Phototransduction

Inner Life of the Cell https://www.youtube.com/watch?v=wJyUtbn005Y



















glutamate molecule vesicle

Photoactivation:

A photon is absorbed by a visual pigment molecule lying in one of the membranous discs contained in the outer segment.

Biochemical cascade:

In the dark there is a steady movement of positively charged ions (cations) into the outer segment, via ionic channels. The visual pigment molecule, activated by the photon, initiates a cascade of events that ultimately closes these channels.

Electrotonic spread:

Normally, the movement of cations into the outer segment is balanced by the outward movement of cations, mainly through the inner segment. The decrease in inward current creates a net outward current, which makes the interior of the cell even more negative. This hyperpolarization of the cell membrane spreads throughout the cell. This is how the information about light absorption spreads to the synaptic terminal.

Steps in phototransduction

Synaptic deactivation:

At the synaptic terminal there are calcium channels that open when the voltage across the cell membrane depolarizes and close when it hyperpolarizes. Thus the hyperpolarization of the cell membrane leads to a decrease in the rate of entry of calcium ions. Free calcium ions are continuously being removed from the cell interior, so a decrease in the rate of entry of calcium leads to a decrease in the internal concentration of free calcium ion.

Decrease in glutamate release:

The synaptic terminal contains vesicles that in turn contain glutamate molecules. In the presence of calcium ions, they are continuously released into the synaptic cleft. Thus a decrease in the internal concentration of calcium ions leads to a decrease in the rate of release of glutamate molecules.



Rhodopsin molecule





Activation of rhodopsin leads to decrease in cGMP concentration



occurs via four intermediate steps

Step I: activated rhodopsin activates G-protein molecules



By means of diffusion, **R*** encounters an inactive G molecule.

The G_{α} part of the G molecule comes to lie over the exposed surface of the activated rhodopsin molecule.

As a result of interacting with \mathbf{R}^* , the GDP molecule held by the G_{α} portion is replaced by a GTP molecule, which converts this subunit to an activated form.

Activation of the G_{α} subunit, now G_{α}^{*} , causes it to separate from both the rhodopsin molecule and the $G_{\beta\gamma}$ portion of the G molecule.

A single activated rhodopsin molecule activates 700 G-protein molecules within 100 ms



Step 2: activated G-protein molecules bind to phosphodiesterase (PDE), exposing catalytic site



Step 3: activated PDE breaks bonds in cGMP, thus converting cGMP to GMP and lowering overall cGMP concentration



If G^*_{α} molecules attach to both ends of a PDE molecule, then two catalytic sites will be activated.

Step 4: decrease in cGMP concentration closes channels



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probability of site filled	J = 0.162





Photocurrent

Inward current in outer segment decreases by 0.7 pA in response to one photon of light



Photovoltage

Charge imbalance created by photocurrent from one photon of light leads to 1 mV hyperpolarization



Synaptic terminal contains 'ribbons' that facilitate migration of vesicles to membrane



Neurotransmitter release



Spontaneous isomerizations determine lower limit of light detection, or visual threshold



starlight

10⁻⁵R*/rod/integration time



daylight



Energy expenditure as a function of light level

