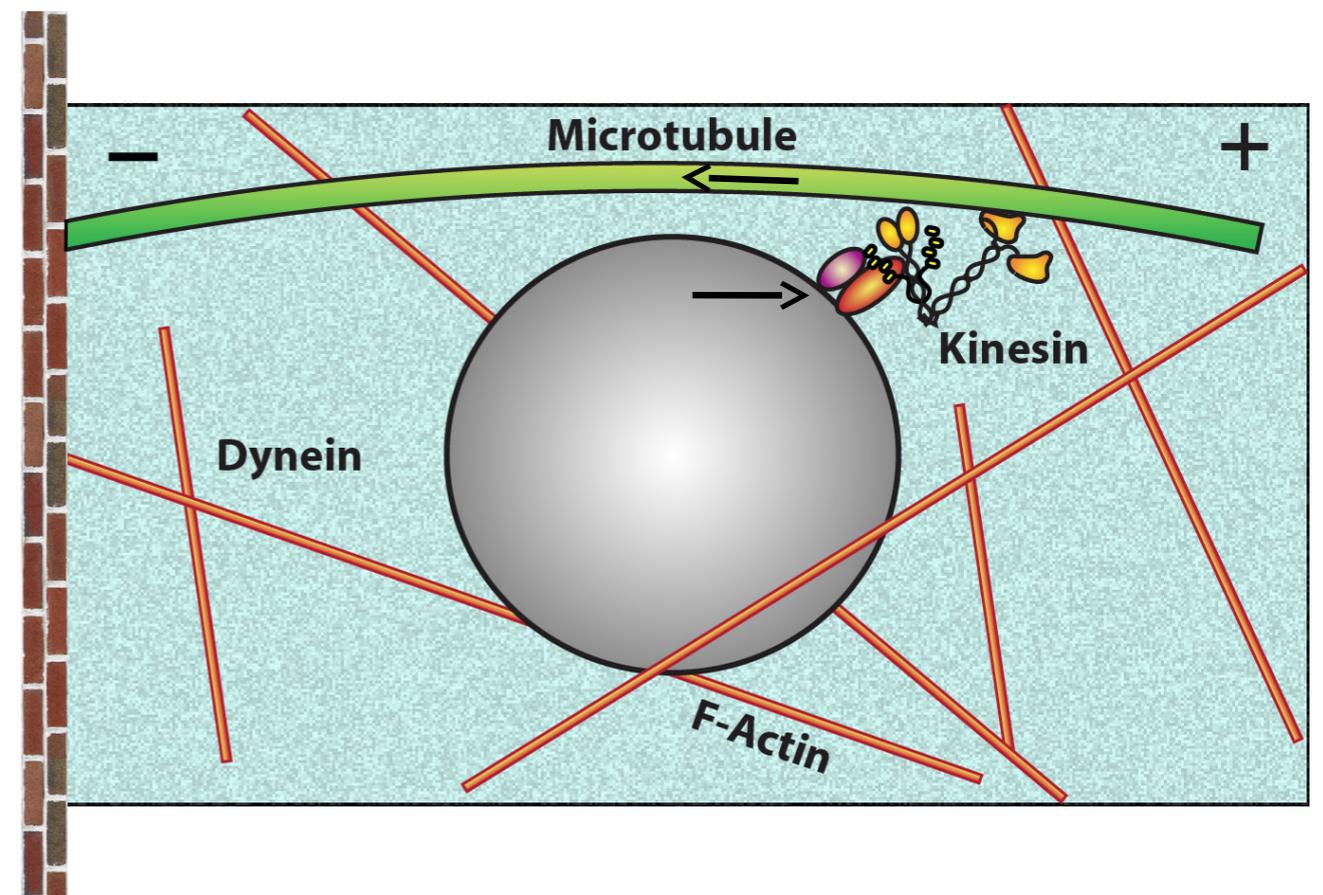
