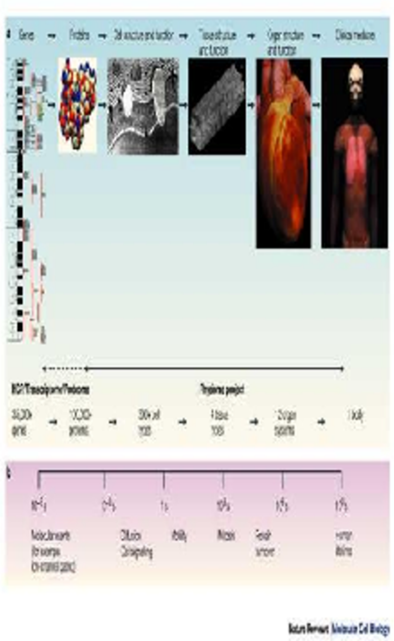
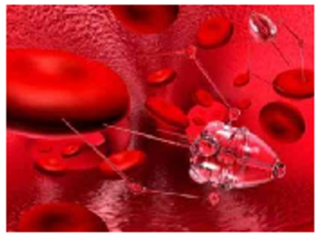
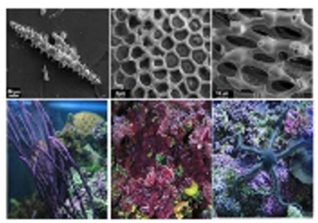
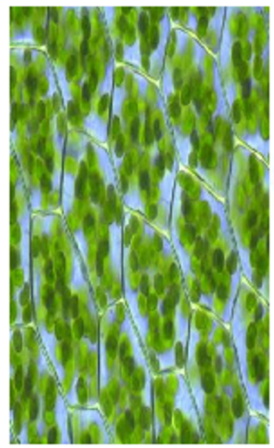
Nano-enabled Biological Tissues



COURTESY: Nature Reviews Molecular Cell Biology, 4, 237-243 (2003).

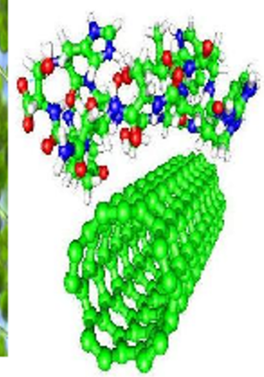






http://laegroup.ccmr.cornell.edu/

http://www.afs.enea.it/project/cmast/group3.php



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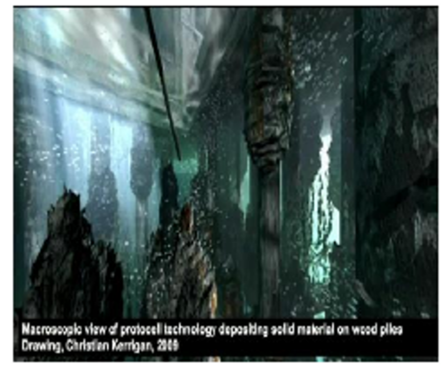
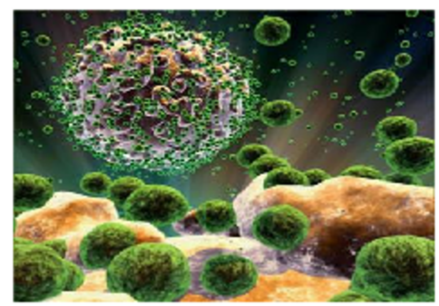




COURTESY: http://library.thinkquest.org/05aug/00736/nanomedicine.htm

By Bradly Alicea

http://www.msu.edu/~aliceabr/

Presented to Nanotechnology and Nanosystems group, Michigan State University. October, 2010.

Abstract

Tissue Engineering

Nano

BioMEMS

Biomechanics

This talk is an attempt to define a new field called "Nano-enabled Biological Tissues". As such, this talk serves as a review of both the theoretical underpinnings and relevant recent results.

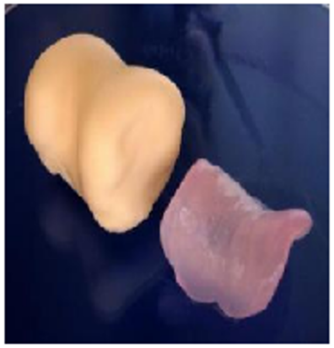
The presentation is divided into several parts:

* in the first section (slides 3-4), the concept of nano-enabled tissues are introduced as a complex system that can be engineered at multiple scales.

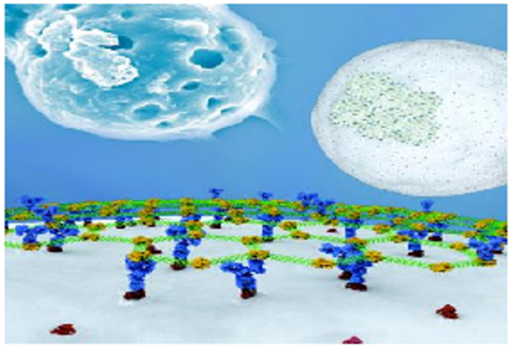
* the second section (slides 5-12) contains three essential ingredients to achieve the technological vision. Current examples of each ingredient are introduced separately.

* in the third section (slides 13-17), additional essential ingredients are considered. This includes strategies for system construction (top-down vs. bottom-up), and additional tools for functionality such as computational intelligence.

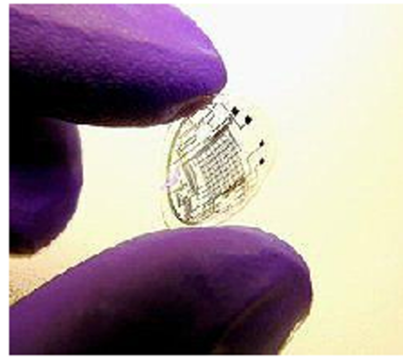
Nanoscale Technology Enables Complexity at Larger Scales.......



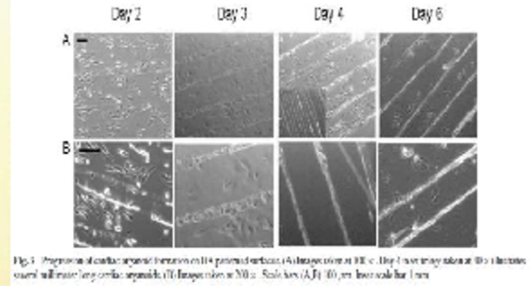
Self-assembled cartilage



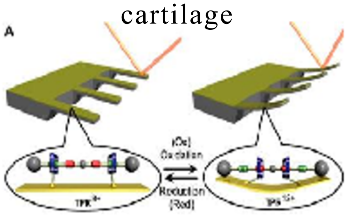
Nano-scale biofunctional surfaces (cell membrane) http://www.nanowerk.com/spotlight/spotid=12717.php



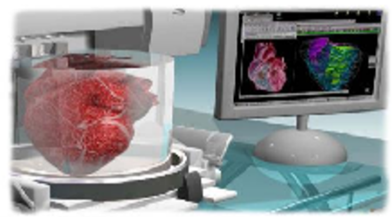
Flexible electronics embedded in contact lens



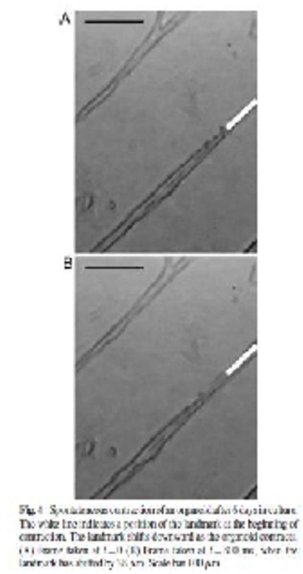
Formation (above) and function (below) of contractile organoids. Biomedical Microdevices, 9, 149 157 (2007).



