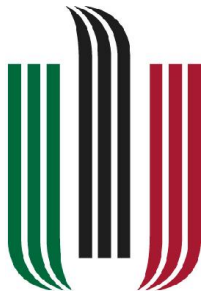


Nanoscale communications



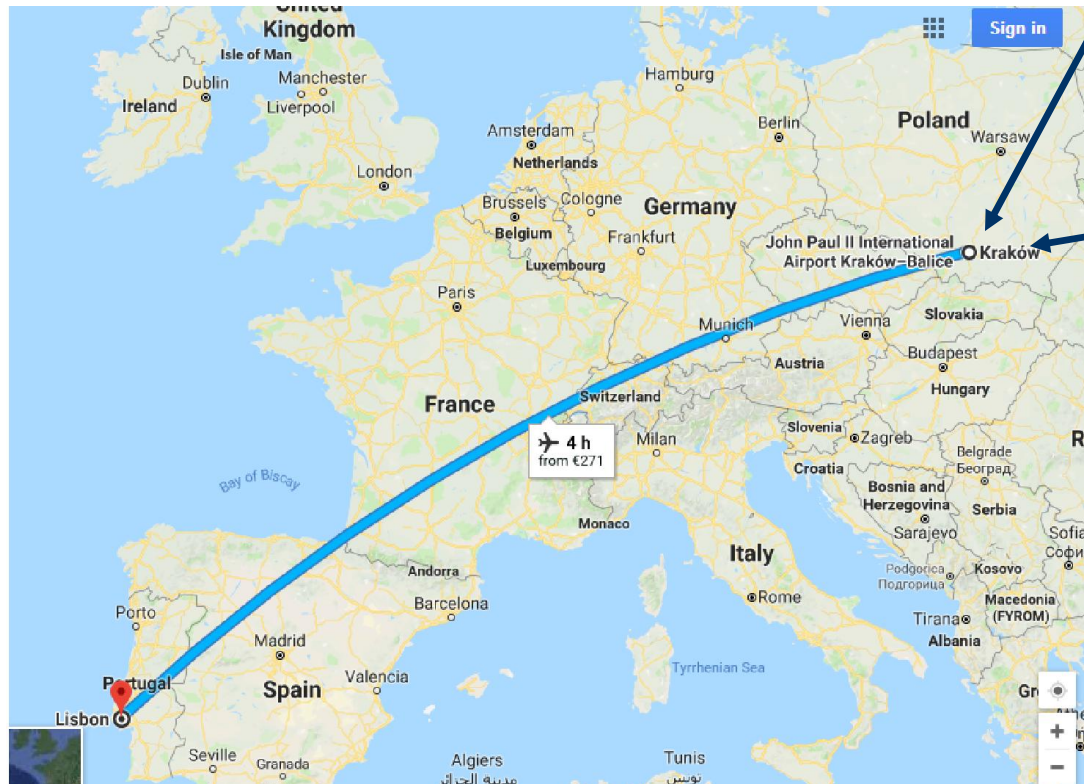
Pawel Kulakowski, AGH Univ. of Sc. And Tech., Krakow, Poland

kulakowski@kt.agh.edu.pl

Where I work



AGH Univ. Of Sc. And Tech.



Jagiellonian Univ.

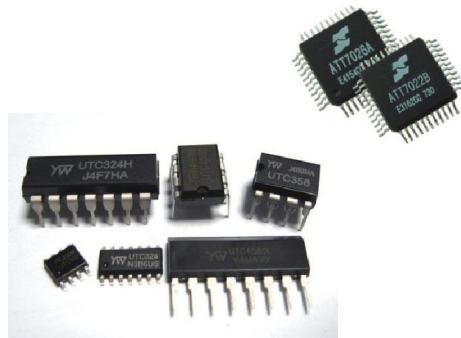


Talk outline

1. Nanocommunications:
motivation and overview of possible approaches
2. Theory of FRET phenomenon
3. Experiments
4. Going from basic science to applications:
simulation results
5. Open issues and conclusions

Nanocommunication origins

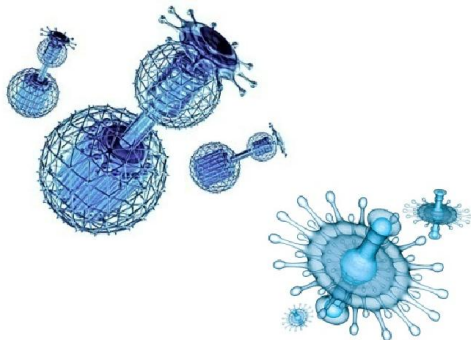
ELECTRONICS



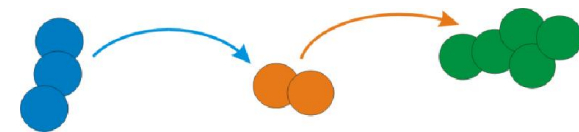
COMMUNICATIONS



NANOTECHNOLOGY



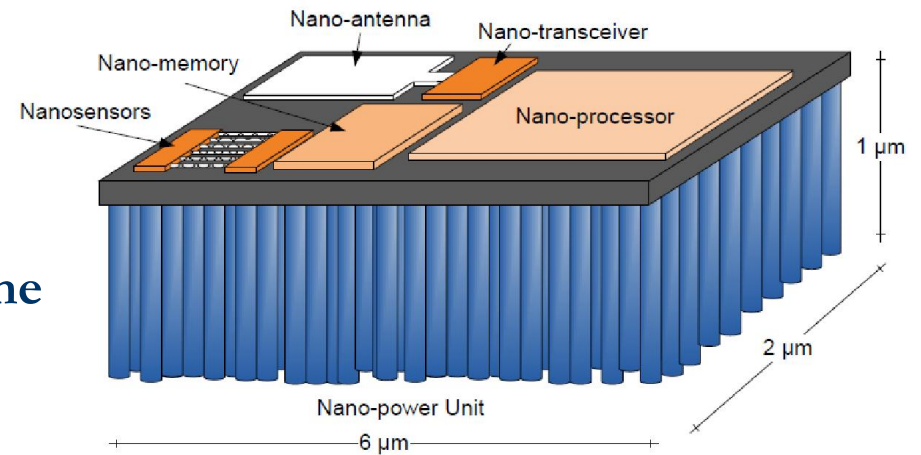
NANO-COMMUNICATIONS



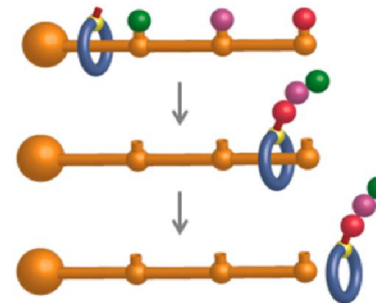
Possible approaches

1. Miniaturization (*top-down*)

- new materials:
carbon nanotubes and graphene
- 0.1-10 THz band



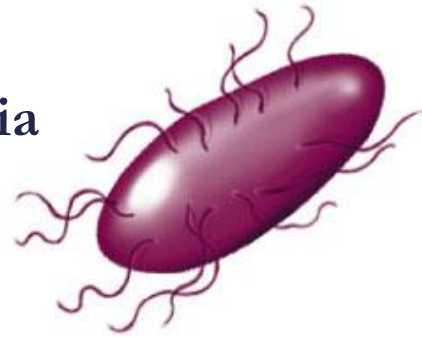
2. Building from single molecules (*bottom-up*)



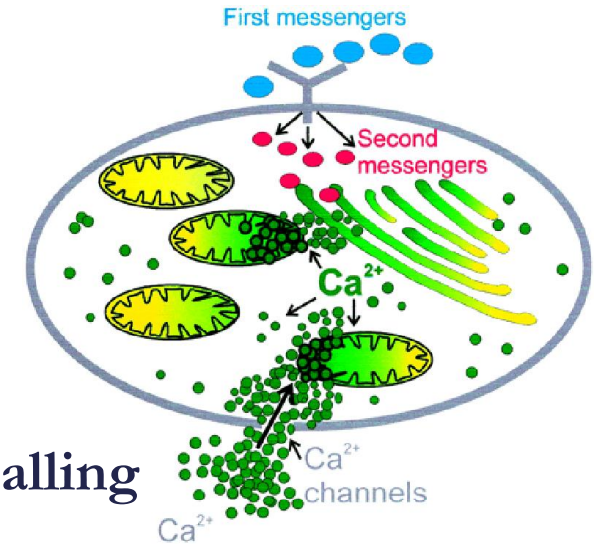
3. Molecular communication (arising from *bio-hybrid approach*)

Molecular mechanisms considered

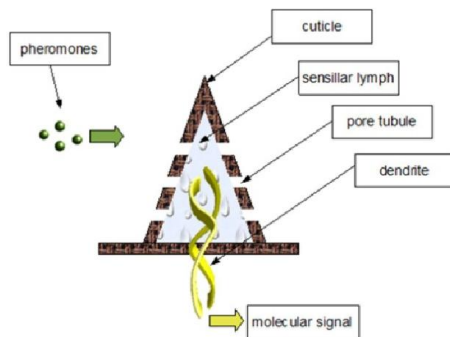
flagellated bacteria



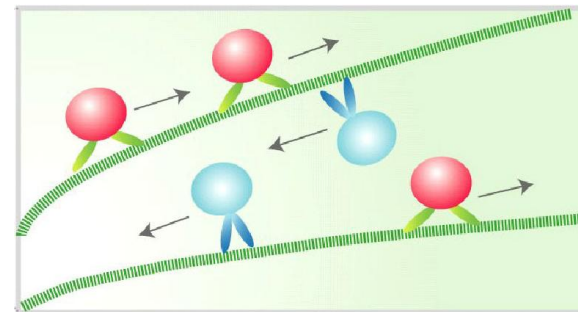
calcium ions signalling



pheromones propagation

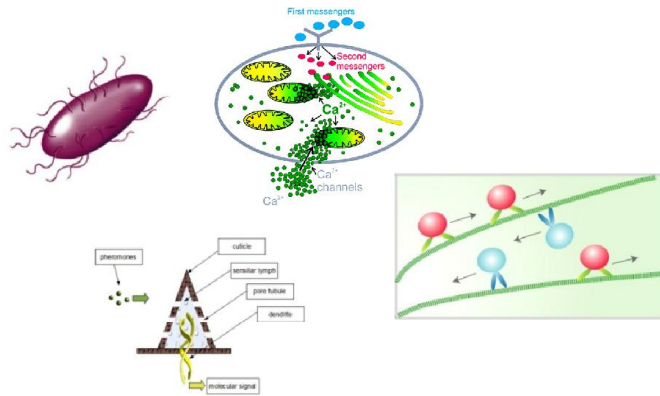


molecular motors



Signal propagation speed in molecular comm.

