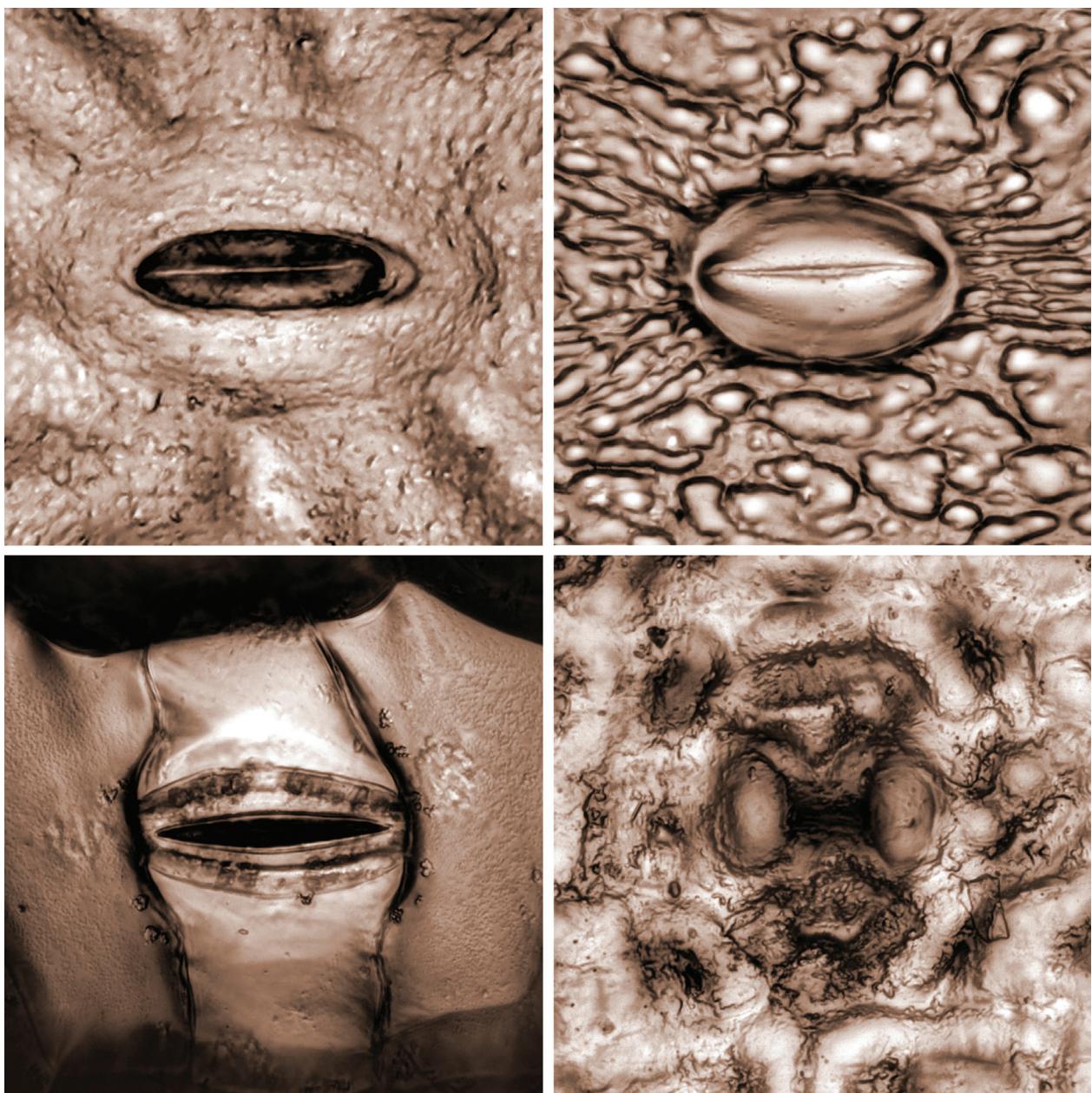


Microscopy

TODAY

Volume 27 Number 1 2019 January



ETHOS

Focused Ion and Electron Beam System

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The *All-New* Hitachi ETHOS SEM-FIB combines ultra-high resolution imaging and elemental analysis at low voltages with ion optics for nm-scale precision processing.



