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Microbeam analysis — Analytical electron microscopy — <u>—</u> Selected area electron diffraction analysis using a transmission electron microscope

<u>Analyse par microfaisceaux — Microscopie électronique analytique — Analyse par diffraction par sélection</u> <u>d'aire au moyen d'un microscope électronique en transmission</u>

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Foreword

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The procedures used to develop this document and those intended for its further maintenance are described in the ISO/IEC Directives, Part 1. In particular, the different approval criteria needed for the different types of ISO document should be noted. This document was drafted in accordance with the editorial rules of the ISO/IEC Directives, Part 2 (see www.iso.org/directives).

ISO draws attention to the possibility that the implementation of this document may involve the use of (a) patent(s). ISO takes no position concerning the evidence, validity or applicability of any claimed patent rights in respect thereof. As of the date of publication of this document, ISO had not received notice of (a) patent(s) which may be required to implement this document. However, implementers are cautioned that this may not represent the latest information, which may be obtained from the patent database available at <a href="https://www.iso.org/patents.com/patents.co

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For an explanation of the voluntary nature of standards, the meaning of ISO specific terms and expressions related to conformity assessment, as well as information about ISO's adherence to the World Trade Organization (WTO) principles in the Technical Barriers to Trade (TBT), see <u>www.iso.org/iso/foreword.html</u>www.iso.org/iso/foreword.html.

This document was prepared by Technical Committee ISO/TC 202, Microbeam, Subcommittee SC 3, Analytical electron microscopy.

ISO/FDIS 25498

This third edition cancels and replaces the second edition (ISO 25498:2018), which has been technically e83428e3511/iso-fdis-25498 revised.

The main changes are as follows:

- Scope has been revised;
- ISO/IEC 17025 has been moved from normative references to -bibliography;
- <u>figure 1</u> has been replaced;
- subclauseSubclause 6.3 has been deleted;

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- <u>subclauseSubclause</u> 8.3.6 has been deleted, the content of 8.3.6 has been moved to 8.3.2;8.3.2;
- subclause 9.2.5<u>Subclause 9.2.5</u> has been added and the following subclause has been renumbered;
- <u>clause 11Clause 11</u> has been revised, <u>subclauses 11.1,11.2,11.3</u> and <u>11.411.4</u> have been added;

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<u>subclauses B.4.1Subclauses B.4.1</u> and <u>B.4.2B.4.2</u> have been added;

— <u>the bibliographyBibliography</u> has been updated and ISO/IEC Guide 98-3 (GUM:1995) has been added.

Any feedback or questions on this document should be directed to the user's national standards body. A complete listing of these bodies can be found at <u>www.iso.org/members.html</u>www.iso.org/members.html

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Introduction

Electron diffraction techniques are widely used in transmission electron microscopy (TEM) studies. Applications include phase identification, determination of the crystallographic lattice type and lattice parameters, crystal orientation and the orientation relationship between two phases, phase transformations, habit planes and defects, twins and interfaces, as well as studies of preferred crystal orientations (texture). While several complementary techniques have been developed, for example microdiffraction, nano-diffraction, convergent beam diffraction and reflected diffraction, the selected area electron diffraction (SAED) technique is the most frequently employed.

This technique allows direct analysis of small areas on thin specimens from a variety of crystalline substances. It is routinely performed on most TEMs in the world. The SAED is also a supplementary technique for acquisition of high-resolution images, microdiffraction or convergent beam diffraction studies. The information generated is widely applied in studies for the development of new materials, improving structure and/or properties of various materials as well as for inspection and quality control purpose.

The basic principle of the SAED method is described in this document. The experimental procedure for the acquirement of SAED patterns, indexing of the diffraction patterns and determination of the diffraction constant are specified. ISO 25498 is intended for use or reference as technical regulation for transmission electron microscopy.

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DRAFT International Standard

Microbeam analysis-_— Analytical electron microscopy — Selected area electron diffraction analysis using a transmission electron microscope

1 Scope

This document specifies the method <u>offor</u> selected area electron diffraction (SAED) analysis using a transmission electron microscope (TEM) to analyse thin crystalline specimens. This document applies to test areas of micrometres and sub-micrometres in size. The minimum diameter of the selected area in a specimen which can be analysed by this method is restricted by the spherical aberration coefficient of the objective lens of the microscope and approaches <u>hundredhundreds</u> of nanometres for a modern TEM.