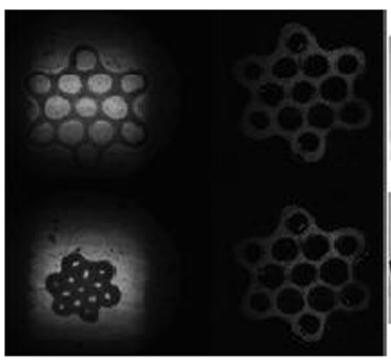
DNA/protein sensor, example of BioNEMS device (left).

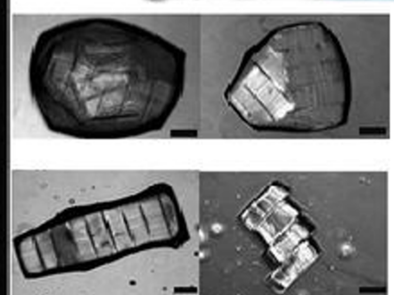


"Bioprinting" to construct a heart (left).

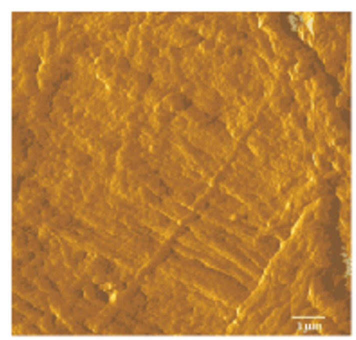


Cells cultured in matrigel clusters



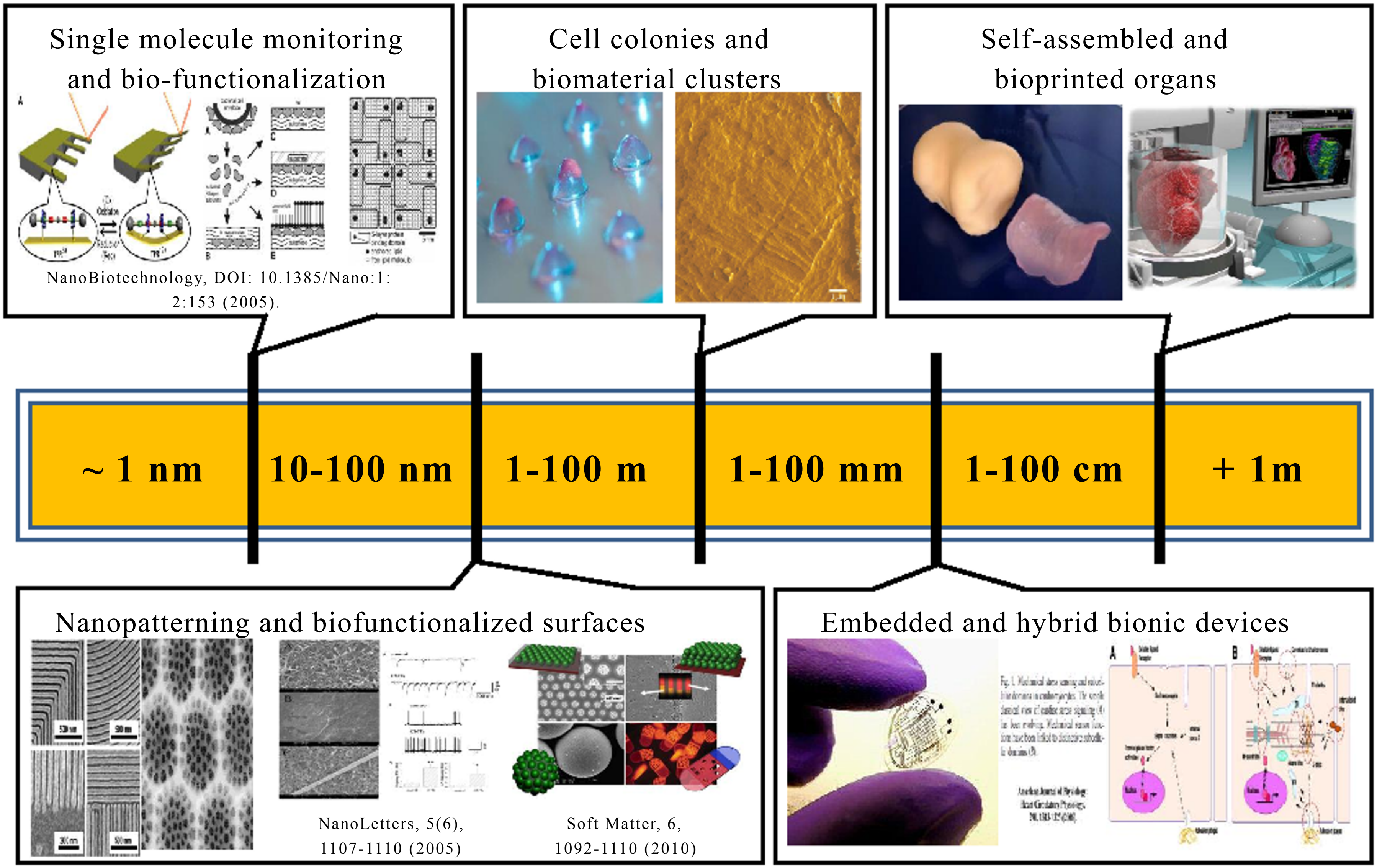


Guided cell aggregation. COURTESY: "Modular tissue engineering: engineering biological tissues from the bottom up". Soft Matter, 5, 1312 (2009).



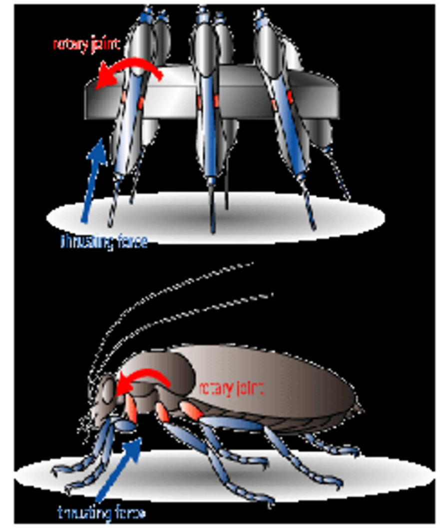
Self-organized collagen fibrils

Role of Scale (Size AND Organization)



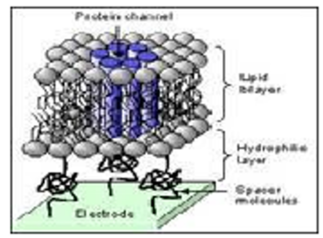
Ingredient I, Biomimetics/ Biocompatibility

Biomimetics: engineering design that mimics natural systems.



