# Lecture 10 Cell communication Part I

### Outline

- I. Overview of cell signaling
- II. Intracellular signaling
- III. General principles of cell surface signaling
- IV. Several methods to study cell signaling
- V. Positive and negative feedback in signaling and signaling kinetics

### What is cell communication

Human:





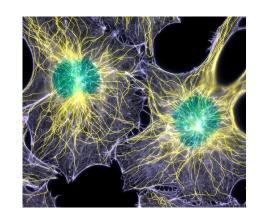
Language Body language

Insect:



Pheromones Touch noise

Cell:



Physical: light, mechanical force, heat
Chemical: proteins, peptides, amino acid
derivatives, nucleotides,
steroids, retinoids,
fatty acid derivatives,
Gases( NO, CO), etc.

### The seadevil- anglerfish

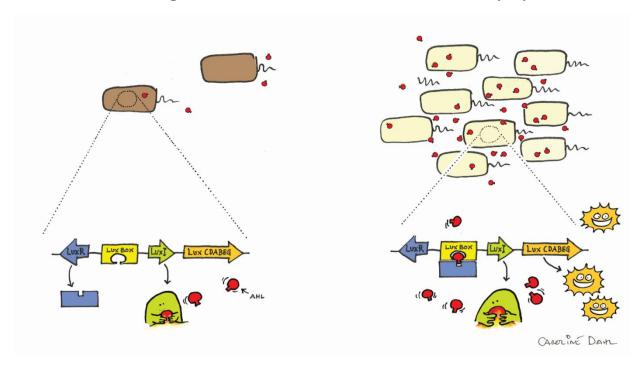




https://video.nationalgeographic.com/video/weirdest-angler-fish

### Quarum sensing in bacteria

#### Chemical signals secretion correlation to cell population

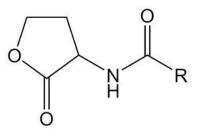


Low cell density

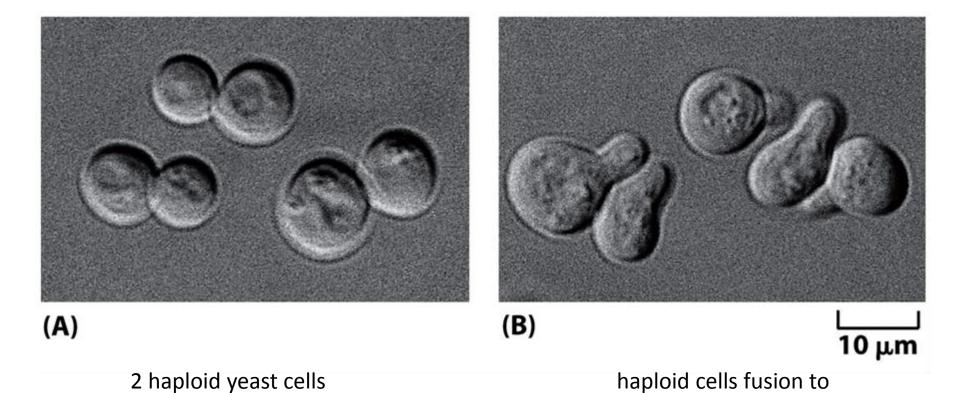
high cell density

The signaling molecule --- AHL: acyl homoserine lactone

酰基高丝氨酸内酯



### Budding yeast mating corresponds to mating factor (a peptide)



become diploid cells

#### I. Overview of cell communication

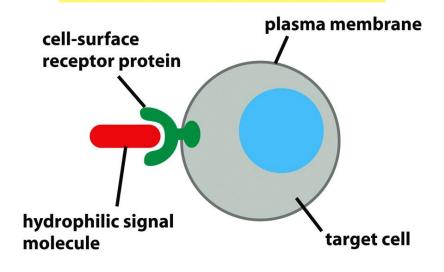
Chemical signaling involves ligands and receptors

Two different types of receptors:

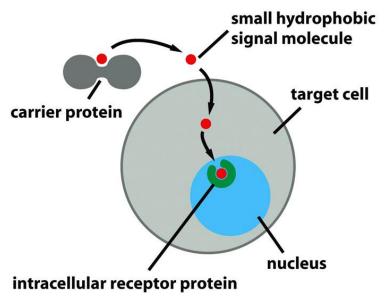
- ♥cell surface receptors
- ♥intracellular receptors

### Two different receptors work differently

#### **CELL-SURFACE RECEPTORS**



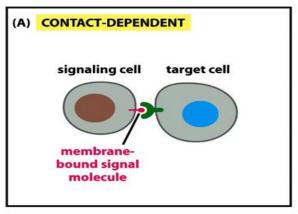
#### **INTRACELLULAR RECEPTORS**

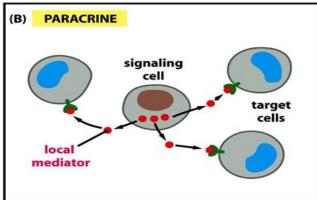


### Types of cell communication

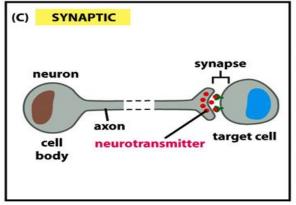
- Cell-cell contact
- Synaptic communication
- Paracrine/autocrine (local environment)
- Endocrine (long distance through blood stream)

