

# **Cell Signaling: signaling molecules and their receptors**

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# Learning Objectives

The students will learn:

- Intracellular signaling pathway
- Intercellular signaling pathways
- Extracellular signaling molecules and receptors
- Intracellular signaling molecules and receptors

# CELL SIGNALING

- **Cell signaling** is part of a **complex system** of communication that governs basic cellular activities and coordinates cell actions.
- The ability of cells to perceive and correctly respond to their microenvironment is the basis of development, tissue repair, and immunity as well as normal tissue homeostasis
- Errors in cellular information processing are responsible for diseases such as cancer, autoimmunity, and diabetes.
- By understanding cell signaling, diseases may be treated effectively and, theoretically, artificial tissues may be created.

# Definitions

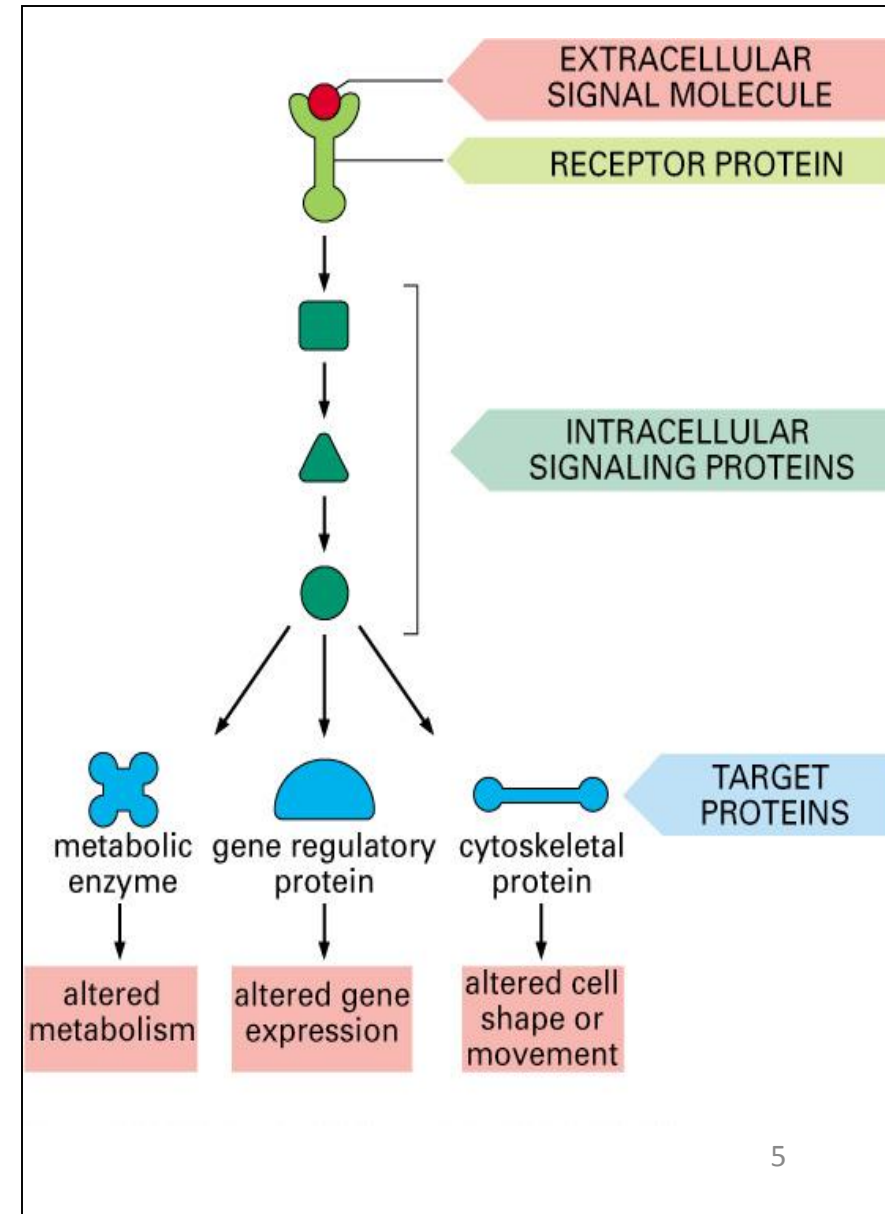
- *Signaling*: Cell-cell communication via signals.
- *Signal transduction*: Process of converting extracellular signals into intra-cellular responses.
- *Ligand*: The signaling molecule.
- *Receptors*: Bind specific ligands and in turn activate one or more intracellular pathways. These pathways depend on intracellular signaling proteins which process the signal and transmit the signal to appropriate intracellular targets. The targets at the end of signaling pathways are called effector proteins.

# A simple intracellular signaling pathway activated by an extracellular signal molecule

The signal molecule usually binds to a receptor protein in the PM of the target cell.

The receptor activates one or more intracellular signaling pathways, involving a series of signaling proteins.

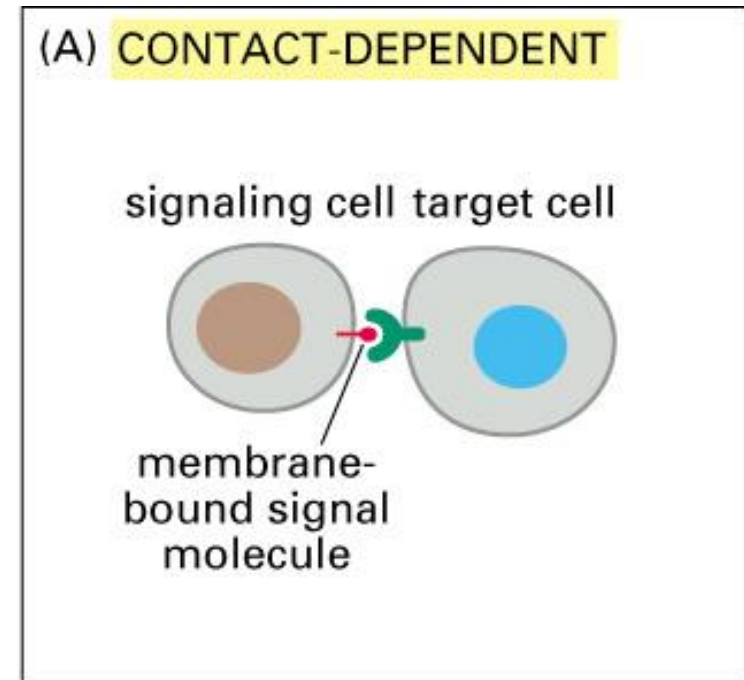
Finally, one or more of the intracellular signaling proteins alter the activity of effector proteins and thereby the behaviour of the cell.



# Four forms of intercellular signaling

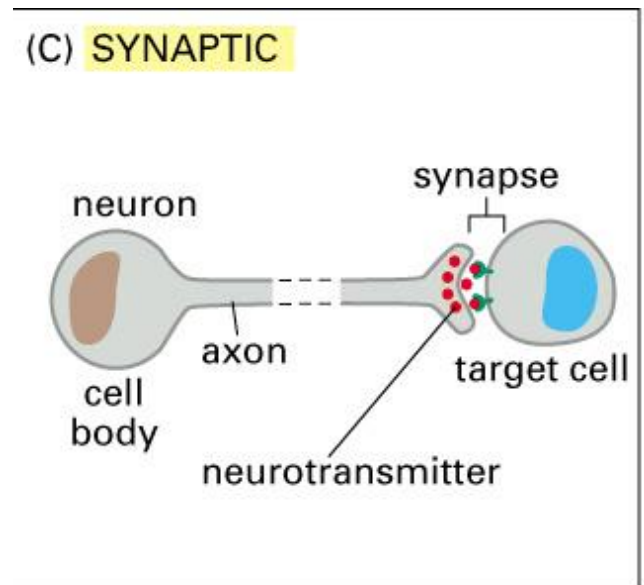
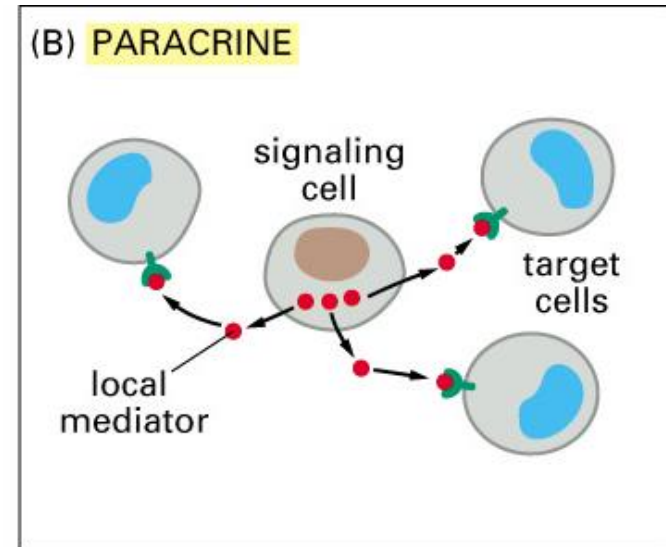
- Cells usually communicate with each other through **extracellular messenger molecules**.

1. **Contact dependent signaling** requires cells to be in direct membrane-membrane contact. This is important during development and in immune responses.



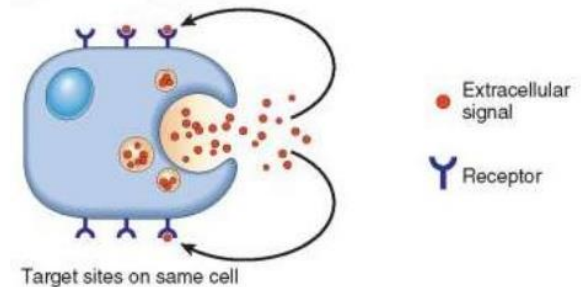
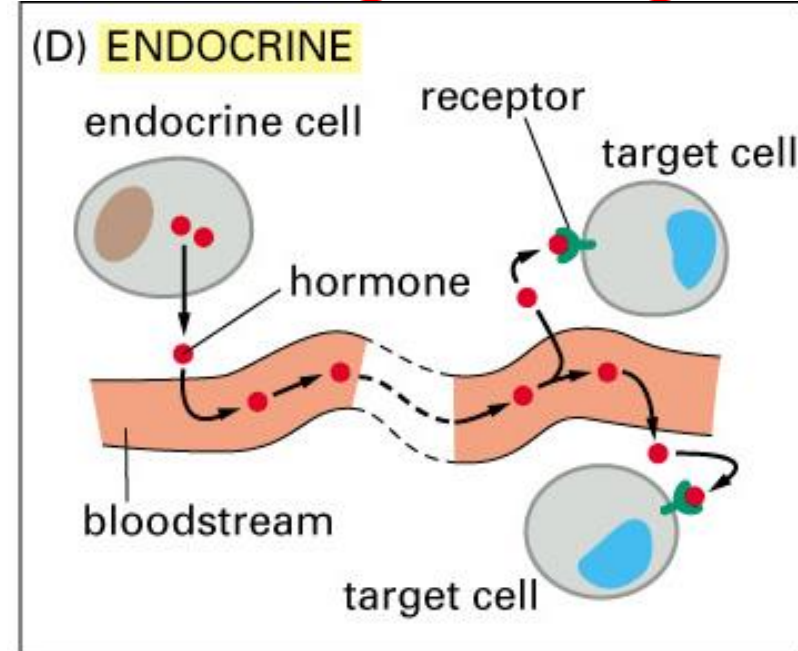
# Four forms of intercellular signaling

- **2. paracrine signaling** depends on local mediators that are released into the extracellular space and act on neighbouring cells. E.g. nerve-muscle
- **3. synaptic signaling** is performed by neurons that transmit signals electrically along their axons and release neurotransmitters at synapses.



