Densities of states of the Falicov-Kimball model off half filling in infinite dimensions

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I. INTRODUCTION

The Falicov-Kimball model was introduced to describe the thermodynamics of metal-insulator transitions in compounds that contained both itinerant and localized quasiparticles. The spinless Falicov-Kimball model is the simplest example of an interacting fermionic system that displays numerous phase transitions. Despite its relative simplicity, the analysis of the model is very complex and still a lot of open problems remain. The model can describe many phenomena such as metal-insulator transition, ferromagnetism, antiferromagnetism, phase separations, etc.

In the dynamical mean-field theory, which is exact in infinite dimensions, the Falicov-Kimball model can be solved exactly by calculating the grand canonical potential and the single-particle Green’s function. The thermodynamics of the Falicov-Kimball model is well investigated. The phase transitions between homogeneous phases or phase separations are investigated using the mentioned exact methods as well as using the effective Hamiltonian at the large-U limit (see Ref. 11 for detailed review of the Falicov-Kimball model).

The problem of an evaluation of the spectral function of localized particles is more complex than the investigation of thermodynamics and the phase transitions. The f spectral function related to the localized particles was calculated at half-filling more than ten years ago. Recently, the exact scheme has been extended to the case of different particle concentrations, but due to computational difficulties the densities of states of localized particles have not been completely investigated off half filling. So, we suggest here an approximate analytical scheme within the dynamical mean-field theory for calculating the f spectral function at different particle concentrations and temperatures.

We consider the asymmetric Hubbard model describing the dynamics of two types of particles (ions, electrons or quasiparticles) as a generalization of the Falicov-Kimball model. The Hamiltonian of the asymmetric Hubbard model in a second quantization has the following form

\[ H = -\sum_{i\sigma} \mu_\sigma n_{i\sigma} + U \sum_i n_{i\uparrow} n_{i\downarrow} + \sum_{ij\sigma} t_{ij}^a a_{i\sigma}^\dagger a_{j\sigma}, \]

where \( n_{i\sigma} = a_{i\sigma}^\dagger a_{i\sigma} \) and the motion of particles is described by the creation (\( a_{i\sigma}^\dagger \)) and annihilation (\( a_{i\sigma} \)) operators. The chemical potentials \( \mu_\sigma \) and the transfer parameters \( t_{ij}^a \) depend on a sort of particles (an electron spin). The value \( U \) describes the local on-site repulsion.

The asymmetric Hubbard model was proposed for the description of mixed-valence compounds. This model can also be used for the investigation of the lattice systems having an ionic conductivity with two types of ions. When a single lattice site can be occupied only by one ion, the nontrivial limit \( U \to \infty \) should be considered. Is this case, various thermodynamic regimes can be realized. The chemical potentials or concentrations of particles of different sorts can be fixed independently. In this context the model can be investigated in the presence of the external field corresponding to the difference between the chemical potentials of different sorts.

A number of methods for describing the strongly correlated electron systems has been developed within the dynamical mean field theory. However, all these methods have various restrictions. The quantum Monte-Carlo method is numerically exact but has severe problems at low temperatures and for high repulsion strength. The exact diagonalization method is restricted to a small number of orbitals. Among the numerical techniques the most reliable one at low temperatures is the numerical renormalization group method. For example, its extension was used for the description of the ground state of the standard Hubbard model.

Besides the numerical approaches the development of analytical approximations for the infinite-dimensional model still remains necessary. The Hubbard model in the large-\( U \) limit was investigated using the non-crossing approximation. Many approximations were developed for the weak-coupling regime, for example, the
Edwards-Hertz approach. The alloy-analogy (AA) and modified alloy-analogy (MAA) approximations do not take into account the effect of scattering processes on forming the energy band and cannot be used for the investigation of spectra of the asymmetric Hubbard model. In the Falicov-Kimball limit, the alloy-analogy (AA) and modified alloy-analogy (MAA) approximations give the density of states of localized particles in the form of a delta function. The scattering processes should be taken into account for a correct description of the broadening of this peak. For example, it was the Hubbard-III approximation that originally included the electron scattering for the half-filled Hubbard model into the theory.

We use and improve the approximate analytical approach originally proposed for the Hubbard model and extended to the asymmetric Hubbard model. In this method the single-site problem is formulated in terms of the auxiliary Fermi-field. The approach is based on the equations of motion and on the irreducible Green’s function for the auxiliary Fermi-field. The approach gives DMFT equations in the approximation which is a generalization of Hubbard-III approximation and includes as simple specific cases the AA and MAA approximations.

The approximation is tested on the infinite-dimensional lattice model is mapped on the single-site problem by the use of the auxiliary Fermi-field and the approximations of the asymmetric Hubbard model. We use the approximation to obtain the densities of states of localized particles for various chemical potentials (concentrations) and temperatures. The dependences of the chemical potentials on the particle concentrations are calculated using the densities of states as well as thermodynamically by calculating the grand canonical potential. To calculate the particle spectra at low temperatures the phase separation should be taken into account.

In Section II, we review the formalism of DMFT with the use of the auxiliary Fermi-field and the approximate analytical scheme based on the projecting technique and the different-time decoupling procedure. In Section III and in Appendix, the exact relations between some Green’s functions are derived, which allows us to find the projecting coefficients using only the single particle Green’s function and the coherent potential. The results are discussed in Section IV, followed by our conclusions in Section V.

