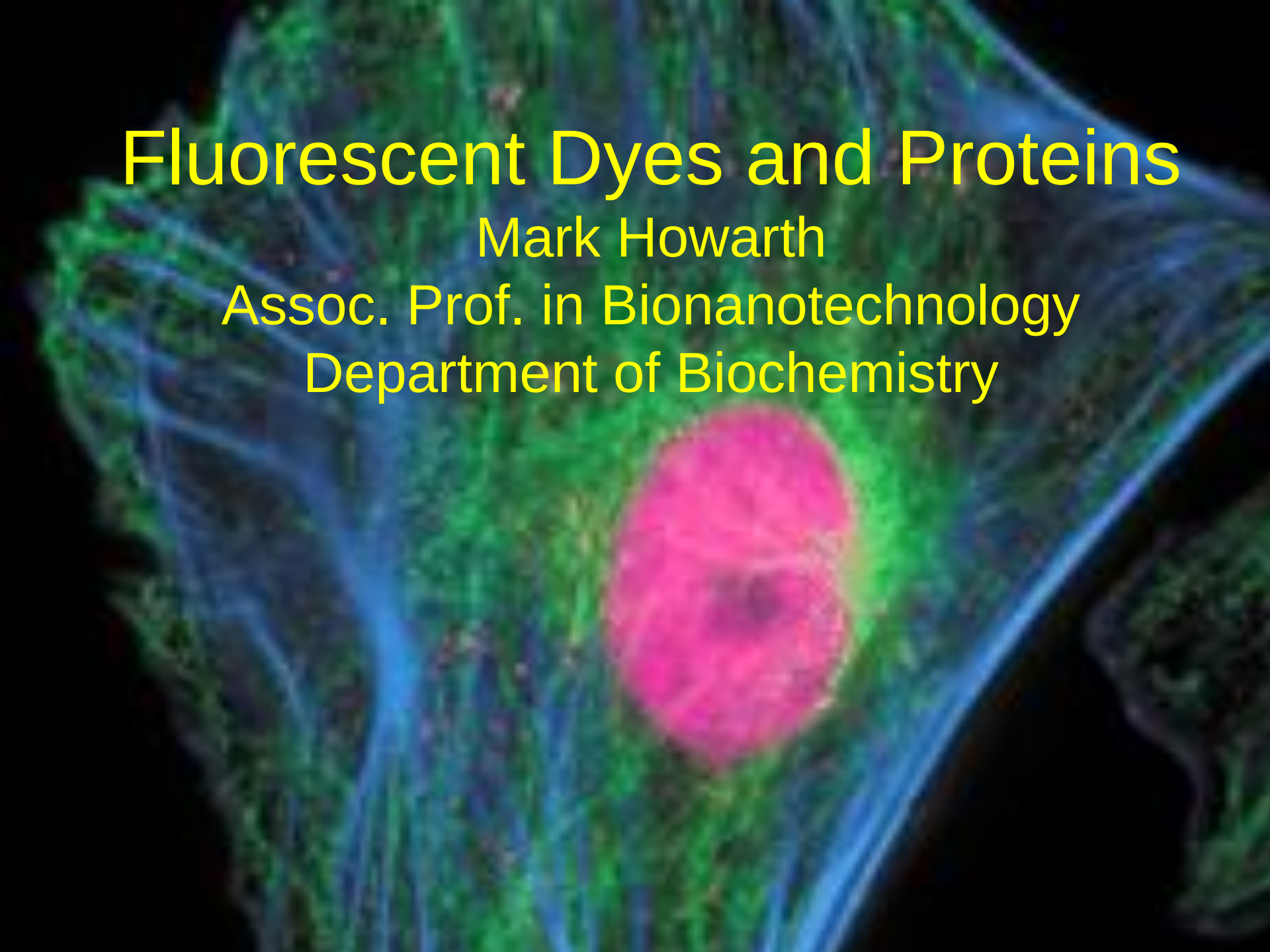
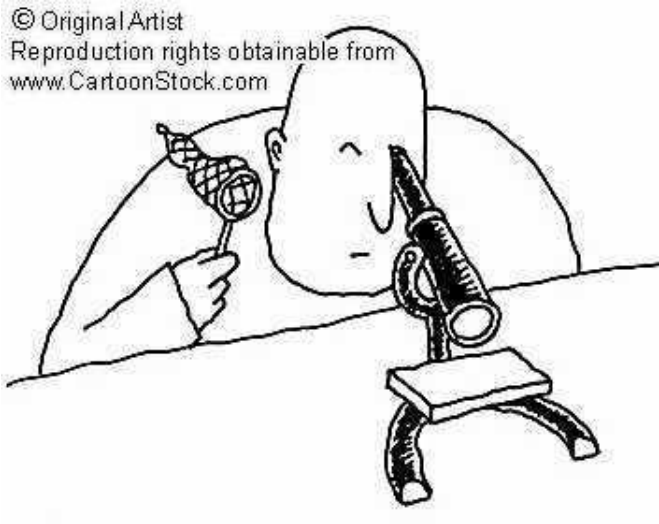


Fluorescent Dyes and Proteins

Mark Howarth
Assoc. Prof. in Bionanotechnology
Department of Biochemistry



Overview

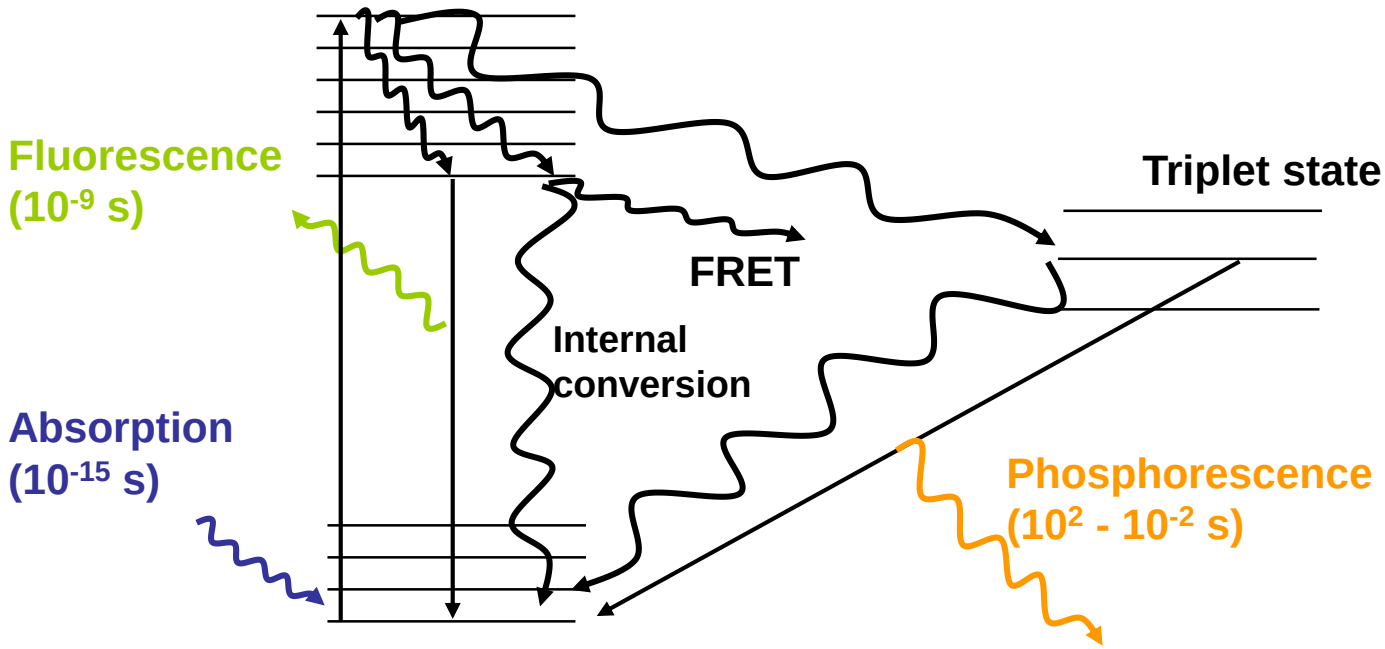
1. What kind of structures are fluorescent

2. How to make and target fluorescent probes

3. Fluorescent probes for cellular structure and function

4. Using light to control cells

Not all energy emitted as fluorescence



$$\text{Quantum yield} = \frac{\text{no. of fluorescent photons emitted}}{\text{no. of photons absorbed}}$$

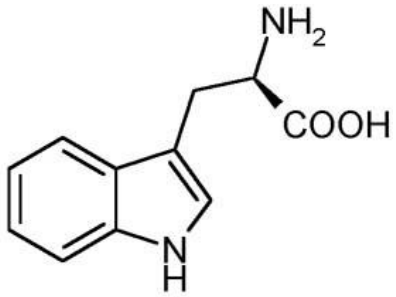
e.g. EGFP QY=0.6 For every 10 photons absorbed, 6 are emitted. (at optimal temp, pH etc.)

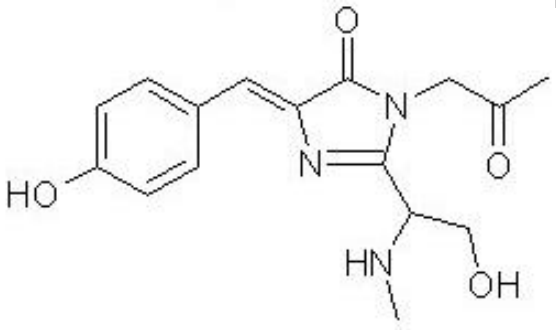
What sort of molecules are fluorescent?

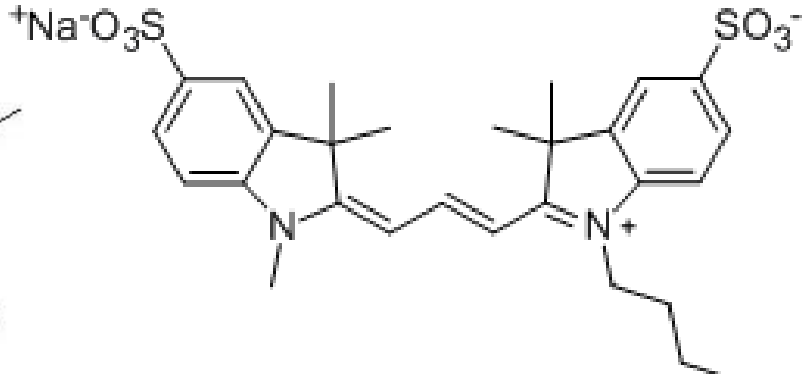
Organic fluorophores
especially
1. Intrinsic fluorophores (source of autofluorescence)
2. Dyes
3. Fluorescent proteins

Inorganic fluorophores
especially
1. Lanthanides
2. Quantum dots

What sort of molecules are fluorescent?
1. Organic fluorophores





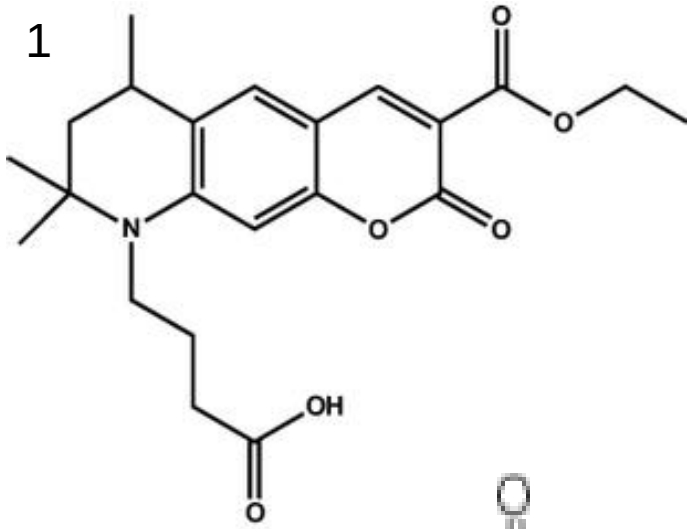


Chemical features:

1. Conjugation
2. Rigidity especially fused aromatic rings
3. Heteroatoms

Relating structure to fluorescence properties

1



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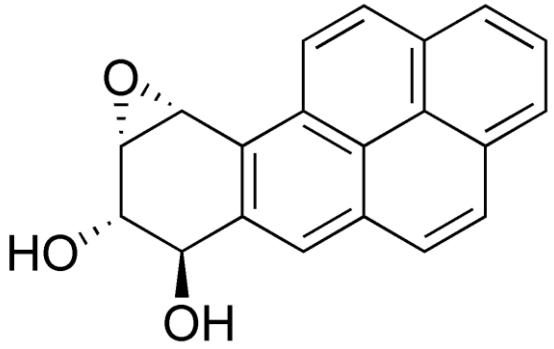

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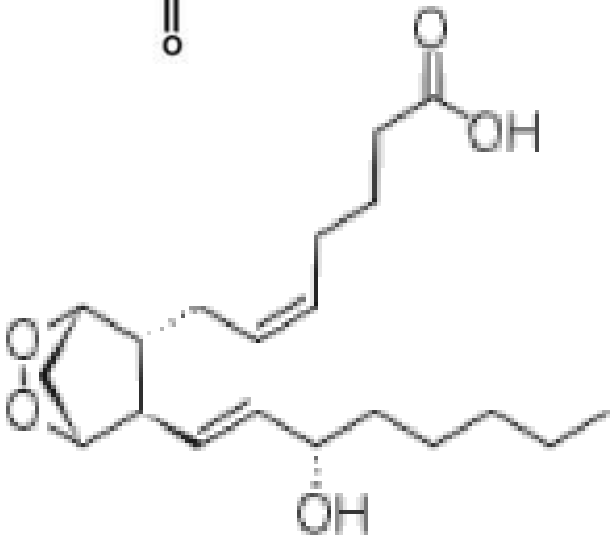

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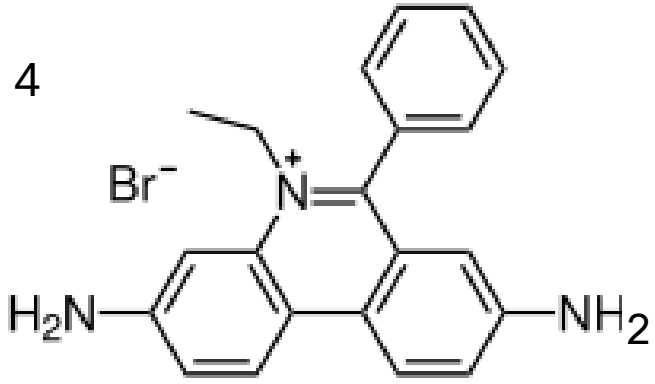


Which two are the best fluorophores?