# Four forms of intercellular signaling

- **4. endocrine signaling** depends on endocrine cells, which secrete hormones into the bloodstream for distribution throughout the body
- **Autocrine signaling**- cell that releases the signal is also the target.



Autocrine Signalling



Each cell is programmed to respond to specific combinations of extracellular signals

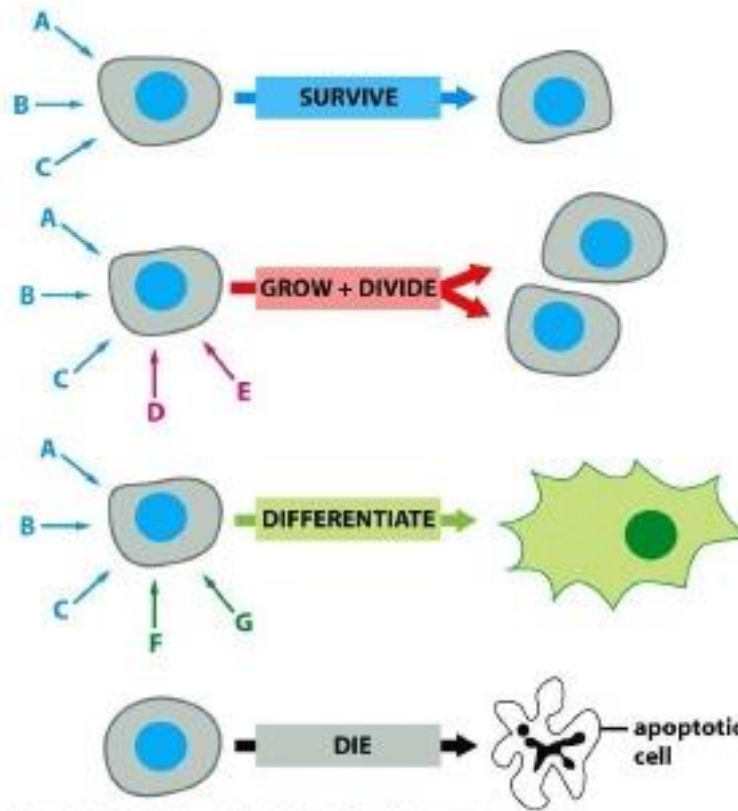


Figure 15-8 Molecular Biology of the Cell 5/e (© Garland Science 2008)

A cell may require multiple signals (A,B,C) to survive. Additional signals to grow and divide (D,E) or differentiate (F,G). If appropriate survival signals are deprived off, the cell undergoes apoptosis.

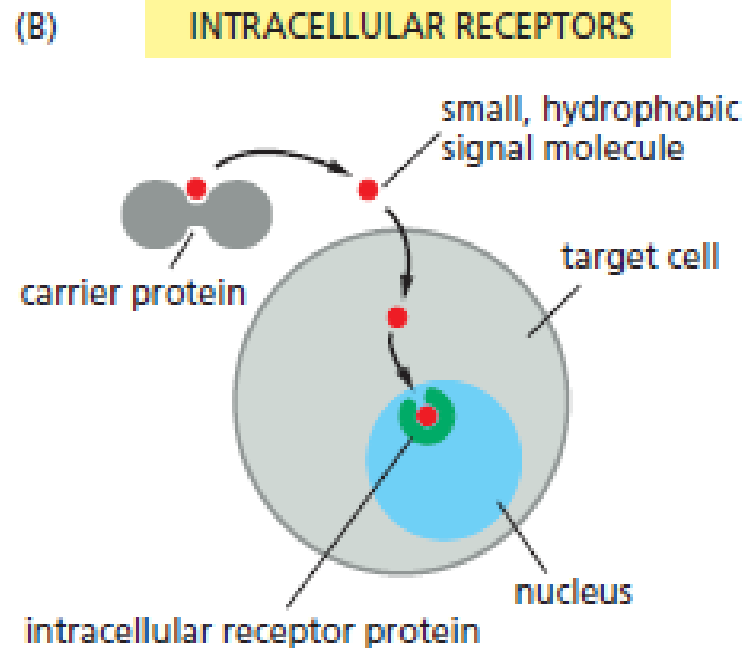
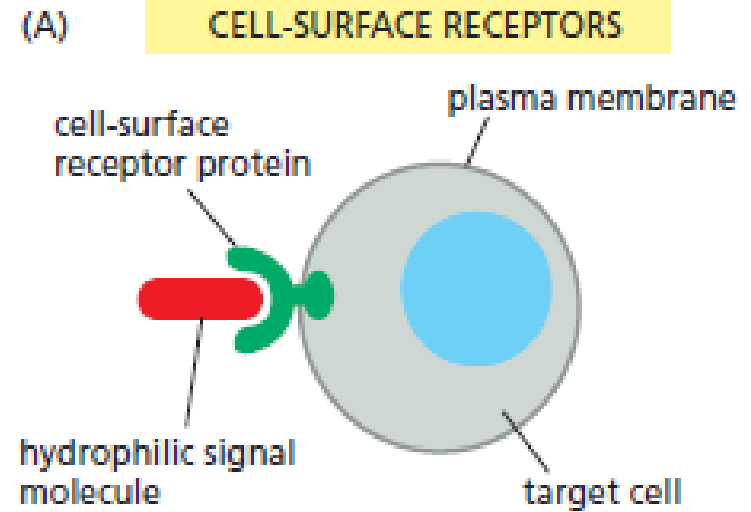
# Extracellular signal molecules bind to specific receptors

- Extracellular signal molecules include proteins, small peptides, amino acids, nucleotides, steroids, retinoids, fatty acid derivatives, NO, CO.
- Target cells respond by means of receptors.
- Receptors are of two types:

# Receptors

1. mostly, receptors are **transmembrane proteins** on the **target-cell surface**. When they bind to Extra cellular molecule (a ligand), and act as signal transducer, they become activated and generate various intracellular signals that alter the behaviour of cell.

2. **Intracellular receptors**-the signal molecule has to be small to diffuse across the PM and bind to receptor proteins inside the target cell-either in the cytosol or nucleus.



# Types of Signaling ligands

Produced by signaling cells, bind to receptors in target cells, act as chemical signals

1. Small Hydrophobic Ligands
2. Other Ligands
3. Water-Soluble Ligands

# Small hydrophobic ligands

- Small hydrophobic ligands can directly diffuse through the plasma membrane and interact with internal receptors. Important members of this class of ligands are the steroid hormones.

# Other Ligands

- Nitric oxide (NO) is a gas that also acts as a ligand. It is able to diffuse directly across the plasma membrane; one of its roles is to interact with receptors in smooth muscle and induce relaxation of the tissue. NO has a very short half-life; therefore, it only functions over short distances.

# Water-soluble ligands

- Water-soluble ligands are polar and, therefore, cannot pass through the plasma membrane unaided. Instead, most water-soluble ligands bind to the extracellular receptors.
- Cell-surface receptors include: **ion-channel, G-protein, and enzyme-linked protein receptors**. The binding of these ligands to these receptors results in a series of cellular changes. These water soluble ligands are quite diverse and include small molecules, peptides, and proteins.

# Signaling molecules

## Types of Signaling Ligands:

### A. Ligands that bind to cell-surface receptors:

1. Neurotransmitters (NT), i.e. norepinephrine, histamine - hydrophilic (charged, polar)
2. Peptide hormones (P), i.e. insulin - can't cross membrane
3. Growth factors (GF),
4. Lipophilic signaling molecules, i.e. prostaglandins

### B. Ligands that bind to intracellular receptors:

lipid soluble hormones that diffuse across the plasma membrane and interact with receptors in the cytosol or nucleus. i.e. steroids, thyroxine, retinoic acid, nitric oxide.

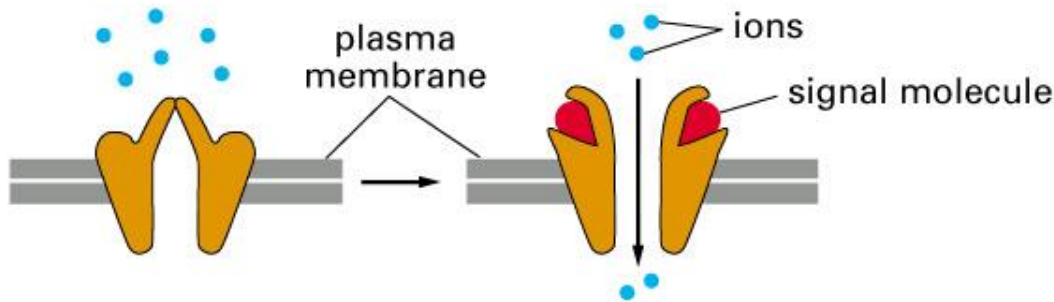


# Cell surface receptors

- **Ion -channel**-linked receptors bind a ligand and open a channel through the membrane that allows specific ions to pass through.
- **G-protein-linked receptors** bind a ligand and activate a membrane protein called a **G-protein**, which then interacts with either an ion channel or an enzyme in the membrane.
- **Enzyme-linked receptors** are cell-surface receptors with intracellular domains that are associated with an enzyme.

