statistical Mechanics*, (Oxford University Press, Oxford, New York, 2001).
[218] R. Luzzi, A. R. Vasconcellos and J. G. Ramos, *Statistical Foundations of Irreversible Thermodynamics* (Teubner-Bertelsmann Springer, Stuttgart, 2000).

[219] R. Luzzi, A. R. Vasconcellos and J. G. Ramos, *Predictive Statistical Mechanics* (Kluwer Publ., Boston, London, 2002).

[220] R. Luzzi, A. R. Vasconcellos and J. G. Ramos, *Int. J. Mod. Phys.* B 14, 3189 (2000).

[221] U. Balucani, M. H. Lee and V. Tognetti, *Phys. Rep.* 373, 409 (2003).

[222] S. Fujita, *Introduction to Nonequilibrium Quantum Statistical Mechanics* (Saunders, Philadelphia, 1966).

[223] M. H. Lee, *Phys.Rev. E* 61, 3571 (2000).

[224] M. H. Lee, *Phys.Rev. E* 62, 1769 (2000).

[225] R. Peierls, in *Transport Phenomena*, G. Kirchenow and J. Marro, eds., (Springer, Berlin, 1974), p. 1.

[226] D. Ya. Petrina, *Stochastic Dynamics and Boltzmann Hierarchy* (Walter de Gruyter, Berlin, New York, 2009).

[227] S. Nakajima and M. Watabe, *Prog. Theor. Phys.* 29, 341 (1963).

[228] J. S. Langer, *Phys.Rev.* 128, 110 (1962).

[229] P. M. Bakshi and E. P. Gross, *Ann. Phys.* 49, 515 (1968).

[230] M. Lax, *Phys.Rev.* 109, 1921 (1958).

[231] W. Kohn and J. Luttinger, *Phys.Rev.* 108, 590 (1957).

[232] J. Luttinger and W. Kohn, *Phys.Rev.* 109, 1892 (1958).

[233] D. A. Greenwood, *Proc. Phys. Soc.* 71, 585 (1958).

[234] G. V. Chester and A. Thellung, *Proc. Phys. Soc.* 73, 745 (1959).

[235] J. Rammer, *Quantum Transport Theory* (Perseus Books, Reading, Mass., 1998).

[236] M. H. Lee, *Phys.Rev.Lett.* 85, 2422 (2000).

[237] M. Ichiyanagi, *Phys. Rep.* 262, 227 (1995).

[238] H. Nakano, *Int. J. Mod. Phys.* B 7, 2397 (1993).

[239] R. Kubo, *Can. J. Phys.* 34, 1274 (1956).

[240] R. Kubo, *J. Phys. Soc. Jpn.* 12, 570 (1957).

[241] R. Kubo, M. Yokota and S. Nakajima, *J. Phys. Soc. Jpn.* 12, 1203 (1957).

[242] R. Kubo, H. Hasegawa and N. Hashitsume, *J. Phys. Soc. Jpn.* 14, 56 (1959).

[243] R. Kubo, in *Lectures in Theoretical Physics*, W. Brittin and P. Durham, eds., (Interscience Publ., New York, 1959) Vol.1, p.120.
[244] R. Kubo, Rep. Prog. Phys. 29, 255 (1966).
[245] K. M. Case, Transport Theory and Statistical Physics 2, 129 (1972).
[246] M. Suzuki, Prog. Theor. Phys. 43, 882 (1970).
[247] M. H. Lee, Phys.Rev.Lett. 87, 250601 (2001).
[248] M. H. Lee, J. Phys. A:Math.Gen. A 39, 4651 (2006).
[249] M. H. Lee, Phys.Rev.Lett. 98, 110403 (2007).
[250] M. H. Lee, Phys.Rev.Lett. 98, 190601 (2007).
[251] M. H. Lee, Acta Phys. Pol. B 38, 1837 (2007).
[252] M. H. Lee, Int. J. Mod. Phys. B 21, 2546 (2007).
[253] M. H. Lee, Acta Phys. Pol. B 39, 1035 (2008).
[254] M. H. Lee, Int. J. Mod. Phys. B 22, 4572 (2008).
[255] M. H. Lee, Int. J. Mod. Phys. B 23, 3992 (2009).
[256] M. H. Lee, Acta Phys. Pol. B 41, 1009 (2010).
[257] R. Kubo, J. Math. Phys. 13, 1454 (1972).
[258] T. Ohta and T. Ohkuma, J. Phys. Soc. Jpn. 77, 074004 (2008).
[259] N. N. Bogoliubov and S. V. Tyablikov, Doklady Acad. Nauk SSSR 126, 53 (1959).
[260] D. N. Zubarev, Uspekhi Fiz. Nauk 71, 71 (1960).
[261] S. V Tyablikov, Methods in the Quantum Theory of Magnetism (Plenum Press, New York, 1967).
[262] L. L. Moseley and T. Lukes, Am. J. Phys. 46, 676 (1978).
[263] H. B. Callen and T. A. Welton, Phys.Rev. 83, 34 (1951).
[264] R. J. Swenson, Physica 29, 1174 (1963).
[265] R. Zwanzig, J. Chem. Phys. 40, 2527 (1964).
[266] J. S. Langer, Phys.Rev. 127, 5 (1962).
[267] S. F. Edwards, Phil.Mag. 3, 33, 1020 (1958).
[268] E. Verboven, Physica 26, 1091 (1960).
[269] T. Izuyama, Prog. Theor. Phys. 25, 964 (1961).
[270] J. R. Magan, Physica 35, 193 (1967).
[271] J. S. Langer, Phys.Rev. 120, 714 (1960).
[272] J. S. Langer, Phys.Rev. 124, 1003 (1961).
[273] L. Smrcka and P. Streda, *J. Physics C: Solid State Phys.* C 10, 2153 (1977).