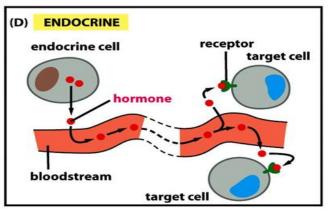
### Four types of cell communication





A and B are short-range





C and D are long-range

### Endocrine versus synaptic

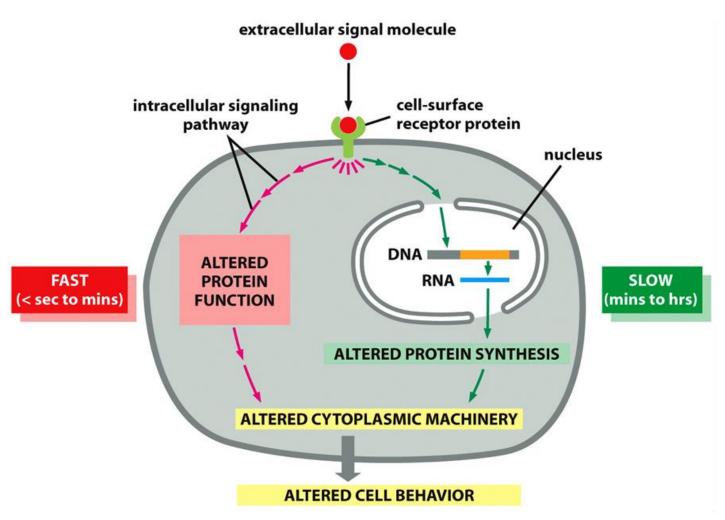
- Endocrine: need diffusion and blood flow, slow
- Synaptic: fast, 100meters/sec
- Endocrine: signals low concentration: 10^-8M and more diffused.
- Synaptic: signals higher concentration: 10^-4M and more precise.

### Effects in signaling can be slow and fast

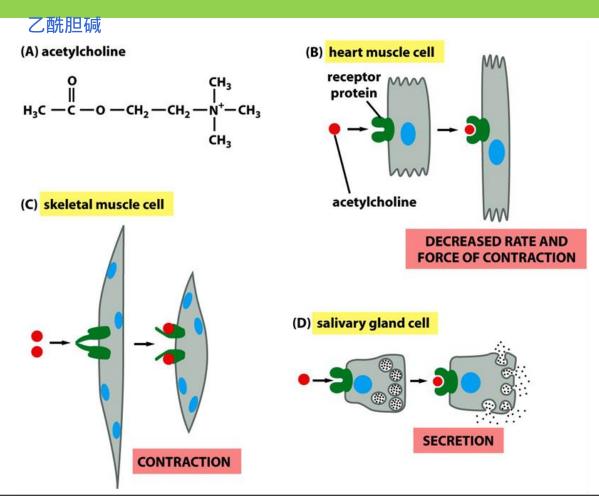
Slow: de novo protein synthesis in transcriptional response

Fast: change in protein behavior

### Slow and fast responses



### The same signals trigger different effects



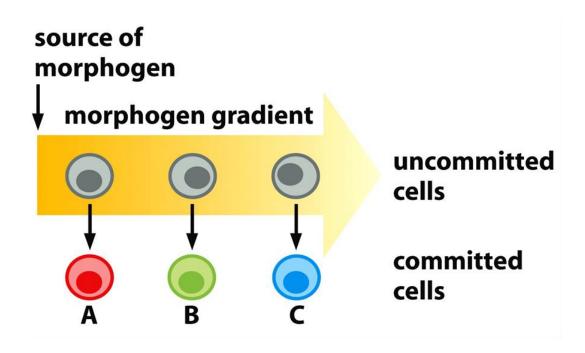
Acetylcholine receptors in heart muscle cells and salivary gland cells are identical However, they result in different effector proteins activation.

Acetylcholine receptors in heart muscle and skeletal muscle are different.

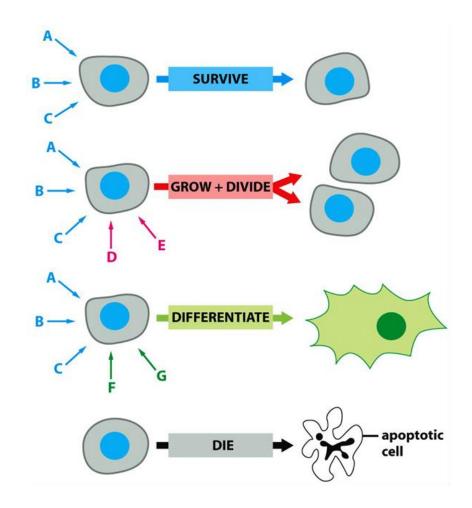
### The same signals trigger different effects

- ♥Same signals act on different receptors
- ♥Same signals act on same Receptors but trigger different effectors
- ◆The same cell type reacts differently to different concentration of signals---the signal in this case is called---morphogen.