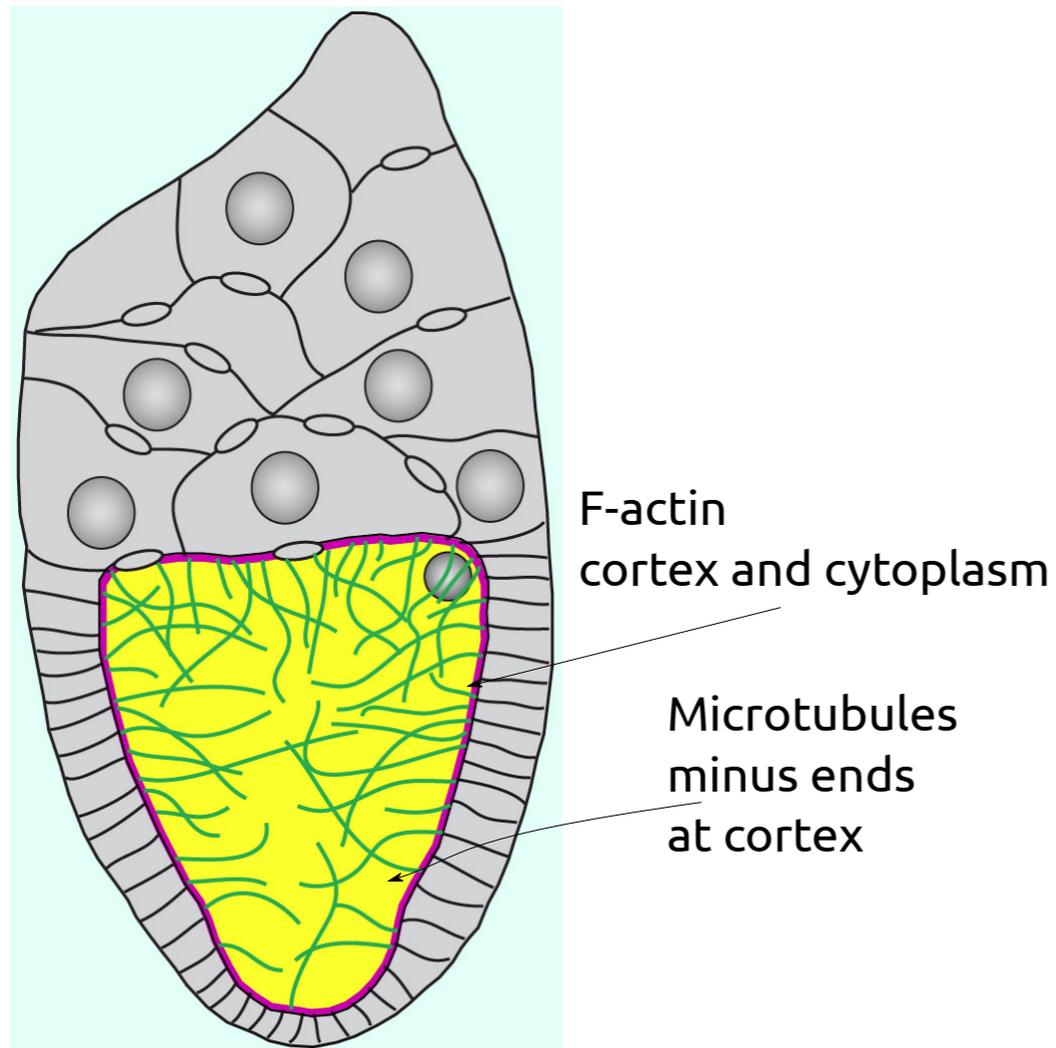