# Ion channel-linked receptors

(A) ION-CHANNEL-LINKED RECEPTORS

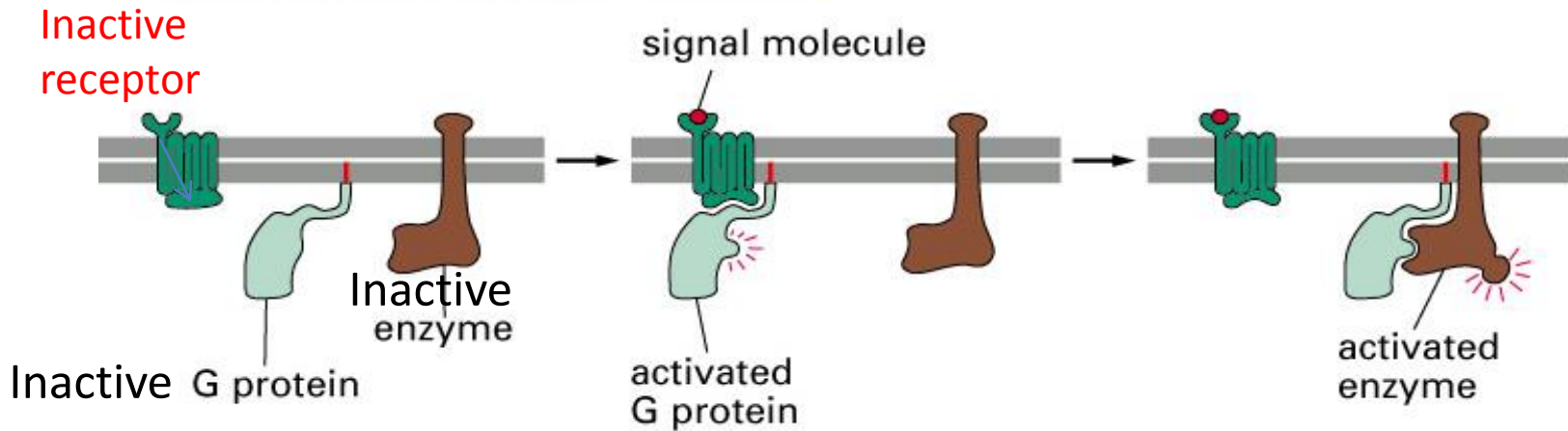


Also called transmitter-gated ion channels or ionotropic receptors. Found in nerve and muscle cells (electrically excitable). This type of signaling is mediated through neurotransmitters.

When a ligand binds to the extracellular region of the ion-channel-linked receptors, there is a conformational change in the receptor protein's structure that allows ions such as sodium, calcium, magnesium, and hydrogen to pass through .

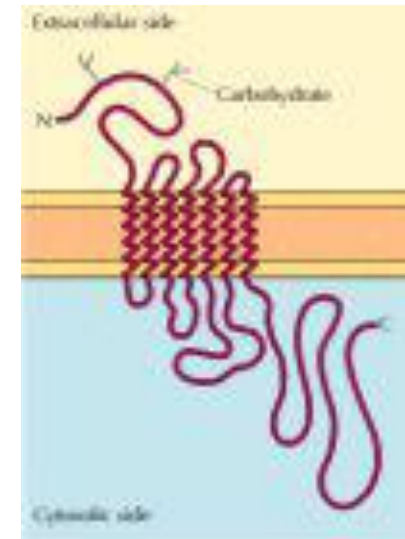
# G-protein-linked receptors

## (B) G-PROTEIN-LINKED RECEPTORS



- The G protein-coupled receptors are characterized by seven membrane-spanning  $\alpha$  helices

- ~45% of all pharmaceutical drugs are known to target GPCRs



# Signal molecules

**Biogenic amines:** Adrenaline, noradrenaline, dopamine,, histamine, acetylcholine

**Amino acids and ions:** Glutamate, Ca<sup>2+</sup>, GABA

**Lipids :** prostaglandins, leukotrienes ( produced in leukocytes by the oxidation of arachidonic acid )

**Peptides / proteins :**

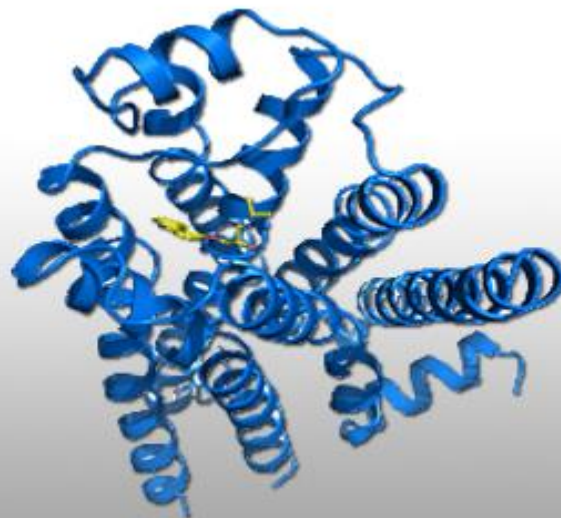
GnRH, angiotensin, bradykinin, thrombin, bombesin, glucagon, calcitonin, vasoactive intestinal peptides, PTH, FSH, LH, TSH

**Nucleotides :** Adenosine nucleotides, adenine nucleotides, uridine nucleotides

**Others :** Light, odorants, pheromones, opiates

# Historical background

- **Robert Lefkowitz** and **Brian Kobilka**: the 2012 Nobel Prize in Chemistry for groundbreaking discoveries that revealed the inner workings of **G-protein-coupled receptors**.



# G-protein-linked receptors

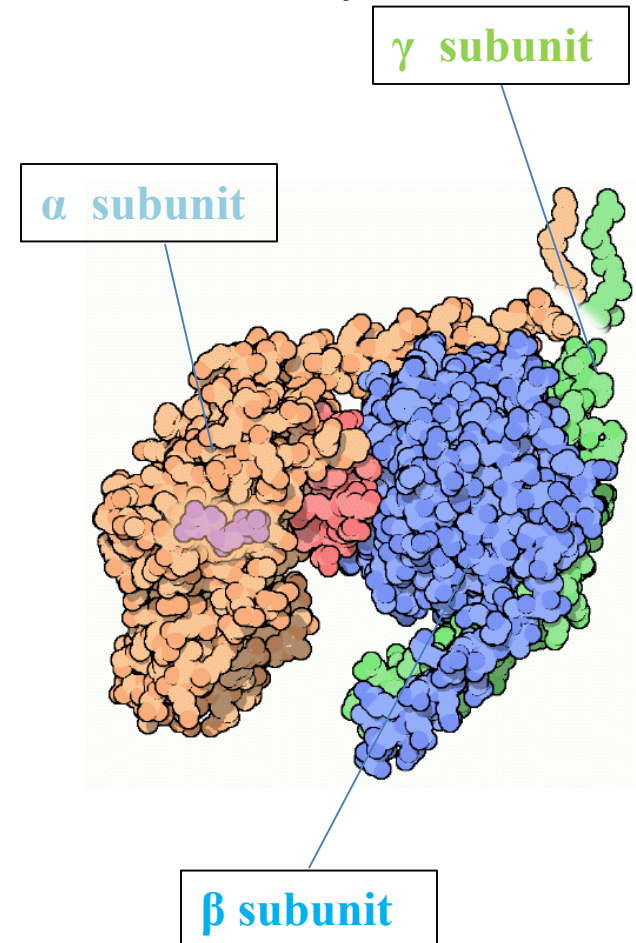
- The binding of ligands to the extracellular domain of these receptors induces a conformational change in the receptor and exposes a binding site for a G protein (bound to the inner face of the plasma membrane). G protein consists of  $\alpha, \beta, \gamma$  subunits.

# heterotrimeric G proteins

➤ Mammalian G protein complexes are made up of

- 20 alpha ( $\alpha$ )
- 6 beta ( $\beta$ )
- 12 gamma ( $\gamma$ ) subunits.

➤ Beta and gamma subunits can form a stable dimeric complex referred to as the beta-gamma complex.

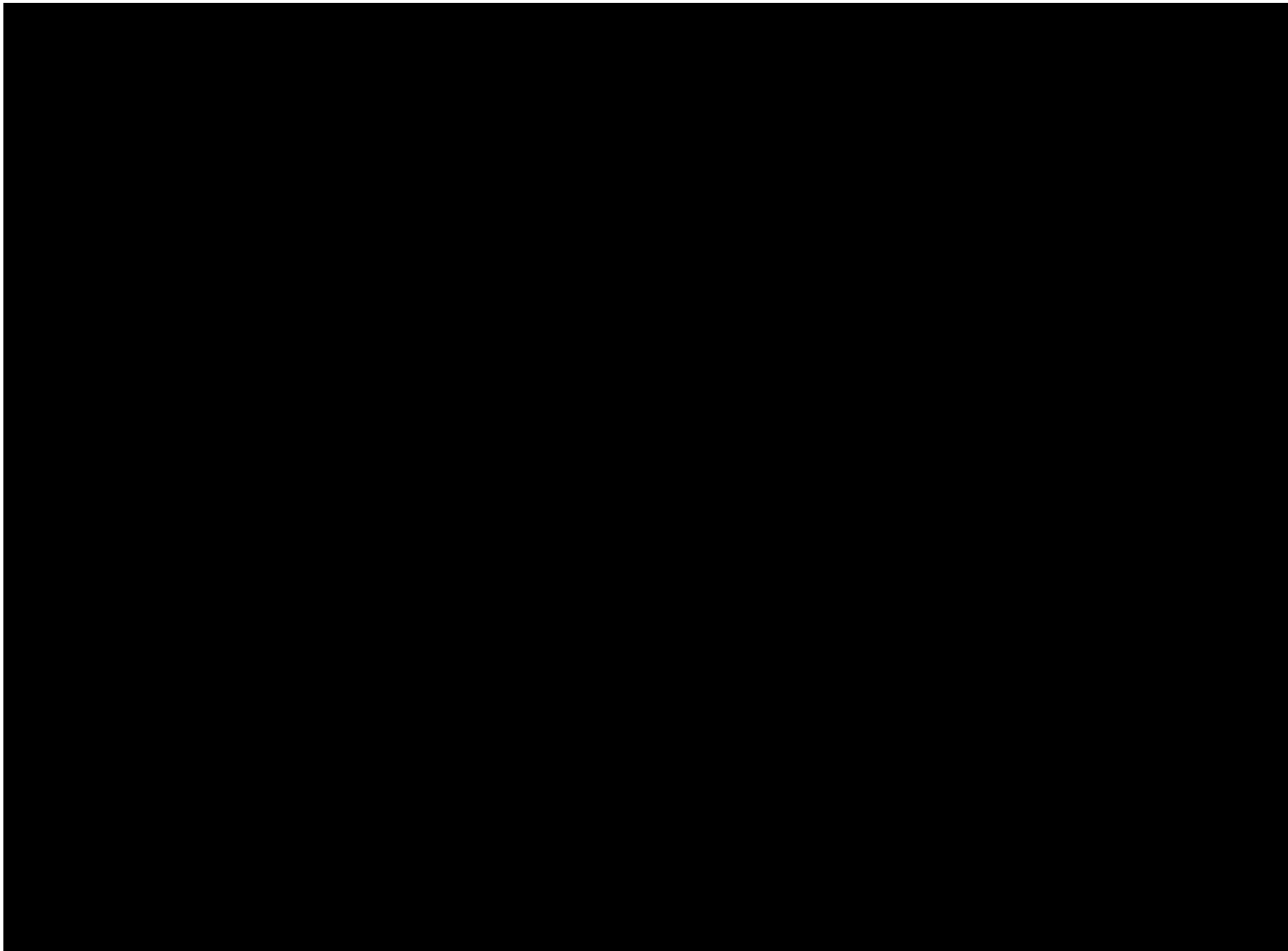


# G-protein-linked receptors

- In the resting state,  $\alpha$  is bound to GDP.
- Ligand binding induces a conformational change in the receptor, such that the cytosolic domain of the receptor interacts with the **G protein** and stimulates the **release of bound GDP from  $\alpha$  subunit** and its **exchange for GTP**.
- The **activated GTP-bound  $\alpha$  subunit** then **dissociates from  $\beta$  and  $\gamma$** , which remain together and function as a  **$\beta\gamma$  complex**.
- **$\alpha$  subunit moves along the inner membrane** and interact with another membrane bound protein 'primary effector' **adenyl cyclase** to elicit an intracellular response.
- The activity of the  $\alpha$  subunit is terminated by **hydrolysis of the bound GTP**, and the inactive  $\alpha$  subunit (now with GDP bound) then reassociates with the  $\beta\gamma$  complex, ready for the cycle to start anew.