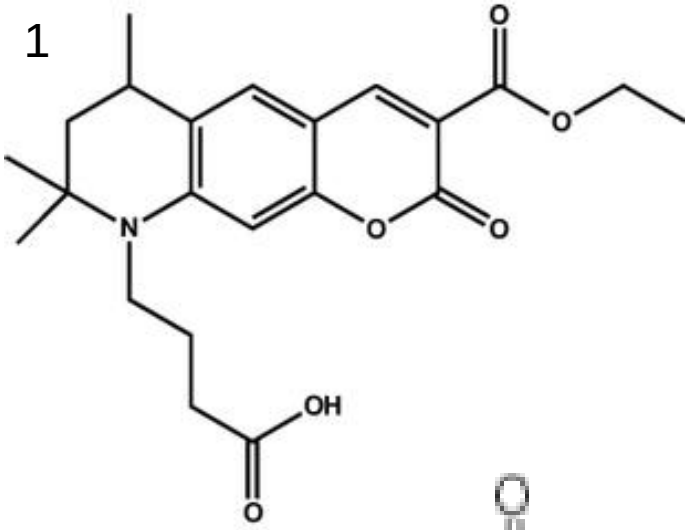
A. 1,2

B. 1,4

C. 2,3

D. 2,4

1



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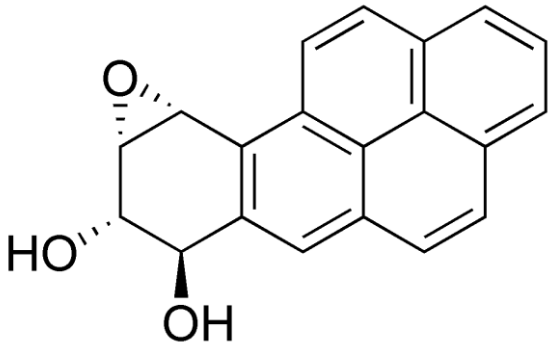

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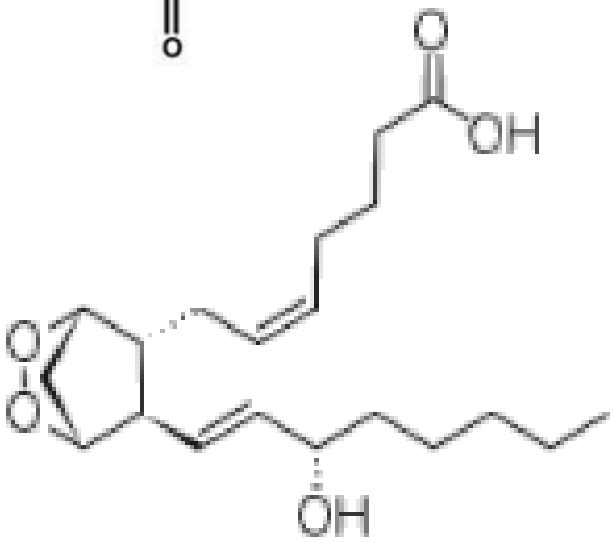

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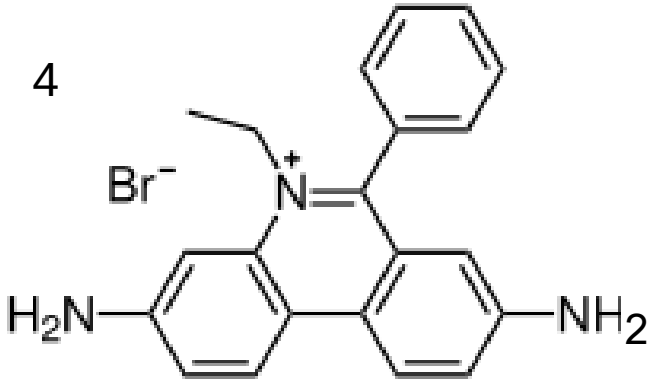
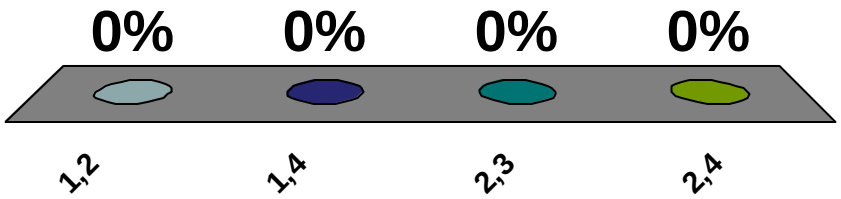
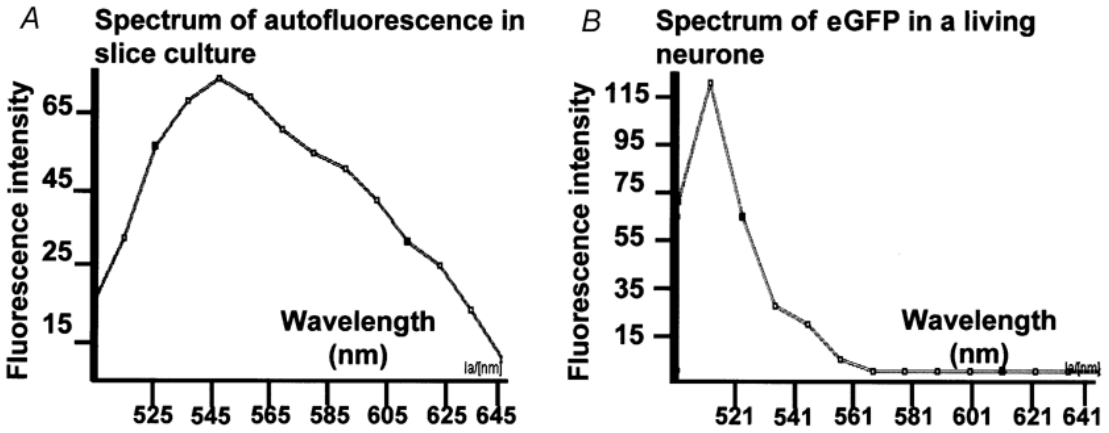
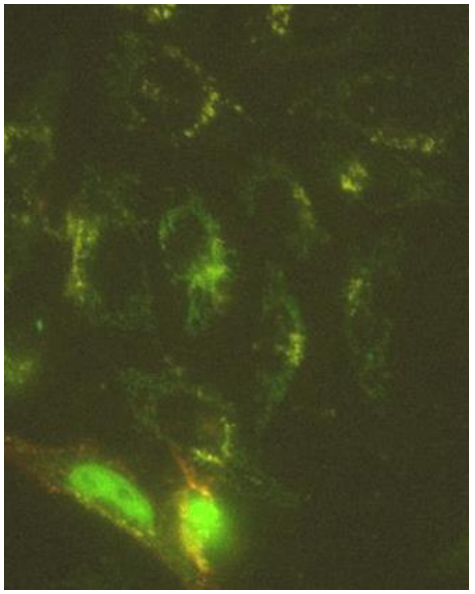




What sort of molecules are fluorescent?
1. Endogenous organic fluorophores





Most common autofluorescent molecules:

Flavins, NADH, NADPH, elastin, collagen, lipofuscin

Avoiding autofluorescence:

choose dye emitting in red with big Stokes shift

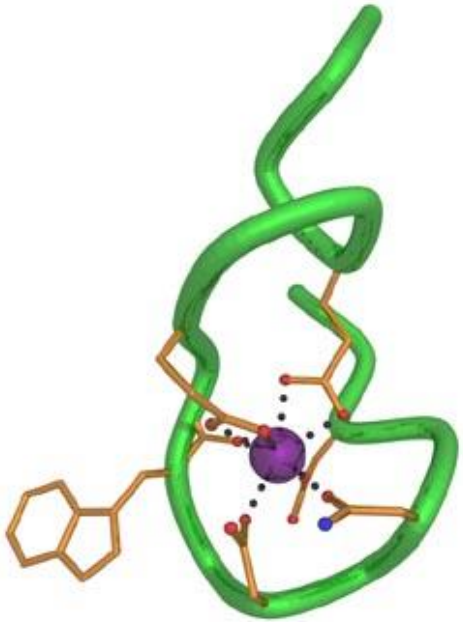
add quencher (Crystal violet)

time-gate fluorescence

What sort of molecules are fluorescent?
2. Inorganic fluorophores

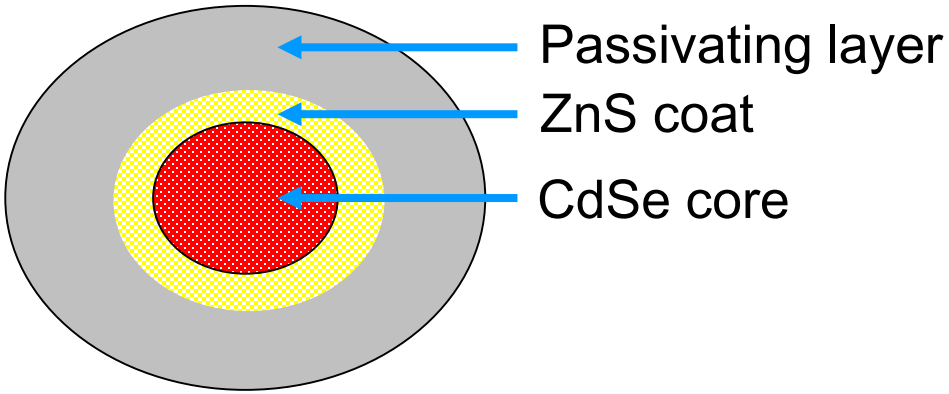
Lanthanides

Peptide sequence
binds Tb^{3+} and protects
from quenching by water



Curr Opin Chem Biol. 2010;14(2):247-54. Lanthanide-tagged proteins--an illuminating partnership. Allen KN, Imperiali B.

Quantum dots



+ bright, photostable, narrow emission
- large (~20 nm), expensive,
hard to target specifically

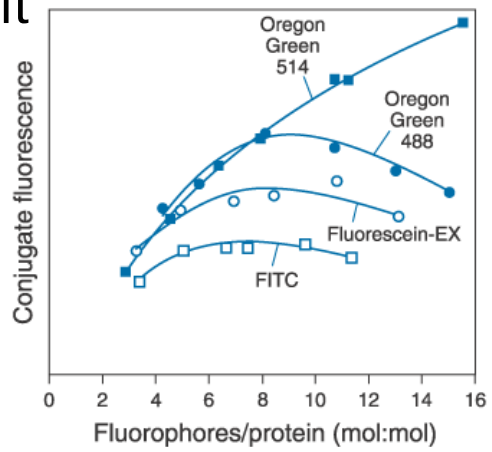
Michalet X, et al. Quantum dots for live cells, in vivo imaging, and diagnostics. Science. 2005 307(5709):538-44.

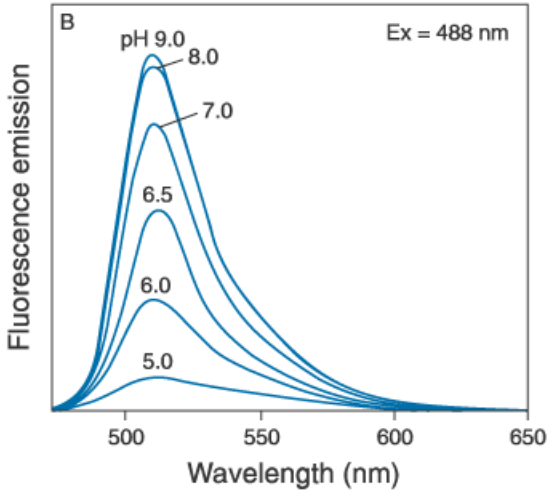
How good is a fluorophore?

1. Excitation and emission appropriate
   - background worse in UV + with small Stokes shift
   - good match to filters on your microscope
   - look at other fluorophores at same time
2. Bright
   - see small numbers of fluorophores,
   - low self-quenching, high QY and absorbance
3. Stable to photobleaching
   - exciting light damages fluorophore
4. Non-toxic
5. Environment-insensitive (especially to pH)
6. Little non-specific binding
7. Small
8. Little blinking

(9. Cost)

Green dye self-quenching

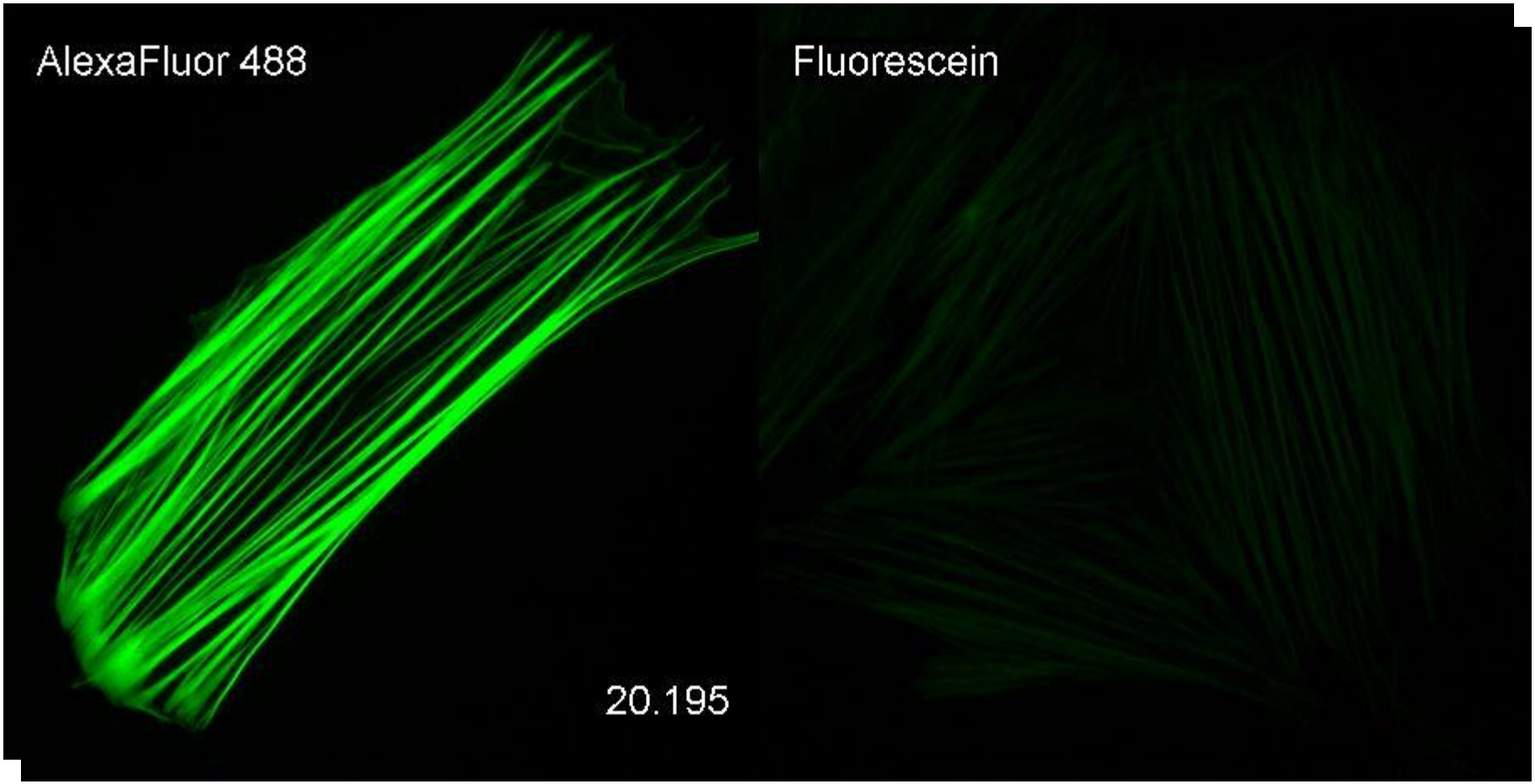




Fluorescein pH sensitivity

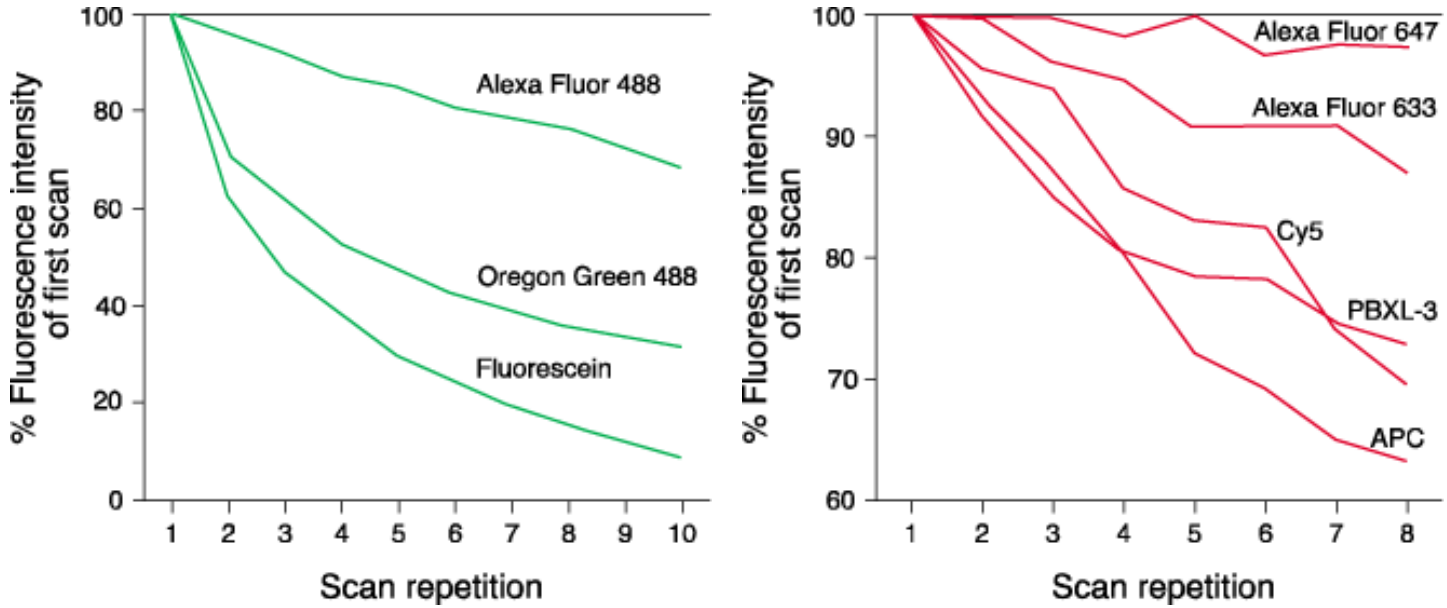
Alexa Fluor 488 vs Fluorescein Bleaching

