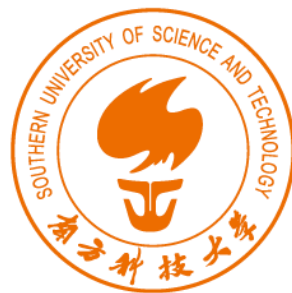


A novel biophysical model for radical pair mechanism in birds' magnetoreception



2019年5月21日

骆锦威 11410163



Contents

- Background
- Three Possible Mechanisms
- OCC Model
- Simulations
- Predictions and Discussion

Background

First discovery in magnetoreception

Magnets Interfere with Pigeon Homing

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Communicated by Donald R. Griffin, October 8, 1970

ABSTRACT Magnets glued to the backs of experienced pigeons often resulted in disorientation when the birds were released from distances of 17-31 miles (27-50 km) under total overcast, whereas no such disorientation occurred during similar releases under clear skies. The magnets did, however, often cause disorientation when first-flight birds were released under sun, and there was some indication of disturbance to experienced pigeons released under sun at longer distances.

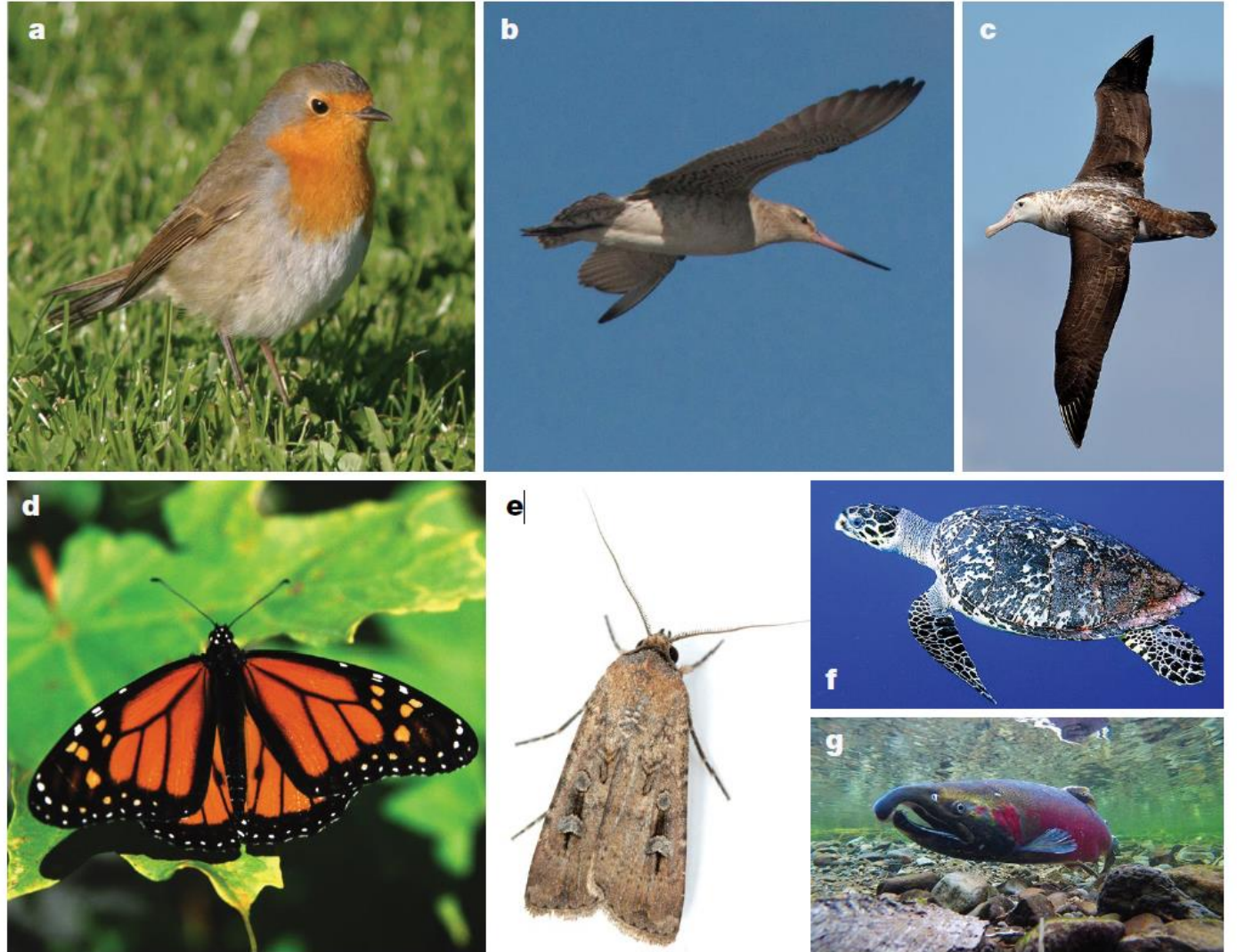
Modified after Ref.[1]

Magnetic compass is ubiquitous across species

Vertebrates: birds, fishes, amphibians, reptiles, mammals.

Invertebrates: mollusks, crustaceans, insects.

- *Erithacus rubecula*
- *Limosa lapponica*
- *Diomedea exulans*
- *Danaus plexippus*
- *Agrotis infusa*
- *Eretmochelys imbricate*
- *Oncorhynchus kisutch*



Modified after Ref.[2]

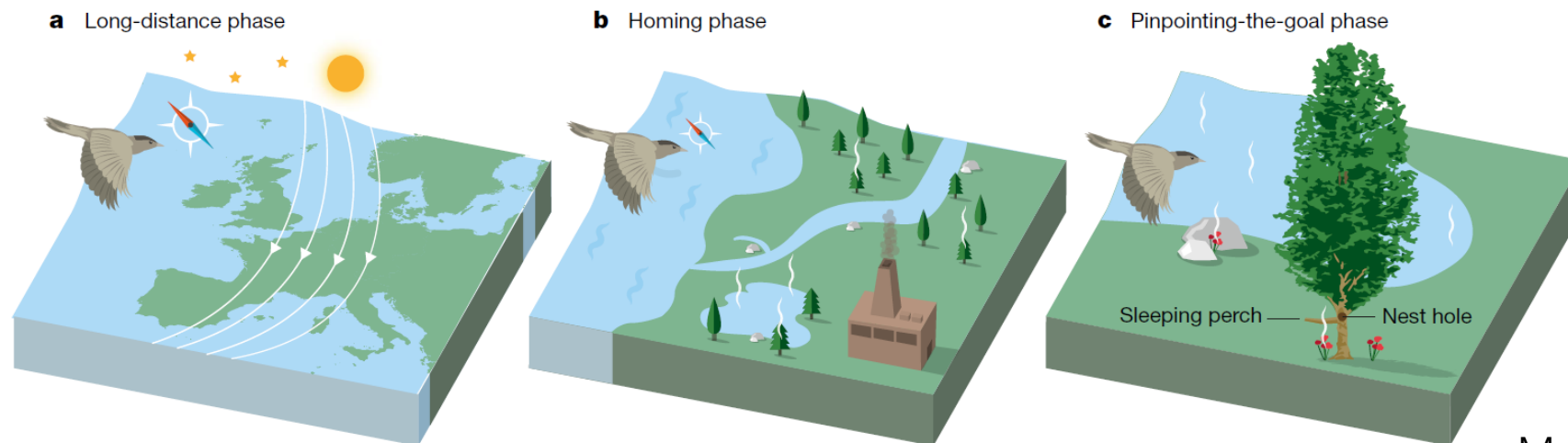
Multisensory cues for bird navigation

- **Long-distance navigation – Migration maps**

Magnetic compass, sun compass, star compass

- **Homing and pinpointing – Local orientation**

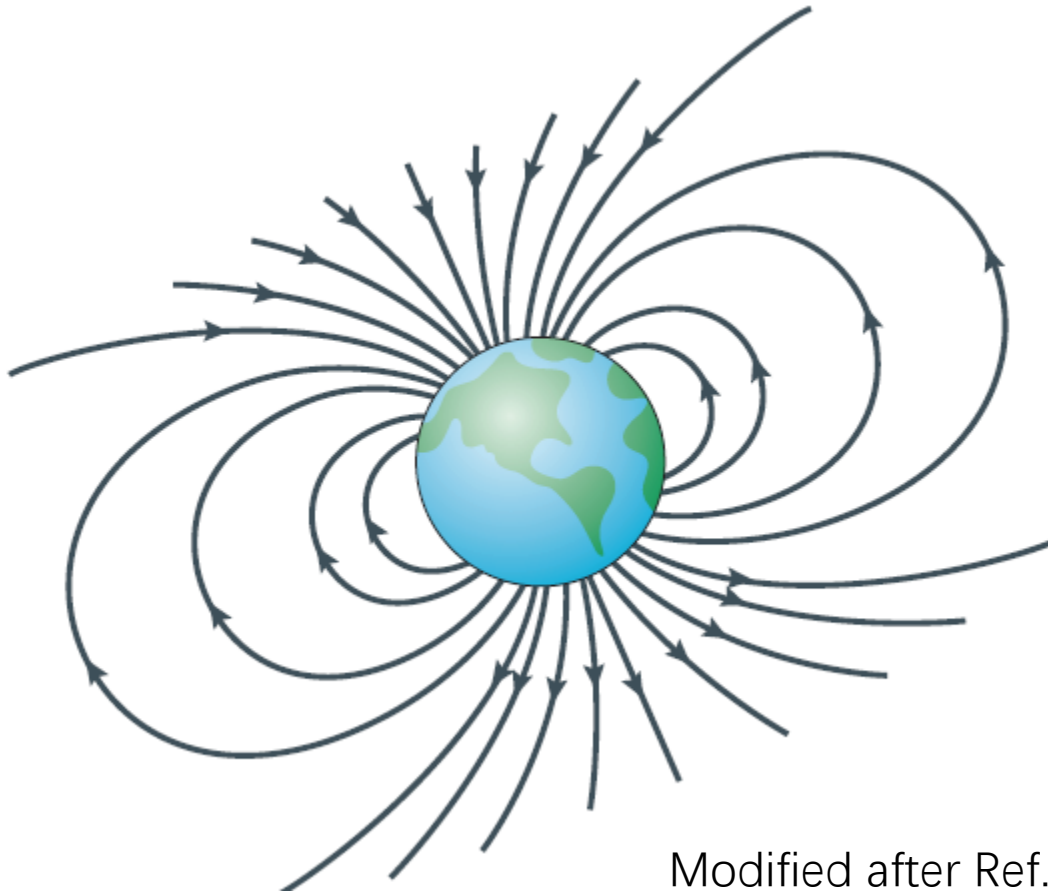
Odors, borders, coast lines, sound, landmarks such as a tree, a small hill, a specific coral



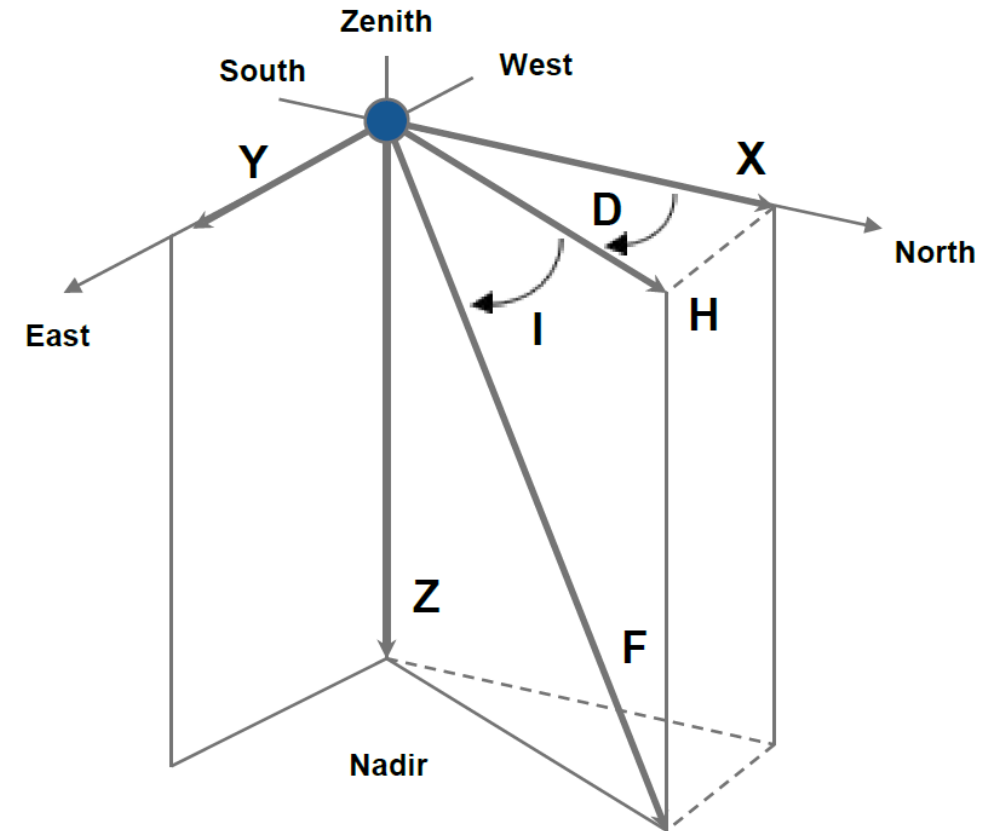
Modified after Ref.[2]

Earth magnetic fields

- Field strength: 25~60uT



Modified after Ref.[3]



Magnetic field information

Inclination

Polarity/Declination

Intensity

Mechanisms

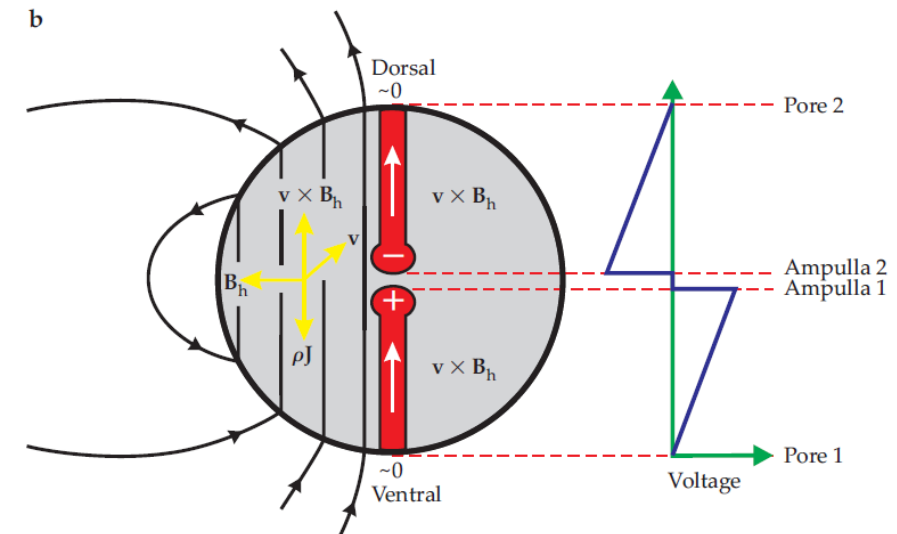
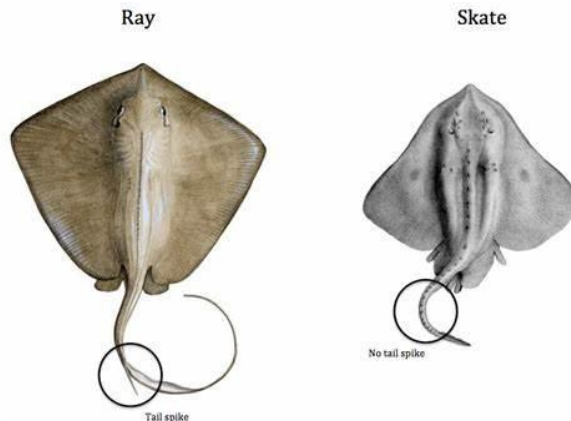
Three possible mechanisms

- Electromagnetic induction
- Magnetite based magnetoreception
- Radical based magnetoreception

Difficulties for verification

- No specific organ is found
- No unique behavioral phenotype
- Multisensory cues coupling
- Adaptation and acclimation
- Multi origins or single origin?

-

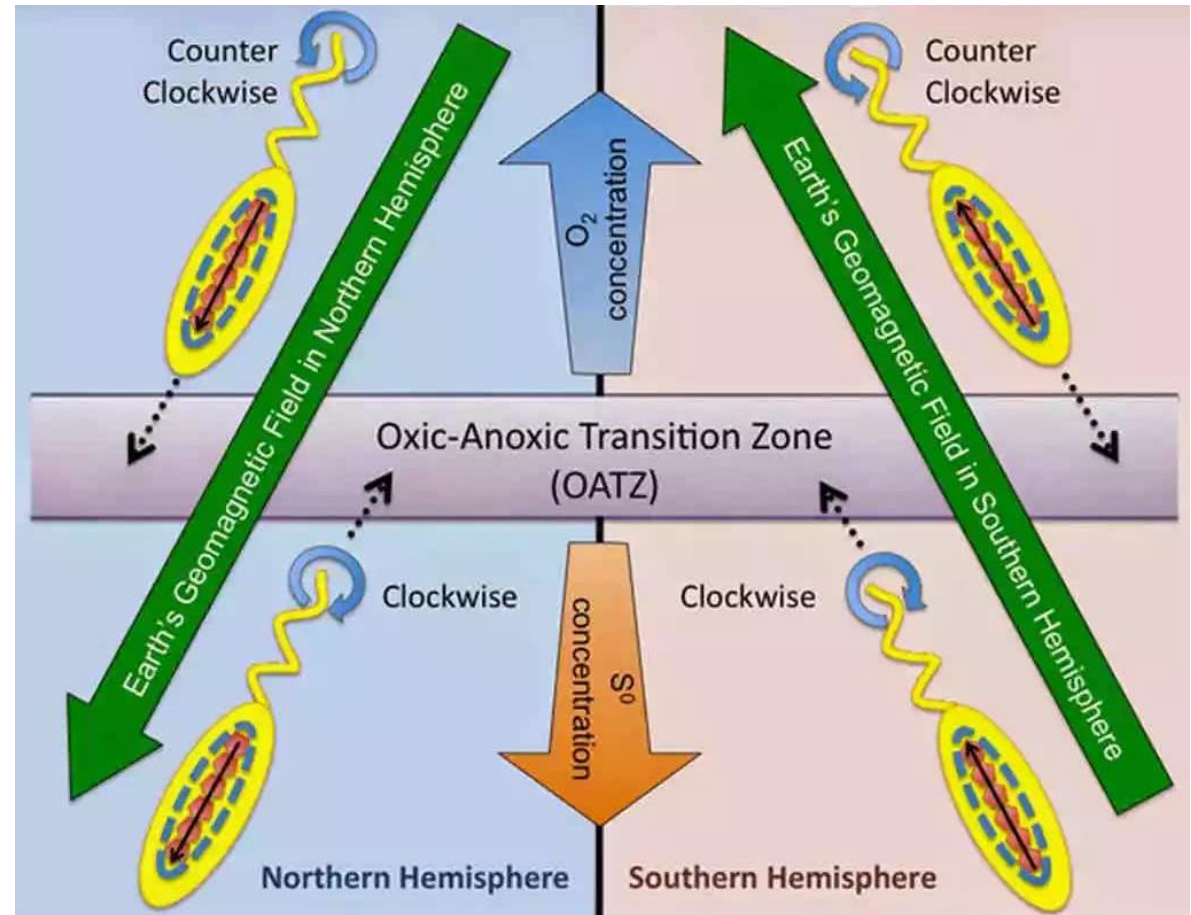
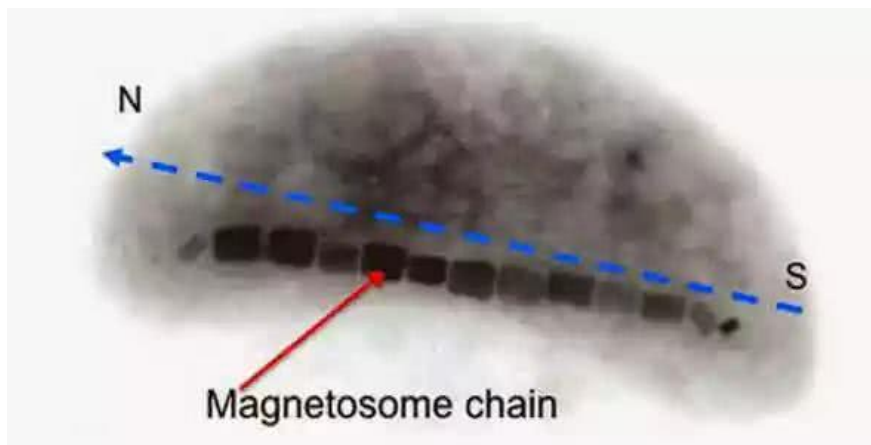


