



Understanding initiation mechanisms and controlling properties of microtubule spools

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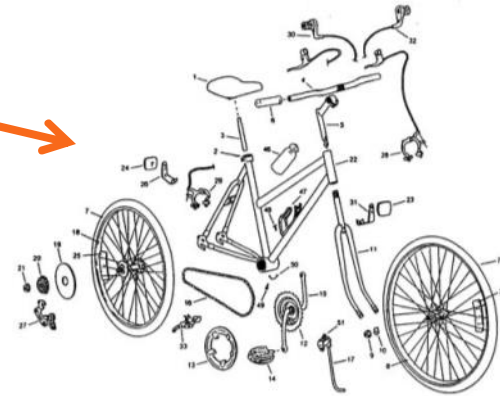
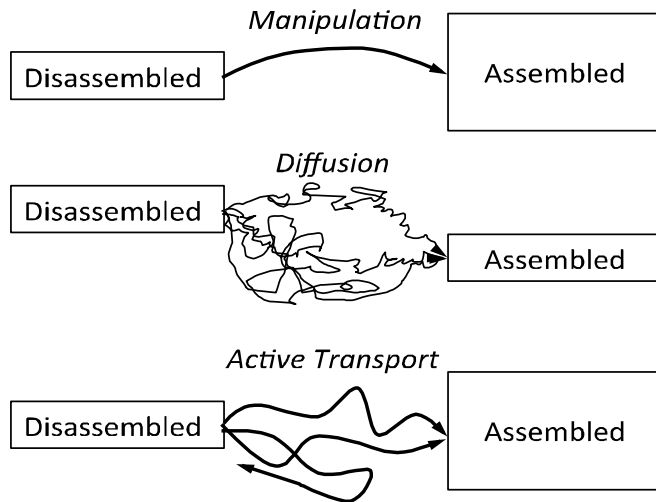
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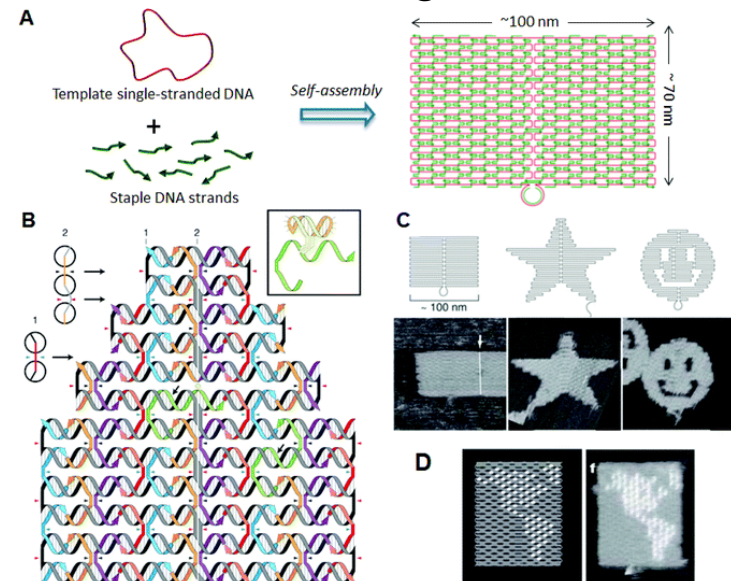


Active self-assembly phenomena



<https://www.youtube.com/watch?v=3Rup3EdA0kw>

DNA origami

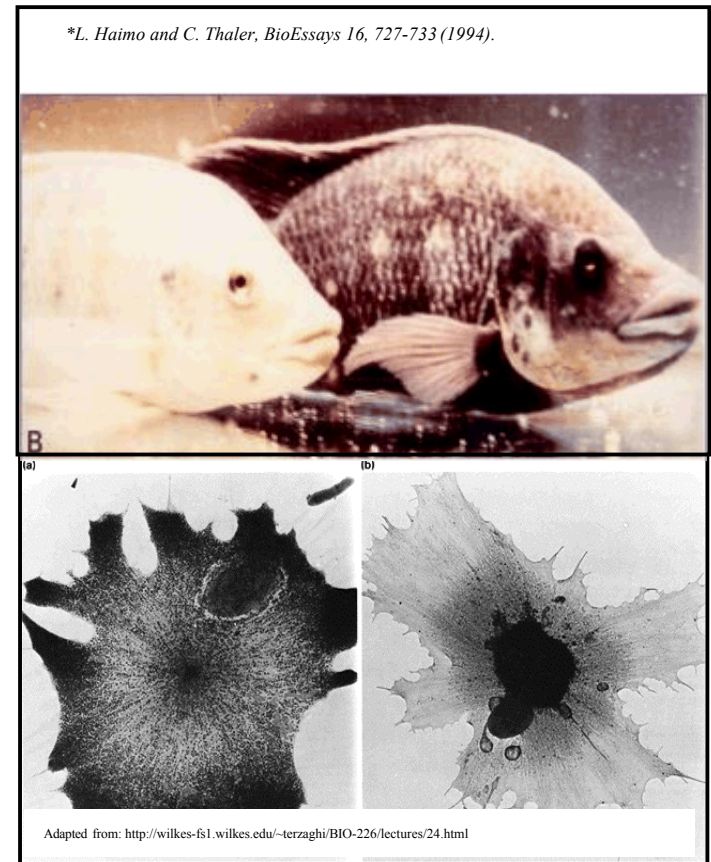


Endo et al, *Biomat. Sci.*, 2013

Biological molecular motors

- Active transport systems:
 - Key to many dynamic physiological processes
 - Nature evolved motor proteins to transport materials in static, nanofluidic environments because fluid flow at those length scales becomes problematic (e.g., pressure-driven fluid flow is hard in nano-channels)
 - Biomotors do not require external power – rather they convert chemical energy (low mass/weight) into mechanical work with great efficiency
 - Biomotors function autonomously without the need for a “user” to control function
 - Ability to transport in complex solutions (e.g., blood, saliva)

