Dielectrophoretic Trapping and Electrical Conductivity of DNA Origami Structures

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Motivation

- Top-down approach (resolution limit, packing density)
- Nanoelectronics / molecular electronics
- Bottom-up approach (self-assembly)
- Biomolecules, CNTs, ... (1 nm 10 μm)
- DNA: superior self-assembly properties

- Controlled positioning on the chip and connecting to other circuitry
- Manipulation of single molecules or larger constructions needed
- Use DEP!



Dielectrophoresis (DEP)



- DEP-force: $F = (p \bullet \nabla)E$ $= (\alpha/2) \nabla (E^2)$
- Brownian motion: $F = k_B T/(2r)$



Basics of DEP Trapping of DNA

- Negatively charged DNA
- Counterion cloud in the aquatic buffer: (Hepes/NaOH, low σ)
- Polarizability increases due to the counterion cloud
- Gold fingertip-type electrodes (e-beam lithography)
- Linker molecule (S-Au bond)





S. Tuukkanen et al. Nanotechnology 18, 295204 (2007).



DNA Origami

Paul W. K. Rothemund, Nature | Vol 440 | 16 March 2006



DNA Origami Structures

- DNA origami as a nanobreadboard
 - non-periodic patterning!
- How to connect origami to outer world?
- Does origami structure conduct electricity?

Example of patterning: Biotin-modified staple strands + streptavidin

A. Kuzyk, K.T. Laitinen, and P. Törmä, Nanotechnology, 20, 235305 (2009).







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DEP Trapping of DNA Origami Structures



- Origami annealing buffer (10x TAE Mg++) changed to the buffer of lower conductivity (Hepes + NaOH + Magnesium acetate)
- Origami structures are ligated
 - Polynucleotide Kinase + DNA Ligase
- Strands with thiol groups are incorporated at each side to allow attachment to gold electrodes



More Results of DEP Trapping

- Height of origami structure: 2 nmDNA sustains its natural form
- Drastic dependence on the ACfrequency and the applied voltage
- Trapping yield: ~10 %



A. Kuzyk, B. Yurke, J.J. Toppari, V. Linko, and P. Törmä *Dielectrophoretic Trapping of DNA Origami*, Small **4**(4), 447-450 (2008).

Electrical Measurements: DC-conductivity

- Rectangular structures
- Relative humidity ~90 %
- Non-linear *IV*-curves
- DC-resistance: 2 GΩ 10 GΩ
 for control sample: over 10 GΩ
- Current vs. Humidity
 - ionized water molecules main charge carriers
- Linker molecule has a large resistance
- No information about the electronic conductivity



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AC Impedance Spectroscopy

- RH = 90 %
- Frequency range: 0.01 Hz -100 kHz
- Amplitude: 50 mV
- Control sample
 - -> Modified Randles circuit
- DNA origami structure
 - -> $R_{\rm DNA},\,R_{\rm C}$ and $Q_{\rm C}$ have to be added

a)

b)



- $W_{diff} = diffusion of ions$
- R_{ct} = charge transfer resistance through double-layer
- C_{dl} = double-layer capacitance
- $C_e = self-capacitance$
- \bullet $R_{\rm c}$ and $Q_{\rm c}$ modify the double-layer



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More AC-IS Results

- Linker molecule does not conduct directly, but there is a charge transfer process in AC
 Measurement of electronic conductivity
- R_{DNA} ~70 MΩ
 Electronic conductivity
- R_{ct} dominates in DC, R_s, R_{ct} ~ GΩ
 R_s + R_{ct} ≈ 10 GΩ (DC-path)
- W_{diff} increases due to mobile counterions around DNA
- Over 90 % relative humidity increases diffusivity, Q_C -> W



 $Z_{CPE} = 1/[Q(j\omega)^n]; Z_W = 1/[W(j\omega)^{1/2}]$

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V. Linko, S.-T. Paasonen, A. Kuzyk, P. Törmä, and J.J. Toppari, *Characterization of the Conductance Mechanisms of the DNA Origami by AC Impedance Spectroscopy*, to appear in Small (2009).

Conclusions

- DEP is a feasible tool for controlled positioning and immobilization of DNA origami structures
 - First example of trapping of a complex selfassembled molecular structure
 - DEP as a bridge between top-down and bottom-up approaches
- AC Impedance spectroscopy:
 - Individual structures measured
 - Nature of conductivity (electronic and ionic)
 - High impedance of linker molecules
 - (DC-measurement is not enough)
 - Conductivity is sufficiently low (for applications: locally conducting parts by doping?)
 - Good approximation for pure dsDNA also



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