II. FORMALISM

In the dynamical mean-field theory the infinite-dimensional lattice model is mapped on the single-site problem

\[ e^{-\beta H} \rightarrow e^{-\beta H_{\text{eff}}} = e^{-\beta H_0} \times T \exp \left[ - \int_0^\beta d\tau \int_0^\beta d\tau' \sum_\sigma J_\sigma(\tau - \tau') a_\sigma^\dagger(\tau) a_\sigma(\tau') \right] \tag{2} \]

with the coherent potential \( J_\sigma(\tau - \tau') \) which has to be self-consistently determined from the conditions

\[ G_\sigma(\omega_n) = \frac{1}{\Xi_\sigma^{-1}(\omega_n) - J_\sigma(\omega_n)} \tag{3} \]

\[ G_\sigma(\omega_n, k) = \frac{1}{\Xi_\sigma^{-1}(\omega_n) - \epsilon_\kappa} \tag{4} \]

\[ G_\sigma(\omega_n) = G_\sigma^{\text{ir}}(\omega_n) = \frac{1}{N} \sum_k G_\sigma^\sigma(\omega_n, k) \tag{5} \]

where \( G_\sigma \) is the one-particle Green’s function, \( \Xi_\sigma \) is the total irreducible Green’s function, \( \epsilon_\kappa \) is the energy of states (DOS) for an infinite-dimensional hypercubic lattice and a semielastic DOS for a \( d = \infty \) Bethe lattice.

It was shown in Ref. \[32\] that the single-site problem can be formulated in terms of the auxiliary Fermi-operators \( \xi_\sigma \) describing the creation and the annihilation of particles in the effective environment. The problem is described by the following Hamiltonian

\[ H_{\text{eff}} = H_0 + \sum_\sigma V_\sigma \left( a_\sigma^\dagger \xi_\sigma + \xi_\sigma^\dagger a_\sigma \right) + H_\xi. \tag{6} \]

The approach does not require an explicit form of the environment Hamiltonian \( H_\xi \). The environment is described by the coherent potential given as the Green’s function for the auxiliary Fermi-field with the unperturbed Hamiltonian \( H_\xi \):

\[ J_\sigma(\omega) = 2\pi V_\sigma^2 \langle \xi_\sigma \xi_\sigma^\dagger \rangle_\omega. \tag{7} \]

The particle creation and annihilation operators are expressed in terms of Hubbard operators

\[ a_\sigma = X^{0\sigma} + \zeta X^{2\sigma} \tag{8} \]

on the basis of single-site states \( |n_A, n_B\rangle \)

\[ |0\rangle = |0, 0\rangle, \quad |A\rangle = |1, 0\rangle, \quad |2\rangle = |1, 1\rangle, \quad |B\rangle = |0, 1\rangle, \tag{9} \]

where the following notations for sort indices are used: \( \bar{\sigma} = B, \zeta = + \) for \( \sigma = A \) and \( \bar{\sigma} = A, \zeta = - \) for \( \sigma = B \). In this representation the local Hamiltonian \( H_0 \) of the asymmetric Hubbard model is

\[ H_0 = - \sum_\sigma \left[ \mu_\sigma \left( X^{0\sigma} + X^{2\sigma} \right) \right] + UX^{22}; \tag{10} \]

and the two-time Green’s function \( G_\sigma(\omega) \equiv 2\pi \langle \langle a_\sigma | a_\sigma^\dagger \rangle \rangle_\omega \) is written as:

\[ G_\sigma = 2\pi \left[ \langle \langle X^{0\sigma} | X^{0\sigma} \rangle \rangle_\omega + \zeta \langle \langle X^{0\sigma} | X^{2\sigma} \rangle \rangle_\omega + \zeta \langle \langle X^{2\sigma} | X^{0\sigma} \rangle \rangle_\omega + \langle \langle X^{2\sigma} | X^{2\sigma} \rangle \rangle_\omega \right]. \tag{11} \]

The Green’s functions in Eq. (11) are calculated using the equations of motion for Hubbard operators:

\[ \frac{d}{dt} X^{0\sigma(2)}(t) = [X^{0\sigma(2)}(t), H_{\text{eff}}]. \tag{12} \]
The commutators [12] are projected on the subspace formed by operators $X^0\sigma$ and $X^2\sigma$:

$$[X^\gamma, H_{\text{eff}}] = \alpha_1^\gamma X^0\sigma + \alpha_2^\gamma X^2\sigma + Z^\gamma. \quad (13)$$

The operators $Z^{0\sigma(2\sigma)}$ are defined as orthogonal to the operators from the basic subspace $^83233$,

$$\{\{Z^{0\sigma(2\sigma)}, X^{0\sigma(2\sigma)}\}\} = 0. \quad (14)$$

These equations determine the projecting coefficients $\alpha_i^{0\sigma(2\sigma)}$ which are expressed in terms of the mean value

$$\varphi_\sigma = \langle \xi_\sigma X^{\sigma0} \rangle + \zeta\langle X^{\sigma2}\xi_{\sigma}^\dagger \rangle. \quad (15)$$