Molecular mechanisms



$< 50 \mu\text{m/s}$ ($5 \times 10^{-5} \text{ m/s}$)

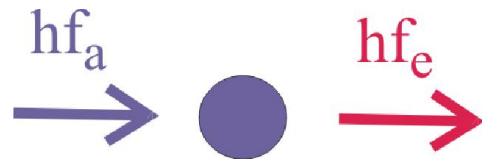
Förster Resonance Energy Transfer (FRET)



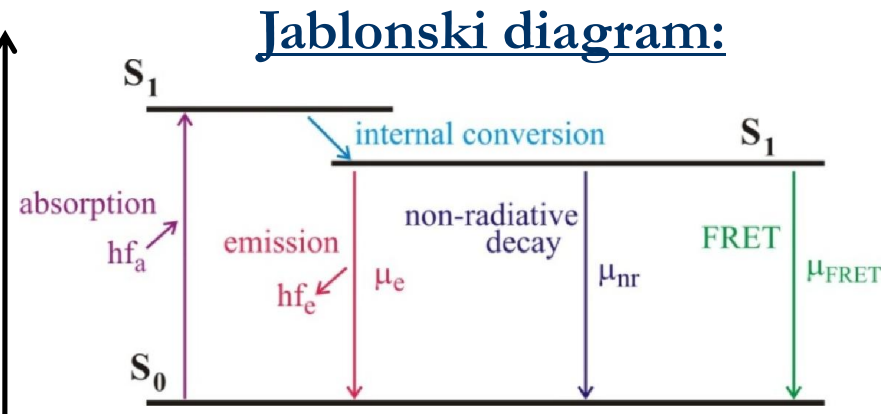
Theory

Förster Resonance Energy Transfer (FRET)

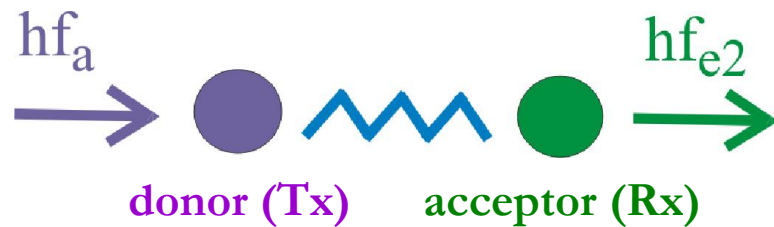
EXCITATION, EMISSION:



Energy ↑



EXCITATION, **FRET**, EMISSION:



FRET efficiency:

$$E = \frac{R_0^6}{r^6 + R_0^6}$$

Energy:

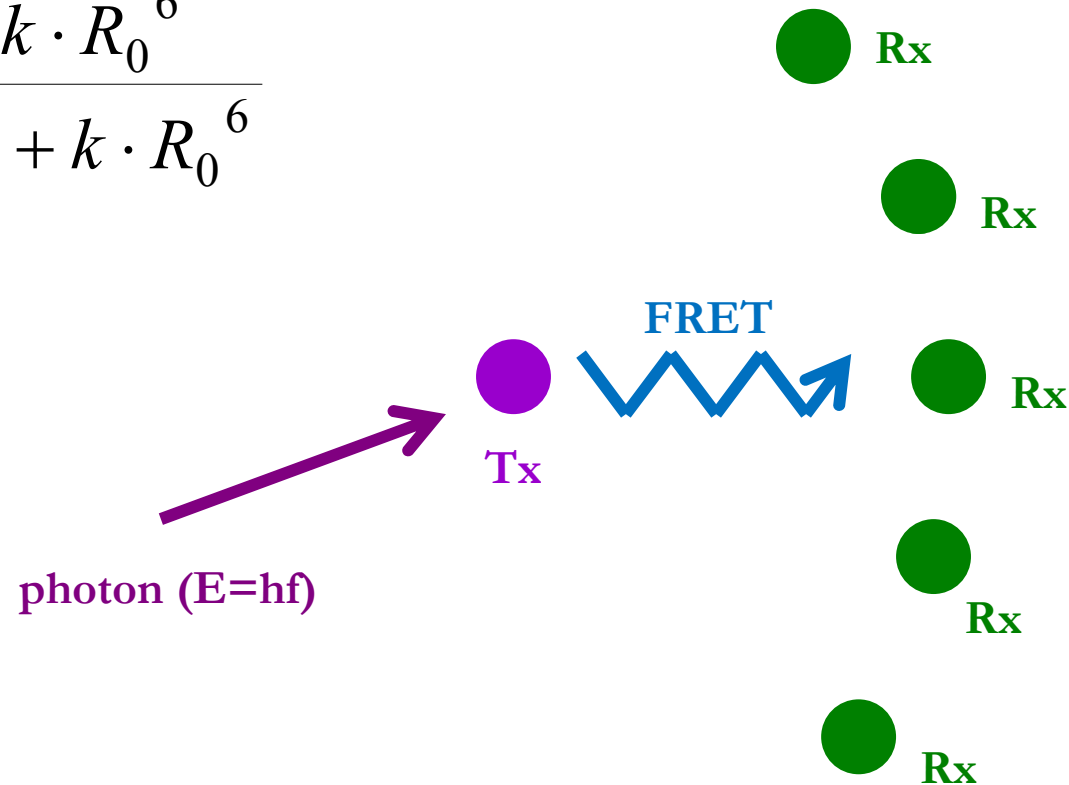
$$hf_a > hf_e > hf_{e2}$$

R_0 (Förster radius) = 3÷9 nm

FRET efficiency

with k equally distant acceptors:

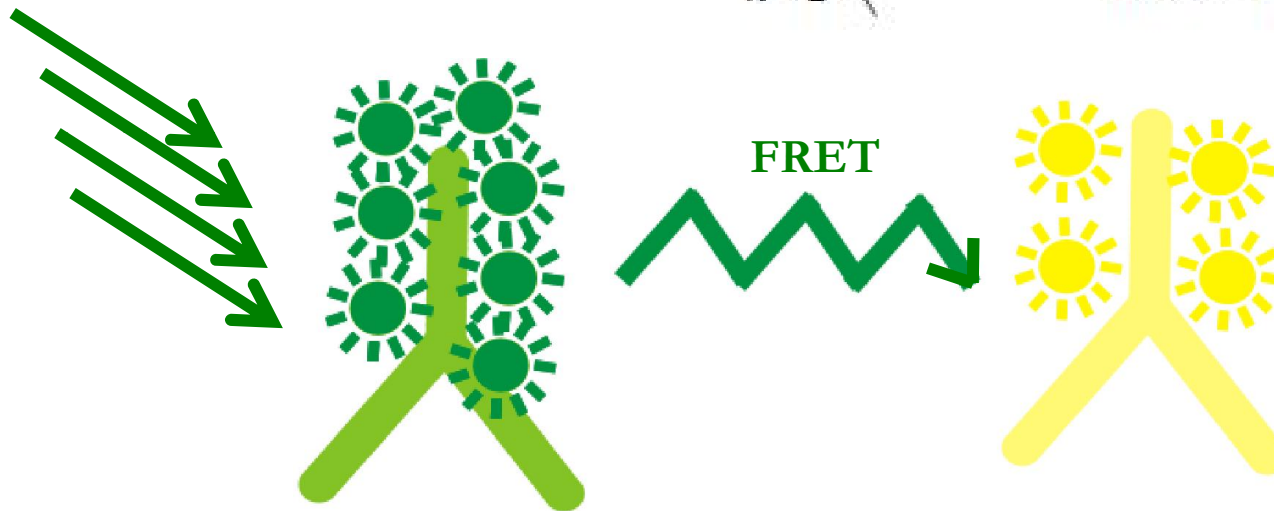
$$E = \frac{k \cdot R_0^6}{r^6 + k \cdot R_0^6}$$



FRET efficiency

with n donors and m acceptors (MIMO):

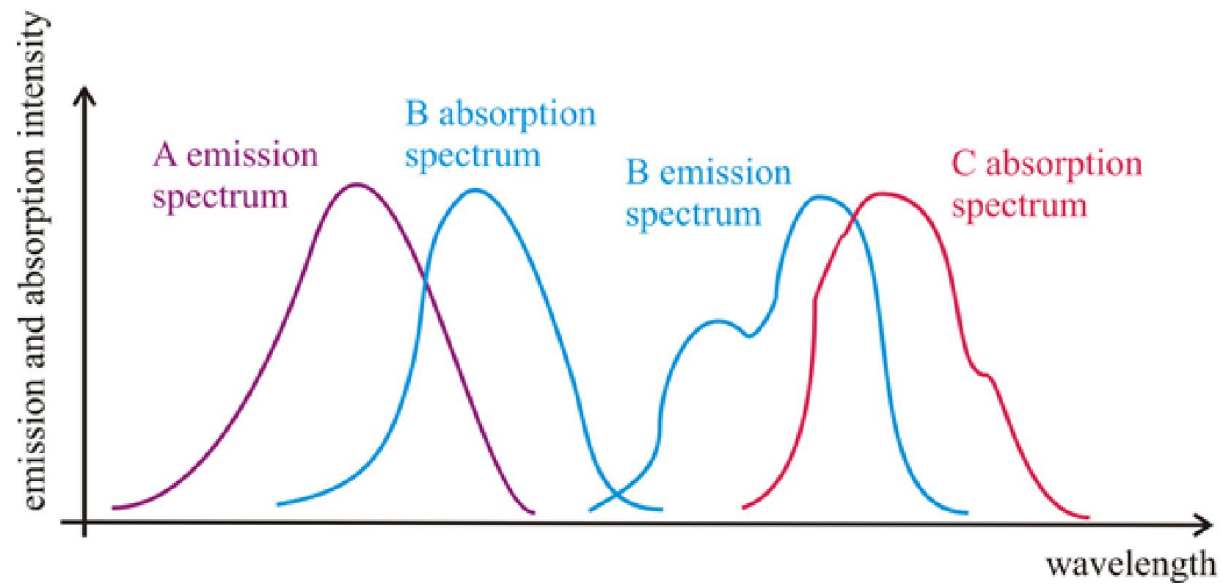
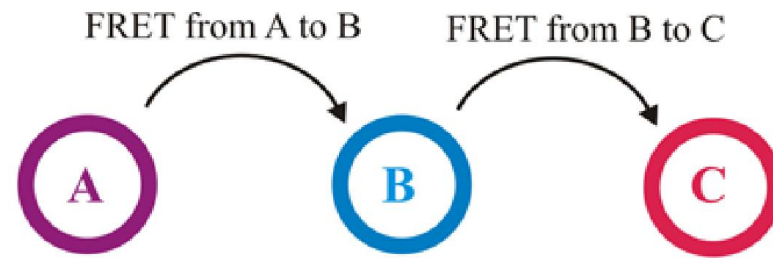
$$E_{n,m} = 1 - \prod_{k=1}^n \left(1 - \frac{R_0^6 \sum_{i=1}^m \frac{1}{r_{ki}^6}}{1 + R_0^6 \sum_{i=1}^m \frac{1}{r_{ki}^6}} \right)$$



