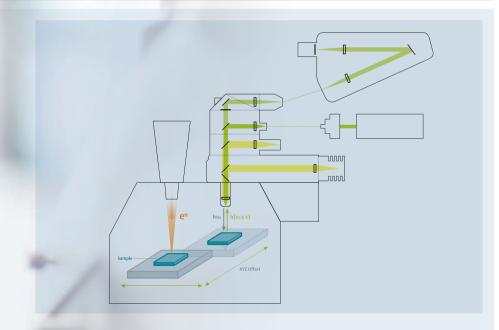
a) SEM image of a geological sample (diorite). b) SEM image overlaid with the Raman image. The different colors in the Raman image illustrate the various molecular compounds. Raman image: 100 x 100 μ m², 300 x 300 pixels = 90,000 spectra, integration time: 34 ms/spectrum.

- c) The corresponding color-coded Raman spectra display each molecular component of the sample. (1) Sample courtesy of Dr. Christine Heim, Geowissenschaftliches Zentrum, University of Goettingen, Germany
- a b

a) SEM image of CVD grown MoS₂ on Si/SiO₂ acquired with the SE (secondary electron) detector. The MoS₂ crystals grow in a triangular shape due to the symmetry of the unit cell of the molecule. Bright areas in the SEM image correspond to borders or overlapping regions of the MoS₂ crystals. b) RISE image of the MoS₂ crystals. This image was obtained by overlaying the Raman image (image parameters: 20 x 20 μ m², 100 x 100 pixels = 10,000 spectra, integration time 0.15 s/spectrum) onto the SEM image. The RISE image demonstrates that the Raman spectra of MoS₂ are extremely sensitive to borders (red) and overlapping (blue) regions. (2)

RISE microscopy done with SEMs from (1) Tescan Brno s.r.o, Czech Republic, or (2) Carl Zeiss Microscopy GmbH, Germany



Principle of RISE Microscopy

For RISE microscopy samples are automatically transferred from one measuring position to the other within the vacuum chamber of the SEM for the entirety of the measurement procedure, thus streamlining the workflow and drastically improving the instrument's ease of use. The beampath of the Raman microscope is shown in light green.

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拉曼成像与扫描电镜联用

RISE Microscopy

显微联用技术的全新时代同时提供化学信息和超结构信息



RISE Microscopy

Fully-integrated Raman Imaging and Scanning Electron Microscopy



Correlative Scanning Electron and confocal Raman imaging for comprehensive sample analysis

A new dimension in imaging: ultra-structural SEM complemented with chemical compound information and molecular Raman imaging



RISE Microscopy is well suited for ...

...Raman newcomers as they will benefit from the intuitive user interface and straighter measurement procedure.

...experienced users as they will appreciate the exceptional correlative microscope performance encompassing the advantages of both techniques in one instrument.

Materials science, nanotechnology, polymer science, geosciences, life sciences, pharmaceutical industry ...

...are among the fields that can benefit from RISE microscopy with its unique imaging capabilities.

WITec Headquarters

for profilometry

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RAMAN IMAGING

Inelastic scattering of light from a molecule.



Example Raman spectrum of cellulose.

The Raman Principle

- Raman spectroscopy is a well-established and non-destructive method to analyze the chemical composition of a sample.
- A Raman spectrum shows the energy shift of the excitation light (laser) as a result of inelastic scattering by the molecules in a sample.
- Each molecule and chemical compound results in a specific Raman spectrum and can be easily identified by its unique Raman 'fingerprint'.

Additional sample information from the Raman spectrum:

- a. Peak intensity: Quantity/amount of a specific compound b. Peak shift: Identification of stress and strain states
- c. Peak width: Degree of crystallinity
- d. Polarization state: Crystal symmetry and orientation









Confocal Raman Imaging

- The WITec confocal Raman microscopy and imaging system combines Raman spectroscopy with confocal microscopy and enables confocal Raman imaging with the information of a complete Raman spectrum at every image pixel and a lateral resolution at the diffraction limit (~200 nm).
- Confocal Raman imaging with unprecedented performance in speed, sensitivity, and resolution
- Outstanding depth resolution ideally suited for 3D image generation and depth profiles
- Ultrahigh-throughput lens-based spectroscopic system for highest sensitivity and best performance in spectral resolution
- Ultra-fast Raman imaging option with only 0.76 ms integration time per spectrum
- Non-destructive imaging technique: no staining of fixation of the sample required

Fully integrated confocal Raman microscope with excellent imaging capabilities and outstanding performance in speed, sensitivity, and resolution.