Using this procedure by differentiating both with respect to the left and to the right time arguments, we come to the relations between the components of the Green’s function $G_{\sigma}$ and scattering matrix $\tilde{P}_\sigma$. In a matrix representation, we have

$$\hat{G}_\sigma = \hat{G}_0^\sigma + \hat{G}_0^\sigma \hat{P}_\sigma \hat{G}_0^\sigma, \quad (16)$$

where

$$\hat{G}_0^\sigma = 2\pi \left( \frac{\langle \langle X^{0\sigma} | X^{0\sigma} \rangle \rangle + \langle \langle X^{2\sigma} | X^{0\sigma} \rangle \rangle}{\langle \langle X^{2\sigma} | X^{2\sigma} \rangle \rangle} \right), \quad (17)$$

and nonperturbed Green’s function $\hat{G}_0^\sigma$ is

$$\hat{G}_0^\sigma = \frac{1}{D_\sigma} \left( \frac{\omega - b_\sigma}{\omega - a_\sigma} - \frac{\omega}{A_0^\sigma} + \frac{\omega}{A_2^\sigma} \right), \quad (18)$$

where

$$A_{pq} = (X^{pp} + X^{pq}), \quad A_{0\sigma} = 1 - n_\sigma, \quad A_{2\sigma} = n_\sigma, \quad (19)$$

$$D_\sigma = (\omega - a_\sigma)(\omega - b_\sigma) - \frac{V^2_\sigma}{A_0^\sigma A_2^\sigma} \varphi_\sigma^2, \quad (20)$$

$$a_\sigma = -\mu_\sigma + \frac{V_\sigma}{A_0^\sigma} \varphi_\sigma, \quad b_\sigma = U - \mu_\sigma + \frac{V_\sigma}{A_2^\sigma} \varphi_\sigma. \quad (21)$$

The scattering matrix

$$\tilde{P}_\sigma = 2\pi \left( \frac{A_0^{-1}}{0} \right) \left( \frac{0}{A_2^{-1}} \right) \times \left( \frac{\langle \langle Z^{0\sigma} | Z^{0\sigma} \rangle \rangle + \langle \langle Z^{2\sigma} | Z^{0\sigma} \rangle \rangle}{\langle \langle Z^{2\sigma} | Z^{2\sigma} \rangle \rangle} \right) \left( \frac{A_0^{-1}}{0} \right) \left( \frac{0}{A_2^{-1}} \right), \quad (22)$$

being expressed in terms of irreducible Green’s functions contains the scattering corrections of the second and the higher orders in powers of $V_\sigma$. The separation of the irreducible parts in $\tilde{P}_\sigma$ enables us to obtain the mass operator $\hat{M}_\sigma = \hat{P}_\sigma|_\text{ir}$ and the single-site Green’s function expressed as a solution of the Dyson equation

$$\hat{G}_\sigma = (1 - \hat{G}_0^\sigma \hat{M}_\sigma)^{-1} \hat{G}_0^\sigma. \quad (23)$$

We restrict ourselves to the simple approximation in calculating the mass operator $\hat{P}_\sigma$, taking into account the scattering processes of the second order in $V_\sigma$. In this case

$$\hat{M}_\sigma = \hat{P}_\sigma^{(0)}, \quad (24)$$

where the irreducible Green’s functions are calculated without allowance for correlation between electron transition on the given site and environment. It corresponds to the procedure of different-time decoupling, which means in our case an independent averaging of the products of $X$ and $\xi$ operators.

Let us illustrate this approximation on the example of calculation of the following irreducible Green’s function:

$$I(\omega) \equiv \langle \langle (X^{0\sigma} + X^{\sigma\sigma}) \xi_\sigma | \xi_\sigma^\dagger (X^{0\sigma} + X^{\sigma\sigma}) \rangle \rangle|_\text{ir}. \quad (25)$$

According to the spectral theorem, this Green’s function is related to the corresponding time correlation function, and according to the different-time decoupling we have

$$\langle \langle (X^{0\sigma} + X^{\sigma\sigma})_t | (X^{0\sigma} + X^{\sigma\sigma}) \rangle \rangle|_\text{ir} \approx \langle (X^{0\sigma} + X^{\sigma\sigma})_t | (X^{0\sigma} + X^{\sigma\sigma}) \rangle \langle \xi_\sigma^\dagger (t) \xi_\sigma \rangle. \quad (26)$$

Calculation of these correlation functions in a zero approximation

$$\langle (X^{0\sigma} + X^{\sigma\sigma})_t | (X^{0\sigma} + X^{\sigma\sigma}) \rangle \approx \langle (X^{0\sigma} + X^{\sigma\sigma})^2 \rangle = A_{0\sigma} \quad (27)$$

leads to the result

$$I(\omega) = A_{0\sigma} \langle \langle \xi_\sigma | \xi_\sigma^\dagger \rangle \rangle \approx \frac{A_{0\sigma}}{2\pi V^2} J_\sigma(\omega). \quad (28)$$

