

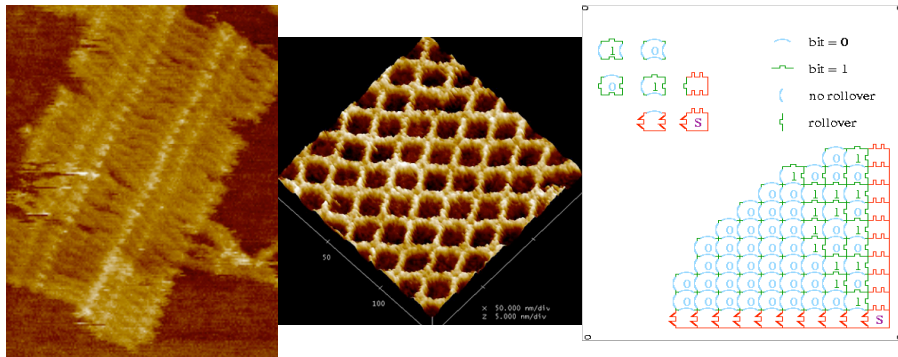
Programmable DNA Lattices: Design Synthesis & Applications

Duke University (PI John Reif)

Subcontracts: NYU (Nadrian Seeman), Caltech (Erik Winfree & Niles Pierce)

Scientific Objectives: *Programmable construction of complex nanostructures of supramolecular (10-100 nm) scale:*

- DNA lattices with complex 2D patterning
- periodic 3D DNA lattices.



Tiling Design for Binary Counter

New Ideas: *Molecular Self-assembly:*

- DNA strands self-assemble into *DNA tile* nanostructures.
- DNA tiles self-assemble into periodic and aperiodic lattices.

Algorithmic Self-assembly:

- DNA tiles form DNA lattices with complex patterning.
- Methodology is *programmable* by choice of DNA tiles.

Nanostructure Templating and Patterning:

- DNA lattice superstructure for other complex nanostructures.
- Other molecular nanostructures attach to specific DNA strands within DNA lattices.

Impact to US defense:

Protein structure determination via host-lattice crystals:

- Periodic 3D DNA lattices capture proteins for X-ray diffraction. *Key spin-off:* Structural characterization of antigens from pathogens.

Patterned DNA Lattice Technology:

- DNA nanostructures can serve as scaffolds for molecular sensors and actuators.

Key Spin-off: identifying pathogens (e.g., bacteria).

- DNA lattices can be used as scaffolds for positioning molecular electronics components into complex circuits.

Key Spin-off: Molecular Nanoelectronics.

Schedule:

Design of novel DNA tiles & lattices and support software

Optimization algorithms for DNA design implemented.

Patterned 2D DNA lattice: modest size (64 tiles)

Periodic 3D DNA lattices: diffracting to 2.5 Å

Characterization of error rates of self-assemblies

Patterned 2D DNA lattices: moderate size (512 tiles)

Periodic 3D DNA lattice: diffracting to 2.5 Å

Self-assembly of 4-bit demultiplexing RAM

Patterned 2D DNA lattices: thousands of tiles

Periodic 3D DNA lattices: diffracting to 2.5 Å

2001 2002 2003 2004