Nature has evolved things better than humans can design them.

* can use biological materials (silks) or structures (synapses).



Biocompatibility: materials that do not interfere with biological function.

* compliant materials used to replace skin, connective tissues.

* non-toxic polymers used to prevent inflammatory response in implants.

Polylactic Acid Coating

Cyclomarin Source

Hydroxyapatite (Collagen)

Parylene (Smart Skin)

Artificial Skin, Two Approaches

Approximating cellular function:

"Tissue-Engineered Skin Containing Mesenchymal Stem Cells Improves Burn Wounds". Artificial Organs, 2008.

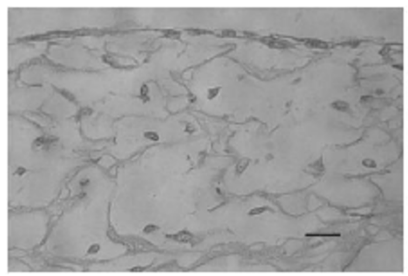


FIG. 2. Hematoxylin–eosin-stained histological section of mesenchymal stem cells (MSCs) grown on collagen-GAG scaffolds. Wavelike collagen bundles and of randomly scattered MSCs can be observed. Scale bars = 100 μm.

Stem cells better than synthetic polymers (latter does not allow for vascularization).

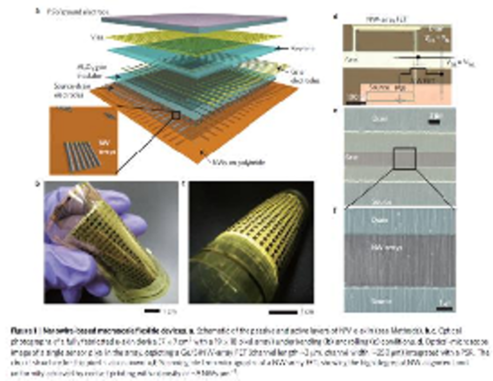
* stem cells need cues to differentiate.

* ECM matrix, "niche" important.

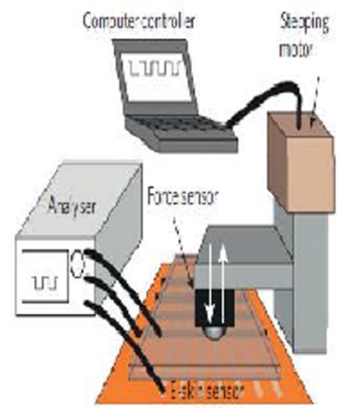
* biomechanical structure hard to approximate.

Approximating electrophysiology:

"Nanowire active-matrix circuitry for low-voltage macroscale artificial skin". Nature Materials, 2010.



a

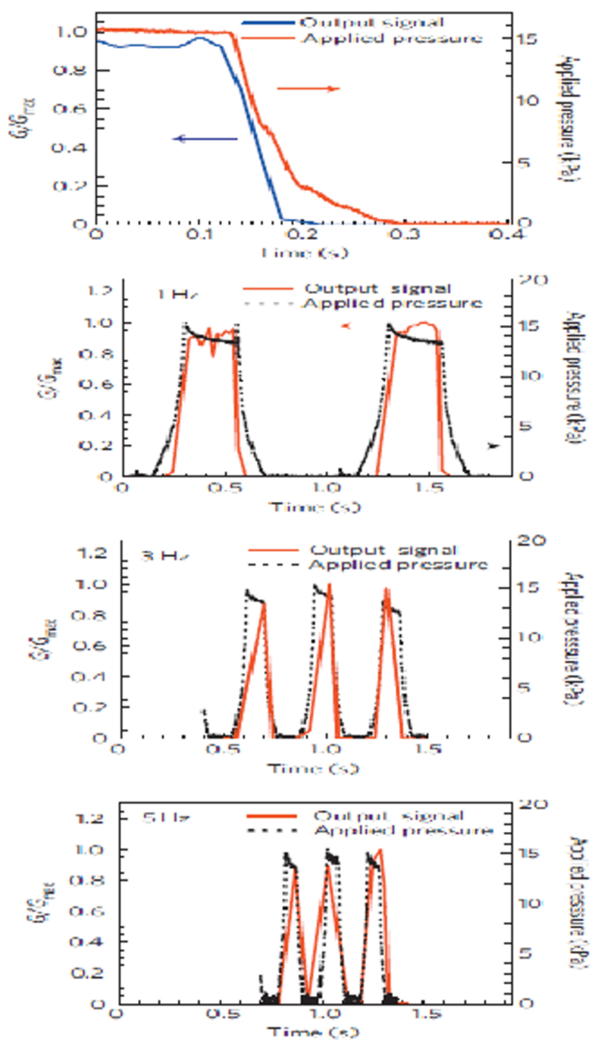


Skin has important biomechanical, sensory functions (pain, touch, etc).

* approximated using electronics (nanoscale sensors embedded in a complex geometry).

* applied force, should generate electrophysiological-like signal.

Artificial Skin Response Characteristics



Results for stimulation of electronic skin:

Output signal from electronic skin, representation is close to pressure stimulus.

* only produces one class of sensory information (pressure, mechanical).

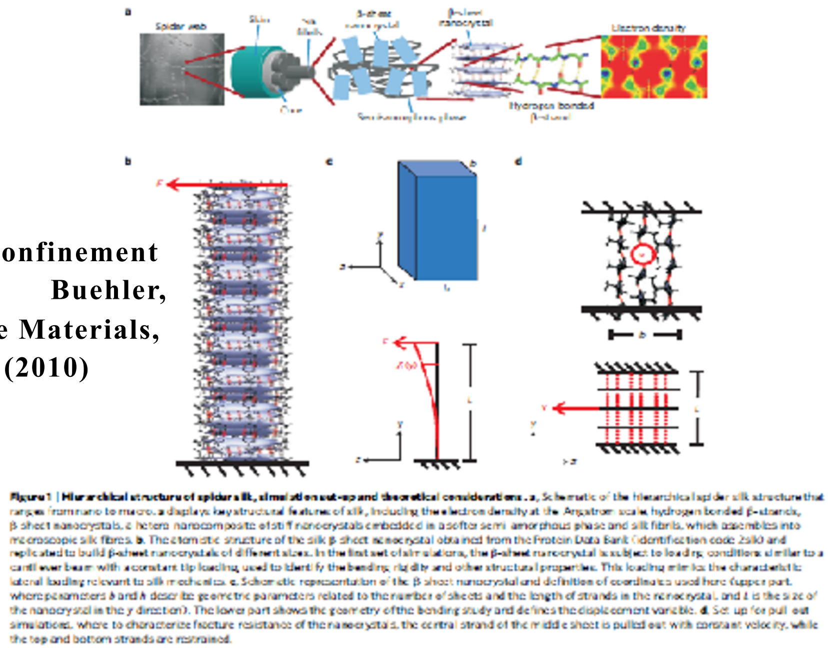
Q: does artificial skin replicate neural coding?

