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## Nano eNabler™

Desktop Molecular Printing System



***“The possible applications are limited only by one’s imagination”***

Jan Hoh Ph.D.  
Johns Hopkins University School of Medicine

# BioForce Nano eNabler™

The Nano eNabler™ is a new and enabling technology for ultramicro- and nanoscale fluid delivery. It is a multifaceted platform capable of delivering attoliter to femtoliter ( $10^{-15}$  to  $10^{-18}$  liter) volumes of solutions containing biomolecules and other materials to defined locations on surfaces with ultramicro (1-20  $\mu\text{m}$ ) feature sizes and nanometer spatial resolution. The ultraminiaturized nature of this technology reduces sample requirements to a bare minimum. For example, the NanoArrayer can create a diagnostic biochip that uses just a few cells, or less than one drop of blood, for critical biomedical analysis. The broad applicability and wide range of compatible materials create many new and exciting opportunities. A few representative examples of printable materials and their applications may be found in the table below.



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## Material Deposited

Antibodies and other proteins	Biosensors, biomedical devices, molecular screening, cell biology, nanobiology
Nucleic acids	Gene chips, genomics R&D, biosensors
Viruses	Biosensors, diagnostics, nanodevices
Adhesives	MEMS, nanodevices
Colloidal particles	Electronics, nanodevices, materials R&D
Quantum dots	Optical devices, diagnostics, materials R&D
Etchants, solvents	MEMS, electronics, microfabrication

## Representative Applications

## Specifications:

- Power Requirements:
  - AC 120V/60Hz 4.0A
  - AC 240V/50Hz 2.0A
- Dimensions: 51 cm x 37 cm x 33 cm
- Controller Dimensions: 54 cm x 54 cm x 64 cm
- Instrument Weight: 18.14 kg (40 lb)
- Controller Weight: 38.6 kg (85 lb)
- X,Y Stage Travel Range: 50 mm x 50 mm
- X,Y Stage Resolution: 20 nm
- Z Stage Range: 45 mm
- Z Stage Fine Resolution: 100 nm
- Laser-Based Force Feedback
- Controllable Humidity Range: 25 - 80% RH
- Motorized Optical Microscope (150X to 1000X) w/ Video Capture
- Integrated Pentium 4 3.0GHz, 512MB RAM
- Windows XP Professional
- Flat Screen LCD Monitor
- Partial List of Compatible Surfaces:
  - ★ Silicon, Glass
  - ★ Gold, Other Metals
  - ★ Alkanethiol Monolayers
  - ★ Silanes
  - ★ PDMS
  - ★ Hydrogels
  - ★ Nitrocellulose
  - ★ Plastics and Other Polymers

"The versatility of this method makes it a powerful tool for nearly any lab"

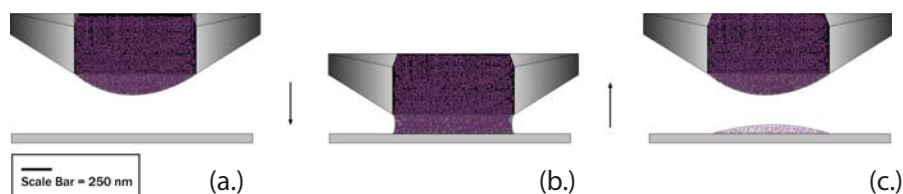
## The FEMTO Process

### Fluidics Enhanced Molecular Transfer Operation

The FEMTO process enables the direct deposition of virtually any molecule onto virtually any surface. Small molecules, biomolecules, reactive solutions, and even nanoparticles can be printed onto a surface. In fact, because this is a fluid transfer, the size or molecular weight of the material has no effect on the process. Rapidly print thousands of spots with attoliter to femtoliter volumes and diameters from 1-20 microns. Need to draw lines instead? FEMTO's continuous fluid flow can handle that too.

The key to the FEMTO process is the Surface Patterning Tool (SPT), with its microfluidic channels constantly delivering a fresh supply of liquid to be transferred onto the surface. Multi-channel SPTs can be easily loaded to allow parallel printing of a single compound, or multiplexed printing of several different molecules.

As depicted in the illustration below, fluid flows from the reservoir, down the channel, to the end of the cantilever where the channel narrows to a tiny gap (a). Upon contact with the surface, a small volume of liquid held in the gap by surface tension is directly transferred to the surface in an event typically requiring less than 100 msec (b). Capillary fluid flow down the channel instantly replenishes the volume at the gap, and the SPT is ready to print the next feature (c).