Modified after Ref.[4]

Magnetite based Magnetoreception

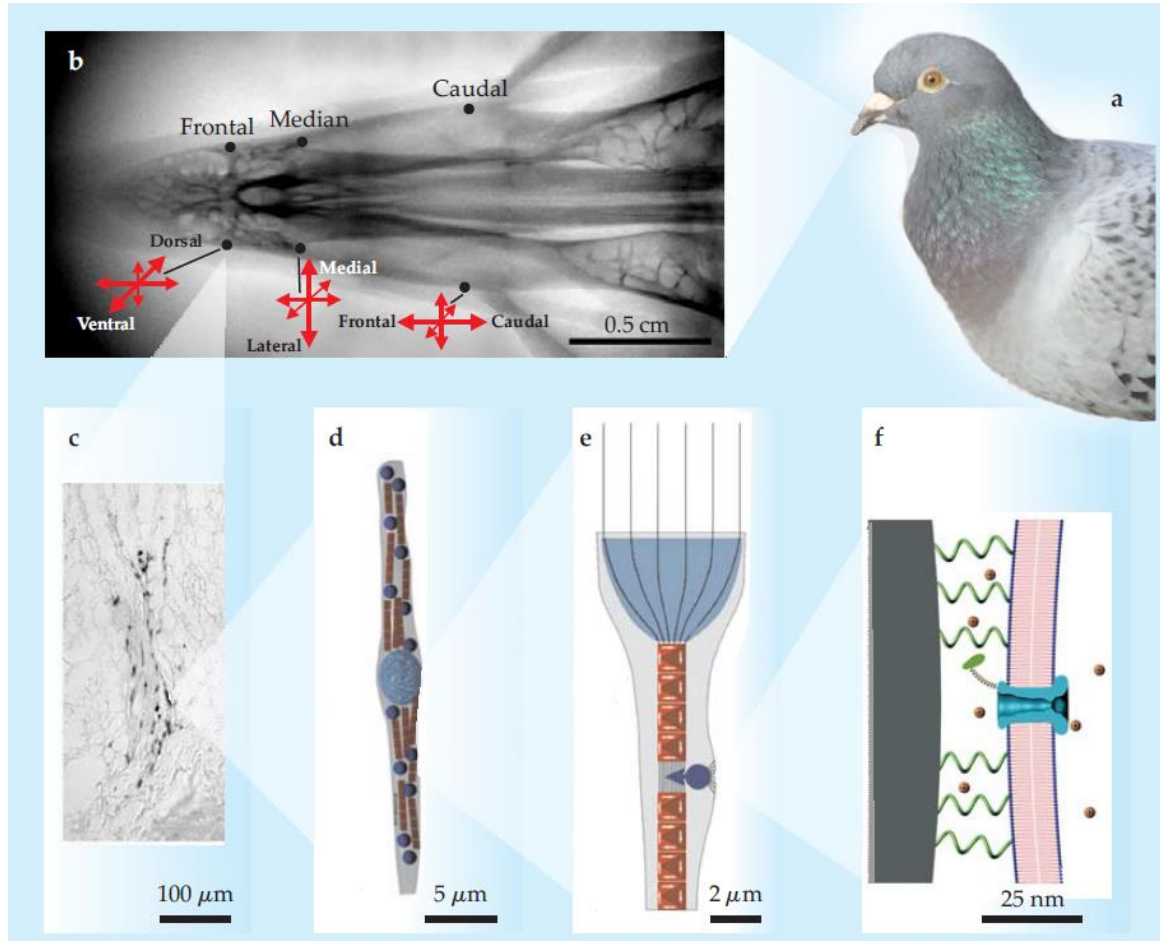
Magnetic force -> Physical rotation

- Magnetotactic bacteria
- A magnetosome chain:
- 15-20 Fe_3S_4 , Fe_3O_4 crystal
- 30-100nm for each crystal

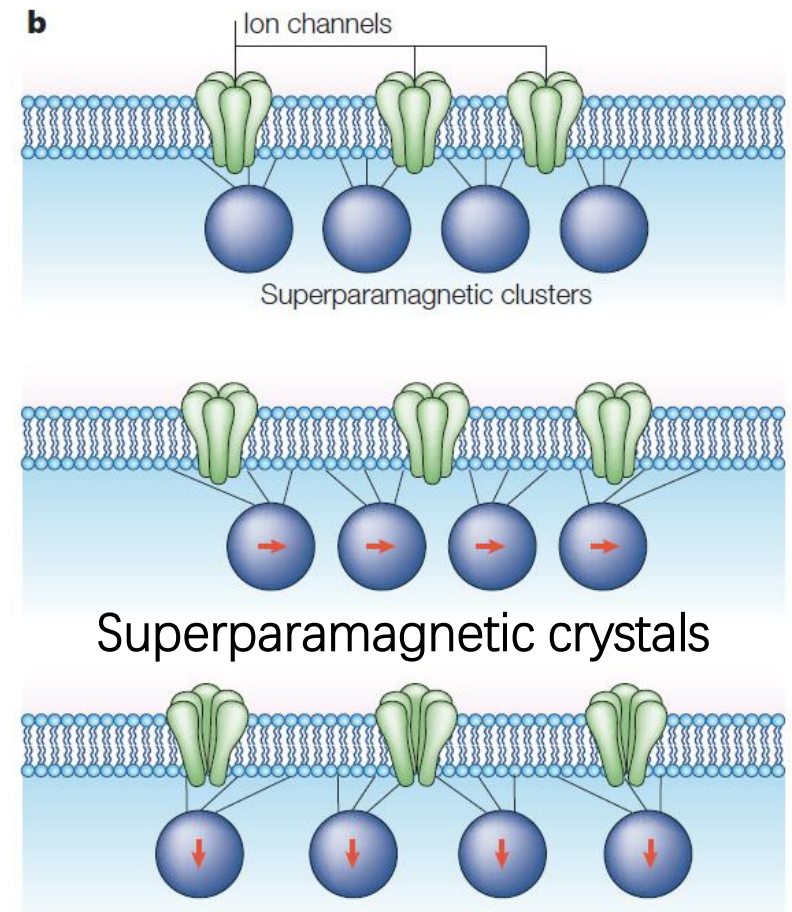


Magnetite based Magnetoreception

Magnetic information -> Pressure signal -> Electric signal



Modified after Ref.[4]

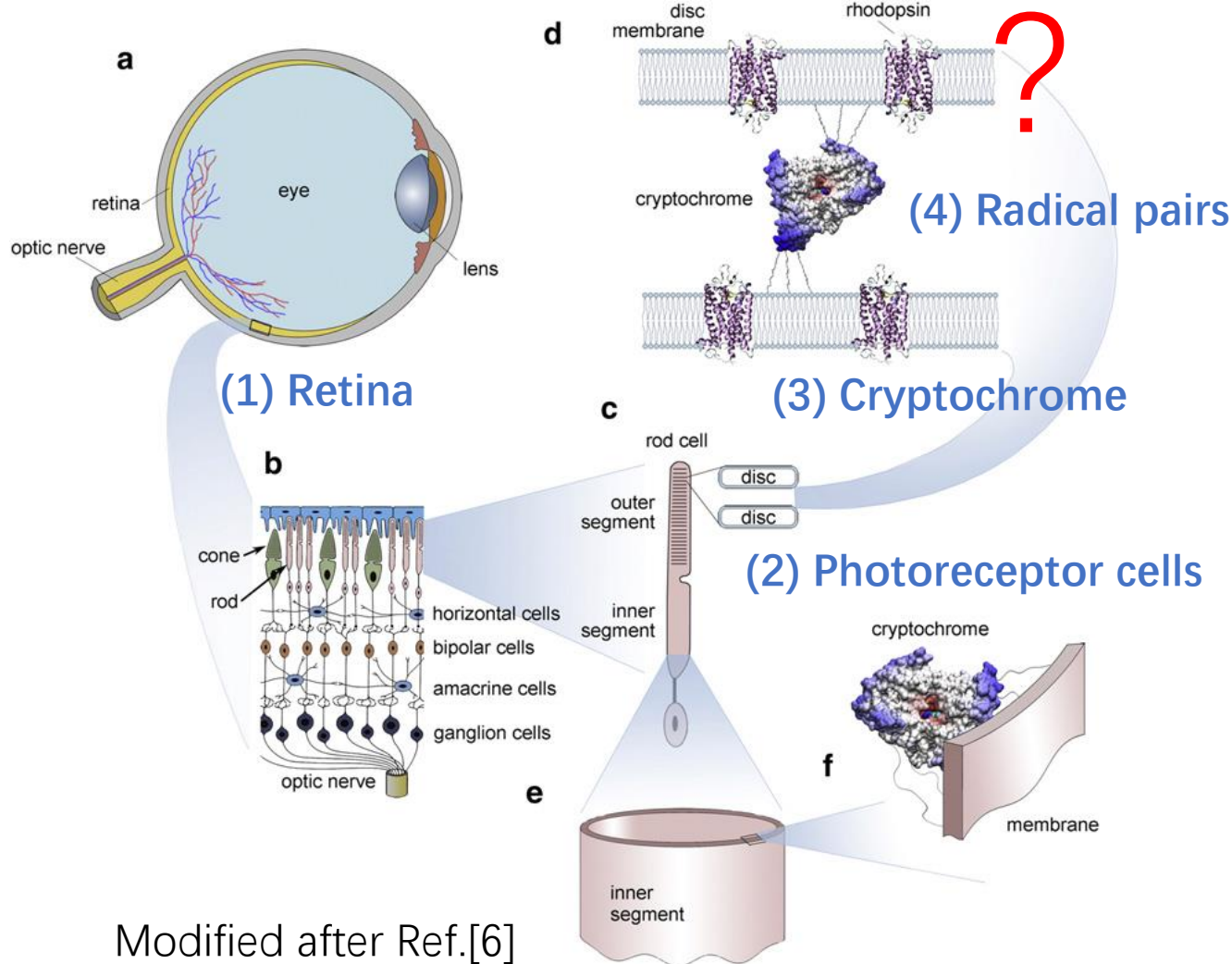


Modified after Ref.[3]

Radical based magnetoreception

Magnetic information -> Chemical signal -> Electric signal

(5) Visual pathway



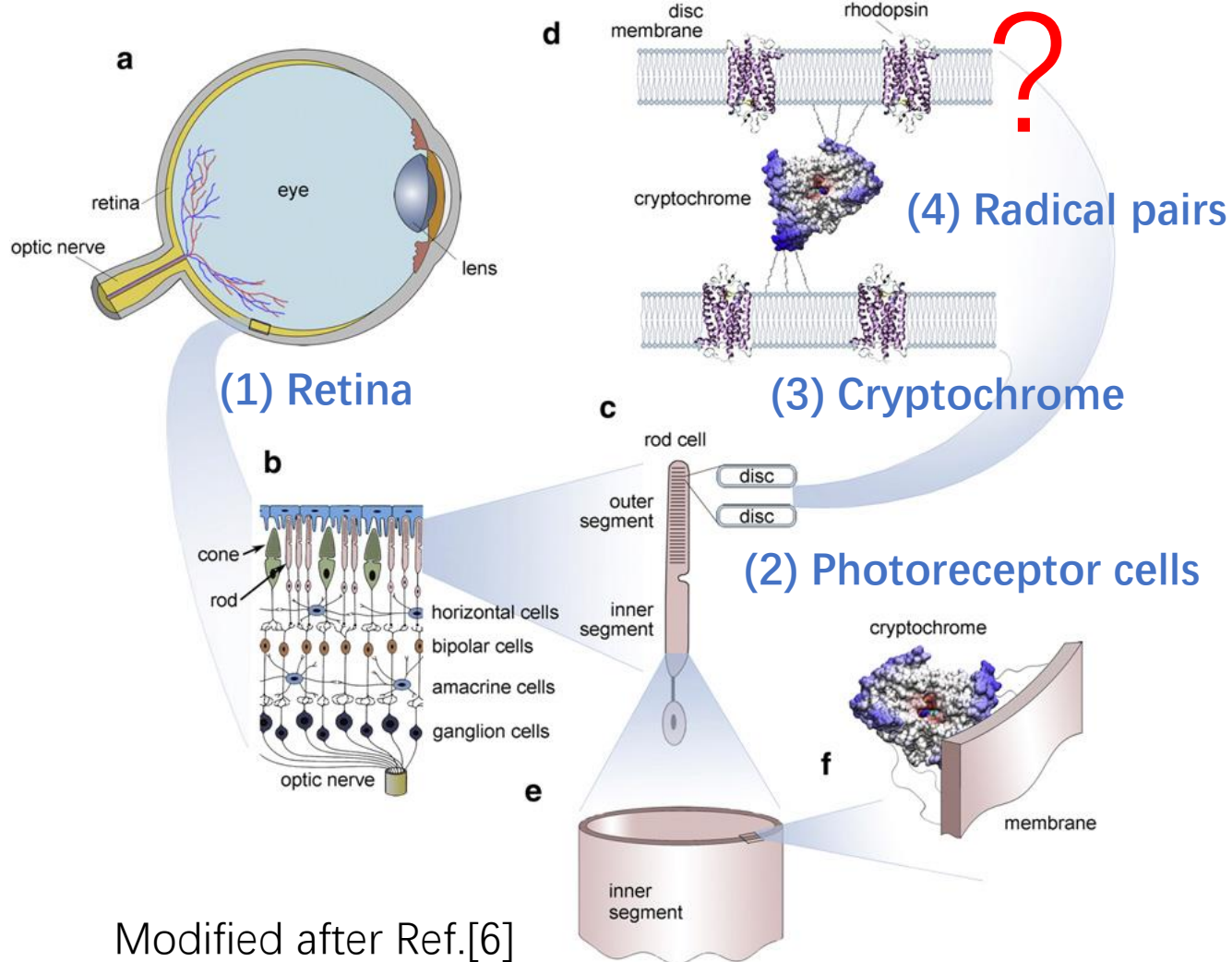
- 1) A **neural connection** between night vision and magnetic sensing in European robin.[12]
- 2) Different wavelength of incident light can **disrupt magnetic orientation** of European robin.[13,14]
- 3) The magnetic compass of European robin is a **inclination compass with $<5^\circ$ precision**, which can be explained by radical pair mechanism instead of any other hypothesis.[15,16]
- 4) Cryptochrome is the **only protein family** that can form a **light-induced radical pair**.

Modified after Ref.[6]

Radical based magnetoreception

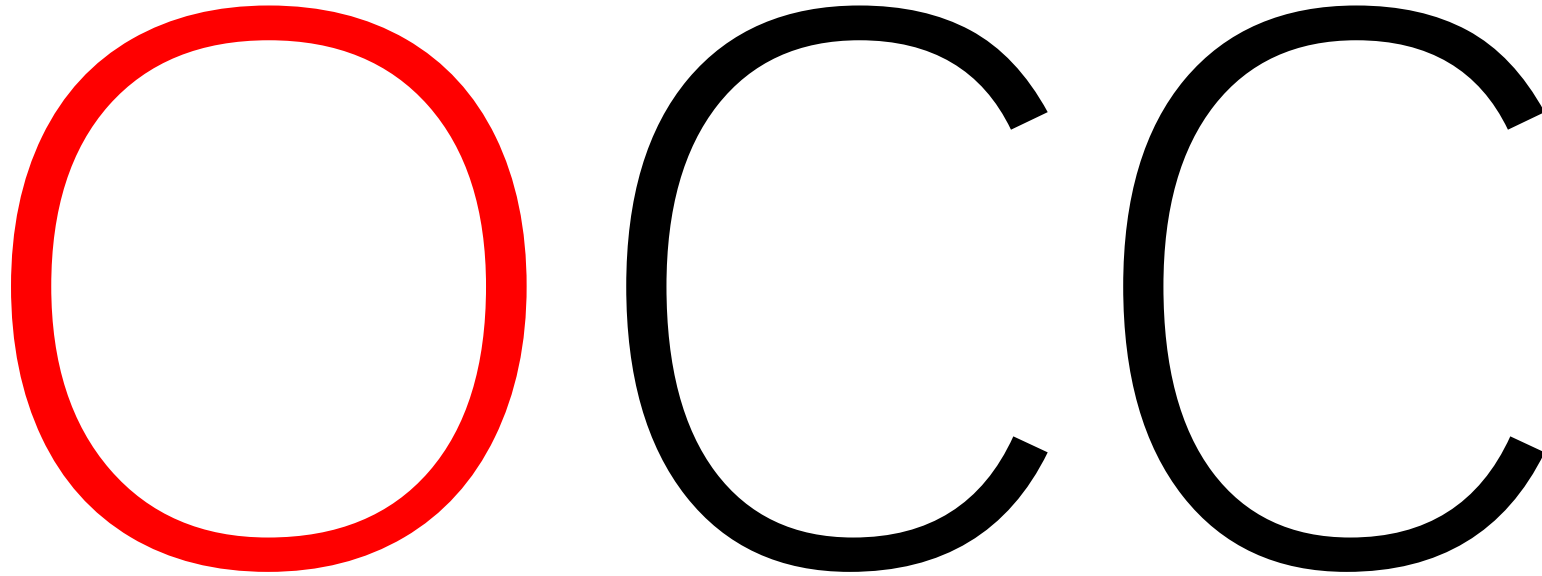
Magnetic information -> Chemical signal -> Electric signal

(5) Visual pathway



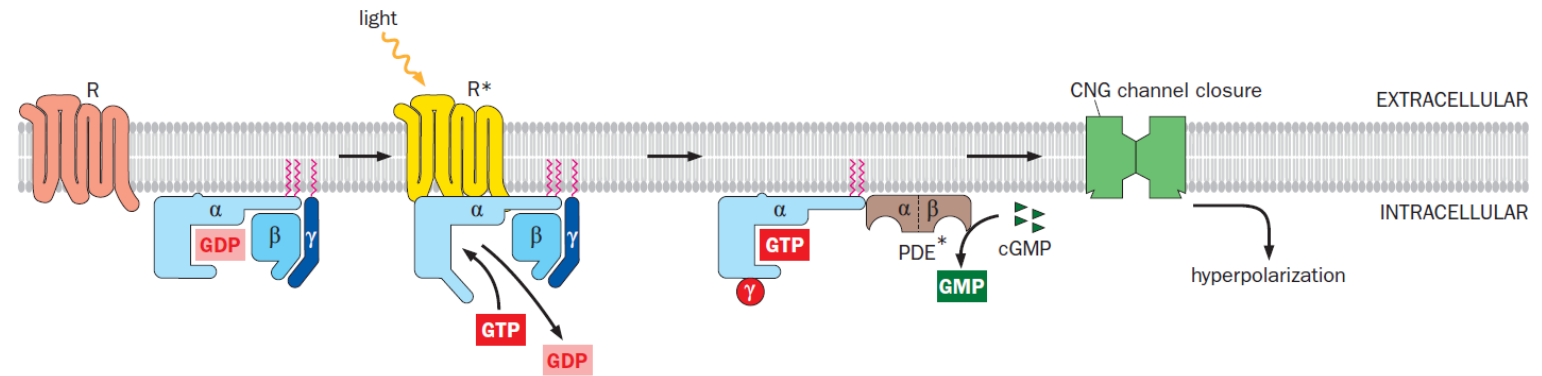
- 5) **Cry4 protein**, a member of cryptochrome family, is found recently located on the **outer segment** in cones in birds' retina.[17]
- 6) A popular hypothesis of (4)-(5) is an noncanonical biological pathway of visual system that can modulate night vision. **No such signaling pathway** is found till now.
- 7) We propose that no downstream signaling involved and instead a photon competition plays a key role. A novel biophysical model:

Opsin-Cryptochrome competition model(OCC model).

