

The MMS Scope

Minnesota Microscopy Society

Local affiliate of the Microscopy Society of America and the Microanalysis Society

April 2022

In this Issue:

Spring Symposium

Join us for the return of the in-person Spring Symposium on May 20 from 7:30am to 3:00pm. The theme this year is Sample Preparation.

Register soon and plan to participate.

MMS Member Dues

Please remember to submit your annual 2022 membership dues.

Past Events

View highlight slides from the virtual Winter Research Extravaganza, Hold the Mayo!

MINNESOTA MICROSCOPY SOCIETY ANNUAL SPRING SYMPOSIUM

FRIDAY, MAY 20, 2022



SYMPOSIUM SCHEDULE – ★ in person! ★ 7:30 a.m. – 3:00 p.m.

7:30 – 8:15 a.m. Registration, continental breakfast, vendor displays

8:15 – 8:30 a.m. Welcome

8:30 – 9:30 a.m. Katherine Igowsky, MNBCA

Sample Preparation in the Forensic Laboratory

9:30 - 10:30 a.m. Vern Robertson, JEOL

How Hydrocarbon Contamination and Sample Prep Techniques Affects Imaging and Microanalysis on an FEG

SEM/EPMA, Especially at Low kVs

10:30 – 11:30 a.m. Break and vendor displays 11:30 a.m. – 12:30 p.m. Lunch and vendor displays

12:30 – 12:45 p.m. Business meeting

12:45 - 1:45 p.m. Mary Buckett, 3M

From Smart Phones to Silicones — Sample Preparation from

an Industrial Lab Perspective

1:45 - 2:45 p.m. Shiba Dandpat, Lumicks USA

Real-time imaging of dynamic single-molecule interactions:

from DNA binding proteins to molecular motors

2:45 p.m. Raffle and social time



LOCATION

Science Museum of Minnesota Discovery Hall St. Paul, MN

➤ <u>Map</u>



120 W Kellogg Blvd, St Paul, MN 55102



REGISTRATION

Cost: \$75 for MMS members

Fee includes continental breakfast, boxed lunch (choose from 4 options at registration link below), and coffee/snacks.

➤ <u>Click to register</u>

PLEASE NOTE:

Attendees must follow the Science Museum's current COVID-19 safety measures and requirements to enter the building. During the event, well-fitted face masks worn over the nose and mouth will be required while in Discovery Hall except when eating and drinking.

"Sample Preparation" continued



SPEAKER BIOS & ABSTRACTS

Katherine Igowsky (Katie) has worked for the Bureau of Criminal Apprehension (BCA) for 17 years. She started out at the Minnesota BCA working in conjunction with the Regional FBI Mitochondrial DNA program as a hair examiner. She continues to work for the BCA in the Trace Evidence Section and serves as the technical leader. She analyzes and compares trace materials in criminal cases such as glass, tape, fibers, hair, using various microscopic techniques, including polarized light microscopy (PLM), Fourier transform infrared microscopy (FTIR), microspectrophotometry (MSP) and scanning electron microscopy (SEM/EDS). In addition to her laboratory duties, Katie also processed crime scenes as a member of the BCA crime scene team.



Katherine Igowsky

Sample Preparation in the Forensic Laboratory

Materials such as tape, hairs, glass, fibers, etc., are examined to establish connections in crimes. They can help establish that a suspect was at a crime scene, that a suspect was in contact with a victim, that the suspect's vehicle was at the crime scene, etc. These materials are analyzed using various microscopic techniques including polarized light microscopy (PLM), Fourier transform infrared (FTIR) microscopy, microspectrophotometry (MSP), and scanning electron microscopy (SEM/EDS). Preparation of this wide range of materials requires many different sample preparation techniques. This presentation will give an overview of some of the materials tested in the forensic laboratory, instrumental techniques used, sample preparation techniques, and how these types of analyses are used to solve crimes.