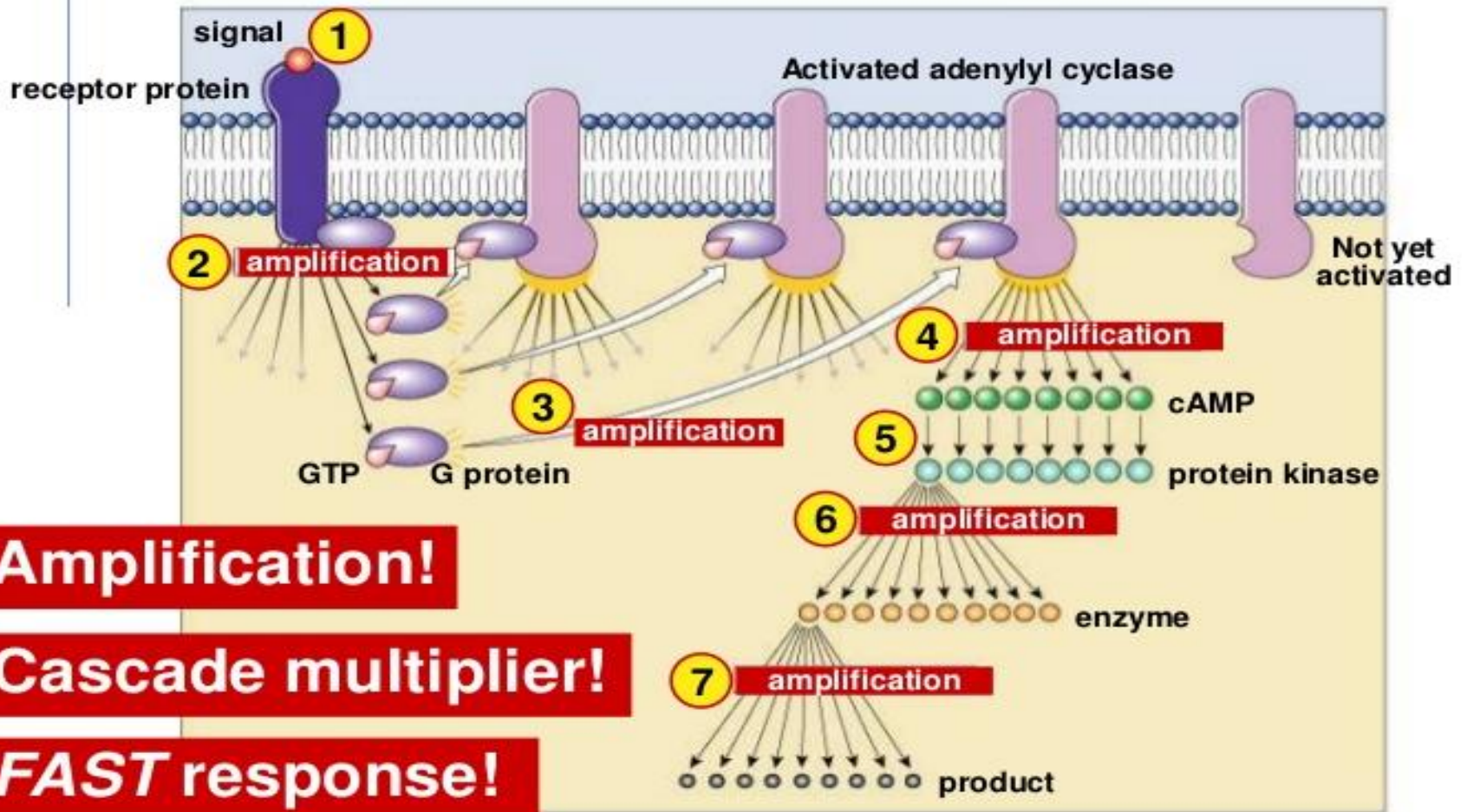


- 'primary effector' creates a second messenger which may activate a 'secondary effector'-protein kinase
- the G protein associated with the epinephrine receptor is called  $G_s$  because its  $\alpha$  subunit *stimulates adenyl cyclase*.



# G<sub>s</sub> cAMP Dependent Pathway

## Signal Amplification through G proteins



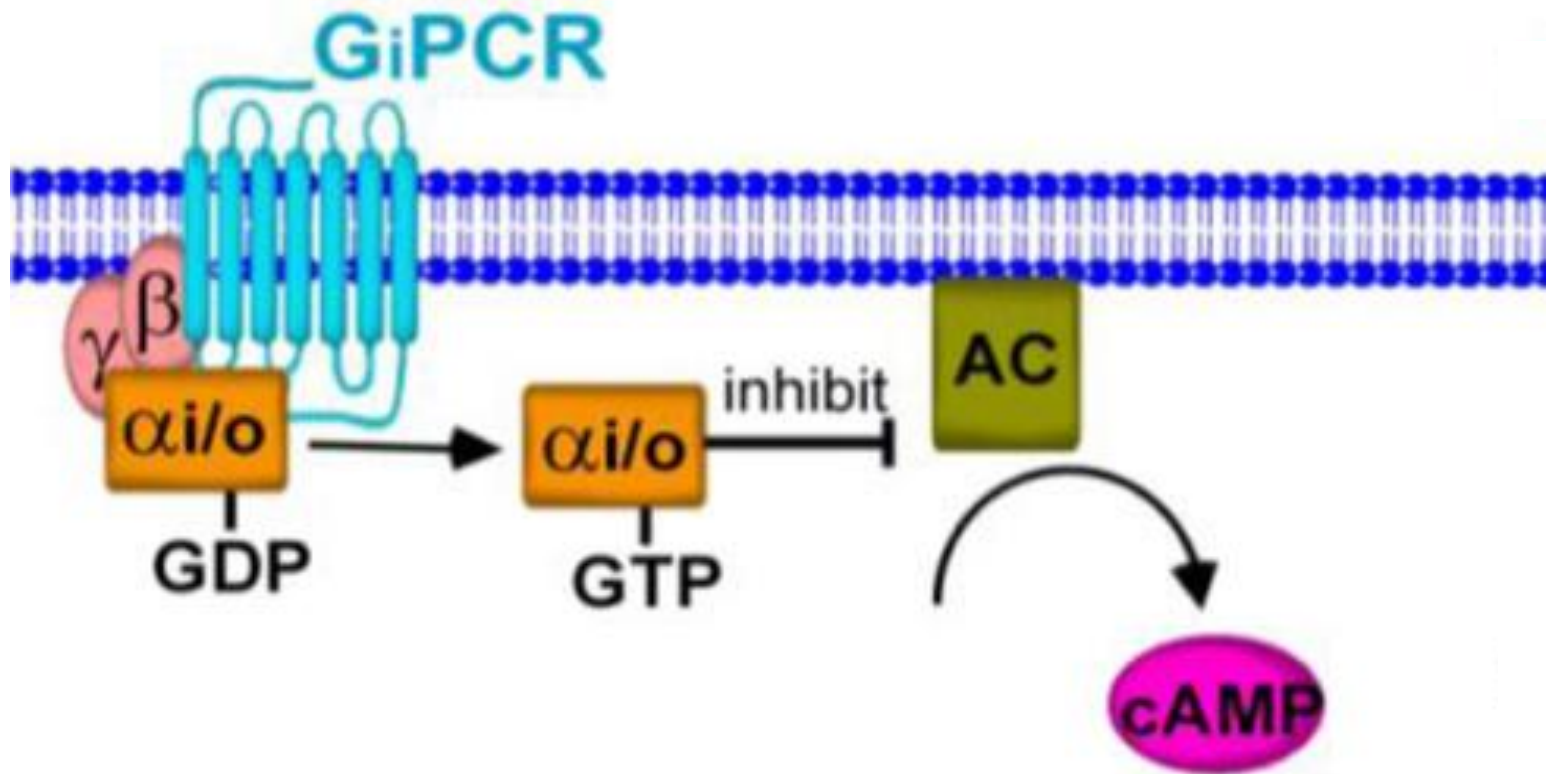
**Amplification!**

**Cascade multiplier!**

**FAST response!**

- In addition to regulating target enzymes, both the  $\alpha$  and  $\beta\gamma$  subunits of some G proteins directly regulate ion channels. Example: action of the neurotransmitter acetylcholine on heart muscle.
- Heart muscle cells have a different acetylcholine receptor, which is G protein-coupled.
- This G protein is designated  $G_i$  because its  $\alpha$  subunit *inhibits adenyl cyclase*. In addition, the  $G_i$   $\beta\gamma$  subunits act directly to open  $K^+$  channels in the plasma membrane, which has the effect of slowing heart muscle contraction.