2x Real Time



Alexa Fluor Dyes – Photostability

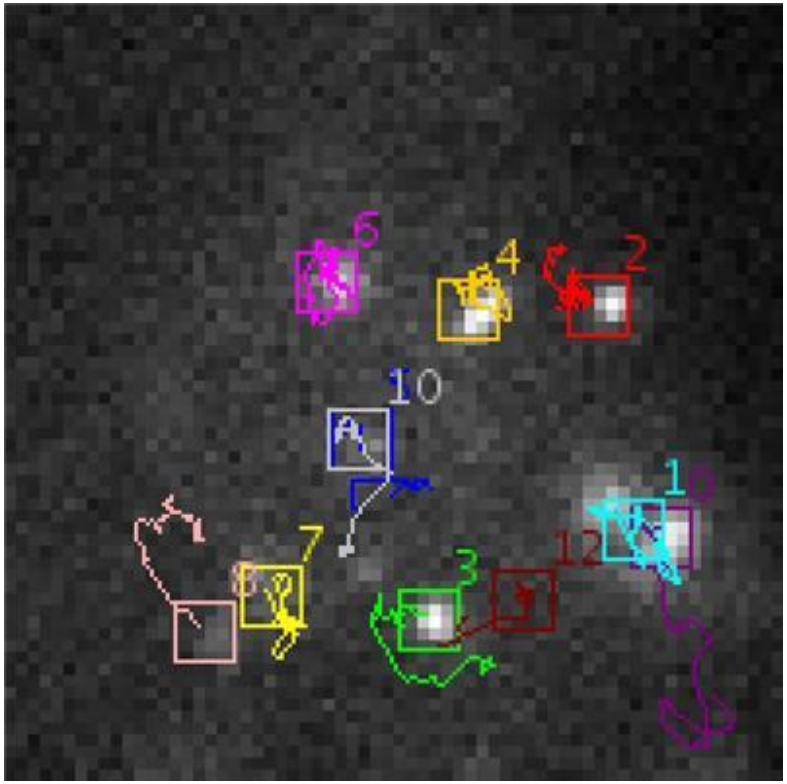
Laser-scanning cytometry
EL4 cells
biotin-anti-CD44 + streptavidin conjugates

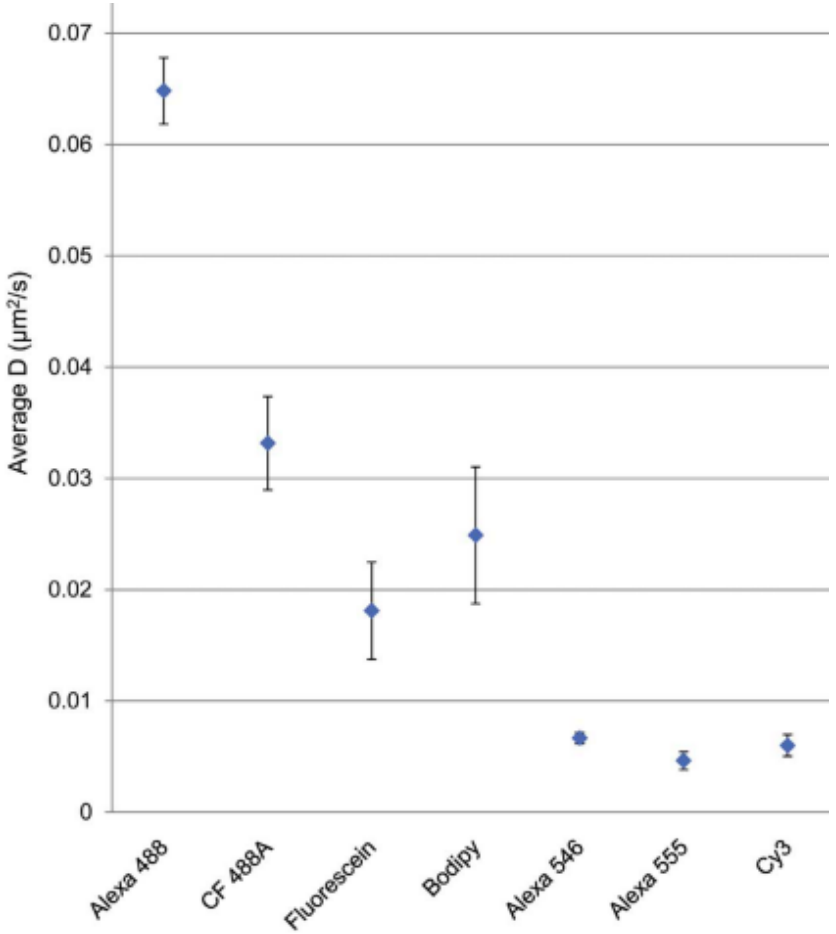


Fluorescein is the commonest dye
but has poor photostability.
Also consider Atto dyes (Sigma) and Dyomics dyes

Dye affects non-specific binding and receptor mobility

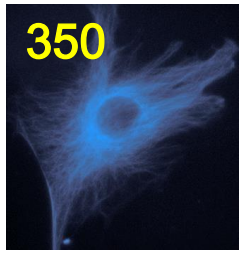
Dye makes a big difference to non-specific binding
Diffusion coefficient for dye-labelled Affibody against EGFR varies 10-fold with hydrophilic versus more hydrophobic dye!
L. Zanetti-Domingues et al.
PLoS one 2013

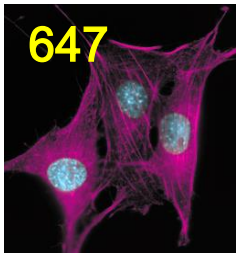




Multiplexing- four main colours

Excitation wavelengths:





Emission wavelengths:

Blue green orange/red far red

350 400 450 500 550 600 650 700

DAPI/UV

FITC

TRITC

FAR RED

Alexa Fluor® 350
Coumarin, AMCA

Alexa Fluor® 488
Fluorescein (FITC)
Cy2

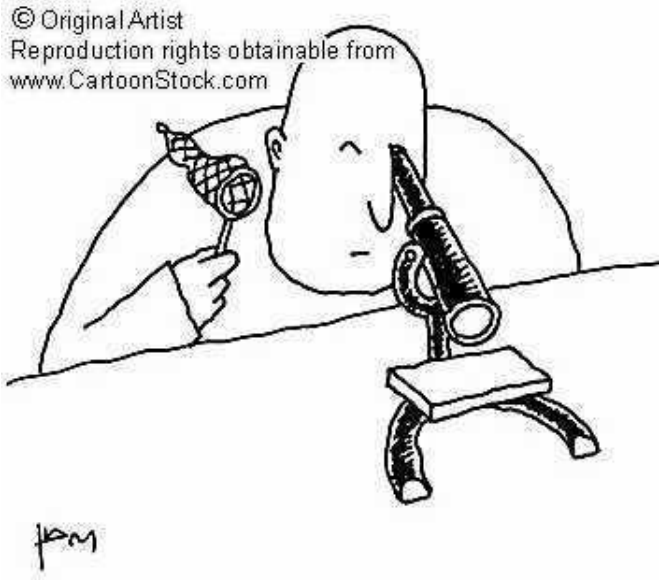
Alexa Fluor® 555
Rhodamine,
TAMRA, TRITC
Cy3

Alexa Fluor® 647
Cy5, APC

Alexa Fluor® 594
Texas Red, Cy3.5

Colour Selection ♦ Brightness ♦ Photostability

Overview



1. What kind of structures are fluorescent

2. How to make and target fluorescent probes

3. Fluorescent probes for cellular structure and function

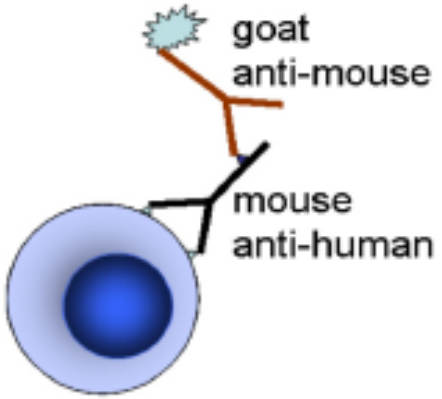
4. Using light to control cells

Antibodies for cellular imaging

Live cells

Label plasma membrane and secretory pathway
Penetrate plasma membrane
(microinjection, electroporation, pinosome lysis, streptolysin, cell permeable peptides, ester cage)

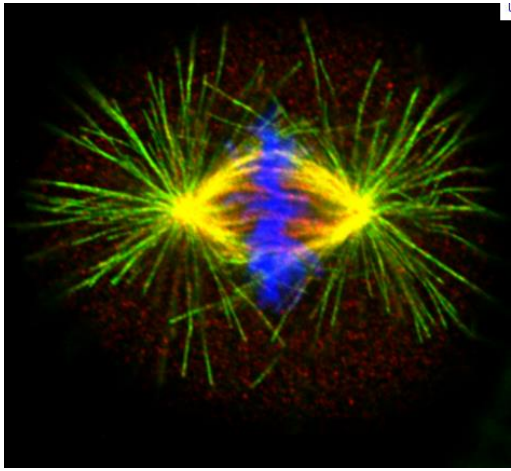
Get dynamics, avoid fixation artifacts



Fixed cells

Permeabilise

Still can give enormous amount of useful information

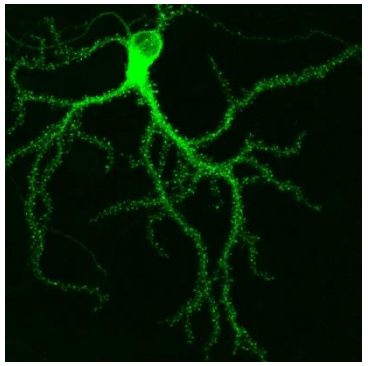


Not just antibodies for targeting

Other types of targeting agents:

Proteins
(especially antibodies, but also transferrin, insulin, EGF etc.)
Peptides (MHC class I pathway, proteasome function)
RNA (mRNA, molecular beacons, aptamers, siRNA)
DNA
lipids, lipoproteins
drugs

?



How to dye: it is easy

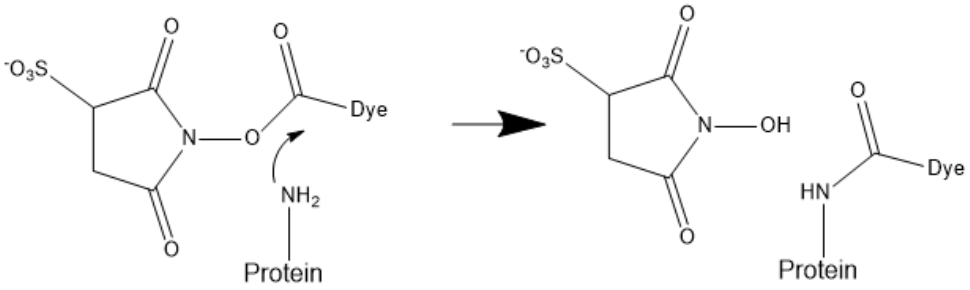
Multiple ways to modify proteins
(see Molecular Probes catalogue)

Most common ways are to modify:

1. Lysine

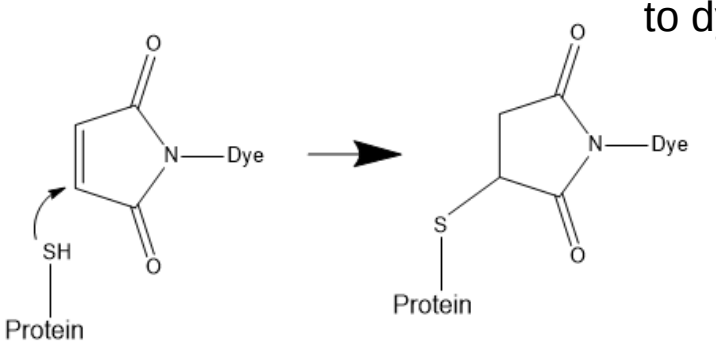
or

2. Cysteine



sulfoNHS-dye

Amide bond to dye



maleimide-dye

Thioether bond to dye

A Add dye to protein for 3 hr
B 1cm Sephadex column to remove most free dye (10 min)
C Dialyse away rest of free dye (24 hr)

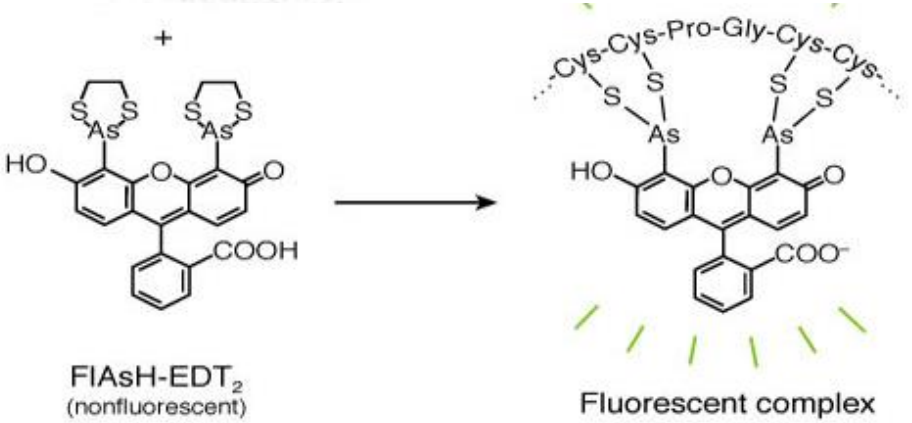
Site-specific protein labelling methods

1. Binding domain

SNAP-tag (19 kDa NEB), HaloTag (34 kDa, Promega)

2. Binding peptide

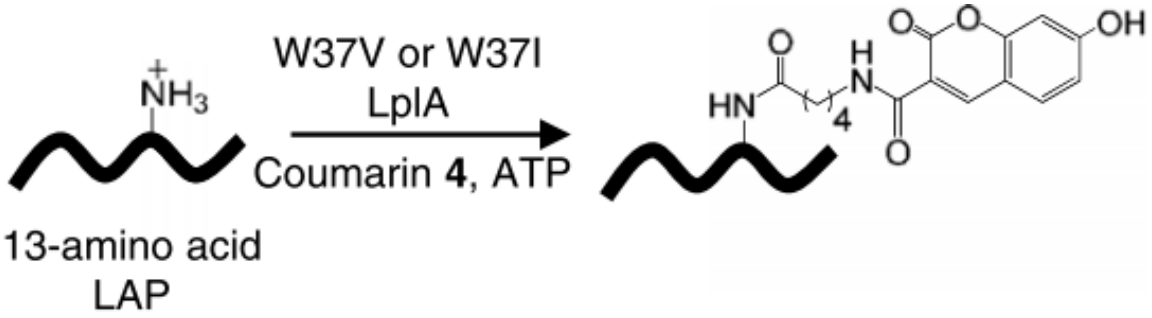
FlAsH (Invitrogen)



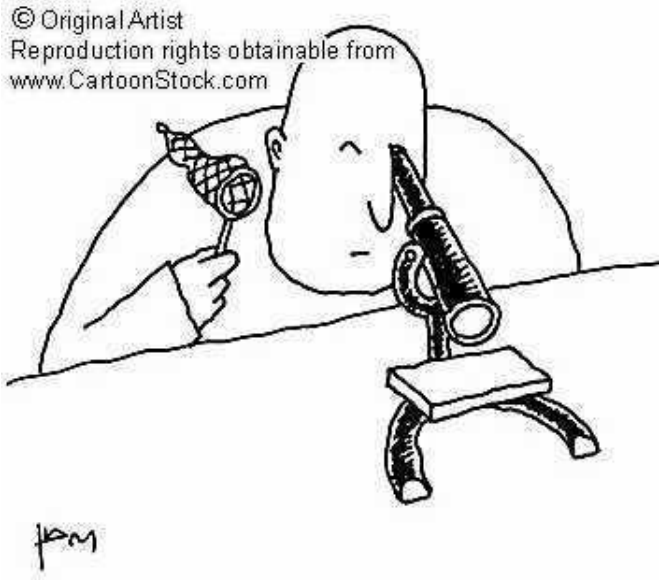
3. Enzymatic ligation to peptide

PRIME

AY Ting PNAS 2010



Overview



1. What kind of structures are fluorescent

2. How to make and target fluorescent probes

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Putting the signal in context: nuclear labelling