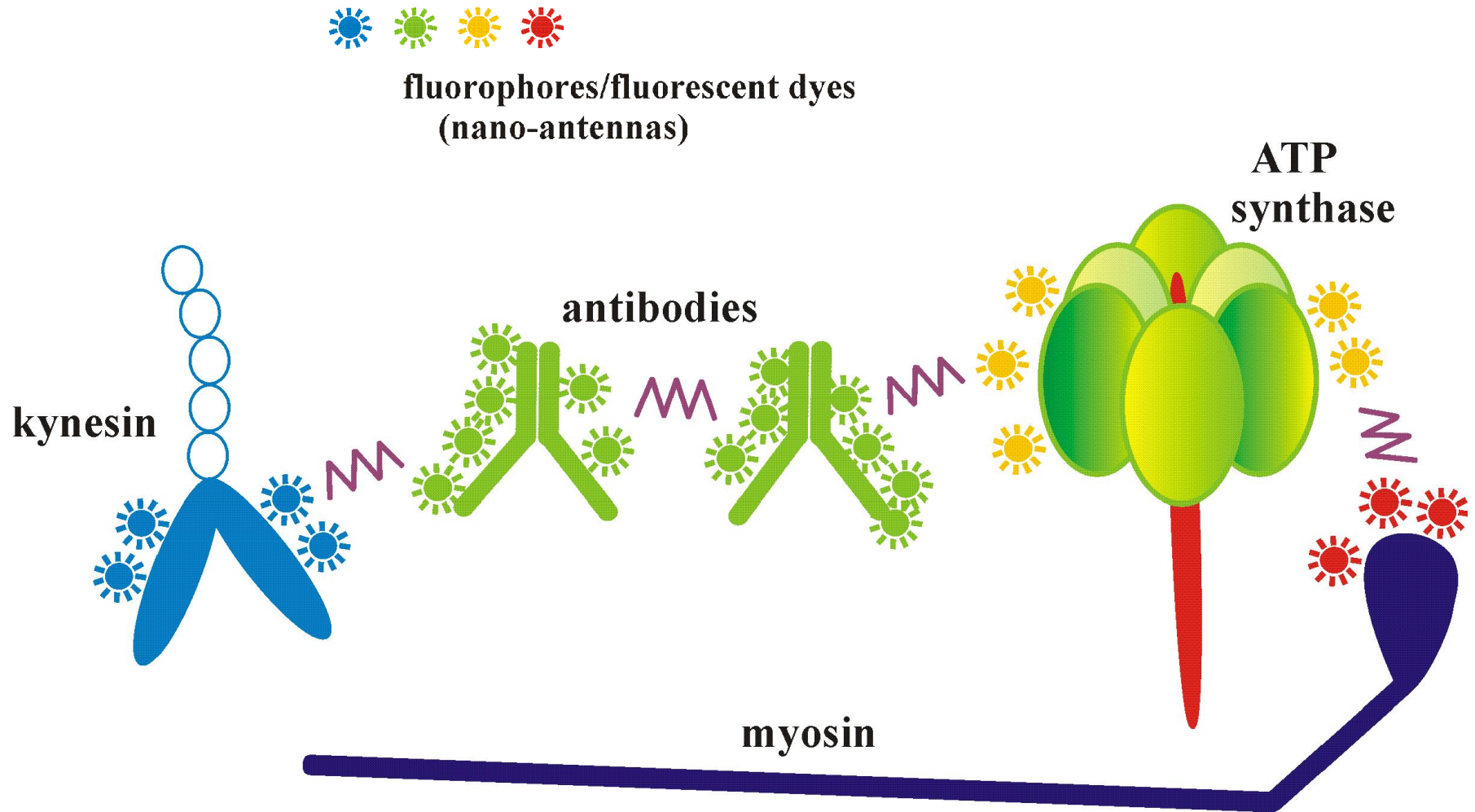
ON-OFF modulation scheme:

- bit '0' = no signal
- bit '1' = excitation of all Tx's, FRET

Multi-hop FRET



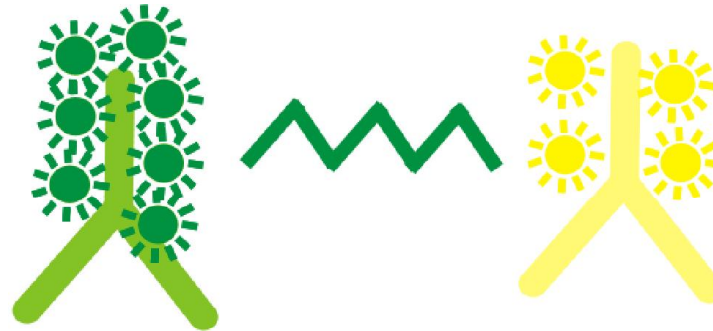
Communicating nanomachines



Transmission ...

ON-OFF modulation scheme

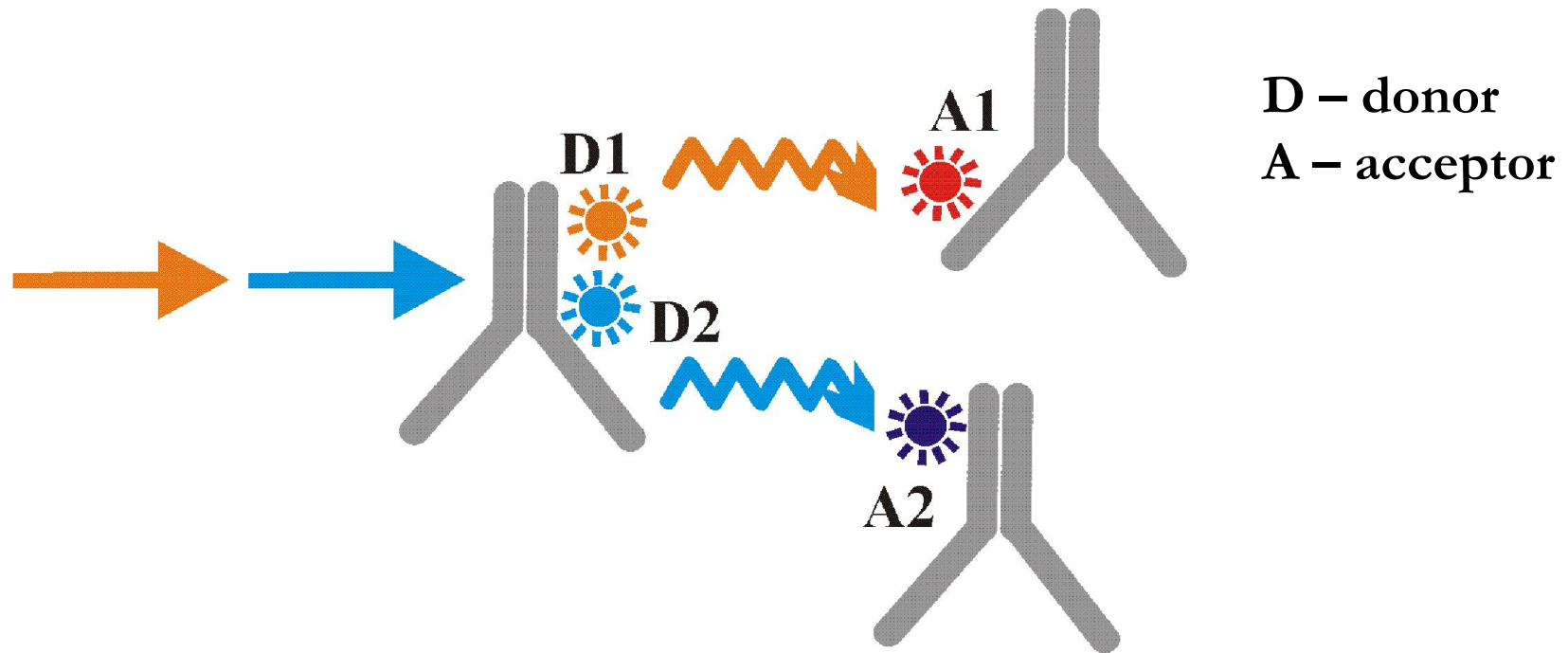
MIMO-FRET



... and routing?

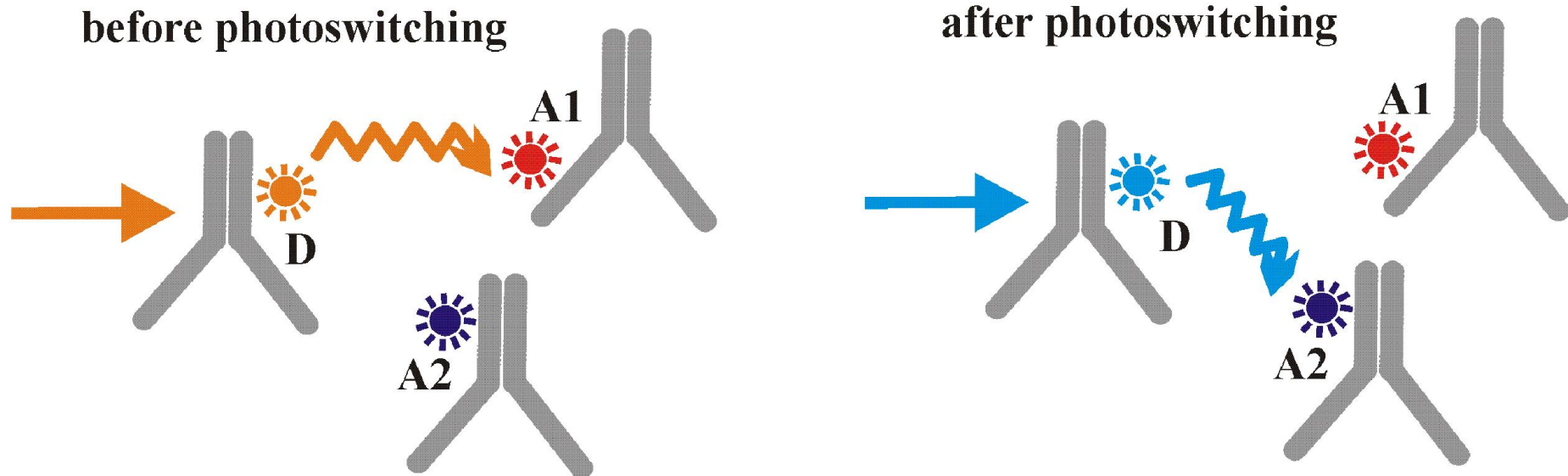
Routing options in FRET-based nano-networks

1. Proteins with a few different fluorescent dyes:



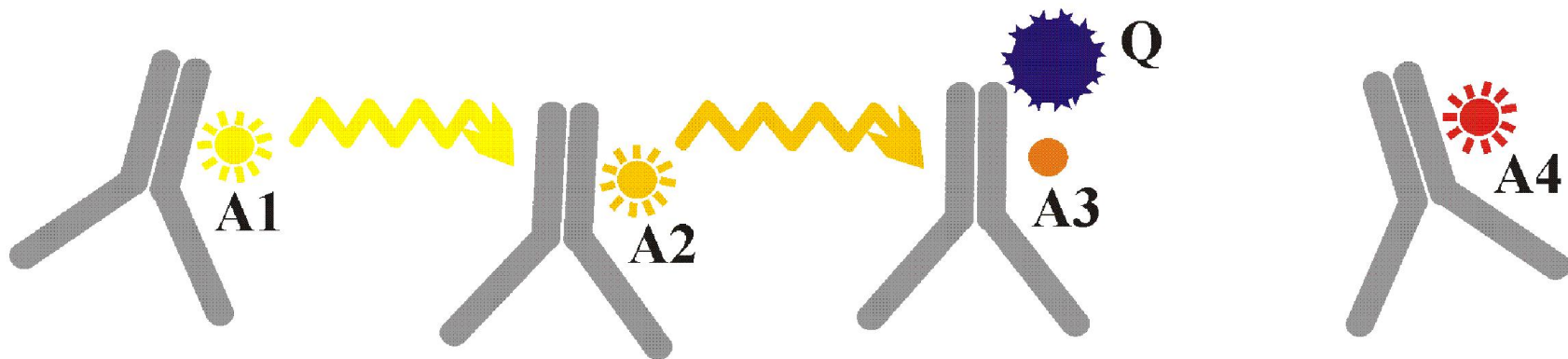
Routing options in FRET-based nano-networks