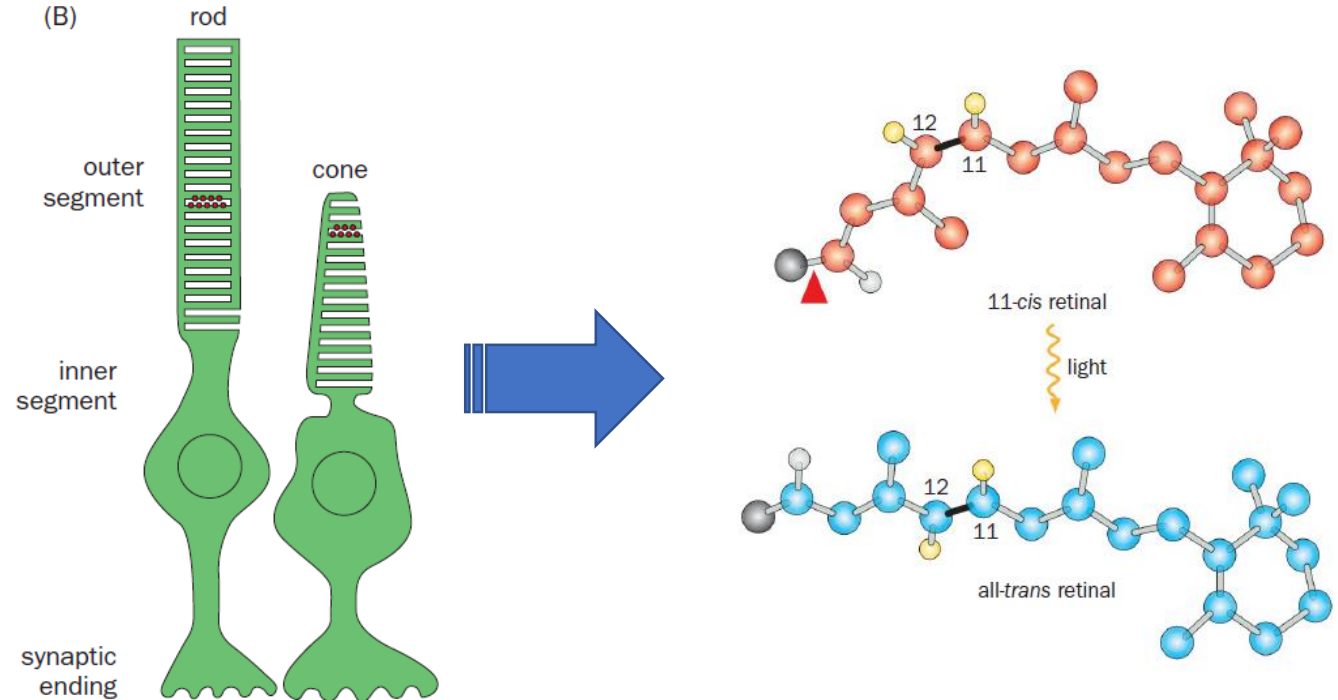
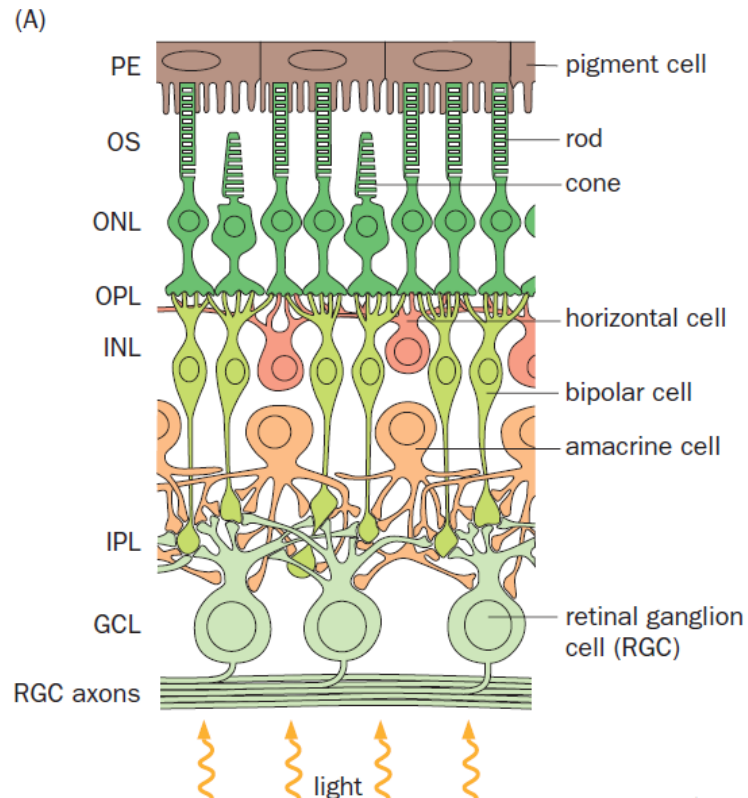


Opsin, Visual system

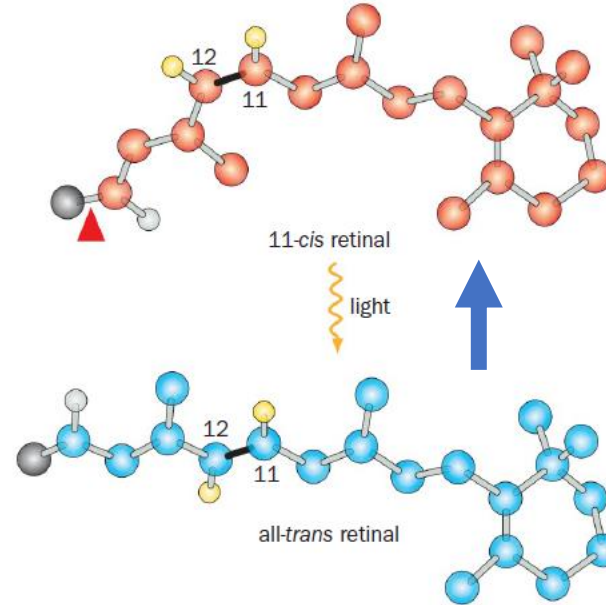
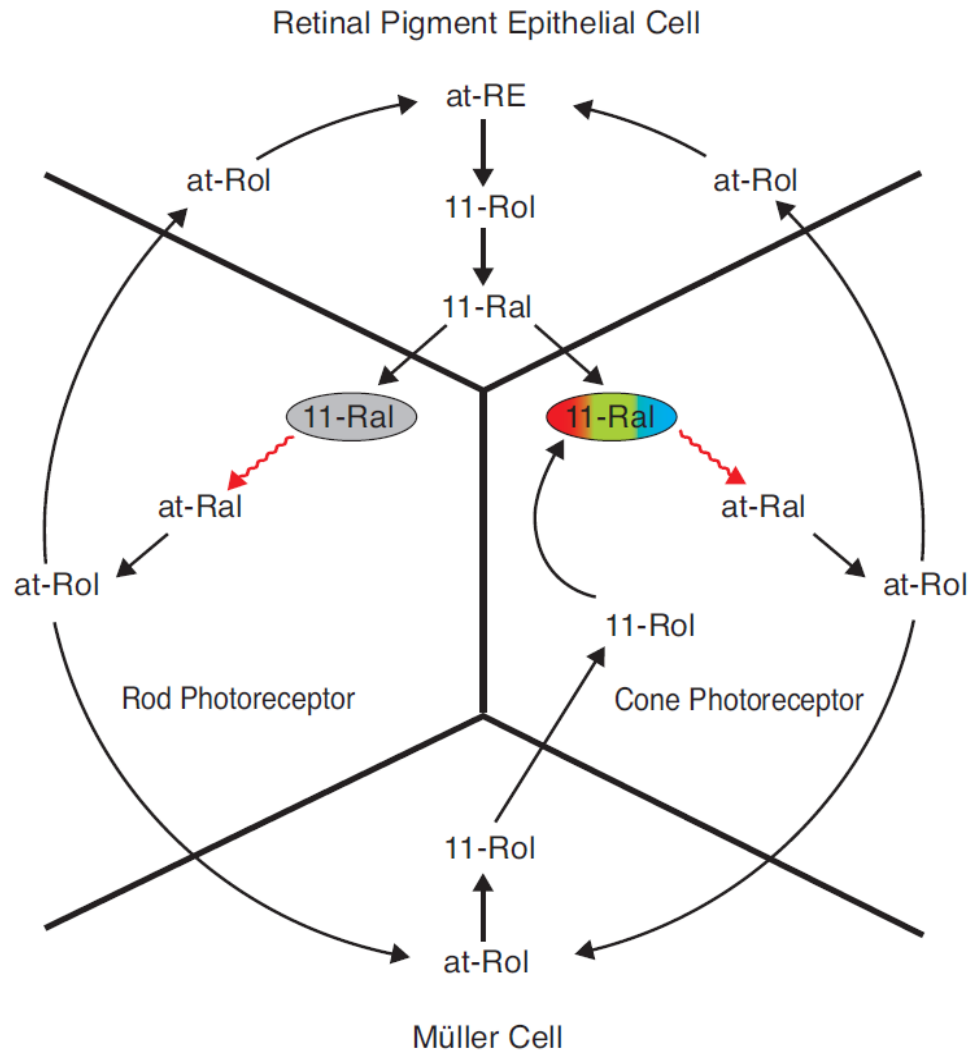
Light perception



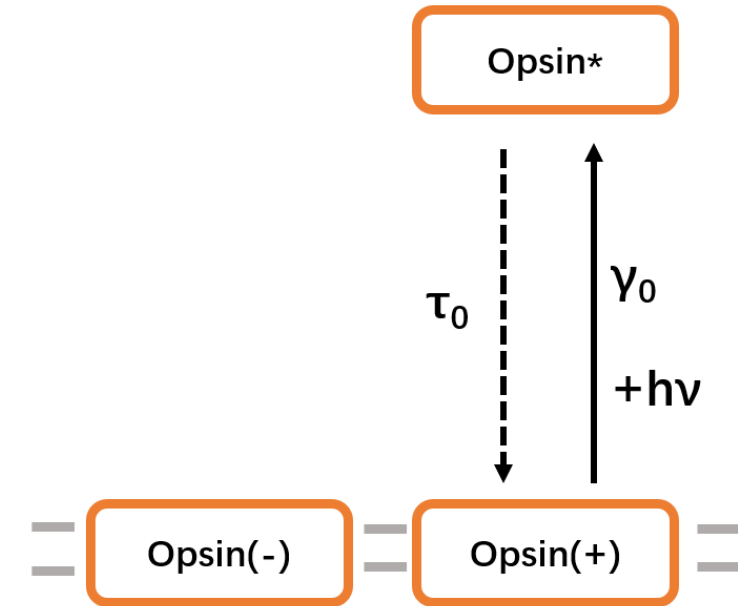
Phototransduction cascade



A cone visual cycle: cis-retinal \leftrightarrow trans-retinal



Retinal in a rhodopsin



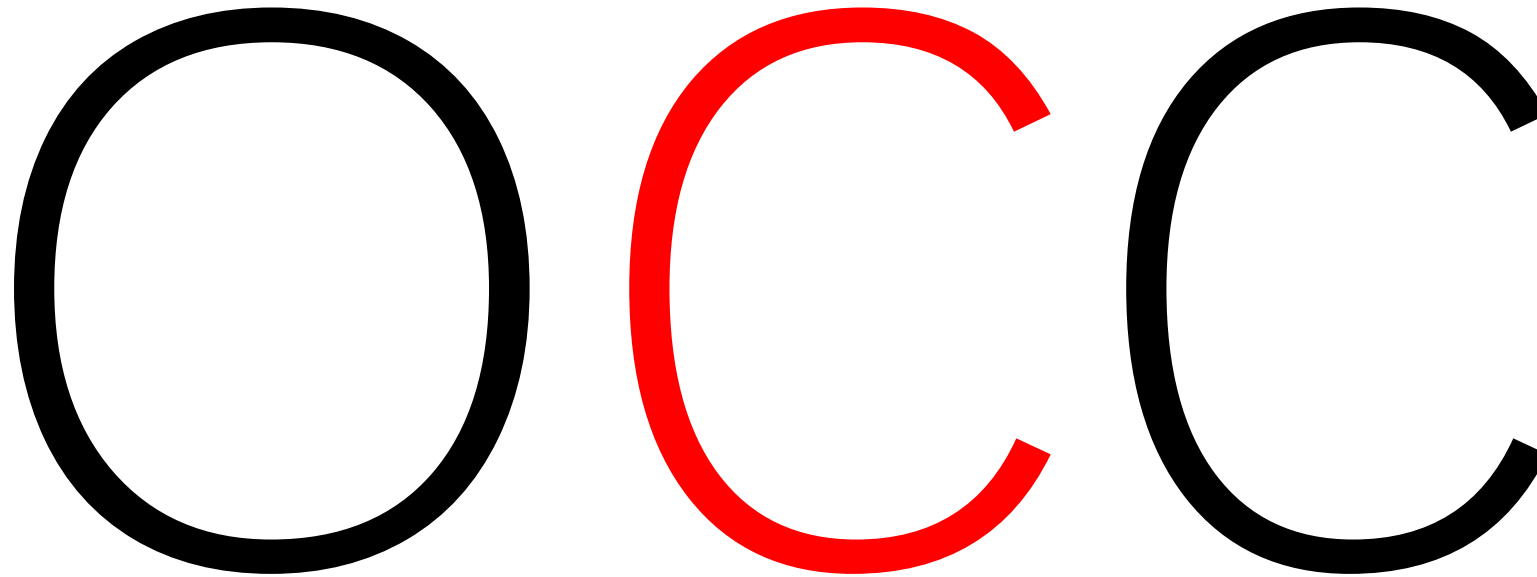
Abbreviations: at-RE, all-*trans*-retinyl esters; at-Ral, all-*trans*-retinal; at-Rol, all-*trans*-retinol; 11-Ral, 11-*cis*-retinal; 11-Rol, 11-*cis*-retinol.

Modified after Ref.[7,8]

Photoreceptor proteins responsible for vision

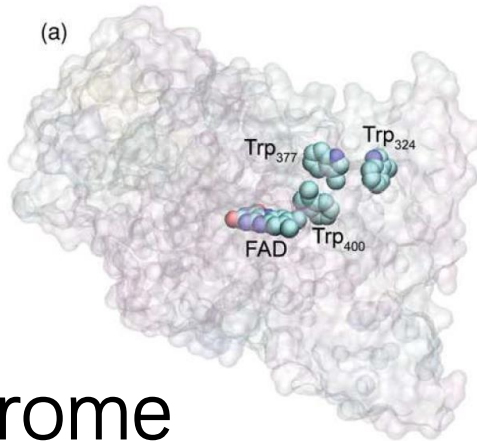
Name	Abbr.	Photo receptor	λ_{\max}	Color
Long-wave sensitive	LWS	Cone	500–570 nm	Green, yellow, red
Short-wave sensitive 1	SWS1	Cone	355–445 nm	Ultraviolet, violet
Short-wave sensitive 2	SWS2	Cone	400–470 nm	Violet, blue
Rhodopsin-like 2	Rh2	Cone	480–530 nm	Green
Rhodopsin-like 1 (vertebrate rhodopsin)	Rh1	Rod	~500 nm	Blue–green

Cryptochrome can absorb one photon with wavelength ranging from 400-565nm.

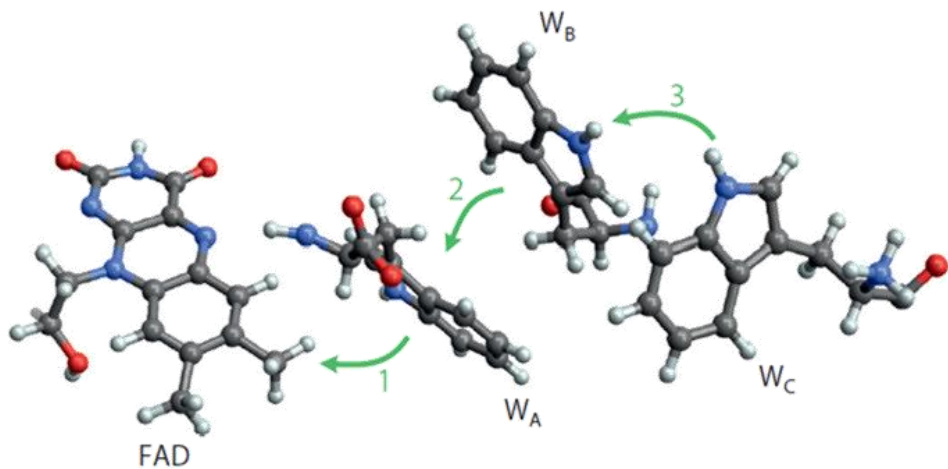


Cryptochrome, Radical pairs

Principle of radical pair mechanism

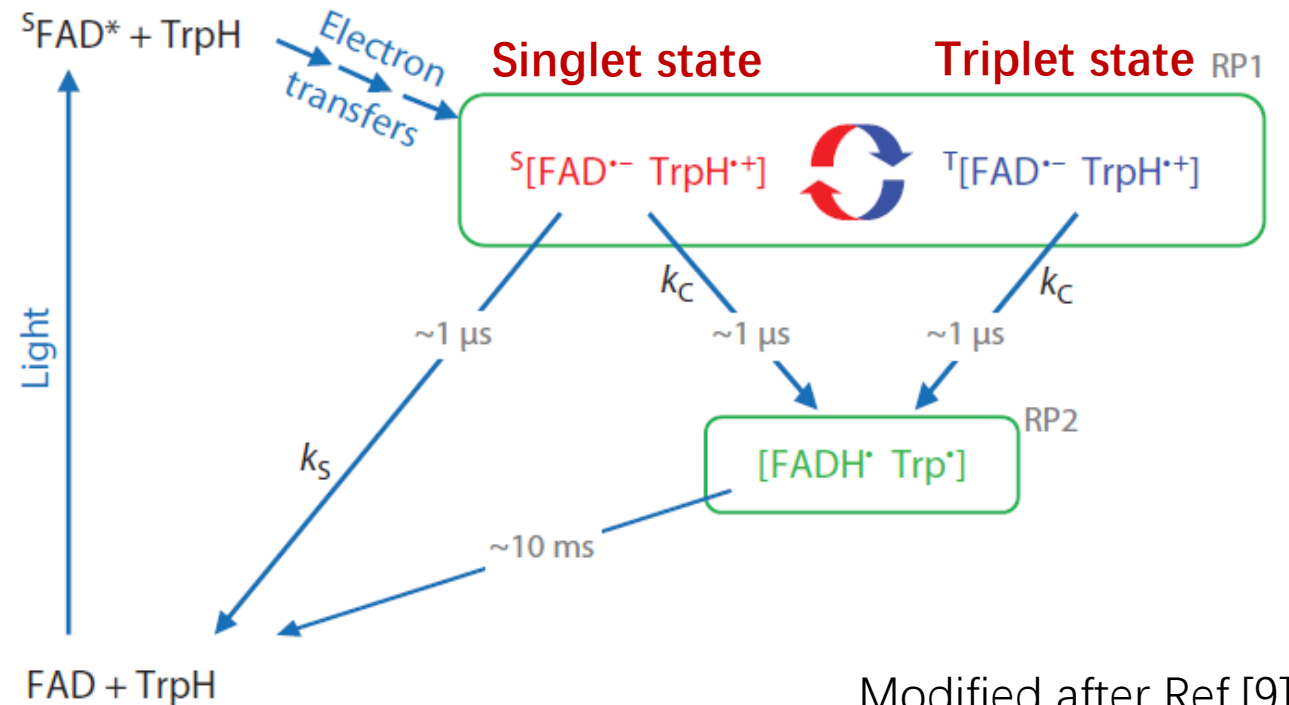
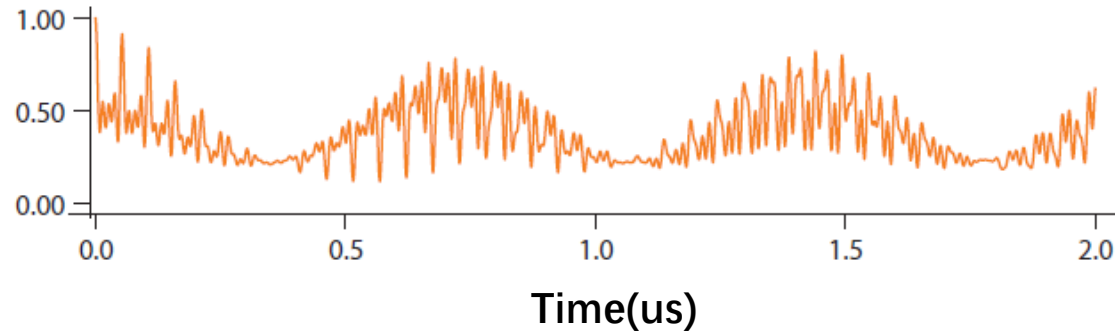