(follow DNA even when nucleus breaks down)

Fixed cells:

Intercalate into DNA

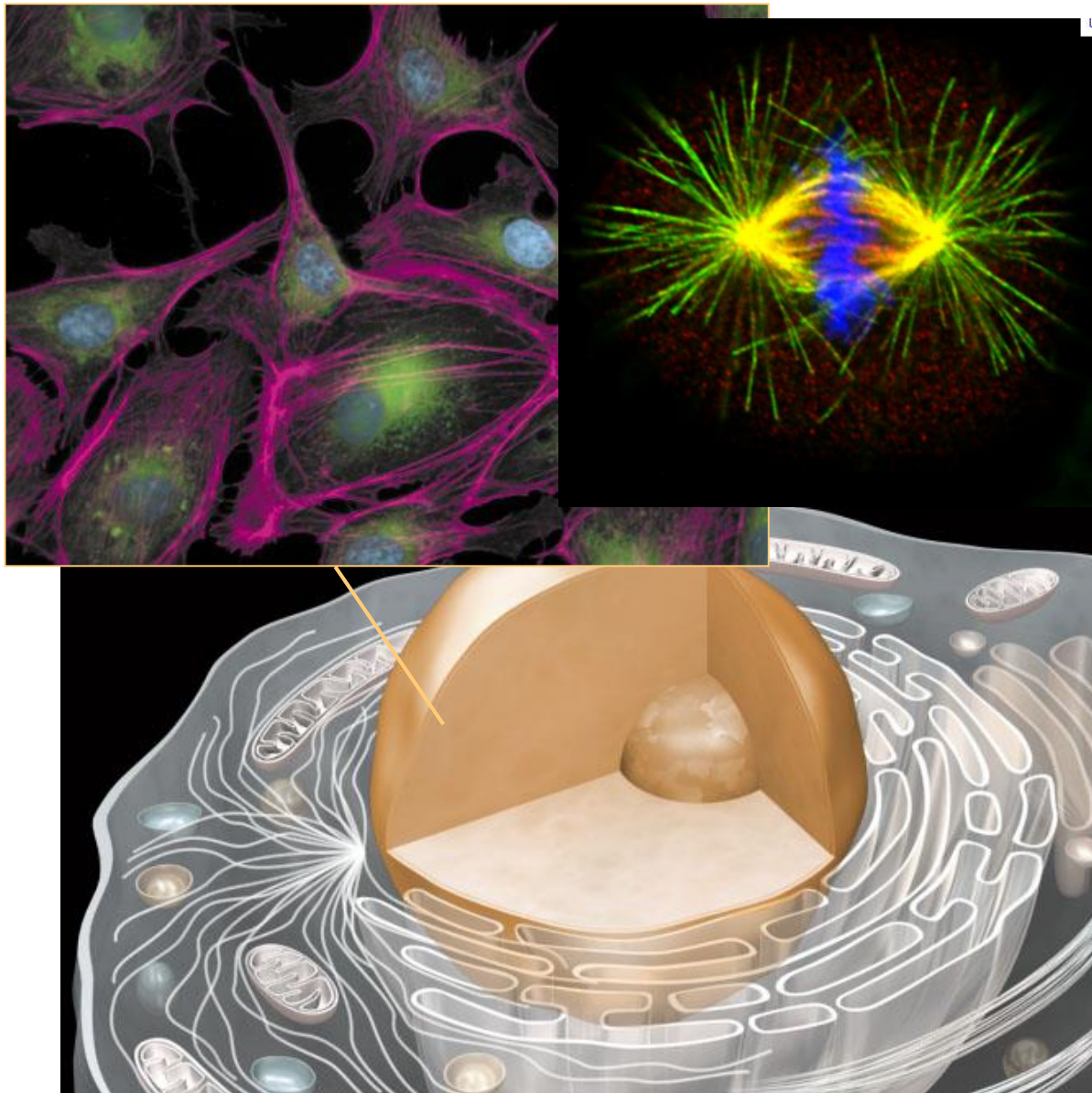
DAPI

(well away from other channels)

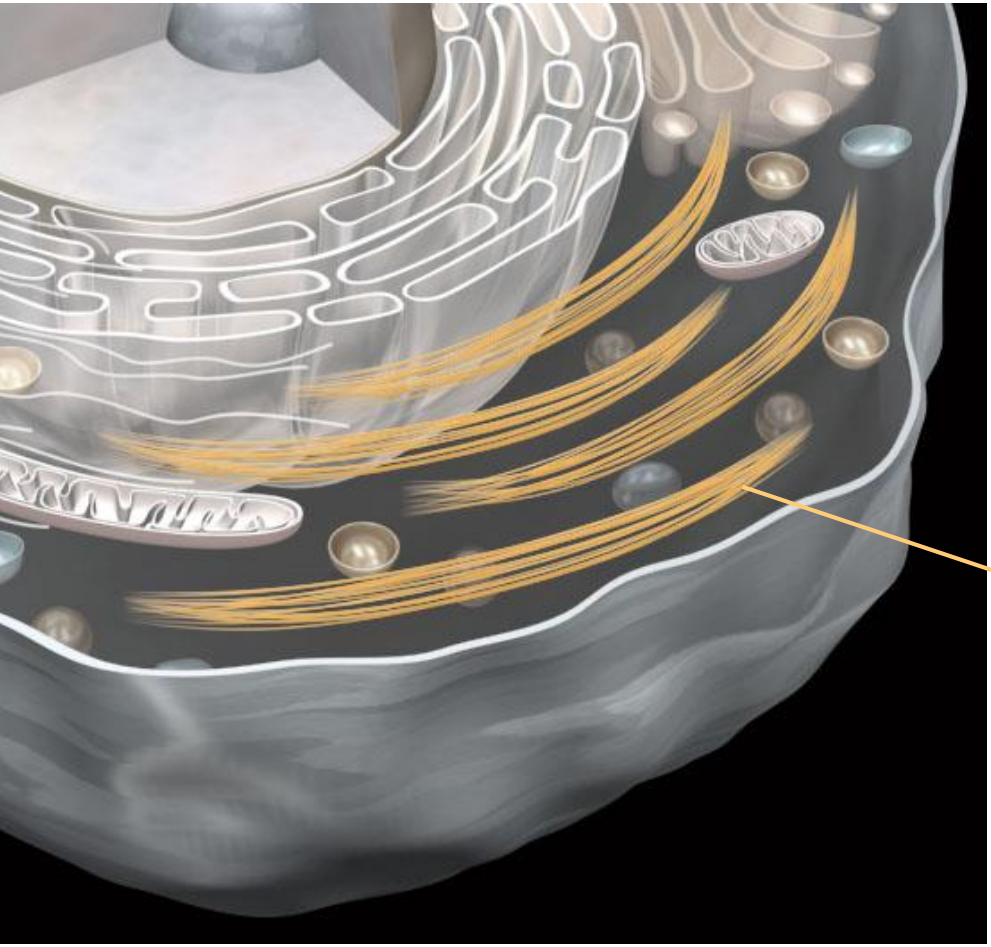
Hoechst 33342

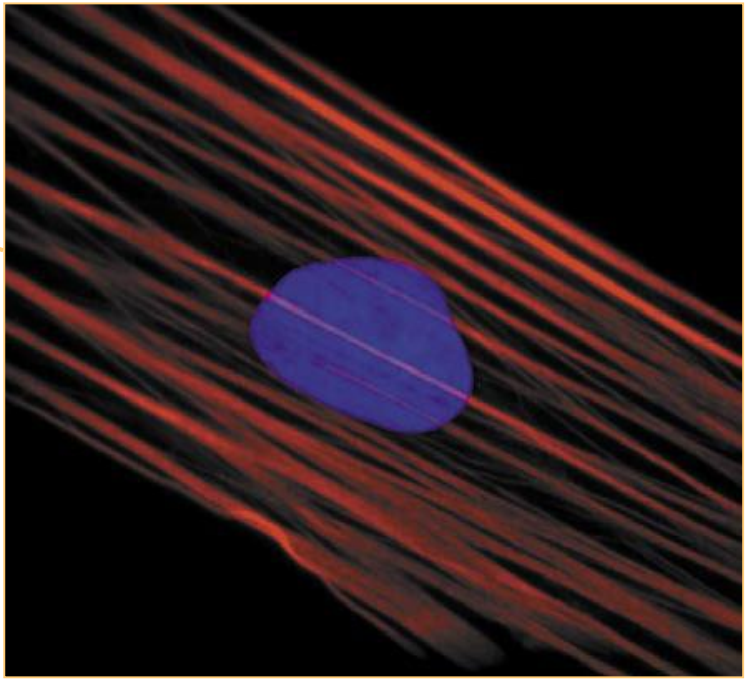
Live cells:

histone H2B-GFP



Putting the signal in context: actin labelling





Fixed cells: phalloidin-dye

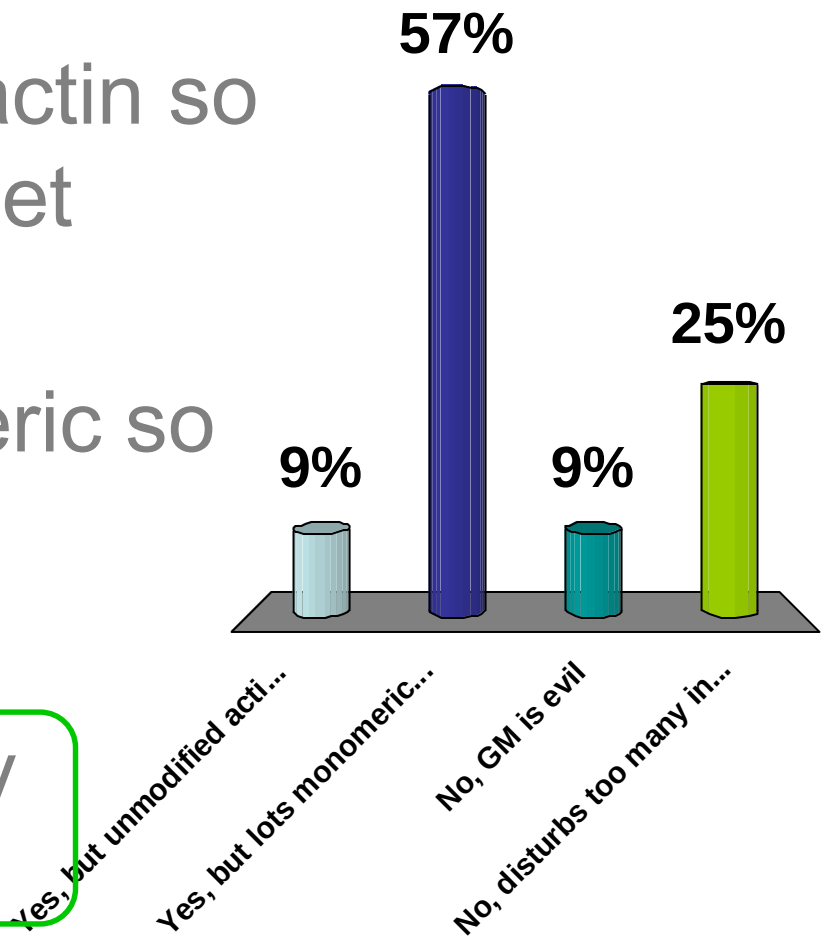
Can you image the actin cytoskeleton in live cells by fusing actin to a fluorescent protein?