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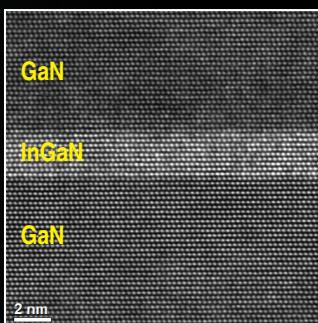
- Large analytical specimen chamber for numerous accessories
- Automated operation including macro processing
- Time sharing mode for dual simultaneous live imaging and processing
- Real-time analytical 3D segmentation capability

Advanced Microsampling

- Sample orientation control with Anti Curtaining Effect (ACE) technologies
- 4-axis lift-out function for advanced TEM specimen preparation

Triple-beam system yielding highest-quality results

- Low acceleration voltage processing with noble gas ion beam
- Selectable ion species (argon/xenon)



200 kV ADF STEM Image of processed lamella by Triple Beam Ar Ion at 1kV

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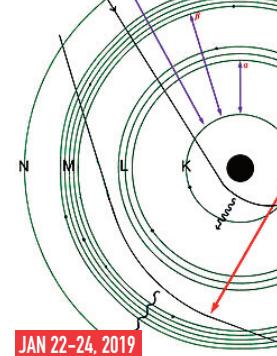
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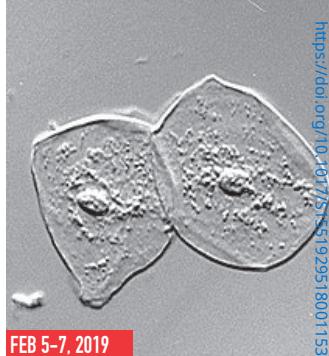
JAN 15-17, 2019