Cryptochrome



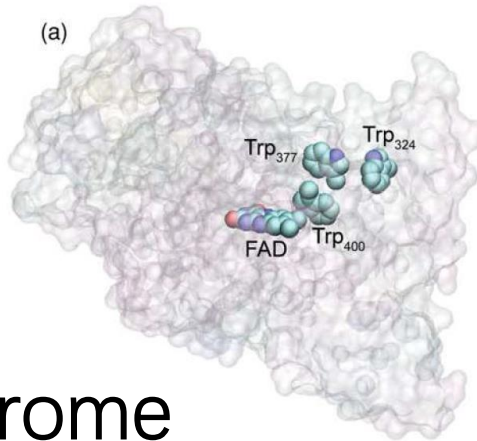
FAD - Flavin adenine dinucleotide
Trp(W) - Tryptophan

Singlet Fraction



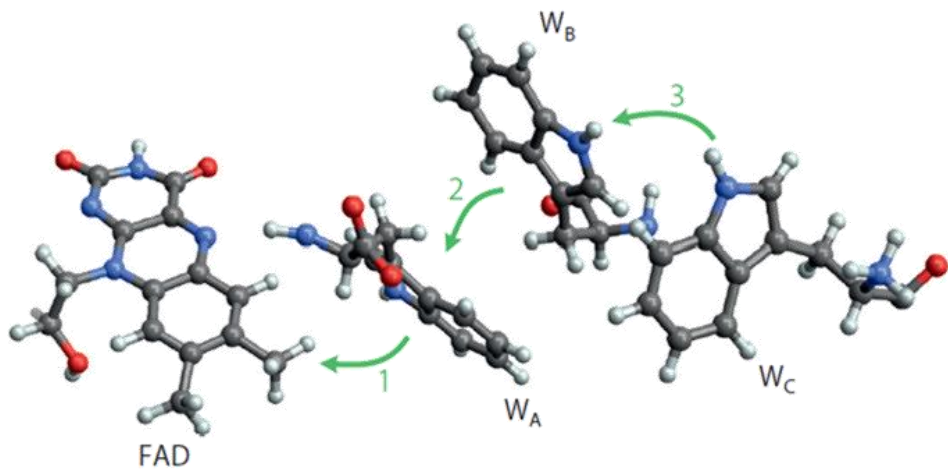
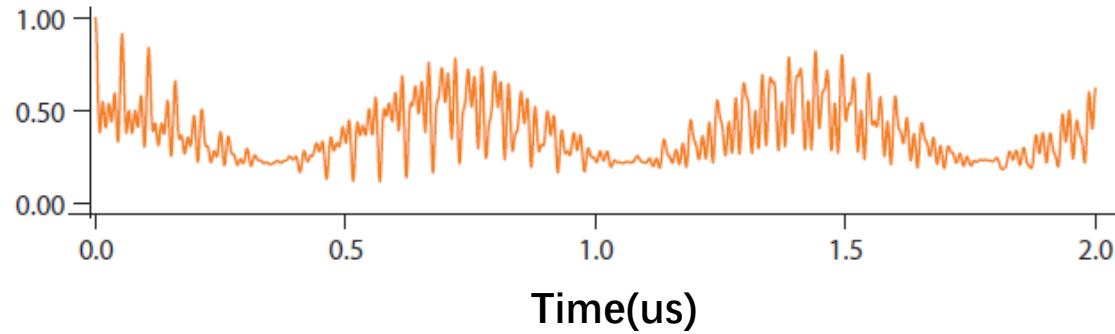
Modified after Ref.[9]

Principle of radical pair mechanism

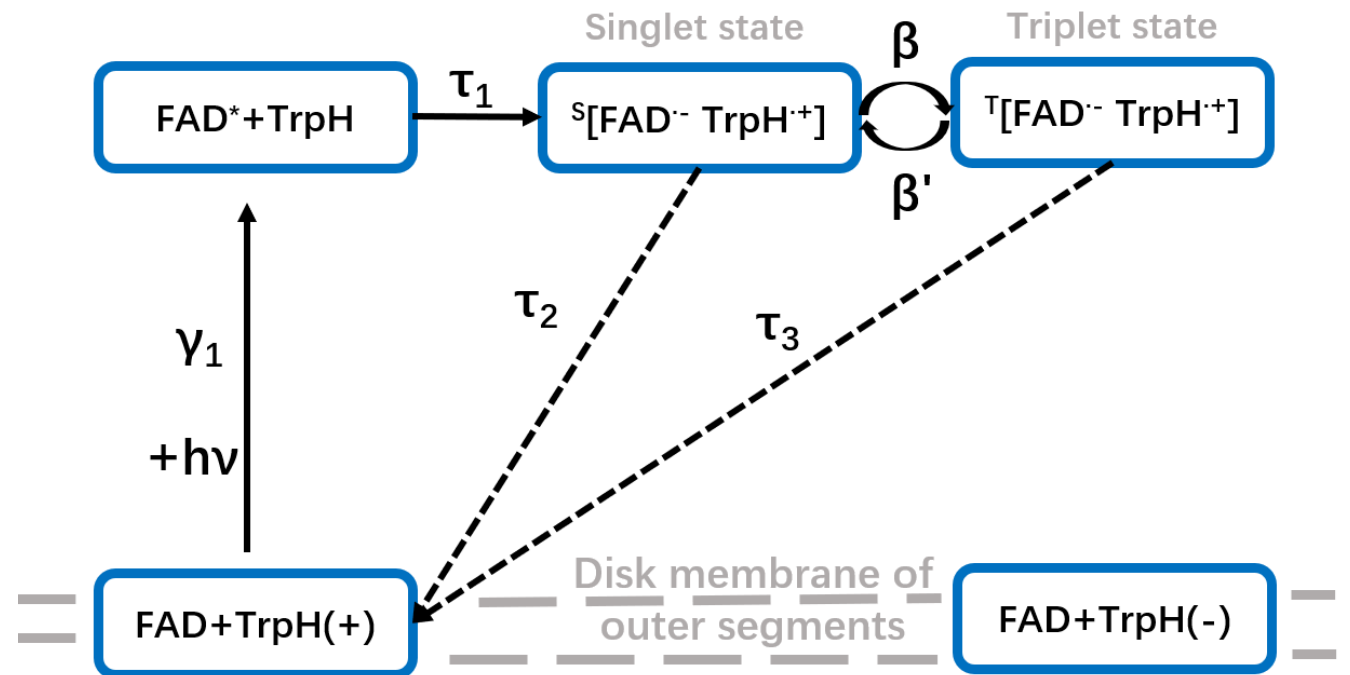


Cryptochrome

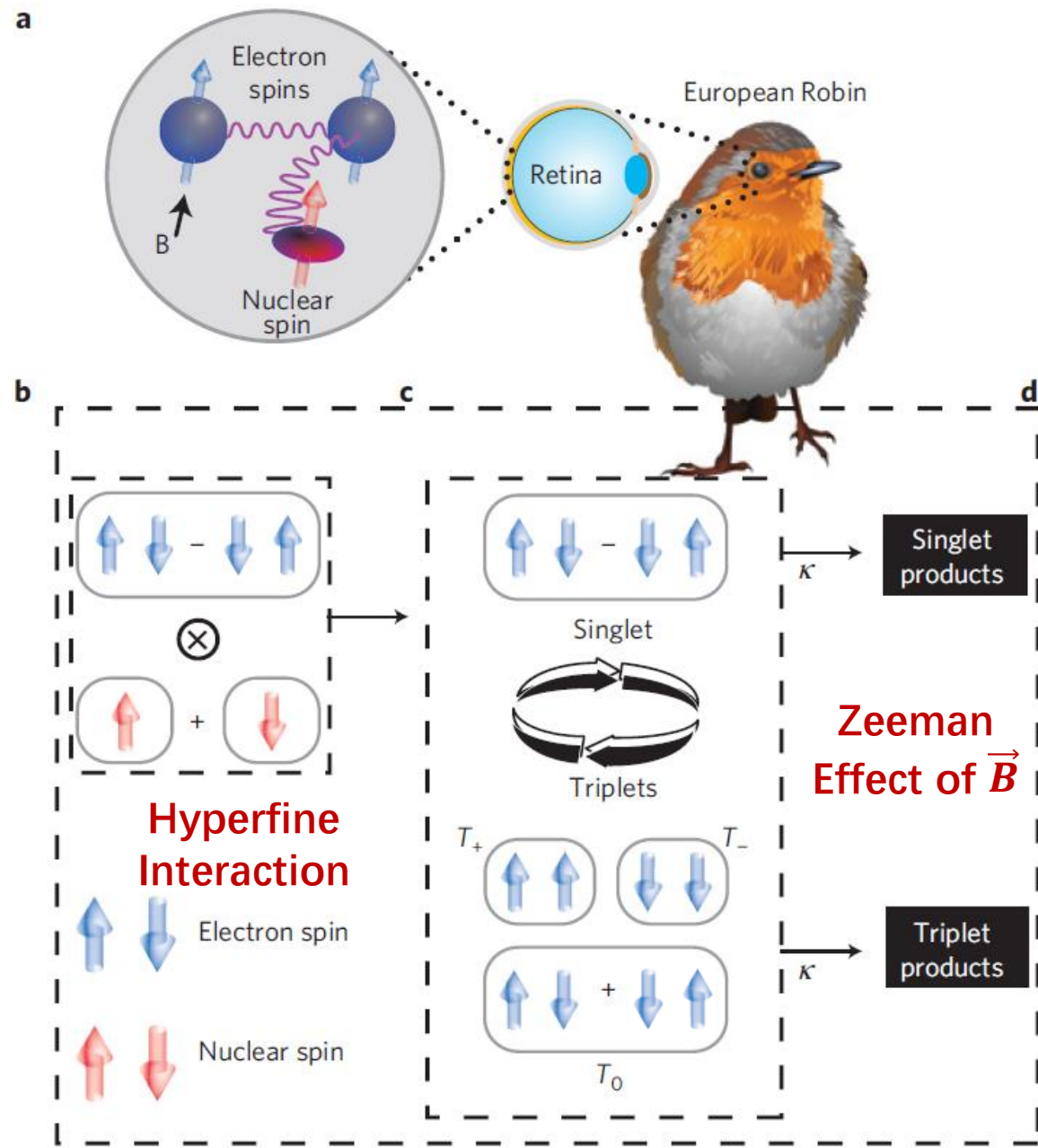
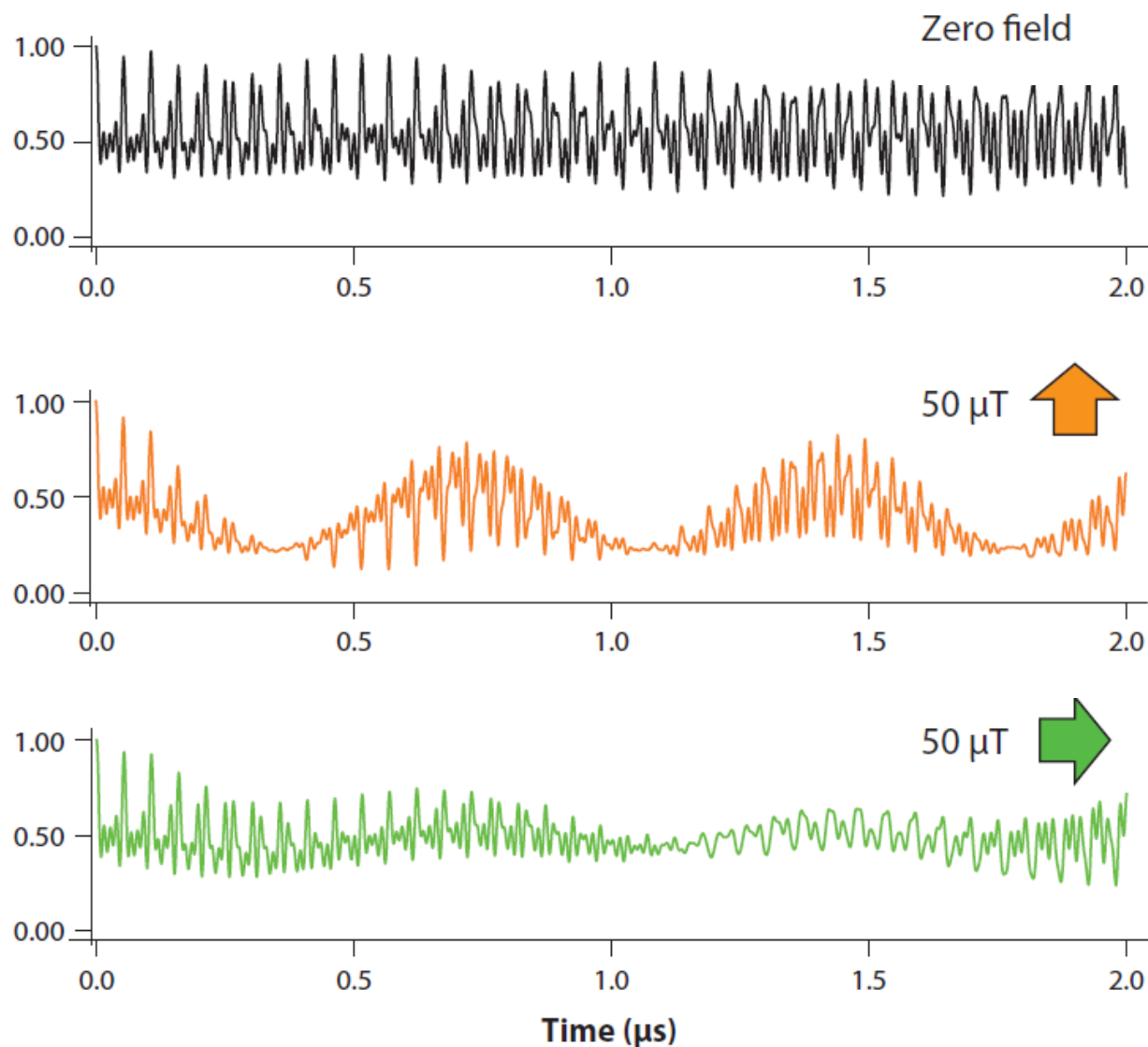
Singlet Fraction



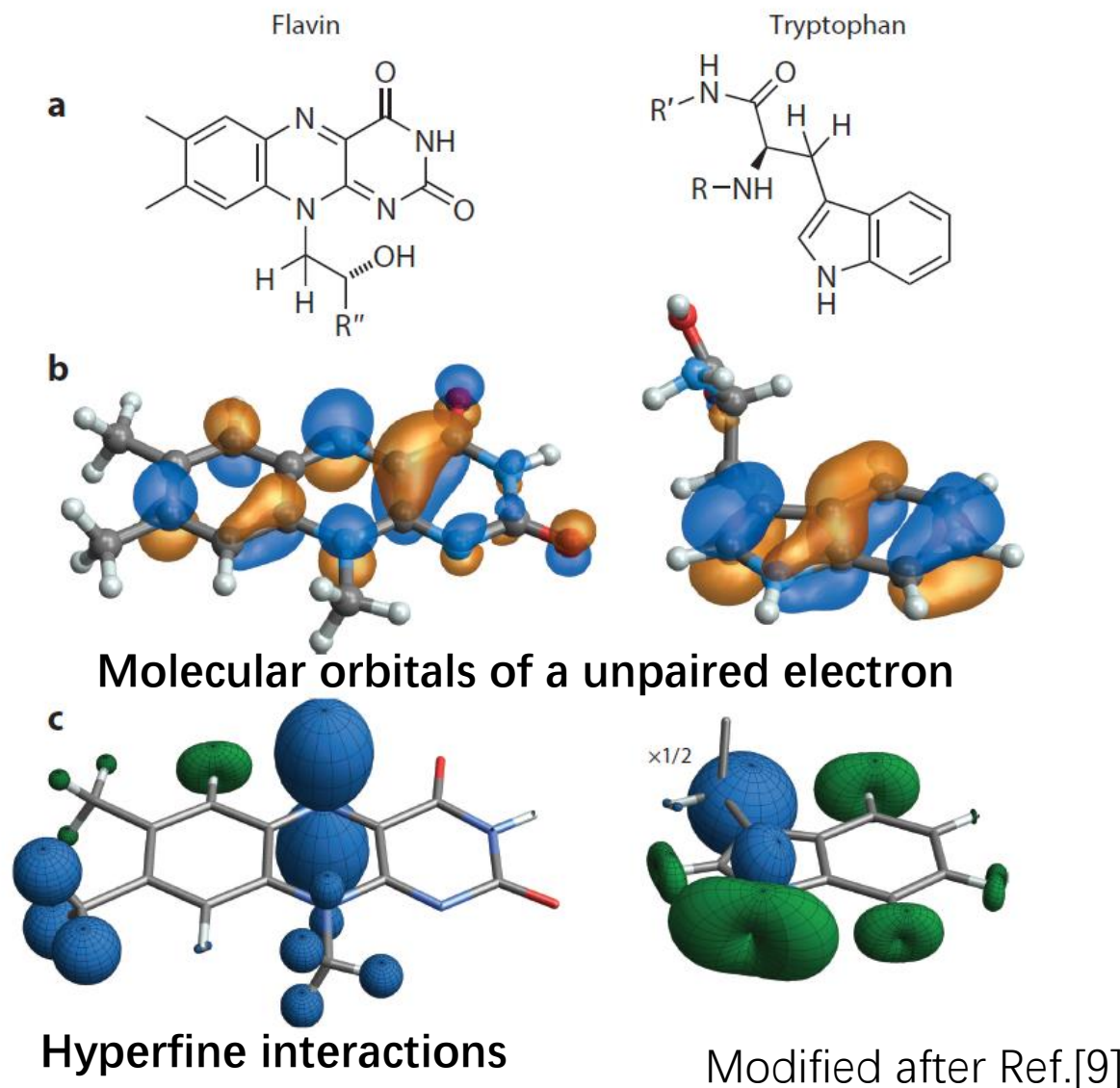
FAD - Flavin adenine dinucleotide
Trp(W) - Tryptophan



Spin dynamics



Quantum effect



The fractional yield of singlet and triplet state product is $\phi_S = 1 - \phi_T$ is an integral of the real part of $T(t)$ which is the singlet fraction at arbitrary time t .

$$\phi_T = k_T \int_0^\infty T(t) dt$$

$$T(t) = \text{Tr}[Q^T \rho(t)]$$

Q^T is the projection operator of triplet state while $\rho(t)$ is the density matrix of the radical pair at arbitrary time. The triplet state fraction is a trace operation of the density matrix projected on the triplet state product.

$$\rho(t) = \frac{1}{N} e^{-\frac{iHt}{\hbar}} \rho(0) e^{\frac{iHt}{\hbar}}$$

Assume there is only singlet product at time 0, where $\rho(0) = Q^S e^{-kt}$, here to make it simple, we let $k = k_S = k_T$. N is the number of nuclear spin.