## The FEMTO Process

### How is FEMTO Different from Other Methods?

Each method of surface modification has its own set of advantages and disadvantages. PDMS microcontact printing ( $\mu$ CP) may offer the benefit of parallel printing but at the cost of other very useful features such as multiplexing and the flexibility of on-the-fly revisions. A scanning probe method such as Dip Pen Nanolithography (DPN) is a good choice for creating very small features of a low molecular weight compound, however this technique has drawbacks as well. DPN is a diffusion-limited process, which means the printing speed for larger spots and higher M.W. molecules such as proteins can be very slow. Other common complaints from DPN users include a limited selection of suitable surfaces and a lack of reproducibility. Techniques employing glass nanopipettes offer the speed of a direct fluid flow process such as FEMTO, but the nanopipettes can be difficult to load and their closed design makes them prone to blockage. FEMTO's open channels ensure that your liquid will always have a path to the surface for maximum reliability.

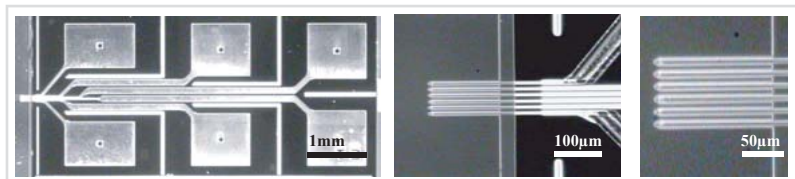
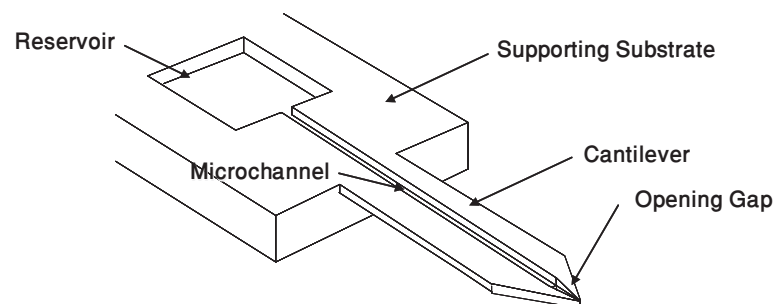
	FEMTO	DPN	Nanopipettes	$\mu$ CP
Parallel Printing	★	●	●	●
Multiplexing	★	●		
Rapid Revisions	★	●	●	
Versatility	★		●	●
Speed	★		●	●
Small Molecules	★	●	●	●
Large Molecules	★		●	●
Reliability	★			●

*"It is the world's smallest pipetter"*

## The Surface Patterning Tool (SPT™)

### Disposable Print Cartridges

The SPT™ (Surface Patterning Tool) is the "ink cartridge" for the Nano eNabler™. Each SPT is a microcantilever-based microfluidic sample handling and delivery device. SPTs contain either a single microcantilever print head or multiple microcantilevers for simultaneous printing of multiple molecular species or materials. The integrated microfluidic network transports fluid samples from reservoirs located on the SPT through microchannels to the distal end of the cantilever. Thousands of spots can be printed with one load. SPTs can be used to print materials that include biological samples such as proteins, DNA, RNA, and whole viruses, as well as non-biological samples such as chemical solutions, colloids and particle suspensions. SPTs are disposable, eliminating any requirement for labor intensive cleaning and risk of cross contamination inherent in re-use. For some experiments, SPTs may be cleaned in the BioForce UV TipCleaner for re-use. BioForce Nanosciences Inc. supplies a variety of different SPTs to meet a broad range of customer needs and can also provide custom SPT design and fabrication.



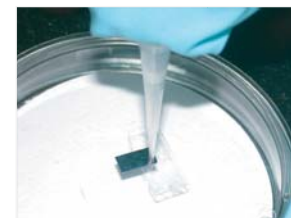
Optical images of 6 cantilever SPT

## The Surface Patterning Tool (SPT™)

For creating arrays of biomolecules, the preferred method of loading the deposition tool will depend upon the ultimate density of the nanoarray. Arrays of a single compound can be created using either front-loading or back-loading of the deposition tool.

### Back Loading

Back-loading involves pipetting a small volume (generally less than 0.1 µl) of sample into one of the wells etched into the top surface of the silicon deposition tool substrate. The sample will fill the well and flood the channel that runs down the length of the cantilever. Several thousand features may be printed with a back-filled SPT.



