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Surface chemical analysis electron spectroscopy and X-1., photoelectron spectroscopy – Methods used to determine peak intensities and information required whimique des surfaces – Spectroscopie des électrons Aug "élédebédectrons per rayons X – Méthodes used "élédebédectrons sité des pics et informations required "élédebédectrons sité des pics et informations required

et spectroscopie de photoélectrons par rayons X — Méthodes utilisées pour la détermination de l'intensité des pics et informations requises

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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 documents 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).

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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</u> .org/iso/foreword.html.

This document was prepared by Technical Committee ISO/TC 201, *Surface chemical analysis*, Subcommittee SC 7, *Electron spectroscopies*.

This third edition cancels and replaces the second edition (ISO 20903:2011), which has been technically revised. The main changes compared to the previous edition are as follows:

- subclause <u>6.3</u> has been replaced to include modern methods for dealing with co-existing chemical states;
- minor editorial changes have been introduced for clarity.

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>.

Introduction

An important feature of Auger electron spectroscopy (AES) and X-ray photoelectron spectroscopy (XPS) is the ability to obtain a quantitative analysis of the surface region (\approx 1 nm to 10 nm) of a solid sample. Such an analysis requires the determination of the intensities of spectral components.

ify if in methods in a method There are several methods of peak-intensity measurement that are applicable to AES and XPS. In practice, the choice of method will depend upon the type of sample being analysed, the capabilities of the instrumentation used, and the methods of data acquisition and treatment available.

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Surface chemical analysis — Auger electron spectroscopy and X-ray photoelectron spectroscopy — Methods used to determine peak intensities and information required when reporting results

Scope 1

This document specifies the necessary information required in a report of analytical results based on measurements of the intensities of peaks in Auger electron and X-ray photoelectron spectra. Information on methods for the measurement of peak intensities and on uncertainties of derived peak areas is also provided.

2 Normative references

The following documents are referred to in the text in such a way that some or all of their content constitutes requirements of this document. For dated references, only the edition cited applies. For undated references, the latest edition of the referenced document (including any amendments) applies.

ISO 18115-1, Surface chemical analysis - Vocabulary - Part 1: General terms and terms used in spectroscopy

Terms and definitions 3

For the purposes of this document, the terms and definitions given in ISO 18115-1 apply.

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

- ISO Online browsing platform: available at https://www.iso.org/obp
- IEC Electropedia: available at http://www.electropedia.org/

Symbols and abbreviated terms 4

- Α peak area
- AES Auger electron spectroscopy
- b number of channels over which intensities are averaged to obtain a baseline
- eV electron volts
- number of channels in a spectrum n
- XPS X-ray photoelectron spectroscopy
- number of counts in the *i*th channel of a spectrum Уi
- ΔE channel width (in electron volts)
- dwell time per channel (in seconds) Δt
- standard deviation of calculated peak area $\sigma(A)$