### Morphogen in development



# Cell is programmed to respond to specific combinations of signals

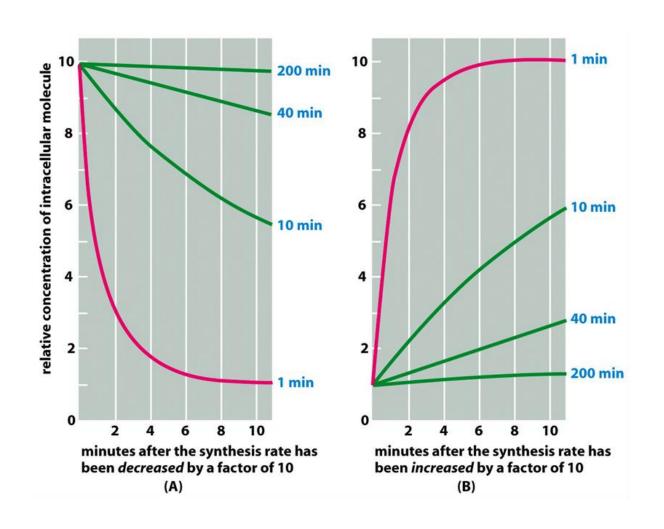


A cell can integrate multiple receptor signaling to dictate individual cell behavior.

## The amount and activity of signaling molecules are important

- Many proteins in signaling have short half lives---ensure quicker response
- •Many signaling proteins have conversion between inactive and active states---quicker response than de novo protein synthesis

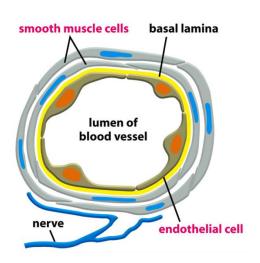
### Proteins that have faster turn over Rate react to stimuli in a faster manner



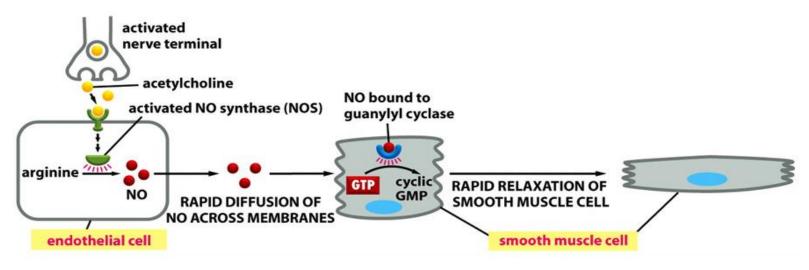
### II. Intracellular receptors

- •Signaling molecules are hydrophobic and can cross plasma membrane.
- •Examples:
  - 1. NO gas, CO gas
  - 2. steroid hormones, thyroid hormones, retinoids, vitamin D, etc.

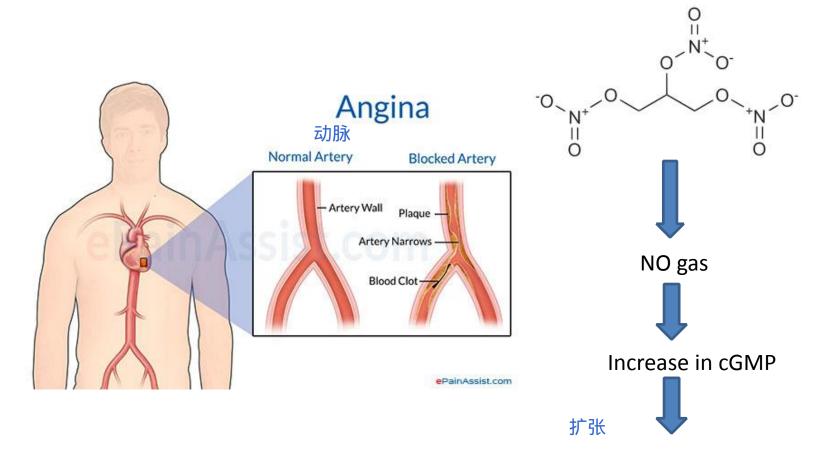
### 1. Signaling of NO in smooth muscle relaxation in blood vessel



NO has a half life of 5-10 sec. It is rapidly converted by water and oxygen into nitrates and nitrites