[274] H. R. Leribaux, *Phys. Rev.* 150, 384 (1966).

[275] R. Karplus and J. M. Luttinger, *Phys. Rev.* 95, 1154 (1954).

[276] T. Tanaka, K. Moorjani and T. Morita, *Phys. Rev.* 155, 388 (1967).

[277] K. D. Schotte, *Phys. Rep.* 46, 93 (1978).

[278] E. G. D. Cohen and M. H. J. Ernst, *Phys. Lett.* 5, 192 (1963).

[279] I. Prigogine and G. Severne, *Phys. Lett.* 6, 177 (1963).

[280] I. Prigogine, P. Resibois and G. Severne, *Phys. Lett.* 9, 317 (1964).

[281] J. M. Luttinger, *Phys. Rev.* 135, A505 (1964).

[282] J. L. Jackson and P. Mazur, *Physica* 30, 2295 (1964).

[283] P. Resibois, *Phys. Lett.* 9, 139 (1964).

[284] P. Resibois, *J. Chem. Phys.* 41, 2979 (1964).

[285] I. Prigogine, *Nonequilibrium Statistical Mechanics* (Interscience Publ., New York, 1962).

[286] G. V. Chester, *Rep. Prog. Phys.* 26, 411 (1963).

[287] R. Peierls, in: *New Directions in Physics*, N. Metropolis, D. M. Kerr and G.-C. Rota, (eds.), (Academic Press, New York, 1987), p.95.

[288] P. Schofield, *Proc. Phys. Soc.* 88, 149 (1966).

[289] P. Schofield, *Phys. Lett.* A 26, 489 (1968).

[290] W. Bernard and H. B. Callen, *Rev. Mod. Phys.* 31, 1017 (1959).

[291] K. Tani, *Prog. Theor. Phys.* 32, 167 (1964).

[292] R. Peterson, *Rev. Mod. Phys.* 39, 69 (1967).

[293] M. Suzuki, *Prog. Theor. Phys.* 53, 1657 (1975).

[294] M. Ichiyanagi, *J. Phys. Soc. Jpn.* 55, 2963 (1986).

[295] M. Ichiyanagi, *Prog. Theor. Phys.* 76, 949 (1986).

[296] N. G. van Kampen, *Physica Norvegica*, 5, 279 (1971).

[297] E. G. D. Cohen, *Physica* A 118, 17 (1983).

[298] G. Jacucci, *Physica* A 118, 157 (1983).

[299] K. M. van Vliet, *J. Stat. Phys.* 53, 49 (1988).

[300] N. I. Chernov, G. L. Eyink, J. L. Lebowitz, and Ya. G. Sinai, *Phys. Rev. Lett.* 70, 2209 (1993).