### Front Loading

Front loading of SPTs is often convenient for arraying compounds in a relatively small number of domains (100s). The solution of interest is placed on a coverslip or similar surface (far less than 1 µl is required). The user then mounts the loading slide or coverslip onto the Nano eNabler™



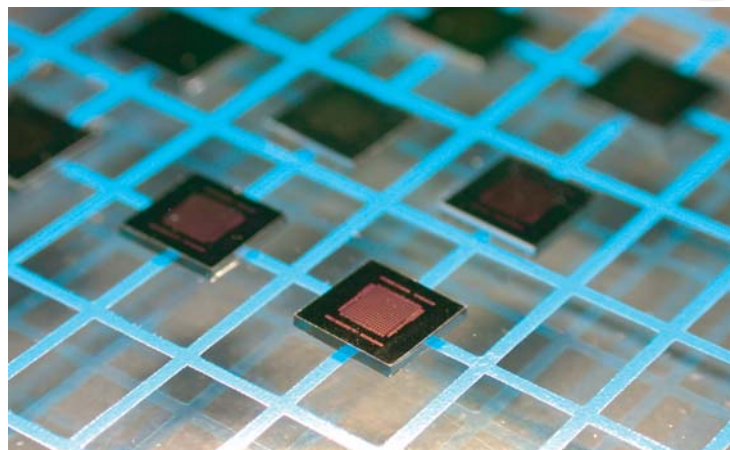
sample platform. The SPT is positioned above the loading spot. The Nano eNabler's "Find Surface" command is then used to bring the deposition tool and loading surface into contact. Finally, the precision X,Y control of the NanoArrayer is used to touch the SPT to the solution of interest, and spontaneously fill the SPT micro-channel.

“...it will completely change the way you think about surface patterning”

## Sindex™ Chips - The Printing Surface

### Ideal Patterning Substrates

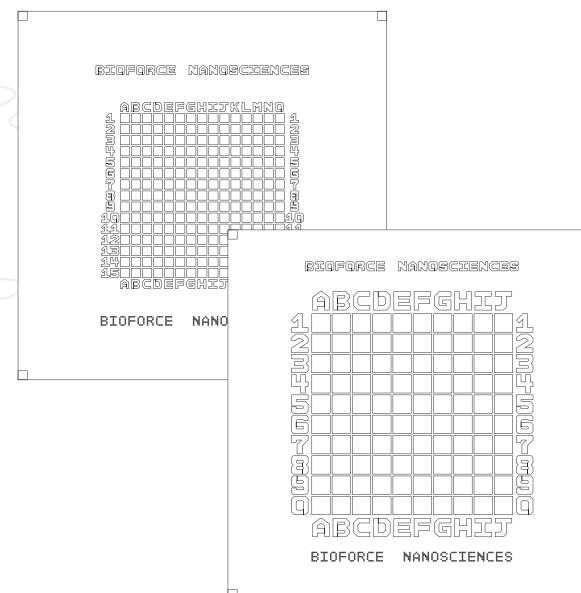
The Sindex™ Chips provide ideal printing surfaces for the NanoArayer™. These 4x4 mm silicon substrates contain topographically defined pads that are arrayed within an alphanumeric indexing system. The pads are flat and smooth, making them compatible with conventional (e.g., fluorescence) and more exploratory (e.g., atomic force microscopy) readout mechanisms. The indexing system allows precise location and relocation of specific positions on the chip. The surface can be coated with different metals and treated by a variety of approaches to render it chemically reactive.



## Sindex™ Chip Options

BioForce Sindex™ Chips come with a variety of surface chemistry options, including a self-assembling monolayer (SAM) of alkanethiolate molecules; CH<sub>3</sub>, COOH, OH, NHS and more.

There are two pattern options. The Sindex 15-100 offers a 15 x 15 pad array with 100 µm square pads. The Sindex 10-200 offers a 10 x 10 pad array with 200 µm square pads.