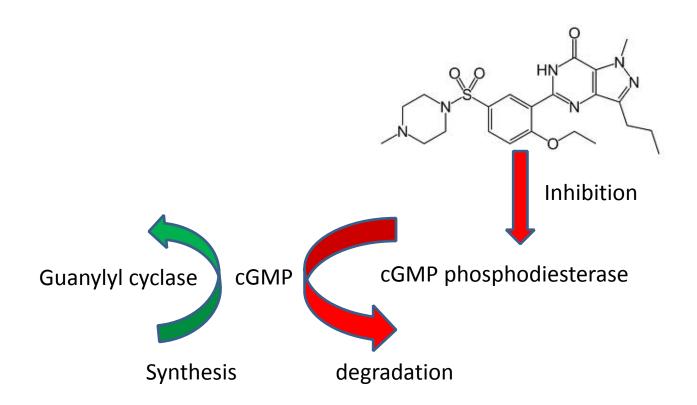


### Mechanism of nitroglycerin in treating angina pectoris



Dilation of blood vessel Reduce workload of heart

### Mechanism of Sildenafil---commercial name Viagra



Accumulation of cGMP causes prolonged blood vessel dilation

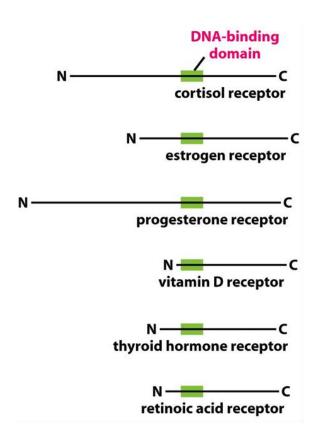
### 2. Signaling via nuclear receptor

- ◆ Steroid hormones (made from cholesterol):
  - ♥ cortisol (secreted from cortex to adrenal gland)
  - ♥Sex hormones (estradiol, testosterone, progesterone
  - ♥ Vitamin D (synthesized in the skin under sunlight)
  - ♥ molting hormone ecdysone (insects)
- ♦ Thyroid hormone: (made from tyrosine)
- ♦ Retinoids ( made from vitamin A)

# Some nongaseous signal molecules that bind to intracellular receptors

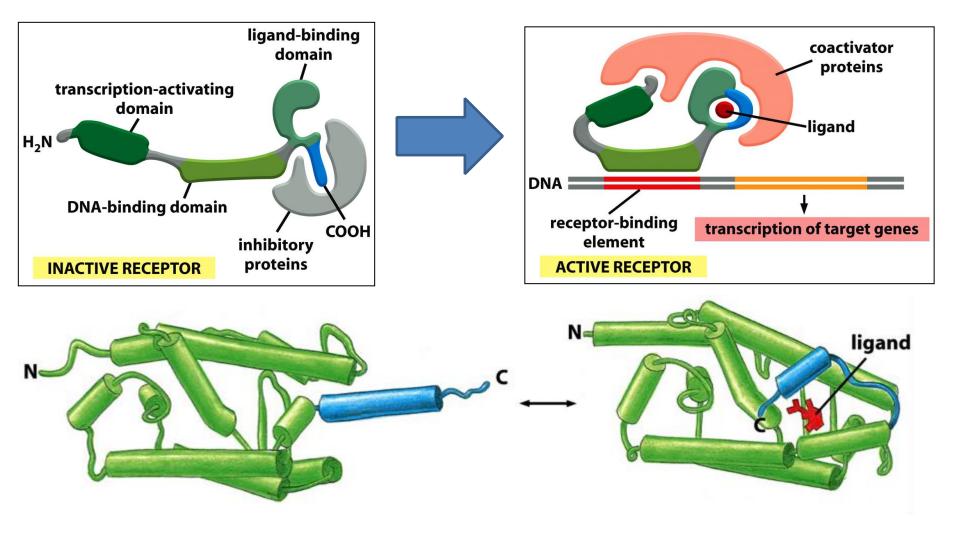
### Common features of nuclear receptors

- Work either as homodimer of heterodimer
- Serve both as ligand receptor and gene transcription factor

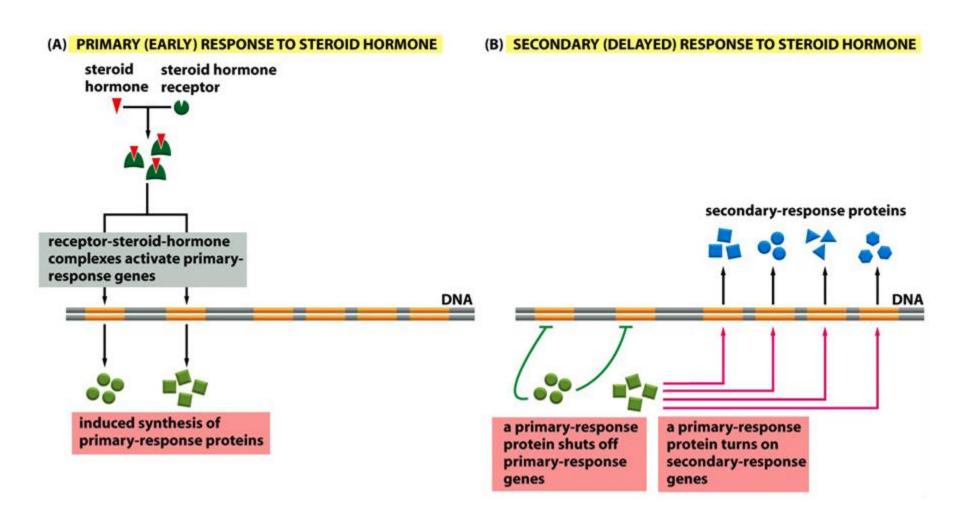


All these receptors have DNA binding domain and gene transactivation domain

### A model for how nuclear factor works



## Hormone receptors trigger both primary and secondary responses

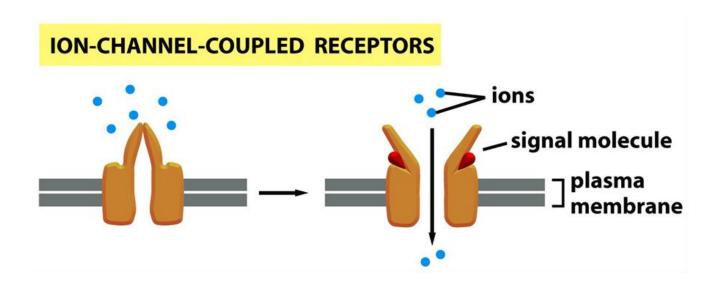


### III. Cell surface receptor signaling

- ♥ Ion-channel coupled receptor
- ♥ G-protein coupled receptor
- Enzyme-coupled receptorOther types