# Cytoplasmic streaming in *Drosophila* Oocytes

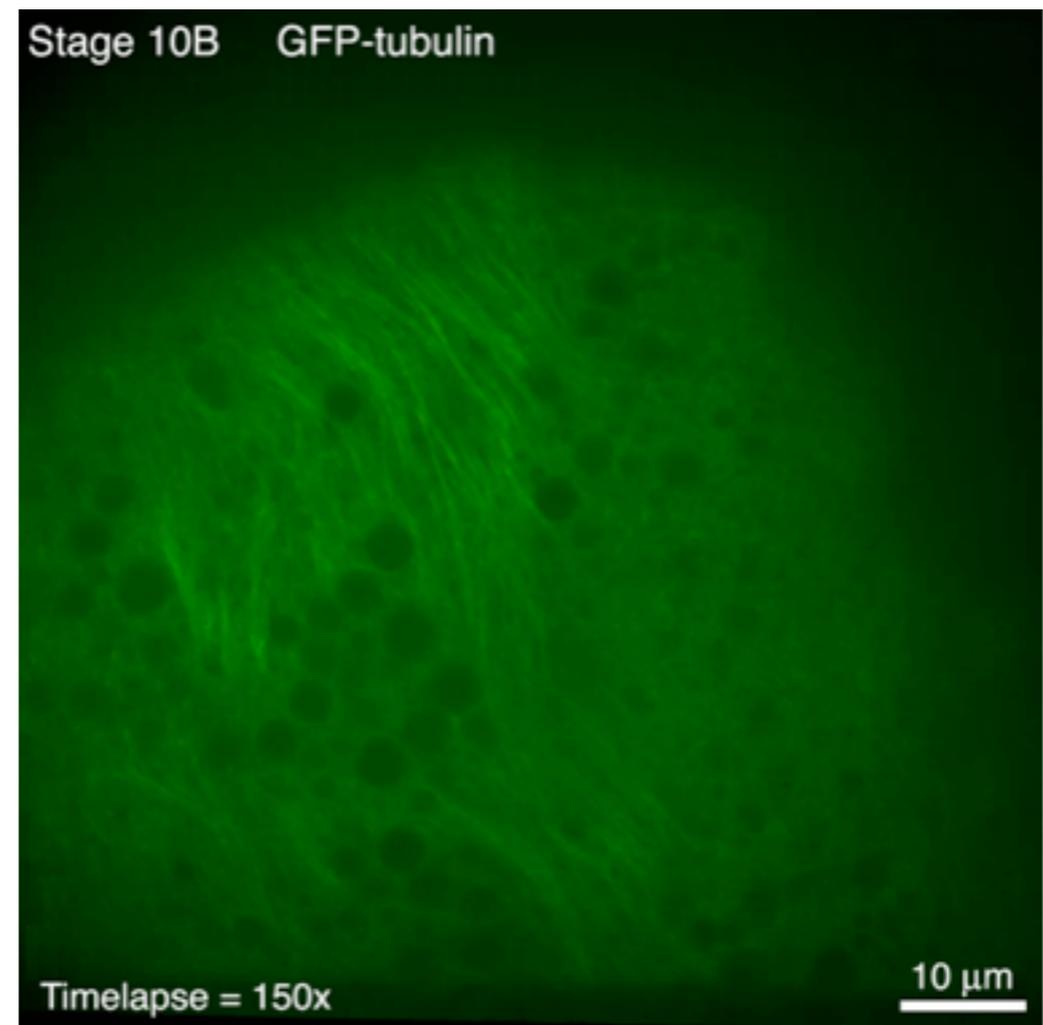
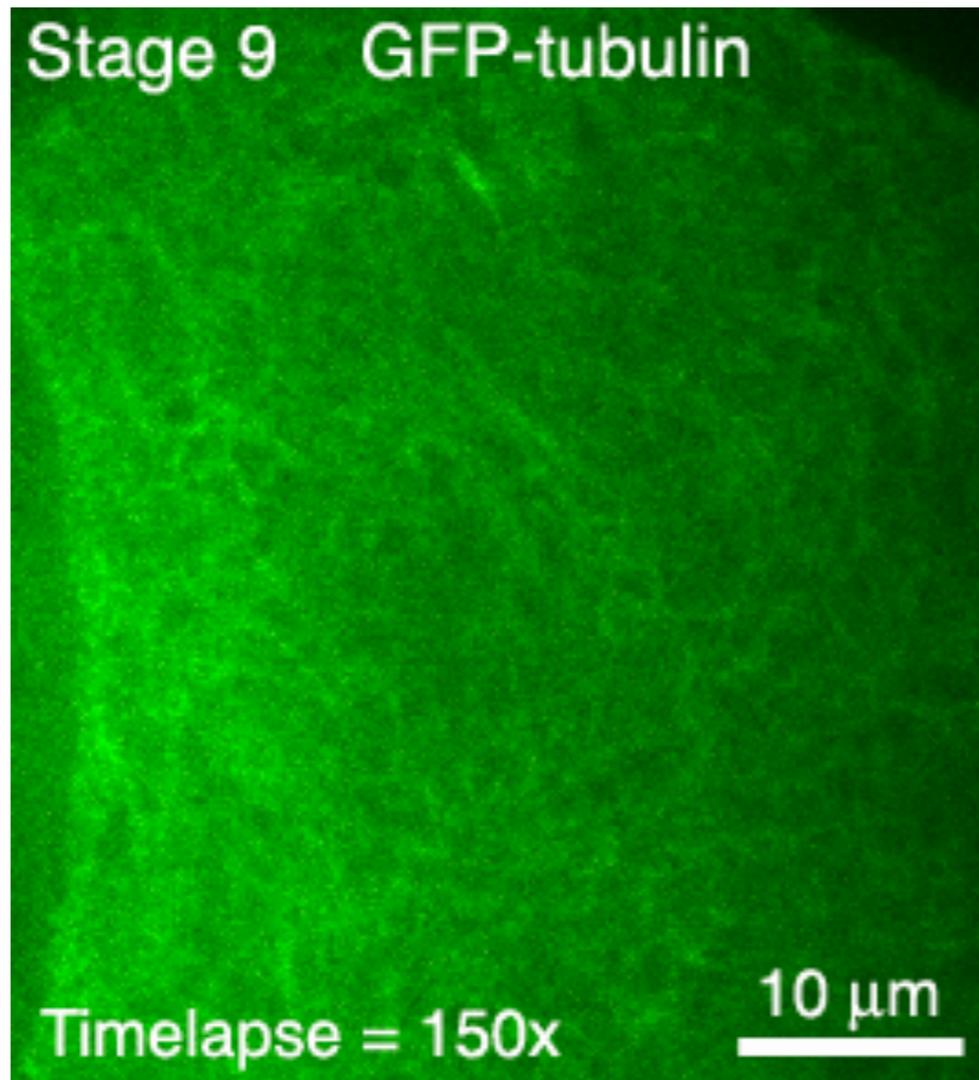
Josh Deutsch, Matt Brunner Corey Monteith, Anthony  
Bielecki, Bill Saxton UCSC Physics and Biology