[301] N. I. Chernov, G. L. Eyink, J. L. Lebowitz, and Ya. G. Sinai, *Comm. Math. Phys.* 154, 569 (1993).
[302] J. Naudts, J. V. Pule and A. Verbeure, *J. Stat. Phys.* 21, 279 (1979).
[303] D. J. Evans and D. J. Searles, *Adv. Phys.* 51, 1529 (2002).
[304] U.M.B. Marconi, A. Puglisi, L. Rondoni and A. Vulpiani, *Phys. Rep.* 461, 111 (2008).
[305] E.M. Sevick, R. Prabhakar, S.R. Williams and D. J. Searles, *Annu. Rev. Phys. Chem.* 59, 603 (2008).
[306] V. P. Kalashnikov and D. N. Zubarev, *Physica* 59, 314 (1972).
[307] A. L. Kuzemsky, *Int. J. Mod. Phys.* B 19, 1029 (2005).
[308] V. Christoph and A. L. Kuzemsky, Calculation of Transport Coefficients in Solids by Generalized Kinetic Equations, in *Proc. 12-th Annual Intern. Symp. on Electronic Structure of Metals and Alloys*, ed. P. Ziesche (TU Dresden, Dresden, 1982), p. 170.
[309] V. Christoph and A. L. Kuzemsky, *phys. stat. sol. b* 111, K1 (1982).
[310] V. Christoph and A. L. Kuzemsky, *phys. stat. sol. b* 120, K219 (1983).
[311] F. Garcia-Moliner, *Proc. Roy. Soc.* A 249, 73 (1958).
[312] P. B. Allen, *Phys. Rev.* B 17, 3725 (1978).
[313] F. J. Pinski, *Phys. Rev.* B 21, 4380 (1980).
[314] Z. Fisk and G. W. Webb, *Phys.Rev.Lett.* 36, 1084 (1976).
[315] D. W. Woodward and G. D. Cody, *Phys. Rev.* 136, A166 (1964).
[316] R. W. Cohen, G. D. Cody and J. J. Halloran, *Phys.Rev.Lett.* 19, 840 (1967).
[317] Z. Fisk and A. C. Lawson, *Solid State Commun.* 13, 277 (1973).
[318] P. B. Allen, W. E. Pickett, K. M. Ho and M. H. Cohen, *Phys.Rev.Lett.* 40, 1352 (1978).
[319] B. Chakraborty and P. B. Allen, *Phys.Rev.Lett.* 42, 736 (1979).
[320] P. B. Allen, in *Superconductivity in d- and f-Band Metals* eds. H. Suhl and M. B. Maple (Academic Press, New York, 1980), p. 291.
[321] P. B. Allen and B. Chakraborty, *Phys. Rev.* B 23, 4815 (1981).
[322] R. Fogelholm, O. Rapp and G. Grimvall, *Phys. Rev.* B 23, 3845 (1981).
[323] W. Schiller and G. Langowski, *J. Physics F:Metal Phys.* F 12, 449 (1982).
[324] V. Christoph and W. Schiller, *J. Physics F:Metal Phys.* F 14, 1173 (1984).
[325] B. Nikolic and P. B. Allen, *Phys. Rev.* B 63, 020201 (2000).
[326] O. Gunnarsson and J. E. Han, *Nature* 405, 1027 (2000).
[327] M. Calandra and O. Gunnarsson, *Phys.Rev.Lett.* 87, 266601 (2002).
[328] O. Gunnarsson, *Alkali-Doped Fullerides: Narrow-Band Solids with Unusual Properties*, (World Scientific, Singapore, 2004).
[329] P. B. Allen, *Physica* B 318, 24 (2002).

[330] O. Gunnarsson, M. Calandra and J. E. Han, *Rev. Mod. Phys.* 75, 1085 (2003).

[331] K. Vafayi, M. Calandra and O. Gunnarsson, *Phys. Rev.* B 74, 235116 (2006).

[332] M. Gurvitch, *Phys. Rev.* B 24, 7404 (1981).

[333] M. Gurvitch, *Physica* B 135, 276 (1985).

[334] M. Gurvitch, *Phys.Rev.Lett.* 56, 647 (1986).

[335] H. Wiesmann, M. Gurvitch, H. Lutz, A. Ghosh, B. Schwartz, M. Strongin, P. B. Allen and J. W. Halley, *Phys.Rev.Lett.* 38, 782 (1977).

[336] P. J. Cote and L. V. Meisel, *Phys.Rev.* B 25, 2138 (1982).

[337] M. Janssen, *Fluctuations and Localization in Mesoscopic Electron Systems*, World Scientific Lecture Notes in Physics - Vol.64, (World Scientific, Singapore, 1999).