When the size of an analysed specimen area is smaller than that the spherical aberration coefficient restriction, this document can also be used for the analysis procedure. However, because of the effect of spherical aberration and deviation of the specimen height position, some of the diffraction information in the pattern can be generated from outside of the area defined by the selected area aperture. In such cases, the use of microdiffraction (nano-beam diffraction) or convergent beam diffraction, where available, can be preferred.

This document is applicable to the acquisition of SAED patterns from crystalline specimens, indexing the patterns and calibration of the camera constant.

2 Normative references

There are no normative references in this document.

3 Terms, definitions and abbreviated terms abbreviations

For the purposes of this document, the following terms and definitions apply. IS 25498

ISO and IEC maintain terminology databases for use in standardization at the following addresses:

- ISO Online browsing platform: available at https://www.iso.org/obp
- — IEC Electropedia: available at <u>https://www.electropedia.org/</u>https://www.electropedia.org/

3.1 Terms and definitions

3.1.1 3.1.1

Miller notation

indexing system for crystallographic planes and directions in crystals, in which a set of lattice planes or directions is described by three axes coordinate

3.1.2 3.1.2

Miller-Bravais notation

indexing system for crystallographic planes and directions in hexagonal crystals, in which a set of lattice planes or directions is described by four axes coordinate

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3.1.3 3.1.3

interplanar spacing

-d_{hki}

 d_{hkl} perpendicular distance between consecutive planes of the crystallographic plane set (*hkl*)

3.1.4 3.1.4

reciprocal vector

*G*_{hkl}

 g_{hkl} vector in the reciprocal lattice

Note-1-to-entry:-_The reciprocal vector, $-g_{hkl}$, g_{hkl} is normal to the crystallographic plane (*h k l*) with its magnitude inversely proportional to *interplanar spacing* $-d_{hkl}$ (3.1.3). $-d_{hkl}$ (3.1.3).

3.1.5 3.1.5

R vector -R_{hkl}

R_{hkl}

coordinate vector from the direct beam, 000, to a diffraction spot, hkl, in a zone diffraction pattern

Note 1-_to-_entry:_See Figure 1.Figure 1.

3.1.6 3.1.6

camera length

effective distance from the specimen to the screen or recording device in a transmission electron microscope in diffraction mode-

3.1.7 3.1.7

camera constant diffraction constant

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 $L\lambda$ product of the wavelength of the incident electron wave and *camera length* (3.1.6)(3.1.6)

[SOURCE: ISO 15932:2013, 3.7.1]

3.1.8 3.1.8

bright field image

image formed using only the non-scattered beam, selected by observation of the back focal plane of the objective lens and using the objective aperture to cut out all diffracted beams

[SOURCE: ISO 15932:2013, 5.5]

3.1.9 3.1.9

dark field image

image formed by a diffracted beam only by using the objective aperture for selection or by collecting the diffracted beams with an annular dark-field detector

[SOURCE: ISO 15932:2013, 5.6]

2-

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3.1.10 3.1.10

energy-dispersive X-ray spectrometry

EDS

analytical technique which enables the elemental analysis or chemical characterization of a specimen by analysing characteristic X-ray emitted by the matter in response to electron irradiation

[SOURCE: ISO 15932:2013, 6.6]

3.1.11 3.1.11

eucentric position specimen position at which the image exhibits minimal lateral motion resulting from specimen tilting

3.1.12 3.1.12

selected area- (selector) aperture

moveable diaphragm that is used to select only radiation scattered from a specific area of the specimen to contribute to the formation of a diffraction pattern

[SOURCE: ISO 15932:2013, 3.2.3.5]

3.1.13 3.1.13

Bragg angle 0₽

 $\theta_{\mathbf{B}}$

angle between the incident beam and the atomic planes, at which diffraction takes place