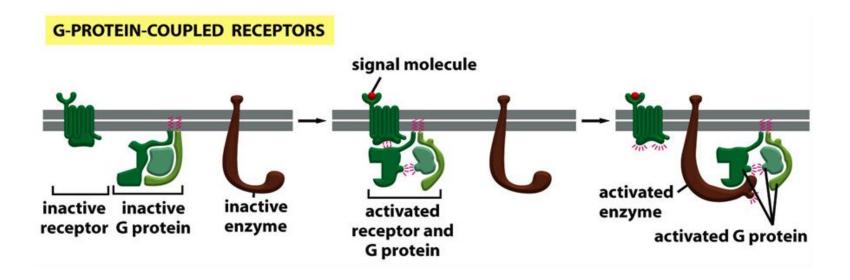
### The major three classes of cell surface receptors

1.



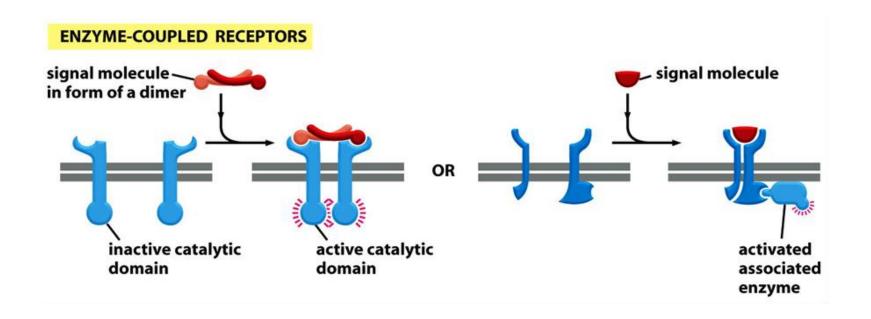
### The major three classes of cell surface receptors

2.



### The major three classes of cell surface receptors

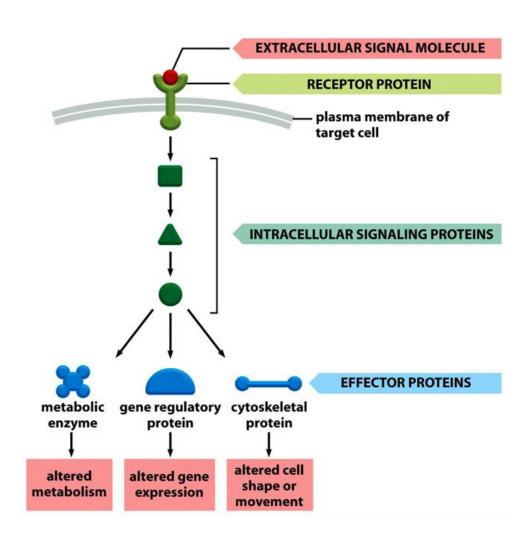
3.



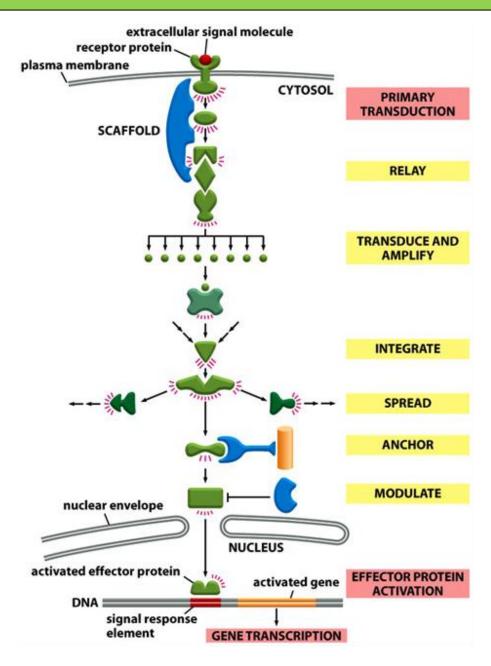
### The concept of second messenger

- ♦ The first messenger---extracellular signals
- ◆ Second messenger--- small molecules generated in large numbers after receptor activation. They are either hydrophilic or lipid diffusing.
- ♦ Second messenger work on effector proteins and relay signals.
  - **♥**cAMP
  - **♥**cGMP
  - **♥**Ca2+
  - ♥diacylglycerol (DAG)
  - ♥Inositol triphosphate (IP3)

