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Micro-XRF

Sample Preparation of Nanocomposites and Nanomaterials by *Ultramicrotomy*

^a Powerful **Alternative**

Join us at the **EMS Microscopy Academy** and learn the latest techniques to reveal internal structures of composites and polymers being investigated with transmission electron microscopy (TEM) and scanning transmission electron microscopy (STEM).

Sample preparation workflow will be illustrated using the Leica EM UC7 Ultramicrotome, its EM FC7 Cryochamber, and the RMC PowerTome Ultramicrotome. Differences between FIB (Focussed Ion Beam) and ultramicrotomy samples will also be covered.

Who can benefit from this alternative?

- Composite and polymer research companies - especially from the automotive and aviation industries
- Materials scientists already working with ultramicrotomy
- FIB users preparing TEM lamellas

For more information, or to sign up for a workshop, please visit our website...

www.emsdiasum.com

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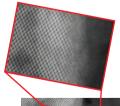
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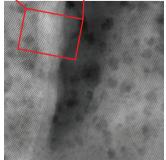


Applications...

Zeolite USY30 Crystal morphology STEM analysis

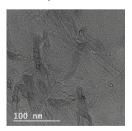


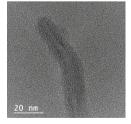
The mesopores (2-50 nm) and the crystalline micro-pores (0.7 nm) are clearly visualized.



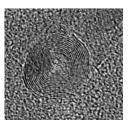
Tom Willhammar, Sara Bals, **EMAT Antwerpen** Cambridge University Press

Epoxy loaded with amino-functionalized CNTs TEM analysis

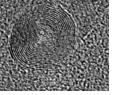




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Mert Kurttepeli, Sara Bals, **FMAT Antwernen**

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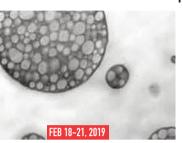
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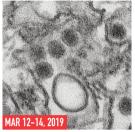
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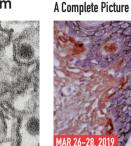
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Materials Ultramicrotomy Workshop



Biological TEM Workshop: A Complete Picture

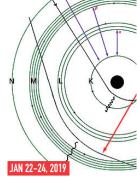


Aurion Immunogold Silver Staining

JAN 15-17, 2019

Sample Preparation for

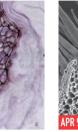
Semiconductor Devices:



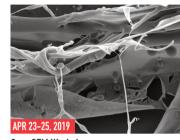
X-Ray Microanalysis Workshop: A Complete Picture



Introduction to Microscopy Techniques Workshop



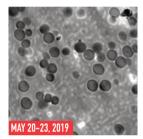
Biological SEM Workshop: A Complete Picture



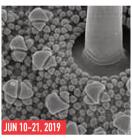
Cryo SEM Workshop



Automated and Rapid Specimen Processing for Electron Microscopy Workshop



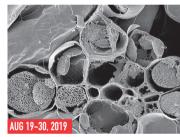
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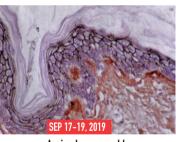
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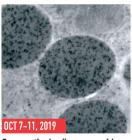
Microscopy: The Complete Image



Microscopy: The Complete Image



Aurion Immunogold Silver Staining



Cryosectioning/Immunogold Workshop



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Biological TEM Workshop: A Complete Picture

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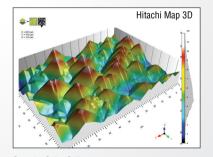
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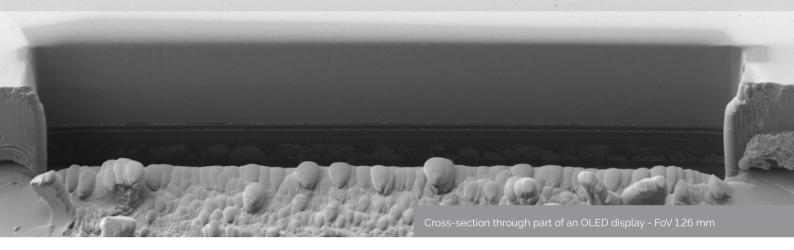
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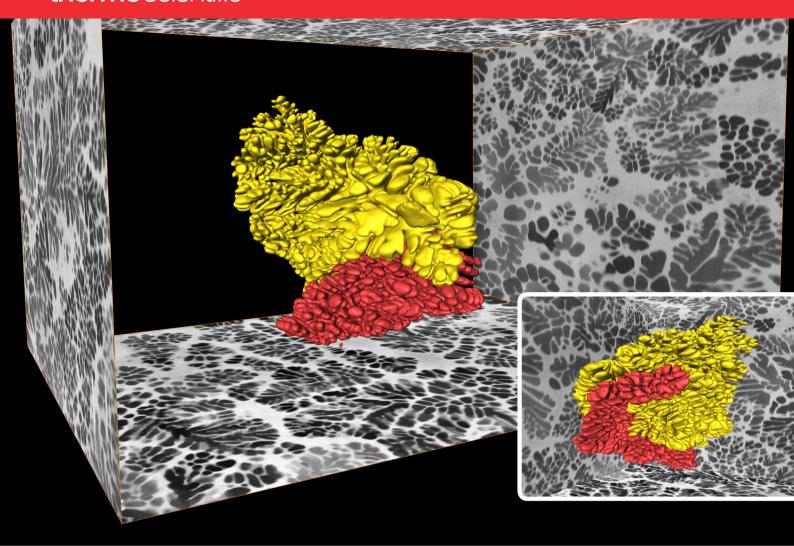
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Thermo ScientificTM AvizoTM 3D visualization of two large adjacent crystalline dendrites of a bulk-metallic-glass matrix composite $(Zr_{s8.5}Ti_{14.3}Nb_{s.2}Cu_{s.1}Ni_{4.9}Be_{11.0})$. Data was obtained by large volume serial sectioning tomography using the Thermo ScientificTM HeliosTM PFIB DualBeamTM microscope. The sectioned block is about $90 \times 80 \times 70 \ \mu m^3$. Sample from The University of Tennessee, USA. Images courtesy of The University of Manchester.