* patterned responses over time (rate-coding) may be possible.

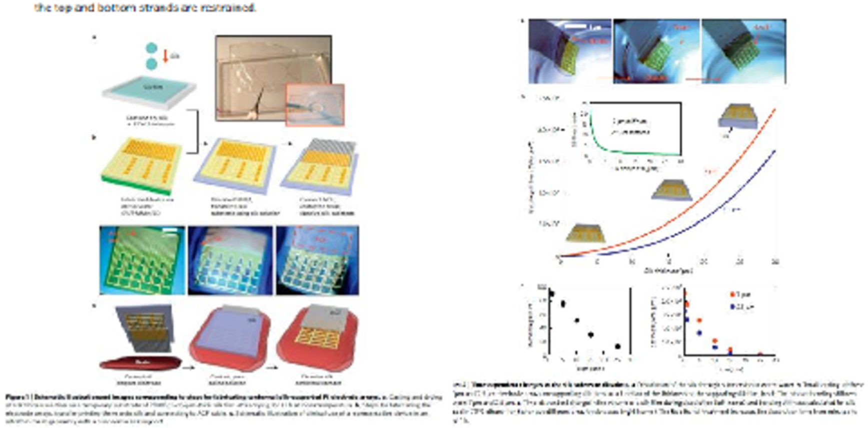
* need local spatial information (specific to an area a few sensors wide).

* need for intelligent systems control theory at micro-, nano-scale.

Silk as Substrate, Two Approaches



Nanoconfinement M. Buehler, Nature Materials, 9, 359 (2010)



Bio-integrated Electronics. J. Rogers, Nature Materials, 9, 511 (2010)

Nanoconfinement (Buehler group, MIT):
* confine material to a layer ~ 1nm thick (e.g. silk, water).

* confinement can change material, electromechanical properties.

Bio-integrated electronics (Rogers group, UIUC):
Silk used as durable, biocompatible *substrate for implants, decays in vivo:*
* spider web ~ steel (Young's modulus).

* in neural implants, bare Si on tissue causes inflammation, tissue damage, electrical interference.

* a silk outer layer can act as an insulator (electrical and biological).

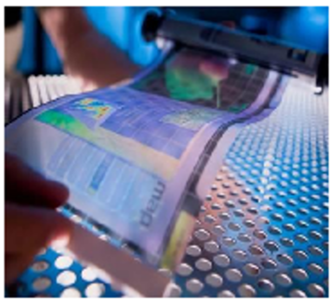
Ingredient II, Flexible Electronics

Q: how do we incorporate the need for compliance in a device that requires electrical functionality?

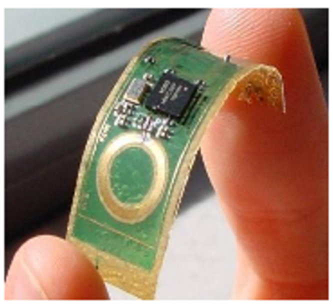
* tissues need to bend, absorb externally-applied loads, conform to complex geometries, dissipate energy.

A: Flexible electronics (flexible polymer as a substrate).

Flexible e-reader



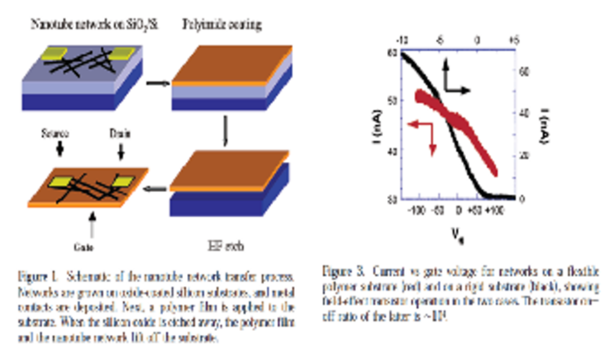
Flexible circuit board



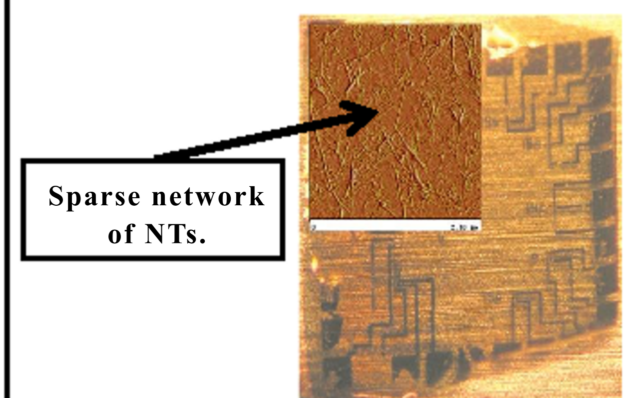
Nano version (Nano Letters, 3(10), 1353-1355 - 2003):

* transistors fabricated from sparse networks of nanotubes, randomly oriented.

* transfer from Si substrate to flexible polymeric substrate.



Nano Letters, 3(10), 1353-1355 (2003)



E-skin for Applications

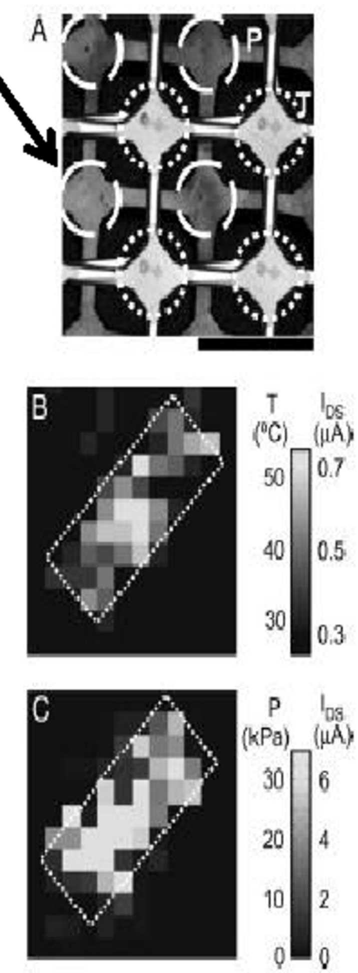
Organic field effect transistors (OFETs):
* use polymers with semiconducting properties.

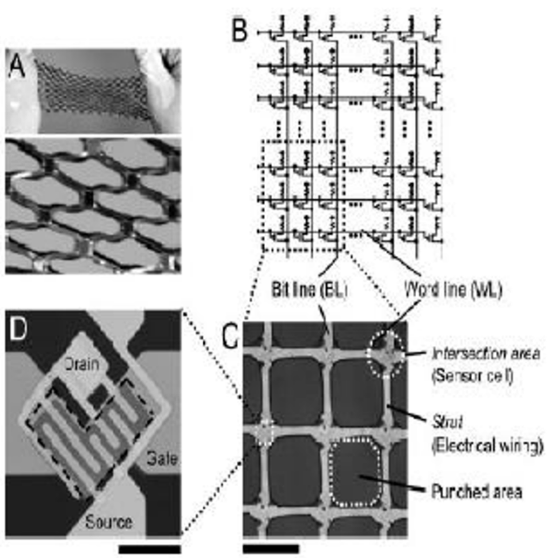
Thin-film Transistors (TFTs):
* semiconducting, dielectric layers and contacts on non-Si substrate (e.g. LCD technology).