A. Yes, but unmodified actin so abundant that won't get much signal

B. Yes, but lots monomeric so fuzzy signal

C. No, GM is evil

D. No, disturbs too many interactions

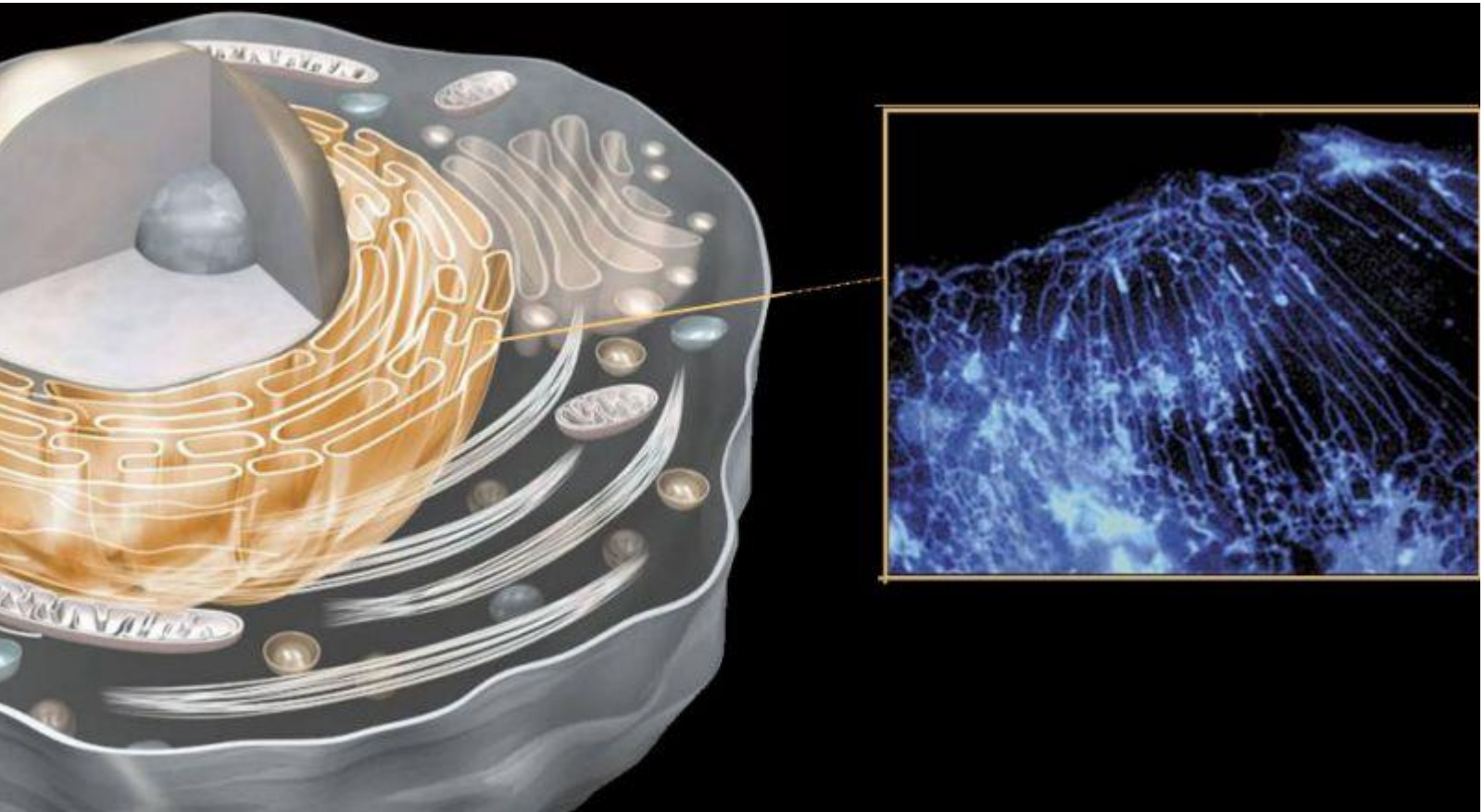


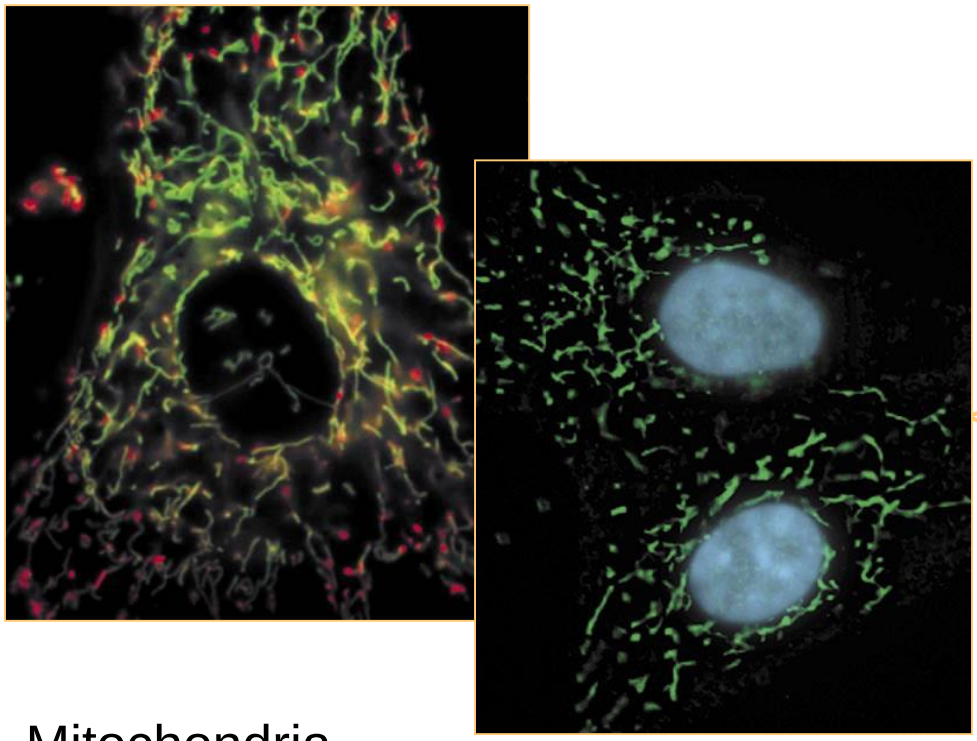
Endoplasmic Reticulum

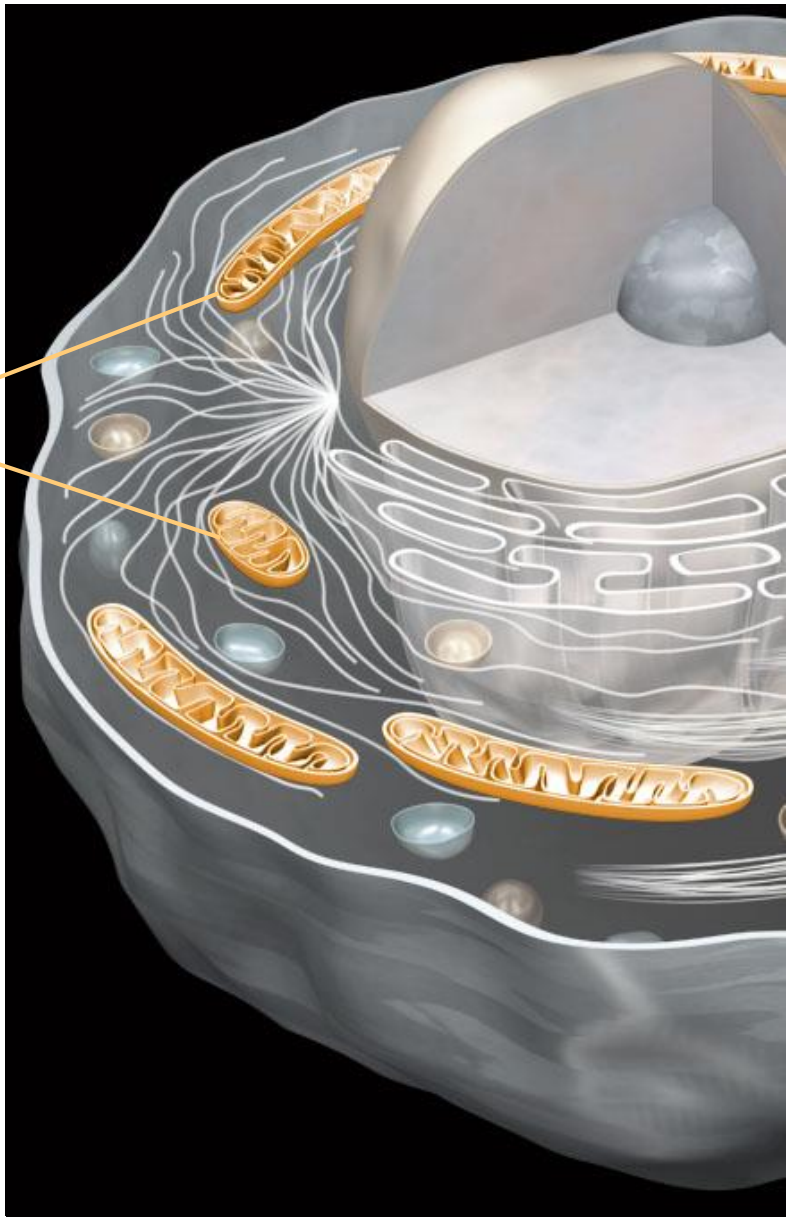
ER-Tracker™ Blue-White DPX

antibody to calnexin

Live cells: ss-GFP-KDEL







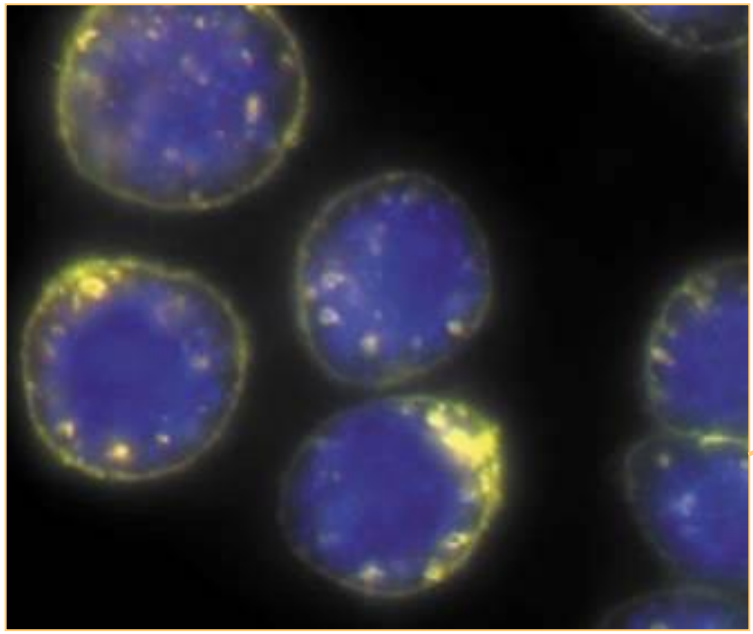
Mitochondria

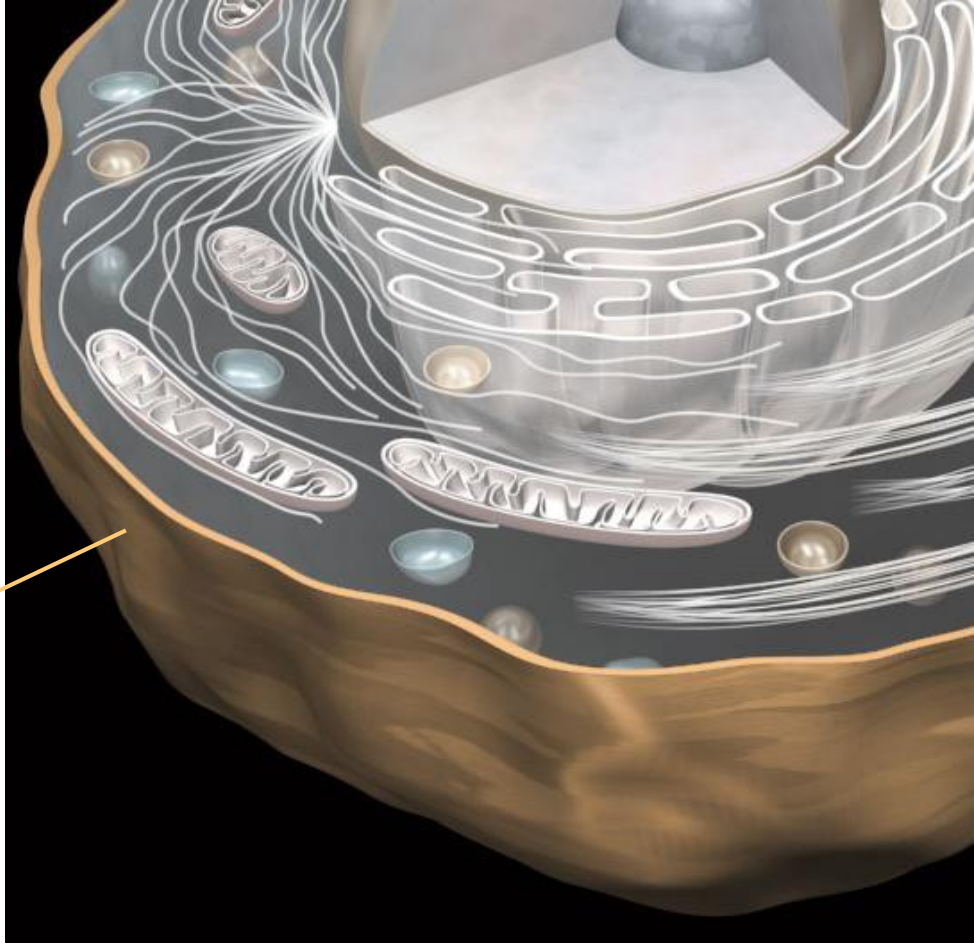
Fixed cells: anti–cytochrome oxidase subunit I Ab

Live cells: MitoTracker® Red/Green/Orange

JC-1 (red J-aggregates at high conc., red to green depends on membrane potential)

Mitochondrial targeting sequence-GFP

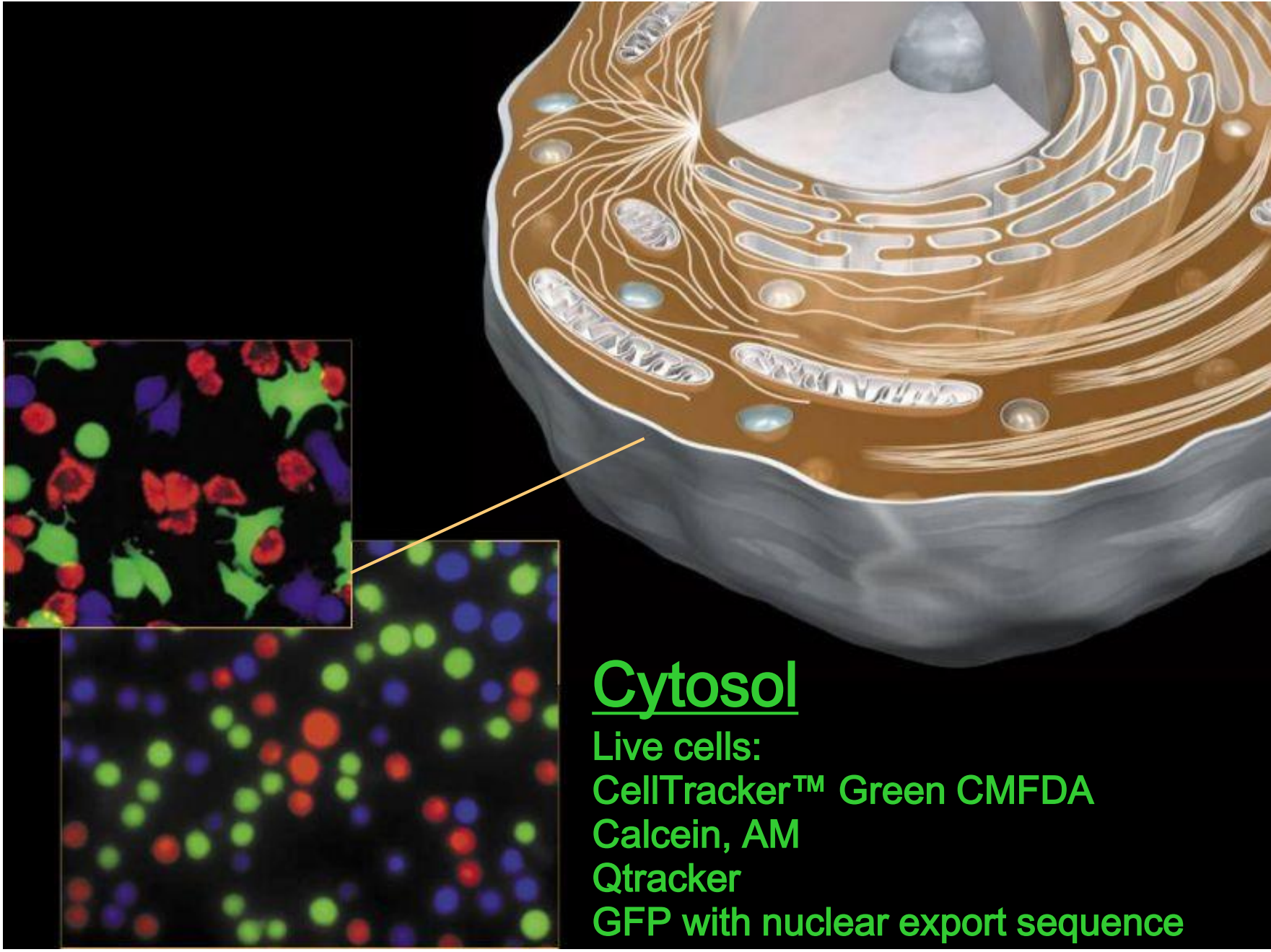




Lipid Rafts

BODIPY® FL C_5-ganglioside GM1

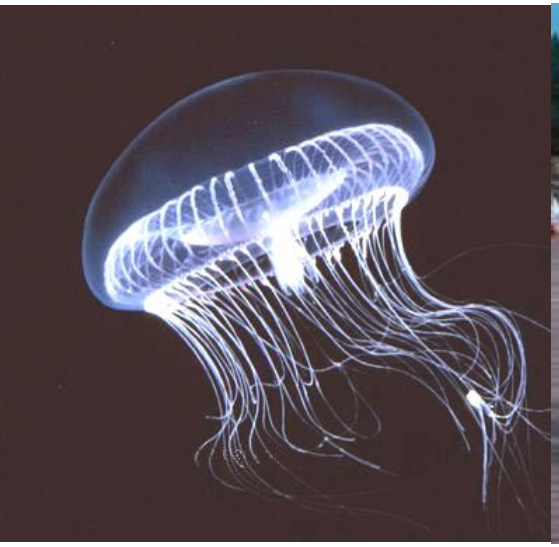
Fluorescent Cholera Toxin subunit B (CT-B)



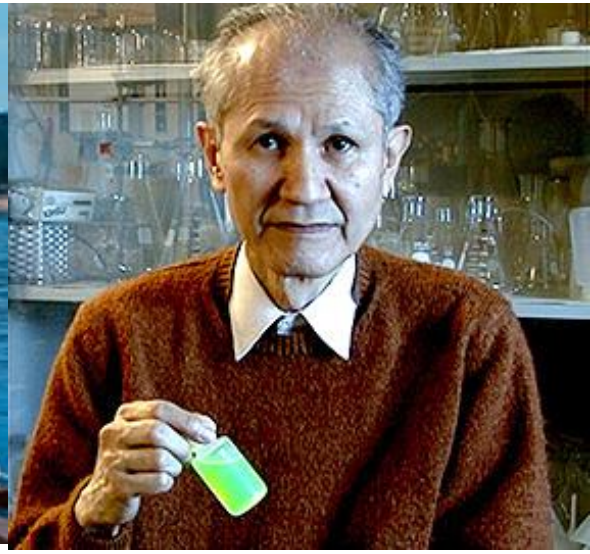
Cytosol

Live cells:
CellTracker™ Green CMFDA
Calcein, AM
Qtracker
GFP with nuclear export sequence

The breakthrough of fluorescent proteins from jellyfish



Aequorea victoria



Osamu Shimomura

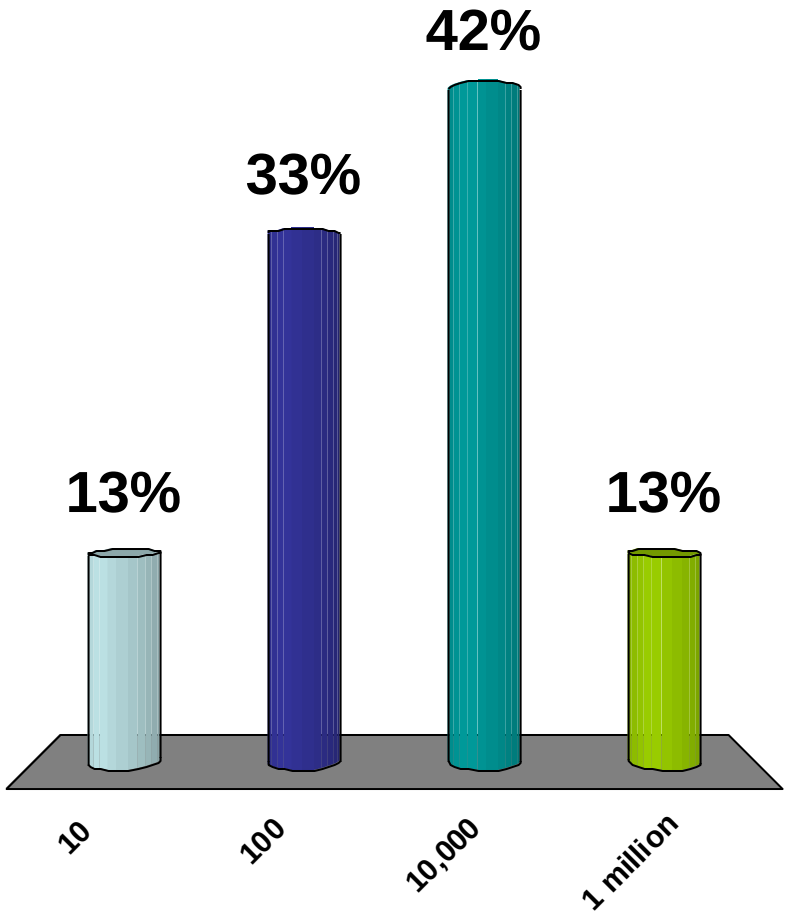
How many jellyfish had to be caught to get enough GFP to analyse?

