

Summer School in NCSR "Demokritos" 13 – 17 September 2010, Athens, Greece "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 other invited experts
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**

The summer school offers: classroom and laboratory experience on:
micro and nano-technology processes / materials / applications
Targeted in: Nanobiotechnology, Nanomedicine

Who should attend

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

Maximum number of attendants: 20

Fees: 1000 Euro

(includes handouts, coffee-breaks, lunches, school dinner,
excursion, NO accommodation)

Deadline: July 30th 2010

Syllabus

Section 1: Principles of biochemistry, cell biology, physics and microelectronics.

- 1.1:** Introduction to nanotechnology and nanobiotechnology
- 1.2:** Cell biology principles
- 1.3:** Structure of biological macromolecules
- 1.4:** Microelectronic Materials and Device Technology
- 1.5:** Introduction to Biosensors

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

Laboratory 2.1.1: Fabrication of microfluidic devices on plastic substrates by
+2.1.2 soft lithography and deep polymer plasma etching



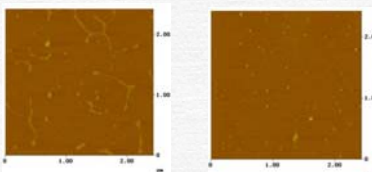
PMMA Capillaries

Laboratory 2.1.3: SPM Techniques for molecular devices

Unit 2.2: Nanomaterials for bio-applications, Characterization, Imaging

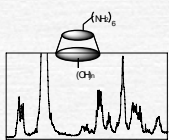
- 2.2.1:** Drug Delivery and Targeting Systems – Focus on Liposomes
- 2.2.2:** Drug Delivery and Targeting Systems – Focus on cyclodextrin delivery, studied by NMR and XRD
- 2.2.3:** Magnetic nanoparticles for bioapplications
- 2.2.4:** Scanning Probe Microscopy in Nanobiotechnology

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



Atomic Force Microscopy
Formation of DNA nanoparticles
of ~40 nm diameter

Laboratory 2.2.2: Intracellular visualisation of Porphyrin-Cyclodextrin conjugates as PDT agents/chemotherapeutic drug carriers by confocal microscopy



Unit 2.3: Molecular and Cellular biology and Applications

- 2.3.1:** Gel-based protein analysis methods
- 2.3.2:** Non-gel based protein analysis methods
- 2.3.3:** Binding Assays and Immunosensors
- 2.3.4:** DNA and Protein arrays: fabrication, detection and applications

Laboratory 2.3.1: Protein separation by two-dimensional electrophoresis

Laboratory 2.3.2: Mass spectrometry

Laboratory 2.3.3: Fabrication of protein microarrays using nanoplatter

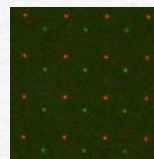
Laboratory 2.3.4: Fabrication of protein microarrays using lithography

Laboratory 2.3.5: Fluorescence detection of protein arrays

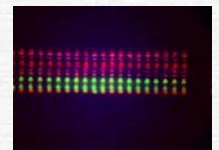
Laboratory 2.3.6: Bioinformatics basic theory & laboratory

Laboratory 2.3.7: Structural Bioinformatics: Molecular Simulations and Visualization

Laboratory 2.3.8: State of the art fluorescence imaging & confocal microscopy of biological samples



Fluorescence picture of the rabbit γ -globulins 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 μ m.

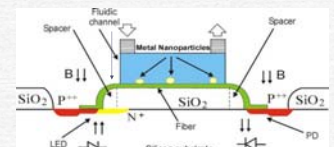
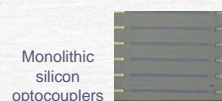


Twelve rows of different protein spots fabricated in 12 successive lithographic steps

Section 3: Towards Integrated Nanobiotechnology systems

3.1: Principles of Integrated Biosensing Devices

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



Laboratory 3.2: Demonstration of a capillary fluoroimmunosensor