Using the above procedure we can obtain the final expressions for the mass operator and the total irreducible part:

$$\Xi^{-1}_\sigma(\omega) = \left[ \frac{A_{0\sigma}}{\omega + \mu_\sigma - \Omega_\sigma(\omega)} + \frac{A_{2\sigma}}{\omega + \mu_\sigma - U - \Omega_\sigma(\omega)} \right]^{-1} \quad + \Omega_\sigma(\omega), \quad (29)$$

where

$$\hat{\Omega}_\sigma(\omega) = J_\sigma(\omega) - \frac{R_\sigma(\omega)}{A_{0\sigma} A_{2\sigma}} + \frac{V_\sigma \varphi_\sigma(\omega)}{A_{0\sigma} A_{2\sigma}}, \quad (30)$$

and

$$R_\sigma(\omega) = \frac{\langle X^{\sigma\sigma} + X^{\bar{\sigma}\bar{\sigma}} \rangle}{2} \left[ \frac{\langle X^{\sigma\sigma} - X^{\bar{\sigma}\bar{\sigma}} \rangle}{2\pi} \int_{-\infty}^{+\infty} \frac{\omega' \operatorname{Im} J_\sigma(\omega' + i\epsilon)}{\omega - \omega' - \mu_\sigma + \mu_\bar{\sigma}} \frac{\beta\omega'}{2} \right] \left[ \frac{\langle X^{0\sigma} + X^{2\sigma} \rangle}{2} \int_{-\infty}^{+\infty} \frac{\omega' \operatorname{Im} J_\sigma(\omega' - i\epsilon)}{\omega - \omega' + \mu_\sigma + \mu_\bar{\sigma} - U} \frac{\beta\omega'}{2} \right] \quad (31)$$

and
The approach used here for the approximate solution of the single-site problem can be called as the generalized Hubbard-III (GH3) approximation. It becomes the standard Hubbard-III approximation in the case of the usual Hubbard model at half-filling with spin degeneration \( n_\up = n_\d = 1/2, \mu_\up = \mu_\d = U/2, \langle X^{00} \rangle = (X^{22}), \langle X^{\sigma\bar{\sigma}} \rangle = (X^{\sigma\bar{\sigma}}) \):

\[
\Xi_\sigma^{-1}(\omega) = \left[ \frac{1/2}{\omega + U/2 - 3J_\sigma(\omega)} + \frac{1/2}{\omega - U/2 - 3J_\sigma(\omega)} \right]^{-1} + 3J_\sigma(\omega). \tag{32}
\]

The function \( R_\sigma(\omega) \) describes band forming for particles of sort \( \sigma \) by the motion of particles of another sort \( \bar{\sigma} \) (scattering processes). The neglect of this contribution \((R_\sigma(\omega) = 0)\) gives MAA approximation. If we put \( R_\sigma(\omega) = 0 \) and \( \varphi_\sigma = 0 \), the system is described within the simple AA approximation.

In the limit of infinite repulsion \( U \) the following solution of the single-site problem is obtained:

\[
G_\sigma(\omega) = \frac{1 - n_\sigma}{\omega + \mu_\sigma - \frac{V_{\sigma\bar{\sigma}}}{1-n_\sigma} - J_\sigma(\omega) + R_\sigma(\omega)}; \tag{33}
\]

\[
R_\sigma(\omega) = -\frac{n_\sigma + n_\bar{\sigma}}{2} J_\sigma(\omega + \mu_\sigma - \mu_\bar{\sigma}) - \frac{n_\sigma - n_\bar{\sigma}}{2\pi} \int_{-\infty}^{+\infty} \frac{\text{Im} J_\sigma(\omega' + i\beta) d\omega'}{\omega - \omega' - \mu_\sigma + \mu_\bar{\sigma}} \tanh \frac{\beta\omega'}{2}. \tag{34}
\]

The constant \( \varphi_\sigma = \langle \xi_\sigma X^{\sigma 0} \rangle + \zeta \langle X^{\sigma 2\bar{\sigma}} \rangle \) can be calculated using the exact relation given in the next section. The average particle concentrations are calculated using the imaginary part of the Green’s functions (density of states):

\[
\rho_\sigma(\omega) = -\frac{1}{\pi} \text{Im} G_\sigma(\omega + i\epsilon). \tag{35}
\]

The self-consistency conditions (a set of equations \( 43 - 46 \)) relate the coherent potential \( J_\sigma \) to the Green’s function \( G_\sigma \). For the Bethe lattice with a semielliptic density of states

\[
\rho_\sigma^{\text{Bethe}}(\epsilon) = \frac{2}{\pi W_\sigma^2} \sqrt{W_\sigma^2 - \epsilon^2}, \quad |\epsilon| < W_\sigma \tag{36}
\]

we have

\[
J_\sigma(\omega) = \frac{W_\sigma^2}{4} G_\sigma(\omega). \tag{37}
\]

In the case of the Falicov-Kimball model the unperturbed bandwidth is zero \((2W_B = 0, J_B(\omega) = 0)\) for localized particles, and the approach gives the exact equation for the Green’s function of itinerant particles \( G_A(\omega) \). The density of state \( \rho_A(\omega) \) on the Bethe lattice is nonzero for \( |\omega + \mu_A| < W_A\sqrt{1-n_B} \):

\[
\rho_A(\omega) = \frac{2}{\pi W_A^2} \sqrt{W_A^2(1-n_B) - (\omega + \mu_A)^2}. \tag{38}
\]