### Relay of signals from cell surface receptors



### Diagram to show various functions for the signaling proteins

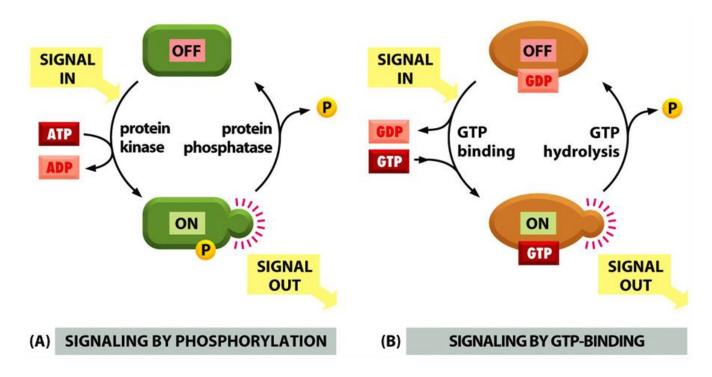


### Functions for intracellular signaling proteins

- Relay signals to the next component
- Act as a scaffold to bring two signaling proteins more quickly and efficiently
- transform the signal into a different form.
- Amply the signal it receives---signaling cascade
- Integrate signals from two or more pathways
- Spread signals from one pathway to another---crosstalk
- Anchor signaling proteins to a specific structure
- Modulate the activity of signaling proteins

### important types of switches to regulate protein activity

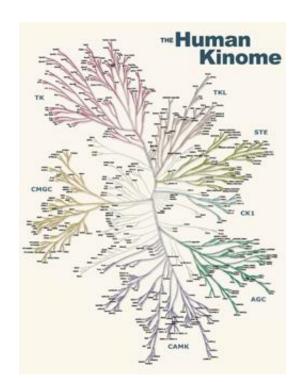
- Protein phosphorylation
- GTP-binding
- cAMP or Ca2+ binding
- Ubiquitination, etc.



# Protein phosphorylation

- ♥It is one major way of post-translational modification to regulate protein activity
- ♥>30% of all human genome proteins can be phosphorylated
- ♥>520 human kinases (kinome) and >150 protein phosphatases
- ♥Two categories: Serine/Threonine kinase; Tyrosine kinase
- ♥ Protein kinases are major therapeutic targets in human diseases

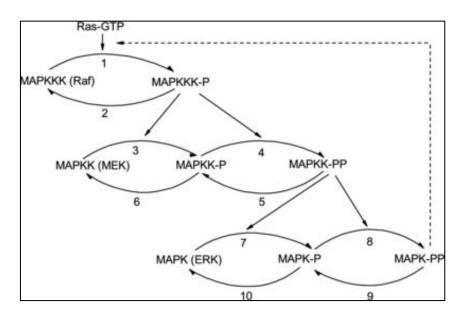
e.g.: Acute leukemia --- Gleevec targets BCR-ABL kinase



# Phosphorylation cascade

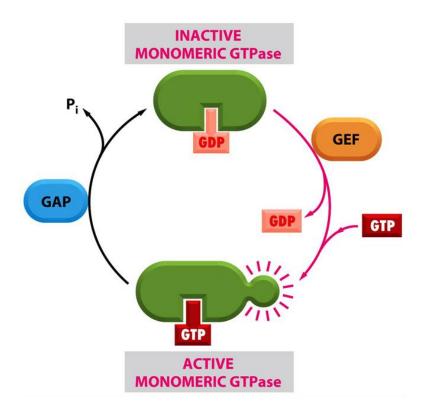
Signaling protein itself is a kinase which can phosphorylate and activate downstream effectors

For example: Ras-Raf-MAP kinase pathway



# GTP-binding proteins (G-proteins)

- •Large trimeric GTP-binding proteins
- •Small monomeric GTPase



# Characteristics of signal transduction

- Specifity
- Efficiency
- Reversibility
- Saturation
- High binding affinity

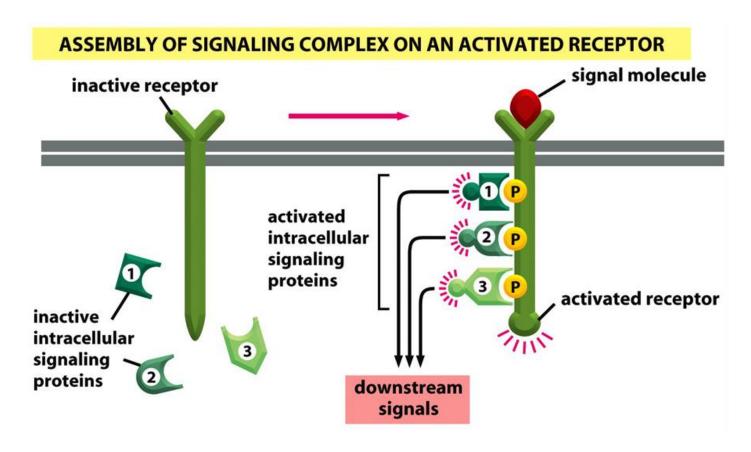
# How to achieve high speed and specificity in signaling

(1)

### PREFORMED SIGNALING COMPLEX ON A SCAFFOLD PROTEIN inactive receptor signal molecule activated receptor **CYTOSOL** plasma membrane scaffold protein inactive activated intracellular intracellular signaling proteins signaling proteins downstream signals