Once positioned on the

scan table, the sample is

automatically transferred

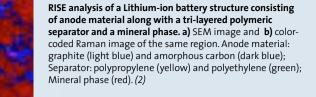
and re-positioned between

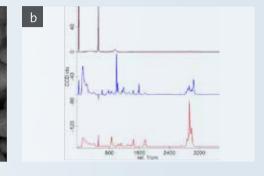
the measuring procedures.

The GM chamber is designed to integrate the Raman extension for RISE measurements for extended analytical potential.

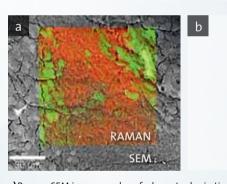
a) RISE image of labeled micro-particles. The SEM image was acquired with the backscattered electron detector and reveals patterned micro-spheres. A small area was measured in Raman imaging mode (image parameters: 30 x 30 μ m², 150 x 150 pixels = 22,500 spectra, integration time 0.05 s/ spectrum) and overlaid onto the SEM image, leading to the color patterned area in the RISE image. (1)

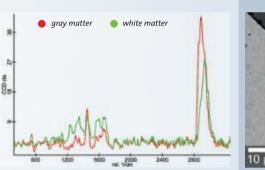
APPLICATIONS



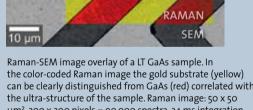


b) The two different colors correspond to polyethylene (red) and polystyrene (blue) as highlighted in the Raman spectra acquired from the sample. The substrate for the





a) Raman-SEM image overlay of a hamster brain tissue sample. In the color-coded Raman image the white brain matter is shown in green and the gray brain matter in red. Raman Image: 100 x 100 µm²,



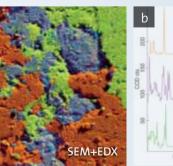
the color-coded Raman image the gold substrate (yellow) can be clearly distinguished from GaAs (red) correlated with the ultra-structure of the sample. Raman image: 50 x 50 μ m², 300 x 300 pixels = 90,000 spectra, 34 ms integration time per spectrum. (1)

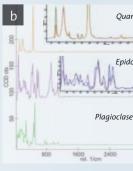
COMPARISON BETWEEN RISE MICROSCOPY AND ENERGY-DISPERSIVE X-RAY SPECTRO

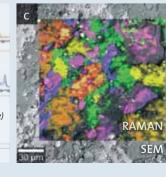
300 x 300 pixels = 90,000 spectra, 50 ms integration time per spectrum. b) The corresponding

Raman spectra reveal the different spectral characteristics of the white and gray brain matter. (1)

RISE Microscopy and EDX analysis of a geological sample: a) Overlaid SEM-EDX image: Three different element groups can be distinguished (Si, O: orange; Si, Al, Fe, Ca: grey; Na: green). **b)** Raman spectral imaging of same sample area (22,500 spectra; integration time: 0.08 s spectrum): Three spectral clusters can be differentiated (quartz, epidote, albite). The inserts show spectral variations through different mineral orientations. c) Overlaid color-coded Raman-SEM image shows the distribution of the molecular compounds quartz in different orientations (yellow/orange) epidote in different orientations (blue/purple) albite (green). (1)







SCANNING ELECTRON MICROSCOPY

RISE Microscopy – The Instrument

The microscope combines all features of a stand-alone SEM and a top-class confocal Raman imaging microscope within one instrument.

- Quick and convenient switching between Raman and SEM measurement
- Automated sample transfer from one measuring position to the other
- Integrated software interface for user-friendly measurement control
- Correlation of the measurement results and image overlay
- No compromise in SEM and Raman imaging capabilities



The sample remains inside the vacuum chamber during the complete measurements to ensure a convenient work flow with ease-of-use.

RISE microscopy done with SEMs from (1) Tescan Brno s.r.o, Czech Republic, or (2) Carl Zeiss Microscopy GmbH, Germany