The hamiltonian for two electron spins are H_1 and H_2 respectively. \vec{B} is the magnetic field vector with a specific direction.

$$H = H_1(\vec{B}) + H_2(\vec{B})$$

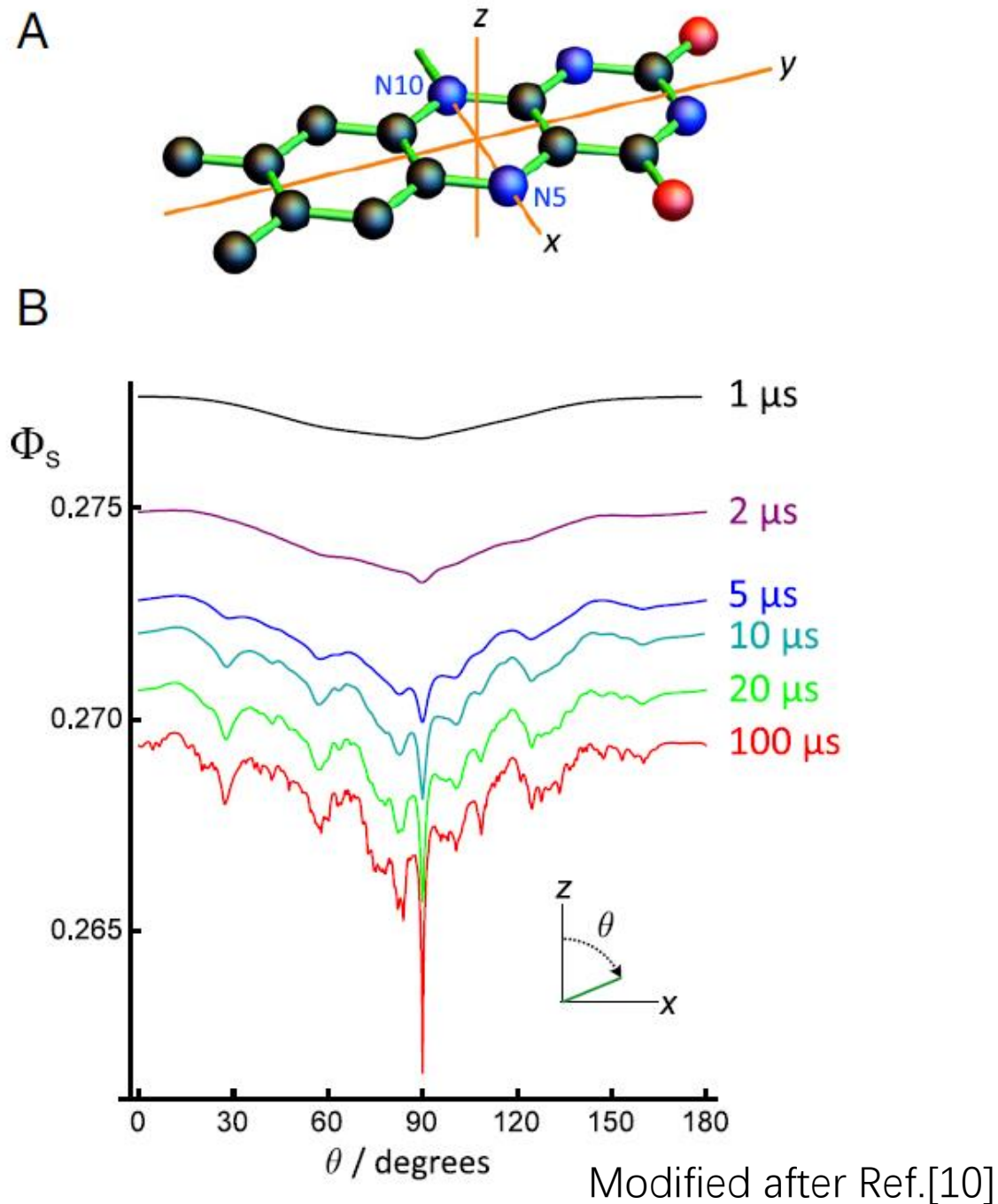
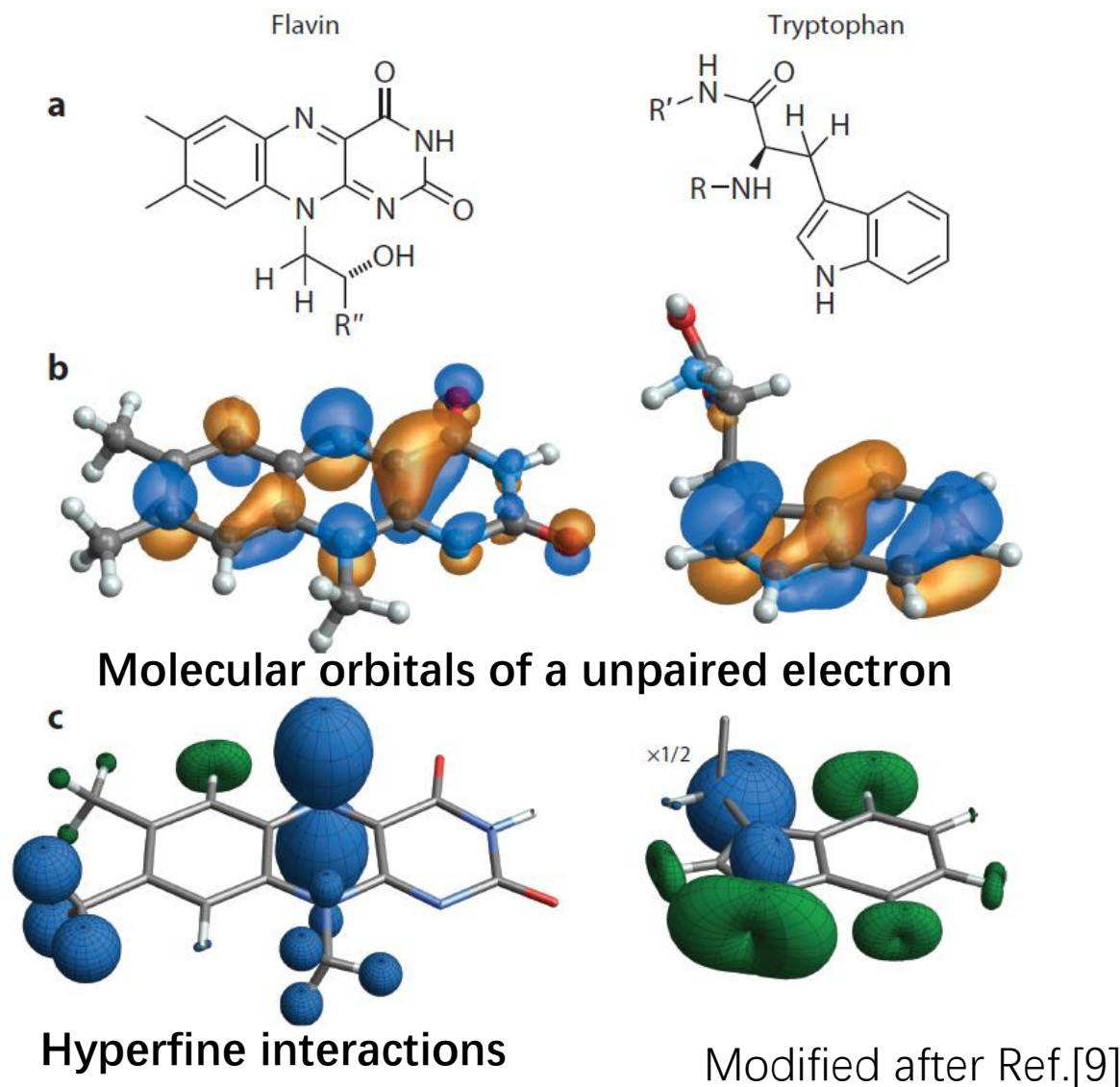
$$H_j = g\mu_B \vec{S}_j \cdot (\vec{B} + A_j \vec{I}_j)$$

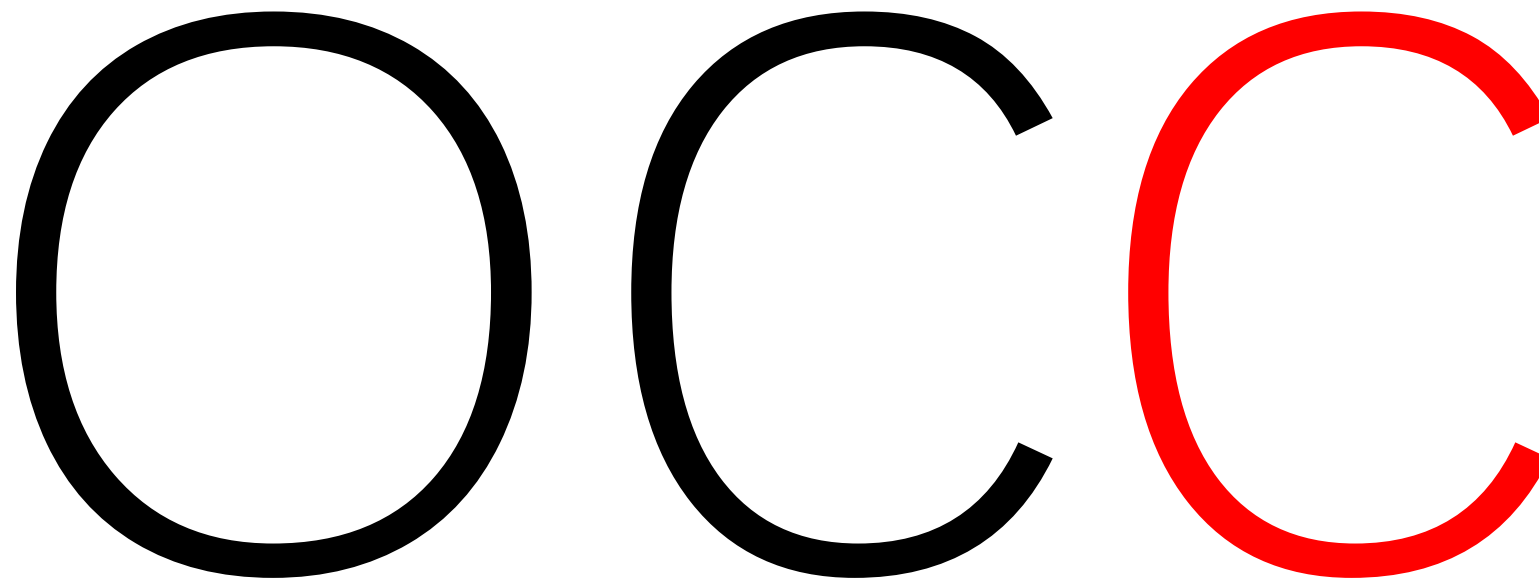
\vec{S}_j is the electron spin operator whereas \vec{I}_j is the nuclear spin operator correlated with the nuclear spin in vicinity. A_j is the anisotropic coefficient of the nuclear spin.

$$T(t) = \phi_S = 1 - \text{Tr}[Q^T \frac{1}{N} e^{-\frac{iHt}{\hbar}} Q^S e^{-kt} e^{\frac{iHt}{\hbar}}]$$

$$\phi_S = 1 - \frac{1}{N} \sum_{mn} Q_{mn}^T Q_{nm}^S \frac{k^2}{k^2 + (\omega_m - \omega_n)^2}$$

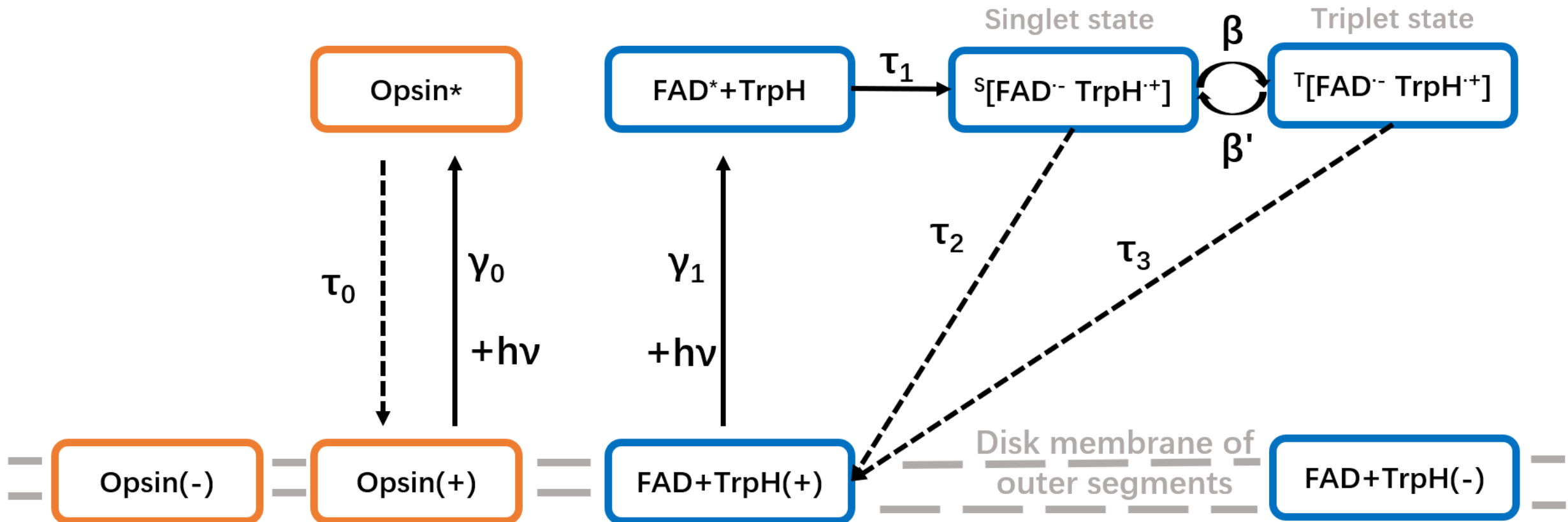
Quantum effect





Photon Competition

Opsin-Cryptochrome Competition model

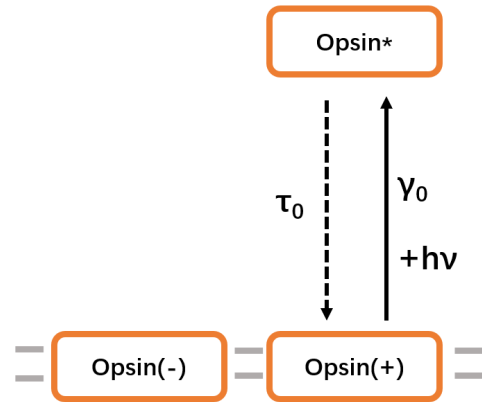


(+) represents the ground state of a molecule absorbed by one photon; (*) represents the excitation state of a molecule after photon absorption; (s) and (t) represent the singlet and triplet state of a molecule.

Model formulation

$$N_{op}^{(+)} + N_{FAD}^{(+)} = N_{photon} \quad (9)$$

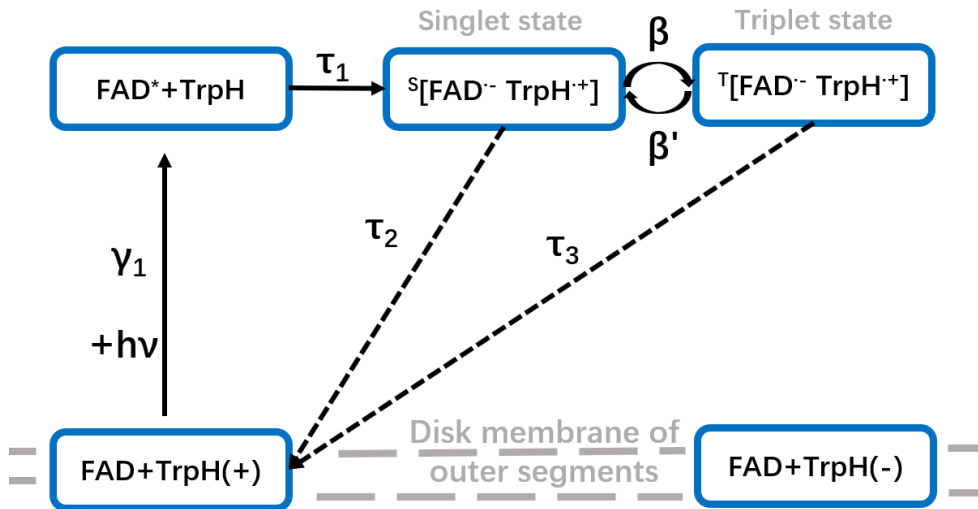
$$R_{opsin} = \frac{N_{op}^{(+)}}{N_{photon}} \quad (10)$$



$$N_{op}^{(+)} + N_{op}^* + N_{op}^{(-)} = C_1 \quad (1)$$

$$\frac{d(N_{op}^{(+)} + N_{op}^{(-)})}{dt} = -\gamma_0 N_{op}^{(+)} + \frac{1}{\tau_0} N_{op}^* \quad (2)$$

$$\frac{dN_{op}^*}{dt} = \gamma_0 N_{op}^{(+)} - \frac{1}{\tau_0} N_{op}^* \quad (3)$$



$$N_{FAD}^{(+)} + N_{FAD}^* + N_{FAD}^{Singlet} + N_{FAD}^{Triplet} + N_{FAD}^{(-)} = C_2 \quad (4)$$

$$\frac{d(N_{FAD}^{(+)} + N_{FAD}^{(-)})}{dt} = -\gamma_1 N_{FAD}^{(+)} + \frac{1}{\tau_2} N_{FAD}^{Singlet} + \frac{1}{\tau_3} N_{FAD}^{Triplet} \quad (5)$$

$$\frac{dN_{FAD}^*}{dt} = \gamma_1 N_{FAD}^{(+)} - \frac{1}{\tau_1} N_{FAD}^* \quad (6)$$

$$\frac{dN_{FAD}^{Singlet}}{dt} = \frac{1}{\tau_1} N_{FAD}^* - \frac{1}{\tau_2} N_{FAD}^{Singlet} - \beta N_{FAD}^{Singlet} + \beta' N_{FAD}^{Triplet} \quad (7)$$

$$\frac{dN_{FAD}^{Triplet}}{dt} = \beta N_{FAD}^{Singlet} - \beta' N_{FAD}^{Triplet} - \frac{1}{\tau_3} N_{FAD}^{Triplet} \quad (8)$$

A stationary solution: let all derivatives=0

$$N_{op}^{(+)} + N_{FAD}^{(+)} = N_{photon} \quad (9)$$

$$R_{opsin} = \frac{N_{op}^{(+)}}{N_{photon}} \quad (10)$$

$$N_{op}^{(-)} + B * N_{op}^{(+)} = C_1 \quad (11)$$

$$N_{FAD}^{(-)} + A * N_{FAD}^{(+)} = C_2 \quad (12)$$

$$(A - B)N_{op}^{(+)^2} + (C_1 + C_2 - N_{photon}(A - B))N_{op}^{(+)} - N_{photon}C_1 = 0 \quad (15)$$

$$R_{opsin} = \frac{-(C_1 + C_2 - N_{photon}(A - B)) + \sqrt{(C_1 + C_2 - N_{photon}(A - B))^2 + 4N_{photon}C_1(A - B)}}{2N_{photon}(A - B)} \quad (16)$$

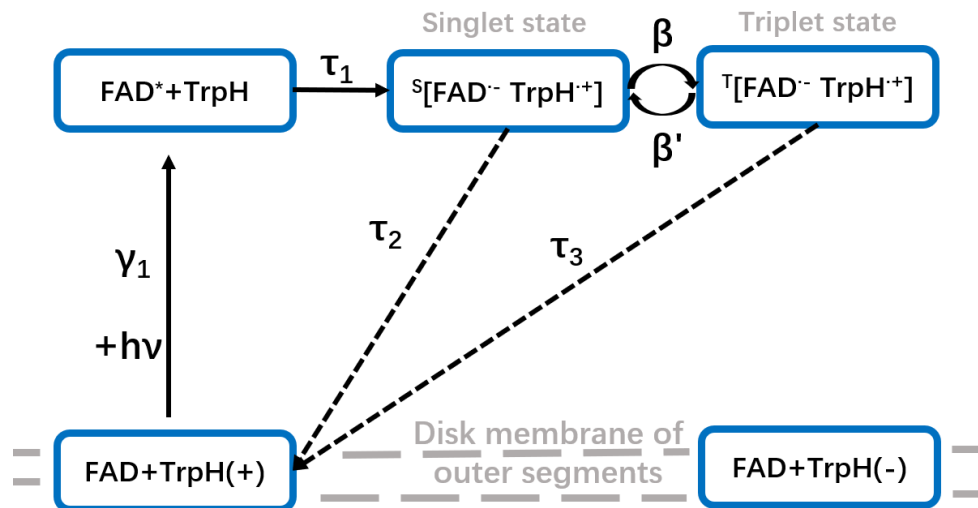
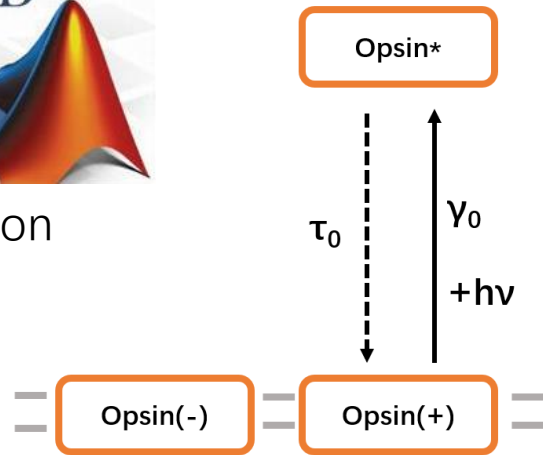
$$A = 1 + \gamma_1 \left(\tau_1 + \frac{1}{\frac{\beta'}{\tau_2\beta} + \frac{1}{\tau_2\tau_3\beta} + \frac{1}{\tau_3}} + \frac{1}{\frac{1}{\tau_2} + \frac{\beta}{\tau_3\beta' + 1}} \right) \quad B = \tau_0\gamma_0 + 1$$