"Sample Preparation" continued



SPEAKER BIOS & ABSTRACTS

Vern Robertson has been with JEOL USA for more than 32 years and was appointed EPMA/Surface Analysis Product Manager in early 2016 and will continue as SEM Technical Sales Manager, providing in-house and in-the-field technical product support and customer applications support. Vern served as the senior SEM Applications Specialist at JEOL beginning in 1986. He was appointed National Laboratory Manager in 2004, and FEG SEM Product Manager in 2005. Vern received his B.Sc. in Geology from the University of New Hampshire. His prior industrial experience includes eight years of consulting in an independent testing lab specializing in industrial and environmental problem solving, with responsibilities including polarized light optical microscopy, and atomic emission and absorption spectroscopy SEM with EDS/WDS and X-ray diffraction. Vern was a recent member of the MAS (Microscopy).



Vern Robertson

EDS/WDS and X-ray diffraction. Vern was a recent member of the MAS (Microanalysis Society) Council.

How Hydrocarbon Contamination and Sample Prep Techniques Affects Imaging and Microanalysis on an FEG SEM/EPMA, Especially at Low kVs

As the scale of things we would like to image and analyze with an SEM or EPMA become smaller and smaller, there is a huge advantage to using an FEG SEM or FEG EPMA at low accelerating voltages and higher beam currents to maintain X-ray counting statistics, and reduce the excitation volume of where the BSE and X-rays are emitted from. The advantage is that you do not have a significant loss of probe diameter at those conditions. Using a low kV with a lower beam current allows one to image at an ultra-high resolution and surface sensitivity.

All of these benefits come with some serious complications and limitations. The first of which is carbon (often referred to as hydrocarbon) contamination. The other factors include sample preparation and sample conductive coatings. In this talk I will present ways to clean the C contamination that may be on the samples, which is sample dependent, and the best ways to prevent C contamination in the first place. I will also address the side effects of the sample prep procedures.

"Sample Preparation" continued



SPEAKER BIOS & ABSTRACTS

Mary Buckett is currently a Lead Research Specialist in the Corporate Research Analytical Laboratory at 3M Company. She has worked her whole career at 3M in the Transmission Electron Microscopy (TEM) Lab, primarily developing characterization methods for new materials technology, competitive analysis, and protecting intellectual property. Mary has particular expertise in sample preparation methods for all types and combinations of materials. She is the recipient of 13 patents and 21 records of invention at 3M, and has coauthored 78 technical publications. Prior to 3M, Mary was at Argonne National Laboratory (Chicago, IL), and was an NSF Fellow at the National Research Institute for Metals (Tsukuba, Japan). She received a B.S. in materials science and engineering from the University of Wisconsin-Madison, and a Ph.D. in materials science and engineering from Northwestern University (Evanston, IL).



Mary Buckett

From Smart Phones to Silicones – Sample Preparation from an Industrial Lab Perspective

From an industrial analytical lab's point-of-view, there is constant pressure to optimize the quality vs. throughput timing of our work. It's not a stretch to realize that each of these considerations has the potential of derailing an entire effort. So, how can we minimize the risk of this happening? One critical, yet under-appreciated, consideration is the sample preparation step. Get this step right on the first attempt and you have a good chance of producing a good quality result while minimizing the throughput time, and ultimately keeping the analytical costs manageable.

I'd like to share some of the learnings from working 28 years in 3M's Corporate Research Analytical Laboratory. Sample preparation – especially the use of microtomy – from the first handoff to the final form that goes into the analytical instrument (TEM, SEM, AFM, IR, SIMS, etc.) will be covered for investigations of new technology/product developments, competitive teardowns, defect analysis, and quantitative metrology.

"Sample Preparation" continued



SPEAKER BIOS & ABSTRACTS

Shiba Dandpat works as an Application Scientist for LUMICKS. He has a background in RNA biochemistry and single-molecule biophysics. Prior to joining joining LUMICKS, Shiba received his Ph.D. in chemistry from the University of Michigan at Ann Arbor, working on bacterial RNA structures for understanding their role in gene regulation.



Shiba Dandpat

Real-Time Imaging of Dynamic Single-Molecule Interactions: From DNA Binding Proteins to Molecular Motors

Biological processes emerge from mechanisms at the molecular scale. While biophysical techniques (e.g., X-ray crystallography, cryoEM) with bulk biochemical assays (e.g., enzymatic reactions, fluorescent reporters) have helped to better understand emergent structure-function relationships, the complete picture of molecular-scale mechanisms is often missed. In addition, the existing techniques often do not provide tools to observe and manipulate a biological system simultaneously and understand mechanisms from the molecular to the cellular level.

