



Centrum pro výzkum  
toxických látek  
v prostředí

# BIOMARKERS AND TOXICITY MECHANISMS

## 08 – Mechanisms

### Signalling and regulation

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[www.recetox.cz](http://www.recetox.cz)

Tento projekt je spolufinancován Evropským sociálním fondem a státním rozpočtem České republiky.



INVESTICE DO ROZVOJE VZDĚLÁVÁNÍ

# Cell communication & regulation: a target for toxicants

... especially sensitively regulated processes are highly susceptible to toxicants

→ toxicity to REGULATIONS & SIGNALLING

## Hierarchy in signalling

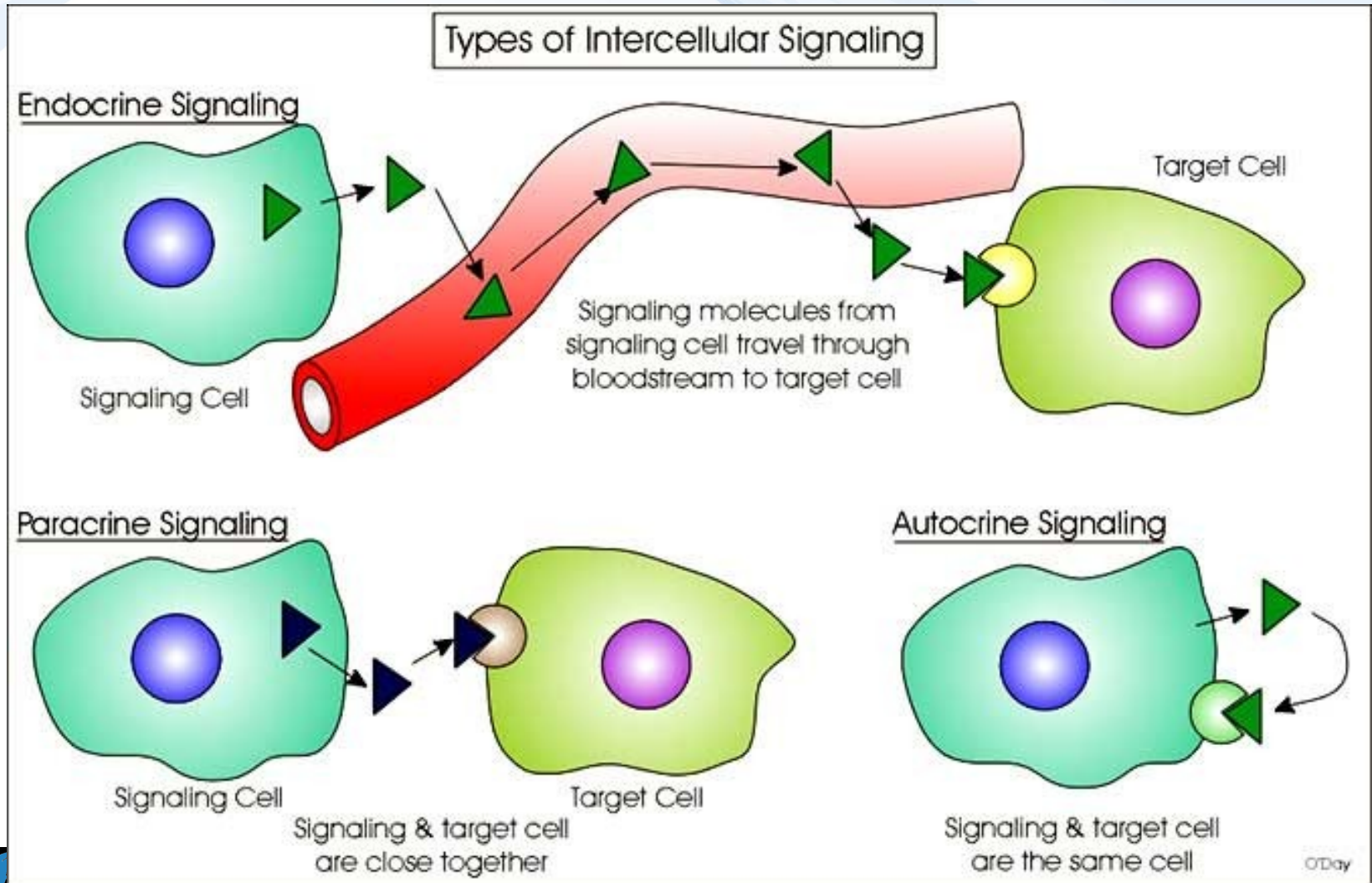
- **systems**: neuronal  $\leftrightarrow$  endocrine
- **cell-to-cell**
  - hormonal & neuronal signal transmission
  - contact channels
- **intracellular** signal transduction