Simulation

Parameter initiation



Simulation



- **Singlet-triplet oscillation β coefficient:**

$$\beta' = 0; \beta = \frac{\Delta(\Phi T)}{\Delta(\tau_2)} \text{ in terms of angle}$$

- **Reciprocal of rate constant(lifetime):**

$$\tau_0 = 4.2\text{ms}; \tau_1 = 1\text{ns};$$

$$\tau_2 = 1\mu\text{s}; \tau_3 = 10\text{ms};$$

- **Rate constant of light absorption:**

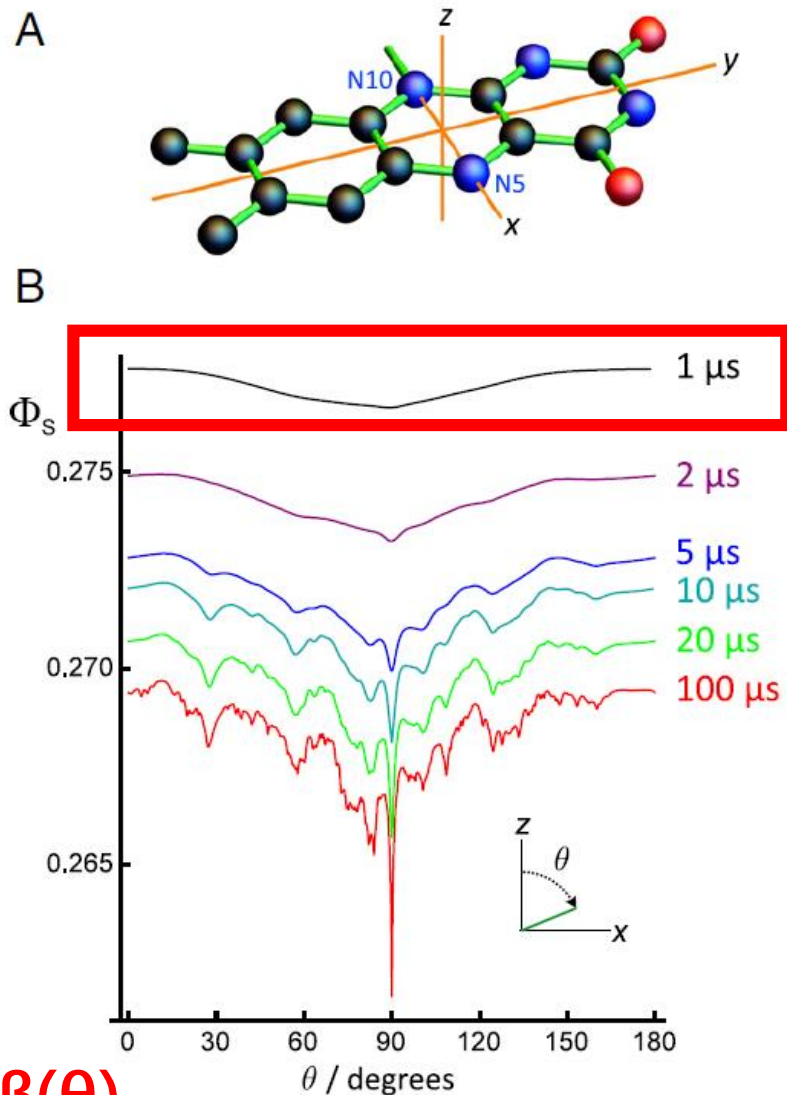
$$\gamma_0 = \gamma_1 = 10^9/\text{s};$$

- Assume the **protein number** of opsin and cryptochrome is the same: $C_1 = C_2$

- **Incident photon number N_{photon}**

$$\text{where } N_{\text{photon}} : C_1 = 10^{-3}$$

Parameter initiation

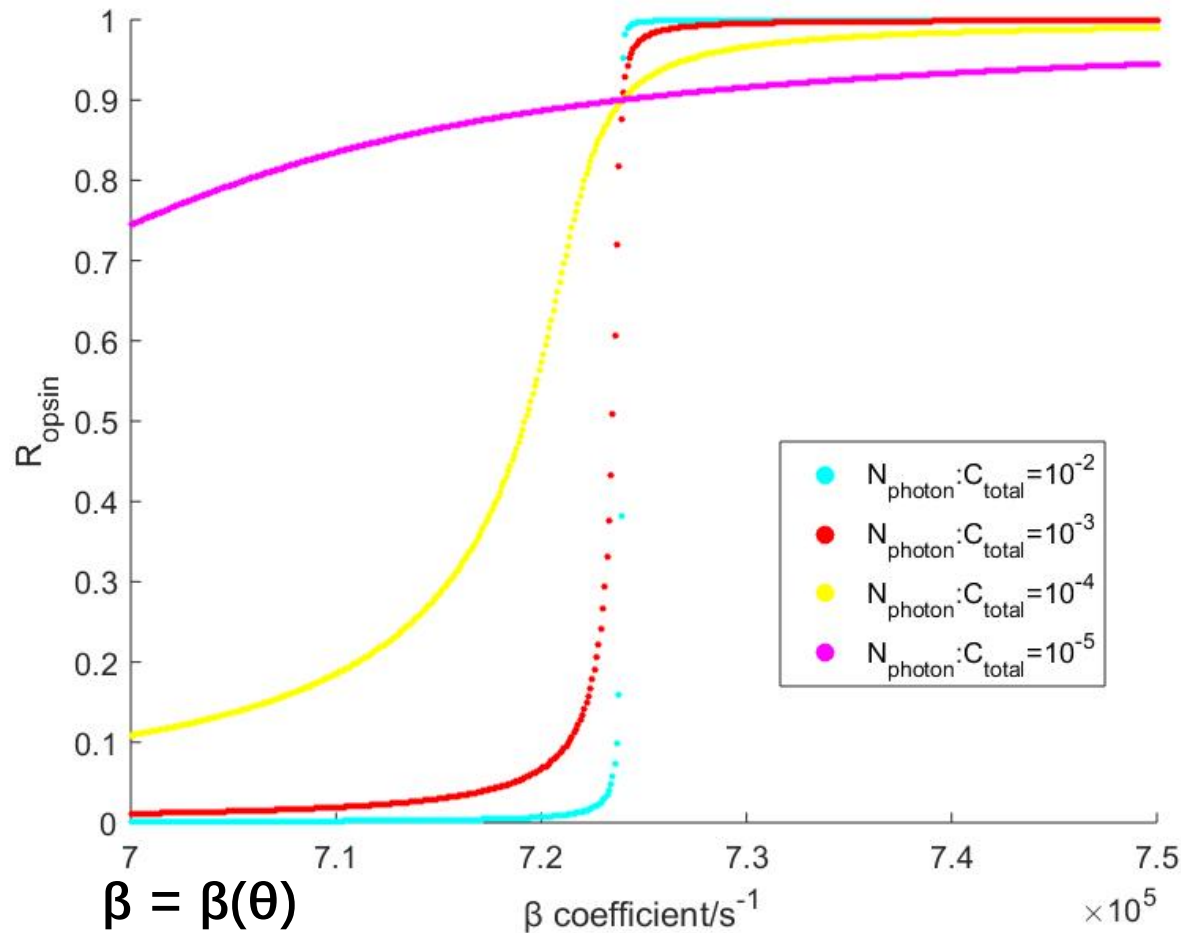


$$\beta = \beta(\theta)$$

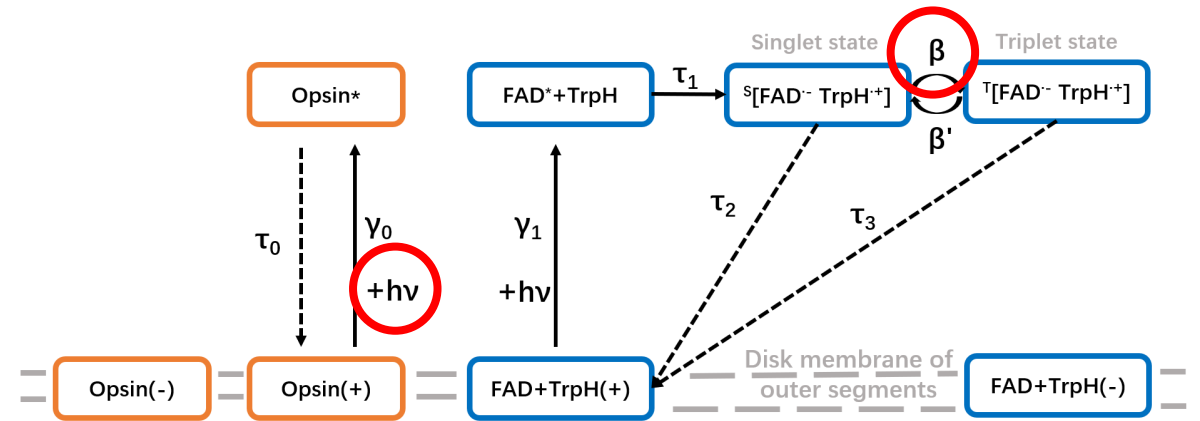
- Singlet-triplet oscillation β coefficient:
 $\beta' = 0$; $\beta = \frac{\Delta(\Phi T)}{\Delta(\tau_2)}$ in terms of angle
- Reciprocal of rate constant(lifetime):
 $\tau_0 = 4.2\text{ms}$; $\tau_1 = 1\text{ns}$;
 $\tau_2 = 1\mu\text{s}$; $\tau_3 = 10\text{ms}$;
- Rate constant of light absorption:
 $\gamma_0 = \gamma_1 = 10^9/\text{s}$;
- Assume the protein number of opsin and cryptochrome is the same: $C_1 = C_2$
- Incident photon number N_{photon}
 where $N_{\text{photon}}:C_1 = 10^{-3}$

Secondary amplification effect

$$R_{opsin} = \frac{-(C_1 + C_2 - N_{photon}(A - B)) + \sqrt{(C_1 + C_2 - N_{photon}(A - B))^2 + 4N_{photon}C_1(A - B)}}{2N_{photon}(A - B)}$$

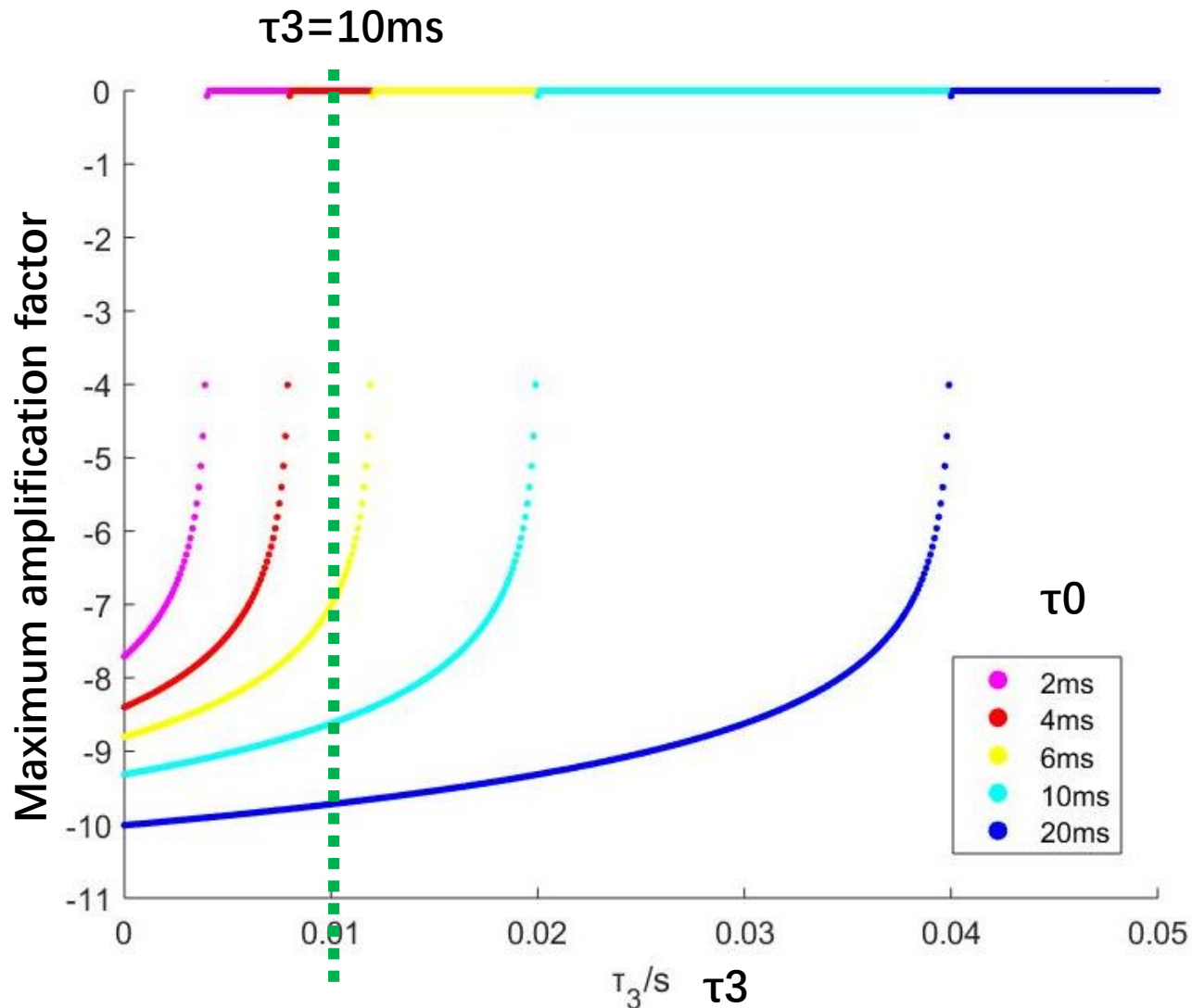


$R_{opsin} \in (0, 1)$ representing the percentage of incident photons that are absorbed by rhodopsin.

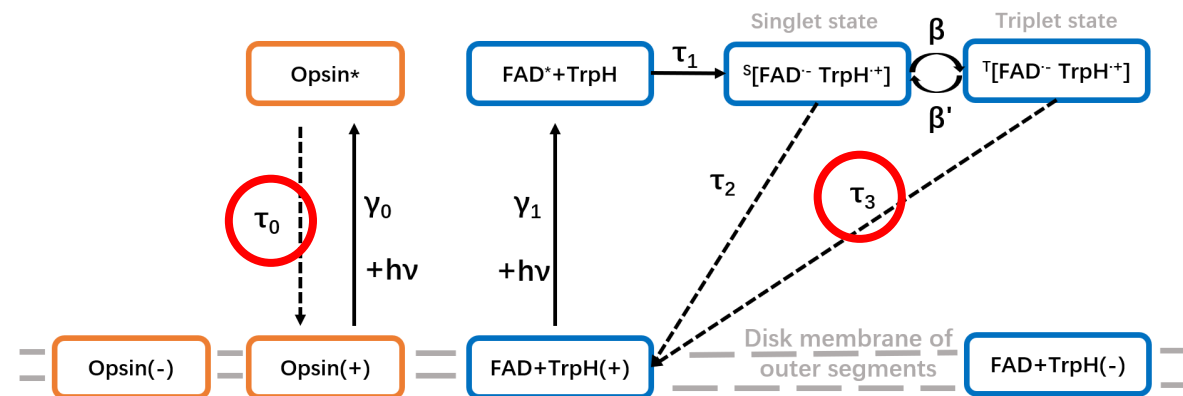


- Ratio of incident photons and protein numbers are critical for amplification **indicating a photon threshold** for magnetoreception.

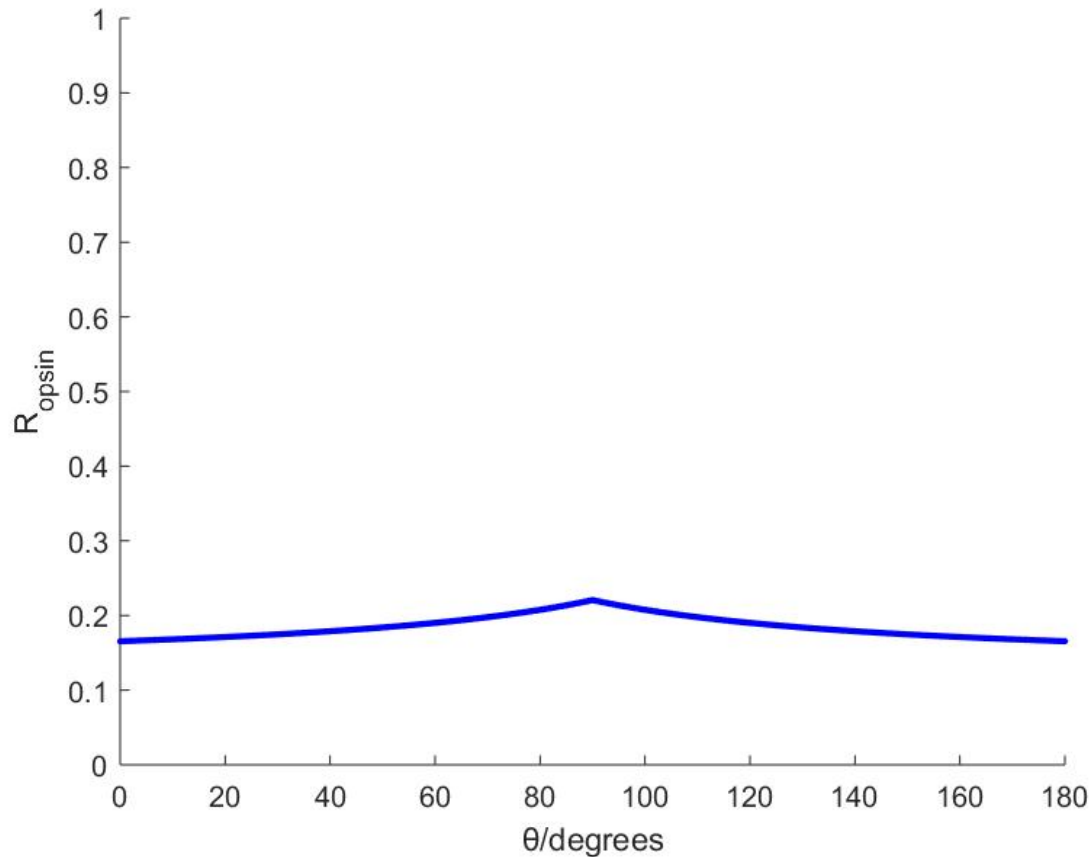
Secondary amplification effect



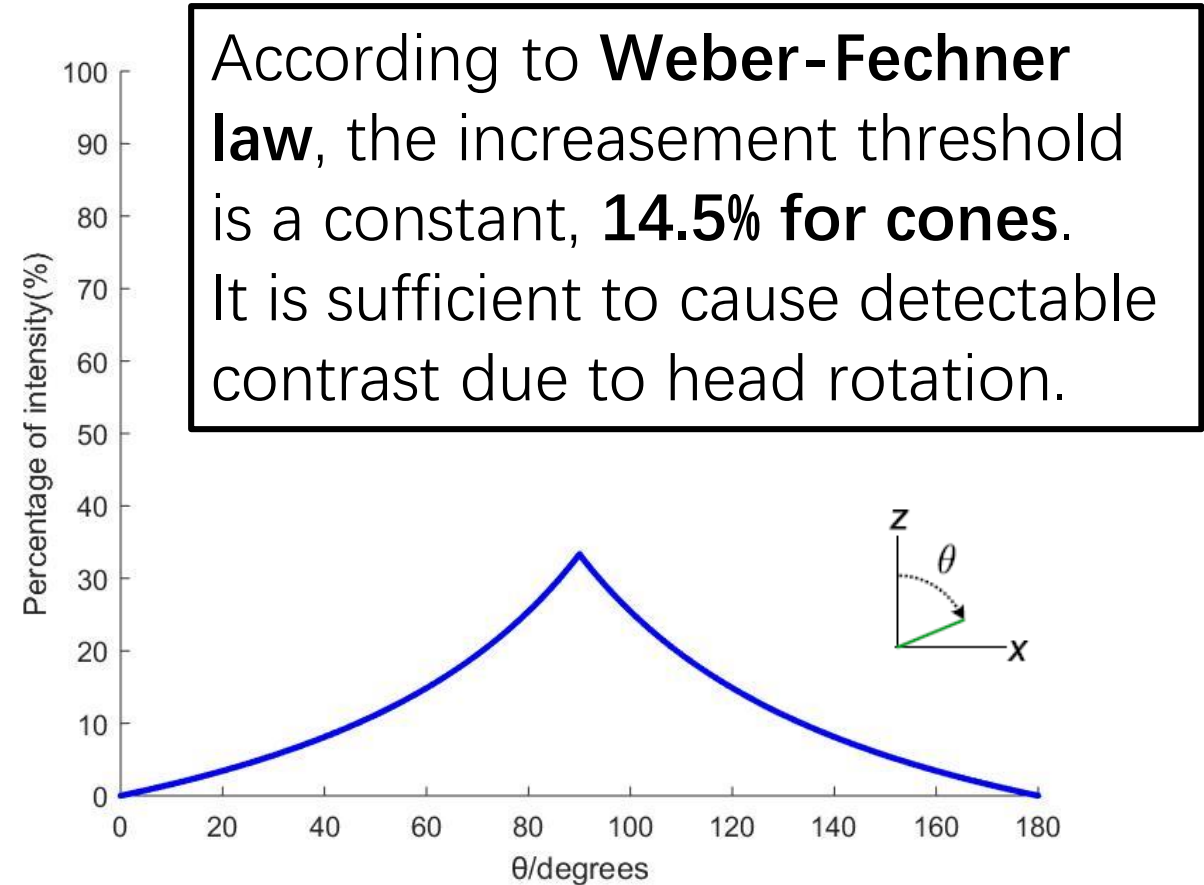
- The quantity of τ_0 is critical for the amplification effect.
- The secondary amplification effect is a key step to **convert a physical cause into biological consequence**.