In this talk, I will introduce a single-molecule imaging tool, the C-Trap, which enables users to observe and manipulate biological systems in real-time. This technology combines two Nobel prize-winning ideas (optical tweezers and super-resolution STED microscopy), integrated with easy-to-use microfluidics, that allows users to perform single-molecule studies on a broad range of biological systems. I will present some case studies to highlight the role of C-Trap in looking at DNA/RNA-proteins interactions, proteins/RNA-structure dynamics, nucleic acid/protein condensates, and single-cell studies. The presented case studies and workflows serve as a framework for measuring and visualizing complex and dynamic biological systems, that would be extremely challenging to study using conventional biophysical techniques. Although most of the current applications fall into the biophysics realm, a variety of applications are emerging in material science, nanoscience, and broader biology field every day.



MEMBER DUES

The Society is collecting dues. Please remember to submit your 2022 calendar year membership dues.

Thank you!



PROJECT MICRO

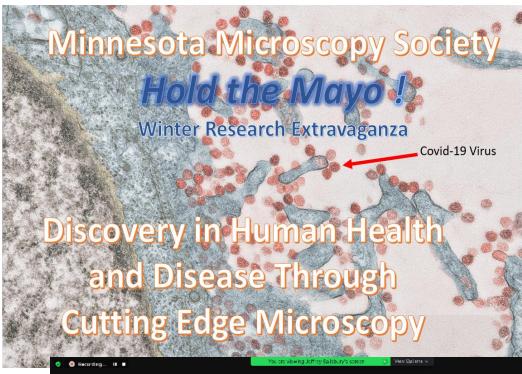
No updates: Project Micro is on hold.

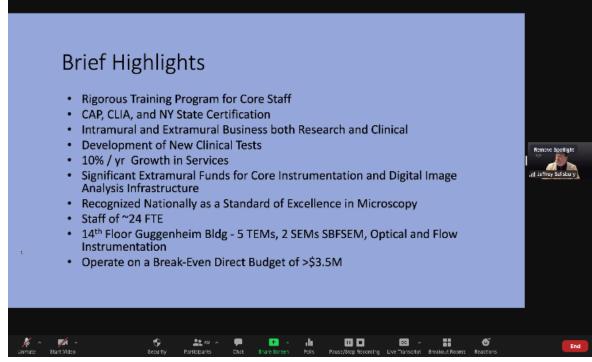
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PAST EVENT HIGHLIGHT: Virtual Mayo Tour & Talks

On February 24, the Minnesota Microscopy Society hosted *Hold the Mayo!* – a Virtual Winter Research Extravaganza. Attendees joined online for a virtual tour of the Mayo Clinic Microscopy Facility and presentations on dissection of brain architecture and chemistry using state-of-the-art microscopy. The event was hosted by our next MMS president, **Jeffrey Salisbury**, **Scientific Director**, **Mayo Microscopy and Cell Analysis Core**, and it also included talks from members of the Mayo research staff.



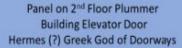




Hold the Mayo! - 2

Microscopy in Mayo's Architecture







Henry Plummer Physician and Architect of the Group Practice



Panel from Foundation House Balfour Hall-Stained Glass showing an Investigator at the RCA 1A TEM



Virtual Tour of Microscopy Facility Jeffrey L. Salisbury, Ph.D.

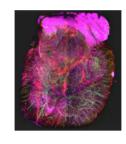
Scientific Director - Mayo Microscopy and Cell Analysis Core



"Chronic Alcohol Exposure Compromises Attention and Avidity for Natural Rewards: A 3D Brain-wide Remodeling of Neural Activation." Lee R. Peyton, Ph.D.

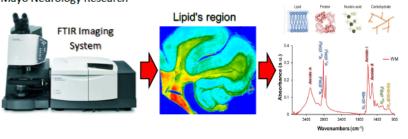
Mayo Molecular Pharmacology and Experimental Therapeutics

(Right) Optical section of a mouse brain stained with FITC-albumin-gelatin (green) and nuclear stain To-Pro 3 (magenta) imaged using a 2.5x objective with a Zeiss Lightsheet 7.





"IR Microspectroscopy as a Tool to Study Biochemical Changes in Neurological Disorders" Alex Gakh, Ph.D. and Yong Guo, Ph.D. Mayo Neurology Research





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Ev Osten

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