# INTER-cellular signals

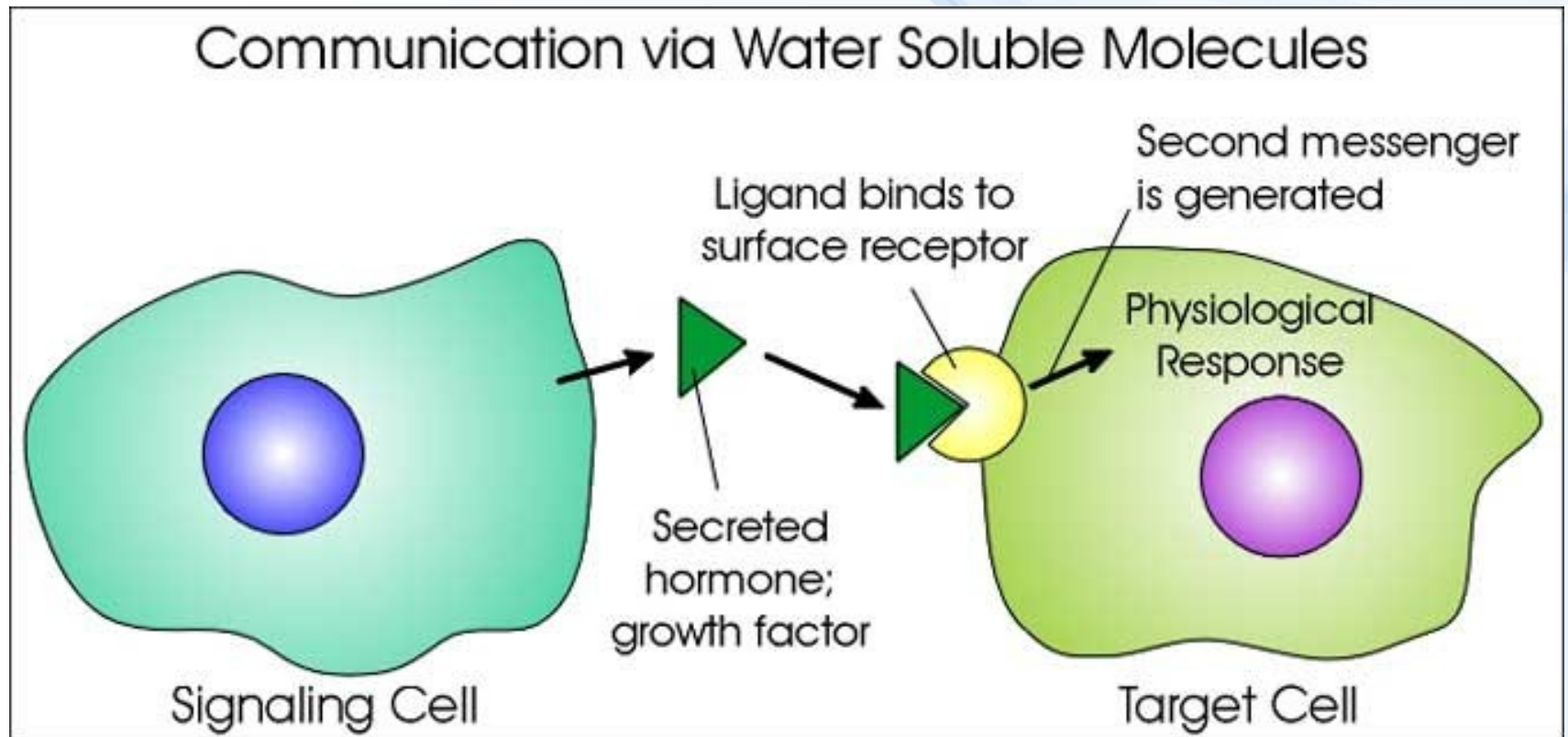
## Overview



# Cell to cell communication & regulation: a target for toxicants

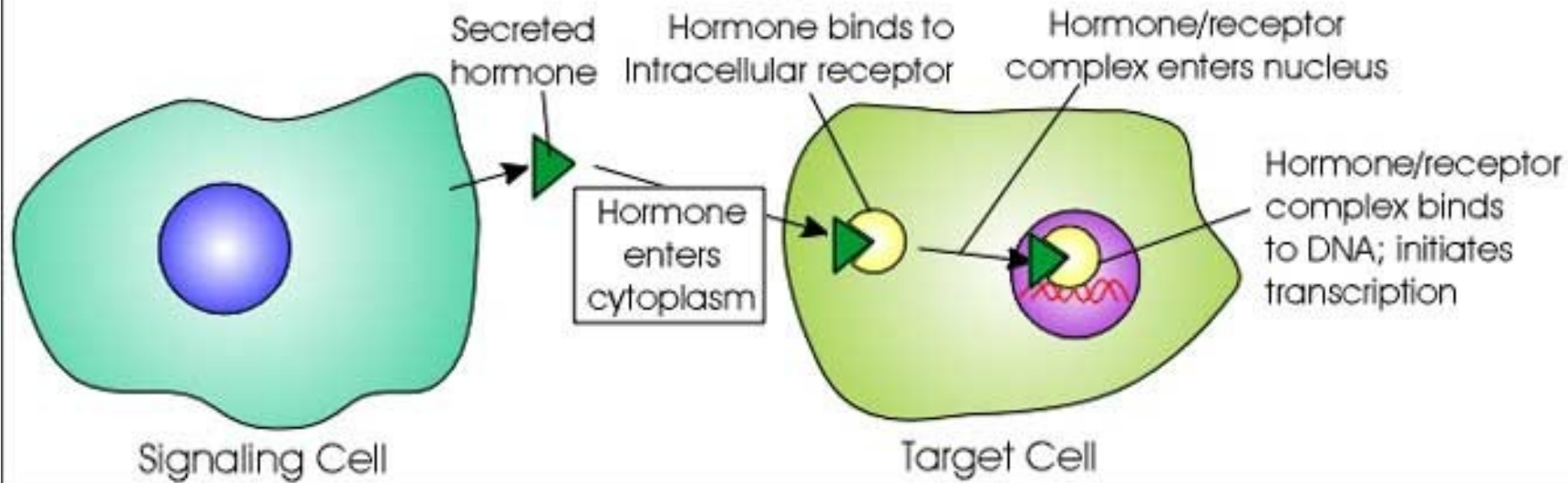


# Cell to cell communication (1)



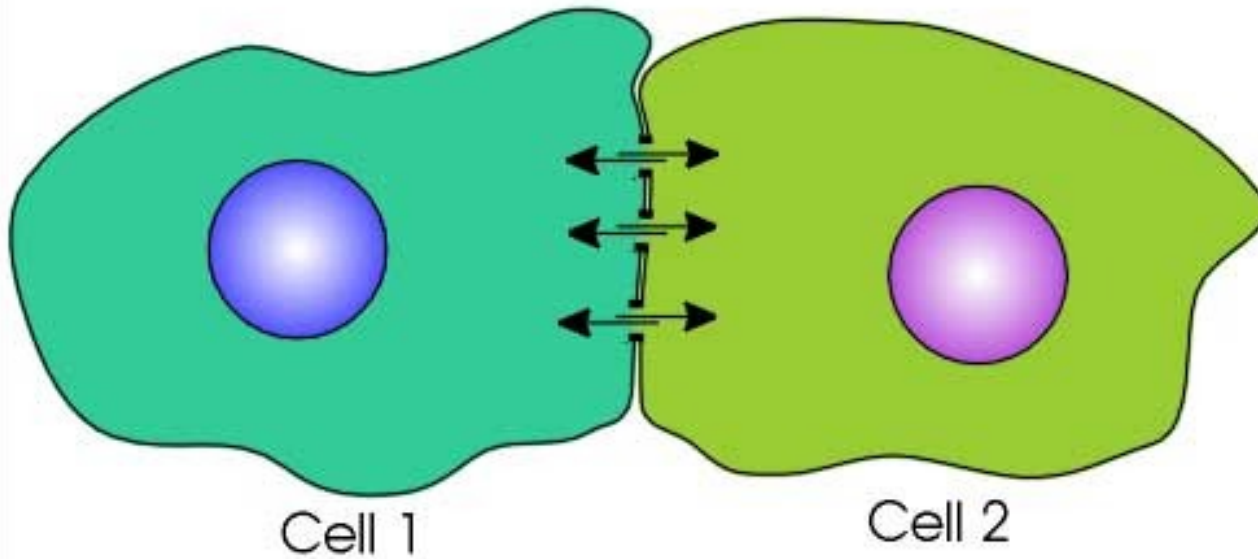
# Cell to cell communication (2)

## Communication via Lipid Soluble Molecules

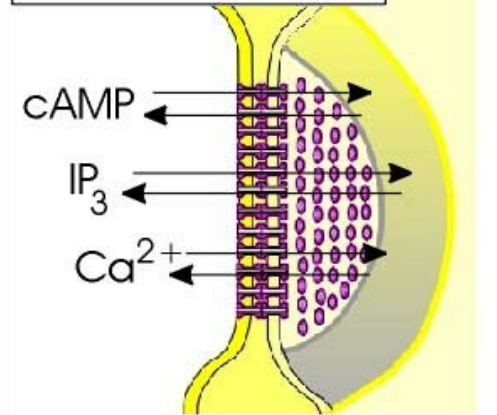


# Cell to cell communication (3)

## Communication via Cellular Continuities



## Gap Junction

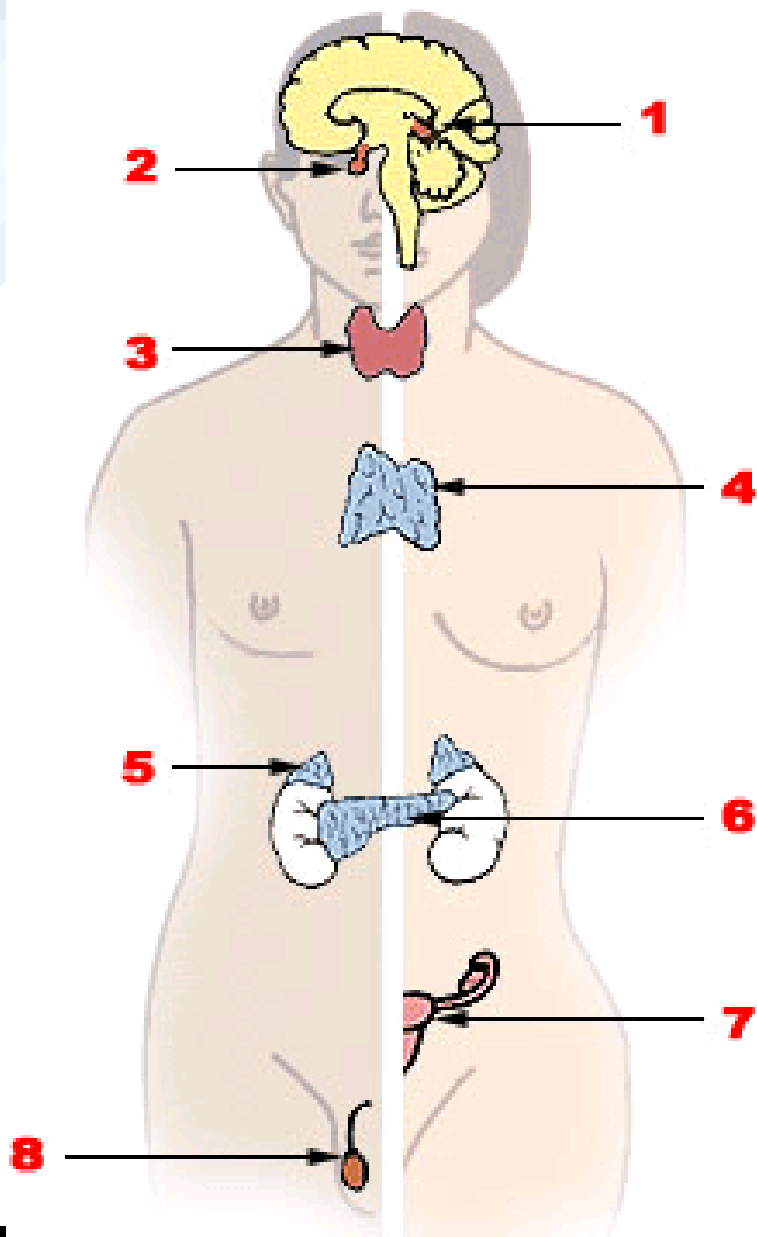


# INTER-cellular signals

Hormones

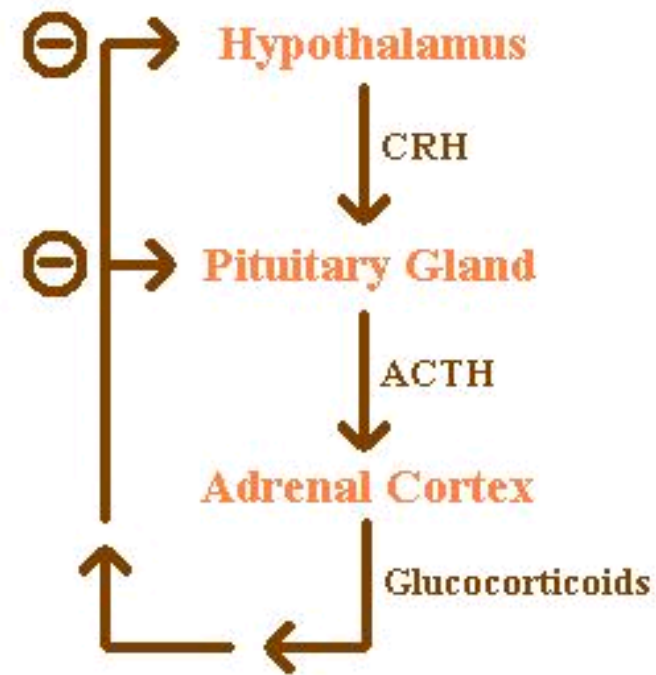






## Endocrine system:

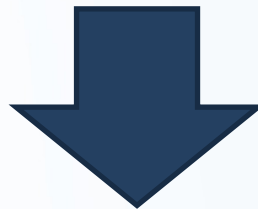
1. Pineal gland, 2. Pituitary gland, 3. Thyroid gland, 4. Thymus, 5. Adrenal gland, 6. Pancreas, 7. Ovary, 8. Testis



Example: feedback loop

## FUNCTIONS OF HORMONES

- \* stimulation or inhibition of growth
- \* mood swings
- \* induction or suppression of apoptosis  
(programmed cell death)
- \* activation or inhibition of the immune system
- \* regulation of metabolism
- \* preparation for fighting, fleeing, mating ...
- \* preparation for a new phase of life  
(puberty, caring for offspring, and menopause)
- \* control of the reproductive cycle  
.... etc.