Biology: Adaptive and responsive materials properties

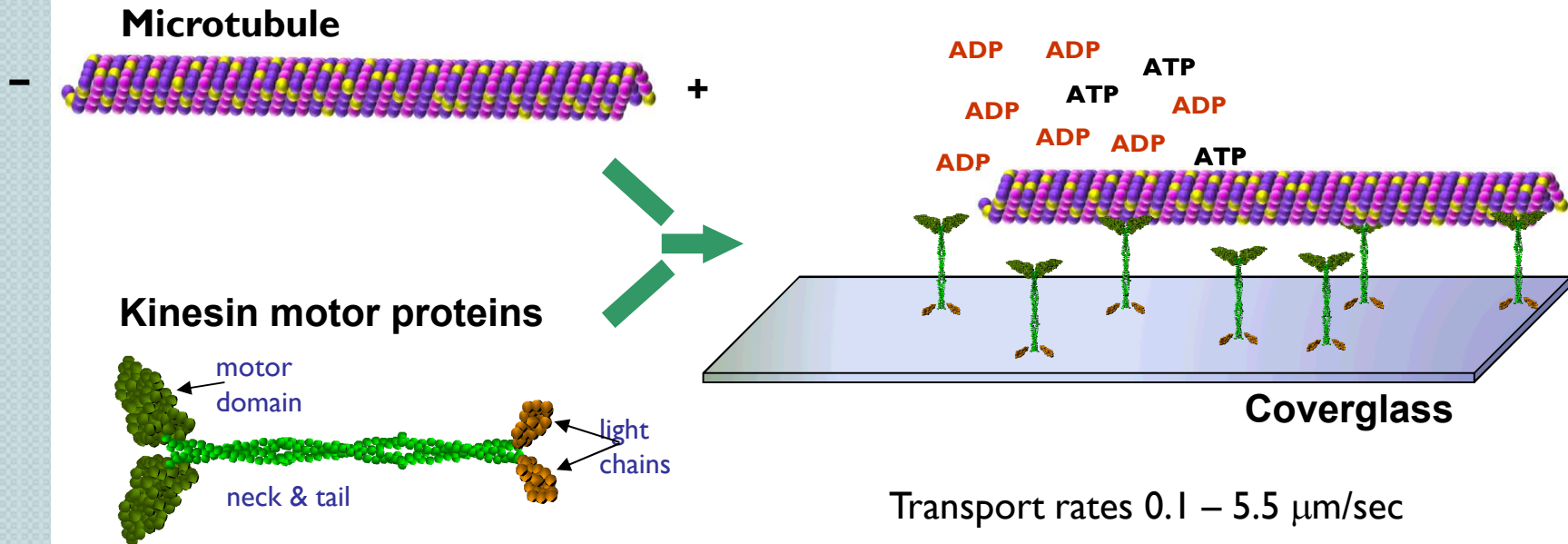


Nanotechnology: Dynamic materials assembly

Kinesin and Microtubules

Biomolecular motors & active transport:

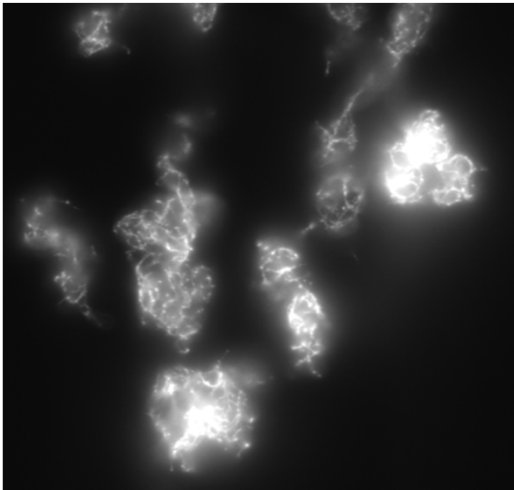
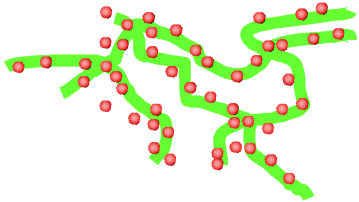
- Convert chemical energy (ATP) to mechanical work (40 pN•nm) with high catalytic efficiency (>50%)



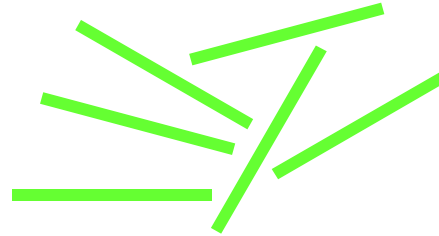
Goal: To utilize biomolecular motor-powered systems to develop novel nanomaterials and nanofluidic systems with biomimetic functionality

Active self-assembly: spools

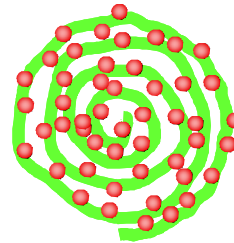
QDs + biotinylated MTs



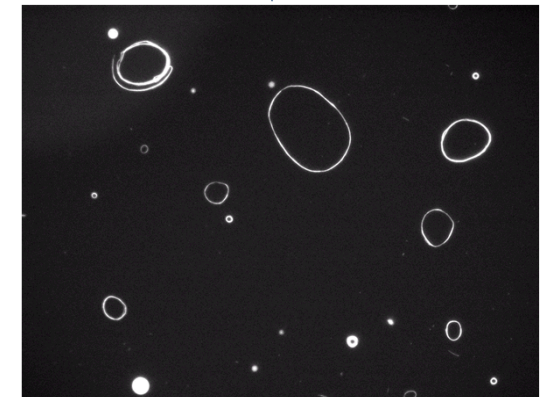
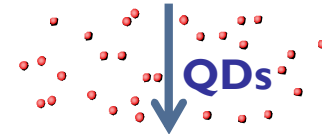
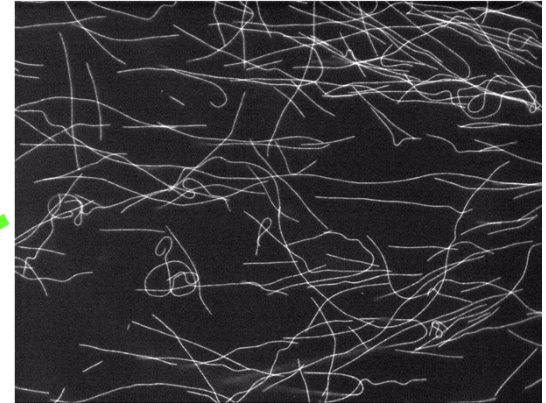
Randomly assembled MTs and QDs (equilibrium)



Energy input via ATP hydrolysis
~50 kJ/mol ATP



Gliding biotinylated MTs

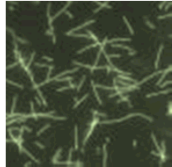


Actively assembled MTs and QDs (+ energy)

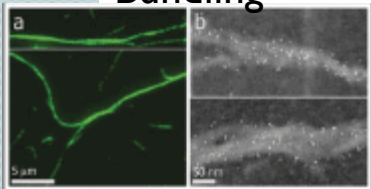
chemical energy → mechanical work → active assembly