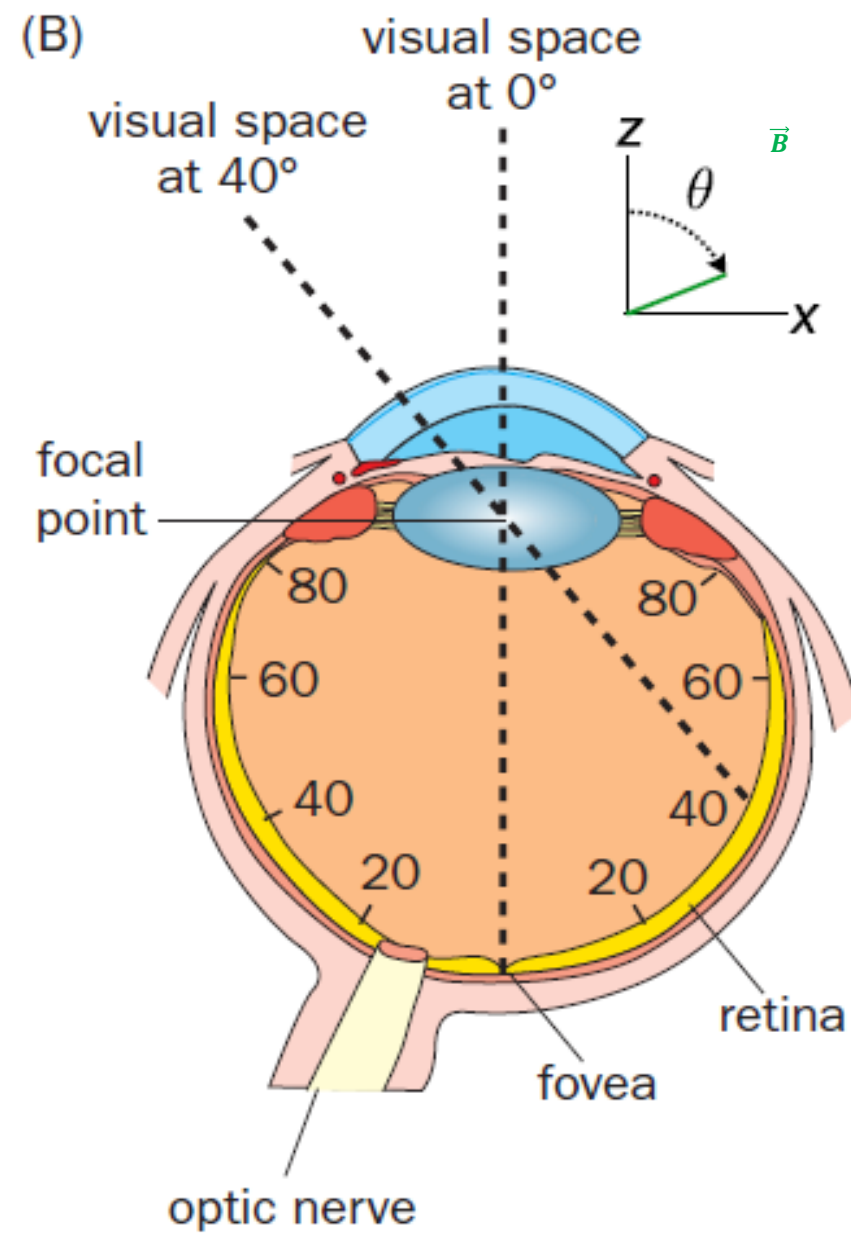
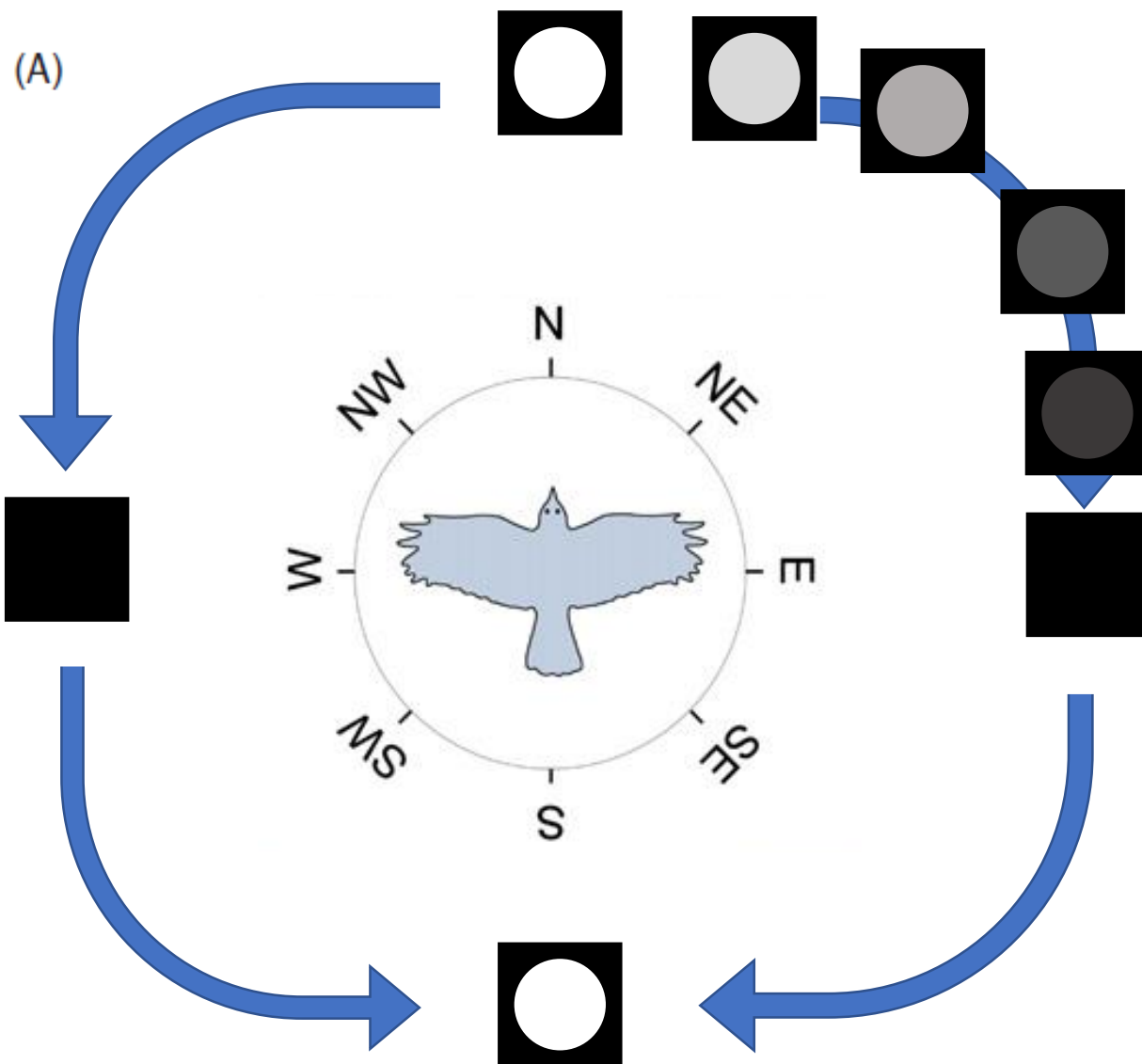
Photon competition simulation result



$$R_{\text{opsin}} = R_{\text{opsin}}(\beta) \text{ and } \beta = \beta(\theta)$$



$$\text{Increasement threshold} = \frac{\Delta I}{I}$$

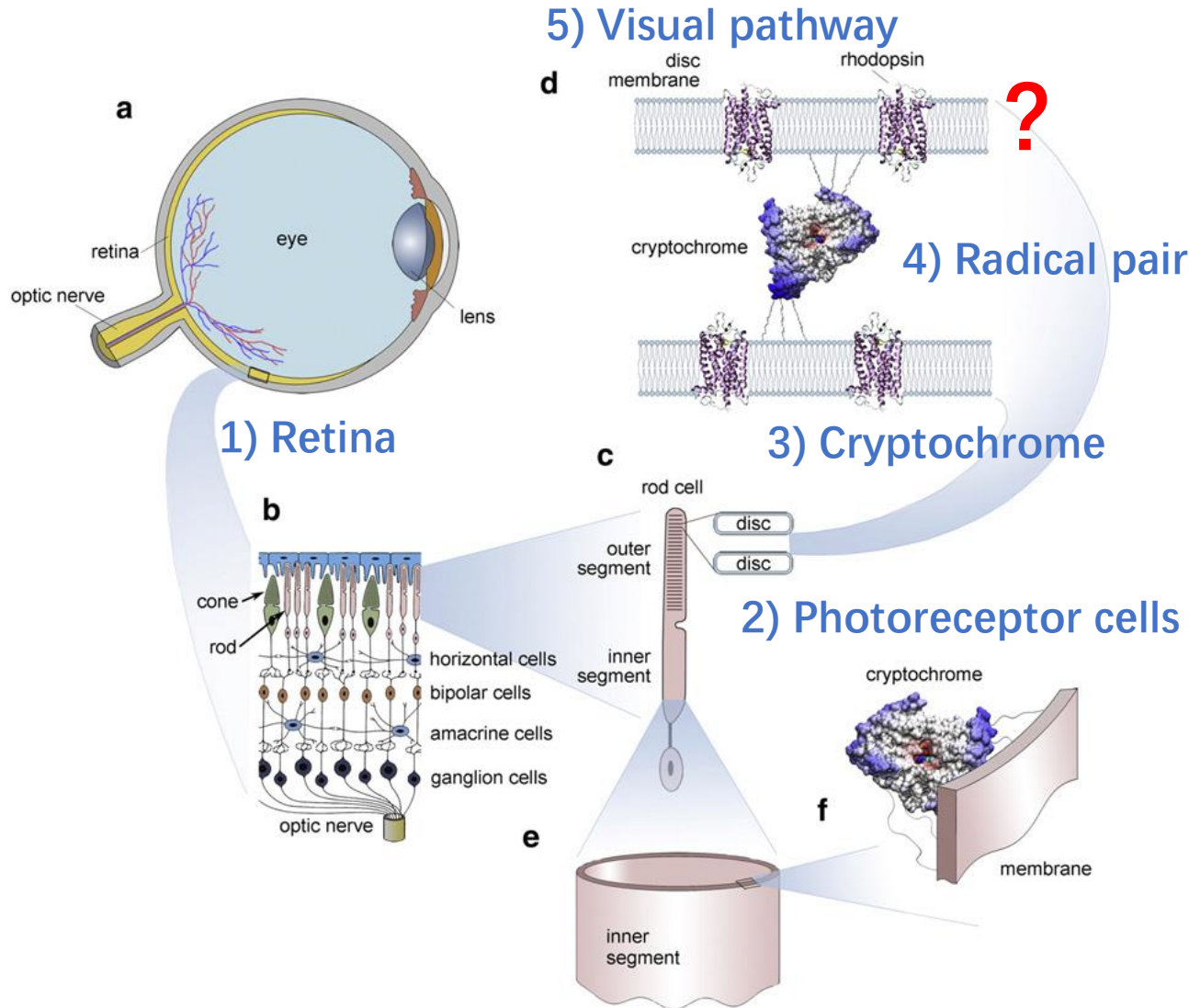


Experimental predictions

- Our model give a prediction that **cone cells are magnetic sensors** with two properties:
 - (1) **Incident photon threshold** for magnetic sensing.
 - There are lower and upper photon thresholds for competition of opsin and cryptochrome. Controlled light intensity of background can be a on-off effect for magnetoreception.
 - (2) **Disruption/Analogy** of the magnetic sensing.
 - Under zero magnetic field, generate an artificial light pattern resembling the earth magnetic field effect.
 - Use changing spatial intensity of incident light on the retina to disrupt the existing light pattern caused by earth magnetic field .

Summary

? ➡ OCC?



- A popular hypothesis of (4)-(5) is an noncanonical biological pathway of visual system that can modulate night vision. **No such signaling pathway** is found till now.
 - We propose that no downstream signaling involved and instead a photon competition plays a key role.
- A novel biophysical model:

Opsin-Cryptochrome competition model(OCC model).

Experiments to verify this model are proposed.

Discussion

- Our model only works under **the radical-based magnetoreception in migratory birds** under **total overcast and under dim light**.
- European robin, pigeon and turtle seems use a inclination magnetic compass whereas other animals use intensity or polarity compass.

Parameter setting:

- Recovery time of retinal: τ_0
 - Singlet-triplet oscillation β
 - Singlet product τ_2
 - Triplet products τ_3
1. Decoherence time – molecular motion
 2. Photon threshold and visual pattern
 3. Protein distribution
 4. Number ratio of cry4 and opsin protein
 5. Intensity perception of rods and cones

References

- [1] Keeton W T . Magnets Interfere with Pigeon Homing[J]. Proceedings of the National Academy of Sciences of the United States of America, 1971, 68(1):102-106.
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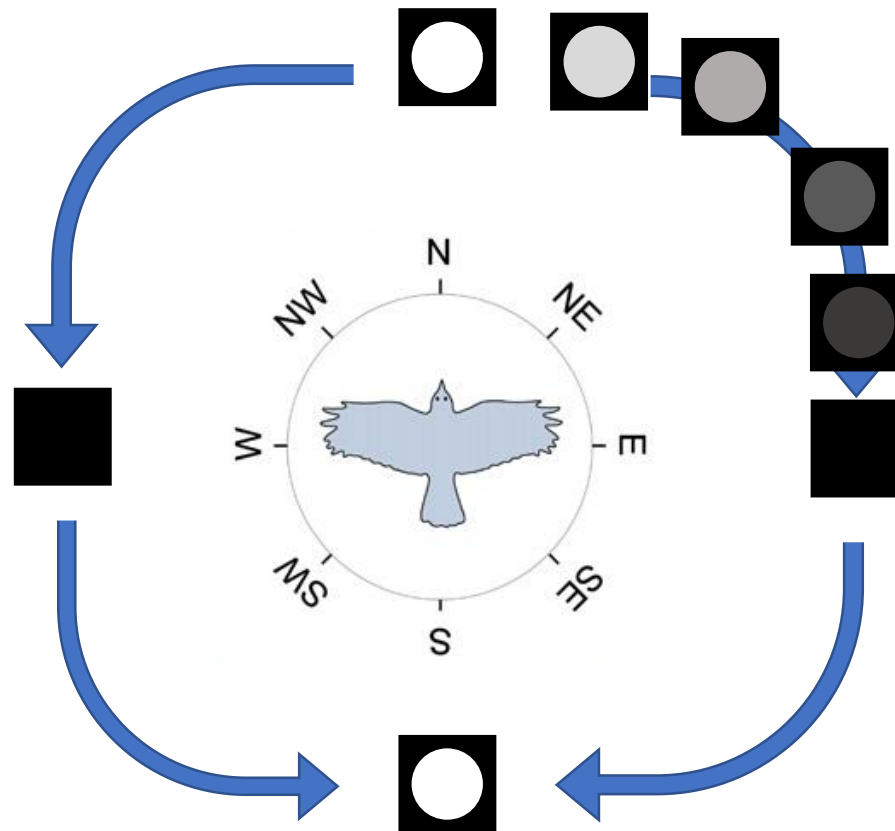
Jiansheng Wu,
Assistance professor
Department of Physics
SUSTech



