Electrical conductivity determination of semiconductors by utilizing photography, finite element simulation and resistance measurement

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ABSTRACT

A new method is developed to measure precisely and reliably the electrical conductivity of a block-shaped semiconductor specimen using four-wire technique with electrodes in arbitrary shape and position. No effort for accurate electrode preparation is necessary anymore. This method may be especially applied to measure the conductivity of ceramics at high temperatures, when typical spring-contacts or clamp-contacts are not possible and instead wound wires are used for electrically contacting the specimen. The method comprises the following: An image of the specimen is processed to a 3D model. By applying a finite element simulation on this 3D model, a form factor (also called geometry factor) that considers the effect of the non-infinitesimally small electrodes is calculated. Together with the measured resistance (preferably in four-wire technique), the actual conductivity of the sample is derived. Experimental results confirmed the validity of the proposed method. As a limitation of the method, the conductivity of the specimen should be within the range of 0.01 Sm⁻¹ and 10⁶ Sm⁻¹.

Introduction

Depending on the sample geometry, various methods [1, 2] are developed to measure the conductivity of semiconductors. Basically, the four-point probe technique is often applied to exclude contact resistances. [3, 4]. As for disk-like samples of arbitrary shape, Van der Pauw method [5] is applicable, if the sample is homogeneous and has no holes. Electrodes have to be infinitesimally small and have to be placed at the edge of the sample. In order to further reduce the measurement error, ASTM F76-08 suggests, for instance, specimens should have the shape of a clover-leaf [6]. One drawback of the Van der Pauw method is that cutting brittle semiconductors, e.g., oxide ceramics, into the required clover-leaf shape is difficult and time-consuming. Especially for ceramics
to be characterized at high temperatures (i.e., when determining the conductivity vs. oxygen partial pressure, as it is typical for defect chemical considerations [7, 8]), spring-contacts or clamp-contacts are hardly possible. In that case, it is common practice to use block-shaped or plate-like specimens that are sometimes referred to as “bar geometry.” Four wires (for four-wire technique) are wound around the specimens and the wires are connected to the specimens with a noble metal paste. For both the Van der Pauw method and the bar geometry, four-wire technique is appropriate, to exclude contact resistances. A scheme is given in Fig. 1a. Very often the blocks are thin in one dimension to form rectangular-shaped small plates, see, e.g., [9–11]. Under the assumption that the electrodes are parallel and infinitely small, the equipotential surfaces can be considered as flat and equidistant. Thus, the electrical field is homogeneous everywhere in the specimen. The conductivity $\sigma$ can be directly derived from the current $I$ passing through electrode $E_1$ and $E_4$, and the potential difference $U_{23}$ between electrode $E_2$ and electrode $E_3$, the distance $L$ between electrode $E_2$ and $E_3$, and the area of the cross section $A$, using

$$\sigma = \frac{I}{U_{23}} \cdot \frac{L}{A} = \frac{I}{U_{23}} \cdot K$$

(1)

In this simple geometry, the form factor (or geometry factor) $K$ can be written as $K = L/A$.

In order to prepare specimens that meet the assumptions above, sometimes even laser engraving is used to make narrow slits where the electrode should be fitted into [11]. Since the specimens usually have dimensions in the mm range, e.g., 10 mm × 3 mm × 3 mm [8], or 15 mm × 5 mm × 0.3..1 mm [10], fitting the thin wires into the slits is conducted under a microscope and their thorough preparation is labor-intensive. Some typical setups from sample as shown in literature are given in Fig. 2. Figure 2c is a typical example for a very precisely prepared specimen, although only in two-wire technique.

In most cases, the slits are not engraved and a conductive paste is used to guarantee a good electrical contact of the electrodes with the specimen. Because the conductive paste is usually applied manually with a small brush, the electrodes are broadened and they have irregular shapes, see, e.g., the deliberately sloppily prepared specimen in Fig. 1b. There, the homogeneous electric field near the electrodes is distorted. Since the electrodes are neither infinitely thin nor parallel to each other, the distance $L$ is no longer trivial to define, and hence, the form factor $K$ cannot be simply set to $K = L/A$ as in Eq. 1. Using FE (finite element) simulation on a simplified example, it is shown that the relative error in the measurement of the conductivity can be at least 20%, if $L$ is taken as the distance between the centerline of the two inner electrodes (see Part I of the Supplementary Information).

Therefore, in order to eliminate the error due to the false determination of $L$ and in order to allow using more or less arbitrarily shaped, especially wide and not well-defined electrode geometries so that the time for precise electrode preparation can be reduced, a new method combining 3D imaging or photography combined with finite element (FE) simulation is developed in this work. It also considers the effect of non-infinitesimally small electrodes.