Chemicals interfering with  
various hormonal functions  
→ **diverse impacts (effects)**

## FATE OF HORMONES: target for toxicants

Toxic compounds can affect “hormone signalling” at various levels (highlighted):

1. Biosynthesis of a particular hormone in a particular tissue
2. Storage and secretion of the hormone
3. Transport of the hormone to the target cell(s)
4. Recognition of the hormone by an associated cell membrane or intracellular receptor protein.
5. Relay and amplification of the received hormonal signal via a signal transduction process -> cellular response.
6. The reaction of the target cells is recognized by the original hormone-producing cells (negative feedback loop)
7. Degradation and metabolism of the hormone

More details will be discussed  
in the lectures dedicated to  
nuclear receptors

# Toxicity to hormone regulation = ENDOCRINE DISRUPTION

## ED & EDCs (endocrine disrupting compounds)

= major problem in environmental toxicology

Effects at **all levels of hormonal action** have been demonstrated

→ *synthesis, transport, site of action ....*

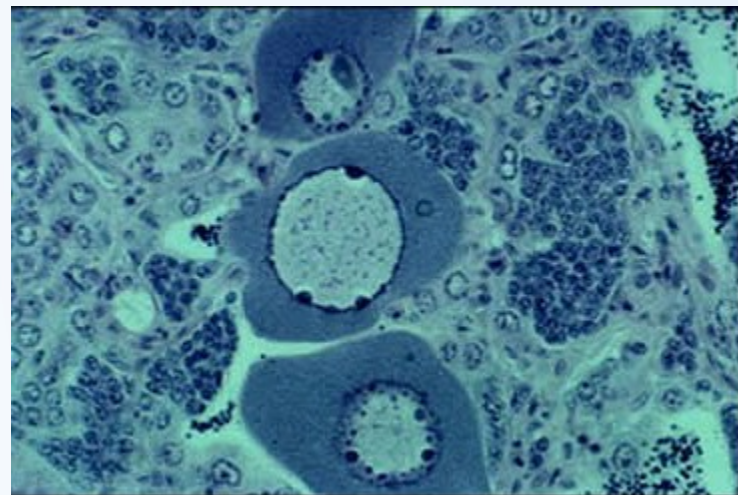
- **Multiple effects** due to ED (! Not only „xenoestrogenicity“ & feminization)

→ *immunotoxicity, developmental toxicity*

**(ED - WILL ALSO BE DISCUSSED FURTHER)**

## Example of ED - Intersex roach testis

containing both oocytes and spermatozoa,  
caused by exposure to environmental oestrogens



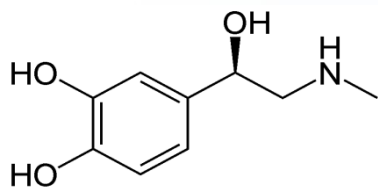
# Types of hormones in vertebrates

## Amine-derived hormones

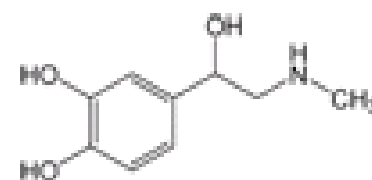
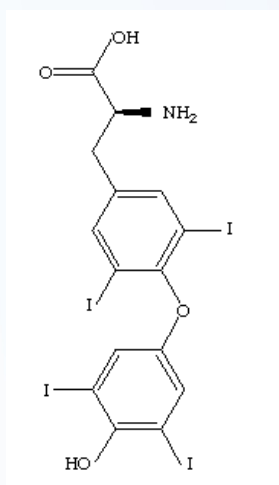
structure: derivatives of the amino acids tyrosine and tryptophan.  
Examples - catecholamines and thyroxine.

(small molecules - **similar to organic toxicants** → **TOXIC EFFECTS**)

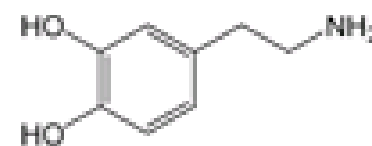
Adrenalin



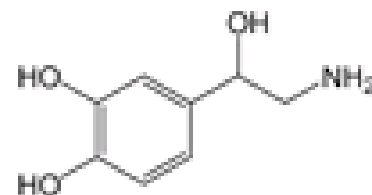
Thyroxin



Epinephrine



Dopamine



Norepinephrine



# Types of hormones in vertebrates

## Peptide hormones

structure: chains of amino acids.

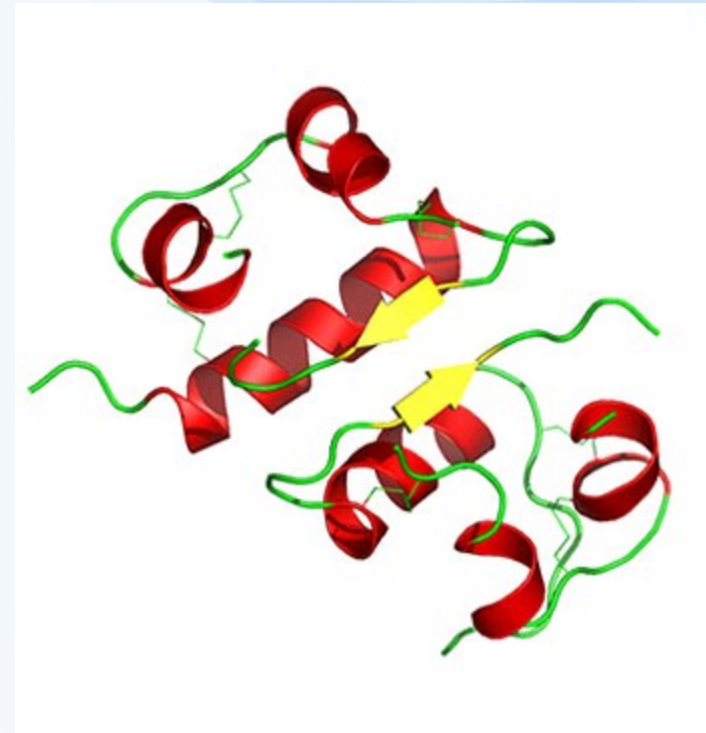
- small peptides: TRH and vasopressin;
- large proteins: insulin, growth hormone, luteinizing hormone, follicle-stimulating hormone and thyroid-stimulating hormone etc.

*Large molecules;*

*receptors on surfaces of the cells*

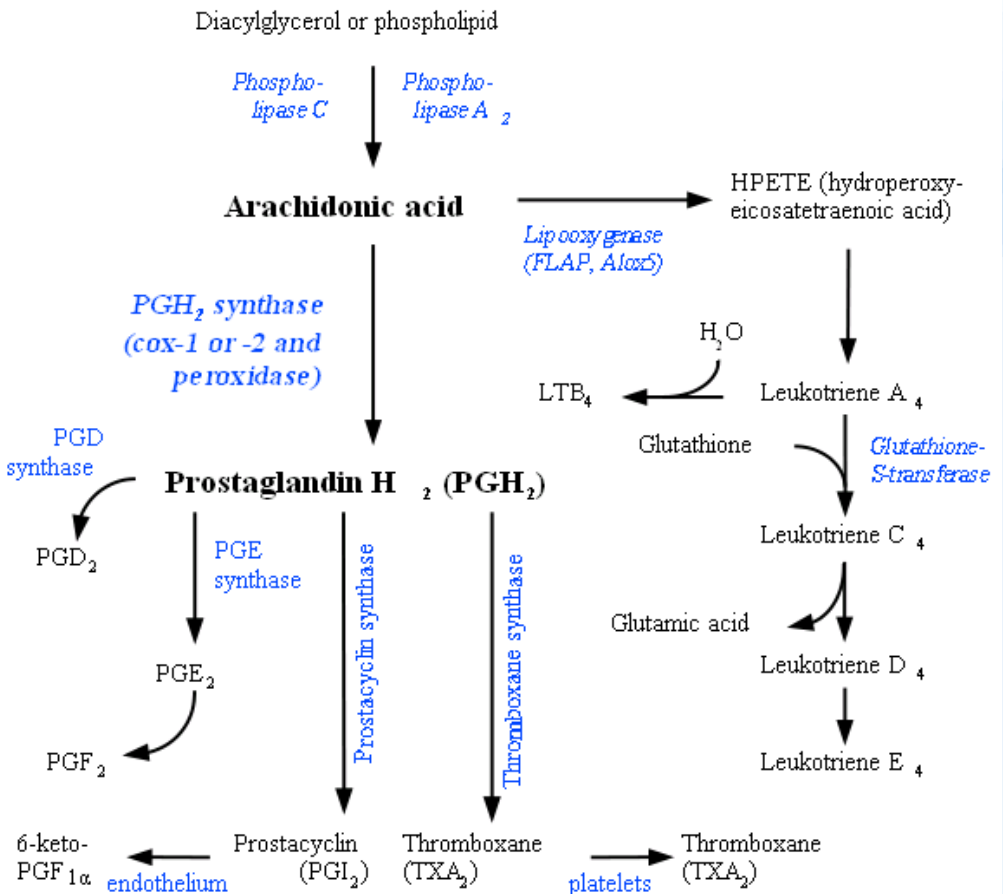
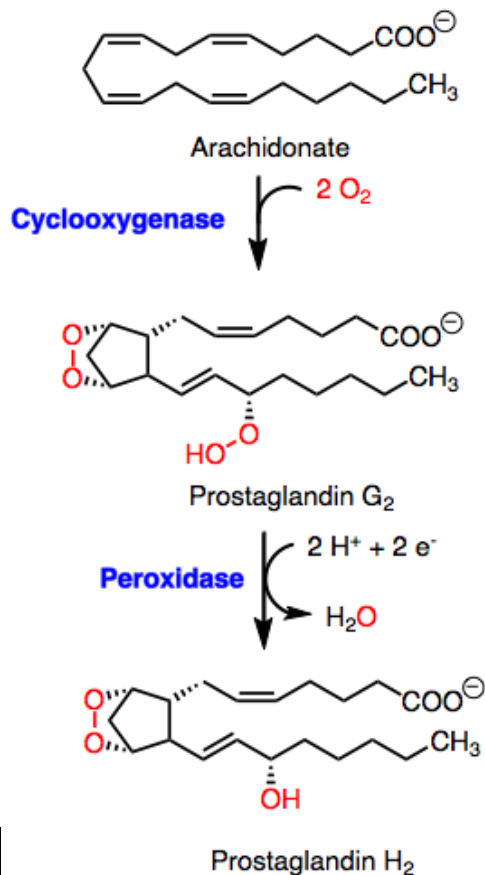
*(Interactions with toxic chemicals **less likely**)*