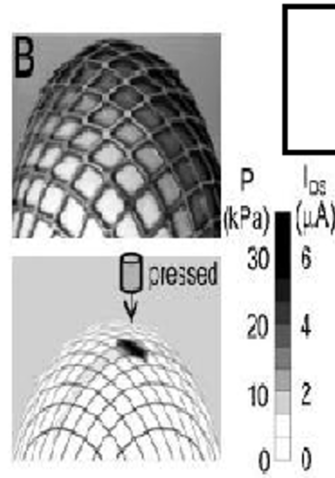
* in flexible electronics, substrate is a compliant material (skeleton for electronic array).

Embedded array of pressure and thermal sensors

PNAS, 102(35), 12321 12325 (2005).







PNAS, 102(35), 12321 12325 (2005).

Conformal network of pressure sensors

Create a bendable array of pressure, thermal sensors.

Integrate them into a single device (B, C on right).

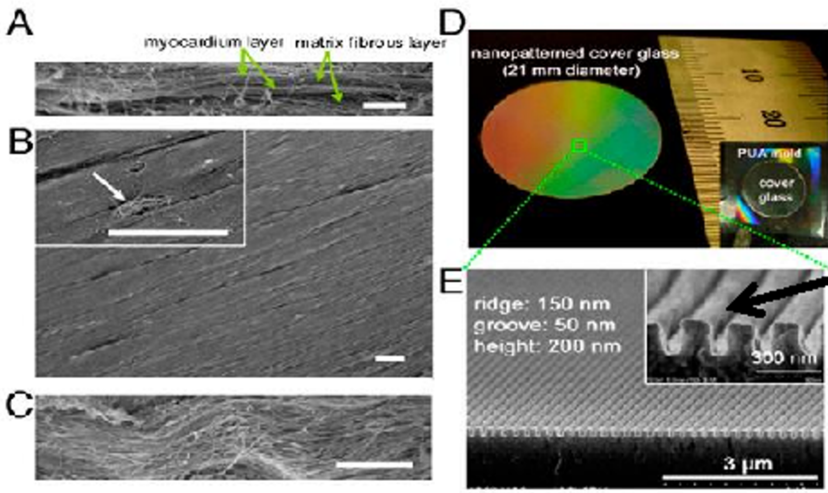
Ingredient III, Nanopatterning

Q: how do we get cells in culture to form complex geometries?

We can use nanopatterning as a substrate for cell monolayer formation.

* cells use focal adhesions, lamellapodia to move across surfaces.

* migration, mechanical forces an important factor in self-organization, self-maintenance.



Gratings at nanoscale dimensions

PNAS 107(2), 565 (2010)

Alignment and protrusions w.r.t nanoscale substrate

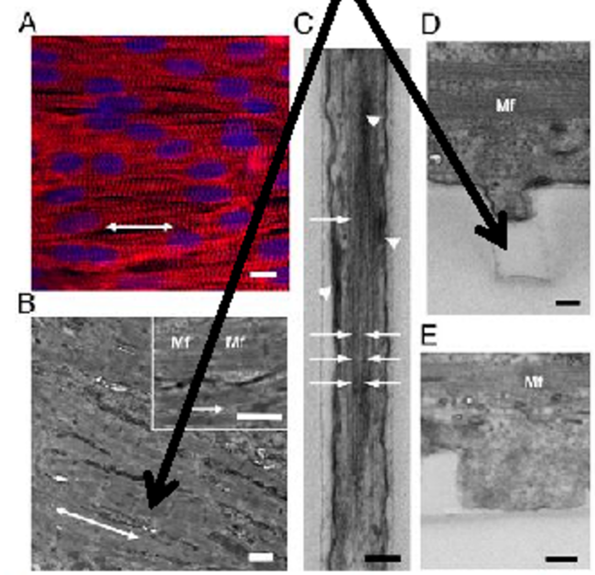


Fig. 3. Cell and cytoskeleton alignment and striations. (*A*) Immunofluorescent images of sarcomeric α-actinin (in red) of NRVMs cultured on the ANFS. Cell nuclei are shown in blue. (*B*) Cross-sectional TEM images of the engineered myocardial tissue grown on the ANFS showing aligned Mf with elongated sarcomeres. Double-headed arrows in (*A*) and (*B*) denote the direction of anisotropic nanopatterns consisting of ridges and grooves. (*C*) An enlarged view of actin bundles (white arrows) and focal adhesions (dark and thick lines indicated by white arrowheads) preferentially formed in parallel to the individual ridges and grooves of the ANFS. (*D*–*E*) Representative cross-sectional view of the PEG sidewalls showing the lower extent of cell protrusion into (*D*) a 400-nm-wide groove than of that into (*E*) an 800-nm-wide groove. [Scale bar: 10 μm in (A); 1 μm in (*B*); 200 nm in (*C*–*E*).]

MWCNTs as Substrate for Neurons

Multi-Wall CNT substrate for HC neurons: Nano Letters, 5(6), 1107-1110 (2005).

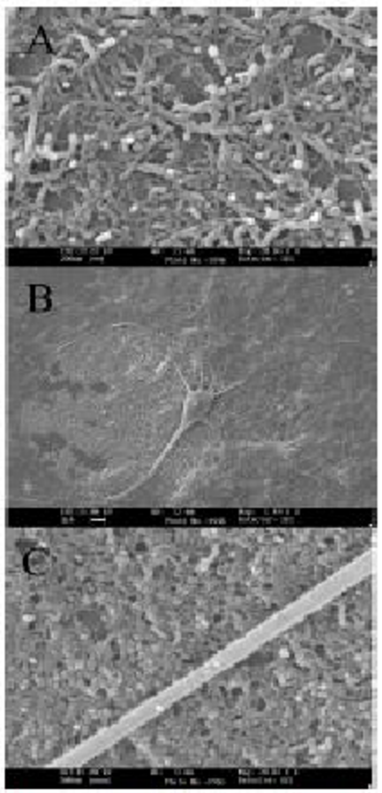


Figure 1. Purified multiwalled carbon nanotubes (MWNT) layered on glass are permissive substrates for neuron adhesion and survival. (A) Micrographs taken by the scanning electron microscope showing the retention on glass of MWNT films after an 8-day test in culturing conditions. (B) Neonatal hippocampal neuron growing on dispersed MWNT after 8 days in culture. The surface structure, composed of films of MWNT and peptide-free glass, allows neuron adhesion. Dendrites and axons extend across MWNT, glia cells, and glass. The relationship between dendrite and MWNT is very clear in the image in (C), were a neurite is travelling in close contact to carbon nanotubes.