3.2 Abbreviated terms and symbols

BCC	body-centred cubic structure	

- FCC face-centred cubic structure
- HCP hexagonal close-packed structure
- selected area electron diffraction SAED
- TEM transmission electron microscope 713de-bc0e-4451-be7c-dc83428e3511/iso-fdis-25498 (hkl) Miller indices of a specific set of crystallographic planes

- Miller indices which denote a family of crystallographic planes $\{hkl\}$
- Miller indices of a specific crystallographic direction or a zone axis [uvw]
- Notation for a set of planes in the reciprocal lattice $(uvw)^*$

NOTE- The normal of the reciprocal plane (uvw)* is parallel to the crystallographic zone axis [uvw]

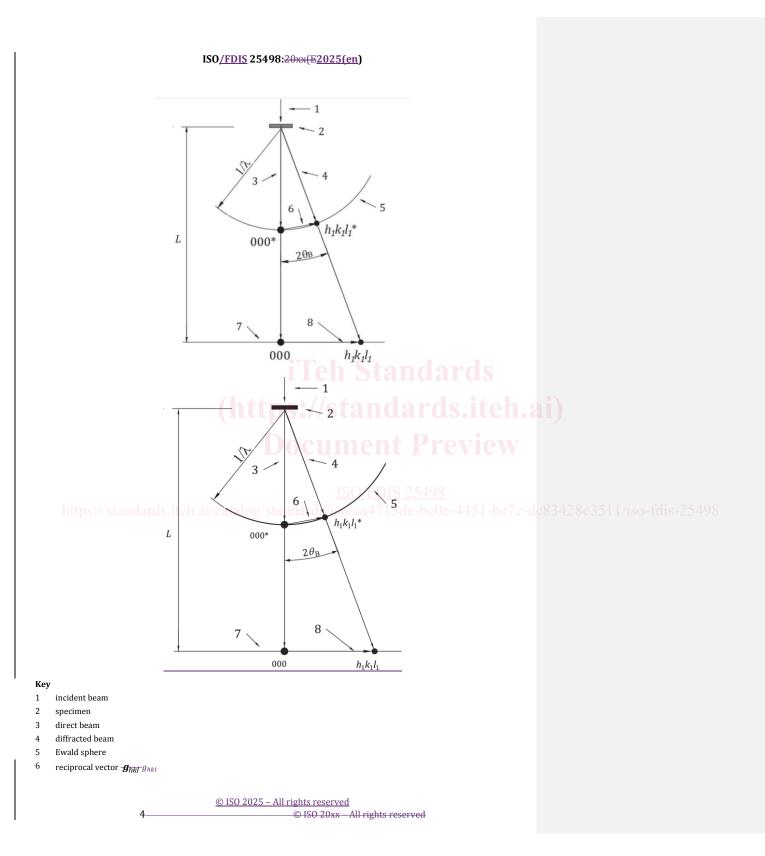
Principle 4

4.1 General

When an energetic electron beam is incident upon a thin crystal specimen in a transmission electron microscope, a diffraction pattern will be produced in the back focal plane of the objective lens. This pattern is magnified by the intermediate and projector lenses, then displayed on a viewing screen and recorded (see Reference [3], [4], [5]). [3], [4], [5]). This pattern can also be displayed on a monitor if the TEM is equipped with a digital camera system.

The geometric relationship of the parameters for selected area electron diffraction (SAED) technique can be understood through the Ewald sphere construction, which is illustrated in Figure 1. Figure 1.

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7 diffraction pattern

8-Rhkl Rhkl vector

L is the diffraction camera length;

 $\theta_{\mathbb{B}}\underline{\theta}_{\mathbb{B}}$ is Bragg angle;

 λ is the wavelength of the incident electron beam.