Chip Type	Array No.	Pad size µm	Spacing µm	Etch depth nm	Chip size, mm	Chip thickness, µm	Material
Sindex-15-100	15	100	20	200	4	480	Si
Sindex-10-200	10	200	20	200	4	480	Si

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*"it is an incredibly exciting platform for patterning molecules"*

## Software Interface

### Flexible Software Interface

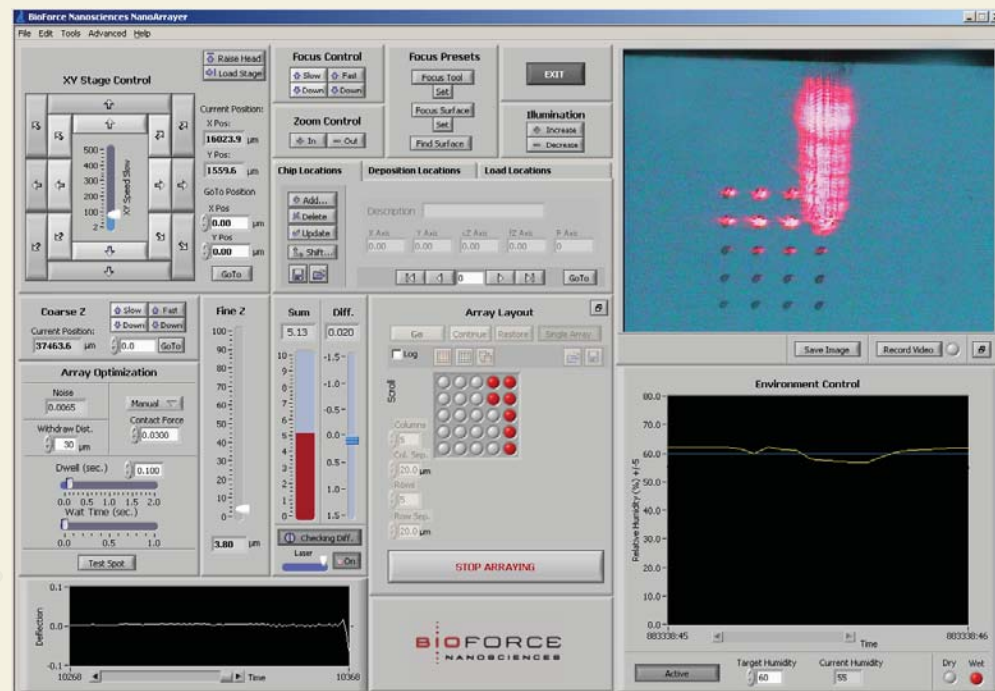
The Nano eNabler™ System utilizes Windows™ based BioForce authored software that allows the user to control nearly every aspect of the instrument. The modular nature of the software creates opportunities for rapid creation of user-specific functions.

### Software Controlled:

- X,Y Movement
- Z Movement
- Optical Focus and Zoom
- Optical Illumination
- Laser Intensity
- Environmental Control
- Integrated Video

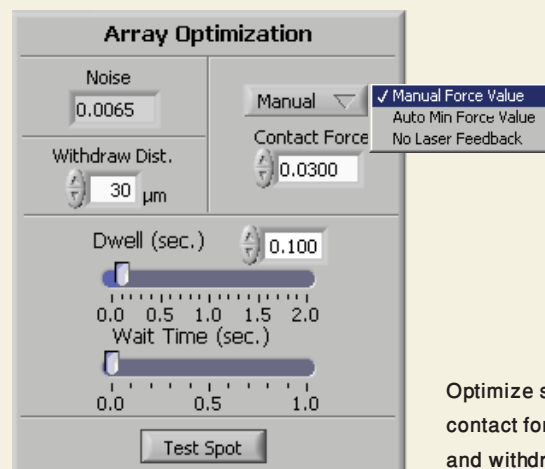
### Features:

- Design & Spot Complex Pattern Layouts on the Fly
- Save Load & Deposition Locations, Pattern Layouts
- Definable Dwell & Wait Time for Spot Size Optimization
- Multi-Chip Automated Arraying
- Automatic Find Surface
- Contact Force Modes:
  - Automatic Contact Force
  - Manual Contact Force Value
  - No Laser Feedback
- Save Video/Photos from Integrated Video
- On Video X,Y,Z Navigation

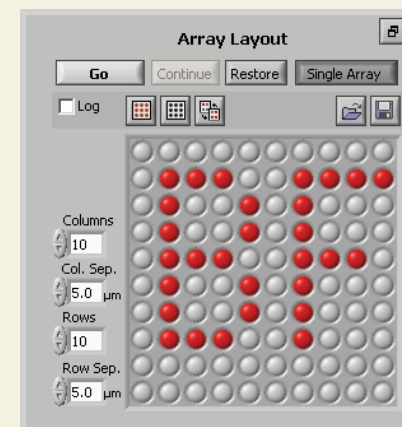


Screenshot of Nano eNabler Software Interface

Easily design and save complex pattern layouts, adjust column and row separation



Optimize spot size by adjusting contact force, dwell time, wait time, and withdraw distance