Example - insulin



# Types of hormones (signal molecules) in vertebrates

## Lipid derived “hormones” (1) - from linoleic acid, arachidonic acid - prostaglandins



# Types of hormones in vertebrates

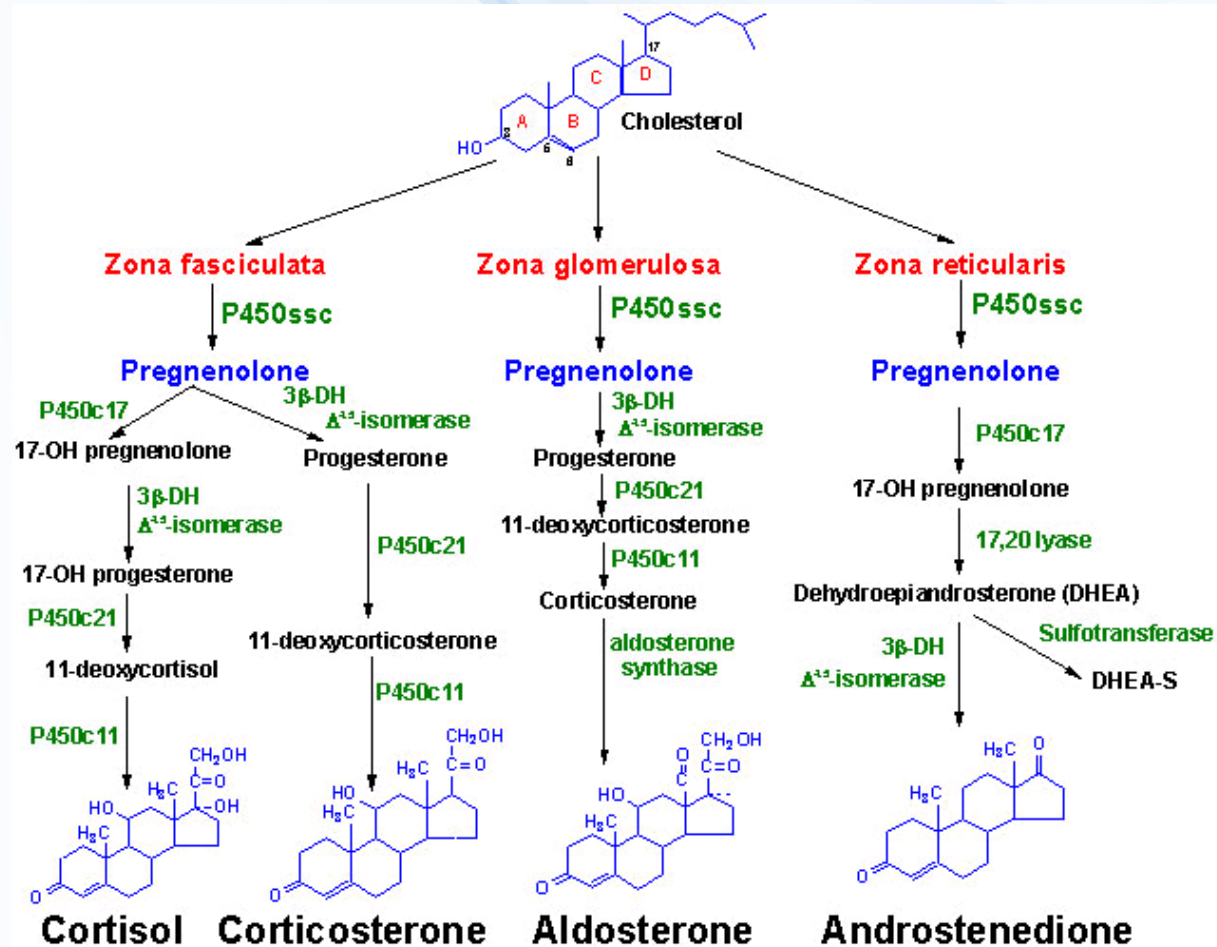
## Lipid derived hormones 2 - steroid hormones

\* Small molecules - similar to organic toxicants:

→ several compounds **interfere with steroid hormones** → **toxicity !!!**

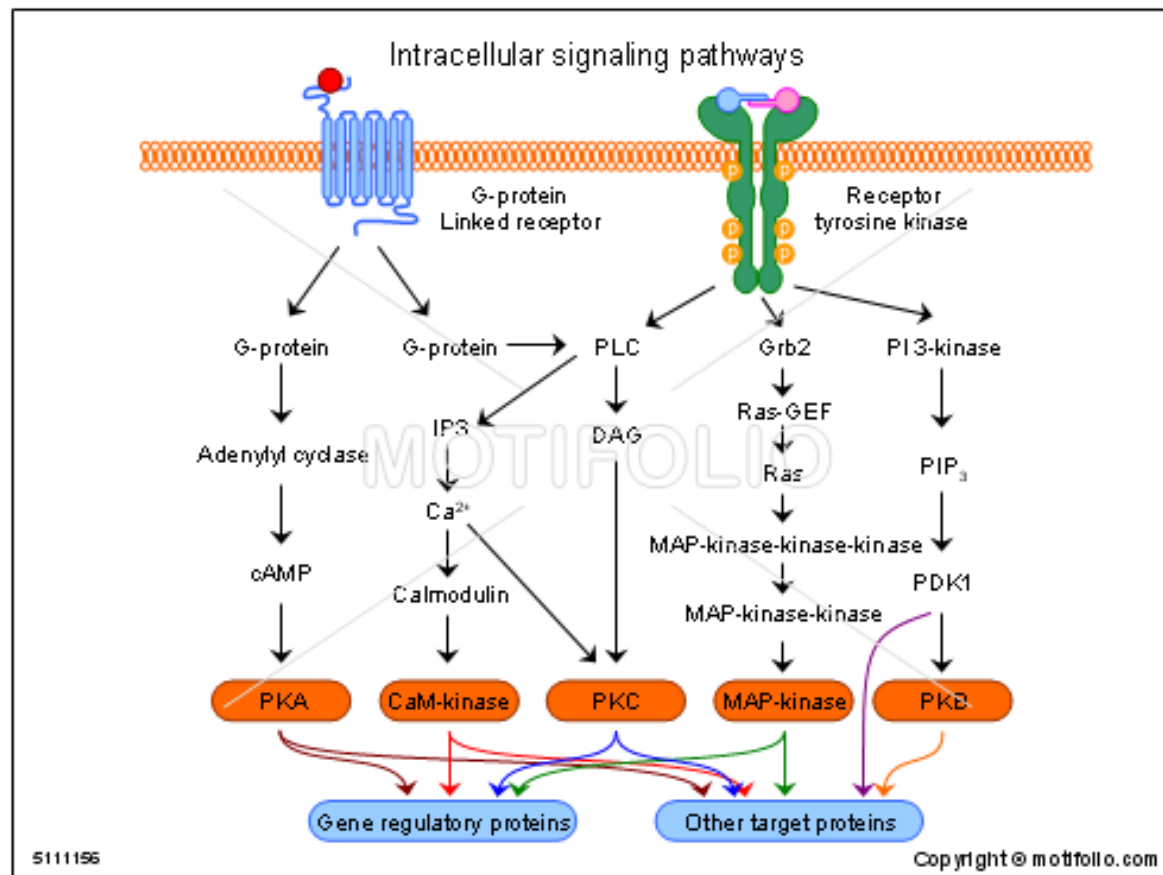
Derived  
from cholesterol

Examples:  
testosterone,  
cortisol,  
estradiol ...