A. 10

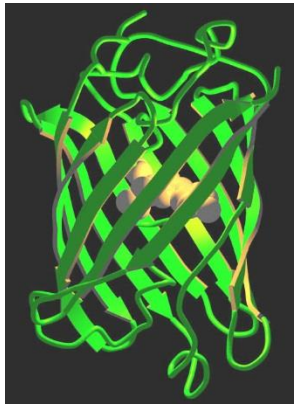
B. 100

C. 10,000

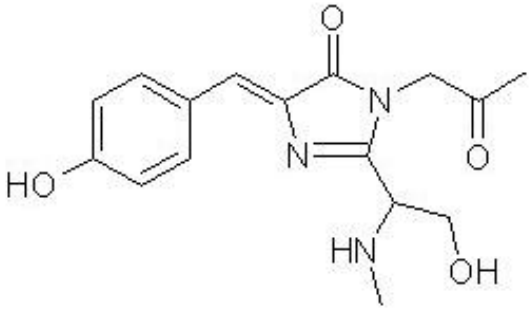
D. 1 million



The breakthrough of fluorescent proteins for live cell imaging

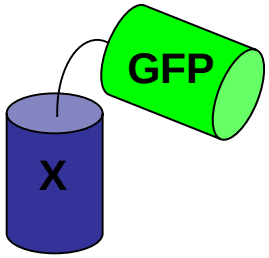


GFP fold
β-can

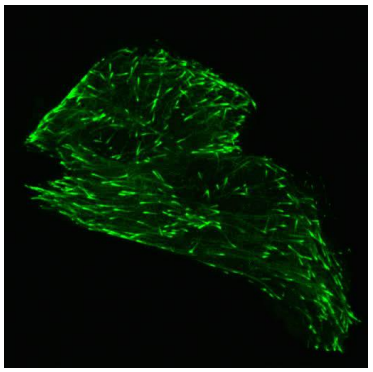


GFP chromophore
from Ser-Tyr-Gly

Link GFP sequence to gene of your favourite protein

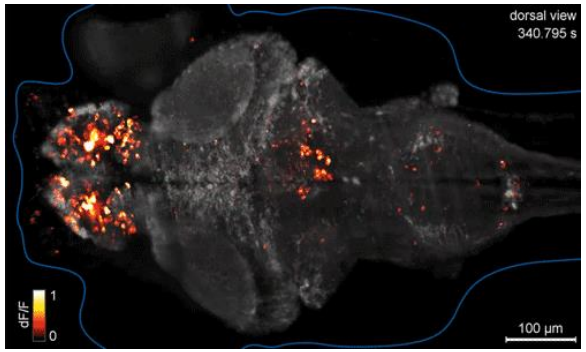


GFP folds
and becomes
fluorescent



GFP lights up your
favourite protein in cell

Fluorescent proteins are more than just labels

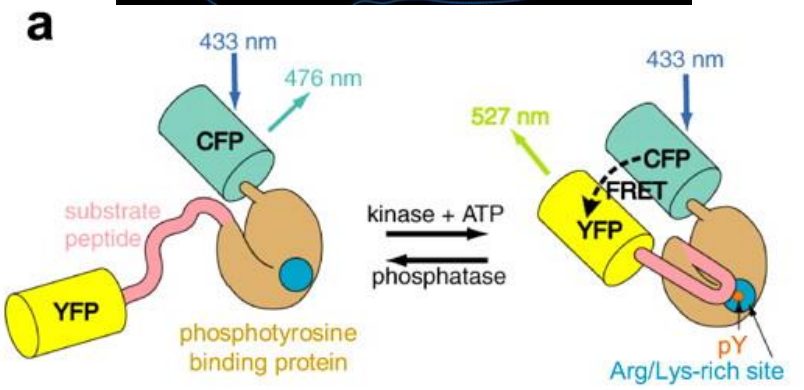


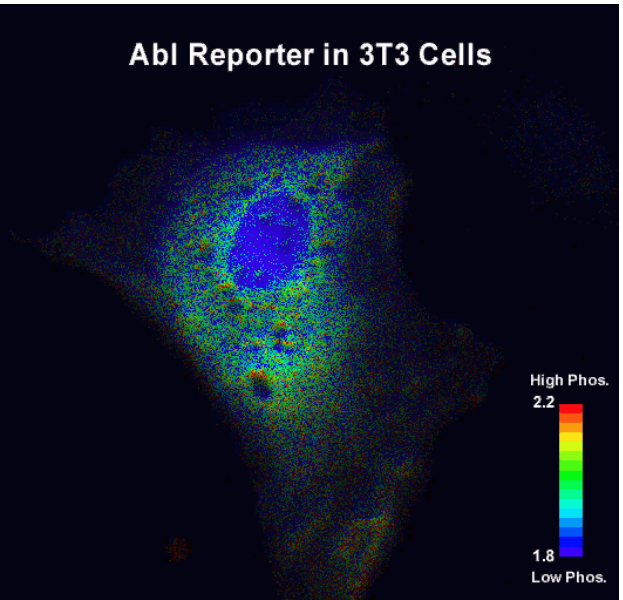
Photoactivation/Photoswitching
- PA-GFP, Dronpa, Eos

Reporting on environment
- Ca^{2+}, phosphorylation, cAMP, cGMP, pH, neurotransmitters, voltage, cell cycle, redox

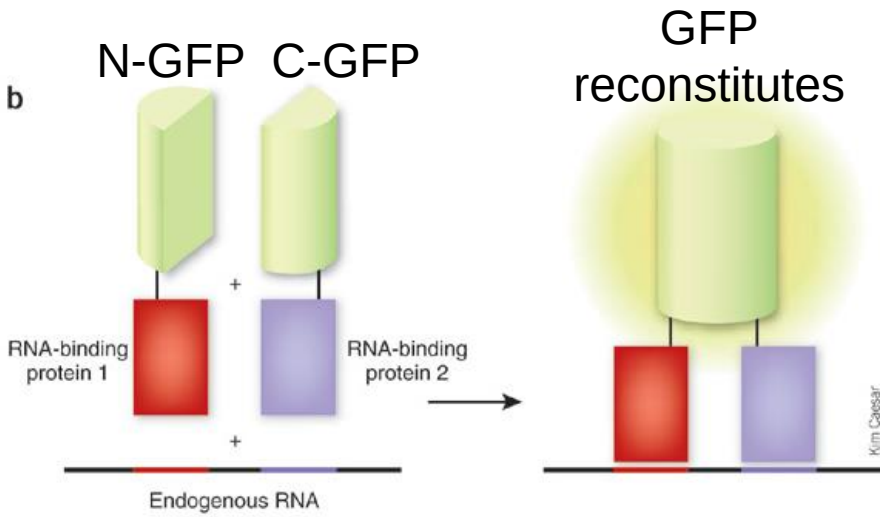
Reporting on protein-protein interaction
- CFP/YFP FRET, split fluorescent proteins

Modifying environment
- Singlet oxygen generation, Channelrhodopsin





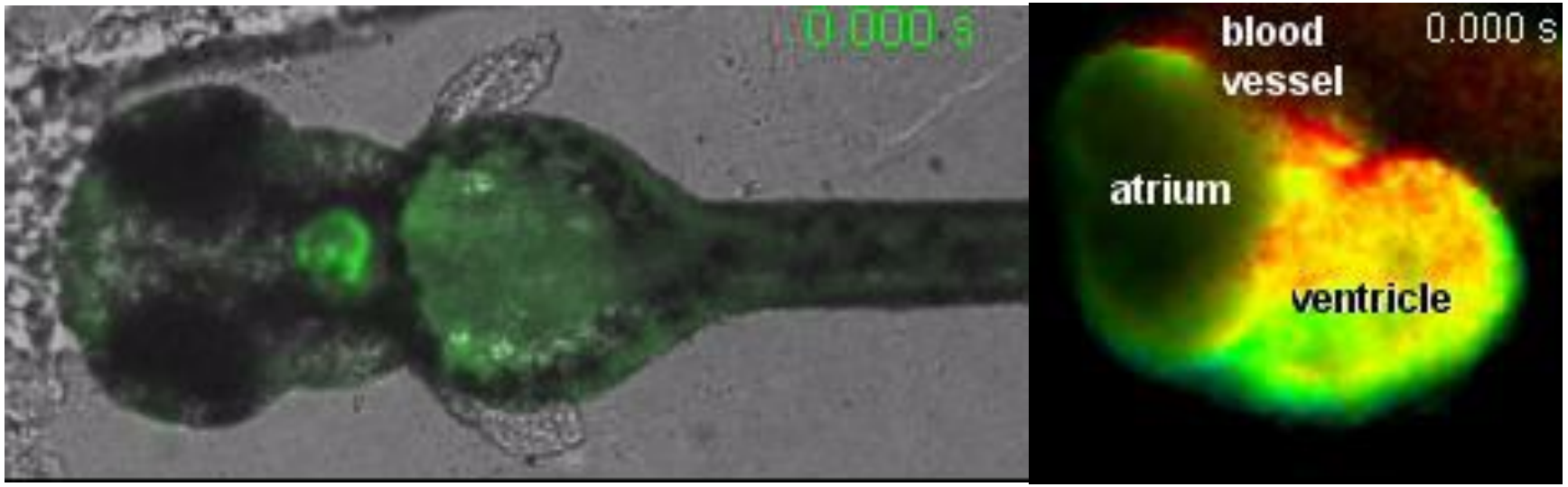
<u>Targeting advantage</u> to defined compartment, cell-type, developmental stage



Sensing voltage with fluorescent protein

Mermaid FRET voltage-sensor
by FP fusion to voltage-sensing phosphatase

Expressed in zebrafish heart
Non-invasive testing of mutant phenotypes
and drug cardiotoxicity.
Tsutsui, Miyawaki J Physiol 2010



FRET sensor ratio crucial

best is YC2.60 cameleon: 600%,
if <20% then lost in cellular noise

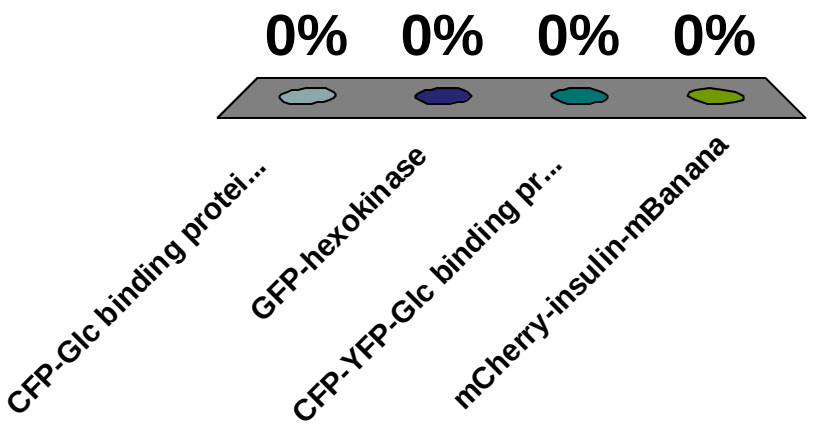
Which of these constructs would be best as a FRET sensor of glucose levels?