Sample Preparation for Semiconductor Devices:
A Complete Picture



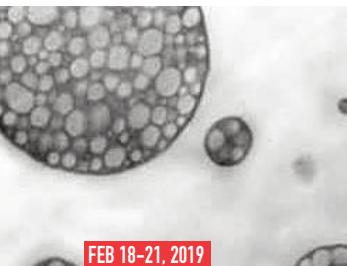
JAN 22-24, 2019

X-Ray Microanalysis Workshop:
A Complete Picture



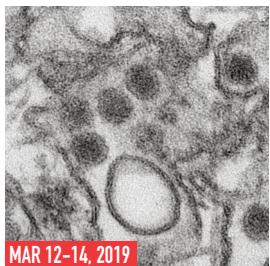
FEB 5-7, 2019

Introduction to Microscopy
Techniques Workshop



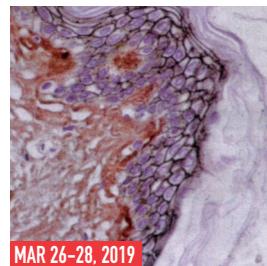
FEB 18-21, 2019

Materials Ultramicrotomy
Workshop



MAR 12-14, 2019

Biological TEM Workshop:
A Complete Picture



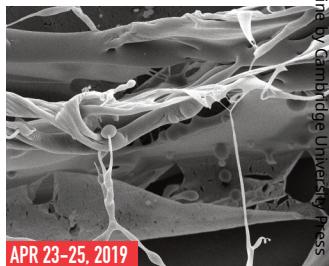
MAR 26-28, 2019

Aurion Immunogold
Silver Staining



APR 9-11, 2019

Biological SEM Workshop:
A Complete Picture



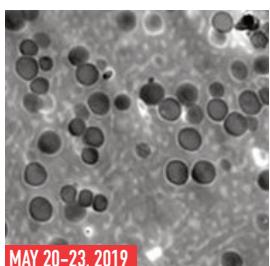
APR 23-25, 2019

Cryo SEM Workshop



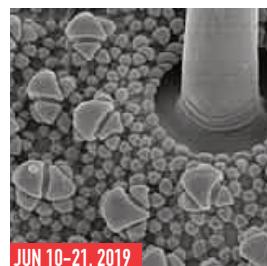
MAY 7-9, 2019

Automated and Rapid Specimen
Processing for Electron
Microscopy Workshop



MAY 20-23, 2019

Materials Ultramicrotomy
Workshop



JUN 10-21, 2019

Microscopy: The Complete Image



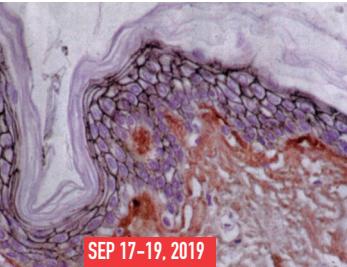
JUL 15-26, 2019

Microscopy: The Complete Image



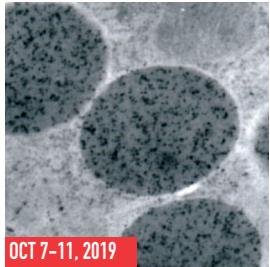
AUG 19-30, 2019

Microscopy: The Complete Image



SEP 17-19, 2019

Aurion Immunogold
Silver Staining



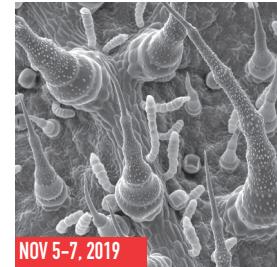
OCT 7-11, 2019

Cryosectioning/Immunogold
Workshop



OCT 22-24, 2019

Introduction to Microscopy
Techniques Workshop



NOV 5-7, 2019

Biological SEM Workshop:
A Complete Picture



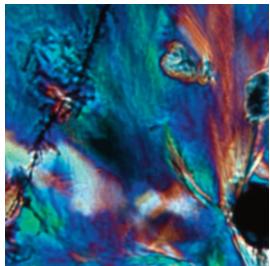
NOV 12-14, 2019

Biological TEM Workshop:
A Complete Picture

Plus: Pharmaceuticals Workshops,
dates to be determined...

- Pharmaceutical Microscopy Workshop
- Pharmaceutical Microscopy Workshop: Applications
- Pharmaceutical Chemical Imaging Workshop
- Pharmaceutical Microscopy Workshop: Polymorphism
- Pharmaceutical Microscopy Workshop: Techniques

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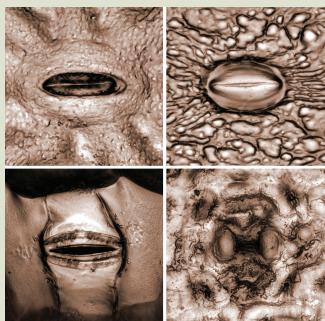
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Stomata of various plants. Clockwise from upper left: *Tradescantia*, *Masdevallia*, *Agave*, and *Paeonia*. Stomata. Image widths are 120 µm, 110 µm, 120 µm, and 70 µm, respectively. Images acquired by light microscopy using photo stacks.

See article by Clark.

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Microscopy Pioneers

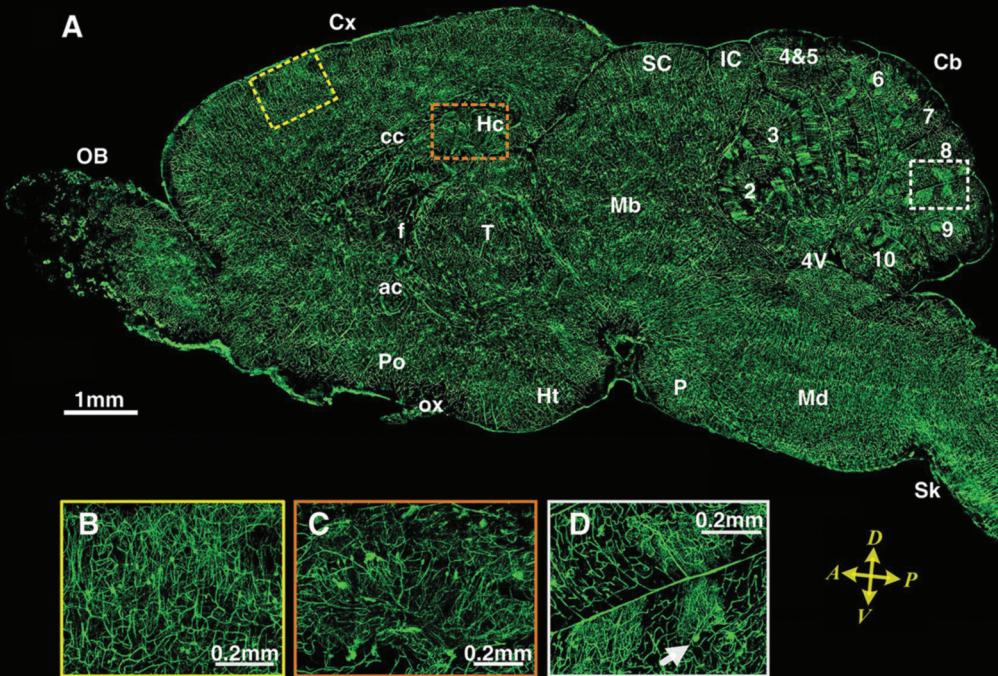
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YOU'LL FIND **DiATOME** AT THE FOREFRONT OF INNOVATION...



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.

DiATOME QUALITY AND INNOVATION APPLIED...

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
Britton Chance Center for Biomedical Photonics, Wuhan National Laboratory for Optoelectronics—Huazhong University of Science and Technology, Wuhan 430074, P. R. China.

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