Spool initiation and growth

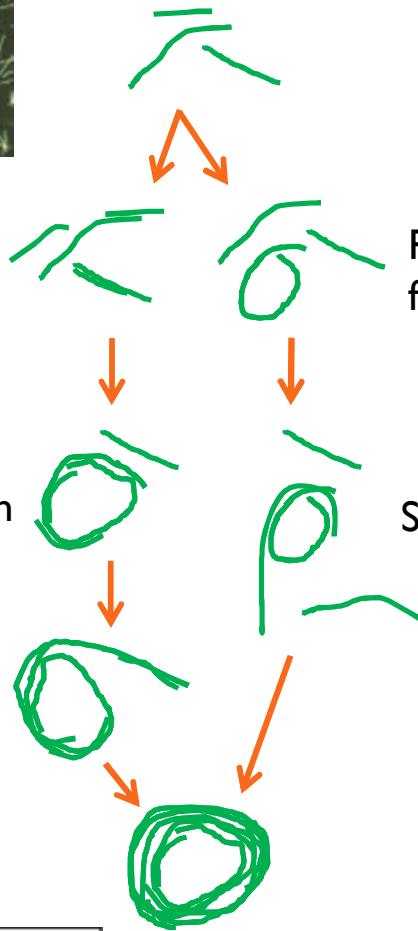
Individual microtubules



Bundling



Ring formation



Ring formation

Spooling

Proposed initiation mechanisms

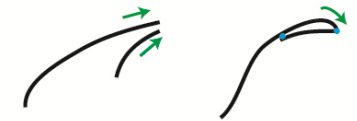
Pinning



Collisions



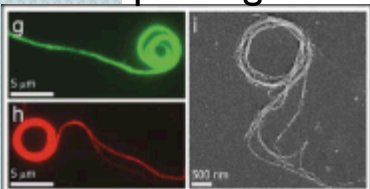
Frozen-in Curvature



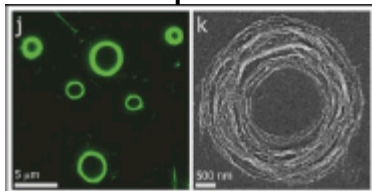
Twist-bend coupling



Spooling



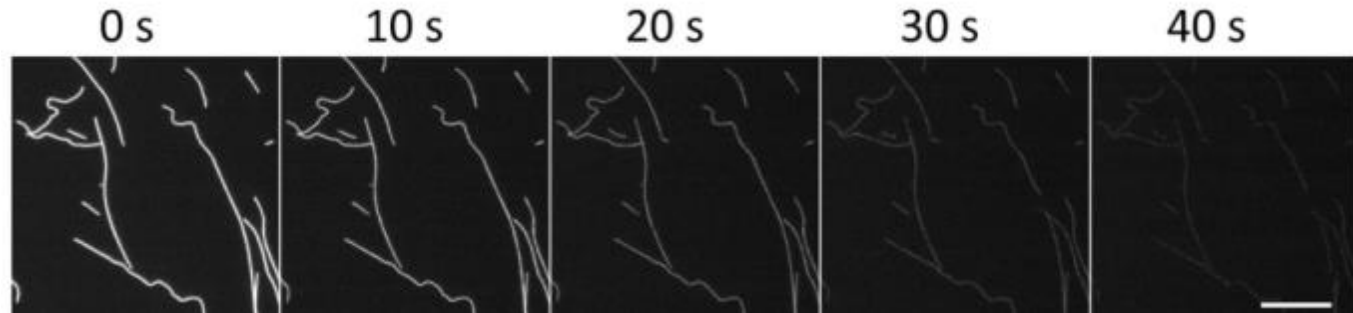
Spool



Initiation mechanism studied primarily by changing experiment parameters, observing effect on resultant spools, and then inferring back to implications for formation.

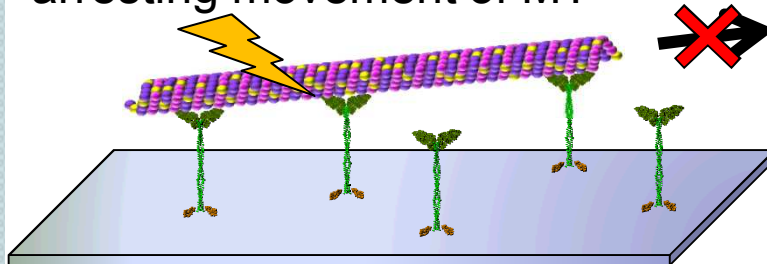
Photodamage

Damage to fluorescent dyes: Decreased brightness

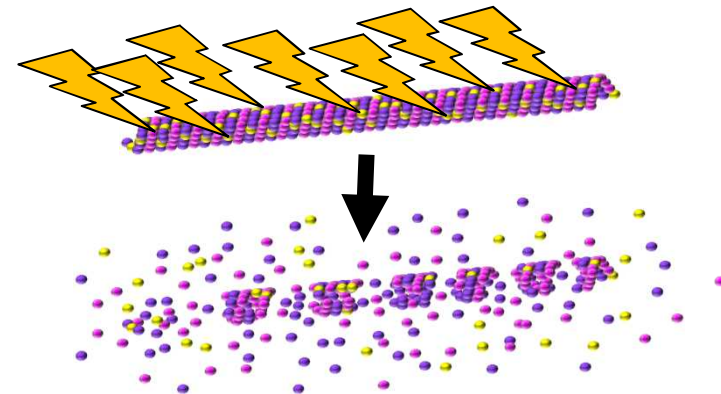


Damage to motors and microtubules

Pinning: fluorophore in excited state or other radical reacts with kinesin, arresting movement of MT

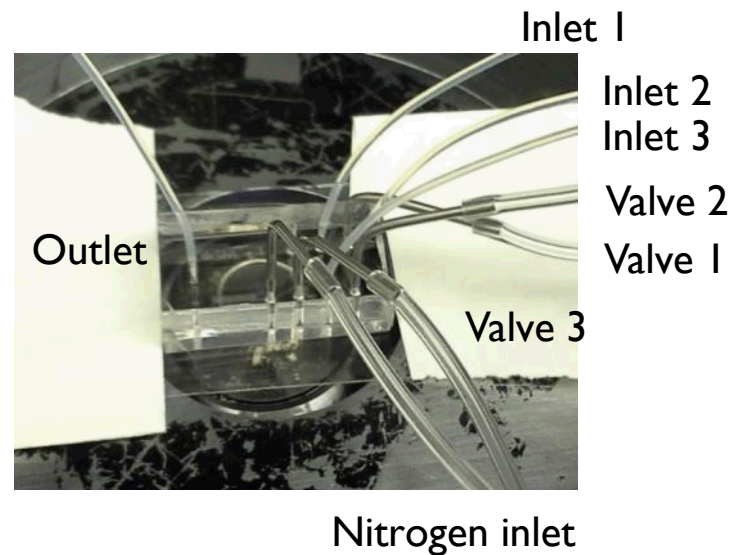
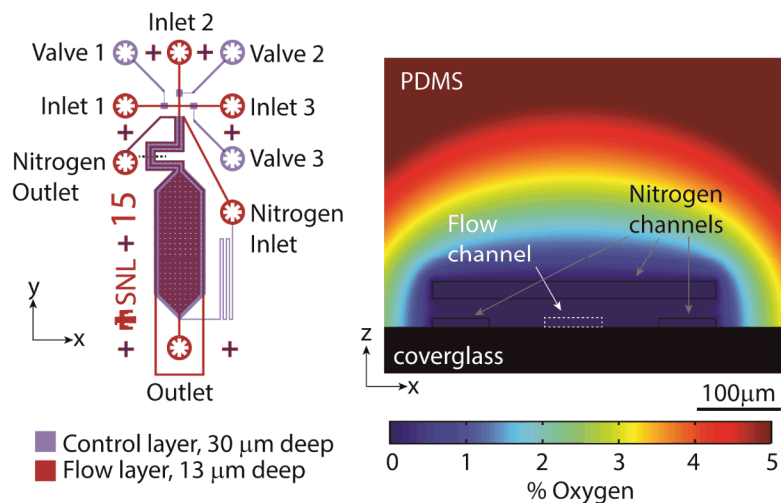