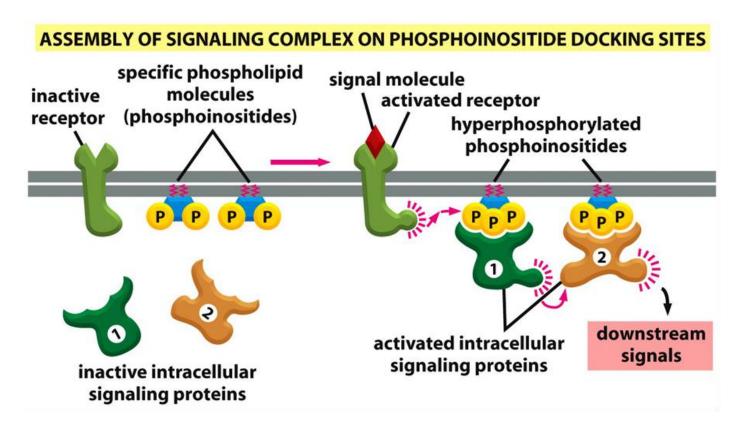
# How to achieve high speed and specificity in signaling

(2)



# How to achieve high speed and specificity in signaling

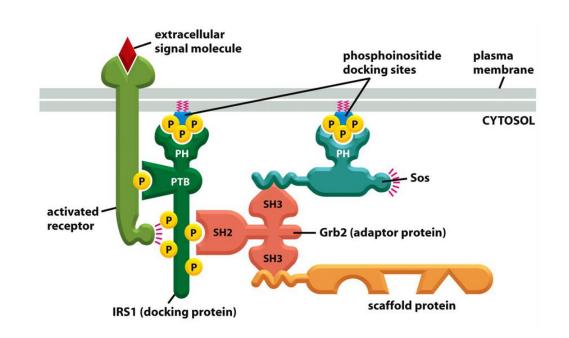
(3)



# Conserved *interaction domains* are important in protein binding

- Scr homology 2 (SH2) domain
- Phosphotyrosine-binding domains (PTB)
- Scr homology 3 (SH3): bind proline rich domain
- Pleckstrin homology (PH): bind phosphoinositides

Diagram to Show how Domains Mediate the Interaction:



Bind phosphotyrosine

# IV. Several methods to study signal transduction

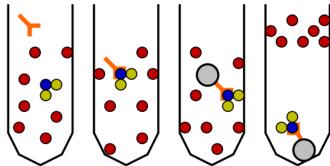
- Protein co-immunoprecipitation
- Western blotting, phospho-specific antibody
- In vitro protein activity studies
- shRNA/siRNA, inhibitors
- Rescue analysis

### 1. co-IP

applications: receptor-ligand interaction

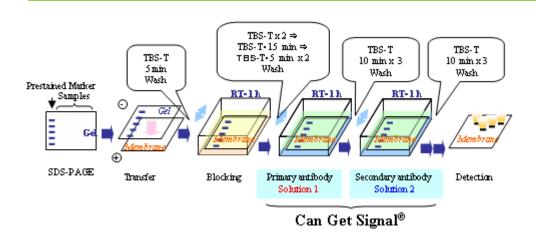
kinase-substrate interaction

other protein interaction partners



- 1. Řesearcher adds antibody.
- 2. Antibody binds target.
- 3. Protein A beads bind antibody.
- 4. Centrifugation sediments beads.

# 2. Western blotting



#### **Detection in Western Blots**

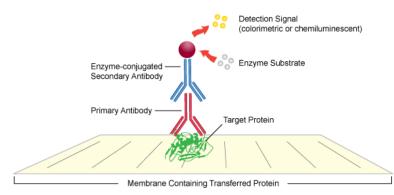
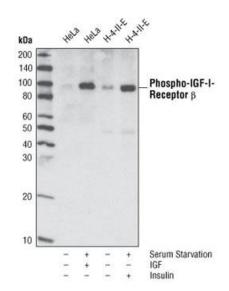


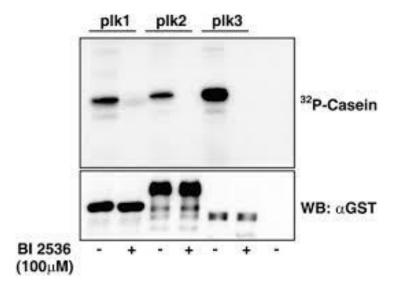
Diagram 2: Illustration of detection in Western Blots.

e. g. IGF receptor activation



# 3. In vitro protein activity assay

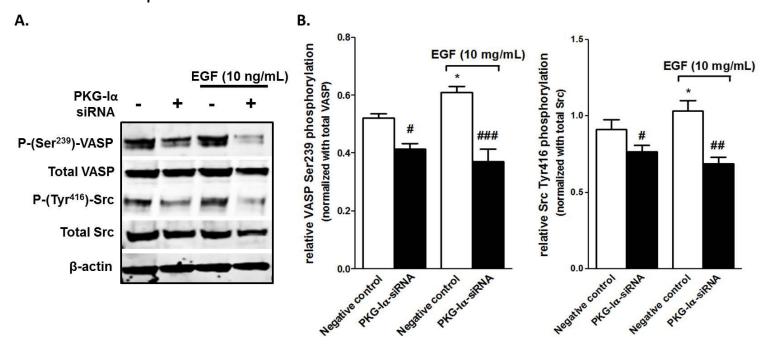
- (1). Purify protein in vitro
- (2). Set up in vitro protein assay with substrates and necessary components such as ATP, etc
- (3). Analyze protein activity by comparing signal strength.