The basic idea comprises
1. to apply electrodes onto the specimen without any special and/or well-defined geometry,
2. to determine the geometry of specimen and electrodes by appropriate means. This may be any type of 3D scanning or tomography; for flat samples with a known thickness as demonstrated here as a first validation, simple photography can be used to determine the electrode areas,
3. to establish a 3D model of the specimen including the electrodes, with the electrodes being defined as equipotential regions,
4. to calculate a form factor $K$ from the derived 3D model with the help of a Finite element (FE) software tool
5. to measure the resistance or conductance (e.g., by applying $I$ and measuring $U$), preferably in four-wire technique to avoid contact resistance issues and
6. to calculate the conductivity $\sigma$ out of the resistance and the derived form factor $K$.

**Theoretical background of the new method**

Some assumptions to correctly calculate the form factor apply. First of all, the specimens must be homogeneous and compact. Furthermore, the conductivity of the electrodes must be at least three magnitudes higher than the conductivity of the specimen. This is a precondition, so that the electrode area can be considered as an equipotential area. For the simulation, let us assume the distance of the two inner electrodes if they were infinitely thin and parallel to each other is $L$, the cross-sectional area where the current flows be $A$ and the conductivity of the specimen in simulation be $\sigma_s$, with index “s” denoting simulation. The resistance of the specimen $R_s$ is expressed using a correction factor $\delta$ due to effect of non-infinitesimally thin electrodes

$$ R_s = \frac{1}{\sigma_s} \cdot \frac{L}{A} = \frac{U_s}{I} \quad (2) $$

Meanwhile, the simulated potential difference of the inner electrodes is $U_s$ and the applied current in simulation is $I$. Since $\delta$, $L$ and $A$ only involve the geometry of the specimen, Eq. (2) can be written using a form factor $K_s$ as (Ohm’s law):

$$ R_s = K_s \cdot \frac{1}{\sigma_s} = \frac{U_s}{I} \quad (3) $$

with $K_s = \delta \cdot \frac{L}{A}$

On the other hand, the measured specimen has an unknown conductivity $\sigma_m$, with index “m” indicating measurement. So, its resistance is

$$ R_m = K_m \cdot \frac{1}{\sigma_m} = \frac{U_m}{I} \quad (4) $$

The measured potential difference between the inner electrodes is $U_m$. Because the currents for simulation and measurement were set to have the same values, one can derive out of Eqs. (3) and (4)

$$ \frac{\sigma_s U_s}{K_s} = \frac{\sigma_s U_m}{K_m} \quad (5) $$

Since the specimen in simulation has the same geometry as the specimen in the actual measurement, Eq. (6) follows
From Eqs. (5) and (6), one obtains

$$K_s = K_m = K$$  \hspace{1cm} (6)

$$
s_{s}U_s = s_m U_m = \text{const.}$$  \hspace{1cm} (7)

$$
s_m = \frac{s_s U_s}{U_m} = \frac{s_s U_s}{IR_m}$$  \hspace{1cm} (8)

Equations (3) and (6) lead to the form factor $K_m$

$$K_m = \frac{s_s U_s}{I}$$  \hspace{1cm} (9)

The described procedure to calculate the conductivity $\sigma$ and the form factor $K_m$ and its validation are summarized in Fig. 3.

For the simulation, the current $I$ and the conductivity $s_s$ are input parameters for the 3D model in the FE software to simulate the potential difference of the inner electrodes $U_s$. $I$ is set to have the same value as the current that will be used later in the four-wire technique. $s_s$ can be set as any positive value, as long as the conductivity of the electrodes in the simulation (here $10^5$ S/cm) is set to be at least three magnitudes higher than $s_s$. Numeric errors may occur if $s_s$ is too small. This problem will be addressed in the discussion part. On the other hand, the resistance of the specimen is measured by four-wire technique. Since the used current $I$ is known, the actual potential difference is calculated by multiplying $I$ with the measured resistance $R_m$, which leads to $U_m$. According to the above said, $s_s U_s = s_m U_m$. Therefore, for the unknown conductivity of the specimen $s_m = \frac{s_s U_s}{U_m}$ and for the form factor of the specimen $K_m = \frac{\sigma_m U_s}{I}$ hold. To validate the calculation, the calculated $\sigma_m$ is put again into the simulation with the current $I$ to simulate a new potential difference $U_{sim}$. This potential difference is then compared with the measured potential difference $U_m$ to see to which extend they agree with each other.

**Method**

This section explains the procedure how the geometry was obtained. It can also be considered as a kind of recipe. The following steps from taking the camera image to building up the CAD (computer-aided design) model that is imported into the FE software should be considered as an example. Therefore, only a few words are spent in the main article, but the steps are comprehensively described in Part II of the Supplementary Information.