Depolymerization: MT catastrophically disassembles

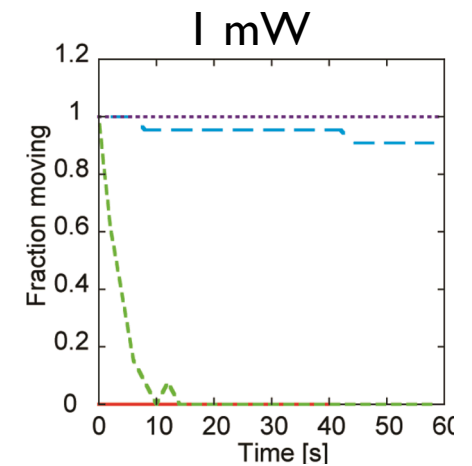
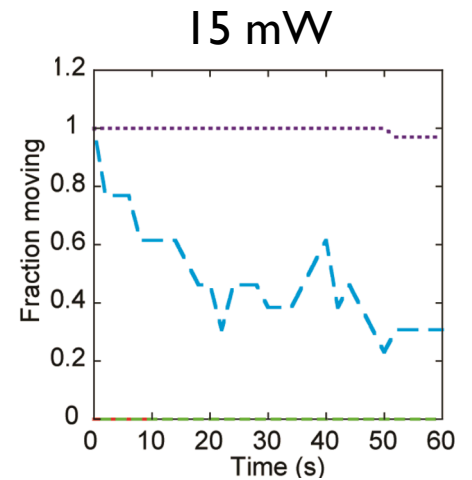
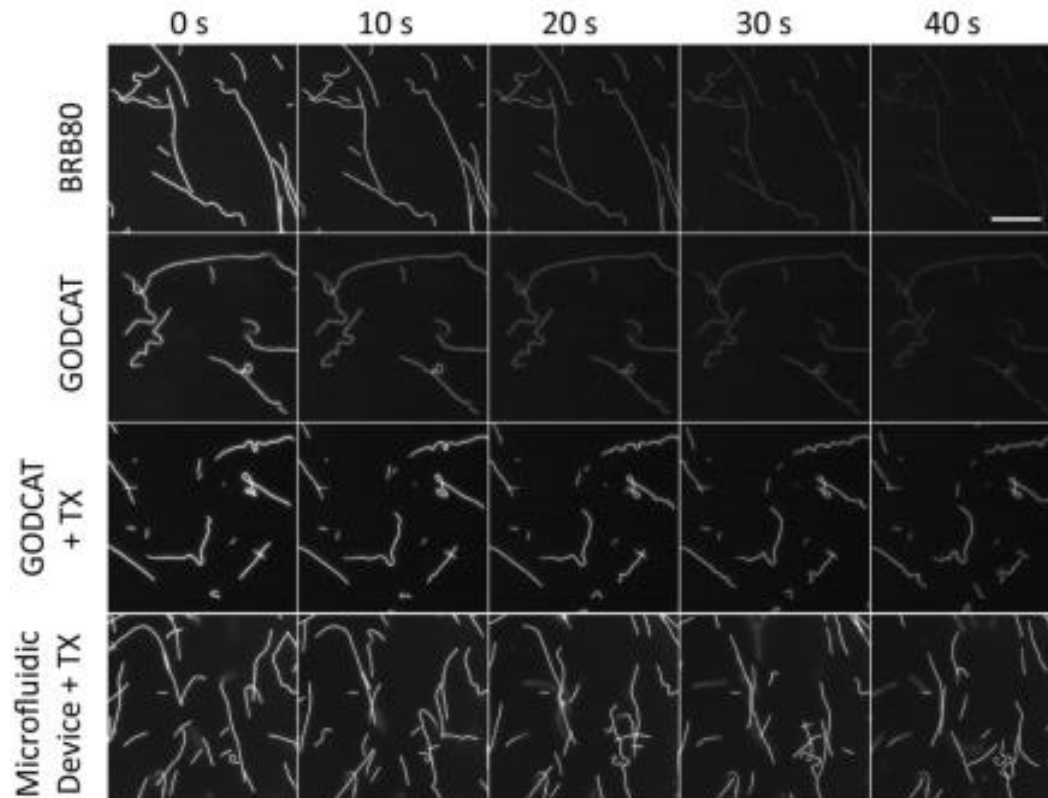


Integrating biomotors into microfluidic devices

- PDMS devices made via soft lithography and molding
 - Rapid prototyping compared to hard materials
 - Multi-layer devices enable integrated valves
 - Permeable to gases
- For lab use to study biomotors
 - Can visualize microtubules while adding QDs
 - Controlled solution exchange and can image during exchange
 - Iterate experiment repeatedly
 - Continuous supply of ATP
 - Improved photostability
 - Less photodamage to motors (less pinning)
 - No enzymatic oxygen scavenger system needed (which stops working and acidifies solution after 1-2 hrs)

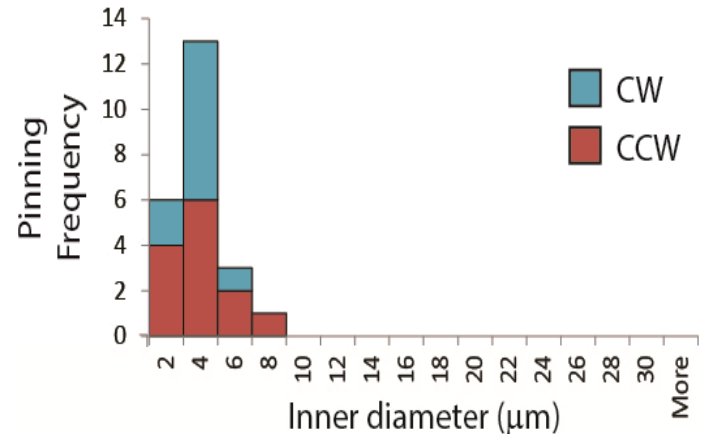
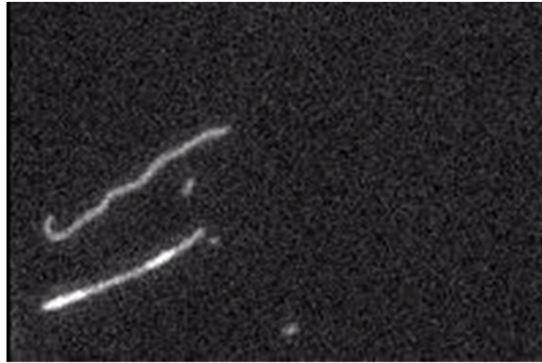


Improved photostability



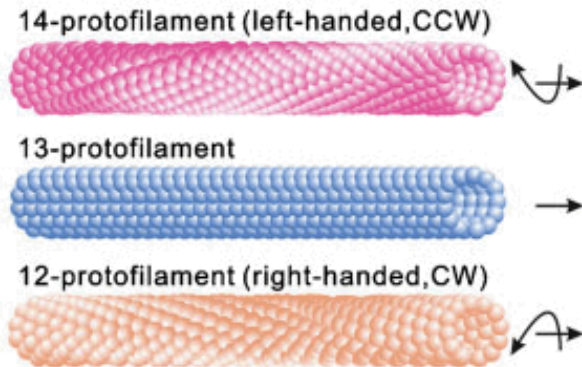
- Plain buffer (BRB80)
- GODCAT
- GODCAT + Trolox
- Microfluidic device + Trolox