# 4. shRNA/siRNA, inhibitors

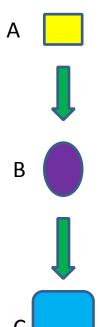
- shRNA- short hairpin RNA, siRNA-small interference RNA
- They work by triggering target mRNA degradation.
- Many enzymes have relatively specific inhibitors.

For example:



# 5. rescue assay

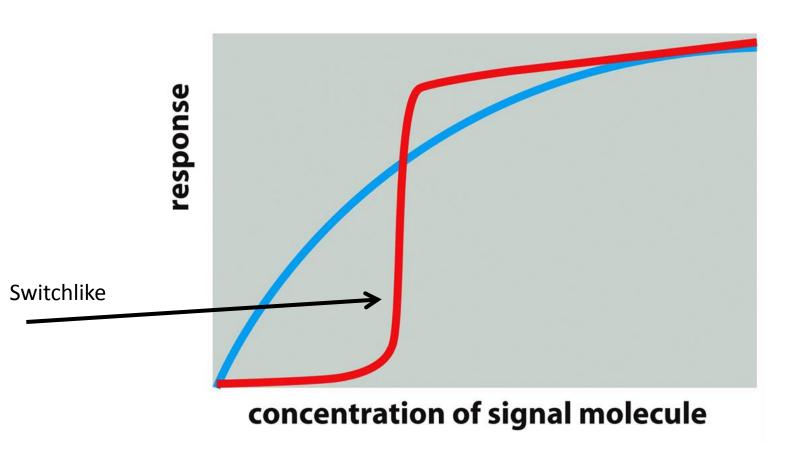
 How to prove one signaling protein locate upstream of downstream of another?



- 1. Deletion of A or B leads to a certain defect
- 2. Expression of activated C can rescue this defect.

# V. signaling kinetics

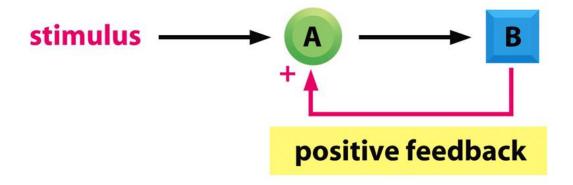
# Signaling can be both *All or none* and *smoothly graded* response

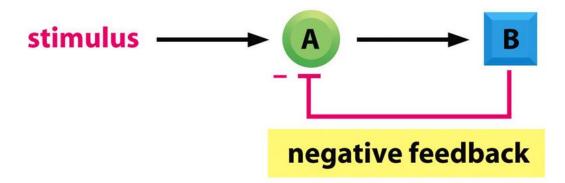


# What causes switchlike responses?

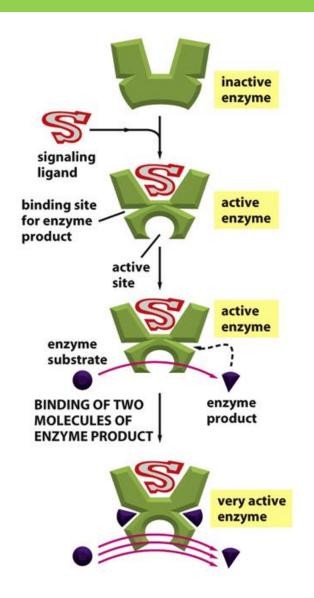
- ♥ All of none could be due to cooperative response(e.g. 4 cAMPs bind to PKA)
- ♥ Or it could be due to concerted effect of a simultaneous inhibition for the opposite reaction.
- ♥ It needs positive feedback response.

# Positive and negative feedback

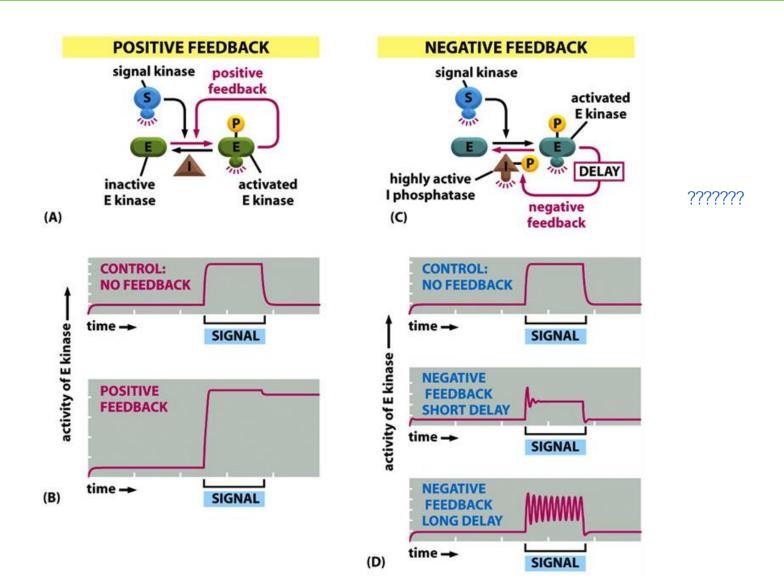




# Positive feedback gives switchlike response



# Different results from positive and negative feedback



# Negative feedback allows adaptation/desensitization for cells

- Detects changes of concentration of signals.
- There are several ways to achieve these:

