

N2L Summer School in NCSR Demokritos Athens, June 25-July 6, 2007 "Methods in micro - nano technology and nanobiotechnology"





Organizer: National Center for Scientific Research "Demokritos", in collaboration with the Foundation of Biomedical Research of the Academy of Athens, and Invited experts (lecturers) from other Nano2Life partners.

Information: www.imel.demokritos.gr



Target

- · Modern Research takes advantage of Micro and Nanotechnology developments
- · Merging areas of research (Nanobiotechnology) demand interdisciplinary skills
- · Necessary for researchers from Life Sciences, Chemistry, and Engineering to acquire skills in Micro and Nanotechnologies, nanomedicine

Establish common language between the various disciplines-promote interdisciplinary research

Content: 2-week intensive summer school

Offers: classroom and laboratory experience on: micro and nano-technology processes / materials / applications Targeted in: Nanobiotechnology, nanomedicine

Section 1: Principles of biochemistry, cell biology, physics and microelectronics. **<u>1.1:</u>** Cell biology principles

1.2: Structure of biological macromolecules

1.3: Microelectronic Materials and Device Technology

1.4: Introduction to nanobiotechnology

Unit 2.1: Micro and Nano-fabrication science and technology

2.1.1 and 2.1.2: Patterning technologies

2.1.3: Patterning of biomolecules and other biological substances

2.1.4: Molecular bioelectronics

Laboratory 2.1.1: Fabrication of microfluidic devices on plastic substrates by lithographic techniques.

Laboratory 2.1.2: Fabrication of microfluidic Devices on Plastic substrates by Lithography and plasma etching techniques





ratory 2.1.3: Electrical characterization of tunnelling devices based on organic molecules or biomolecules

Unit 2.2: Nanomaterials for bio-applications, Characterization, Imaging 2.2.1: Drug discovery and development

2.2.2 and 2.2.3: Drug Release and Delivery Systems - Methods

2.2.4: Bioengineered nanomaterials

2.2.5: Magnetic Nanoparticles for Bioapplications

2.2.6: Imaging with Scanning Probes (AFM, STM, SNOM).

2.2.7: Experimental techniques for magnetic characterization of ferrofluids and Applications of ferrofluids in medicine

2.2.8: Fluoresence and 3D imaging visualization using confocal microscope Laboratory 2.2.1: Drug inclusion in cyclodextrins: monitoring in situ by NMR spectroscopy, X-ray diffraction characterisation of drug inclusion and 3-D visualisation



Laboratory 2.2.2: Liposomes: preparation and characterisation by dynamic light scattering and ζ-potential

Laboratory 2.2.3: Video enhanced optical microscopy and Atomic Force



Liposome-liposome interactions: Correlation of Optical Microscopy and Dynamic Light Scattering results

Liposomes PEG 0% PEG 5% PEG 15%

Who should attend:

Group leaders involved in molecular biology or biotechnology Post Doctoral Fellows, Graduate students with Life Science, Science or Engineering background, medical doctors All those who wish to apply micro-technology in their research

Maximum number of registrants 30 persons

Fees: N2L members:1000 Euro Others: 1400 Euro (includes handouts, coffee-breaks, lunches, school dinner, two excursions, NO accommodation) To encourage Greek participation Demokritos will grant partial scholarships to selected Greek participants oon request on the application form Deadlines: N2L May 7th - Others May 14th

Syllabus

Laboratory 2.2.4:	State of the art	confocal	microscopy	of biological samples	

Laboratory 2.2.5: Magnetic nanomaterials for bio applications

Laboratory 2.2.6: Magnetic Hyperthermia experiment using magnetic nanoparticles

- Unit 2.3: Molecular and Cellular biology and Applications
- 2.3.1: Introduction to proteomics
- 2.3.2: Analysis of biomolecules by mass spectrometry
- 2.3.3: Binding Assays and Immunosensors
- 2.3.4: Protein and DNA arrays
- 2.3.5: Metabolomics
- 2.3.6: Bioinformatics topics with emphasis on software for proteomics
- 2.3.7: Applied Bioinformatics in BioNanoTechnology

Laboratory 2.3.1: Protein Separation by two-dimensional electrophoresis

Laboratory 2.3.2: Protein identification by MALDI-TOF MS, LC-ESI-MS and LC-MALDI-MS

Laboratory 2.3.3: Fabrication of protein microarrays using nanoplotter

Laboratory 2.3.4: Fabrication of protein microarrays using lithography

Laboratory 2.3.5: Fluorescence detection of protein arrays





Twelve rows of different protein spots fabricated in 12 succesive lithographic steps

Fluorescence picture of the rabbit yglobulins and biotinylated-BSA spot arrays after a 2 h immunoreaction with a mixture of AF 546 labeled streptavidin (red spots) and AF 488 labeled anti-rabbit IgG antibody (green spots). The spot size is approximately 4 µm.

Laboratory 2.3.6: Bioinformatics laboratory

- Unit 3.1: Microfluidic and Lab on chip devices
- 3.1: Principles of Integrated Biosensing Devices
- 3.2: Acoustic wave sensors: from device fabrication to biological applications
- 3.3: Lab on chip devices: Principles, applications, opportunities

Laboratory 3.1: Operation of a lab-on-a-chip optical device using model assays and real time measurements

Monolithic silicon optocouplers



Laboratory 3.2: Demonstration of a capillary fluoroimmunosensor