# Composition of Oocyte

Kinesin walks on microtubules with impellers attached



# Slow vs Fast streaming



# Theoretical Model

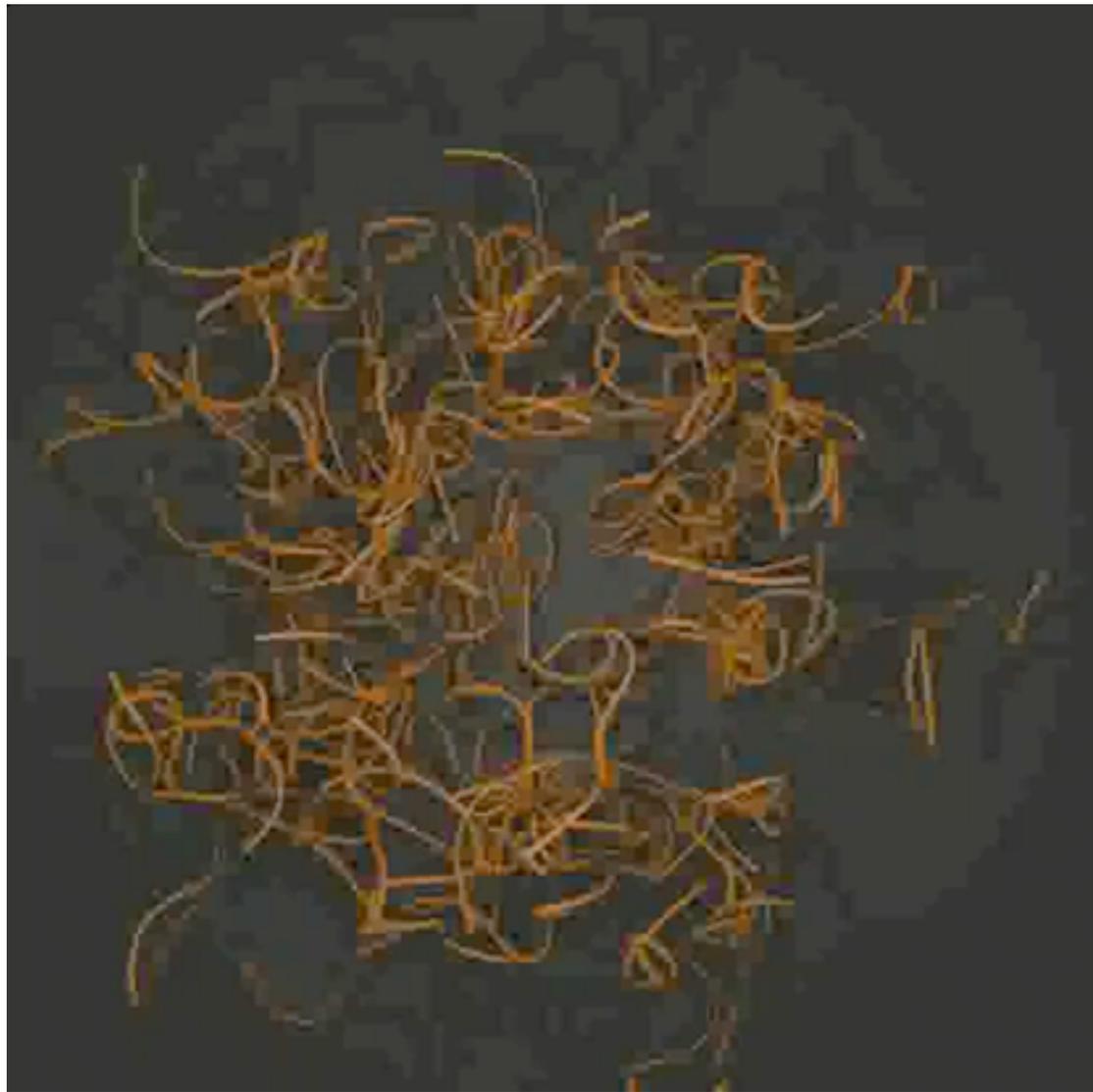
- Model microtubules as elastic rods anchored at minus ends.
- Net effect of kinesin is to apply force tangent to rod.
- Include hydrodynamic coupling.

# Analysis

- Exhibits a non-linear instability giving waves with time and length scales close to those found experimentally.
- Hydrodynamic interactions depend on height above surface, viscosity, and separation of microtubules.

# Simulations with full hydrodynamic interactions.

Shown here for microtubules at two heights above a planar surface ( $h=1$  and  $h = 2$ )



# Photo-mechanical energy conversion using biomimetic materials.

Josh Deutsch  
Dept of Physics  
UCSC

January 24, 2014

Supported by and National Science Foundation CCLI Grant DUE-0942207.

# This Approach

Substitute synthetic engine



## For biological one



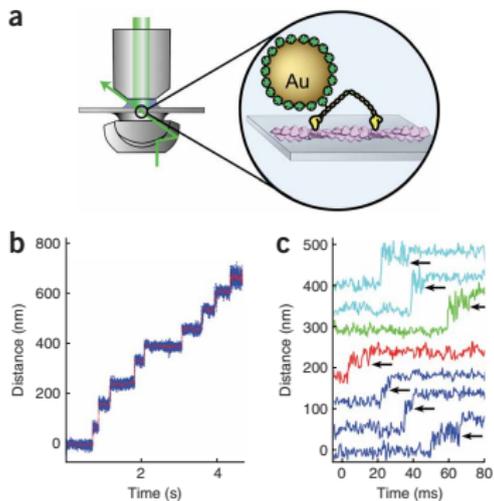
# Engineer muscles using ATP



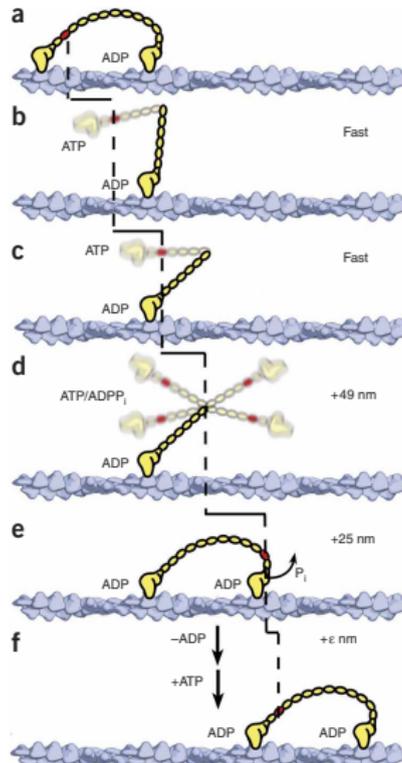
to use photons instead



# How do biological motors work? Myosin V



Dunn and Spudich (2007)



# Model of Myosin V

- Periodic distribution of binding sites.
- Heads bind to these sites at an angle.
- Two rates: Rate end goes from sticky → not sticky and vice versa.
- The chain is semiflexible