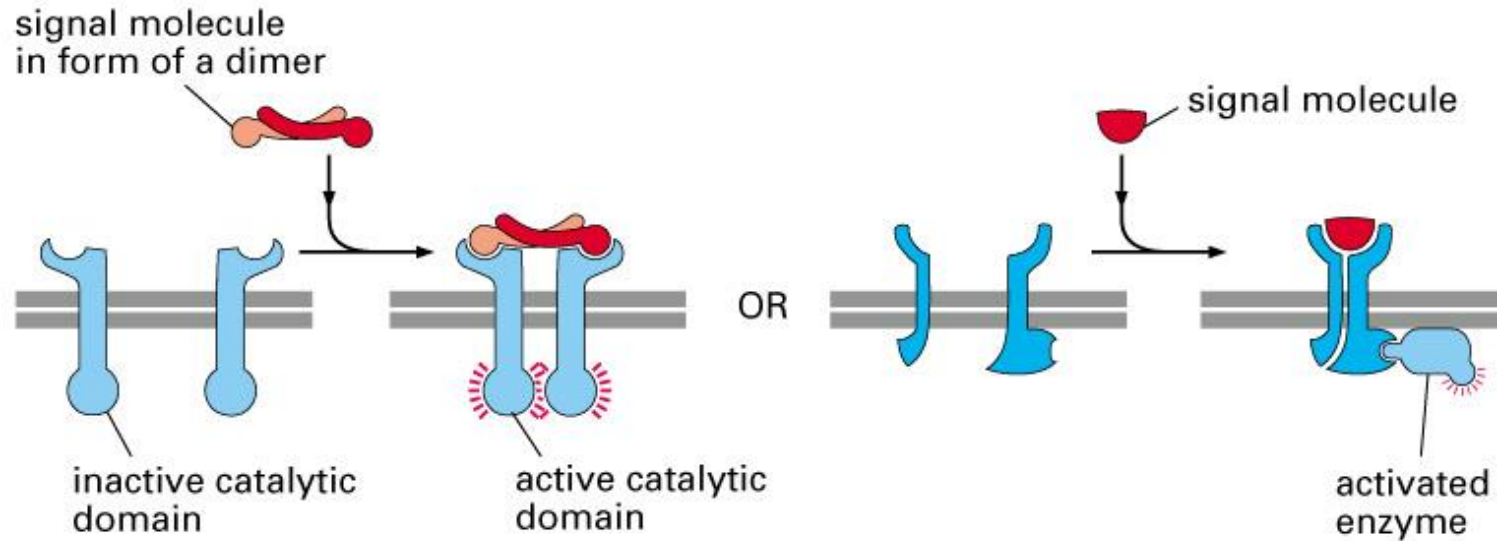
# $G_i$ cAMP Dependent Pathway



Inhibition of AC activity  
Decreased in cAMP level

# Enzyme-linked receptors

## (C) ENZYME-LINKED RECEPTORS



An example of enzyme-linked receptor is the tyrosine [kinase](#) receptor. Signaling molecules bind to the extracellular domain of two nearby tyrosine kinase receptors, which then dimerize. The tyrosine kinase receptor transfers [phosphate](#) groups to tyrosine molecules on the intracellular domain of the receptors and can then transmit the signal to the next messenger within the cytoplasm.

- This family includes the receptors for most polypeptide growth factors, so protein-tyrosine phosphorylation has been particularly well studied as a signaling mechanism involved in the control of animal cell growth and differentiation.

# Intracellular signaling molecules

- These relay signals received by cell surface receptors into the cell interior.
- Some are small molecules often called **2nd messenger** (1st messenger being the extracellular signals). Generated in large nos. in response to receptor activation-
- cAMP and  $\text{Ca}^{++}$  are water soluble and diacylglycerol is lipid-soluble.
- Most intracellular molecules are proteins, which help relay the signal into the cell by either generating 2nd messenger or activating another signaling or **effector protein** in the pathway.
- Many of these proteins behave like molecular switches, mostly activated or deactivated by phosphorylation.



# Second messengers

There are 3 major classes of second messengers:

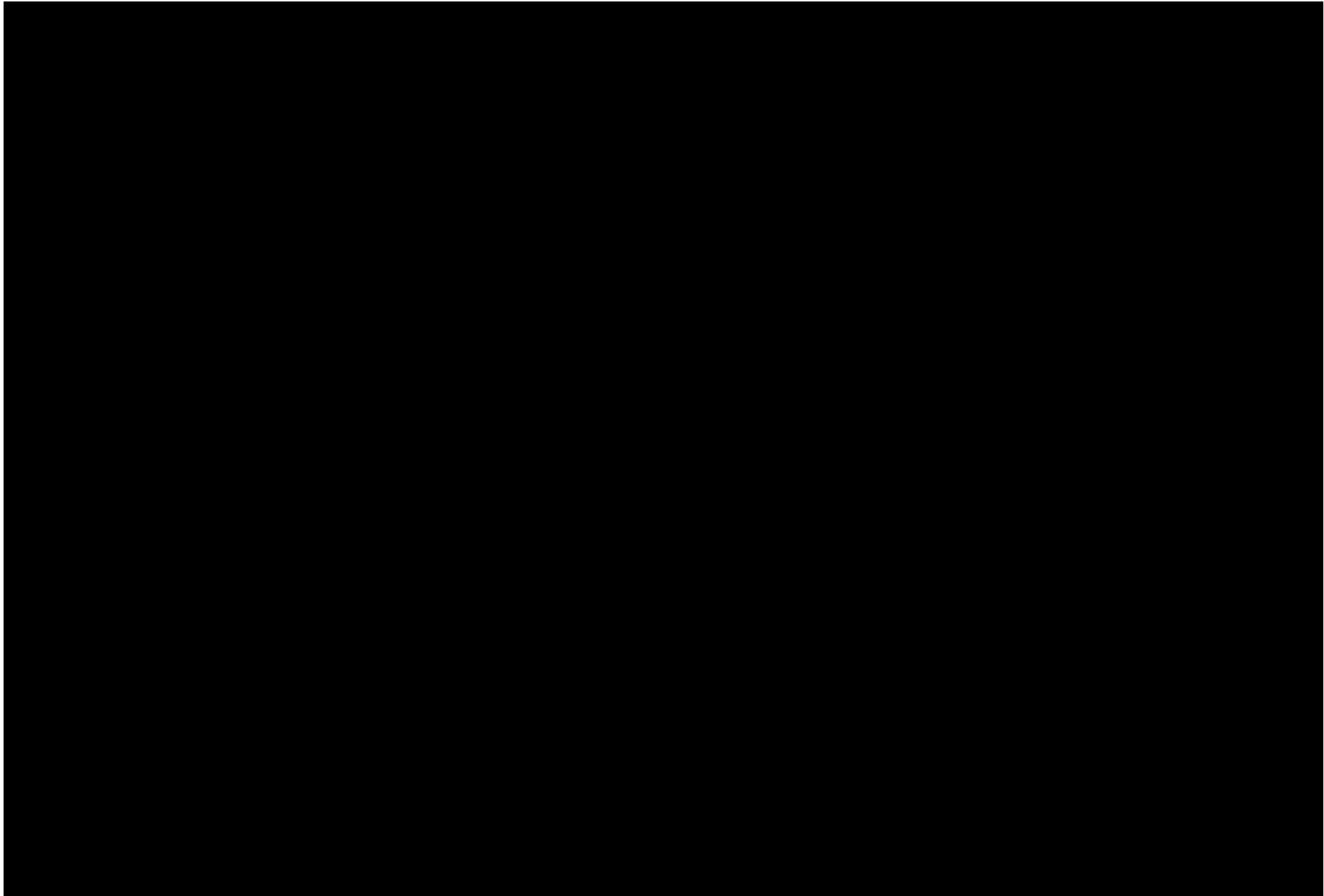
- **cyclic nucleotides** (e.g., **cAMP** and **cGMP**)
- **Inositol trisphosphate** (IP<sub>3</sub>) & **diacylglycerol** (DAG)
- **Calcium ions** (Ca<sup>2+</sup>)

An important feature of the second messenger signaling system is that second messengers may be coupled downstream to multi-cyclic kinase cascades to greatly amplify the strength of the original first messenger signal.

# Second messengers: Cyclic Nucleotides

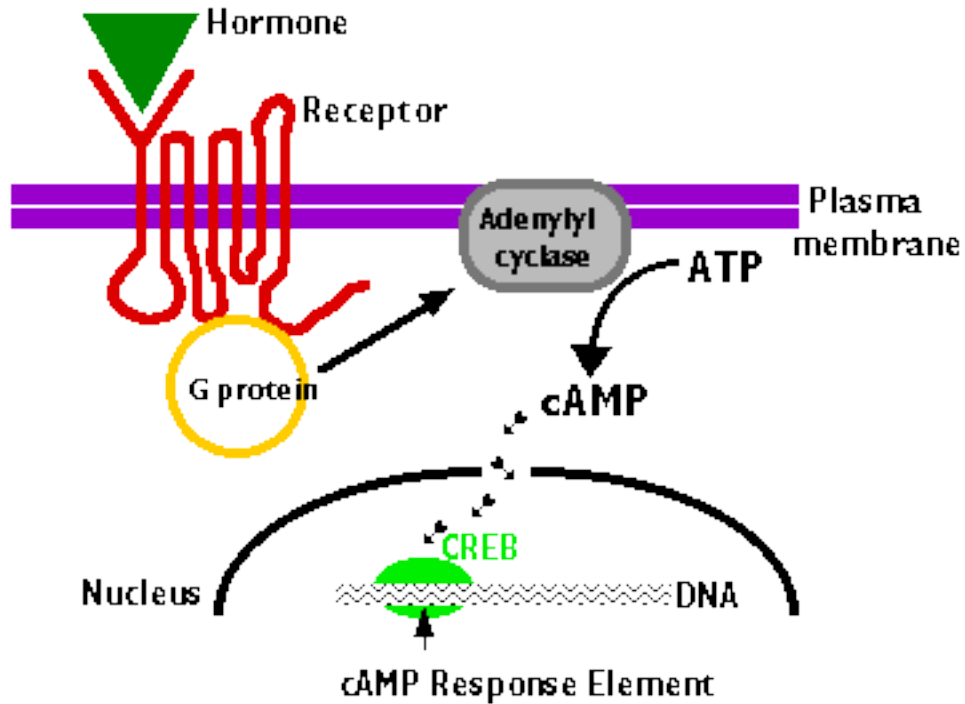
## Cyclic AMP (cAMP)

- Some of the hormones that achieve their effects through cAMP as a second messenger:
- [adrenaline](#)
- [glucagon](#)
- [luteinizing hormone](#) (LH), ACTH, FSH, LH, MSH, PTH
- Cyclic AMP is synthesized from [ATP](#) by the action of the enzyme **adenylyl cyclase**. Binding of the hormone to its receptor activates a [G protein](#) which, in turn, activates adenylyl cyclase.
- The resulting rise in cAMP turns on the appropriate response in the cell by either (or both):
  - changing the molecular activities in the cytosol, often using **Protein Kinase A (PKA)** — a cAMP-dependent protein kinase that phosphorylates target proteins;
  - turning on a new pattern of gene transcription.