Formation mechanism: pinning

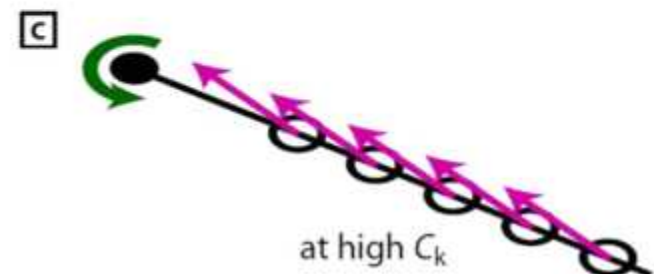
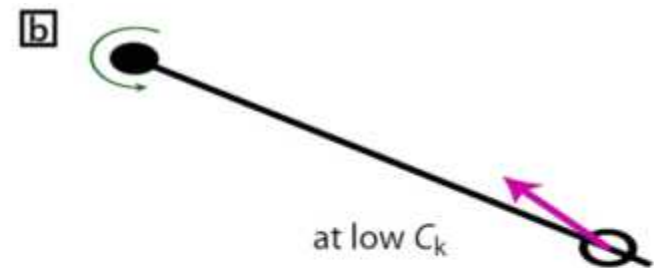
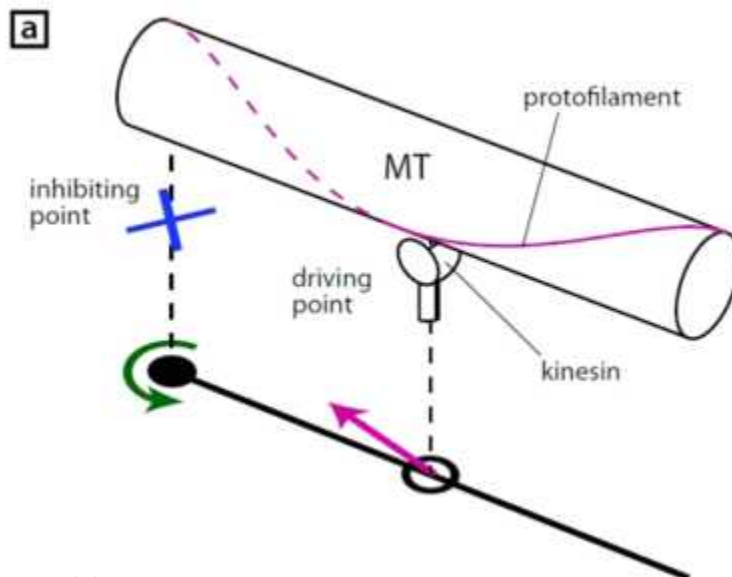


- Diameter $2.7 \pm 1.5 \mu\text{m}$ (mean \pm SD)
- The diameter of rings formed by pinning is determined the buckling radius of a MT
 - contributes to the uniformity of size of rings formed by this mechanism
- Rate at which microtubule encounters a dead motor is a stochastic process
- Pinning is greatly increased by photodamage to motors
- The majority of rings formed by pinning rotated in the CCW direction (62%)

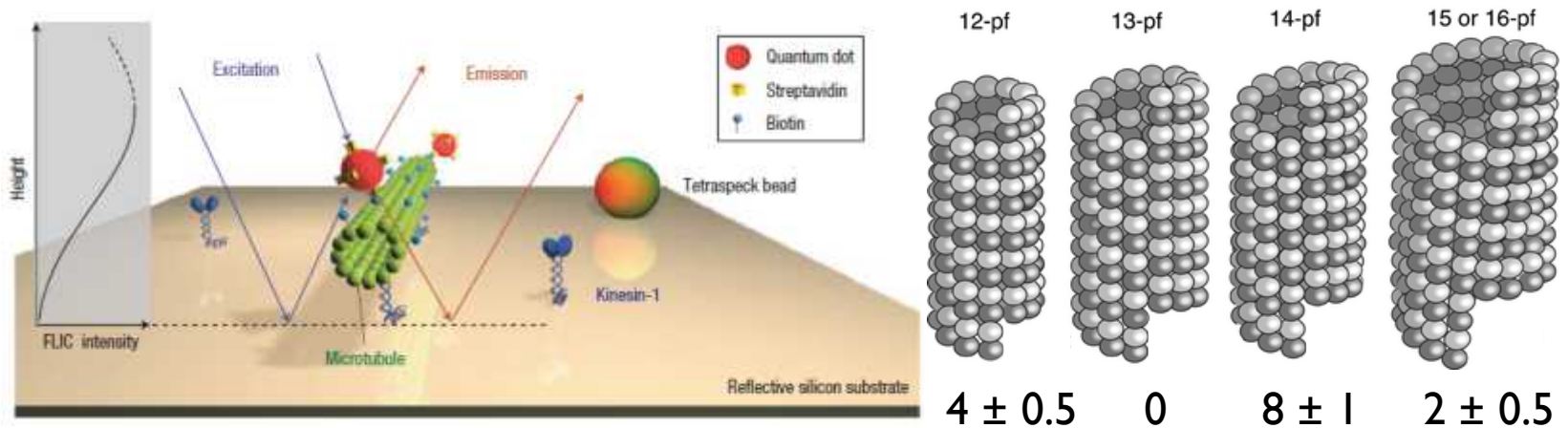
Biased rotation direction



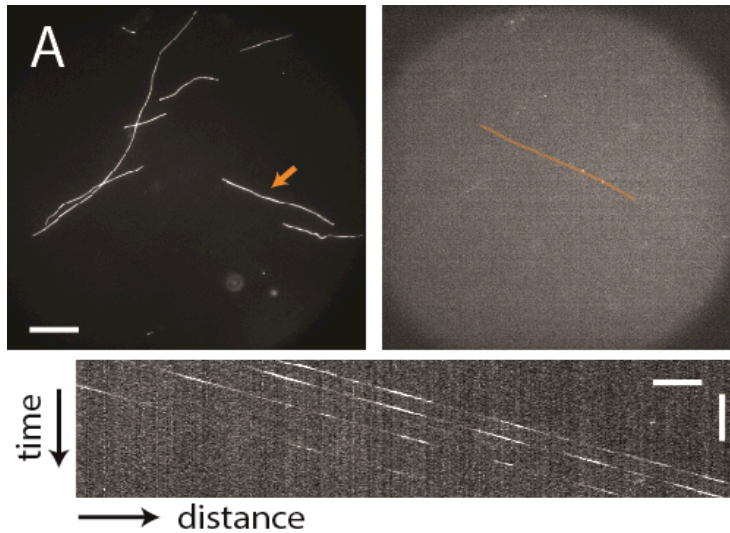
- Microtubules rotate axially depending on protofilament #
- Protofilament # has been observed to affect ring rotation direction
- Wada et al proposed mechanism by which an inhibiting point would produce directional bias in ring rotation direction
 - Amount of bias is dependent on kinesin surface density