# INTRACELLULAR signals



# Intracellular signal transduction: target of toxicants

- **Regulation of cell life = control of major cell functions**
  - metabolism
  - proliferation
  - differentiation
  - death (apoptosis)
  
- **Regulation controlled by complex signalling**
  - "network" of general pathways
  - similar in all cells / different cell-specific effects



# Intracellular signal transduction: target of toxicants

## - Consequences of signalling disruption

- unwanted changes in „homeostatic“ rates among proliferation / differentiation / apoptosis

- cell transformation (carcinogenicity)

- embryotoxicity

- immunotoxicity

- reproduction toxicity

*.... and other chronic types of toxicity*

# Signal transduction - principles

## Two major signalling processes

- **protein-(de)phosphorylation**

ProteinKinases - PKs, ProteinPhosphatases - PPases

- **secondary messengers**

cAMP / IP3, PIP2, DAG, Ca<sup>2+</sup>, AA

## Three major types of signalling

1: Membrane receptors (G-protein, kinases)

→ activation of protein kinase A (PKA):

major messenger: cAMP, MAPKs

2: Membrane receptors

→ activation of membrane lipases → and later proteinkinase C

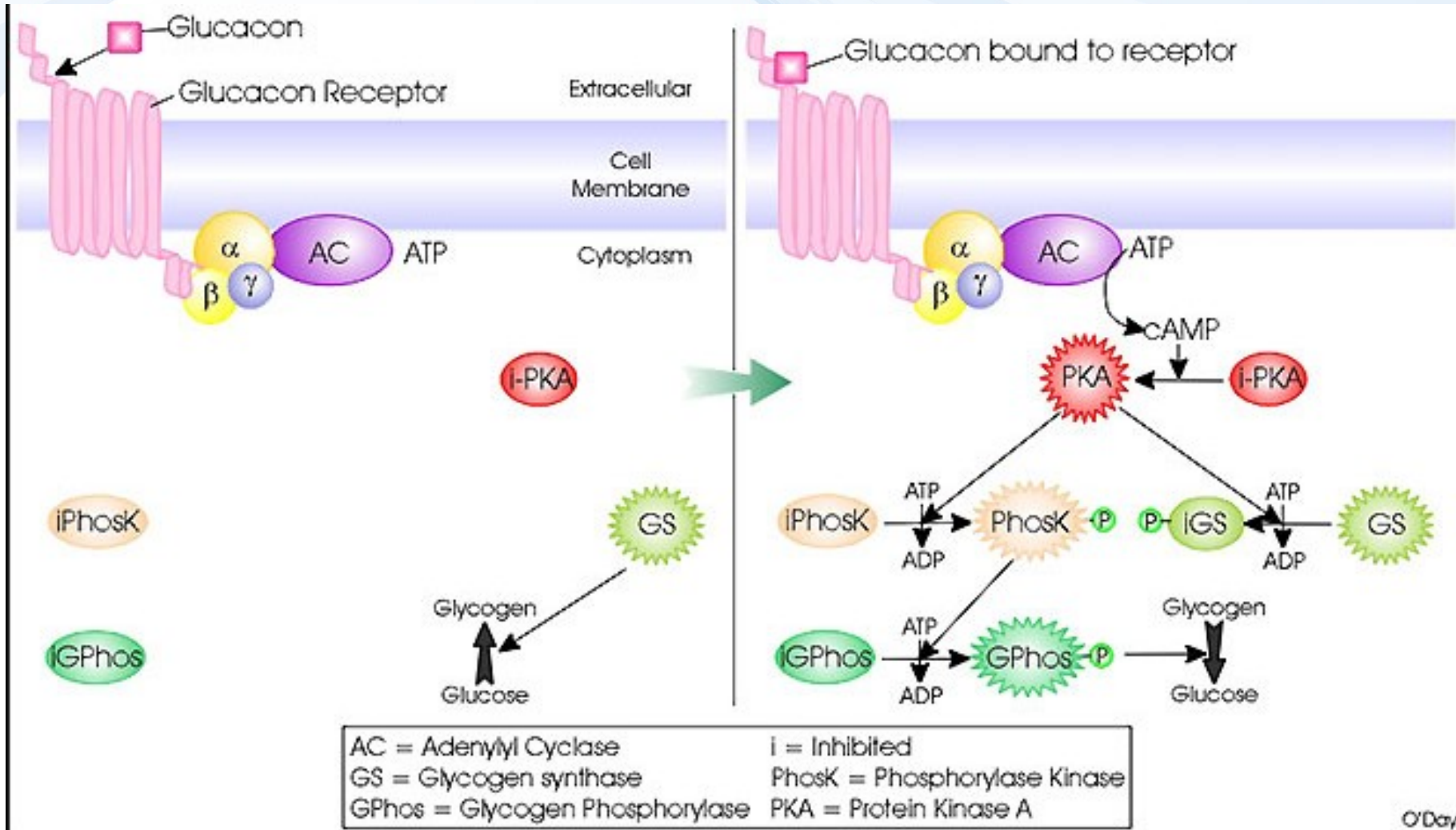
IP3, PIP2, DAG, Ca<sup>2+</sup>, AA

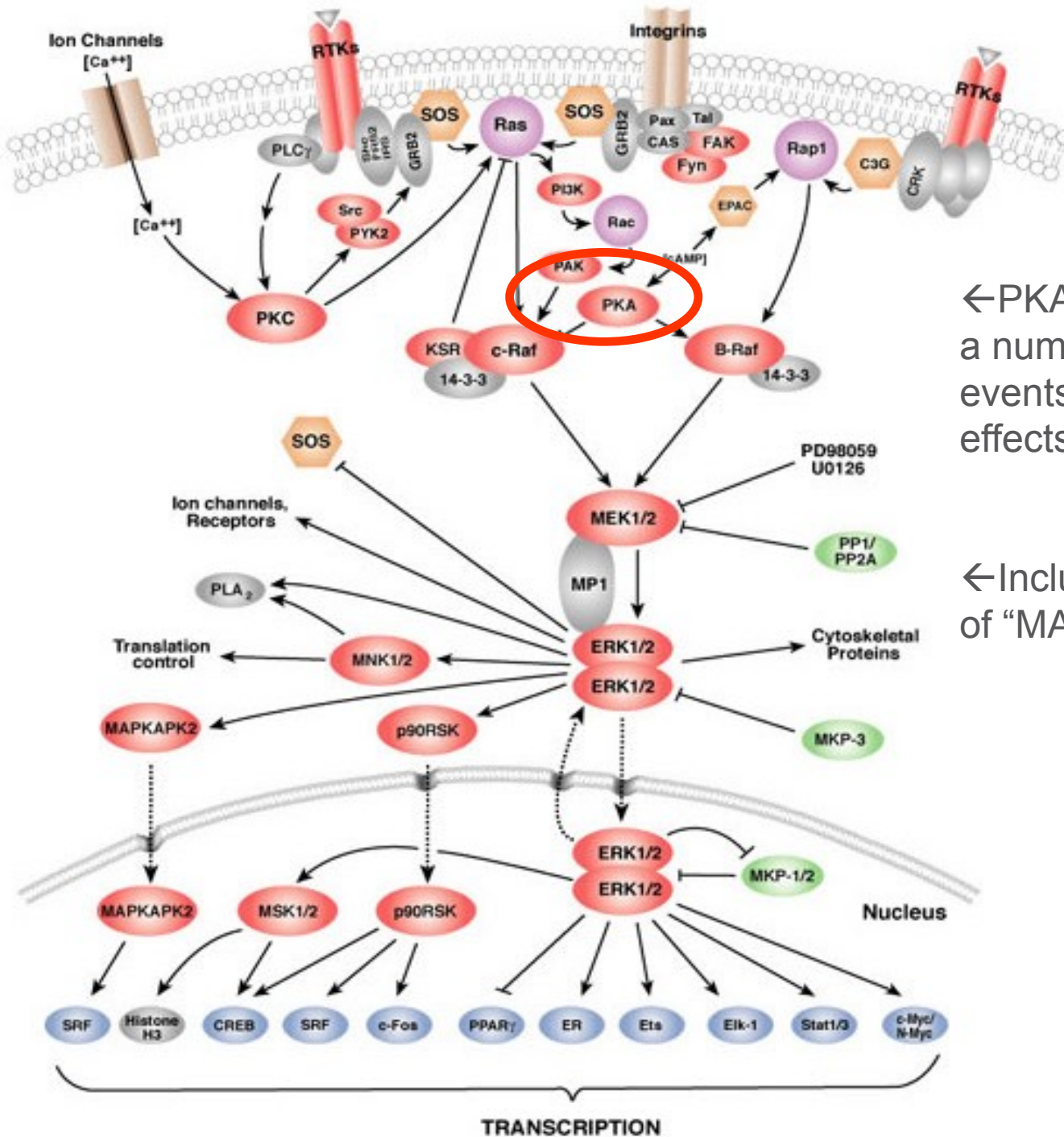
3: **Cytoplasmic (nuclear) receptors** (discussed in detail in other sections)