2. Photoswitchable fluorophores:



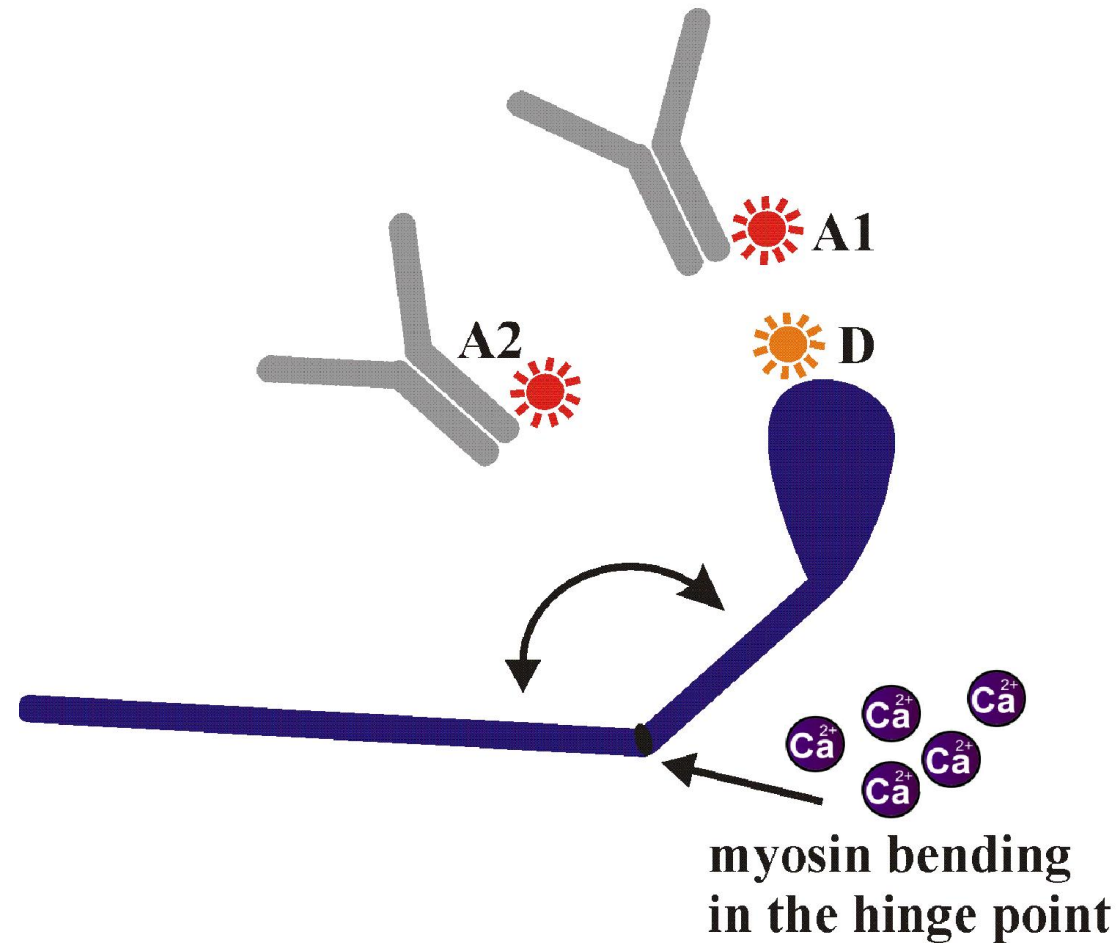
Routing options in FRET-based nano-networks

3. Quenchers:



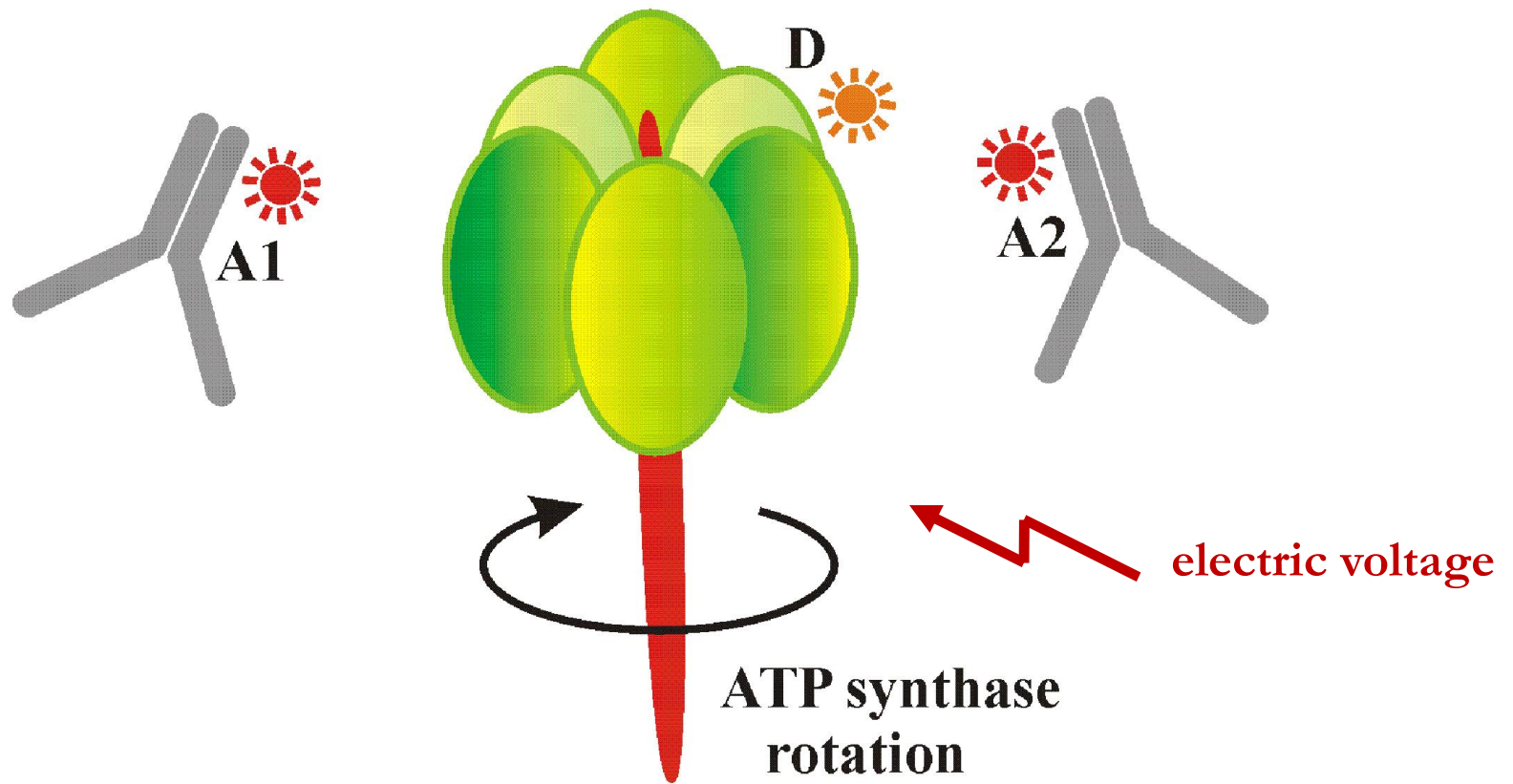
Routing options in FRET-based nano-networks

4. Proteins with changeable shape:



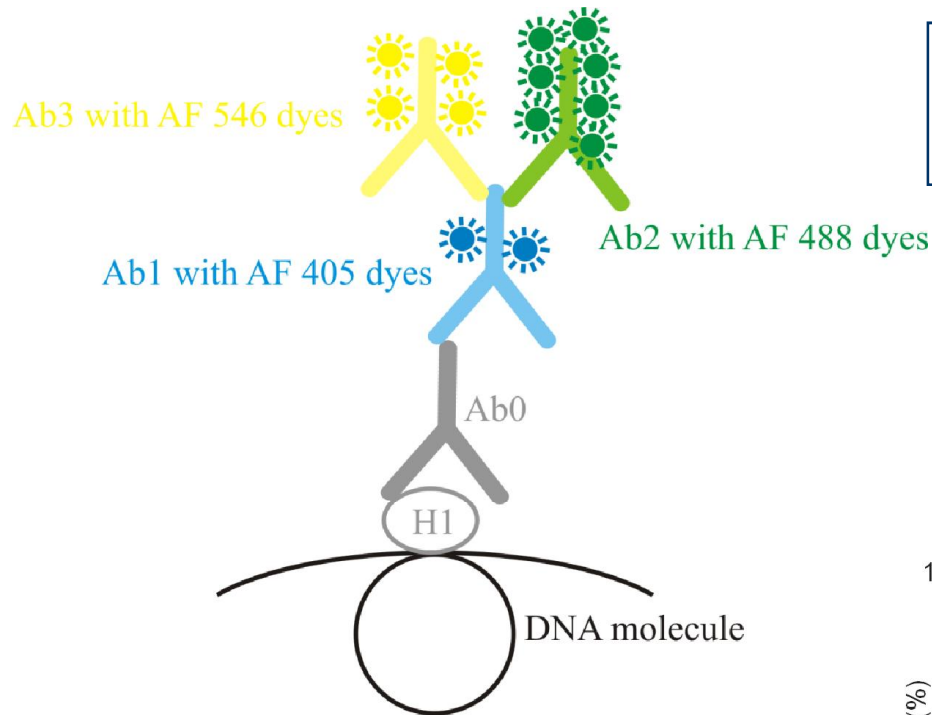
Routing options in FRET-based nano-networks

5. ATP synthase:



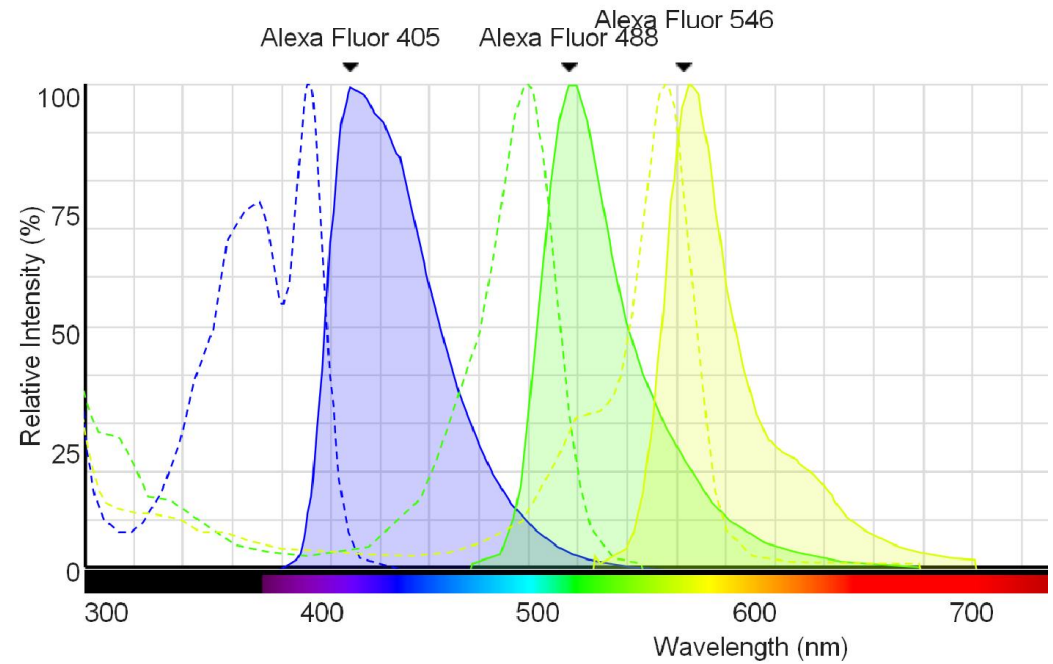
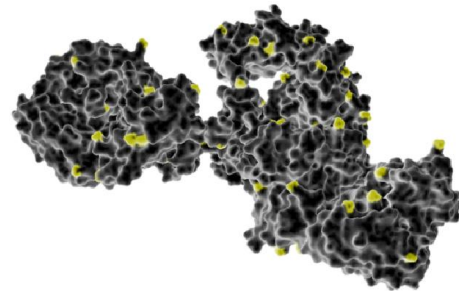
Experiments

A network of antibodies (Ab) and Alexa Fluor (AF) dyes



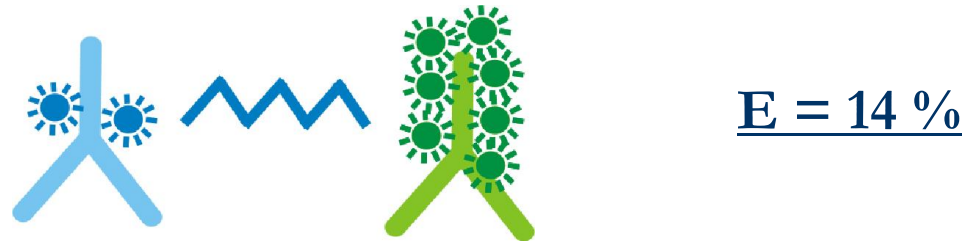
antibody (protein) = nano sensor
AF dye = nano transceiver/antenna

antibodies in the experiment:
immunoglobulin G

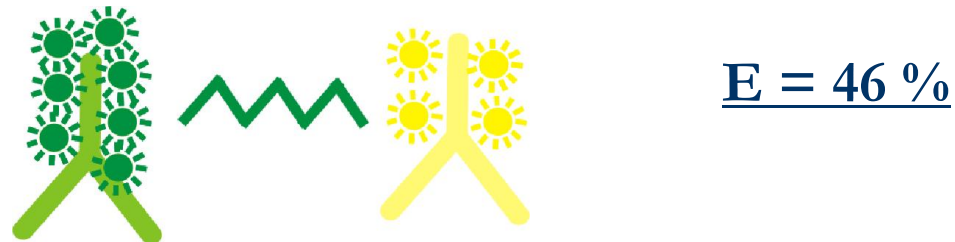


MIMO-FRET measurements

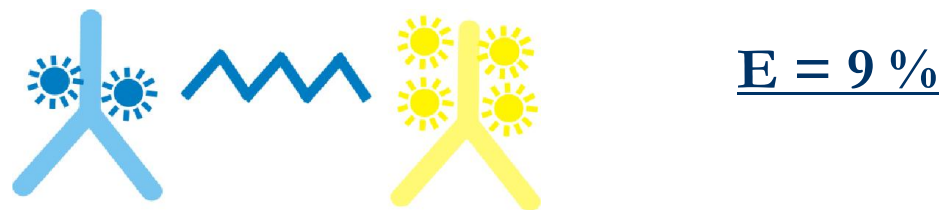
AF405→AF488 - MIMO (1,7):



AF488→AF546 - MIMO (1,4):

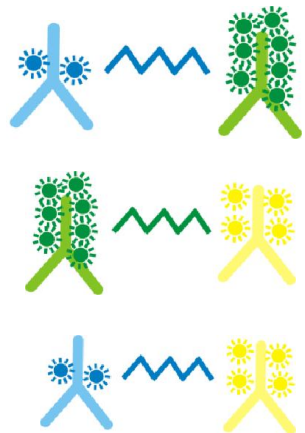


AF405→AF546 - MIMO (1,4):



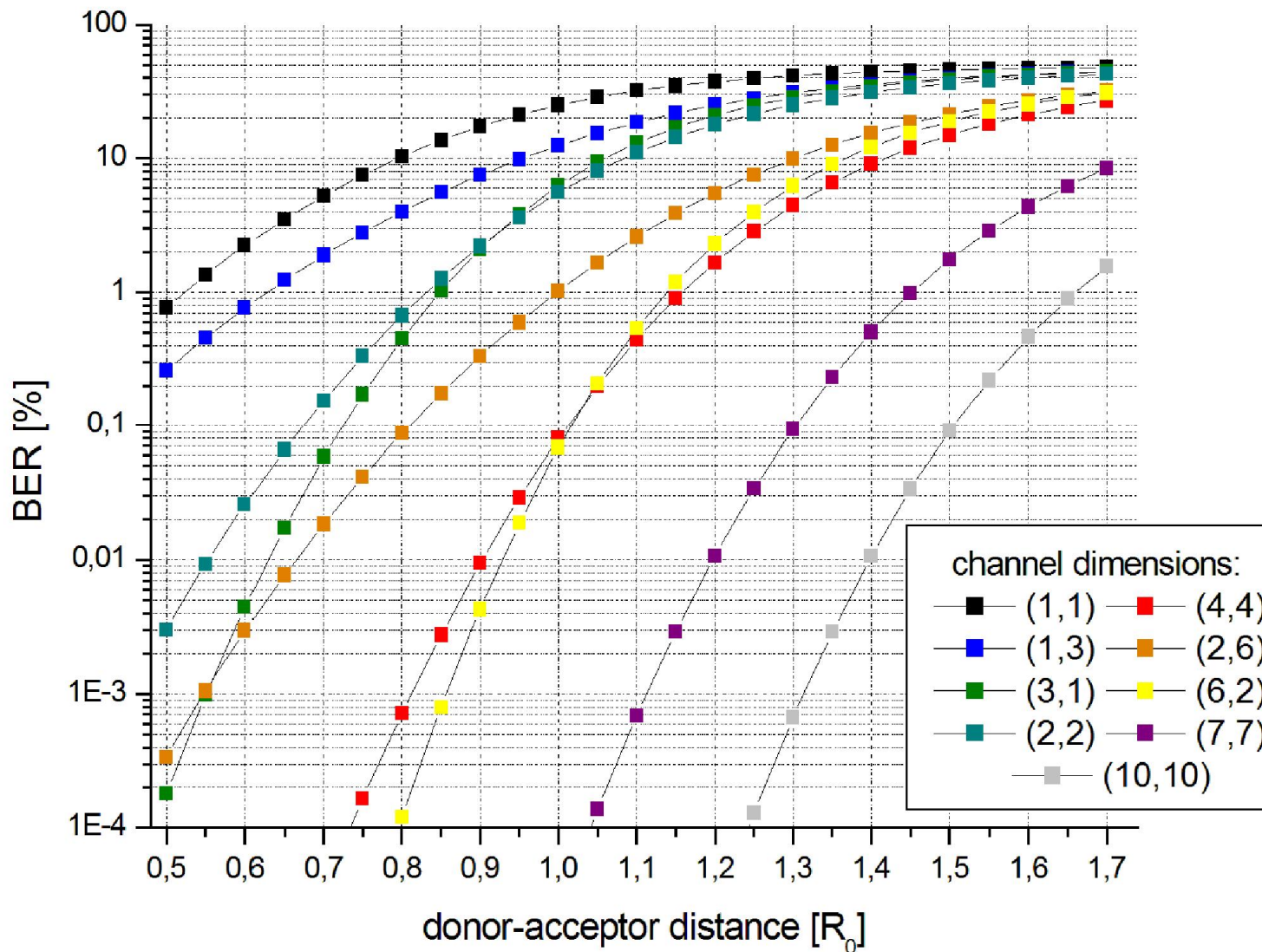
BER calculations for MIMO (1,m) and (n,m)

$$\text{BER}_{n,m} = 0.5 \cdot (1 - E)^n$$

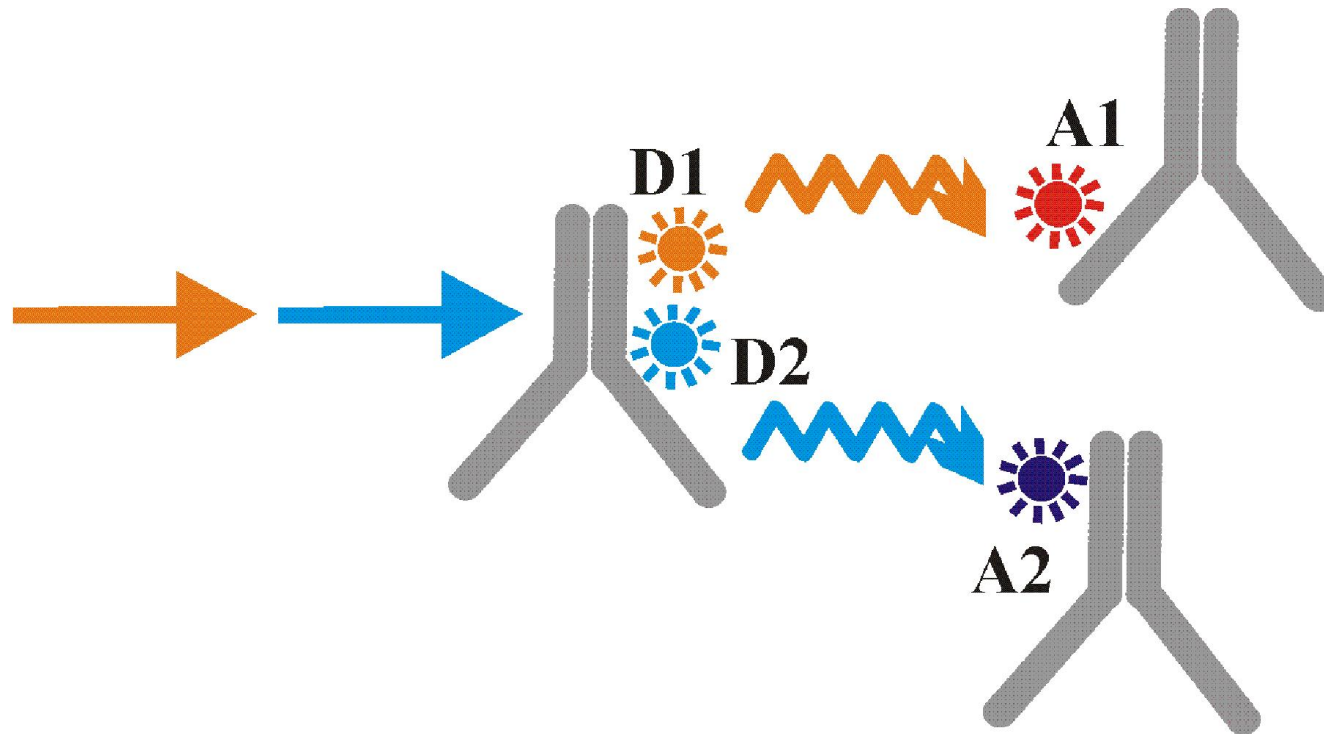


	measured FRET efficiency	BER for (1,m) channel	BER for (n,m) channel
AF405→AF488	14 %	43 %	37 %
AF488→AF546	46 %	27 %	0.7 %
AF405→AF546	9 %	45.5 %	41 %