# Cyclic Nucleotides

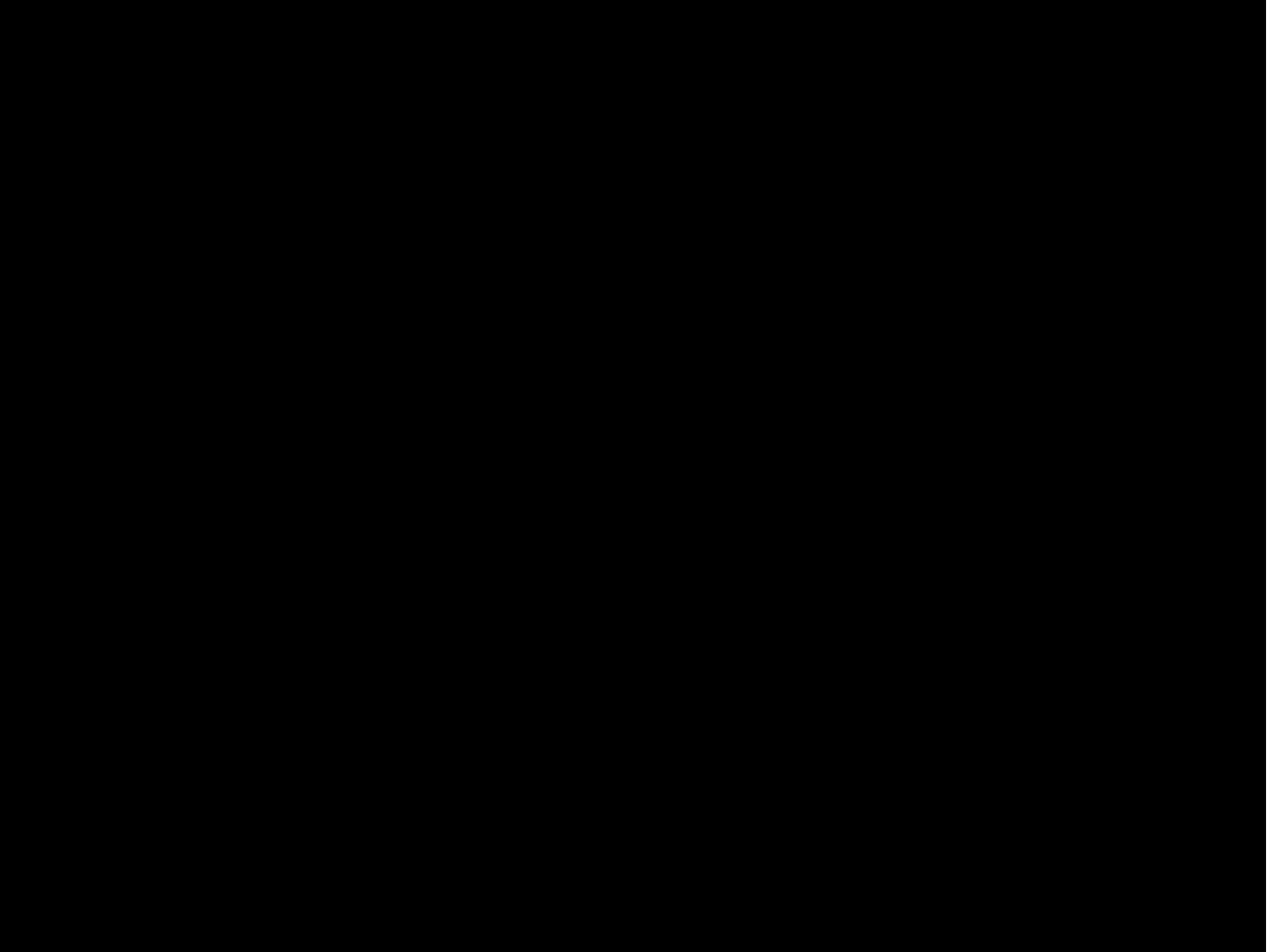
CREB-transcription factor



# Cyclic Nucleotides

## Cyclic GMP (cGMP)

- Cyclic GMP is synthesized from the nucleotide GTP using the enzyme **guanylyl cyclase**. Cyclic GMP serves as the second messenger for
  - atrial natriuretic peptide (ANP)
  - nitric oxide (NO)
- Some of the effects of cGMP are mediated through **Protein Kinase G (PKG)** — a cGMP-dependent protein kinase that phosphorylates target proteins in the cell.



# Second messengers: Inositol trisphosphate & diacylglycerol

Peptide and protein hormones like

- [vasopressin](#),
- [thyroid-stimulating hormone \(TSH\)](#), and
- [Angiotensin](#) and [neurotransmitters](#) like [GABA](#) bind to [G protein-coupled receptors](#) (GPCRs) that activate the intracellular enzyme **phospholipase C (PLC)**. PLC hydrolyzes [phospholipids](#) — specifically phosphatidylinositol-4,5-bisphosphate (PIP<sub>2</sub>) which is found in the inner layer of the plasma membrane. Hydrolysis of PIP<sub>2</sub> yields two products:
- **Diacylglycerol (DAG)** acts via **Protein Kinase C (PKC)** — a calcium-dependent kinase, made available by IP<sub>3</sub>
- **inositol-1,4,5-trisphosphate (IP<sub>3</sub>)**

# Inositol trisphosphate & diacylglycerol

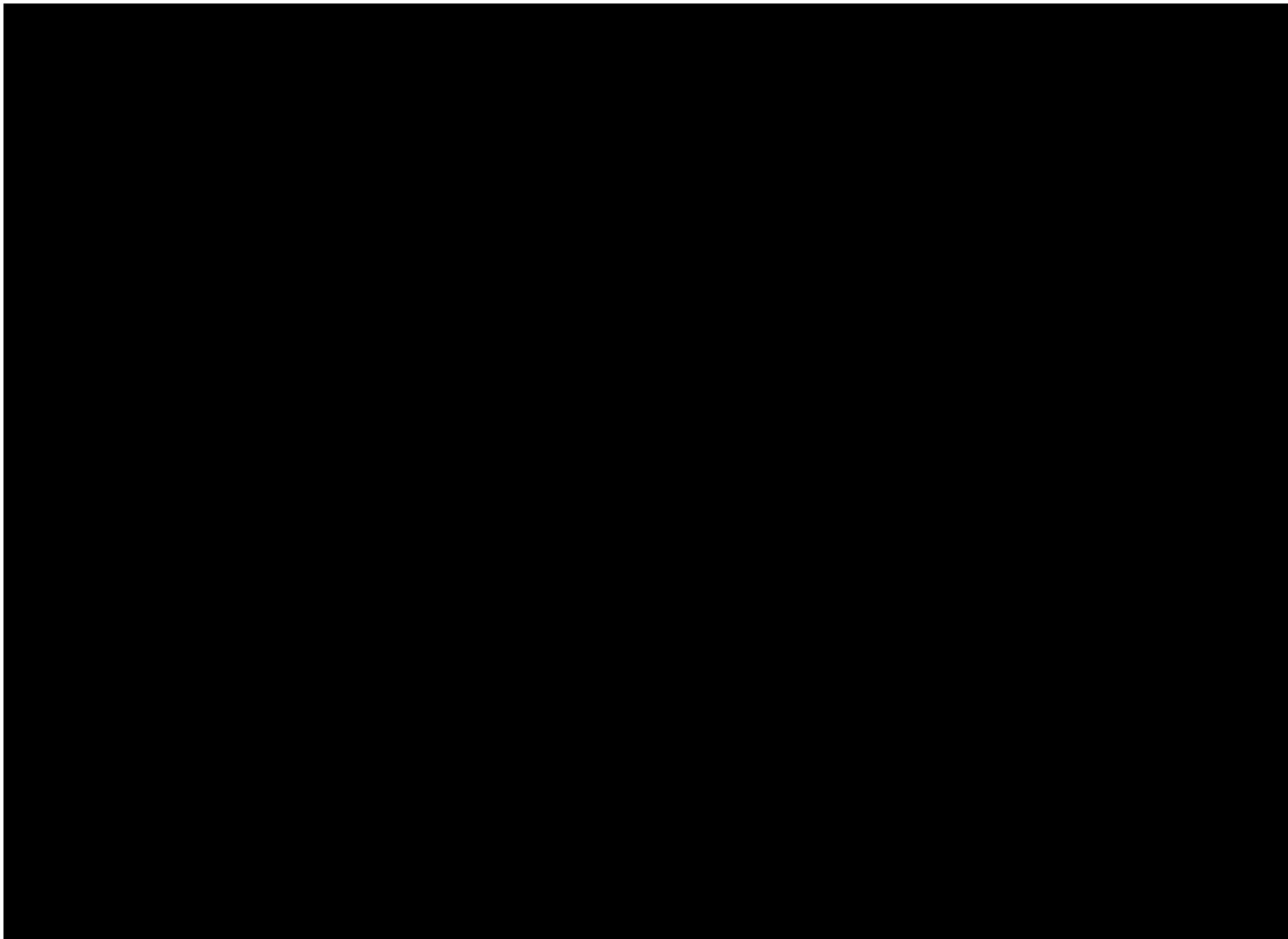
**DAG** remains in the inner layer of the plasma membrane.

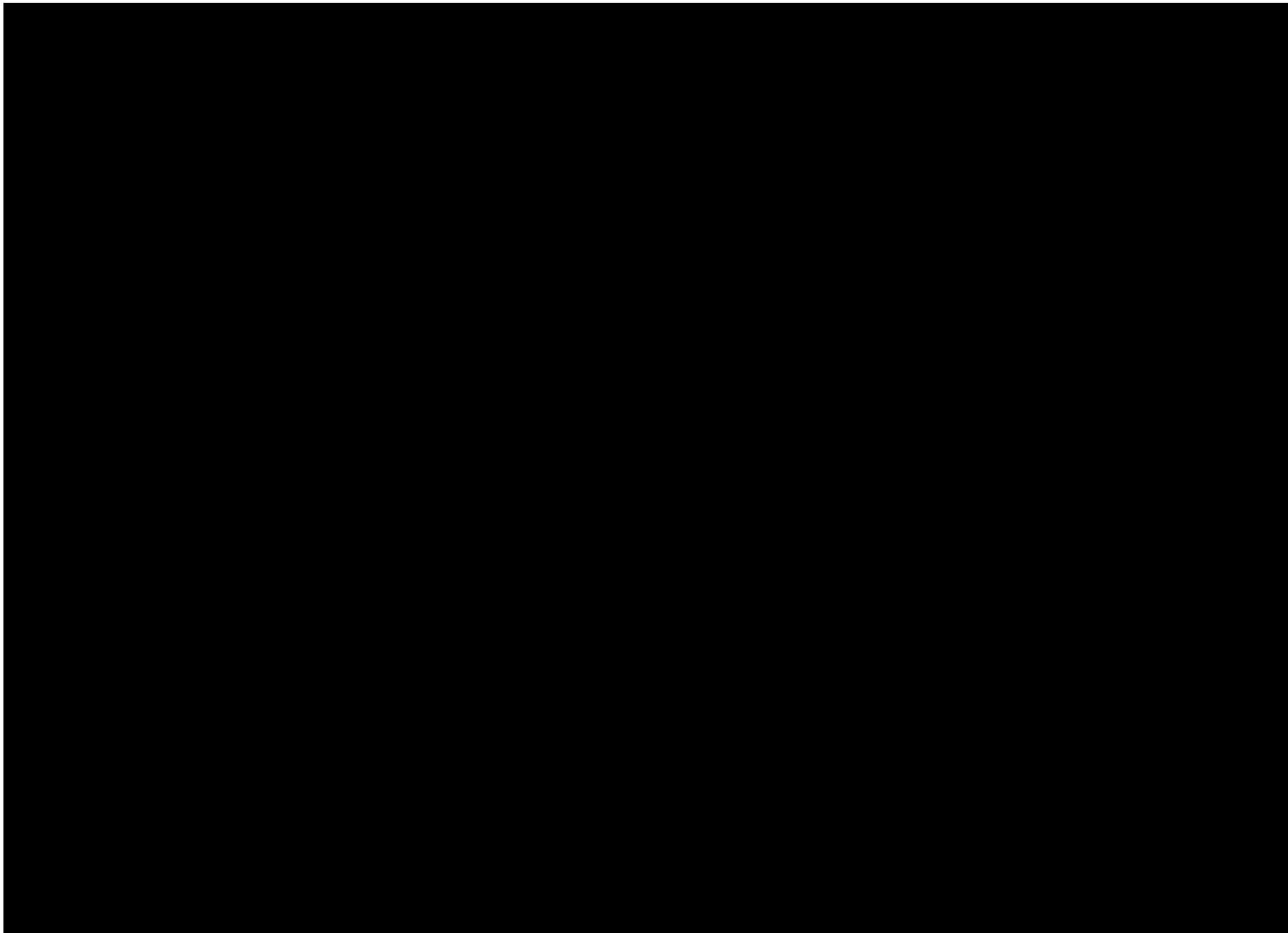
- It recruits **Protein Kinase C (PKC)** — a calcium-dependent kinase that phosphorylates many other proteins that bring about the changes in the cell.  $\text{Ca}^{2+}$  made available by **IP<sub>3</sub>**