# Signalling mechanism 1

→ Activation of adenylate cyclase → cAMP → PKA





← PKA is central to a number of signalling events and following effects

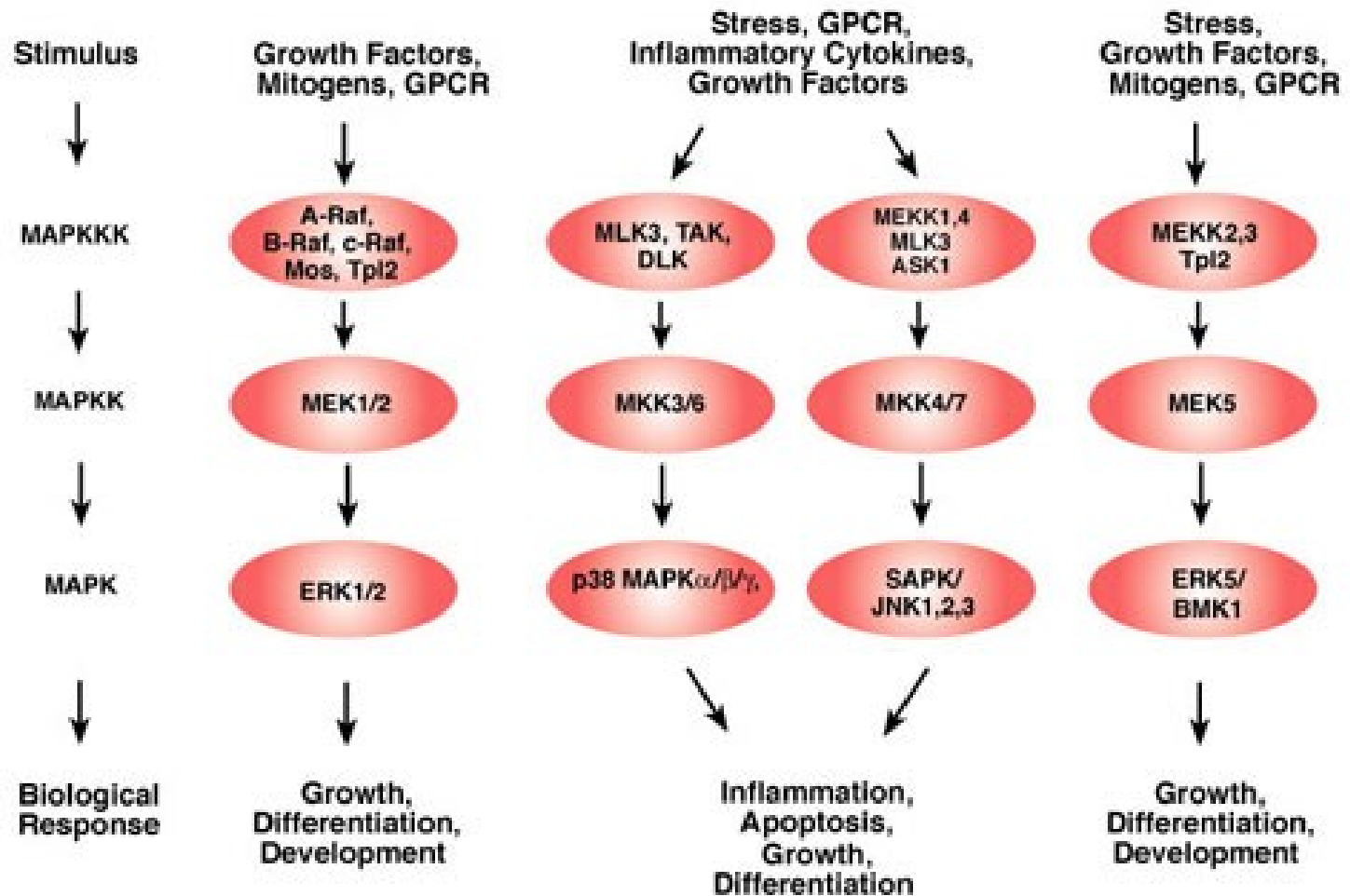
← Including modulation of “MAPKs”

TRANSCRIPTION



# Mitogen Activated Protein Kinases (MAPKs) & dependent effects

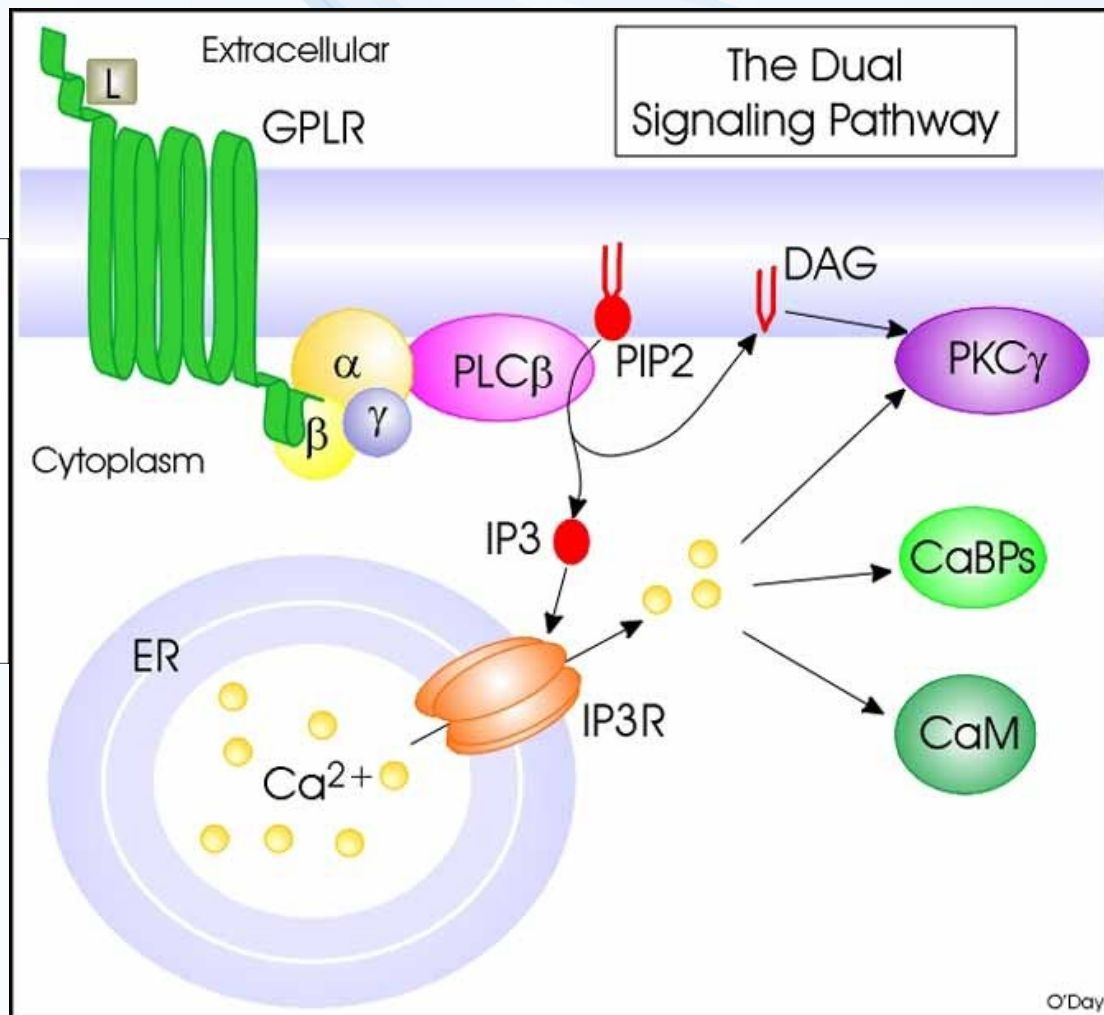
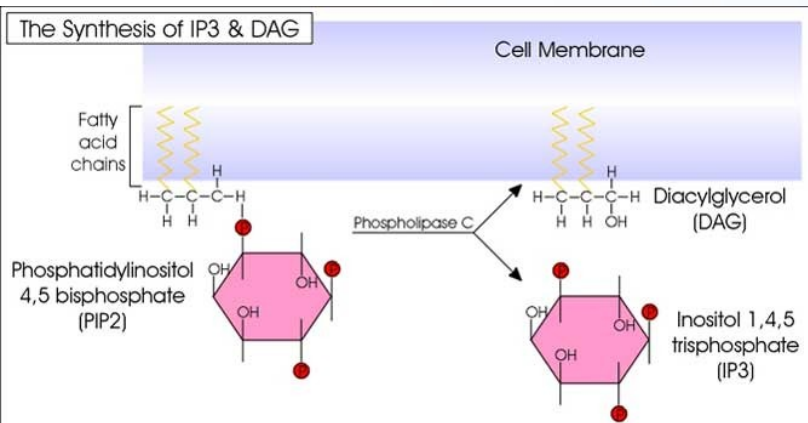
## MAPK signaling cascades