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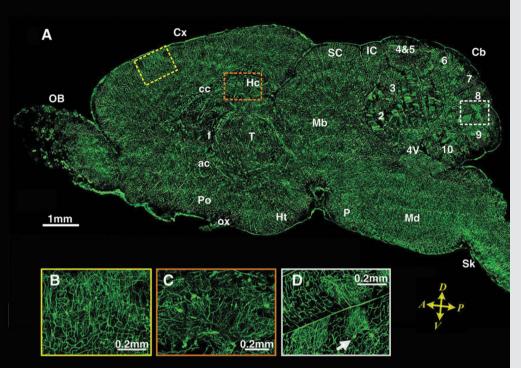
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Creating a High Resolution Atlas of the Mouse Brain...

(A) A sagittal image reconstructed from a stack of 100 virtual sagittal sections (total thickness of 0.1 mm). These sections were transformed from the original coronal sections. The sagittal image was located in the right hemisphere about 0.4 mm lateral to the middle. Almost all major regions of the brain can be seen in this image, e.g., the Olfactory Bulb (OB), Cerebral Cortex (Cx), Hippocampus (Hc), Fornix(f), Anterior Commissure (ac), Thalamus (T), Cerebellum (Cb), Midbrain (Mb), Pons (P), Medulla (Md), Corpus Callosum (cc), Superior Colliculus (SC), Inferior Colliculus (IC), Hypothalamus (Ht), Preoptic Area (Po), Optic Chiasm (ox), 4th ventricle (4V) and nine lobules of the cerebellum (Arabic numerals, 2 to 10). The three regions inside the different colored rectangle in (A) are the positions of (B), (C) and (D), which illustrate the cerebral cortex, hippocampus and cerebellum, respectively. In the reconstruction of sagittal image, no dislocation was observed along the D-V axis, i.e., the coronal sections are inherently aligned along the A-P axis.



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Micro-Optical Sectioning Tomography to Obtain a High-Resolution Atlas of the Mouse Brain

Existing imaging tools have limitations for brainwide mapping of neural circuits at a mesoscale level. In collaboration with DiATOME, researchers developed a Micro-Optical Sectioning Tomography (MOST) system utilizing a DiATOME Diamond Knife that can provide micron tomography of a centimeter-sized whole mouse brain.

Slicing was performed by moving the specimen to generate ribbons, and each ribbon was simultaneously imaged. The illuminating beam passed through a beam splitter, mirror and objective to irradiate the ribbon. The imaging beam collected by the objective and passed through the mirror, beam splitter and tube lens was then recorded by a line-scan CCD.

A 3D structural dataset of a Golgi-stained whole mouse brain at the neurite level was obtained. The morphology and spatial locations of neurons and traces of neurites were clearly distinguished. Researchers found that neighboring Purkinje cells were sticking to each other.

Acknowledgement

Micro-Optical Sectioning Tomography to Obtain a High-Resolution Atlas of the Mouse Brain Anan Li, Hui Gong, Bin Zhang, Qingdi Wang, Cheng Yan, Jingpeng Wu, Qian Liu, Shaoqun Zeng, Qingming Luo

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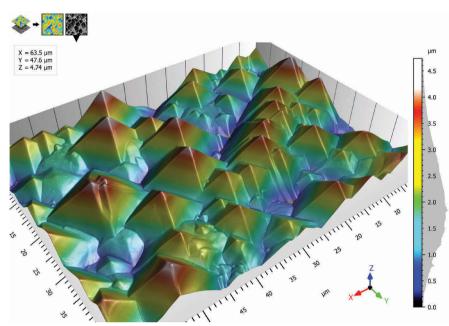




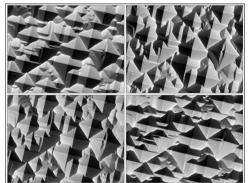
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ISO :	12781		
Flatn	ess Para	ime	ters
FLTt	5.40	μm	Peak-to-valley flatness deviation of the surface
FLTp	2.58		Peak-to-reference flatness deviation

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