[338] R. Harris, M. Shalmon and M. Zuckermann, *J. Physics F:Metal Phys.* F 7, L259 (1977).

[339] F. J. Ohkawa, *J. Phys. Soc. Jpn.* 44, 1112 (1978).

[340] A. B. Chen, G. Weisz and A. Sher, *Phys.Rev.* B 5, 2897 (1972).

[341] D. Markowitz, *Phys.Rev.* B 15, 3617 (1977).

[342] R. Harris, M. Shalmon and M. Zuckermann, *Phys.Rev.* B 18, 5906 (1978).

[343] S. M. Girvin and M. Jonson, *Phys.Rev.Lett.* 43, 1447 (1979).

[344] S. M. Girvin and M. Jonson, *Phys.Rev.* B 22, 3583 (1980).

[345] D. Belitz and W. Schirmacher, *J. Physics C:Solid State Phys.* C 16, 913 (1983).

[346] B. G. Nickel and J. W. Wilkins, *Phys.Rev.* B 5, 5000 (1972).

[347] A. Gonis, *Green Functions for Ordered and Disordered Systems* (North-Holland, Amsterdam, New York, 1993).

[348] P. Soven, *Phys.Rev.* 156, 809 (1967).

[349] D. W. Taylor, *Phys.Rev.* 156, 1017 (1967).

[350] B. Velicky, S. Kirkpatrick and H. Ehrenreich, *Phys.Rev.* 175, 747 (1968).

[351] B. Velicky, *Phys.Rev.* 184, 614 (1969).

[352] R. J. Elliot, J. A. Krumhansl and P. L. Leath, *Rev. Mod. Phys.* 46, 465 (1974).

[353] K. Levin, B. Velicky and H. Ehrenreich, *Phys.Rev.* B 2, 1771 (1970).

[354] A. B. Chen, *Phys.Rev.* B 7, 2230 (1973).

[355] H. Fukuyama, H. Krakauer and L. Schwartz, *Phys.Rev.* B 10, 1173 (1974).

[356] K. Hoshino and M. Watabe, *J. Physics C:Solid State Phys.* C 8, 4193 (1975).
[357] H. C. Hwang and A. Sher, *Phys.Rev.* B **12**, 5514 (1975).

[358] H. C. Hwang, A. Sher and C. Gross, *Phys.Rev.* B **13**, 4237 (1976).

[359] A. B. Chen, A. Sher and G. Weisz, *Phys.Rev.* B **15**, 3681 (1977).

[360] H. C. Hwang, R. A. Breckenridge and A. Sher, *Phys.Rev.* B **16**, 3840 (1977).

[361] K. Hoshino, *J. Physics F:Metal Phys.* F **9**, 1619 (1979).

[362] F. Goedsche, R. Richter and G. Vojta, *phys. stat. sol.* b **91**, 581 (1979).

[363] P. N. Argyres, *Phys.Rev.* B **23**, 2447 (1981).

[364] P. N. Argyres and S. C. Papadopoulos, *Phys.Rev.* B **23**, 2455 (1981).

[365] M. Hwang, A. Gonis and A. J. Freeman, *Phys.Rev.* B **33**, 2872 (1986).

[366] M. Hwang, A. Gonis and A. J. Freeman, *Phys.Rev.* B **35**, 8974 (1987).

[367] M. Hwang, A. Gonis and A. J. Freeman, *Phys.Rev.* B **35**, 8985 (1987).

[368] F. Brouers and M. Brauwers, *J. Physique (Paris)* **36**, L-17 (1975).

[369] M. Auslender and M. I. Katsnelson, *Phys.Rev.* B **76**, 235425 (2007).

[370] A. H. Castro Neto, F. Guinea, N. M. R. Peres, K. S. Novoselov and A. K. Geim, *Rev.Mod.Phys.* **81**, 109 (2009).

[371] N. M. R. Peres, *J. Phys.: Condens. Matter* **21**, 323201 (2009).

[372] T. Ihn, *Electronic Quantum Transport in Mesoscopic Semiconductor Structures* (Springer, Berlin, 2004).

[373] S. Datta, *Quantum Transport: Atom to Transistor* (Cambridge University Press, Cambridge, 2005).

[374] C. Durkan, *Current at the Nanoscale. An Introduction to Nanoelectronics* (World Scientific, Singapore, 2008).

[375] F. A. Buot, *Nonequilibrium Quantum Transport Physics in Nanosystems* (World Scientific, Singapore, 2009).