# Signalling mechanism 2

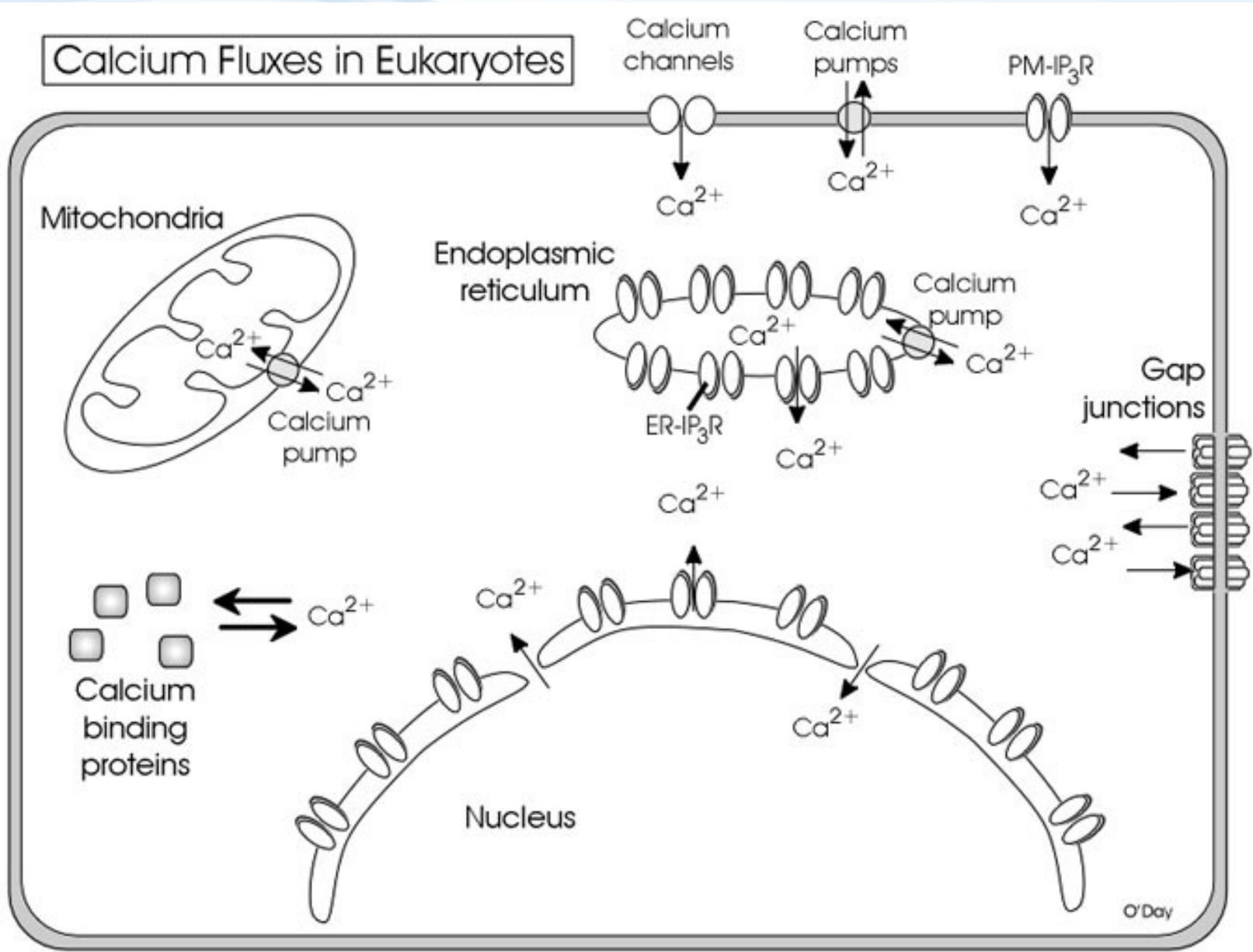
## Activation of Phospholipase C

→ release of PIPs → DAG → PKC / arachidonic acid  
+ IP3 → activation of Ca<sup>2+</sup> signalling





# Calcium Fluxes in Eukaryotes

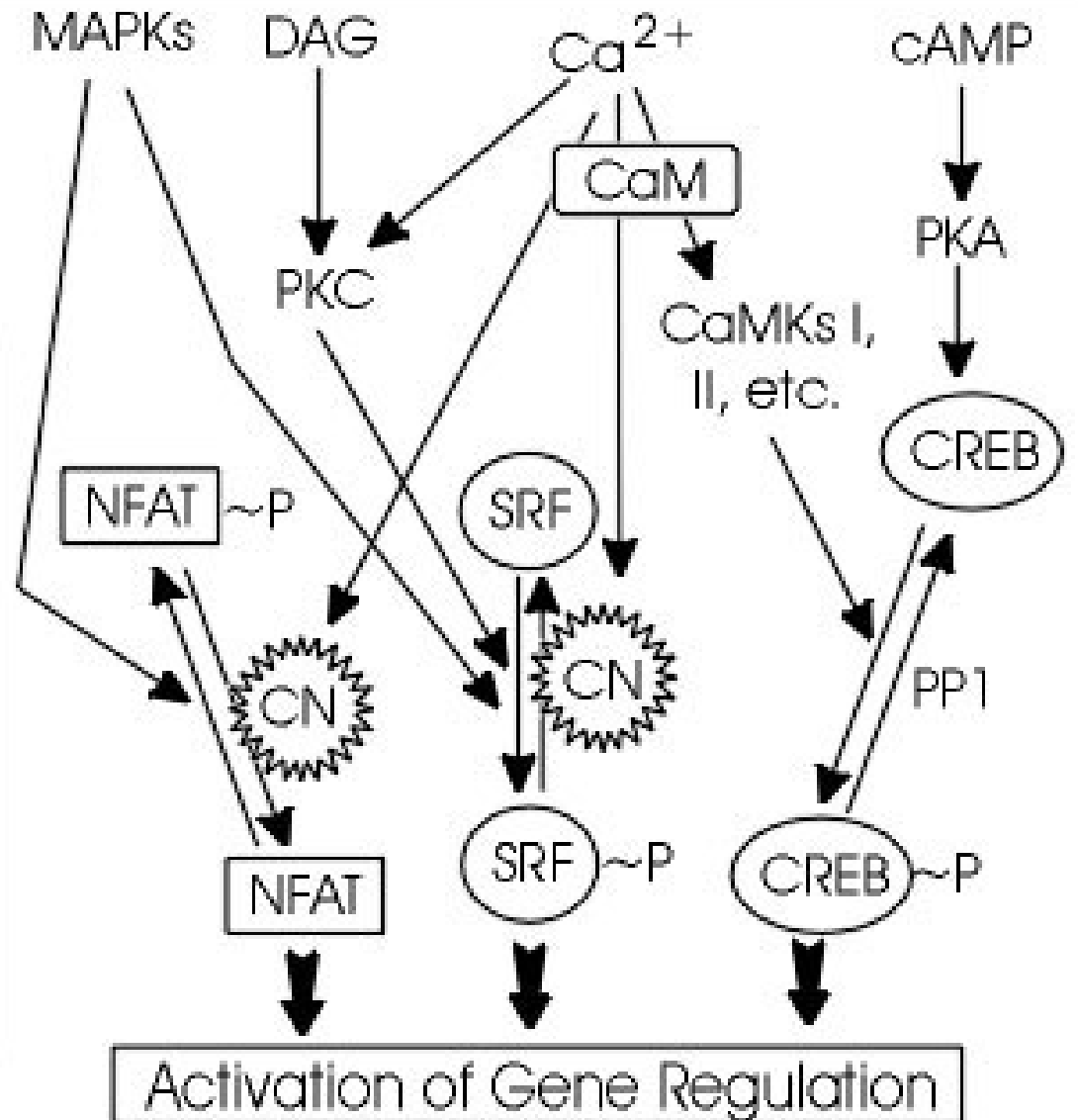


# Different "types" of signalling crosstalk → networks

## Some Signaling Pathways Leading to Gene Regulation

### Transcription Factors

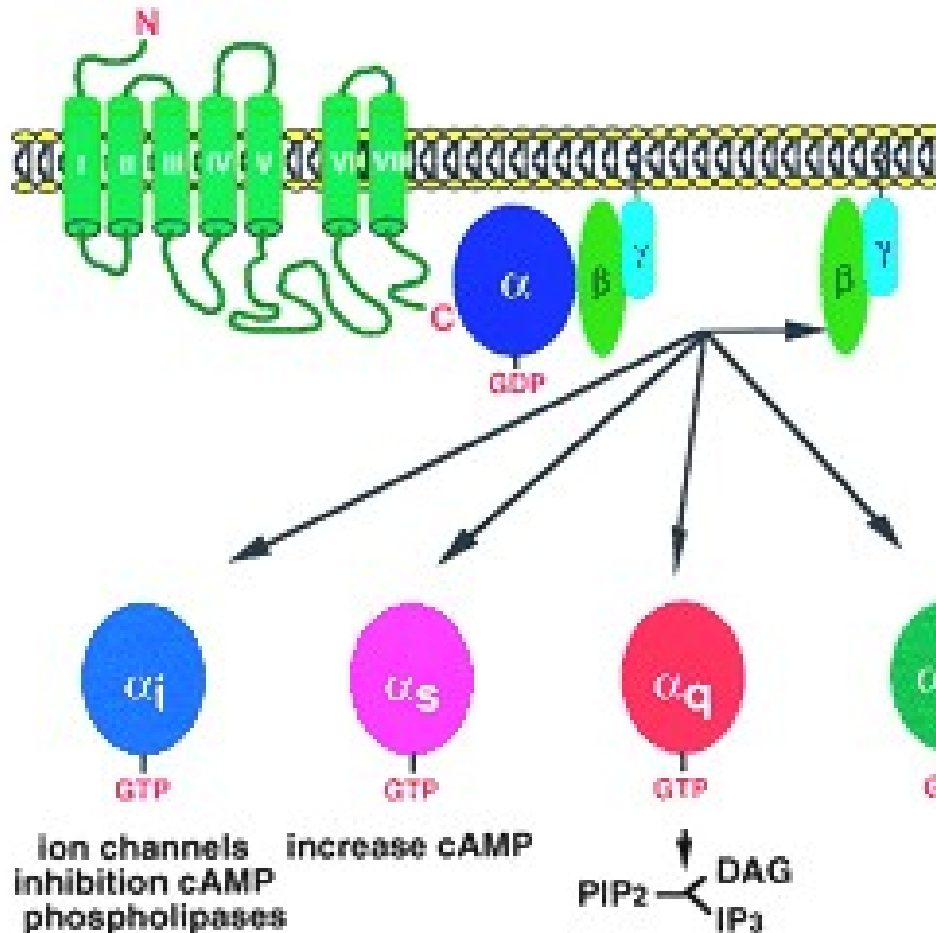
- NFAT** = Nuclear Factor of Activated T-cells
- SRF** = Serum Response Factor
- CREB** = cAMP Response Element Binding protein