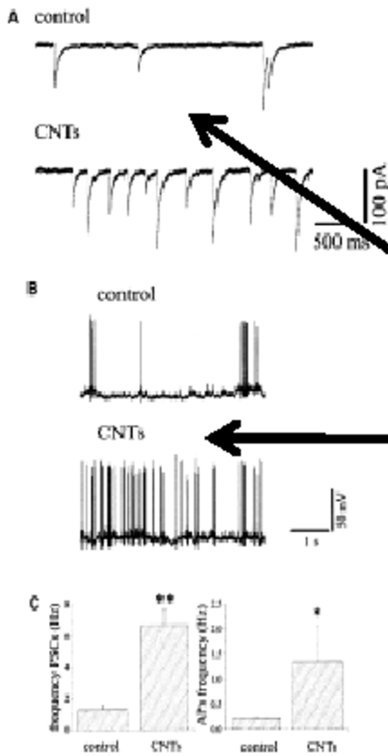
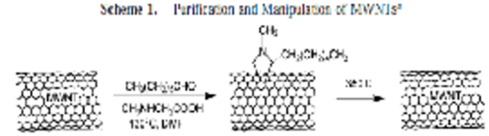


Figure 2. CNT substrate increases hippocampal neurons spontaneous synaptic activity and firing. (A) Spontaneous synaptic currents (PSCs) are shown in both control (top tracings) and in cultures grown on CNT substrate (bottom tracings). Note the increase in PSCs frequency under the latter condition. Recordings were taken after 8 days in culture. (B) Current clamp recordings from cultured hippocampal neurons in control (top tracings) and CNT growth conditions (bottom tracings). Spontaneous firing activity is greatly boosted in the presence of CNT substrates. (C) Histogram plots of PSCs- (left) and APs- (right) frequency in control and CNT cells; note the significant increase in the occurrence of both events when measured in CNT cultures. $^{**}P < 0.0001$ and $^{*}P < 0.05$.

Scheme 1. Purification and Manipulation of MWNTs[a]



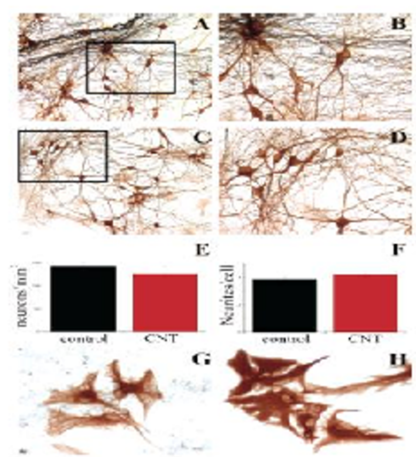
[a] The MWNTs were first functionalized and purified, then deposited on a glass substrate and heated to 350 °C, a process that eliminates the organic part, leaving intact the carbon network.

CNTs functionalized, purified, deposited on glass (pure carbon network desired).

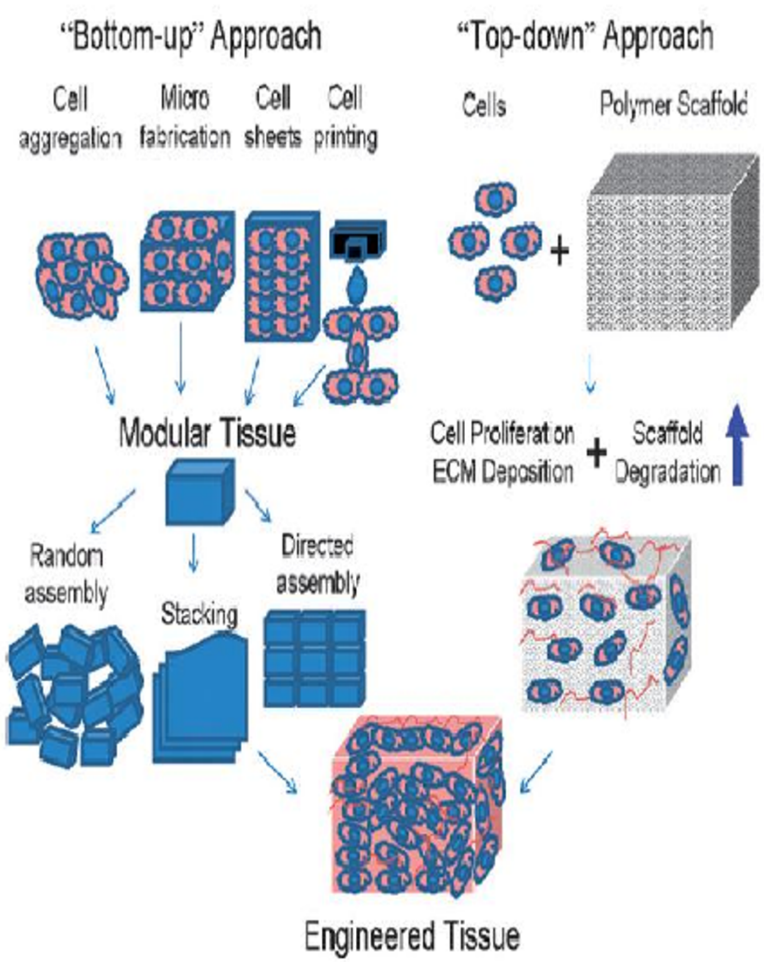
Improvement in electrophysiology: IPSCs (A) and patch clamp (B).

Neuronal density similar between CNTs and control.

* increase in electrical activity due to gene expression, ion channel changes in neuron.

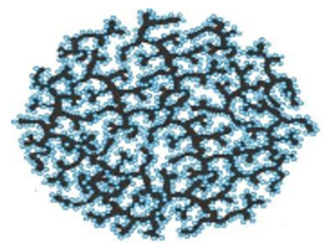


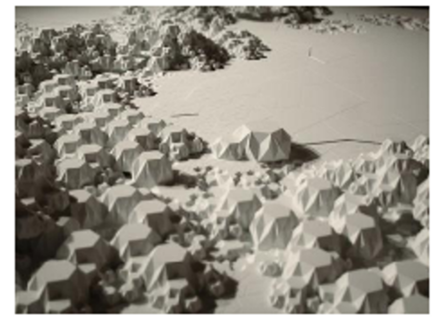
Bottom-up vs. Top-down Approaches



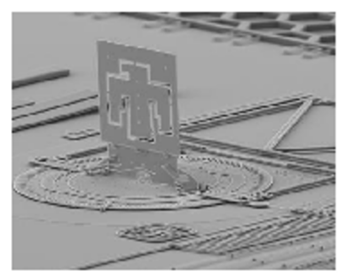
Soft Matter, 5, 13121319 (2009).

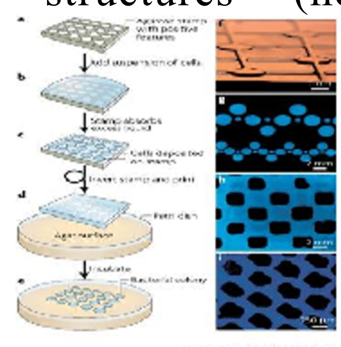
Theoretically, there are two basic approaches to building tissues:





1) bottom-up: molecular self-assembly (lipids, proteins), from individual components into structures (networks, micelles).