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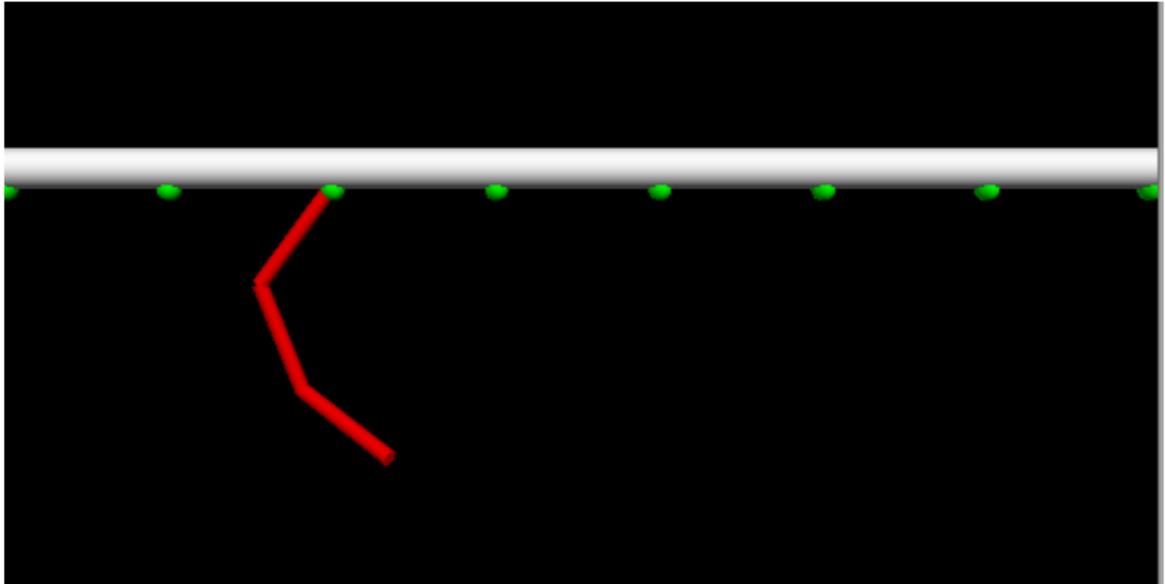


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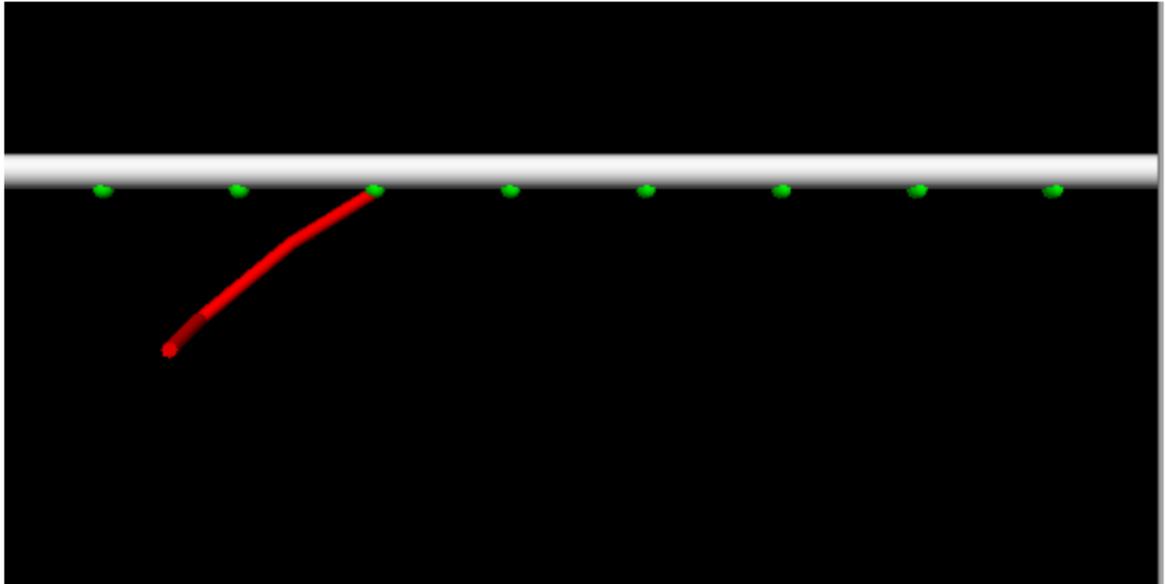
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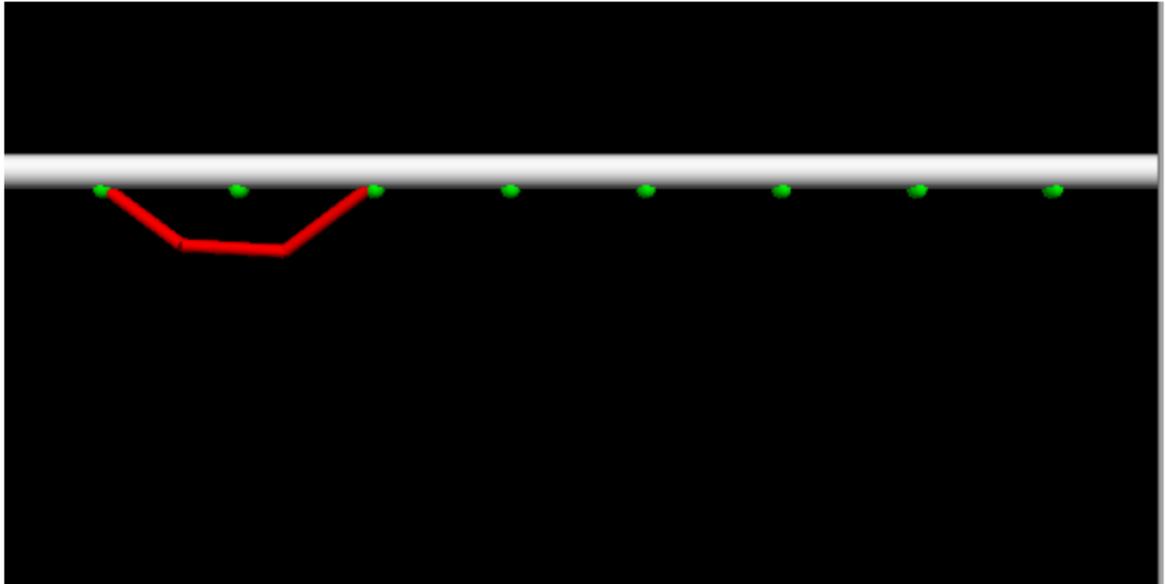
# Model of Myosin V