**IP<sub>3</sub>**

- diffuses through the cytosol and binds to receptors on the endoplasmic reticulum causing the release of calcium ions ( $\text{Ca}^{2+}$ ) into the cytosol.
- The rise in intracellular calcium triggers the response.







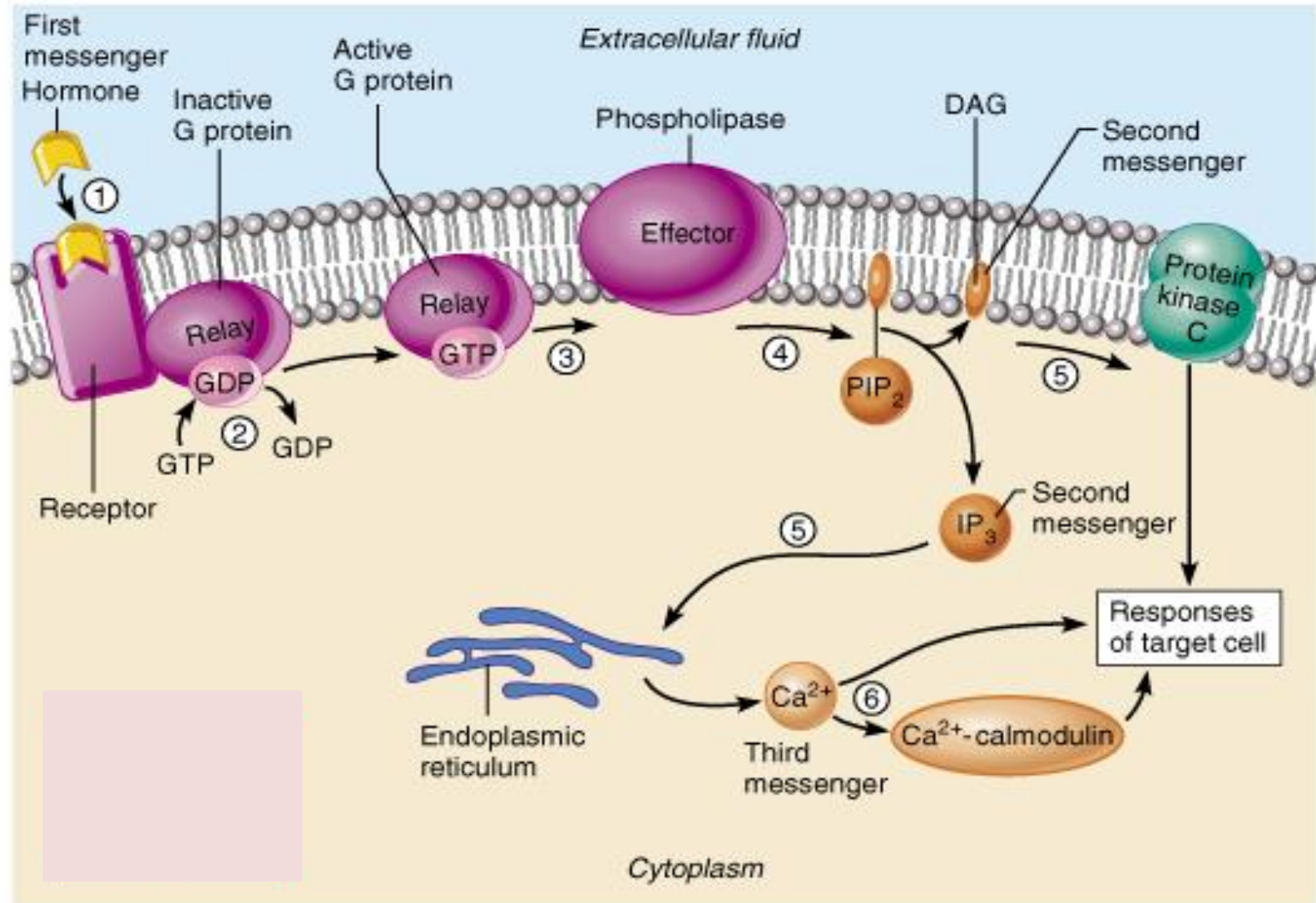
# Second messengers :Calcium ions

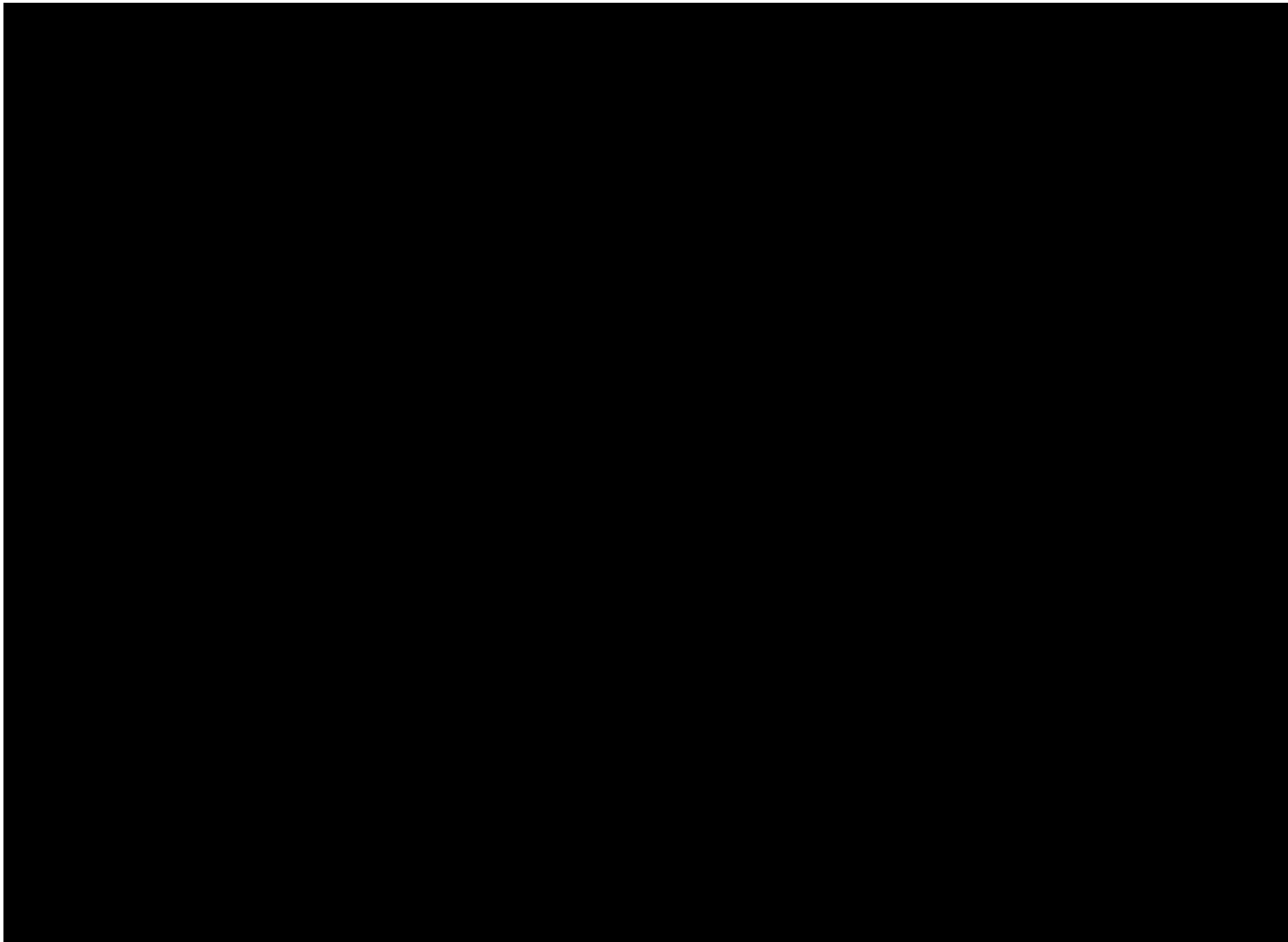
- calcium ions are probably the most widely used intracellular messengers.

In response to many different signals, a rise in the concentration of  $\text{Ca}^{2+}$  in the cytosol triggers many types of events such as

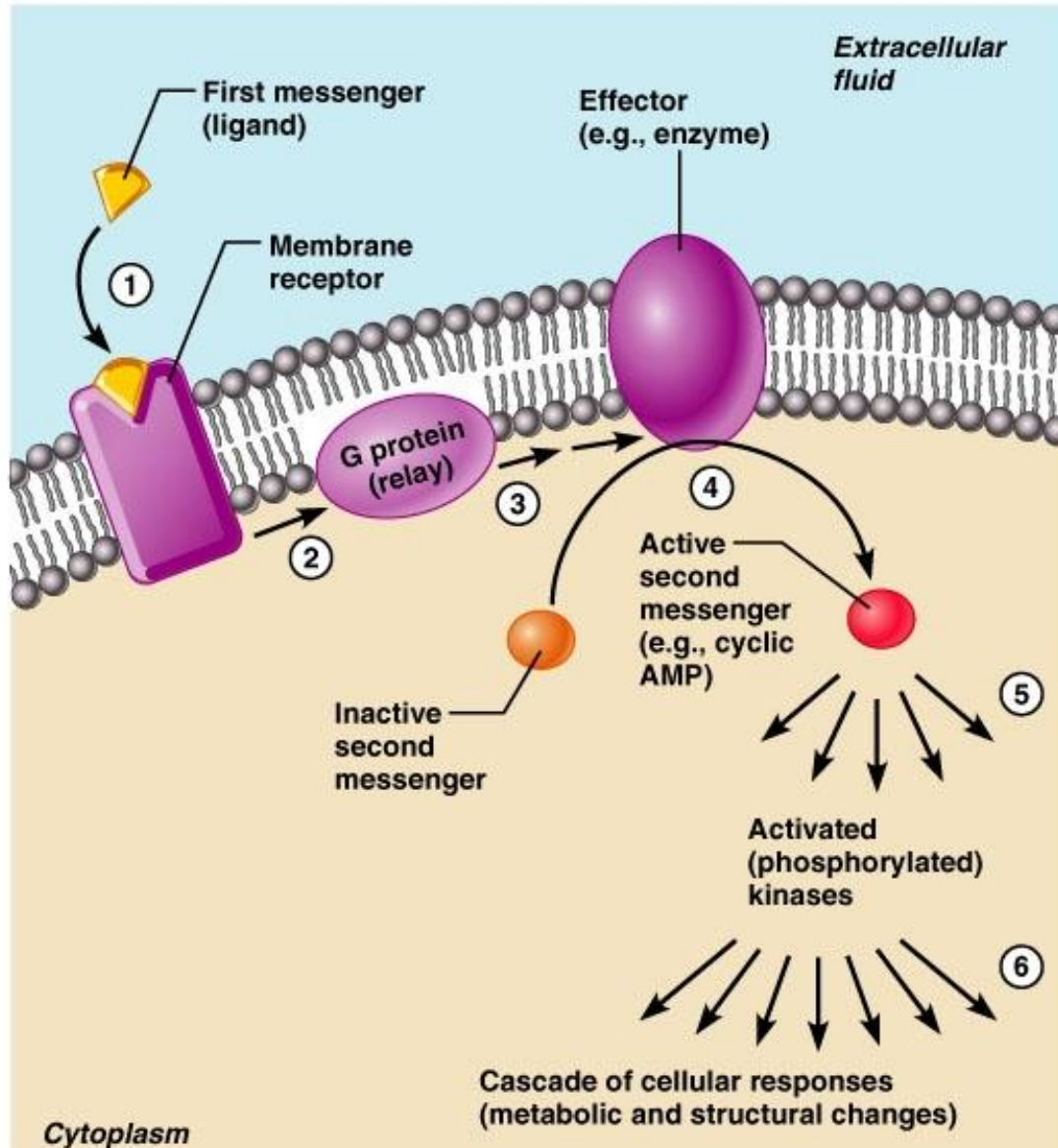
- muscle contraction;
- exocytosis, e.g.,
  - release of neurotransmitters at synapses
- activation of T cells and B cells when they bind antigen with their antigen receptors
- apoptosis
- a variety of biochemical changes mediated by Protein Kinase C (PKC).

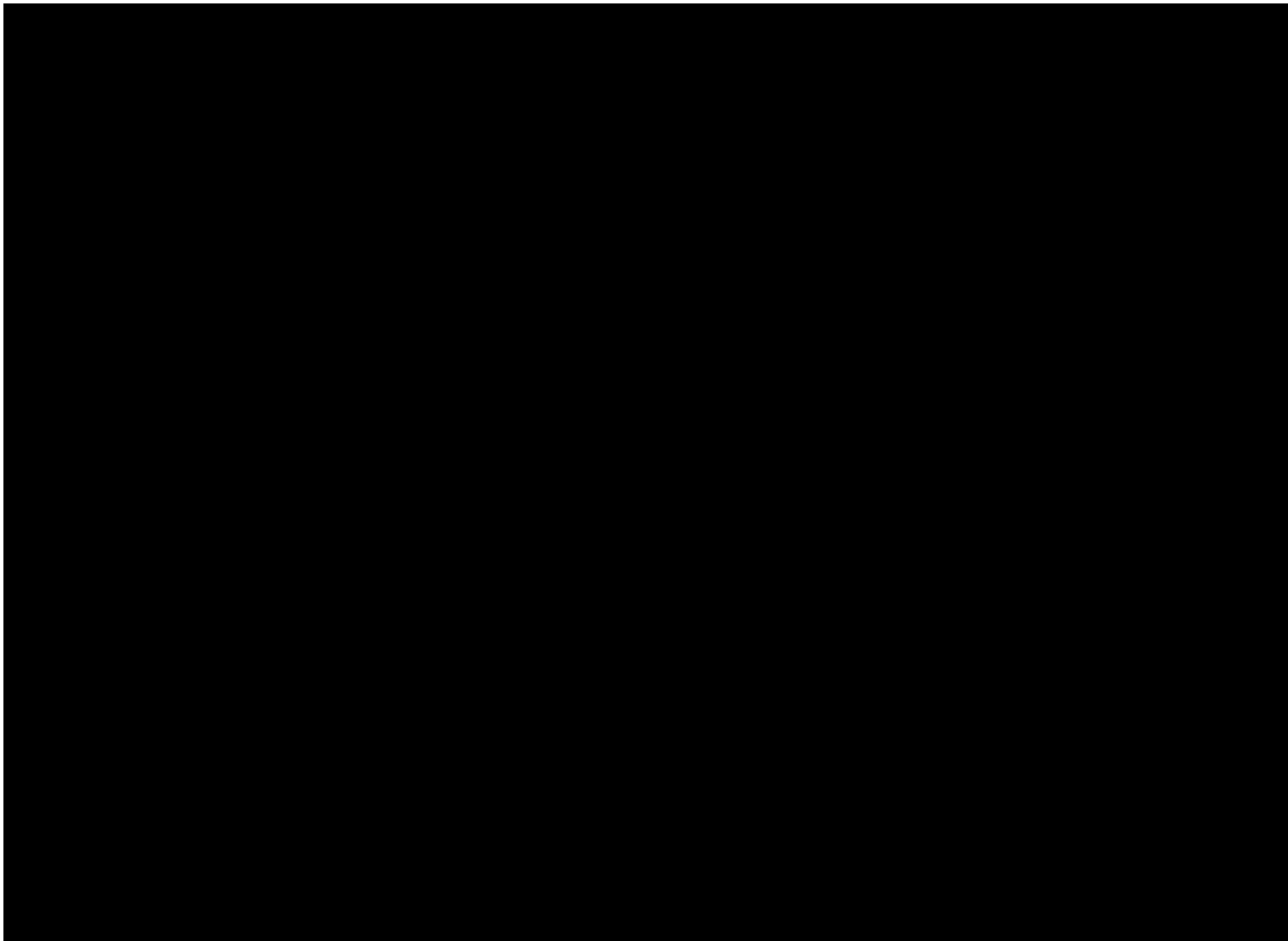
$IP_3$  binds with receptors on smooth endoplasmic reticulum to stimulate the release of calcium ions as a part of the amplification of a hormone's regulation of cellular enzymes.

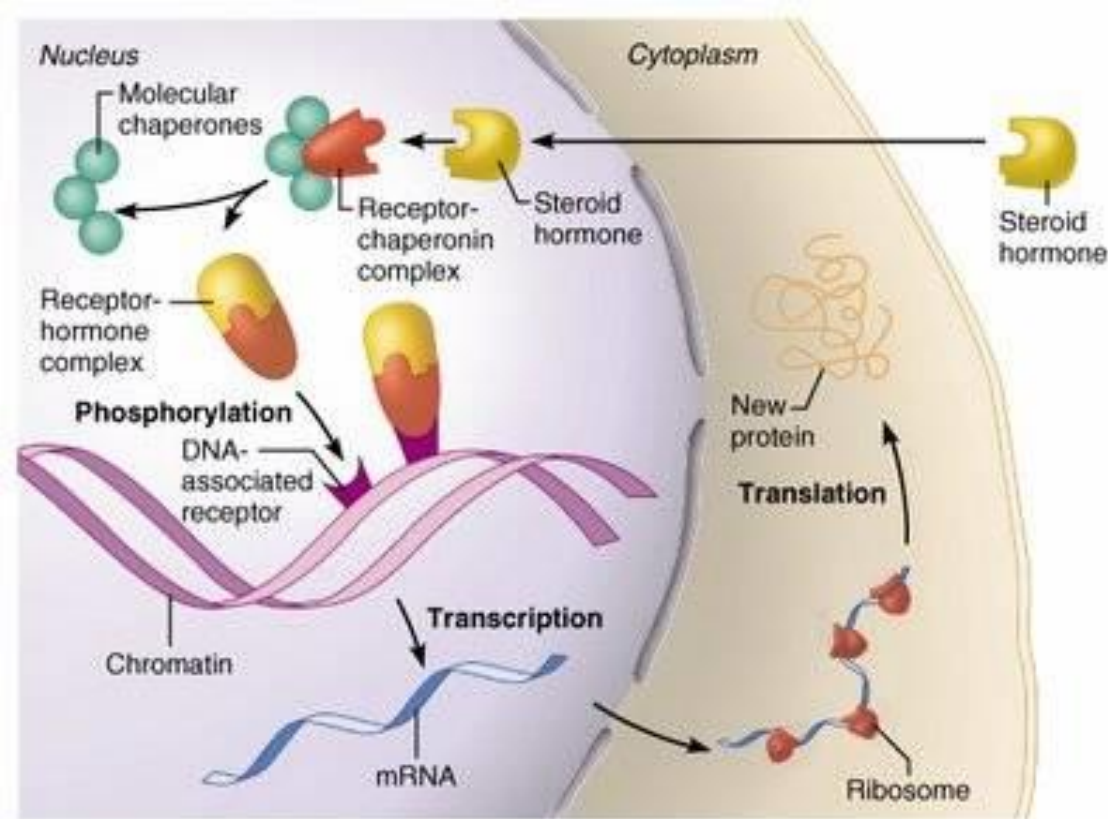




# Hormone stimulated enzyme cascades.







Internal receptors are found in the cytoplasm of the cell and respond to hydrophobic ligand molecules that are able to travel across the plasma membrane. Once inside the cell, many of these molecules bind to proteins that act as regulators of mRNA synthesis to mediate [gene expression](#).

Gene expression is the cellular process of transforming the information in a cell's [DNA](#) into a sequence of [amino acids](#) that ultimately forms a protein. When the ligand binds to the internal receptor, a conformational change exposes a DNA-binding site on the protein. The ligand-receptor [complex](#) moves into the nucleus, binds to specific regulatory regions of the chromosomal DNA, and promotes the initiation of [transcription](#). Internal receptors can directly influence gene expression without having to pass the signal on to other receptors or messengers.



# References

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- Becker, Kleinsmith, Hardin, Bertoni (2009). The World of the Cell. Seventh Edition. Pearson Education
- Youtube videos