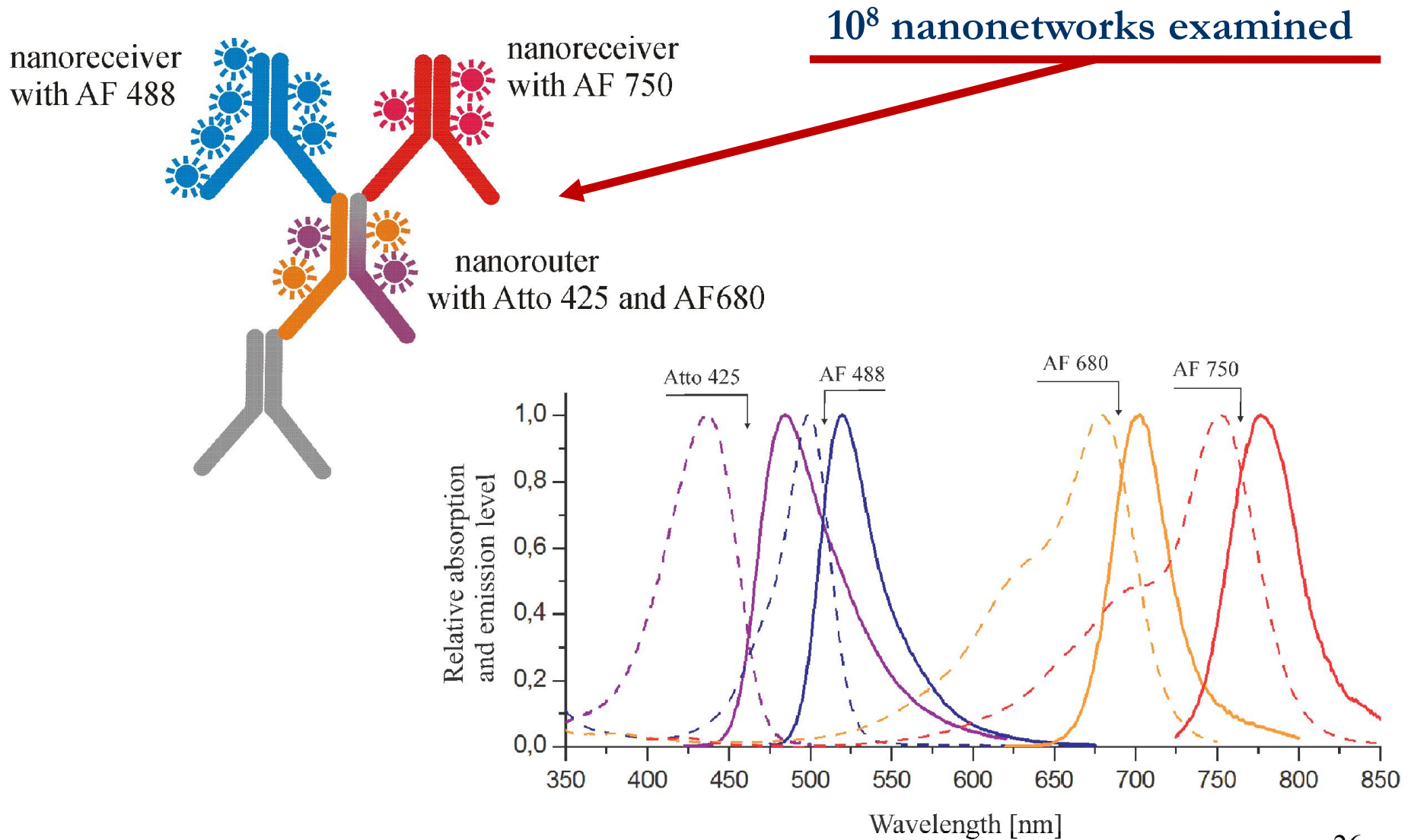
Theoretical BER curves for MIMO (n,m) channels



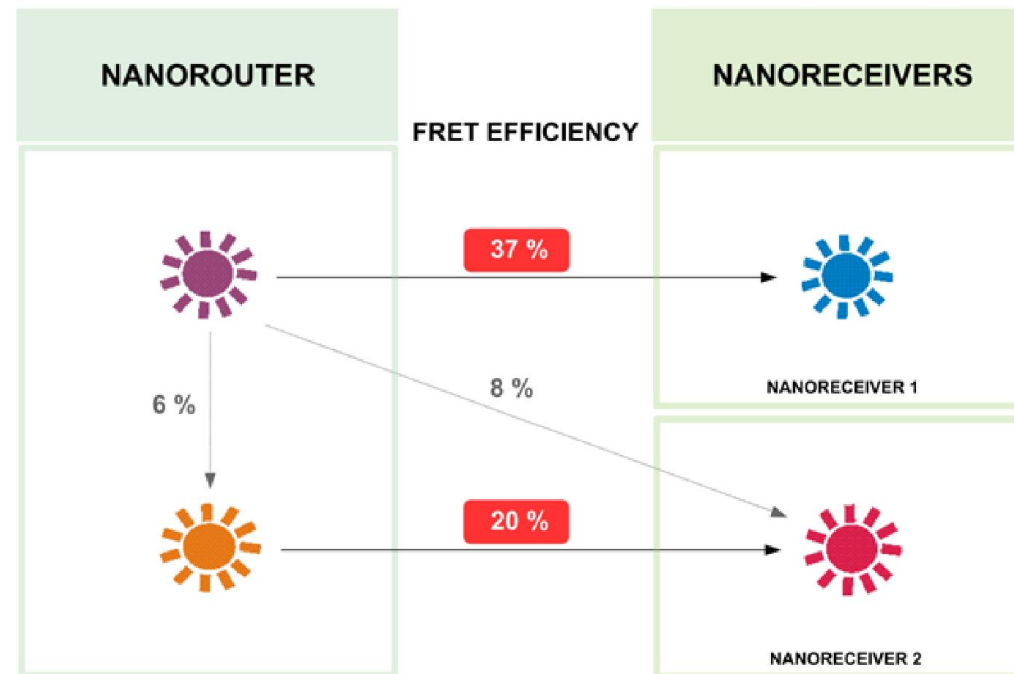
Routing measurements



Measurements with Leica confocal microscope



Measurements with Leica confocal microscope



- ATTO 425
- Alexa Fluor 488
- Alexa Fluor 680
- Alexa Fluor 750

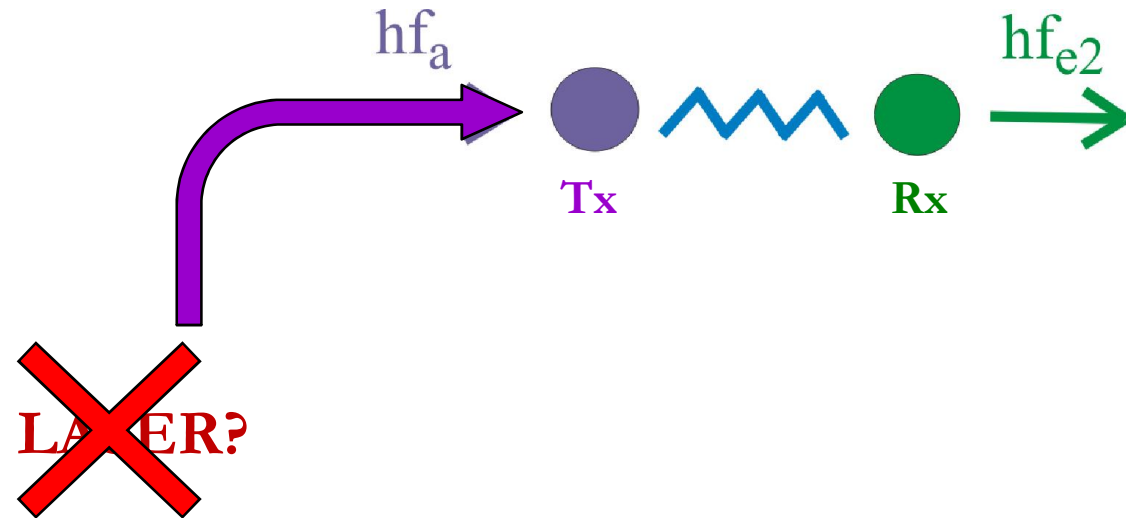
- desired energy transfer
- undesired energy transfer (signal leakage)

Simulations

Going from basic science to applications

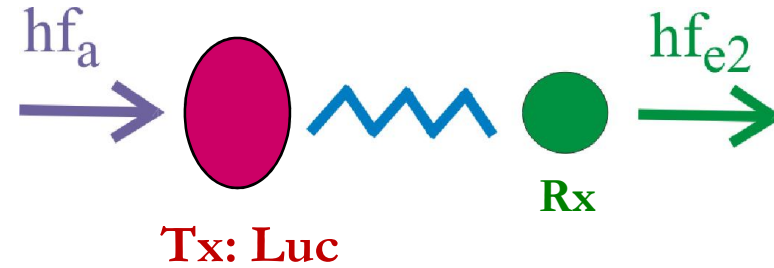
Question 1: How to provide INPUT signals?

EXCITATION, FRET, EMISSION:



Question 1: How to provide INPUT signals?

CHEMICAL
EXCITATION, FRET, EMISSION:



Instead:

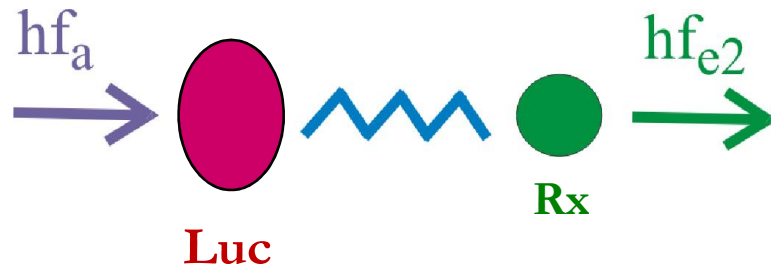
LUCIFERASE

*(a molecule that can be excited
via a chemical reaction)*



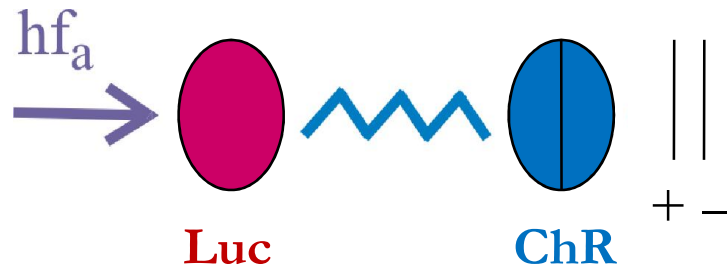
Question 2: How to collect OUTPUT signals?