# Model of Myosin V



# Model of Myosin V



As this simulation shows...

There is little conceptual difference between using light and chemical energy (ATP) to drive this motion.

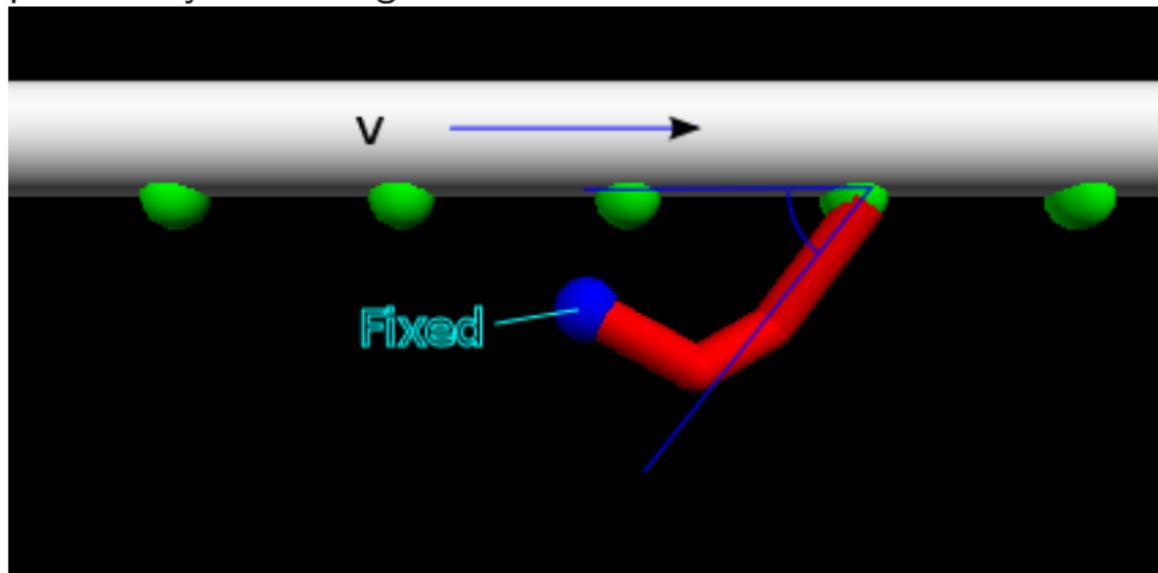
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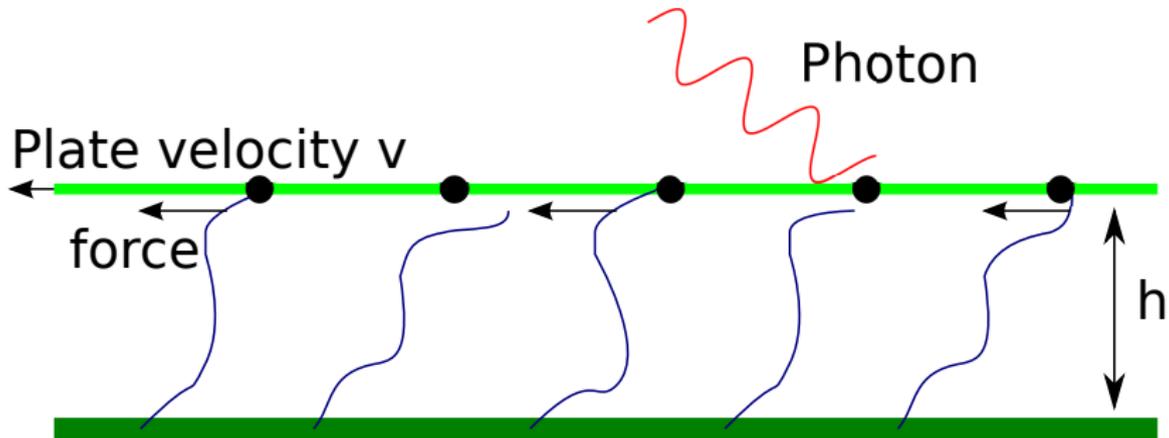
## Simplified model of myosin 2

Now look at a biomotor doing work against a force, either being powered by ATP or light.



As this simulation shows...