A. CFP-Glc binding protein-YFP

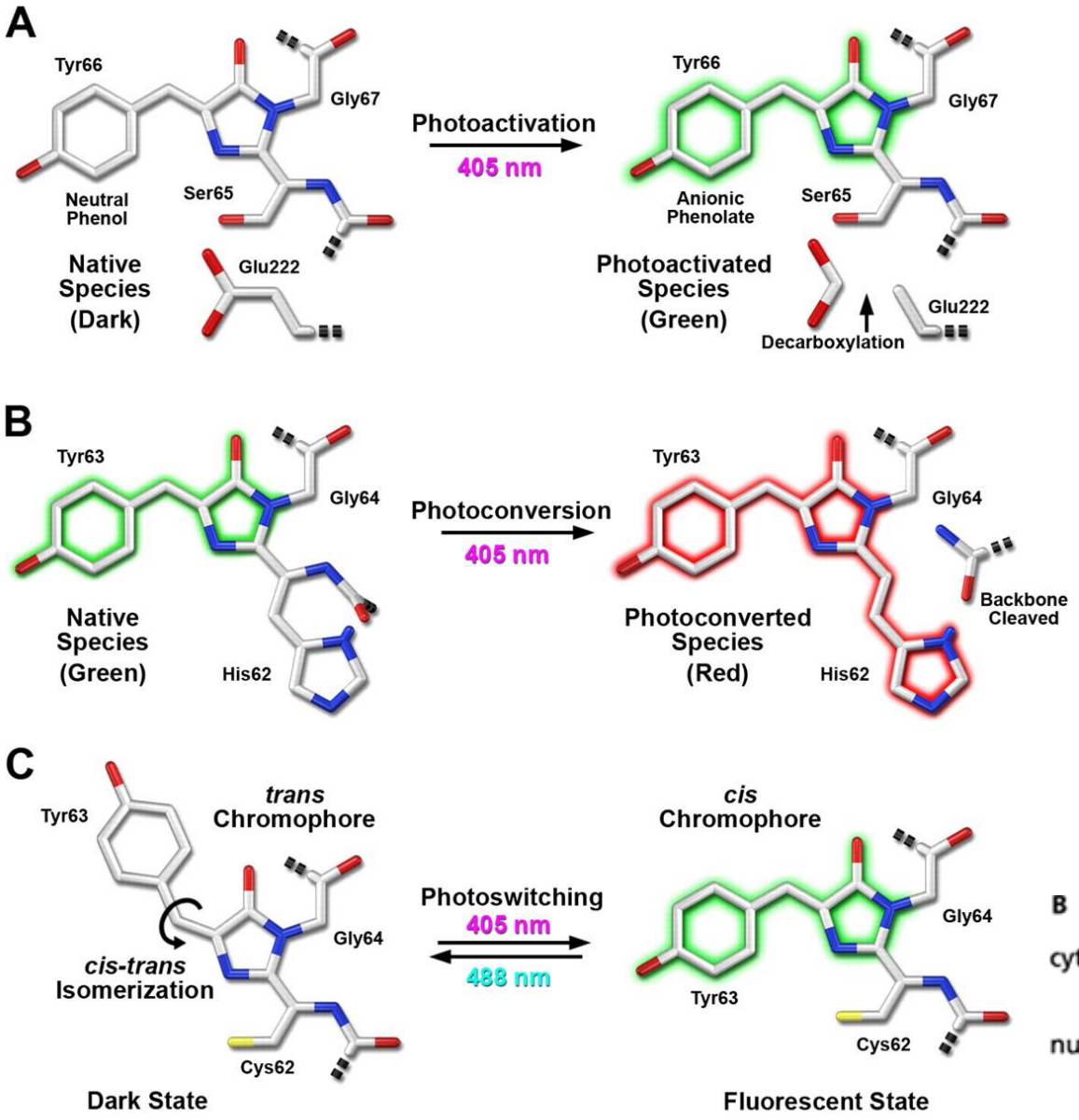
B. GFP-hexokinase

C. CFP-YFP-Glc binding protein

D. mCherry-insulin-mBanana



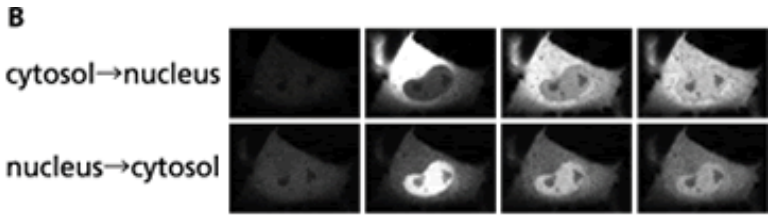
Chromophores in switching



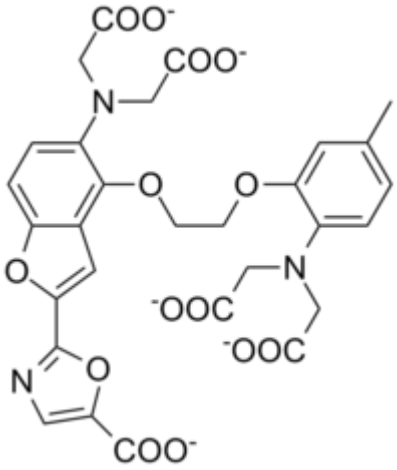
PA-GFP

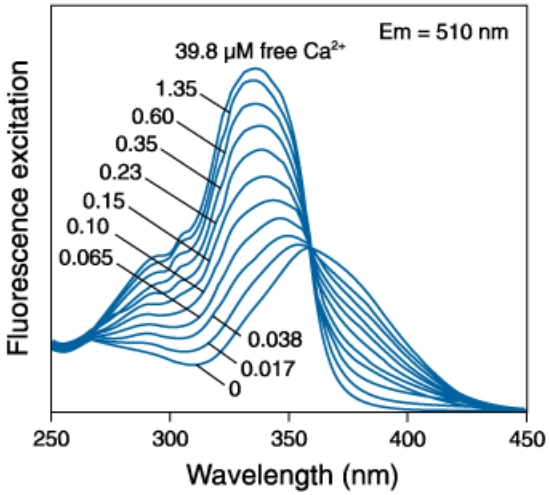
Dendra, Eos

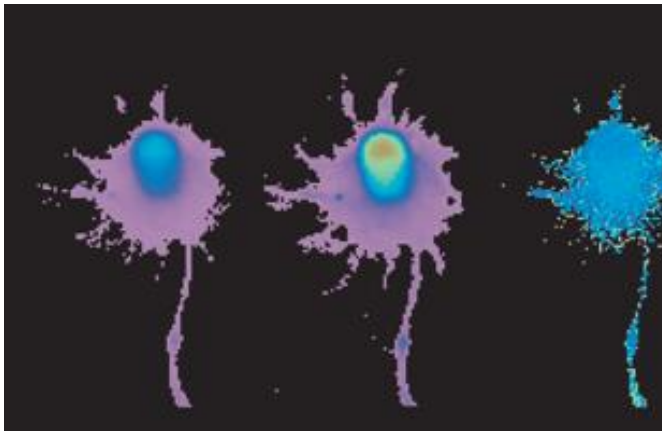
Dronpa



Small molecule fluorescent sensors







Fura-2 sensing calcium

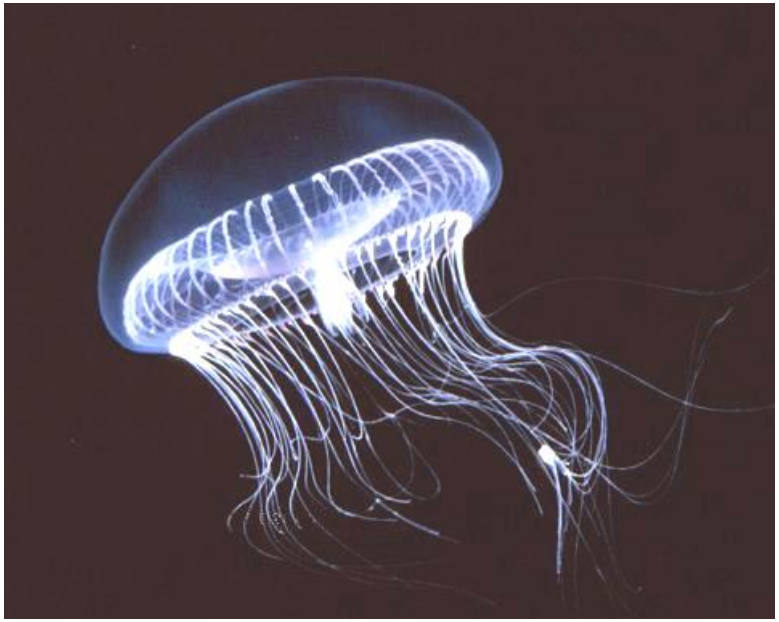
Metal ions: calcium, magnesium, zinc, sodium, potassium, chloride, mercury

pH (also dyes to conjugate to proteins, CyPher from GE, SNARF from Invitrogen)

Reactive oxygen species, nitric oxide

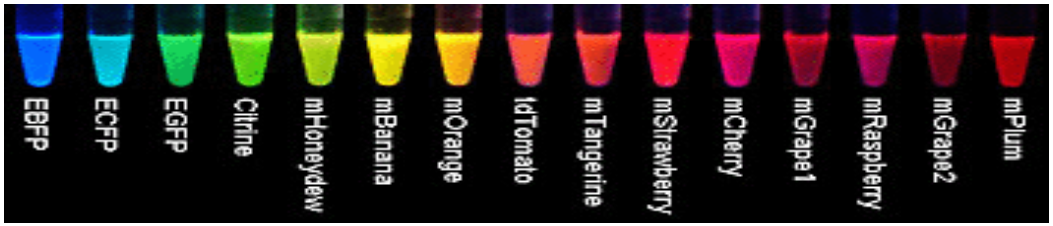
Transmembrane potential

How good is a fluorescent protein?



A. victoria GFP is good for jellyfish,
but not great for cell biologists!

How good is a fluorescent protein?



A. victoria GFP is terrible!

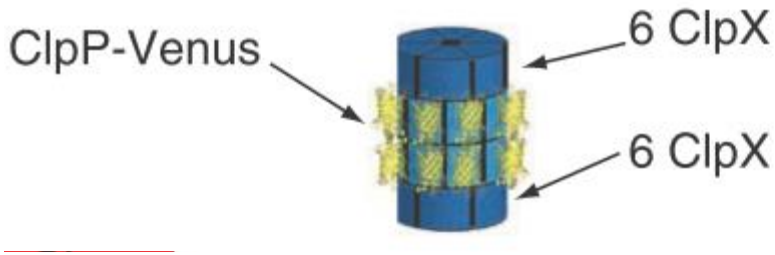
EGFP is OK, but there are now better...

1. Excitation and emission λ — good match to filters on your microscope
   look at other fluorophores at same time
2. Bright — ε x QY Clover, YPet 2.5 x EGFP
   mRuby2 3x mCherry
3. Stable to photobleaching — EBFP bad, mCherry and YPet good
4. Non-toxic — attach on right part of your protein
   all make H_2O_2, FPs can transfer electrons
5. Environment-insensitive — especially to pH, chloride
   CyPet does not fold at 37°C, all need O_2
   Photoactivatable FP did not work in ER
6. Little non-specific binding — fully monomeric, A206K non-dimerising
7. Fast Maturation — Venus 2 min. Red FPs can start off green
   half-time ~15 min mCherry, 100 min TagRFP

You MUST worry about FP multimerization!

Tag multimerizing protein with FP and sometimes see foci-

are these real or caused by the tag?



With hexameric barrel involved in *E. coli* protein degradation,

many commonly used FPs induce artifactual foci

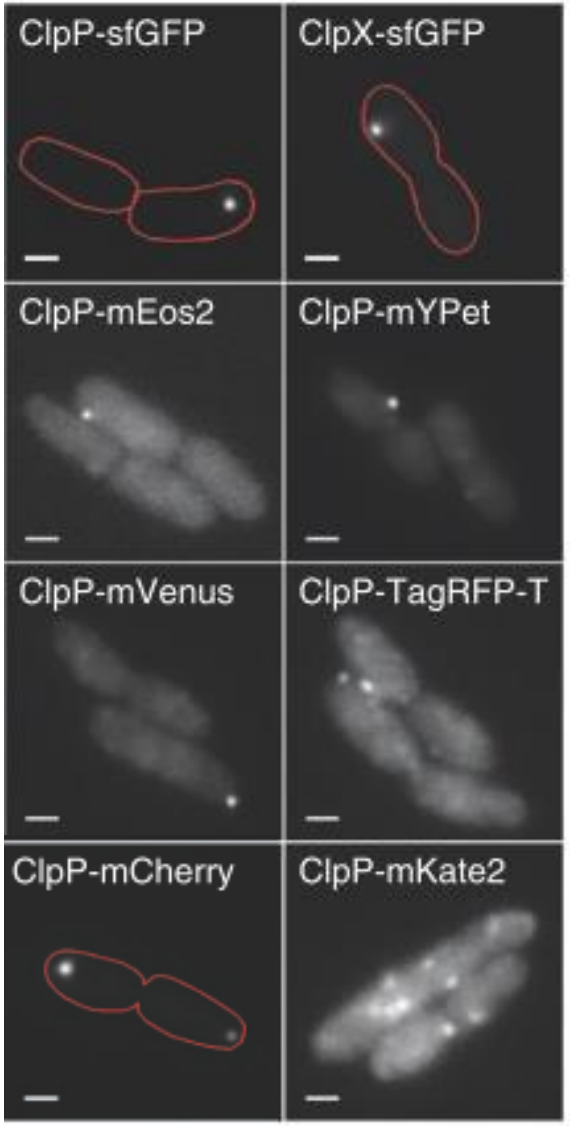
(no cluster with Ab or SNAP-Tag)

as well as affecting daughter cell inheritance of proteolysis ability

mCherry, sfGFP, mYPet poor!

mGFPmut3, Dronpa OK

D. Landgraf et al. Nature Meth 2012



Problems with GFP in cells

- GFP with light can donate electrons to different acceptors
  (FMN, FAD, NAD^+, cyt. c)
  GFP reddens after transfer: photobleaching and phototoxicity
  use DMEM lacking e^- acceptors
(riboflavin or all vitamins) for less bleaching
(HEK 293T happy for 1 week)
effect for EGFP and PA-GFP, not RFPs
Lukyanov Nat Meth 2009

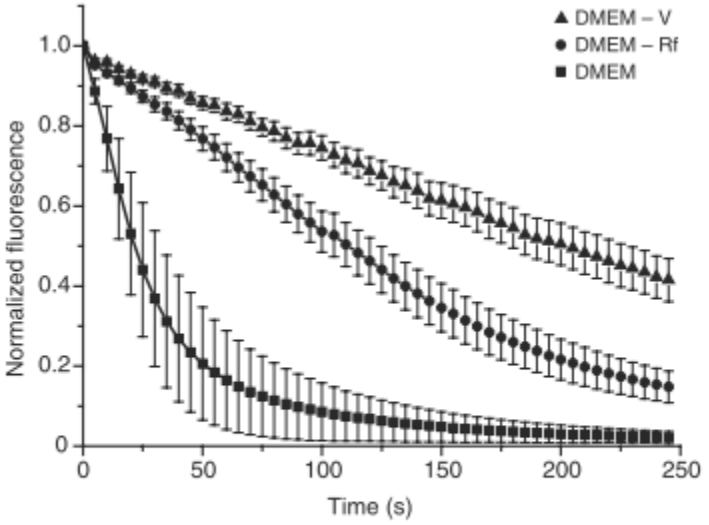
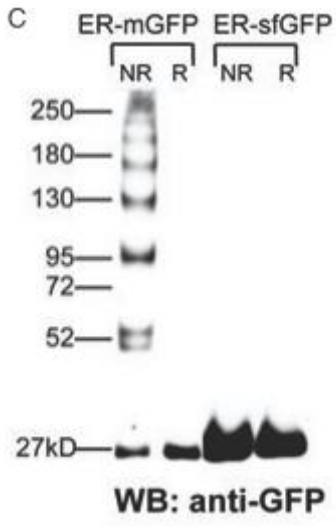


Figure 1 | Influence of cell medium on fluorescent protein photostability. Normalized bleaching curves for EGFP in live HEK293T cells maintained in DMEM, DMEM – Rf or DMEM – V. Error bars, s.d. (n = 20 cells).

- EGFP not good in secretory pathway
  mixed disulfide oligomers in ER and non-fluorescent in *E. coli* periplasm
  (superfolder GFP behaves fine)
  Erik Snapp, Traffic 2011



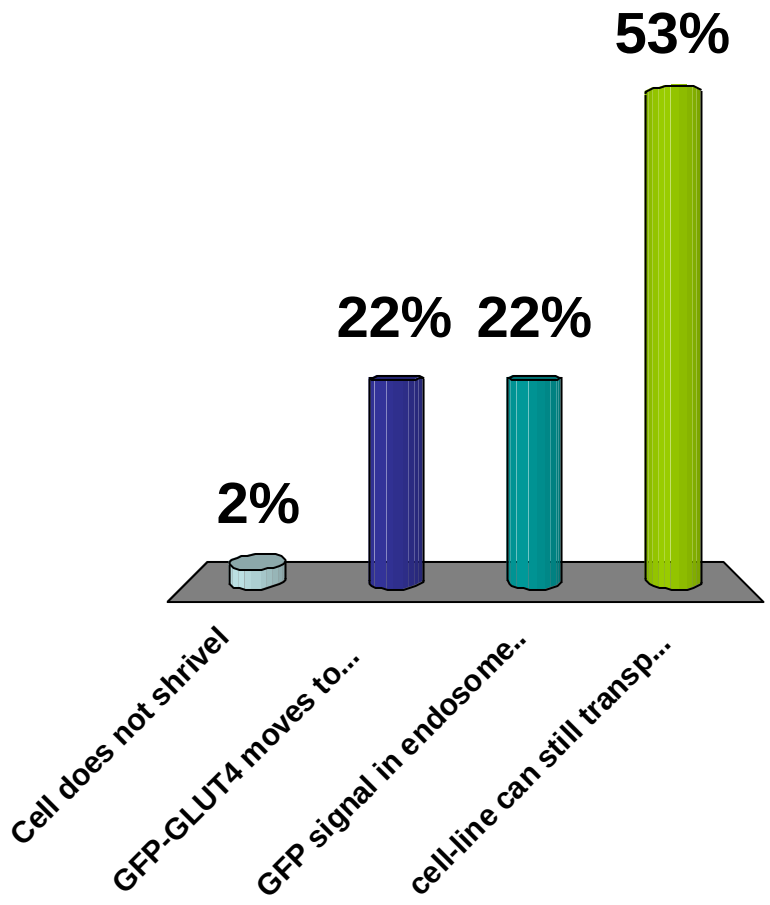
I have transfected GFP-tagged GLUT4 into a mammalian cell. What is the best control that I have not interfered with the system?