Figure 1-_— Ewald sphere construction illustrating the diffraction geometry in TEM

4.2 Spot diffraction pattern

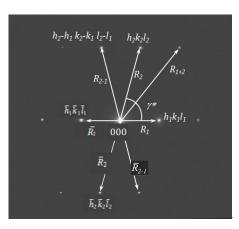
The diffraction pattern of a single crystal appears as an array of "spots", the basic unit of which is characterized by a parallelogram. An example of the spot diffraction pattern is shown in Figure 2. Figure 2. Each spot corresponds to diffraction from a specific set of crystal lattice planes in the specimen, denoted by Miller indices (*hkl*). The vector, $R_{nkl}R_{hkl_{\perp}}$ is defined by the position of the diffracted spot, *hkl*, relative to position on the pattern corresponding to the direct beam, i.e. the centre-spot, 000, of the pattern. It is parallel to the normal of the reflecting plane, (*hkl*). The magnitude of $R_{nkl}R_{nkl}$ is inversely proportional to the interplanar spacing, $d_{nkl}d_{hkl}$, of the diffracting plane, (*hkl*) (see References [4][4] to [9])-[9]). In the context of this document, vectors $R_{n_1k_1l_1}$, $R_{h_2k_2l_2}$, $(R_{h_2k_2l_2}, R_{h_1k_1l_1})$, $R_{h_1k_1l_1}$, $R_{h_2k_2l_2}$, $(R_{h_2k_2l_2}, R_{h_1k_1l_1})$ and $(R_{n_1k_1l_1} + R_{h_2k_2l_2})$ ($R_{h_1k_1l_1} + R_{h_2k_2l_2}$) are simplified as R_1 , R_2 , $R_{2-T}R_{2-1}$ and $R_{1+2}R_{1+2}$ respectively. The included angle between vectors, R_1 and R_2 , is denoted by γ^* . The basic parallelogram is defined by R_1 and R_2 , where they are the shortest and next shortest in the pattern respectively and not along a common line. The spot, $h_2k_2l_2$, is positioned anticlockwise around the centre spot relative to spot, $h_1k_1l_1R_1k_1l_1$.

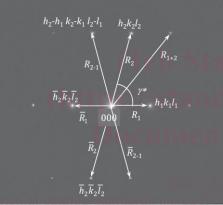
Because the centre-spot is often very bright, it is often difficult to determine the exact centre of the pattern. Therefore, a practical procedure is to establish the magnitude of $|\mathbf{R}_{hkl}| |R_{hkl}|$ by measuring the distance between the spots, *hkl* and $\overline{hkl} hk\bar{l}$ on the diffraction pattern and dividing by two, i.e.

 $\frac{|R_{hkl}|}{2} = \frac{1}{2} \left(\frac{|R_{hkl}| + |R_{hkl}|}{2} + \frac{1}{2} \left(|R_{hkl}| + |R_{h\bar{k}\bar{l}}| \right) \right)$ On the example pattern shown in Figure 2, Figure 2, the

magnitude of R_1 , R_2 and $R_{2-1}R_{2-1}$ is obtained from $\frac{1}{2}(R_1 + \bar{R}_1)$, $\frac{1}{2}(R_2 + \bar{R}_2)^{\frac{1}{2}}(R_1 + \bar{R}_1)$, $\frac{1}{2}(R_2 + \bar{R}_2)$ and $R_3428e3511/iso-fdis-25498$. $\frac{1}{2}(R_{2-1} + \bar{R}_{2-1})^{\frac{1}{2}}(R_{2-1} + \bar{R}_{2-1})$ respectively.

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 Key

 R_1 is the vector from 000 to spot, $h_1k_1h_1$, the shortest vector in the diffraction pattern

 R_2 Is in evector from 000 to spot, $h_2k_2l_2$, the next shortest vector

NOTE The basic parallelogram is constituted by R_1 and R_2 .

Figure 2 — Example of the spot diffraction pattern from a single crystal

The relationship between the interplanar spacing, d_{nkl} , d_{nkl} , and the magnitude of $R_{nkl}R_{hkl}$ for a reflecting plane, (*hkl*), can be approximately expressed as shown in Formula (1) (see References [7][7] and [8]):[8]):

$$\frac{L\lambda = R_{hkl} \times d_{hkl} \left[1 - \frac{3}{8} (R_{hkl} / L)^2 \right] = R_{hkl} \times d_{hkl} (1 - \Delta)$$

$$(1)$$

$$L\lambda = R_{hkl} \times d_{hkl} [1 - \frac{3}{8} (R_{hkl} / L)^2] = R_{hkl} \times d_{hkl} (1 - \Delta)$$

$$(1)$$

where