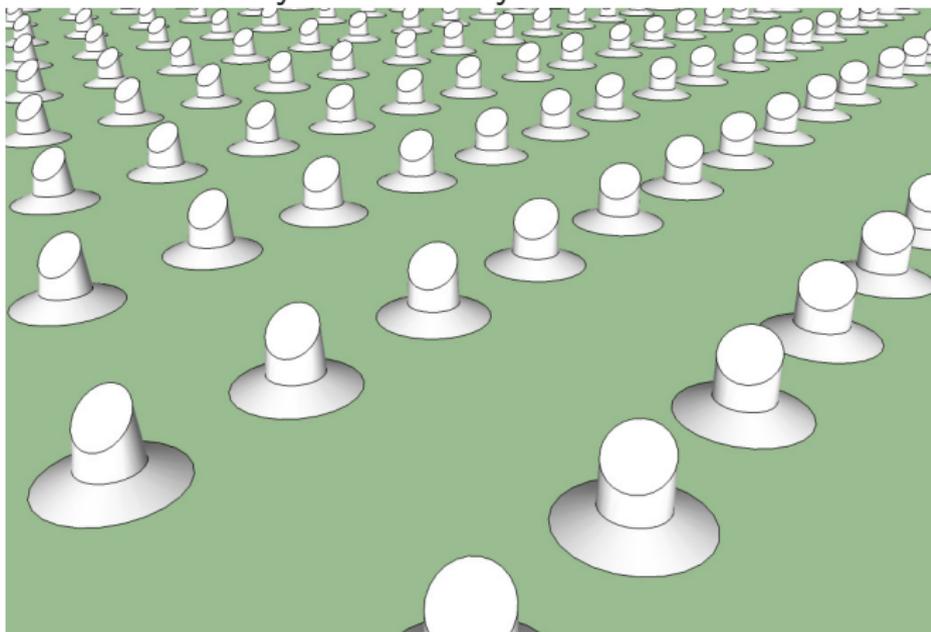
# Prototype photomechanical device



- **A semi-flexible polymer brush attached to a flat plate.**
- A parallel plate right above the brush.
- The parallel plate contains an array of photoreactive binding sites. It is crucial that these polymers bind with the surface in an asymmetric way.
- **At least one of the plates is transparent.** Light causes the unbinding of polymer ends from the photoreactive binding sites.
- **Binding catalyst.** To control the rate at which binding occurs, the binding of the end of the polymer to a binding site can be facilitated by the use of a catalyst.

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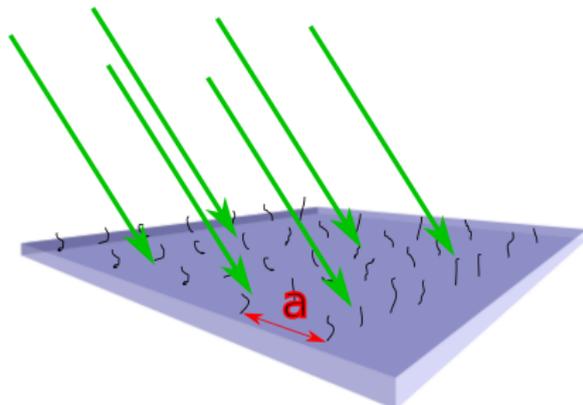
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# Operation in sunlight

Assume

- Solar Flux  $I = 600W/m^2$ ,
- photon energy  $2eV$
- polymer size  $3nm$ ,
- relaxation time  $\approx 10^{-7}s$

$\implies$  photon flux  $\approx 2 \times 10^{21}/(m^2s) \implies$  separation  $a \approx 70nm$ .



## $k_B T \ll$ Photon Energy

Photon energy  $\sim 2eV$ , but  $k_B T \approx 0.025eV$ ,  $80 \times$  smaller. Higher efficiency could be obtained if powerstroke  $\gg k_B T$ .

How can we harness as much of the solar energy as possible?

To harness the energy of a photon, we need to release a lot of potential energy in the motor when a head unbinds. Otherwise most of the photon energy is wasted.

We'll start by modelling the process mathematically.

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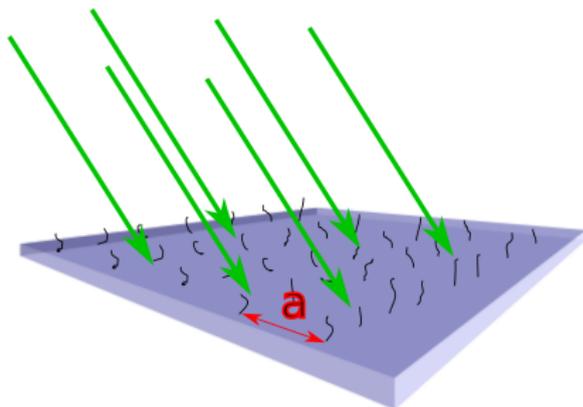
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# How to improve design

If the photon cross section is made arbitrarily low, the efficiency can be adjusted to remain constant.

The energy delivered in one cycle  $\propto kL^2$ , but the larger this is, the lower the cross section.