Inside a photo light box, a scale was placed next to the specimen. The camera (here a camera of a mobile phone) was arranged above the specimens, and the camera lens was kept parallel to the surface of the specimens. The images were taken using the automatic mode. Typical images for two specimens are given in Fig. 4.

The camera images were denoised using a so-called technique “morphological anisotropic diffusion” [12, 13] followed by edge detection using a canny edge detector. The resulted edge maps were processed to vector graphics using centerline tracing technique [14]. After postprocessing such as

![Diagram](image-url)

**Figure 3** Procedure to calculate the conductivity, to derive the form factor $K_m$, and to validate it. The form factor $K_m$ outlined with a pentagon is the output, whereas the current $I$ outlined with a triangle is the input.

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eliminating artifacts, 3D extrusion in the CAD program and rescaling, the CAD models were imported into COMSOL Multiphysics. After assigning the boundary conditions, the conductivity and the form factor were calculated following the procedures illustrated in Fig. 3.

Experiments

To verify the described method above, a ceramic spinel-type NiMn$_2$O$_4$-based block (50 mm x 5 mm x 0.6 mm) provided by Vishay Electronic GmbH, Selb, Germany, was used, with a composition similar to the one described in [15]. Two specimens (sample A, sample B, with a thickness of 0.60 mm each) were cut from the block, and each specimen was contacted with four electrodes using silver wires and silver paint (Ferro, #530042). A typical specimen image is shown in Fig. 4. The resistance of each specimen was measured in a silicon oil bath where the temperature was controlled very precisely at 25 °C (for details of the setup see [16]). Then, the electrodes were removed and newly contacted with other electrode geometries. The experiments were conducted three times. In each experiment, the form of the electrodes was modified. The specimens are denoted as A1, A2, A3 and B1, B2, B3, respectively. They are shown in Fig. 5a. Their respective CAD plots are shown in Fig. 5b.

Results and discussion

In Table 1, the calculated form factors $K_m$ of each specimen are shown. Between A3 and A2, the form factor differs by a ratio of 2.48. In other words, the voltage drop $U_{23}$ would differ by a factor of almost 2.5, when applying the same current $I$.

Although the form factor of the specimens differs significantly within sample A and sample B, the average derived conductivity of the sample A and B is $3.613 \pm 0.249$ S/cm and $4.07 \pm 0.184$ S/cm, respectively. The standard deviation of the conductivity for sample A and sample B is 6.89% and 4.52% respectively, indicating that specimens A1, A2, A3 and specimens B1, B2, B3 should be the same material. The difference in the average conductivity of sample A and sample B might be due to the fact that sample A and sample B were cut from different parts of the bulk material and inhomogeneity in the bulk material. Another explanation for the difference might be the precision of the thickness measurement. The thickness was measured by a caliper at three different positions of the specimen (near the left end, in the middle and near the right end) and an average thickness was taken as the overall thickness of the sample; however, the thickness of the specimens might not be constant and the thickness measurement has a precision of 0.05 mm; this is already approximately 8% of the total sample thickness of 0.60 mm. The measurement uncertainty of the thickness can be reduced by using a micrometer screw gauge rather than a caliper. Other errors might come from distortion of the image due to distortion caused by camera lens and denoising of the image. Nevertheless, the good agreement between measured $U_{23}$
and $U_m$ from the simulation using $\sigma_m$ supports the validity of the proposed method.

As a side remark, one may consider the effect of temperature on the form factor. For simplicity, let us assume the thermal expansion is isotropic and the thermal expansion coefficient is $a$, and then, the form factor at elevated temperatures with temperature difference of $\Delta T = T - 25 \, ^\circ C$ is:

$$K_m(T) = \delta \frac{L(1 + a\Delta T)}{A(1 + a\Delta T)^2} = \frac{K_m(25^\circ C)}{(1 + a\Delta T)}$$  \hspace{1cm} (10)

Assuming a typical value for a thermal expansion coefficient of ceramic material of $a \approx 10^{-6} \, K^{-1}$, and $T = 1025 \, ^\circ C$, i.e., $\Delta T = 1000 \, ^\circ C$, a deviation from the room temperature value for $K_m$ of $\frac{1}{(1 + a\Delta T)} \approx 0.99$ follows. Therefore, the form factors at 1000 $^\circ C$ differ only by ca. 1% from the form factors at 25 $^\circ C$. In other words, the form factors obtained for a specimen at room temperature can be used for the same specimen at high temperatures. Further experiments at high temperatures are necessary to validate this theory.