In this case the equations \( 33, 43 \) give an explicit approximate expression for the Green’s function of localized particles:

\[
G_B(\omega) = \frac{1 - n_A}{\omega + \mu_B - \frac{V_{\sigma\bar{\sigma}}}{1-n_A} + R_B(\omega)}; \tag{39}
\]

where

\[
R_B(\omega) = -\frac{n_B + n_A}{2} J_A(\omega + \mu_B - \mu_A) - \frac{n_B - n_A}{2\pi} \int_{-\infty}^{+\infty} \frac{\text{Im} J_A(\omega' + i\beta) d\omega'}{\omega - \omega' - \mu_\sigma + \mu_\bar{\sigma}} \tanh \frac{\beta\omega'}{2}. \tag{40}
\]

**III. THE PROJECTING COEFFICIENTS IN EQUATIONS OF MOTION FOR HUBBARD OPERATORS**

The average values \( \langle X^{pq} \xi_\sigma \rangle \) are calculated using corresponding Green’s functions according to the spectral theorem

\[
\langle X^{pq} \xi_\sigma \rangle = \int_{-\infty}^{+\infty} \frac{d\omega}{e^{\beta\omega} + 1} \left[ -2 \text{Im} \langle \xi_\sigma | X^{pq} \rangle_\omega \right]. \tag{41}
\]

These Green’s functions can be calculated using the exact relation \( 16, 24 \) derived in Appendix

\[
V_\sigma \langle \langle \xi_\sigma | X^{pq} \rangle \rangle_\omega = J_\sigma(\omega) \langle \langle a_\sigma | X^{pq} \rangle \rangle_\omega. \tag{42}
\]

For the asymmetric Hubbard model we have to calculate the coefficients \( V_\sigma \varphi_\sigma \):

\[
\varphi_\sigma = \langle \xi_\sigma X^{\sigma 0} \rangle + \zeta \langle X^{\sigma 2\bar{\sigma}} \rangle = \langle \xi_\sigma (X^{\sigma 0} + \zeta X^{2\bar{\sigma}}) \rangle \tag{43}
\]

which are expressed as

\[
V_\sigma \varphi_\sigma = \int_{-\infty}^{+\infty} \frac{d\omega}{e^{\beta\omega} + 1} \text{Im} \rho_\sigma^\epsilon(\omega + i\epsilon), \tag{44}
\]

\[
\rho_\sigma^\epsilon(\omega) = 2J_\sigma(\omega) \langle \langle X^{0\sigma} - \zeta X^{\sigma 2\bar{\sigma}} \rangle \rangle_\omega = 2J_\sigma(\omega) \left[ \langle \langle X^{0\sigma} | X^{\sigma 0} \rangle \rangle_\omega - \langle \langle X^{\sigma 2\bar{\sigma}} \rangle \rangle_\omega \right]. \tag{45}
\]

In the limit of infinite on-site repulsion \( U \) the state with double occupation is excluded and we have

\[
\rho_\sigma^\epsilon(\omega) = \frac{1}{\pi} J_\sigma(\omega) G_\sigma(\omega). \tag{46}
\]

For the Falicov-Kimball model it is possible to calculate the exact Green’s functions for itinerant particles and it gives the exact expression for \( V_B \varphi_B \).

Let us note that previously this parameter was calculated approximately using the Green’s functions obtained by means of the linearized equations of motion and neglecting the irreducible parts \( 32, 33 \). In the model with infinite \( U \) this approximate value is

\[
\rho_B^\epsilon(\omega) = \frac{1}{\pi} J_A(\omega) \frac{1 - n_B}{\omega + \mu_B}, \tag{47}
\]
and it corresponds to the approximation of the series where only the first term (the zero approximation for $G_A$) is taken into account.

There is shown a comparison between the approximate and the exact values of $\phi_B$ in Fig. 1. The improvement given by the exact relation allows us to investigate the system at low temperatures where summing up the whole series is essential.

IV. RESULTS

Dependences of chemical potentials $\mu_A$ and $\mu_B$ on the particle concentrations are calculated using corresponding densities of states. DOS of localized particles is obtained as an imaginary part of the Green’s function

$$\rho_B(\omega) = \frac{1}{\pi} \text{Im} G_B(\omega + i\varepsilon). \quad (48)$$

This Green’s function has the correct analytic properties (relations between imaginary and real parts) within the considered approximations (GH3, MAA, AA). The approximate DOS always has the correct sign and the sum rule is fulfilled: the integral of DOS over all frequency is equal to unity for finite $U$ and is equal to $1 - n_\sigma$ for $U \to \infty$ when the upper band tends to infinity.

For the Falicov-Kimball model the density of states of localized particles can be calculated exactly, but numerical results were obtained mostly at half-filling. The constant $\phi_B$ is zero at half-filling ($n_A = n_B = 1/2$) because of the particle-hole symmetry. In this case we have the simple approximate solution of the single-site problem

$$G_B(\omega) = \frac{\omega - 2J_A(\omega)}{\omega^2 - U^2/4 - 2\omega J_A(\omega)}. \quad (49)$$

This result at half-filling is independent of temperature, but for high temperatures and large values of $U$ the approximate scheme reproduces the exact results (Fig. 2).

