

## Biomaterial Surfaces & Interfaces

### Room Naupaka Salon 1-3 - Session BI-MoP

#### Biomaterial Surfaces & Interfaces Poster Session

**BI-MoP-1 Fabrication of Hydrogel-Based Optical Biosensor for Smart Intraocular Lens, Soongeun Kwon, Y. Eom, H. Choi, J. Ahn, S. Park, H. Lim, G. Kim, K. Choi, J. Lee, Korea Institute of Machinery and Materials, Republic of Korea**

Due to the high biocompatibility, facile chemical modification and excellent responsiveness, hydrogel materials have received great deal of attention as wearable or implantable biosensor substrates. To fabricate a hydrogel-based biosensor, a stable bond at the interface of hydrogel and a functional sensing material is essential. In this study, we demonstrated fabrication and application of hydrogel-based optical sensor with a biocompatible micro-grating pattern for implantable medical devices.

To fabricate a functional micro-grating pattern, photolithographic patterning of a photoresist (PR) was performed to define the micro-scale line and spacing pattern. Gold (Au) nanoparticles spin-coated on the PR pattern were patterned by ligand exchange and lift-off process, resulting in an Au micro-grating pattern on a silicon (Si) wafer. The as-fabricated Au micro-grating pattern showed a rabbit ear morphology by controlling the thickness of the PR pattern. Subsequently, molding of a hydrogel precursor into the Au micro-grating pattern on a Si wafer was conducted to transfer the Au micro-grating pattern to the target hydrogel substrate.

The rabbit ear morphology and porous structure of the Au pattern enabled large interfacial contact area between hydrogel precursor and Au nanoparticles, resulting in stable bonding at the interface of Au micro-grating pattern and hydrogel substrate. Due to the biocompatibility of Au and hydrogel, this hydrogel-based biosensor can be used as for implantable medical devices.

As a case study, we demonstrated the application of hydrogel-based optical sensor composed of Au micro-grating pattern for smart intraocular lens (IOL). A pH-responsive hydrogel sensor with Au grating pattern was attached to an IOL to measure the micro-displacement of reactive hydrogel in response to pH changes by optical Moiré pattern detection. With the optical Moiré pattern detection scheme, the proposed hydrogel-based biosensor provides novel implantable optical sensor without external battery, highlighting its potential as a versatile tool for detecting various disease-specific biomarkers.

**BI-MoP-2 Correlative Microscopy Without the Instrument Manufacturer; Using Computer-Readable Fiducial Markers to Navigate Specimens Irrespective of Who Made the Sample Stage, Peter Cumpson, La Trobe University, Australia**

In the diverse field of microscopy, researchers often face challenges in correlating data across different instruments, each with proprietary hardware and software. This work introduces a novel, manufacturer-agnostic solution for correlative microscopy using computer-readable fiducial markers, facilitating seamless navigation and analysis across various microscopy platforms.

Our approach employs laser-etched fiducial markers on sample holders, enabling precise localisation of sample features. This methodology eliminates the need for instrument-specific solutions, significantly enhancing workflow efficiency and accuracy.

We have demonstrated the effectiveness of our system across multiple microscopy techniques, including Scanning Electron Microscopy (SEM), Atomic Force Microscopy (AFM), Energy Dispersive X-ray Spectroscopy (EDX), X-ray Photoelectron Spectroscopy (XPS), and Time-of-Flight Secondary Ion Mass Spectrometry (ToF-SIMS). Our results indicate that this "GPS map for microscopy" not only improves the precision of correlative microscopy but also significantly reduces the time and costs associated with manual sample alignment and calibration.

In collaboration with the National Physical Laboratory (NPL) and University of Durham in the UK we have begun a project to demonstrate and verify the accuracy of this technique. This model considers various scales and imaging modalities, ensuring traceable measurement accuracy and enhancing the reliability of our method.

Our system offers significant advantages for a wide range of applications, from material science and battery research to biomedical and pharmaceutical studies. By enabling precise and consistent navigation across different microscopes, we facilitate interdisciplinary collaboration and accelerate scientific discoveries.

**BI-MoP-3 Supervised MVA and Random Forests for Analysis of GCIB-SIMS Data from Bacteria, John Fletcher, University of Gothenburg, Sweden**

Antibiotic resistance can rapidly spread through bacterial populations via bacterial conjugation. The bacterial membrane has an important role in facilitating conjugation, thus investigating the effects on the bacterial membrane caused by conjugative plasmids, antibiotic resistance, and genes involved in conjugation is of interest. Analysis of bacterial membranes was conducted using gas cluster ion beam-secondary ion mass spectrometry (GCIB-SIMS). The complexity of the data means that data analysis is important for the identification of changes in the membrane composition. Pre-processing of data and several analytical methods for identification of changes in bacterial membranes have been investigated. GCIB-SIMS data from *Escherichia coli* samples were subjected to principal components analysis (PCA), principal components-canonical variate analysis (PC-CVA), and Random Forests (RF) data analysis with the aim of extracting the maximum biological information. The influence of increasing replicate data was assessed, and the effect of diminishing biological variation was studied. Optimized *m/z* region-specific scaling provided improved clustering, with an increase in biologically significant peaks contributing to the loadings. PC-CVA improved clustering, provided clearer loadings, and benefited from larger data sets collected over several months. RF required larger sample numbers and while showing overlap with the PC-CVA, produced additional peaks of interest. The combination of PC-CVA and RF allowed very subtle differences between bacterial strains and growth conditions to be elucidated for the first time. Specifically, comparative analysis of an *E. coli* strain with and without the F-plasmid revealed changes in cyclopropanation of fatty acids, where the addition of the F-plasmid led to a reduction in cyclopropanation.

**BI-MoP-4 Establishing Semi-Oriented Crimped Dual-Sized Fibrous Skeleton for Soft Tissue Engineering Scaffolds, Han Wang, L. Ren, Deakin University, Australia, China; S. Zhao, Deakin University, Australia; H. Yang, Wuhan Textile University, China; L. Kong, Deakin University, Australia**

Soft tissues, such as blood vessels and heart valves, exhibit unique mechanics stemming from their intricate fibrous network architecture. Replicating harmonious structure-function relationships with synthetic analogs remains an unmet challenge. In this work, semi-oriented crimped dual-sized poly(lactic-co-glycolic) acid fibrous membranes with soft-tissue-like mechanical biocompatibility were fabricated by optimizing the organization of the polymer molecular chains and fibers. This is achieved by controlling the entanglement and conformational adjustment of molecular chains. Mechanical properties (modulus:  $39.46 \pm 11.01$  MPa, strength:  $3.19 \pm 0.33$  MPa, toughness:  $0.66 \pm 0.17$  MJ/m<sup>3</sup>) of submicron (145 - 863 nm) and micron (0.66-5.60  $\mu$ m) crimped fiber membranes with an orientation of 38.91-47.38% were not significantly different from those of valve soft tissues. This fibrous skeleton provides inspiration for the development and design of soft tissue scaffolds with superior structure and performance.

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