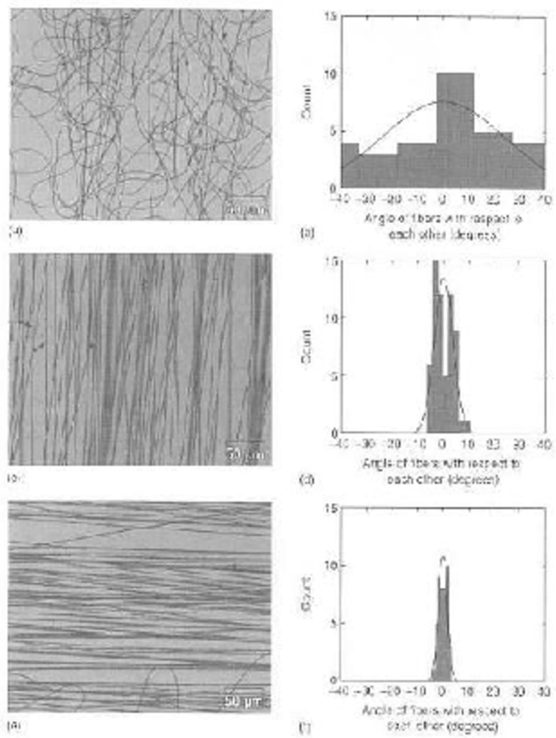
Nature Reviews Microbiology 5, 209-218 (2007).

2) top-down: allow cells to aggregate upon a patterned substrate (CNTs, oriented ridges, microfabricated scaffolds).

Top-down approach: Electrospinning

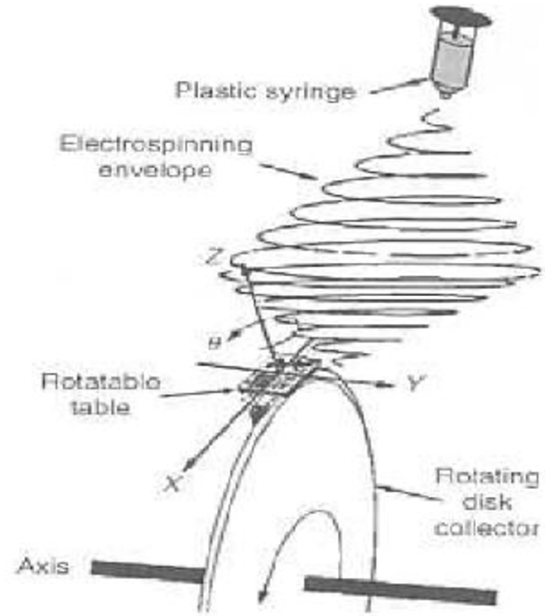
Align nanofibers using electrostatic repulsion forces (review, see Biomedical Materials, 3, 034002 - 2008).

Contact guidance theory:
Cells tend to migrate along orientations associated with chemical, structural, mechanical properties of substrate.



Left: "Nanotechnology and Tissue Engineering: the scaffold". Chapter 9.

Right: Applied Physics Letters, 82, 973 (2003).



Electrospinning procedure:

* fiber deposited on floatable table, remains charged.

* new fiber deposited nearby, repelled by still-charged, previously deposited fibers.

* wheel stretches/aligns fibers along deposition surface.

* alignment of fibers ~ guidance, orientation of cells in tissue scaffold.

Bottom-up approach: Molecular Self-assembly

Protein and peptide approaches commonly used.

Protein approach see review, Progress in Materials Science, 53, 11011241 (2008).

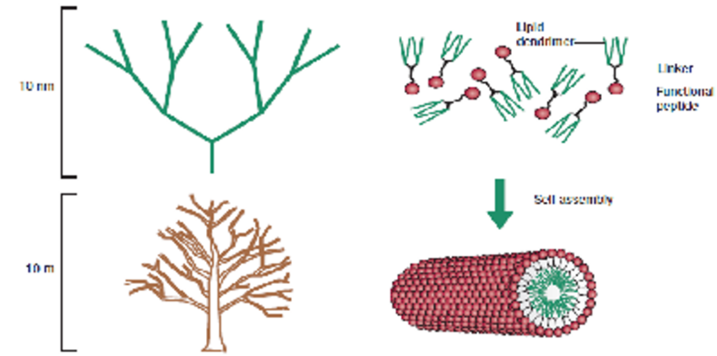
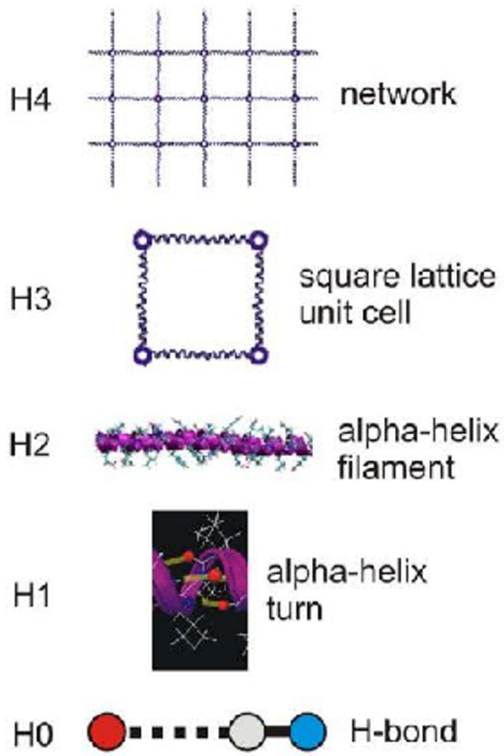


Figure 1 Dendrimers are tree-like molecules that have repeatedly branched structures. The combination of a functional peptide with dendritic lipid groups enables nanoparticles with controlled shapes and sizes to be assembled when the molecules are dissolved in water. The resulting assemblies have a hydrophobic lipid core (green) and a biologically active hydrophilic peptide coating (red).

Nature Nanotechnology, 3, 8 (2008).

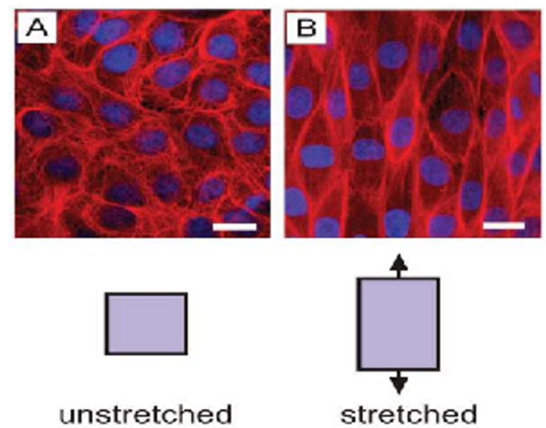


Hierarchical Network Topology, MD simulations. PLoS ONE, 4(6), e6015 (2009).

-helix protein networks in cytoskeleton withstand strains of 100-1000%.

* synthetic materials catastrophically fail at much lower values.

* due to nanomechanical properties, large dissipative yield regions in proteins.