For simplicity we restrict our investigation to the Falicov-Kimball model with the infinite repulsion $U$ on the Bethe lattice. In this case the model describes the system with an average site occupation no more than unity. There is a homogeneous state for temperatures larger than critical $T_c \approx 0.060 W_A$. For lower temperatures ($T < T_c$) there are various types of phase transitions depending on a thermodynamic regime.

First, we consider a possibility of describing the phase transitions using the approximate equations. For this reason, the behavior of dependences of the chemical potential $\mu_B$ on the concentration of localized particles $n_B$ is investigated. The phase transition is indicated by the thermodynamic unstable region where $\partial \mu_B / \partial n_B < 0$. When the critical temperature is approached from below, the dependence $\mu_B(n_B)$ becomes monotonic ($\partial \mu_B / \partial n_B \geq 0$).

In Fig. 3 the approximate curves $\mu_B(n_B)$ are compared with the exact results obtained thermodynamically. The alloy-analogy based approximations give the density of states of localized particles in a form of a noninteracting delta-function, which is correct in the atomic limit $t_A = 0$. Thus, the MAA approximation can give reasonable results only when the chemical potential $\mu_A < 0$ (i.e. the concentration of itinerant particles tends to zero). This approximation shows the presence of a phase transition. However, it overestimates the critical temperature. In Fig. 3 the exact curve at $T = 0.060$ is already monotonic but the MAA approximation shows the region with unstable concentration values, i.e. there is still a phase transition.

The GH3 approximation, unlike the MAA approximation, incorporates the scattering processes forming the energy band of localized particles. This is crucial for the calculation of thermodynamic quantities. The approximate curves coincide with the exact ones in a wide range of temperatures for the concentrations larger than some value which depends on $\mu_A$. In Fig. 3 ($\mu_A = 0$) one can see a good agreement with the exact result at $\mu_A = 0$ for
concentrations \( n_B \gtrsim 0.5 \). At temperatures lower than the critical one the approximation clearly indicates the phase transition and gives the correct value for \( T_c \approx 0.060 \).

Spectra of localized particles are plotted in Fig. 4. In this case temperatures are higher than the critical one, so a homogeneous state is stable. The parameter values are chosen so that the approximation gives the correct thermodynamic \( \mu_B(n_B) \) relations. The energy band of itinerant particles (A) depends only on a concentration \( n_B \) and its width is \( 2W_A\sqrt{1-n_B} \). However, the band of localized particles (B) is generated by scattering processes and its spectral shape depends on the concentration \( n_B \), the chemical potential \( \mu_A \) (or the corresponding concentration \( n_A \)) and temperature.

Depending on the values of the chemical potential of itinerant particles \( \mu_A \) (or \( n_A \)) there are two limiting cases with different properties of the localized particle spectrum. For very small \( n_A \) (negative \( \mu_A \)) the system is close by its behavior to the atomic limit, i.e., the spectrum \( \rho_B(\omega) \) is in the form of a delta peak. The sharp peak is slightly broadened by the scattering of itinerant particles (Fig. 4A, \( \mu_A = -0.4 \)). The contrary case with the nearly filled bands is when the total concentration of particles tends to unity, \( n_A + n_B \to 1 \) (Fig. 4C, \( \mu_A = 0.4 \)). In this case the peak vanishes because the contribution of the coherent potential of mobile particles \( J_A(\omega) \) and the corresponding function \( R_B(\omega) \) becomes larger and the simple pole in \( \rho_B(\omega) \) disappears. The spectrum \( \rho_B(\omega) \) corresponds by the form to the lower Hubbard subband (or the lower subband for the half-filled Falicov-Kimball model obtained in Ref. 12) with the chemical potential located in a gap in the strong coupling limit. The intermediate case with the broad band superimposed by the sharp peak is shown in Fig. 4B.

States with concentration values in some region (where \( \partial \mu_\sigma / \partial n_\sigma < 0 \)) are thermodynamically unstable at low temperatures. So, the presence of phase transitions should be taken into consideration. There are phase transitions between homogeneous phases with a concentration jump in the thermodynamic regime with the fixed chemical potentials \( (\mu_A = \text{const}, \mu_B = \text{const}) \). In this case the density of states changes instantly as the concentration jumps. If one of the concentrations \( n_A \) or \( n_B \) is fixed the homogeneous state is unstable and the phase separation takes place. The phase transitions for the Falicov-Kimball model were investigated in many works (see Ref. 11); the regimes mentioned above were investigated for the Falicov-Kimball model using the exact thermodynamic equations in Ref. 13.

Let us consider the thermodynamic regime with fixed

![Figure 3](image3.png)

**FIG. 3:** The dependence of \( \mu_B \) on \( n_B \) in different approximations is compared with the exact result obtained thermodynamically. The parameter values: \( W_A = 1; W_B = 0; U = \infty \). 1 – exact result; 2 – MAA; 3 – GH3.