CHEMICAL
EXCITATION, FRET, EMISSION:

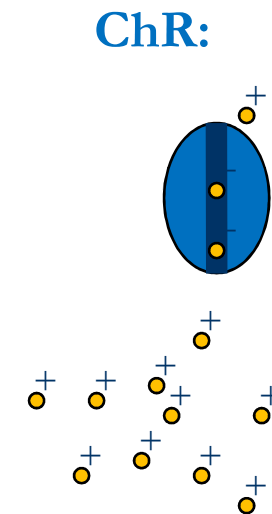
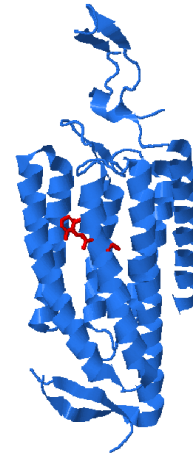


Question 2: How to collect OUTPUT signals?

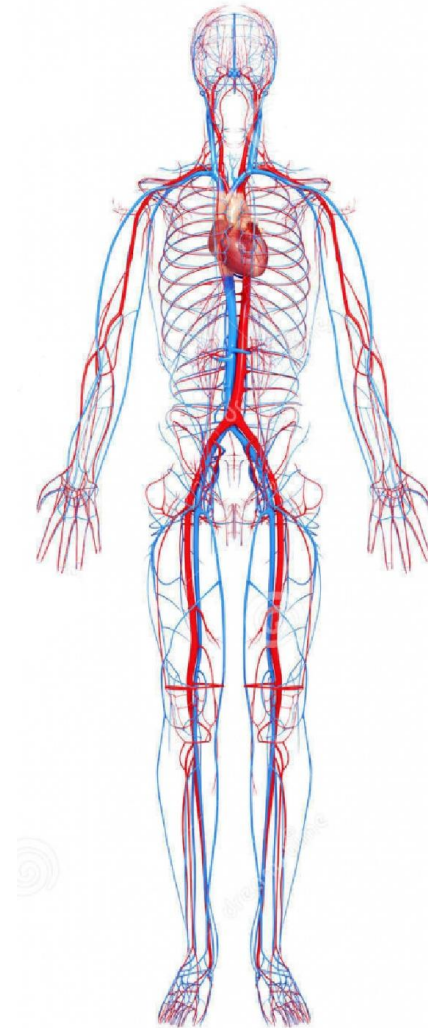
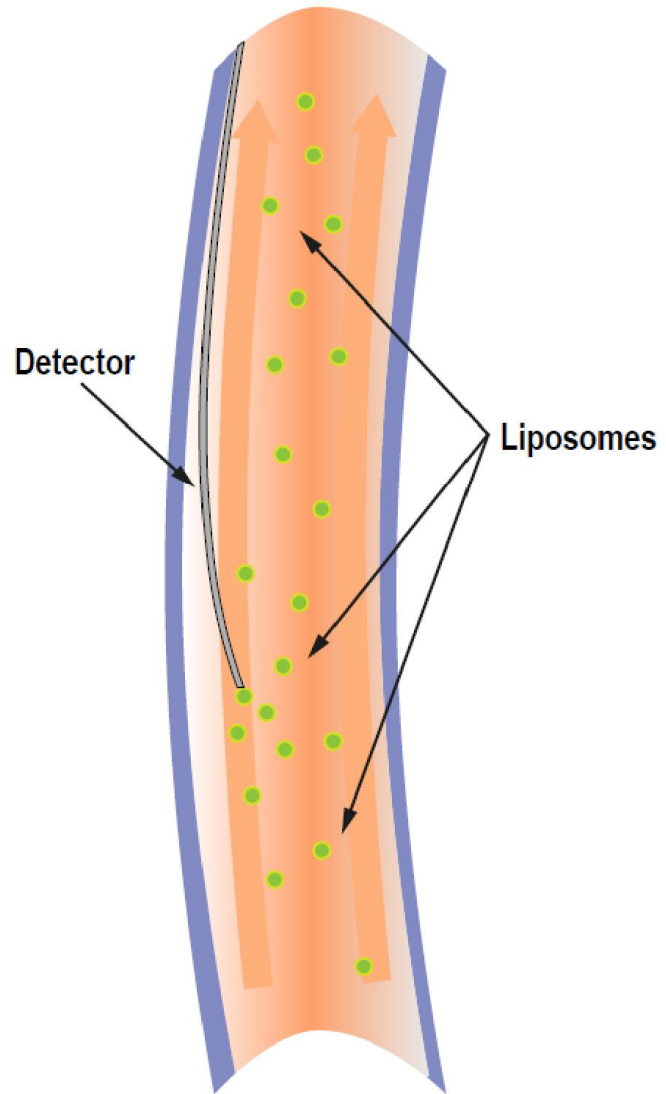
CHEMICAL
EXCITATION, FRET, CHANGE:



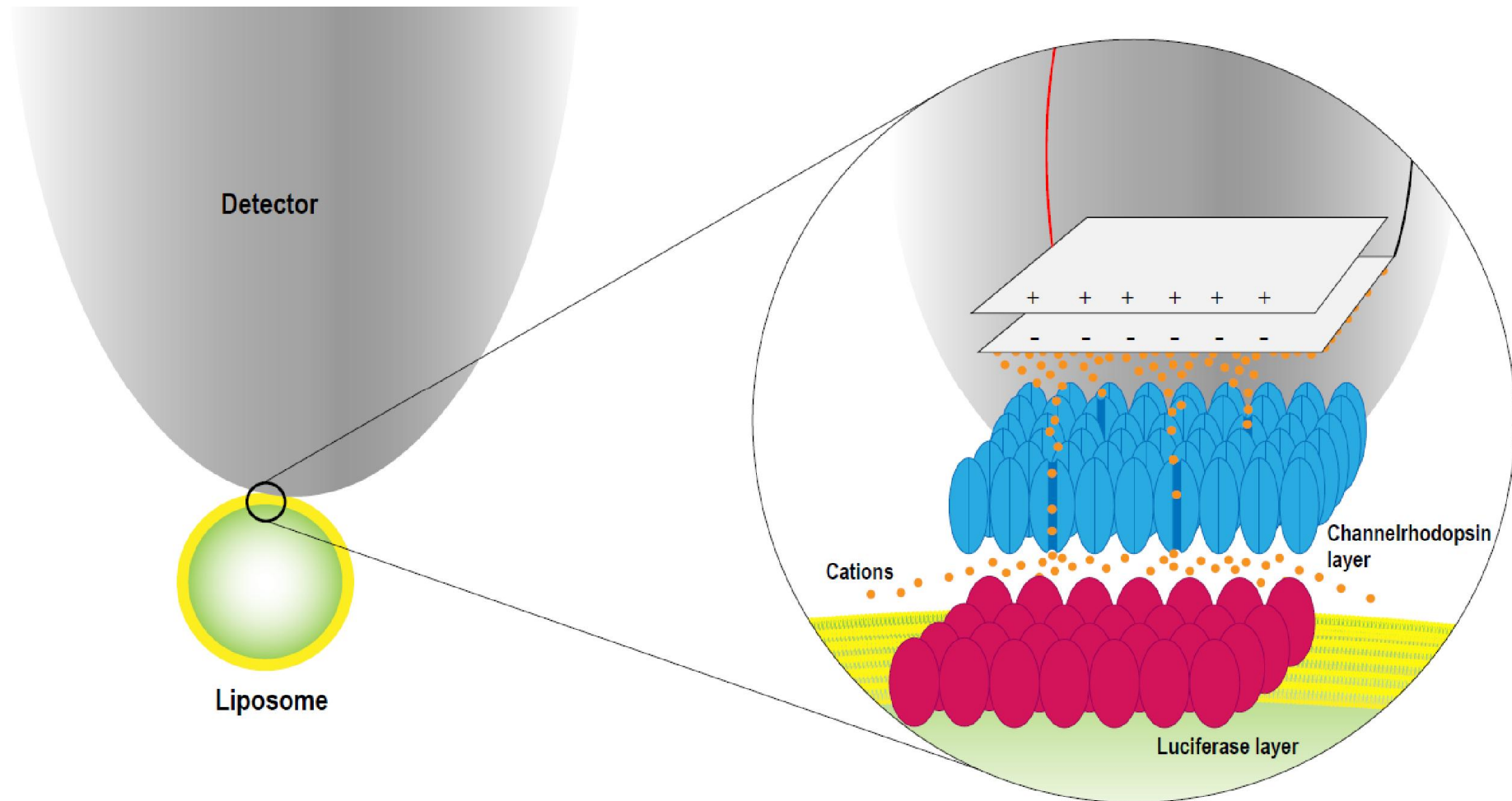
CHANNELRHODOPSIN
*(after excitation,
opens a channel for cations)*



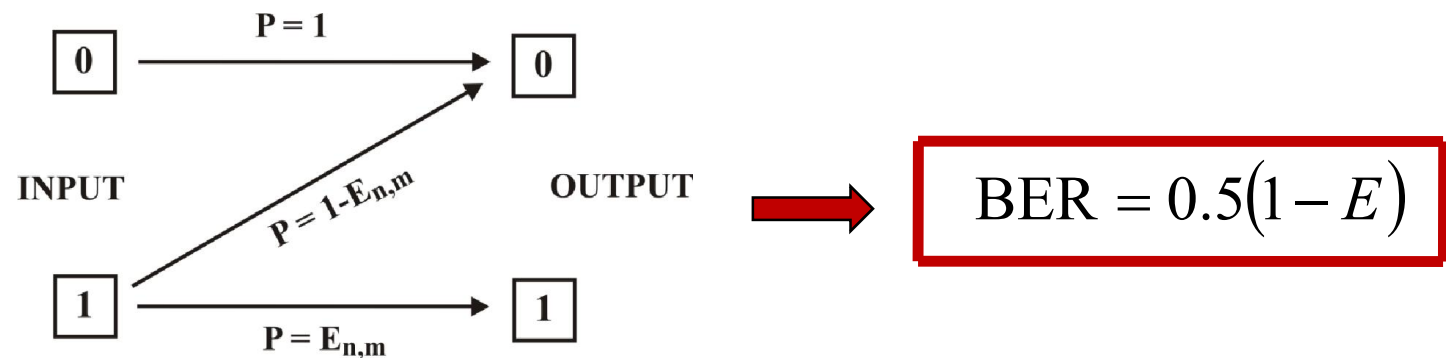
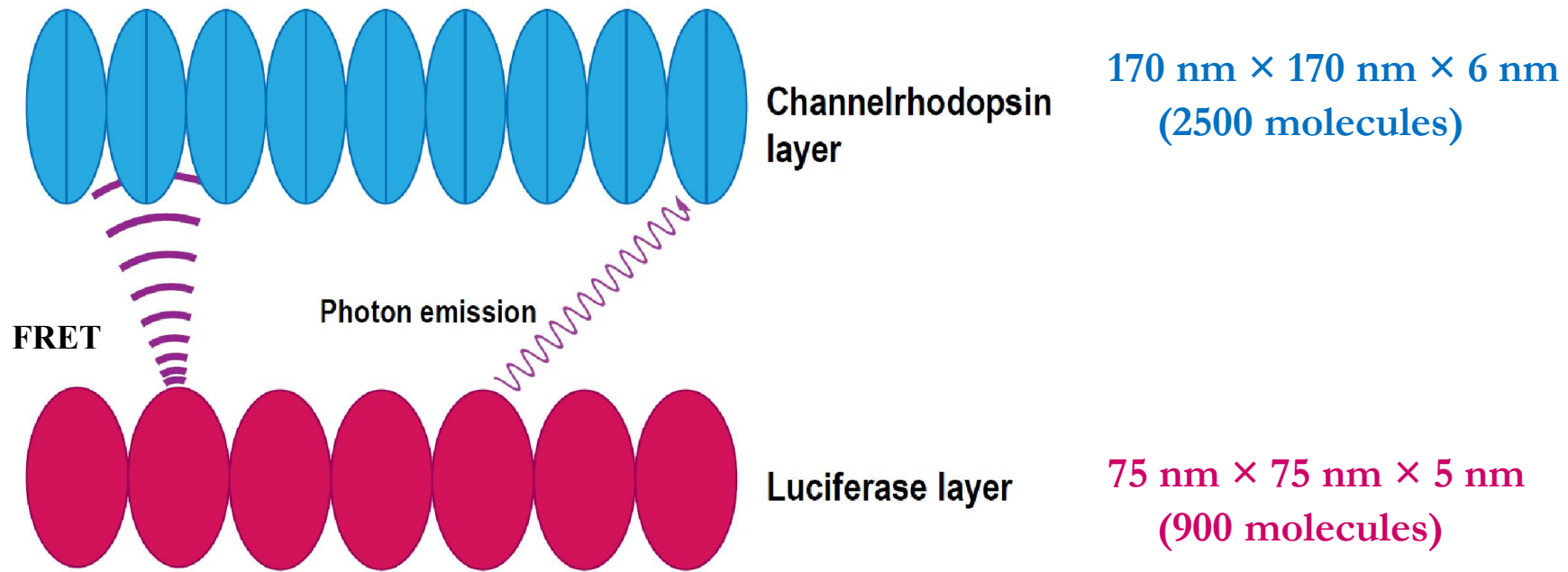
In-body medical system