*"The Nano eNabler is one of those new tools that come along infrequently..."*

## Environmental Control

### Precise Control for Patterning Optimization

Humidity is one of the most important variables to control when attempting to print micron or sub-micron scale features. Insufficient humidity can negatively impact material transfer, and excessive humidity can result in undesirable heterogeneity of feature size. The Nano eNabler™ is equipped with an outer enclosure that provides a barrier between the Nano eNabler™ instrumentation and the ambient room conditions.

## Environmental Control

### Achieving and Maintaining

The Nano eNabler™ environmental control chamber allows complete control of the humidity surrounding the instrument. The front door accommodates full access for initial setup or periodic adjustments. Desired humidity is set through the Nano eNabler™ software interface and achieved using a computer-controlled system of dry gas and humid air flow. Dry conditions are attained by filling the chamber with a user-supplied inert dry gas such as nitrogen. Humid environmental conditions are realized by the novel humidification device attached to the environmental chamber that rapidly creates a saturated  $\text{ddiH}_2\text{O}$  atmosphere (shown below).



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*"It is an innovative instrument for various surface science technologies"*

## Applications

### Application Notes

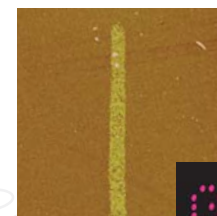
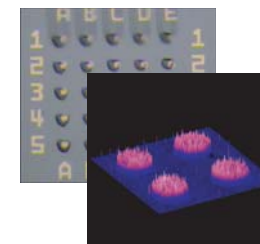
The ability of the Nano eNabler™ to deliver attoliter volumes of materials to defined locations with nanoscale accuracy creates an unprecedented opportunity for discovery science. The open-ended possibilities enabled by the Nano eNabler are rapidly being documented in a series of application notes (available at [www.bioforcenano.com/resources](http://www.bioforcenano.com/resources)). Each Application Note describes a use of the Nano eNabler developed either in-house or by our growing user base. Some examples appear at the right.

### Additional Applications:

- Molecular Detection
  - Chemical and Biosensor Functionalization
    - Cantilever Sensors
    - Nanotube Sensors
    - Optical Waveguide Sensors
  - Printing Inside Microfluidic Channels
- Diagnostics and Pharmaceutical Discovery
  - Small Volume Biomarker Analysis
    - Laser Capture Microdissection Samples
    - Single Cell Expression Profiling
    - Small Animal/Organism Model Systems
    - Pre-Natal Biomarker Screening
  - Patterning Surfaces for Cellular Studies
    - Stem Cell Differentiation/Proliferation
    - Cell Adhesion/Motility/Chemotaxis
    - Tissue Engineering
    - Peptide/MHC Complexes
- Engineering Surface Architectures
  - Direct Patterning of Nanomaterials
    - Quantum Dots, Colloids, Magnetic Nanoparticles
  - Patterning Reactive Solutions
    - Etchants, Resists, Adhesives

### ViriChip - Virus Detection by AFM

A novel biosensor platform enabled by the Nano eNabler™



### Patterning Nanomaterials:

Drawing lines and printing patterns with quantum dots

### Patterning for Protein-Cell Studies

Spots or lines of extracellular matrix proteins or cell signalling molecules can be patterned to enable the study of single cells. Complex patterns of multiple proteins may be constructed such that a single cell interacts with many spots.

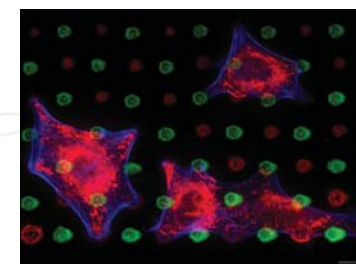


Image courtesy of J. Hoh, Johns Hopkins University

### Protein-Protein Interaction Screening by AFM

The ultraminiaturized size of patterns made by the NanoArrayer enable the use of novel approaches to identify molecular interactions. Here the AFM is used to evaluate a 4 x 4 array of antibody-antigen interaction domains on a biochip surface.