<https://github.com/LokyWei/Biophysical-model>



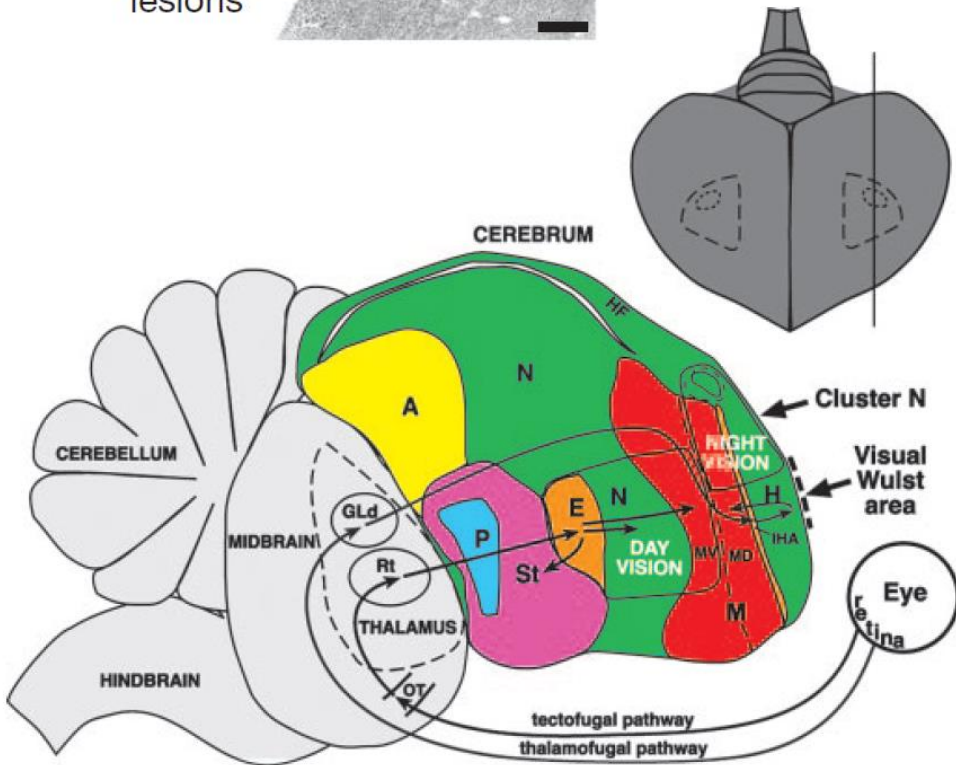
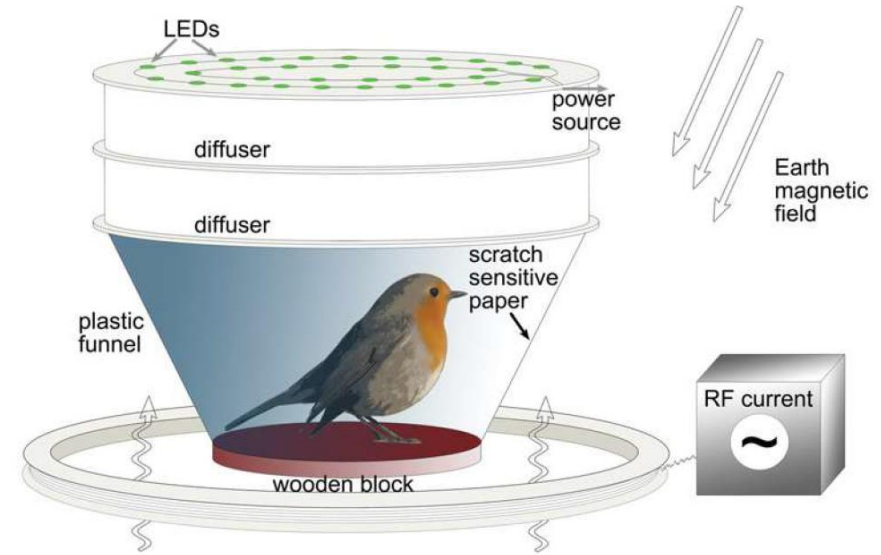
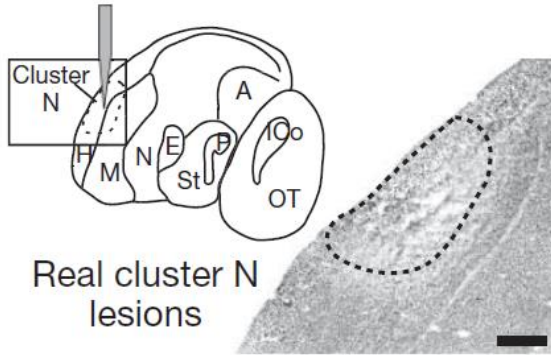
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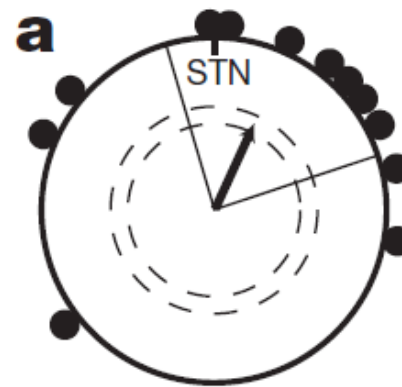
Appendix

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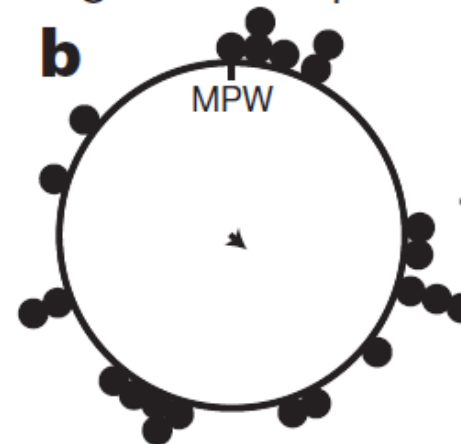
- 1) A neural connection between night vision and magnetic sensing in European robin.



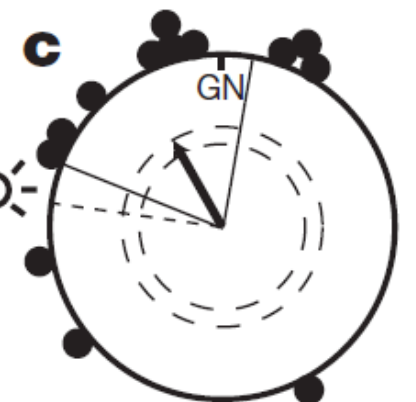
Star compass



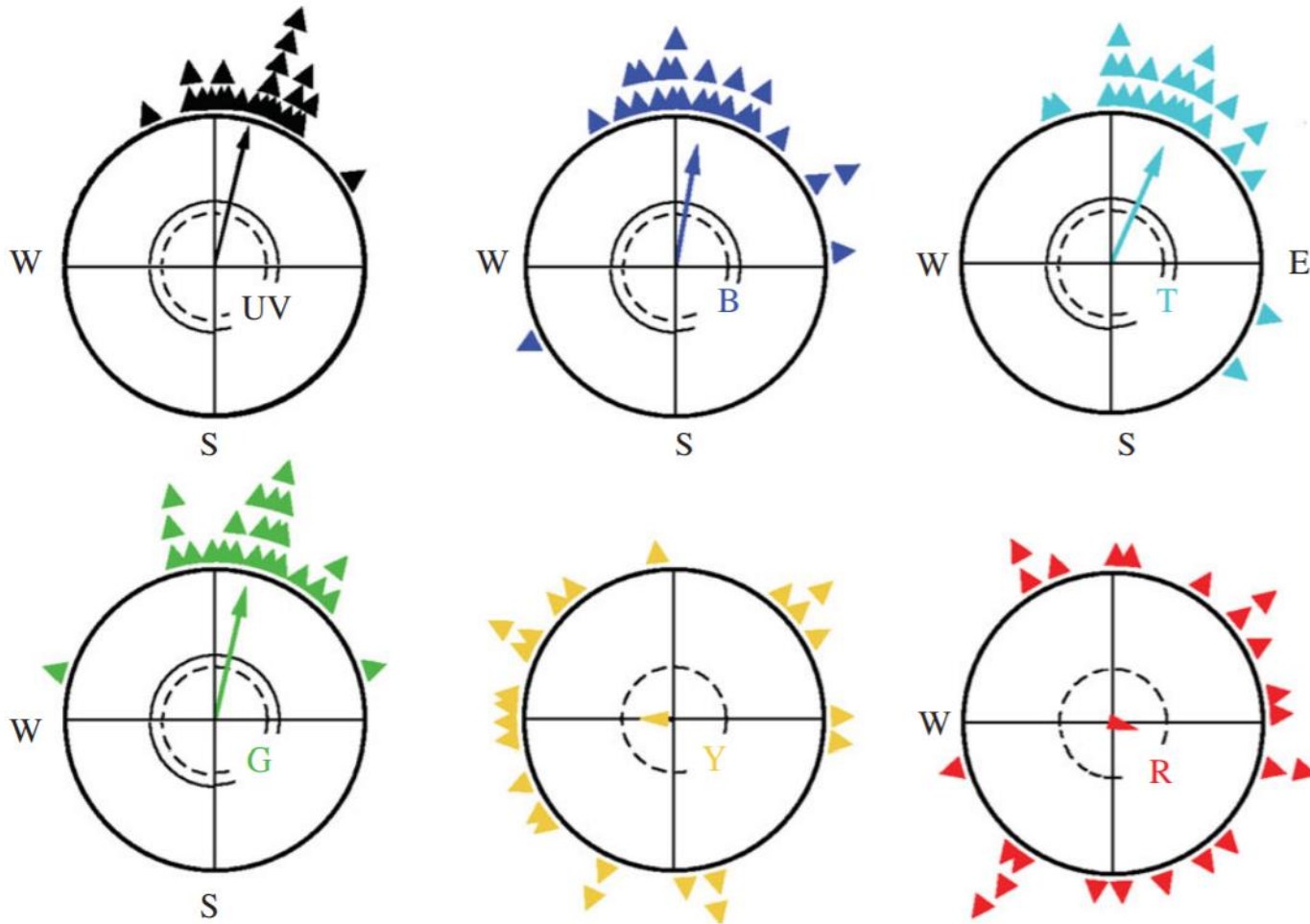
Magnetic compass



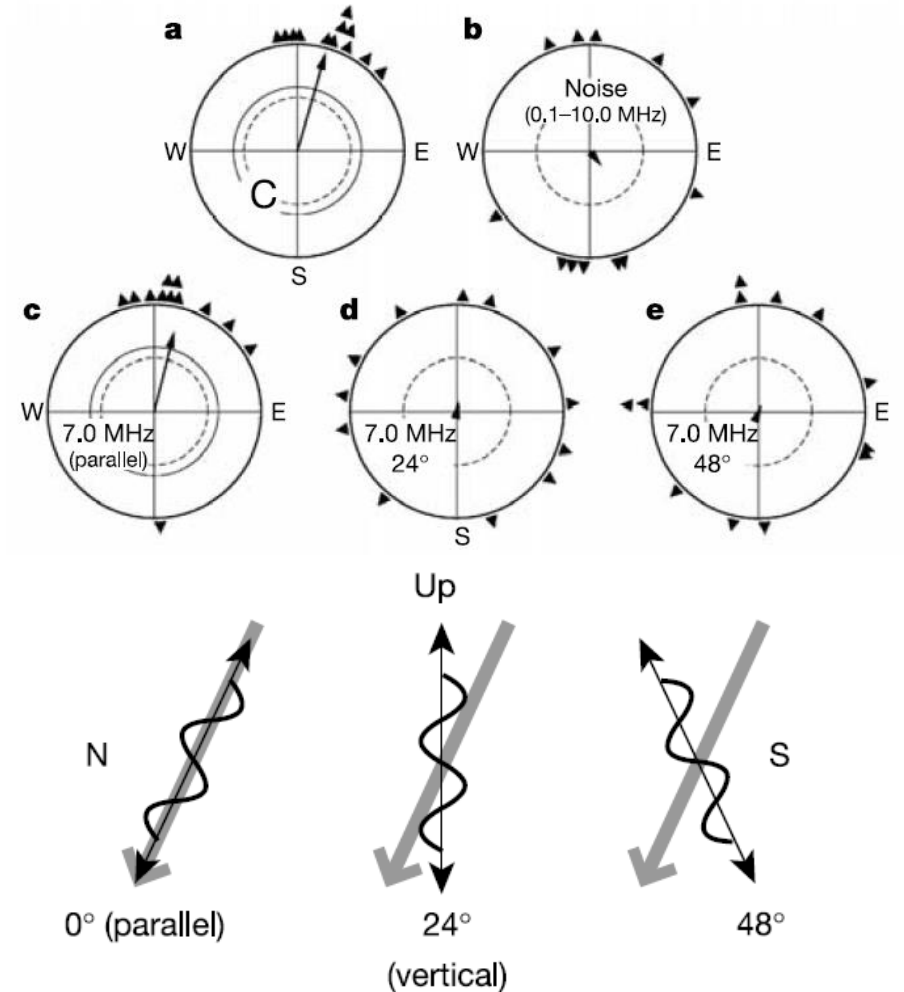
Sunset compass



- 2) Different wavelength of incident light can disrupt magnetic orientation of European robin.

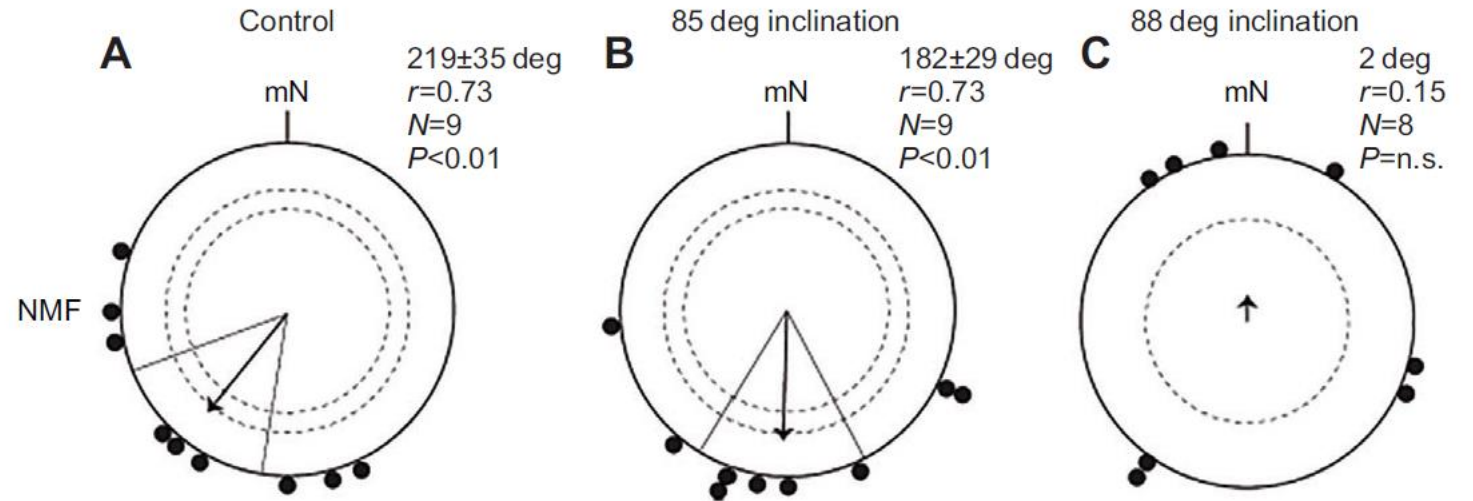
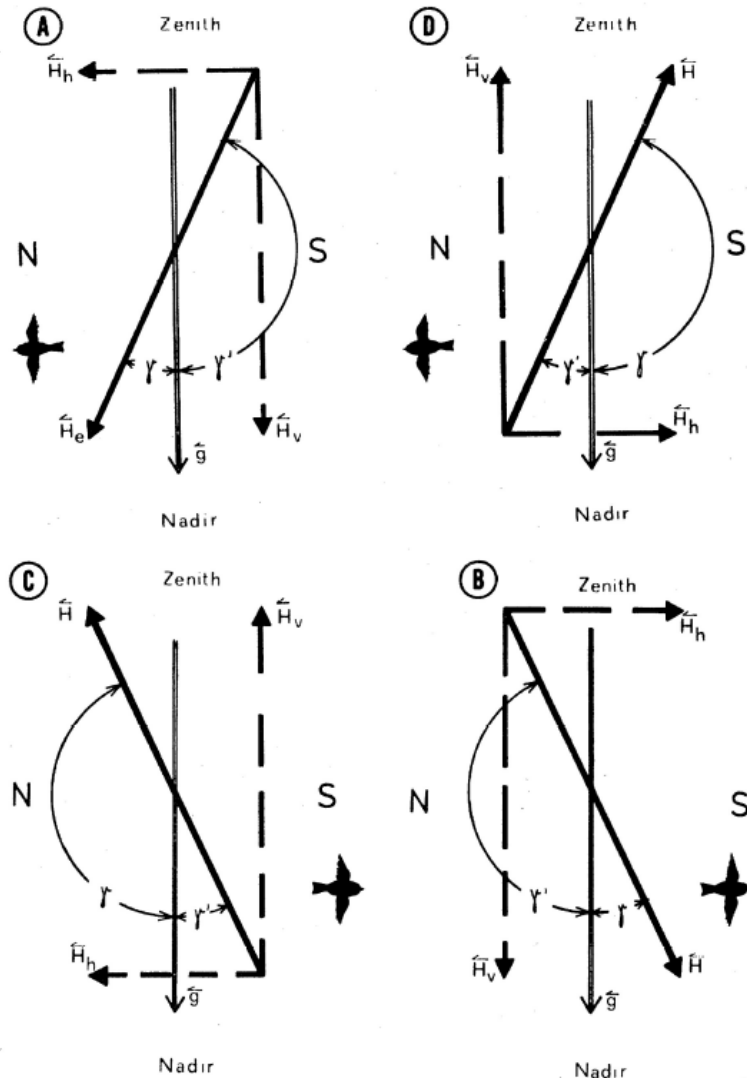


Orientation of European robin under monochromatic light of different wavelengths



Robins were disoriented when exposed to a vertically aligned broadband (0.1–10 MHz) or a single-frequency (7-MHz) field in addition to the geomagnetic field.

- 3) The magnetic compass of European robin is a inclination compass with $<5^\circ$ precision, which can be explained by radical pair mechanism instead of any other hypothesis.

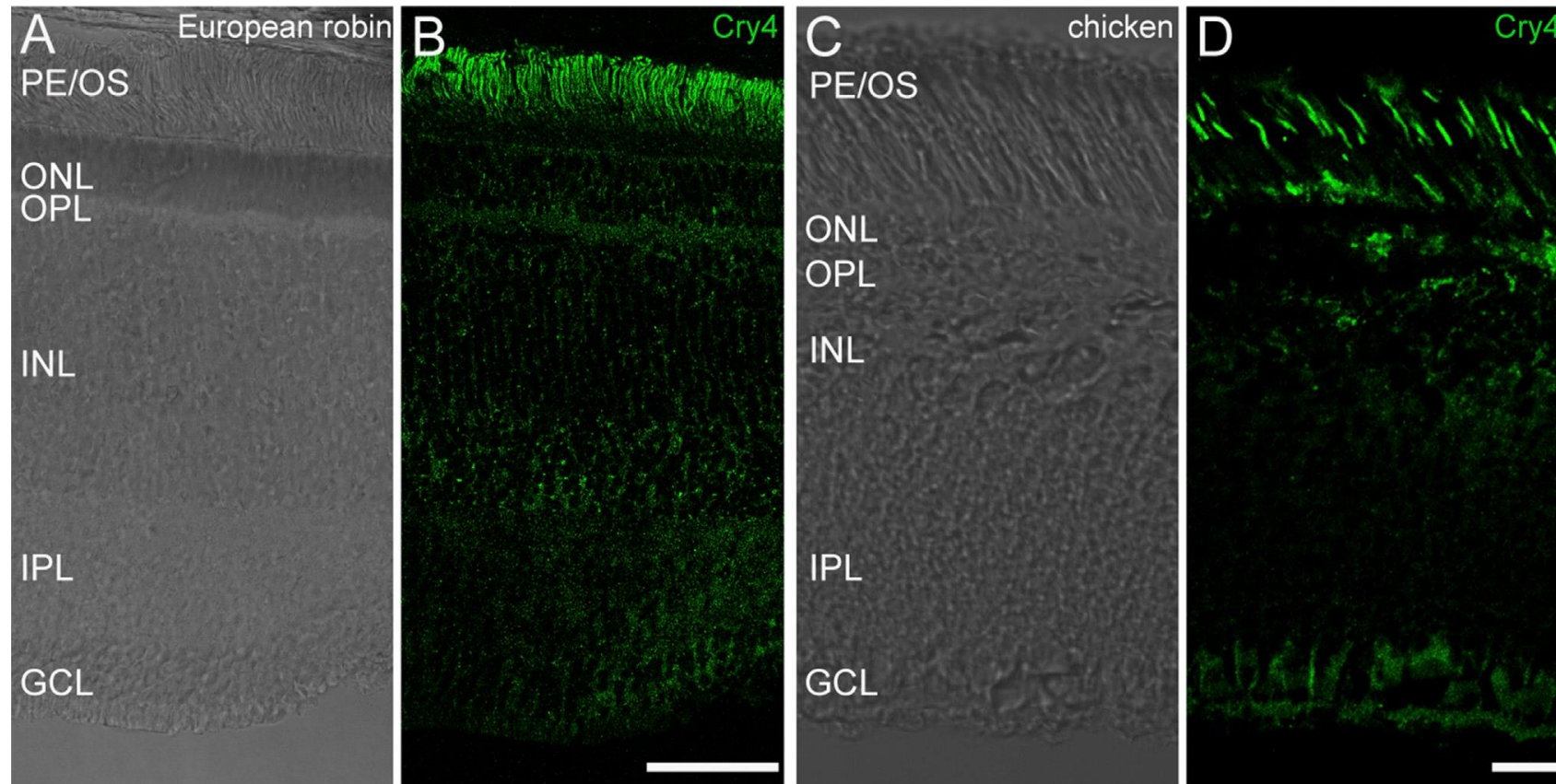


The orientation of individual blackcaps was tested in magnetic fields with 67, 85 or 88 degree inclination

The magnetic compass of European robins **does not use the polarity of magnetic field** for detecting the north direction.

The birds derive their north direction from interpreting the **inclination of the axial direction** of the magnetic field lines in space, where **field lines and gravity vector form the smaller angle**.

- 4) Cry4 protein, a member of cryptochrome family, is found recently located on the outer segment in cones in birds' retina.



Cry4 Is Expressed in the Outer Segments of Specific Photoreceptor Cells in the Retina of European Robin and Chicken