The question may arise, whether this method has limitations. Two limitations are obvious:

1. numerical errors arise if $\sigma_s$ for simulation is set below 0.01 $Sm^{-1}$
2. the procedure is not applicable for sample with a conductivity above $10^6$ $Sm^{-1}$

The first limitation is based on two facts derived from the simulation of the model in Fig. S1. The first fact is that the ratio $\frac{U_s}{U_s^0}$ deviates significantly from 1, when $\sigma_s$ is below 0.01 $Sm^{-1}$ (see Fig. 6). Here, $U_s$ and $\sigma_s$ are values from the case where the conductivity of the specimen in the simulation is 1 $Sm^{-1}$. Since $\frac{U_s}{U_s^0} = const. \ (see \ Eq. \ (7))$, it is to expected that this ratio should be 1. The second fact is that the potential difference is not on the extrapolated line, when $\sigma_s$ is below 0.01 $Sm^{-1}$ (see Fig. 6). However, the potential difference should vary linearly if potential and conductivity are plotted logarithmically (see Equations (11)–(13)).

From Eq. (7), we have:

### Table 1 Form factors of the specimens A1, A2, A3 and B1, B2, B3 as shown in Fig. 5, measured resistances in four-wire technique $R_m = U_{23}/I$ according to Eq. (4) and derived conductivities $\sigma_m$

| Specimen | $K_m/m^{-1}$ at 25 $^\circ C$ | $R_m/\Omega$ at 25 $^\circ C$ | $\sigma_m/10^{-4} \, Scm^{-1}$ at 25 $^\circ C$ |
|----------|-----------------------------|-------------------------------|---------------------------------|
| A1       | 1882                        | 53,704                        | 3.505                           |
| A2       | 2125                        | 56,086                        | 3.789                           |
| A3       | 857                         | 24,931                        | 3.437                           |
| B1       | 1265                        | 32,563                        | 3.885                           |
| B2       | 1861                        | 43,757                        | 4.253                           |
| B3       | 1243                        | 30,499                        | 4.076                           |
\[ U_s \sigma_s = \text{const.} \quad (11) \]

Then, we have:

\[ \log(U_s \sigma_s) = \log(\text{const.}) \quad (12) \]

From here, let us assume \( \log(\text{const.}) = c \), as \( c \) is a constant

\[ \log(U_s) + \log(\sigma_s) = c \quad (13) \]

This limitation can be circumvented by setting \( \sigma_s \) to be \( 10^5 \) times higher (for example, 1000 \( \text{Sm}^{-1} \)). The simulated potential difference should then be divided by \( 10^5 \) for further calculations. However, \( \sigma_s \) should be set at least three orders smaller than the conductivity of the electrodes in the simulation, otherwise the region of the electrode would not be an equipotential area and the assumptions for the calculation are not valid anymore. This is also the reason for the second limitation.

It should also be pointed out that in the shown example, a 2D method is used, since only the surface with the electrodes is photographed and the thickness of the specimen is assumed as constant along the length and width direction. For specimens with non-uniform thickness, the thickness profile should be taken into consideration during the extrusion of the 2D CAD sketch. Moreover, the postprocessing of the edge image into a CAD file is not yet fully automated.

**Conclusion and outlook**

A novel method is presented in this study that enables a precise and reliable measurement of the conductivity of a block-shaped or plate-shaped semiconductor using four-wire-technique whose electrodes have arbitrary form. It can be especially applied for high temperature conductivity measurements of ceramics. This initial work used a 2D model for planar samples. The basic idea is to reconstruct the arbitrary-shaped electrodes and the block-shaped specimen (here obtained by a camera image of the sample) into a CAD file. After that, a form factor (geometry factor) is calculated with the help of FE software. The conductivity can then be derived from the resistance measurement, preferably in four-wire technique. This method can save experimentalist time, as most of the steps in the method is automated and one does not have to spend much time ensuring that the electrodes are thin enough and/or parallel to each other in order to obtain precise results. For further development of this method, the postprocessing of the edge image into a CAD file, the import of the model and the assignment of the boundary conditions in the FE software should be fully automated. One idea for a fully automated postprocessing of the edge image is to use machine learning to detect the desirable features. An application interface can
also be written to import the CAD file into the FE software and to assign the boundary conditions automatically. Moreover, by combining this method with computer topography (CT), it can also be applied for macroporous semiconductor specimens, e.g., conductive oxides. The ultimate goal is to develop a compact application so that conductivity measurement of semiconductors can be reduced to just a four-wire measurement of the resistance and taking an image of the specimens.

Supplementary Information: The online version contains supplementary material available at (http:s://doi.org/10.1007/s10853-021-05949-4).

Acknowledgements

The authors thank Vishay Electronic GmbH, Selb, Germany, for providing the ceramic samples. The authors thank Mr. Robin Werner for supporting sample preparation.

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Funding

Open Access funding enabled and organized by Projekt DEAL.

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