A. Cell does not shrivel

B. GFP-GLUT4 moves to PM with insulin

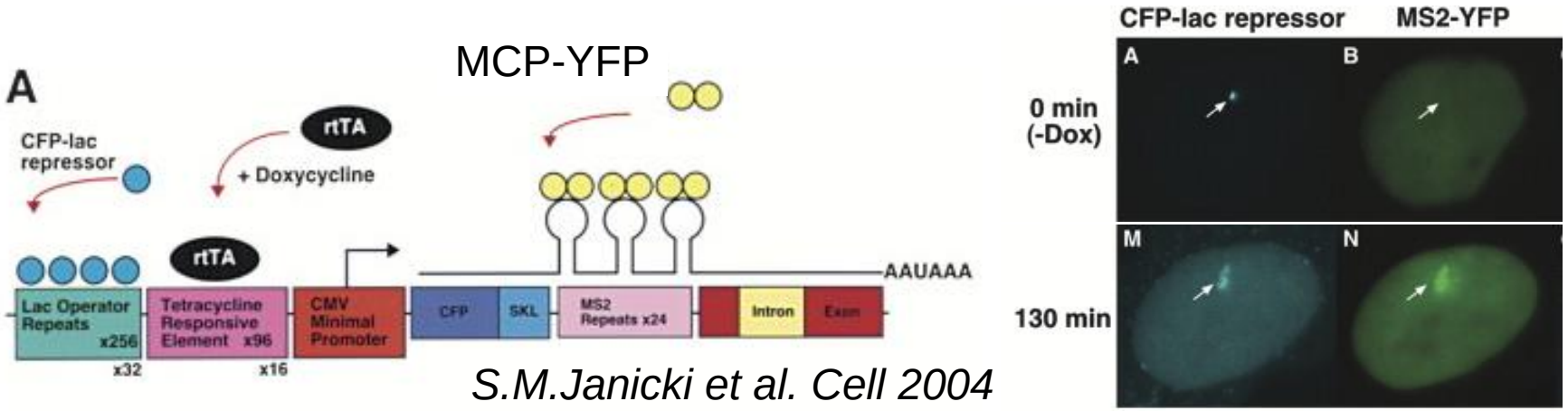
C. GFP signal in endosomes, PM

D. cell-line can still transport glucose



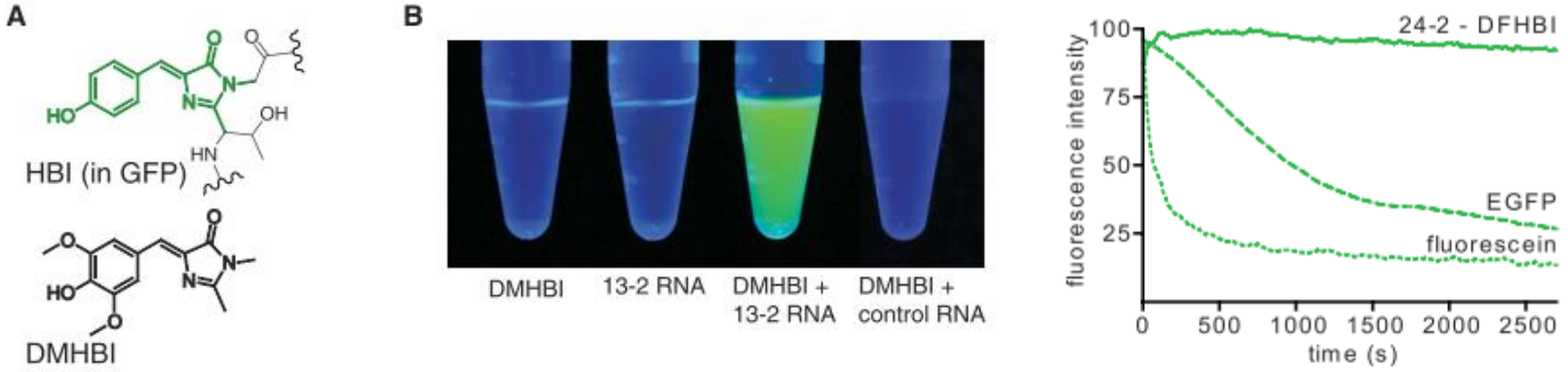
Fluorescent RNA imaging

See single mRNA: MS2 mRNA stem-loops bound by MCP-YFP
See product of translation: mRNA encodes CFP-SKL which goes to peroxisomes

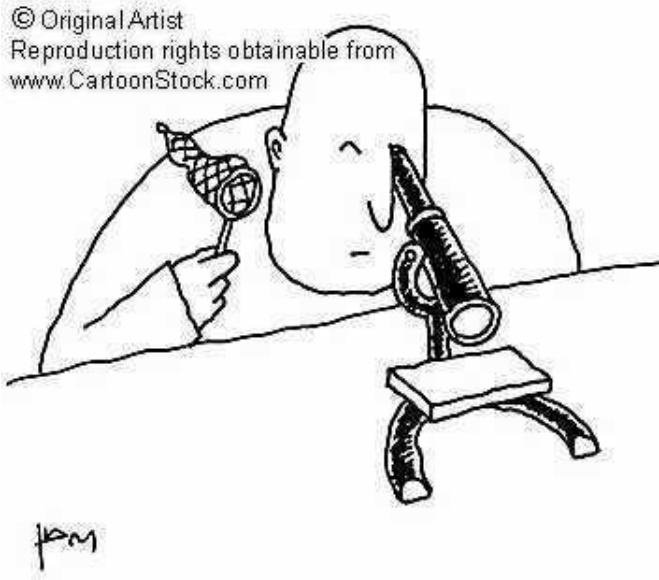


S.M.Janicki et al. Cell 2004

Spinach RNA 60 nt aptamer binds cell-permeable fluorogenic dye
Photostable. Used to label 5S RNA in HEK cells. *Samie Jaffrey Science 2011*



Overview



1. What kind of structures are fluorescent

2. How to make and target fluorescent probes

3. Fluorescent probes for cellular structure and function

4. Using light to control cells

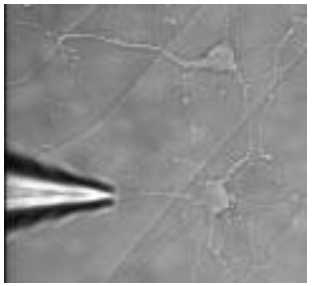
Why use light to control biology?

Light control allows extreme temporal and spatial control.

Temporal control

genes< chemicals < light

min-hr s-min µs-s



Spatial control

chemicals / genes < light

one or many cells 1 µm part of cell

(note micropipettes for precise small molecule delivery)

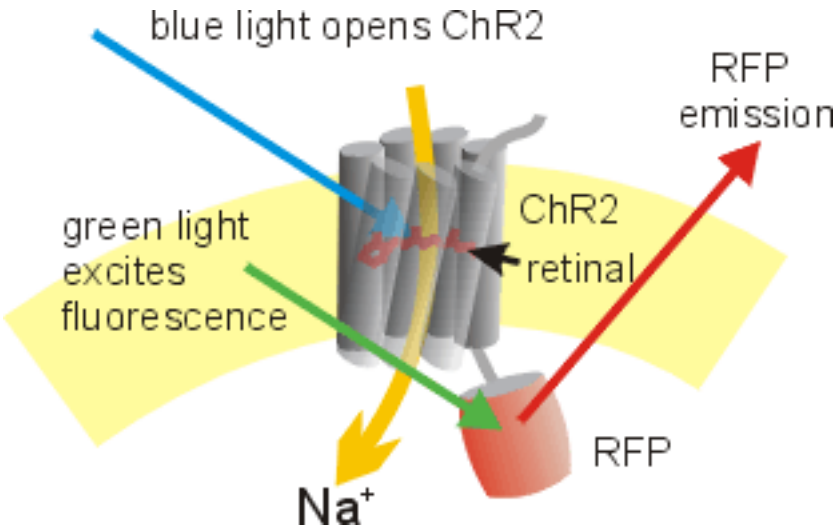
(often combine chemical/light control or gene/light control)
optogenetics/chemogenetics

Limitations of light? $$$$$

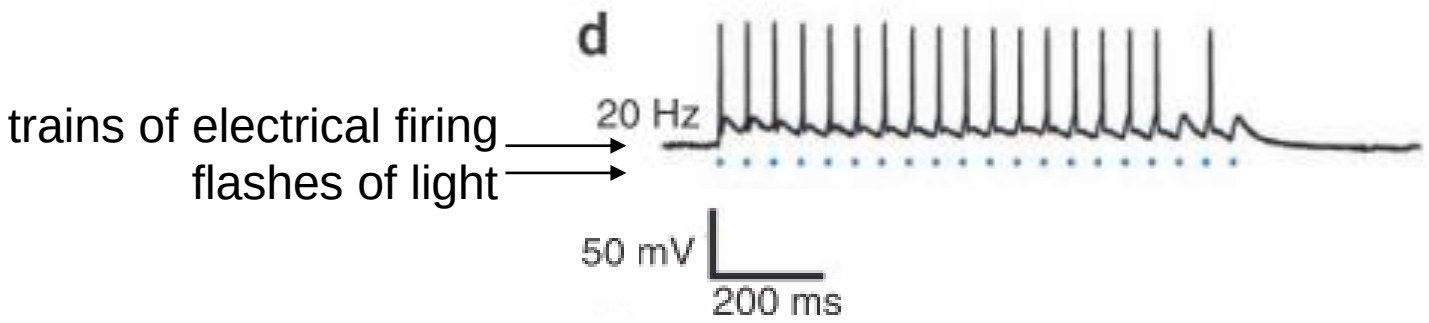
and usually data on one cell at a time

Controlling biology with light: light-gated ion channels

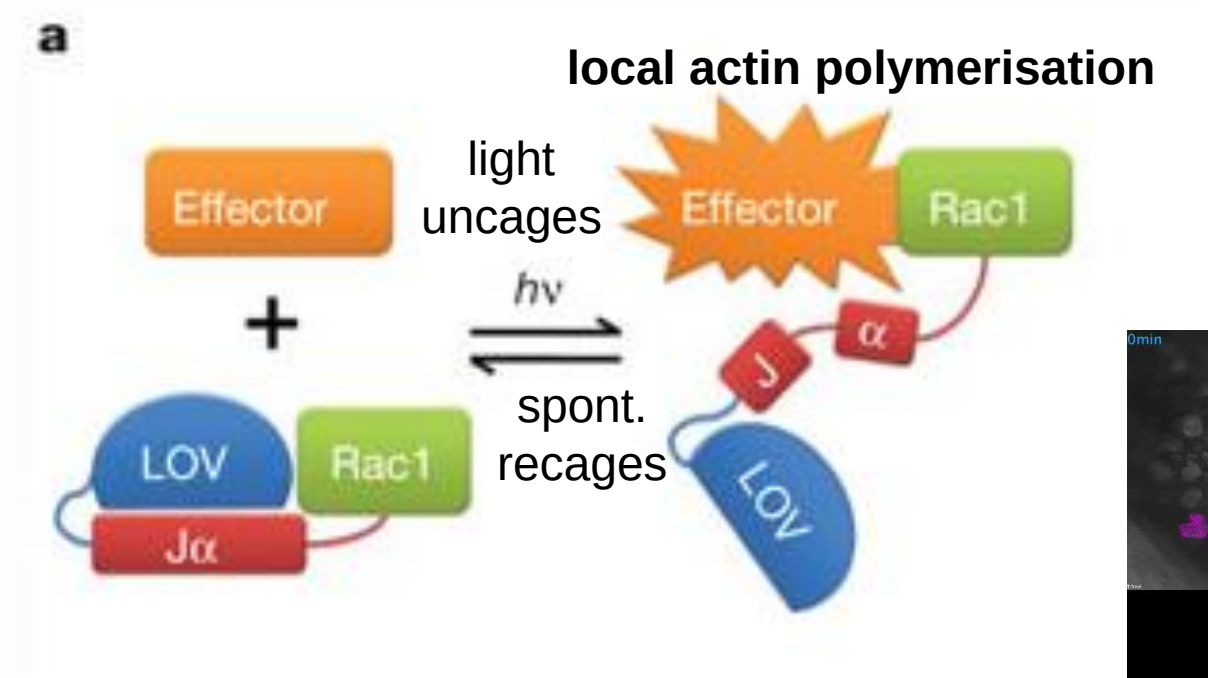
Channelrhodopsin from an alga, like rhodopsin, undergoes retinal isomerisation in response to light, and changes conformation, but opens a Na^+ channel. This allows light to control membrane voltage and trigger neuron firing.

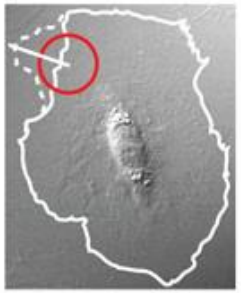


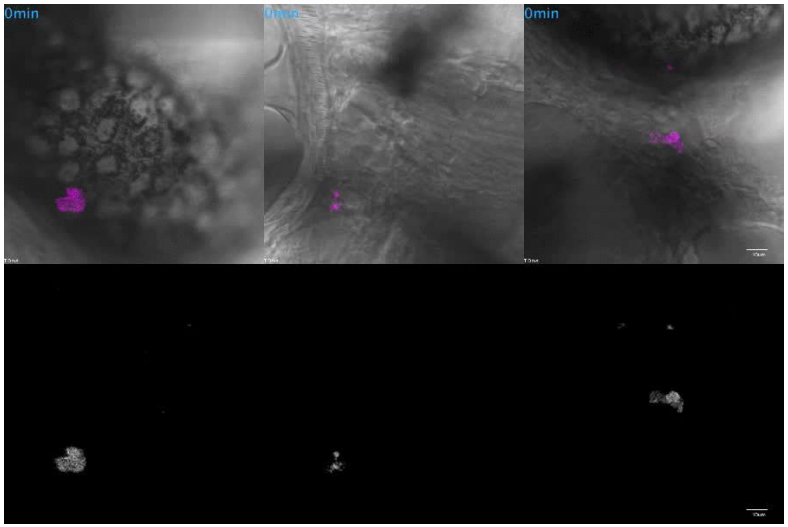
to understand neuronal firing patterns
to control secretion in diabetes
potentially in fixing neural diseases?
e.g. damping down overactivity in epilepsy



Genetically-encoded photoactivation







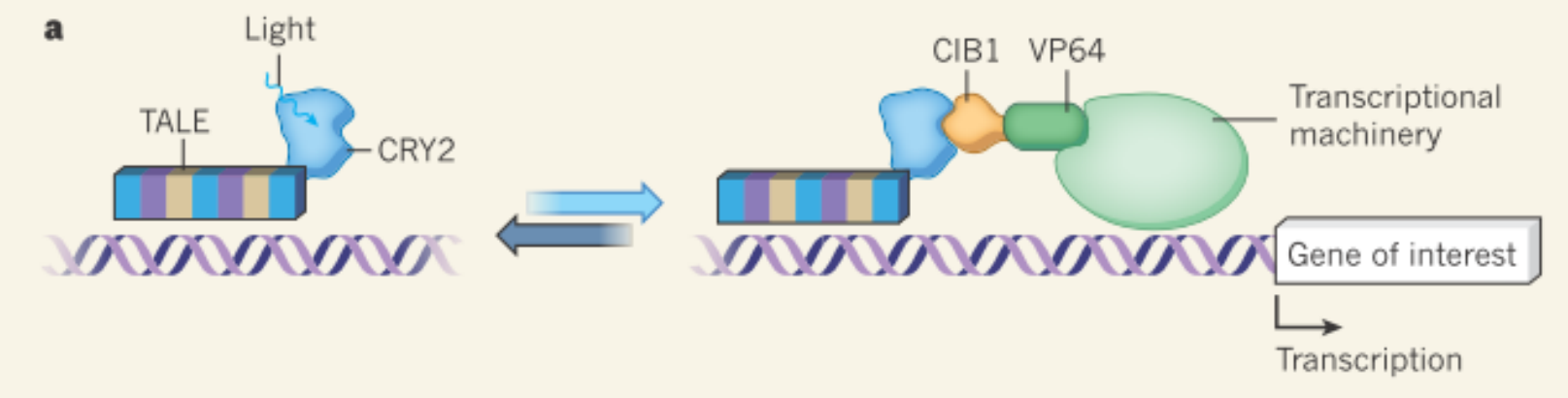
1. Constitutively active Rac mutant
2. Optimise LOV-Rac junction,
3. knockout GTP hydrolysis and GAP/GNDI/GEF interactions

K_d for PAK 2 μM in dark, 200nM in light 10-fold ratio

Interaction of Rac with PAK stimulates cell protrusion and migration.

K.Hahn et al. Nature Sept. 2009

Photoactivation of transcription



1. Transcription-activator-like effector (TALE) for targeting arbitrary DNA sequence
2. 10-fold induction
3. Induction in minutes in animals

Modular design (can also recruit nuclease or repressive domain)

A. Moglich et al. Nature Aug 2013

Conclusions

Choosing the right dye or fluorescent protein can make a big difference for:

- sensitivity
- signal stability
- modification to molecule/cell function
  - by size or multimerization

Fluorescent probes allow more than just following location:

- reporting cellular events
- uncaging biomolecule function
- controlling interactions and ion flux

References

Fluorescence probes
Molecular Probes Handbook, from Life Technologies.
Principles of Fluorescence Spectroscopy 2nd edition, by Joseph R. Lakowicz.

Protein modification
Bioconjugate Techniques, 2nd Edition
by Greg T. Hermanson.
Chemical labeling strategies for cell biology, Marks KM, Nolan GP. Nat Methods. 2006 Aug;3(8):591-6.

Fluorescent proteins
(i) See table at
http://nic.ucsf.edu/FPvisualization/

(ii) as sensors: Designs and applications of fluorescent protein-based biosensors.
Ibraheem A, Campbell RE.
Curr Opin Chem Biol 2010;14:30-6