FLIC of individual MTs

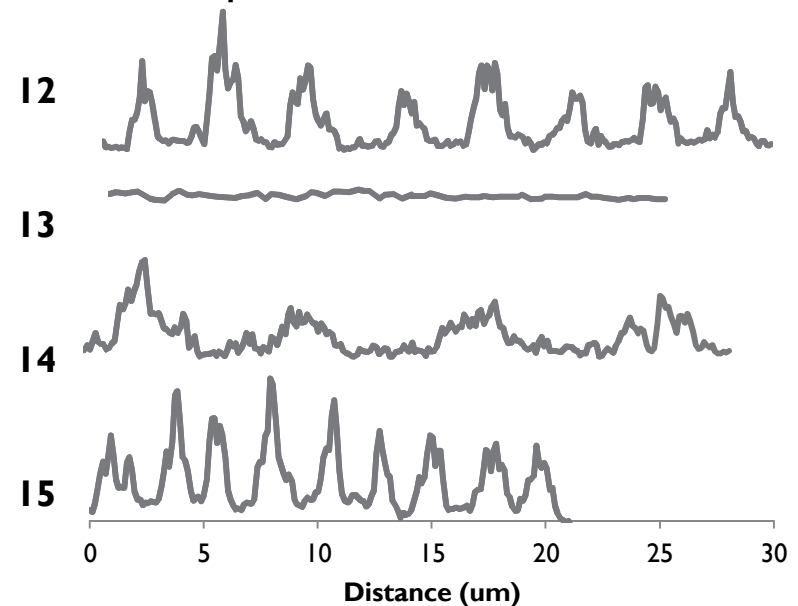


Nitzsche 2008

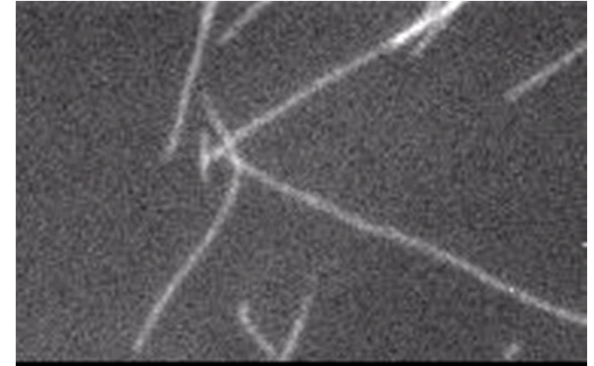
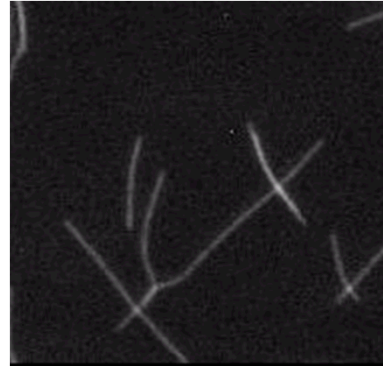


Protofilament #	13	14	15
# of MTs	37	51	14

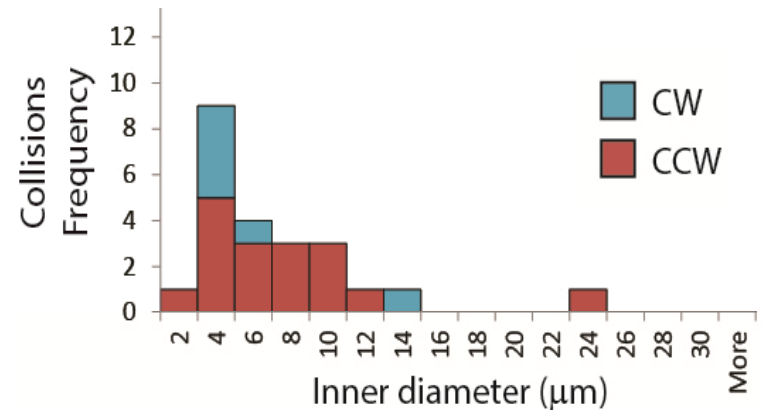
QD intensity traces for various protofilament #'s



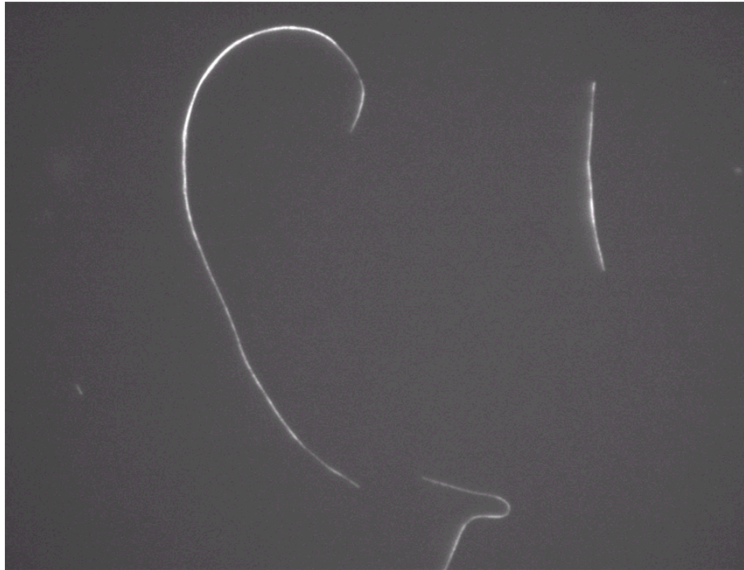
Formation mechanism: collisions



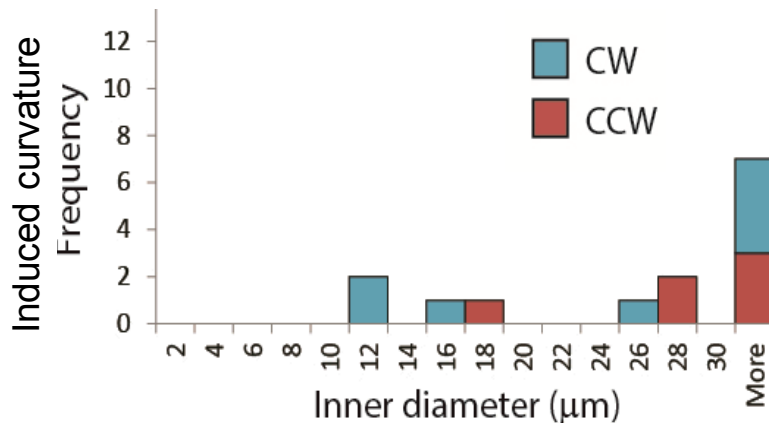
- Diameter $6.2 \pm 4.8 \mu\text{m}$
- Spools formed via collisions displayed a biased rotation direction (74% CCW)
 - Getting stuck on another MT can serve as an inhibiting point



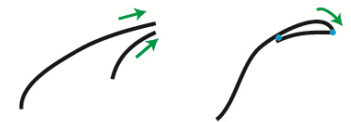
Formation mechanism: induced curvature



- Closing events were rare and thus difficult to capture
- This mechanism wasn't previously observed
- Diameter $33 \pm 20 \mu\text{m}$
- 58% CCW
- Two different mechanisms could underlie induced curvature:



Frozen-in
Curvature

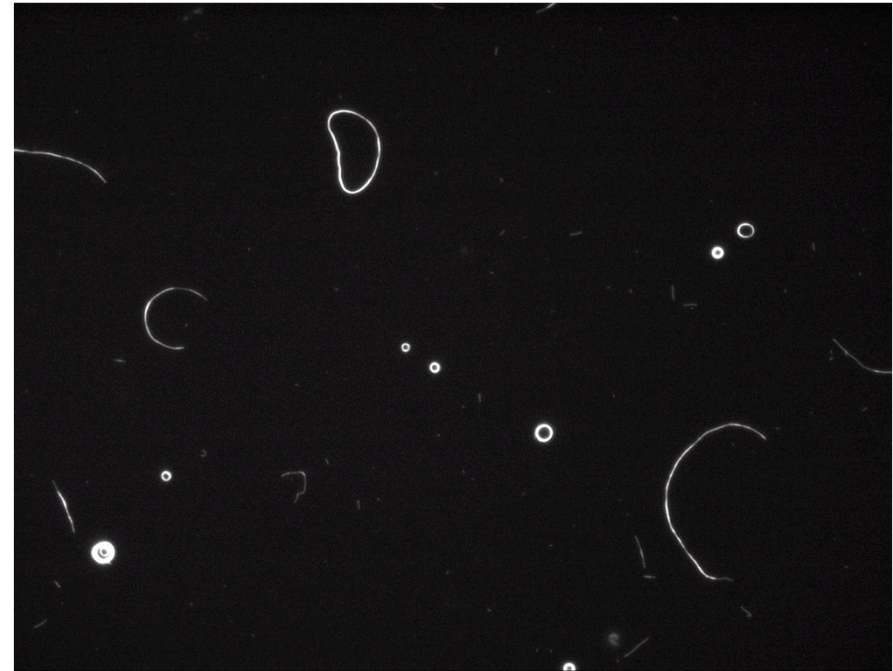


Twist-bend coupling

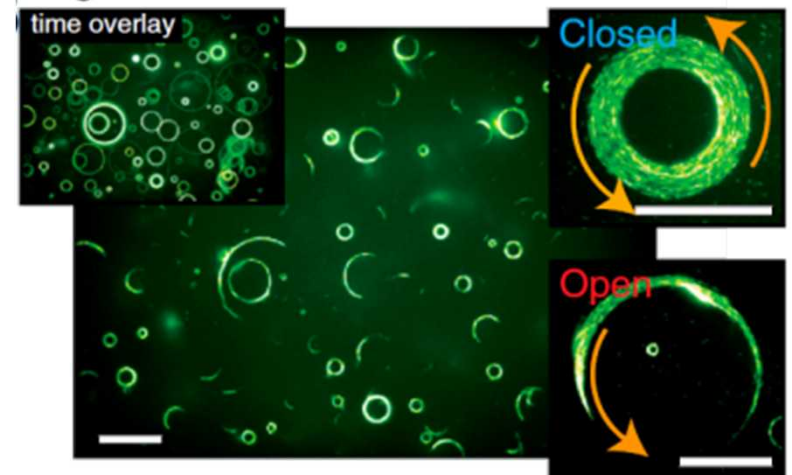


Induced curvature of bundles

- Observed many bundles that followed persistent curved trajectories
- Similar to what was previously seen for myosin-driven actin filaments crosslinked by facsin

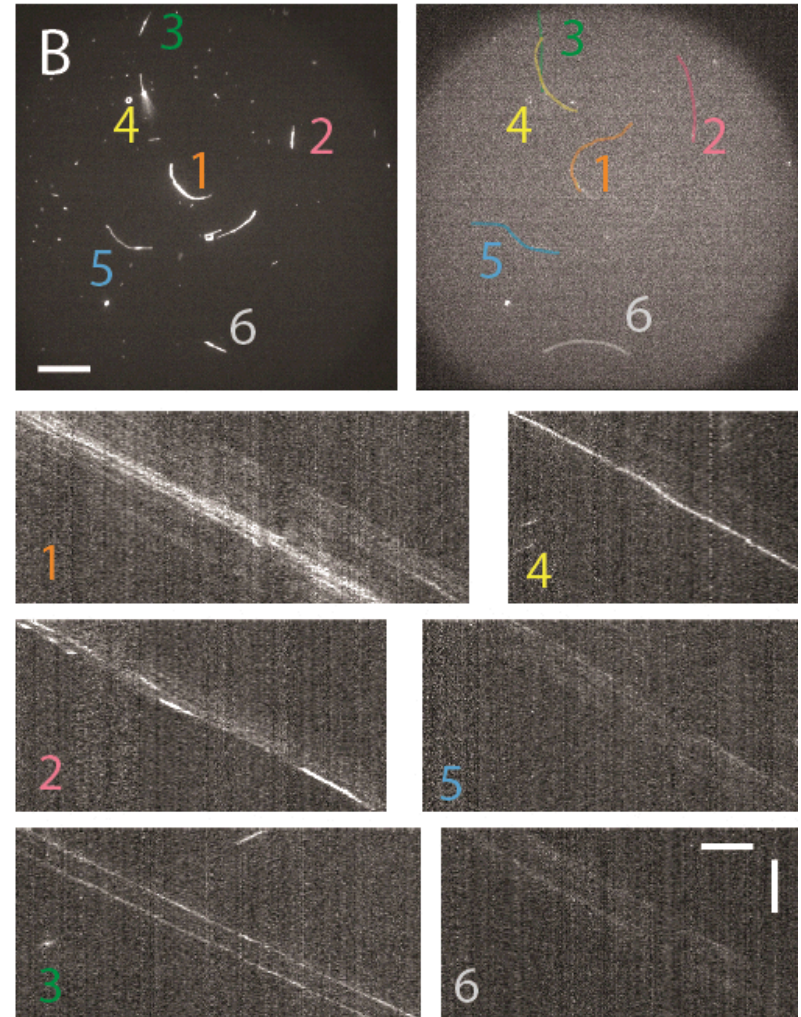


Ring Phase



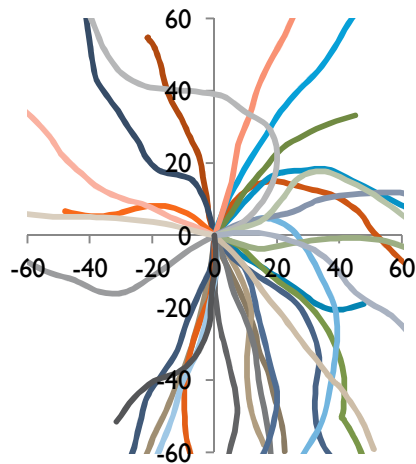
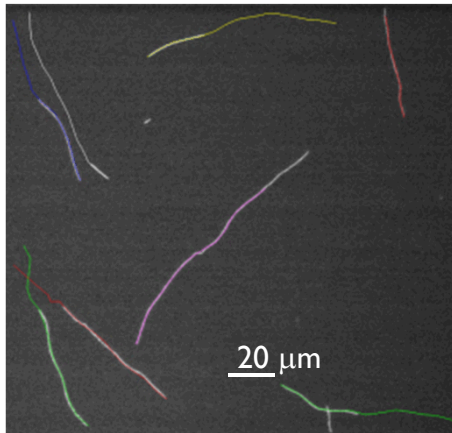
FLIC of bundles

- If twist-bend coupling is responsible to curved trajectory of MT bundles, then bundles should rotate axially as they translate across the surface
- Only ~20% of bundles appear to be rotating
- Does not support twist-bend coupling hypothesis