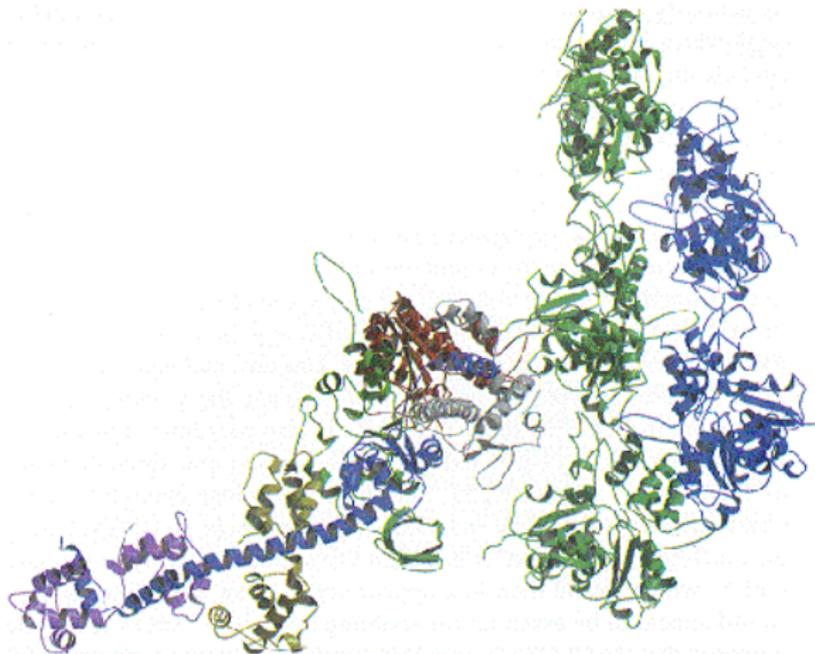
With the parameters we used earlier, the density can be increased by  $\approx 1000 \implies$  cross section of  $1/1000$ , so  
Realistically the work delivered in a power stroke can be  $\sim 7k_B T$ .



# How to build it?

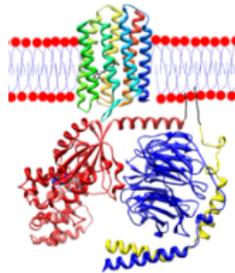


Actin/myosin sounds like a good candidate but it is very complicated and still not well understood.



There are many proteins that respond to light,

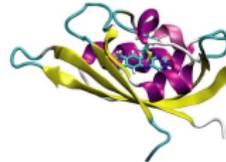
Rhodopsin



DRONPA



LOV domains

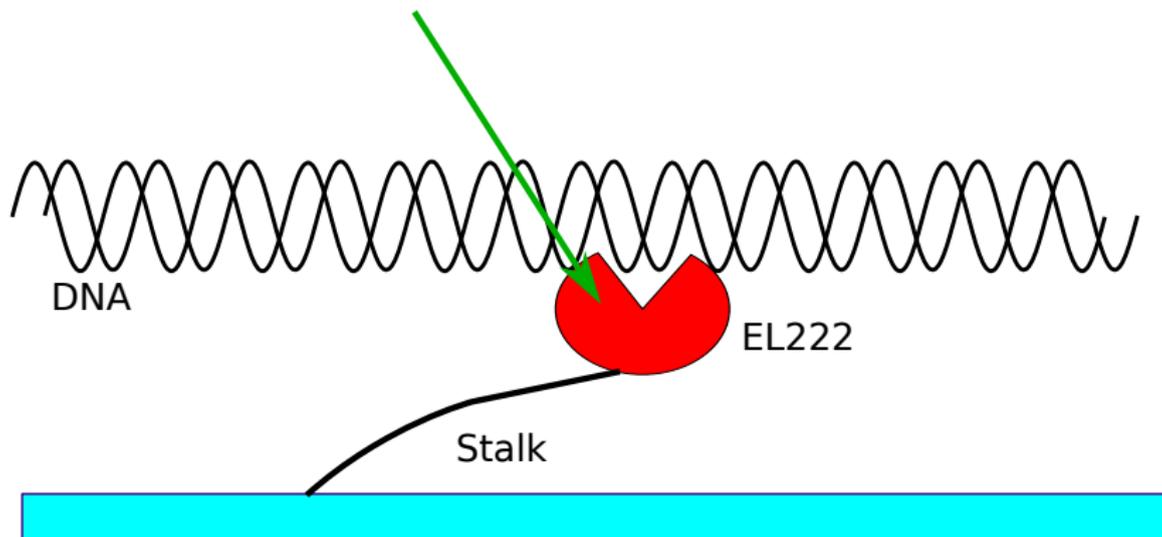


# Structural basis of photosensitivity in a bacterial light-oxygen-voltage/helix-turn-helix (LOV-HTH) DNA-binding protein

Abigail I. Nash<sup>a,1</sup>, Reginald McNulty<sup>b,1</sup>, Mary Elizabeth Shillito<sup>b</sup>, Trevor E. Swartz<sup>c,2</sup>, Roberto A. Bogomolni<sup>c</sup>, Hartmut Luecke<sup>b,3</sup>, and Kevin H. Gardner<sup>a,3</sup>

EL222 is a light regulated DNA binding protein:

- Light state: binds to specific DNA sequence
- Dark state: inhibits binding



## Conclusion: are bio-solar-motors possible?

- There is no theoretical reason prohibiting direct conversion of light to mechanical energy.
- There are many biological systems converting chemical energy (ATP) to mechanical energy.
- We need to substitute ATP for photons.
- The efficiency can be optimized by lowering the cross section per motor and raising their density.
- There are a number of biological systems that already use light to bind or unbind molecules.
- With movement in the experimental direction, the future for solarmotors will be bright!

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