![Figure 4](image4.png)

**FIG. 4:** DOS of localized particles within the GH3 approximation for various \( \mu_A \) and temperatures. \( W_A = 1; W_B = 0; U = \infty \); 1) \( T = 0.06 \), 2) \( T = 0.2 \).
values of \( \mu_A \) and \( n_B \). In the homogeneous state the spectral function of itinerant particles \( \rho_A \) depends on \( n_B \) and \( \mu_A \) (Eq. 35) and is independent of temperature. However, in the phase separated state the spectrum of the whole system can be considered as a superposition of spectra of each component. The homogeneous state is unstable for the concentration values \( n_B \in (n_{B1}, n_{B2}) \) and the system is separated into two different phases with the concentrations of \( B \) particles \( n_{B1} \approx 0.52 \) and \( n_{B2} \approx 0.78 \) at \( T = 0.059W_A \). The concentrations of the components \( (n_{B1}, n_{B2}) \) depend on temperature. Thus, the spectra \( \rho_A \) and \( \rho_B \) are temperature dependent in the segregated phase. In Fig. 5(A) DOS of localized particles is plotted at temperature \( T = 0.059W_A < T_c \). In Fig. 5(B) the spectrum at various temperatures is compared. The bandwidth is larger in the segregated state than in the homogeneous state \( (T > T_c) \) and is temperature dependent.

V. CONCLUSIONS

The approximate analytic approach within DFMT for calculating the single-particle Green’s functions of the asymmetric Hubbard model is developed and improved. This approximation allows one to investigate the model for various concentration values. The method is tested on the Falicov-Kimball model with the infinite on-site repulsion. It is shown that for high enough temperatures or large concentrations of localized particles the approximate approach reproduces exact values of chemical potentials. The approximation can correctly indicate the instability of a homogenous state and the presence of phase transitions.

The generalized Hubbard-III approximation (GH3) partially includes the scattering of particles into the theory and describes the formation of the band of localized particles. In the infinite-\( U \) limit the spectrum of localized particles is obtained for various particle concentrations and temperatures. The form of this spectrum continuously changes from a delta peak to the characteristic form of the lower subband of the spectrum in the Hubbard-III approximation when the chemical potential of itinerant particles increases.

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APPENDIX: EXACT RELATIONS BETWEEN GREEN’S FUNCTIONS

Let us consider the effective single-site problem in terms of the auxiliary Fermi-field

\[
\hat{H}_{\text{eff}} = \hat{H}_0 + \hat{H}_\xi + \sum_{\sigma=1}^{m} V_{\sigma} \left( a_\sigma^\dagger \xi_{\sigma} + \xi_{\sigma}^\dagger a_\sigma \right), \quad (A.1)
\]

\( \hat{H}_0 \) is a single-site Hamiltonian; \( \hat{H}_\xi \) is an auxiliary environment Hamiltonian. The number \( m \) of sorts of itinerant particles can be arbitrary (\( m = 1, 2, 3, \ldots \)). So, the effective Hamiltonian can describe the Falicov-Kimball model (\( m = 1 \)) and the Hubbard model (\( m = 2 \)). The algebra of \( \xi \) operators is defined by the anticommutation relations

\[
\{ \xi_{\alpha}, \xi_{\beta} \} = \{ \xi_{\alpha}^\dagger, \xi_{\beta}^\dagger \} = 0, \quad \{ \xi_{\alpha}, \xi_{\beta}^\dagger \} = \delta_{\alpha,\beta}. \quad (A.2)
\]

It can be proved that the following relations take place

\[
V_{\sigma} \langle \langle \xi_{\sigma} | \hat{A} \rangle \rangle_\omega = J_\sigma(\omega) \langle \langle a_{\sigma} | \hat{A} \rangle \rangle_\omega, \quad (A.3)
\]

\[
2\pi V_{\sigma}^2 \langle \langle \xi_{\sigma}^\dagger | \xi_{\sigma}^\dagger \rangle \rangle_\omega = J_\sigma(\omega) + 2\pi J_{\sigma}^3(\omega) \langle \langle a_{\sigma}^\dagger | a_{\sigma}^\dagger \rangle \rangle_\omega, \quad (A.4)
\]

where \( \hat{A} \) is the arbitrary Fermi-operator that anticommutes with the \( \xi \) operators, and \( J_\sigma(\omega) \) is the Green’s function for the unperturbed Hamiltonian

\[
J_\sigma(\omega) = 2\pi V_{\sigma}^2 \langle \langle \xi_{\sigma}^\dagger | \xi_{\sigma}^\dagger \rangle \rangle_\omega. \quad (A.3)
\]

Proof. Thermodynamic perturbation theory can be formulated based on the interaction representation for the statistical operator with the use of the \( \hat{H}_0 + \hat{H}_\xi \) operator as the zero-order Hamiltonian:

\[
\dot{\rho} = e^{-\beta \hat{H}_{\text{eff}}} = e^{-\beta (\hat{H}_0 + \hat{H}_\xi)} \dot{\sigma}(\beta), \quad (A.5)
\]

\[
\dot{\sigma}(\beta) = e^{-\beta (\hat{H}_0 + \hat{H}_\xi) / \beta} \exp \left[-\int_0^\beta d\tau \hat{H}_{\text{int}}(\tau) \right]. \quad (A.6)
\]