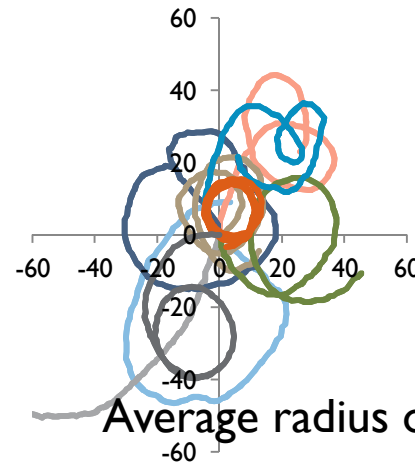
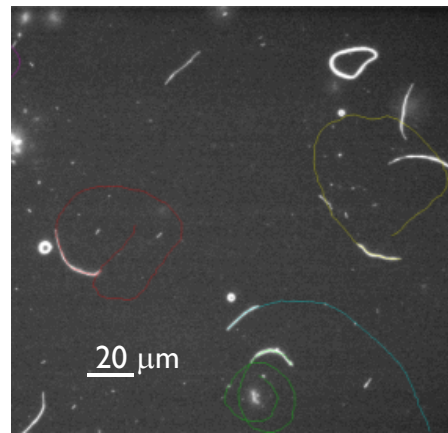


Trajectories

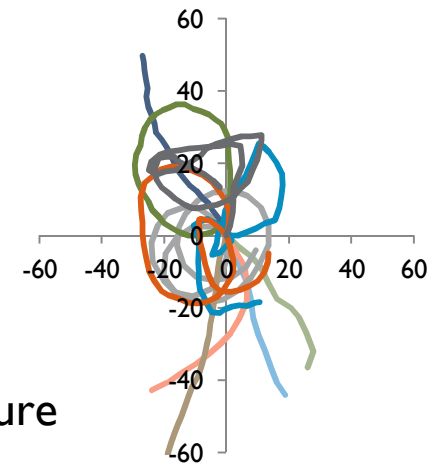
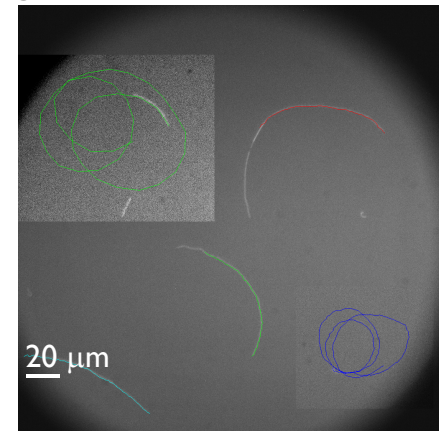
Single MTs



Bundles



Single MTs with QDs



Individual MTs

$51 \pm 23 \mu\text{m}$

Bundles

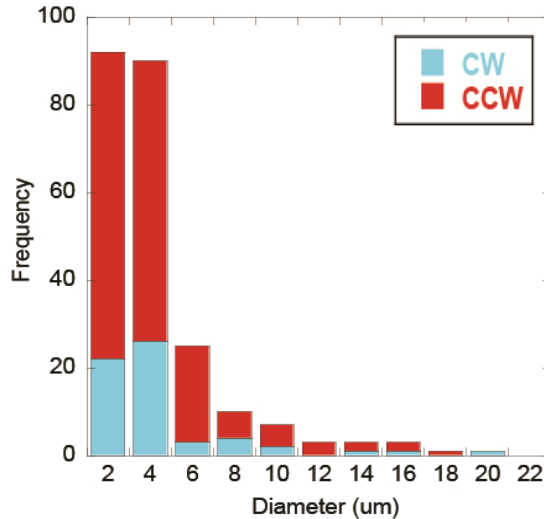
$17 \pm 6 \mu\text{m}$

Individual MTs with QDs

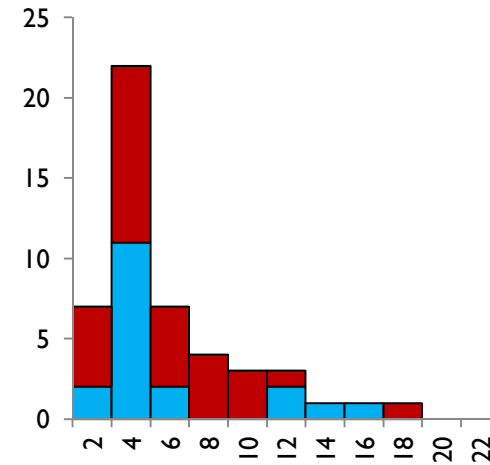
$22 \pm 13 \mu\text{m}$

Comparison between device and flow cell

Conventional glass flow cell



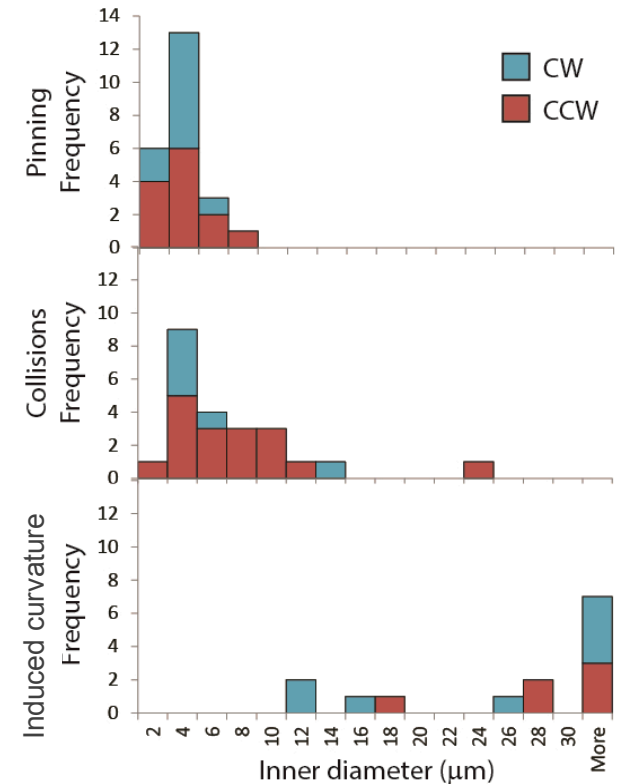
Microfluidic device



- On aggregate, the diameters and rotation direction were consistent in the device and flow cell
 - Device had fewer small (0-2 μm) diameter rings, as expected from decreased photodamage in device

Effect of formation mechanism on spool properties

- Formation mechanism affects properties of spools
- Can tune spool by biasing which formation mechanism dominates
 - Oxidative damage to motors » more pinning and small rings
 - High surface density of MTs » more collisions and medium-sized rings

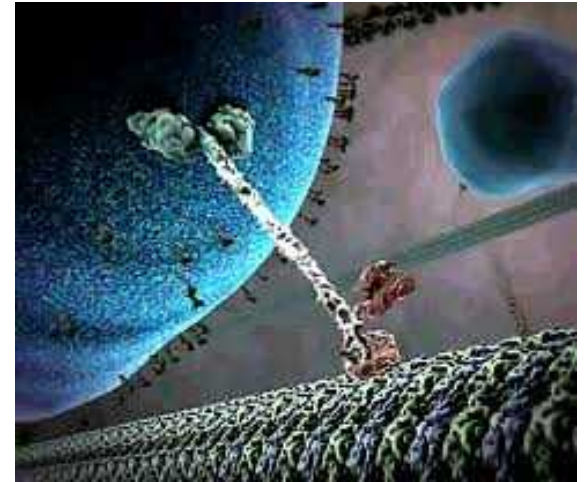


	Pinning	Collision	Induced Curvature
Inner Diameter (μm)	2.7	6.2	31.6
Std dev	1.5	4.8	19.4
%CCW	62	74	58

Acknowledgements

George Bachand

Stephanie Brenner



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