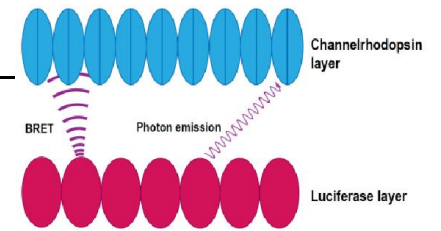
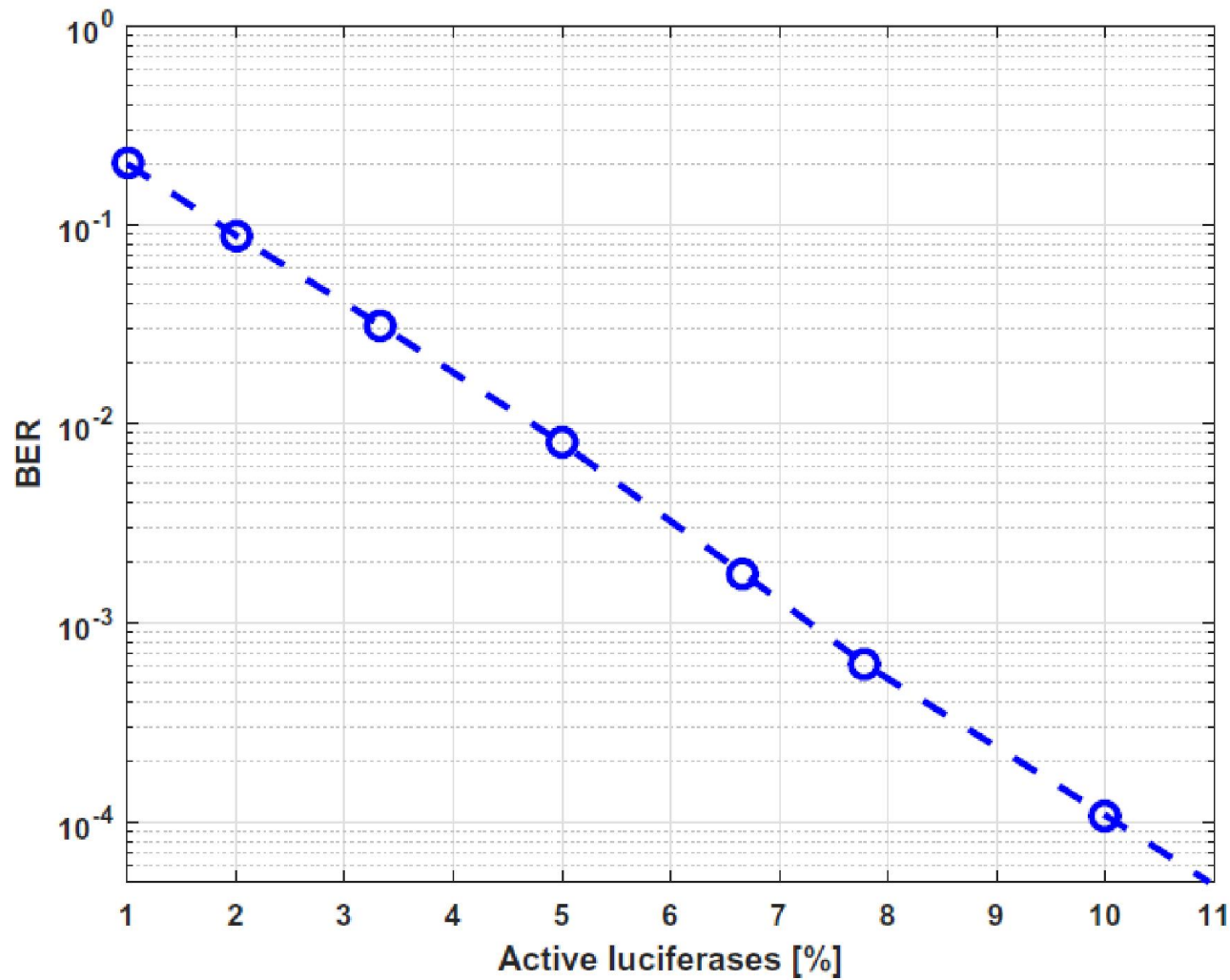
FRET-based communication



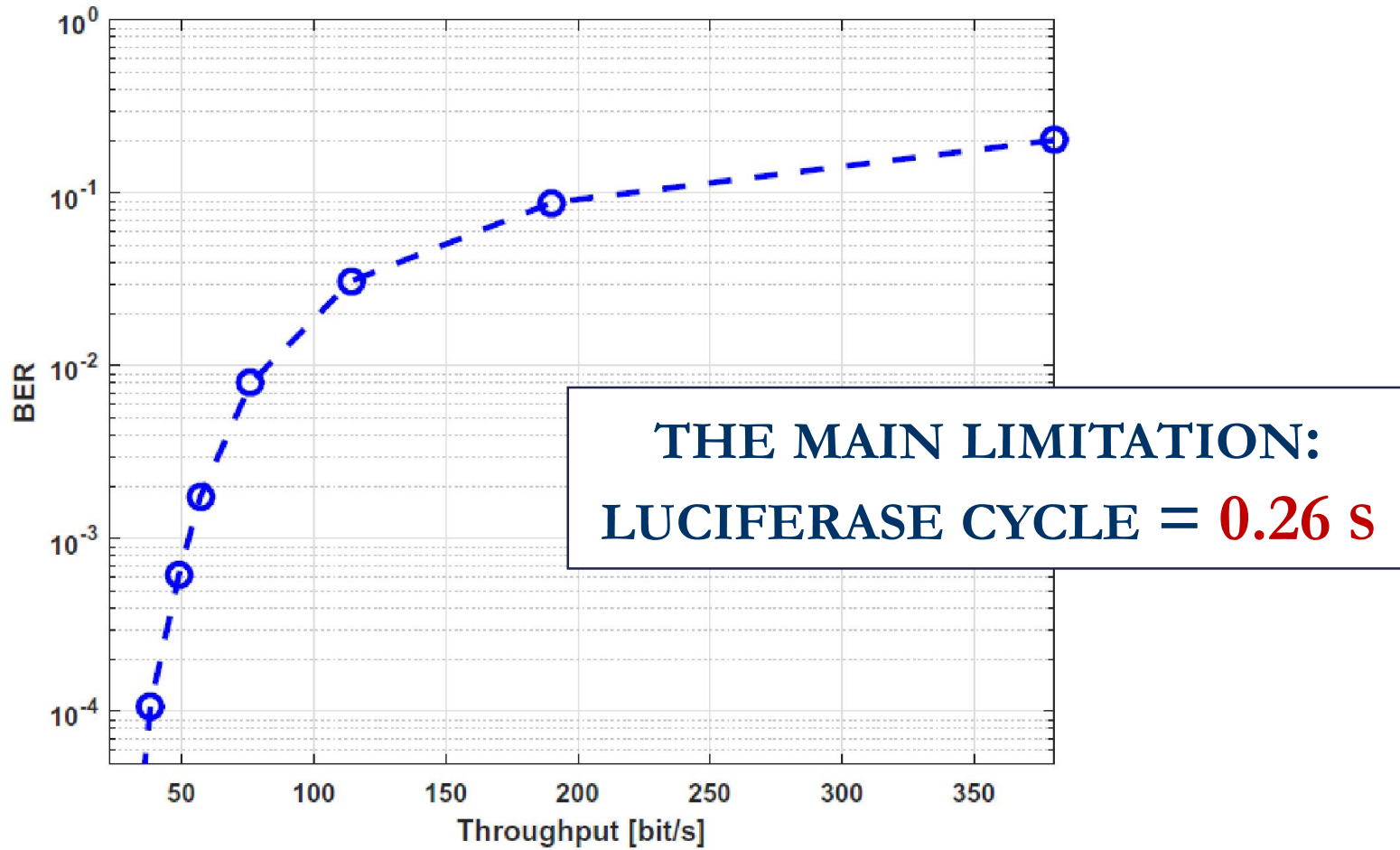
Communication with ON-OFF modulation



Bit error rate



BER vs. throughput trade-off



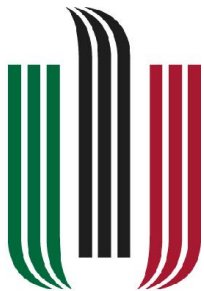
Open issues

1. Routed signal **LEAKAGES**
 - maybe fluorophores in different EM bands?
2. Switching **TIME**
3. **CONTROL** over the transmitter (e.g. providing energy)
4. Nanomachine **MOVEMENTS**
5. Experiments on **SINGLE** nanostructures,
e.g. single molecule **FRET**?

Conclusions

1. Nanocommunications – a very hot topic, but requiring stronger **BIO-TELE** cooperation
2. EM (Terahertz) communications
– suitable rather for **MICRO** world
3. **Molecular** (bio-inspired) approach:
applicable for nanoscale, but quite slow in most cases
4. **FRET**:
fast, but distance limited, MIMO-FRET required
5. Routing is based on features of **specific molecules**
6. Nanocommunications is still **basic science**,
more experiments are welcome!

Thank you for your attention



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