The part of the interaction Hamiltonian describing one sort (\( \sigma \)) of particles is separated off:

\[
\hat{H}_{\text{int}} = V_{\sigma} \left( a_{\sigma}^\dagger \xi_{\sigma}^\dagger + \xi_{\sigma} a_{\sigma} \right) + \hat{B}_{\sigma}. \quad (A.7)
\]
The residue $\hat{B}_\sigma$ commutates with the operators of the chosen sort $\sigma$: $[\hat{B}_\sigma, \xi_\sigma] = [\hat{B}_\sigma, \xi_\sigma^\dagger] = 0$. Thus, at the perturbation expansion the operator $\xi_\sigma^{(1)}$ does not have to be paired with the operator $\hat{B}_\sigma$. Here we introduce notations for the Green’s functions

\[
G_\sigma(\tau - \tau') = \langle \tau | \xi_\sigma^{(1)}(\tau) \xi_\sigma(\tau') | 0 \rangle,
\]

\[
G_\sigma(\tau - \tau') = \langle \tau | a_\sigma^\dagger(\tau) a_\sigma(\tau') | 0 \rangle,
\]

\[
\Phi_\sigma(\tau - \tau') = \langle \tau | \hat{A}_\sigma(\tau) \xi_\sigma(\tau') | 0 \rangle,
\]

\[
G_\sigma(\tau - \tau') = \langle \tau | T \hat{A}_\sigma(\tau) \xi_\sigma(\tau') | 0 \rangle.
\]

The perturbation theory expansion of the scattering matrix $\Phi(\beta)$ gives the following series for the Green’s function $\Phi_\sigma(\tau - \tau')$:

\[
\Phi_\sigma(\tau - \tau') = \frac{V_\sigma^{2l+1}}{(\sigma(\beta))_0} \sum_{p=0}^{\infty} \frac{(-1)^p}{(2p+1)!} \sum_{l=0}^{p} \frac{(2p+1)!}{l!(l+1)!(2p-2l)!} \int_0^\beta d\tau_1 \ldots d\tau_{2p+1}
\]

\[
\times \langle \tau | \hat{A}(\tau) \xi_\sigma(\tau') \rangle \langle a_\sigma^\dagger \xi_\sigma(\tau') \rangle \cdots \langle a_\sigma^\dagger \xi_\sigma(\tau') \rangle \langle \xi_\sigma(\tau') \rangle_{l+1} \cdots \langle \xi_\sigma(\tau') \rangle_{2l+1} \hat{B}_\sigma(\tau_{2l+2}) \cdots \hat{B}_\sigma(\tau_{2p+1}) \rangle_0.
\]

(A.8)

The averaging of $T$ products is performed in the zero-order Hamiltonian according to the Wick’s theorem by the consecutive pairing. We start the pairing procedure from the operator $\xi_\sigma(\tau')$ and it is performed only with the operators $\xi_\sigma^\dagger$. After the first pairing we have the following expression:

\[
\Phi_\sigma(\tau - \tau') = \frac{V_\sigma^{2l+1}}{(\sigma(\beta))_0} \int_0^\beta d\tau'' G_\sigma(\tau'' - \tau') \sum_{p=0}^{\infty} \frac{(-1)^p}{(2p+1)!} \sum_{l=0}^{p} \frac{1}{l!(2p-2l)!} \int_0^\beta d\tau_1 \ldots d\tau_{2p}
\]

\[
\times \langle \tau | \hat{A}(\tau) a_\sigma(\tau'') \rangle \langle a_\sigma^\dagger \xi_\sigma(\tau'') \rangle \cdots \langle a_\sigma^\dagger \xi_\sigma(\tau'') \rangle \langle \xi_\sigma(\tau'') \rangle_{l+1} \cdots \langle \xi_\sigma(\tau'') \rangle_{2l+1} \hat{B}_\sigma(\tau_{2l+2}) \cdots \hat{B}_\sigma(\tau_{2p}) \rangle_0.
\]

(A.9)

Summing up the series in (A.8) gives the Green’s function $G_\sigma(\tau - \tau')$ and we have

\[
\Phi_\sigma(\tau - \tau') = V_\sigma \int_0^\beta d\tau'' G_\sigma(\tau'' - \tau') G_\sigma(\tau''),
\]

or in the Matsubara frequency representation:

\[
\Phi_\sigma(\omega_n) = V_\sigma G_\sigma^{(4)}(\omega_n) G_\sigma(\omega_n).
\]

(A.10)

(A.11)

Finally, in order to obtain the expression (A.3), an analytical continuation of the Green’s functions from the imaginary axis to the real one should be done ($\omega_n \rightarrow \omega + i\varepsilon$, $G_\sigma^{(4)}(\omega_n) \rightarrow 2\pi \langle \langle a_\sigma | \hat{A} \rangle | a_\sigma \rangle$, $\Phi_\sigma(\omega_n) \rightarrow 2\pi V_\sigma \langle \langle \xi_\sigma | \hat{A} \rangle | \omega_n \rangle$, ...). If we put $A = \xi_\sigma^\dagger$, the pairing of the operator $\xi_\sigma(\tau')$ with $\xi_\sigma^\dagger(\tau)$ gives the first term in (A.3). The second term is obtained using the procedure the same as at deriving the relation (A.3).
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