# G-proteins & G-protein coupled receptors – GPCRs

Involved in many functions → triggering different downstream events

## G PROTEIN- COUPLED RECEPTORS



## Biological functions

smell and taste  
(~1000 types of receptors)  
perception of light  
neurotransmission  
function of endocrine  
and exocrine glands  
chemotaxis  
exocytosis  
control of blood pressure  
embryogenesis  
development  
cell growth and differentiation  
HIV infection  
oncogenesis

# Disruption of intracellular signaling - EXAMPLES

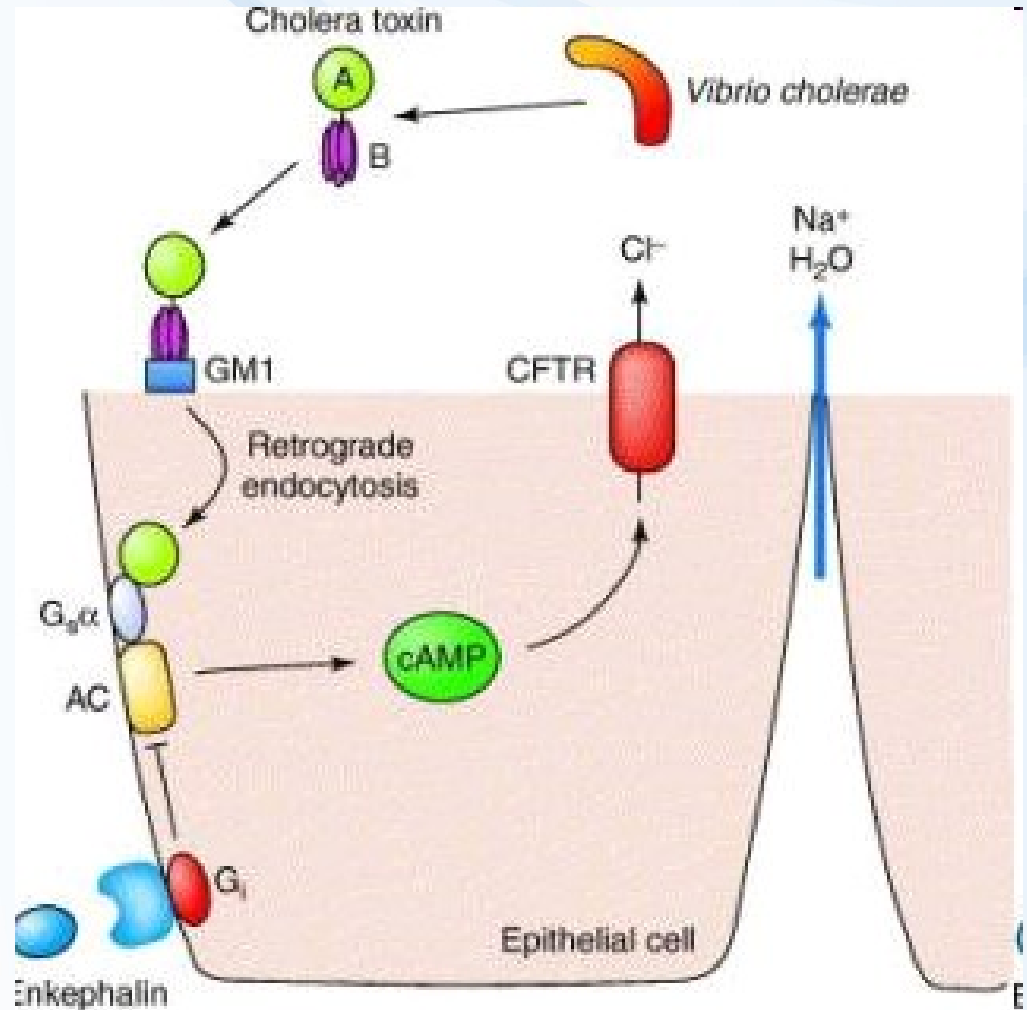
## Cholera toxin

*Vibrio cholerae*

CT acts as **adenylate cyclase** enzyme

→ increasing cAMP levels

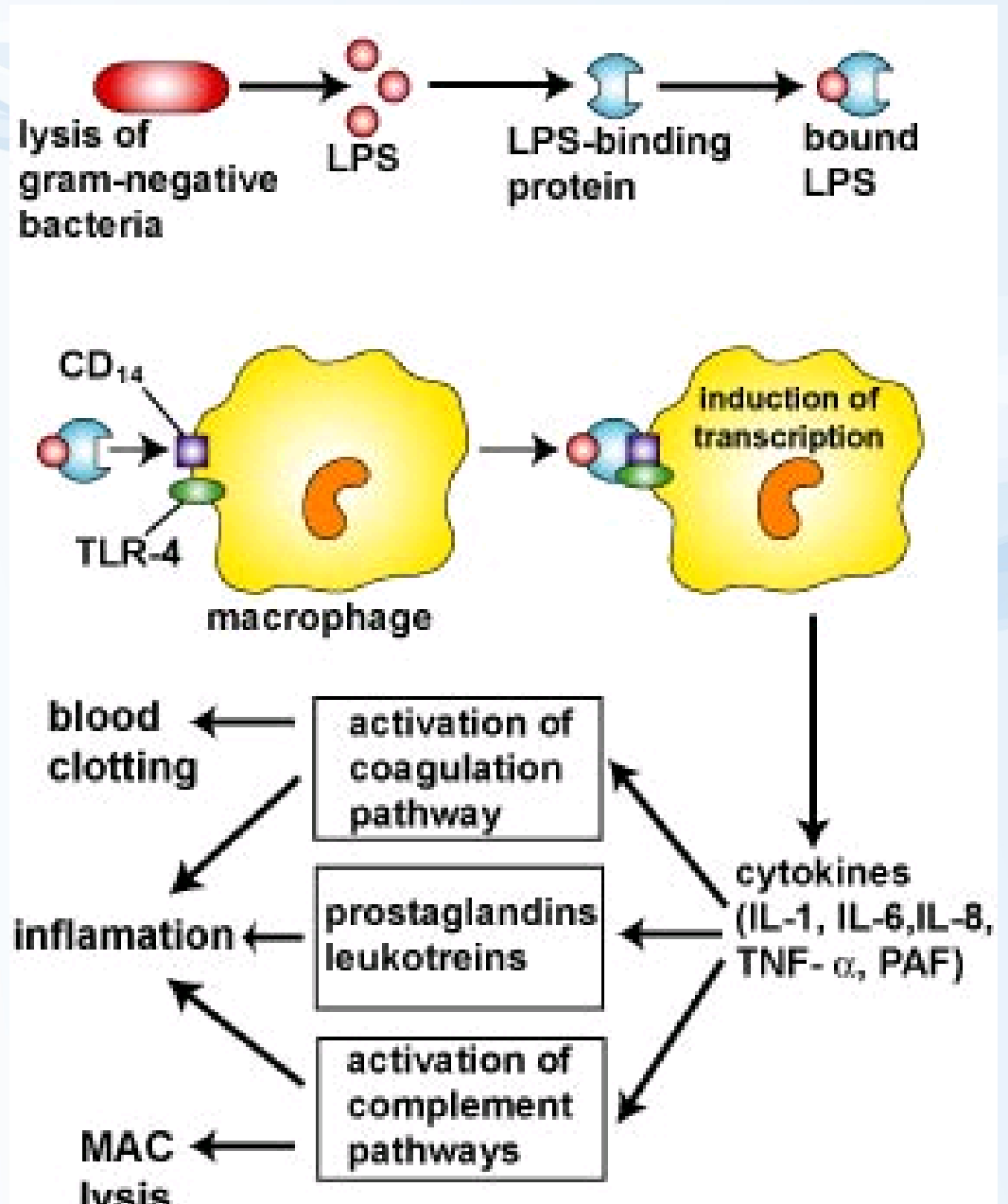
→ TOXICITY



## Example:

Lipopolysaccharides (LPS) from cell walls

→ hyperactivation of intracellular signals → immunotoxicity



## Immunomodulatory Potency of Microcystin, an Important Water-Polluting Cyanobacterial Toxin

Ondrej Adamovsky,<sup>\*,†</sup> Zdena Moosova,<sup>†</sup> Michaela Pekarova,<sup>‡</sup> Amrita Basu,<sup>†</sup> Pavel Babica,<sup>†</sup> Lenka Svihalkova Sindlerova,<sup>‡</sup> Lukas Kubala,<sup>‡</sup> and Ludek Blaha<sup>†</sup>