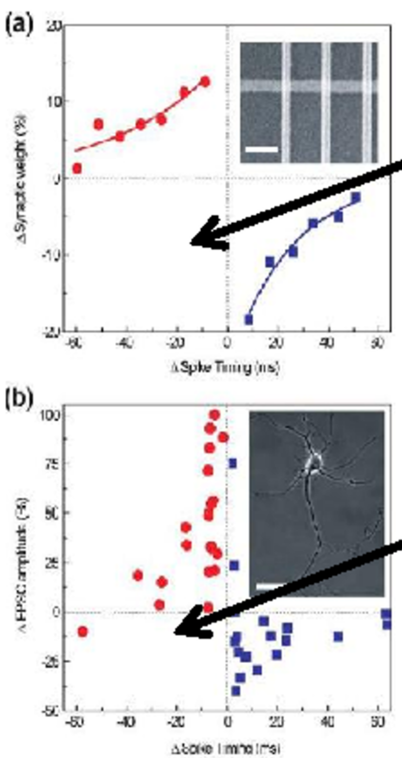


Filament network, in vivo. PLoS ONE, 4(6), e6015 (2009).

Additional Tools: Memristor

Memristor: information-processing device (memory + resistor, Si-based) at nanoscale.

* conductance incrementally modified by controlling change, demonstrates short-term potentiation (biological synapse-like).



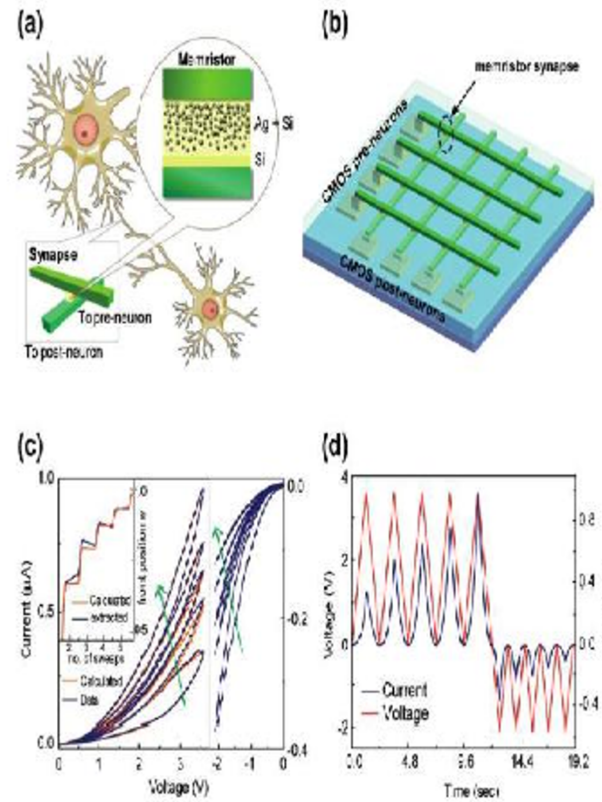
Memristor response

Biological Neuronal response

Learning = patterned (time domain) analog modifications at synapse (pre-post junction).

Array of pre-neurons (rows), connect with post-neurons (columns) at junctions.

* theory matches experiment!



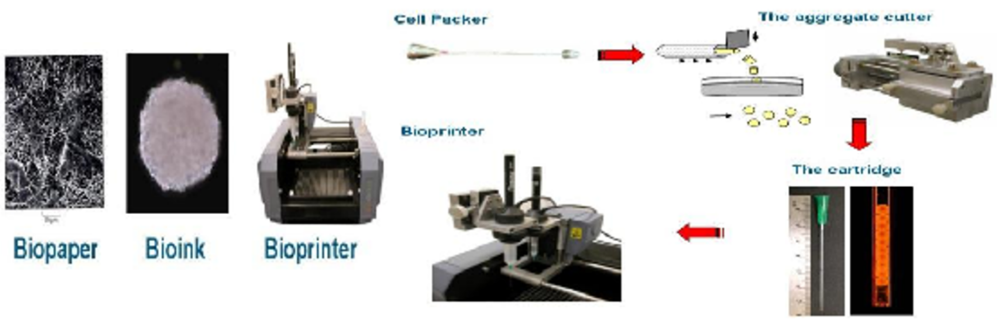
Nano Letters, 10, 12971301 (2010).

Nano Letters, 10, 12971301 (2010).

Additional Tools: Bioprinting

Bioprinting: inkjet printers can deposit layers on a substrate in patterned fashion.

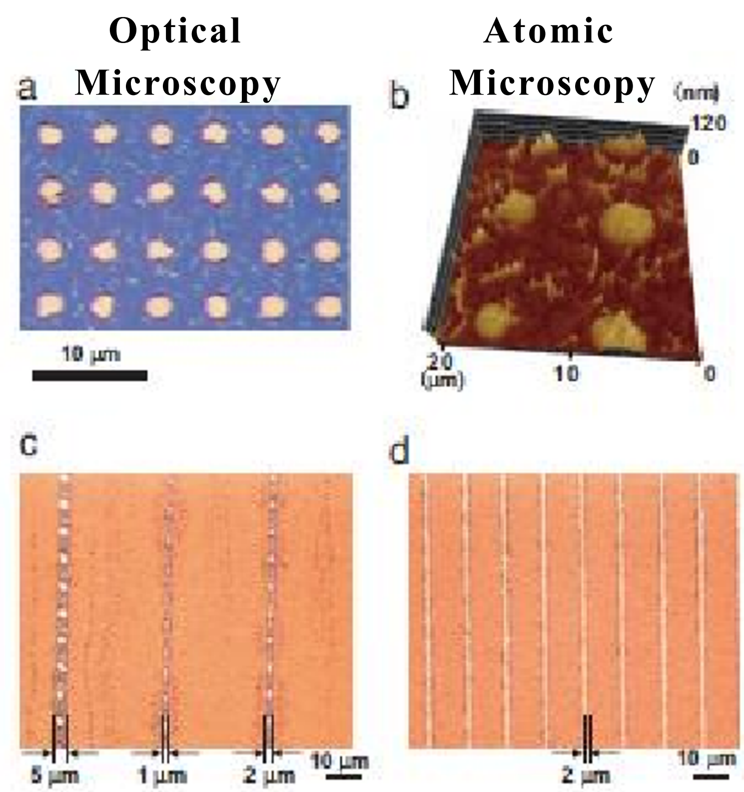
* 3D printers (rapid prototypers) can produce a complex geometry (see Ferrari, M., "BioMEMS and Biomedical Nanotechnology", 2006).



Sub-femtoliter (nano) inkjet printing:
* microfabrication without a mask.

* amorphous Si thin-film transistors (TFTs), conventionally hard to control features smaller than 100nm.

* p- and n-channel TFTs with contacts (Ag nanoparticles) printed on a substrate.



PNAS, 105(13), 4976 (2008).

Conclusions

Nano can play a fundamental role in the formation of artificial tissues, especially when considering:

* emergent processes: in development, all tissues and organs emerge from a globe of stem cells.

* merging the sensory (electrical) and biomechanical (material properties) aspects of a tissue.

Advances in nanotechnology might also made within this problem domain.

* scaffold design requires detailed, small-scale substrates (for mechanical support, nutrient delivery).

* hybrid protein-carbon structures, or more exotic "biological" solutions (reaction-diffusion models, natural computing, Artificial Life)?