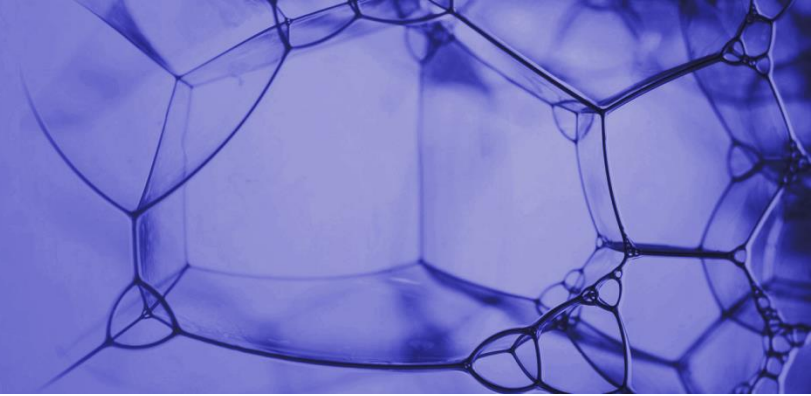


**LOSCHMIDT  
LABORATORIES**



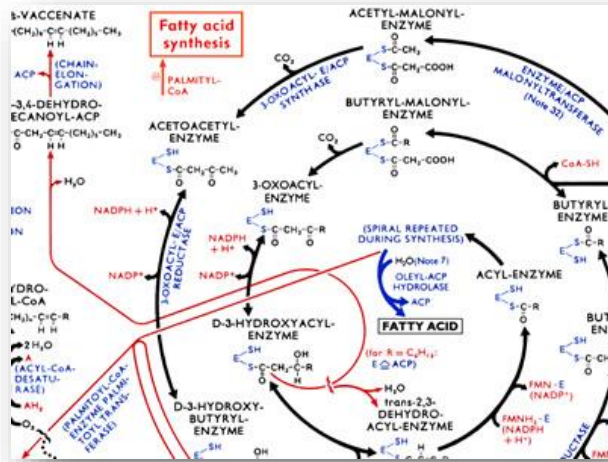
## **7. Molecular Biotechnology in Industry**

# Outline

- ❑ Enzymes and applications
- ❑ Definition of white biotechnology
- ❑ Sustainable development
- ❑ Enzyme sources
- ❑ Industrial production of proteins
- ❑ Enzyme and cells immobilization
- ❑ Examples of biocatalytic applications

# Enzymes

- ❑ natural catalysts (biocatalyst)
- ❑ catalyze chemical reactions in living systems



- **oxidoreductases** - oxidation/reduction
- **transferases** - transfer of functional groups
- **hydrolases** – hydrolytic cleavage
- **lyases** - cleavage of C-C, C-N and C-O bonds
- **isomerases** - racemization, epimerization
- **ligases** - formation of C-C, C-N and C-O bonds

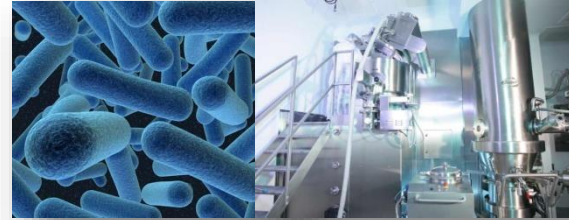
# Enzyme applications

restrictases  
DNA ligases  
polymerases



phosphatases  
peroxidases

amylases  
proteases  
cellulases  
phytases  
lipases



lipases  
nitrilases  
peptidases  
amidases  
aldolases

asparaginase  
DNase  
urokinases  
proteases



cellulases  
ligninase  
lipases

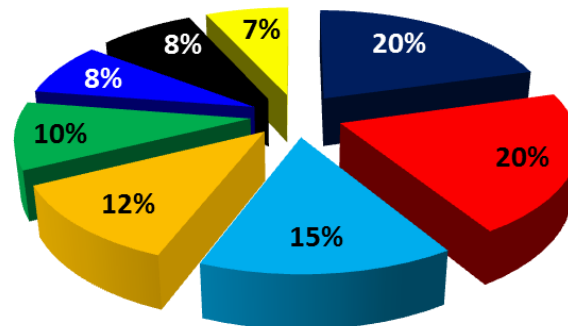
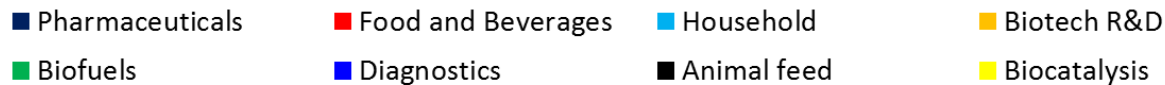
amylase  
cellulases  
catalase



dehalogenases  
choline esterase  
peroxidases

# White (industrial) biotechnology

- ❑ biotechnology incorporated into production processes and products that **involve chemical reactions - biocatalysis**
- ❑ **sustainable** and **environmentally-friendly** industry
- ❑ provide **energy efficiency**, increased **productivity** and better **safety**
- ❑ uses **enzymes** and **micro-organisms** to make products and services in a wide range of industrial sectors



# Enzyme-based technologies

## ADVANTAGES

- ❑ high catalytic efficiency
- ❑ broad substrate specificity
- ❑ high selectivity
- ❑ compatibility of each other
- ❑ reusability
- ❑ sustainability
  - produced from biomass
  - non-toxic and biodegradable
  - operate at mild conditions
  - less byproducts and wastes

## LIMITATIONS

- ❑ cofactor requirement
- ❑ prone to inhibitions
- ❑ highest activity in water
- ❑ less stable
- ❑ low selectivity
- ❑ expensive



# Enzyme sources

## ❑ animal and plant tissues

- thousands years old developed empirically
- pancreas (treatment of hides), calf stomach (cheese-making)
- papaya, pineapple (meat tenderization)
- **content up to 1%** enzyme of tissue weight
- **less competitive** compared to fermentation of microorganism
- **risk of contamination** with prions and viruses harmful to humans

<i>Source</i>	<i>Enzyme</i>	<i>Application</i>
<b>Animal tissues</b>		
Bovine and porcine pancreas	proteases (e.g., trypsin, chymotrypsin), amylases, lipases	digestive enzymes, anti-inflammatory agents, health food additives
Porcine stomach	pepsin	body fortifying agents
Liver and muscle	aldolases	fructose digestion
Porcine kidney	D-aminoacid oxidase	
<b>Plant tissues</b>		
Pineapple stem	bromelain (mixture of proteases)	anti-inflammatory agents, meat tenderizer
Papaya latex	papain (protease)	anti-inflammatory agents
Aspegillus	proteases, lipases, amylases, cellulases	natural food supplements, digestive enzymes

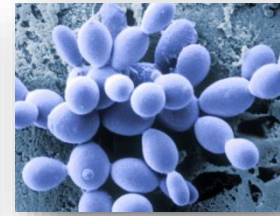
# Enzyme sources

## ❑ wild-type microorganisms

- enzymes from microorganisms long been safely used in food industry
- food processing regulations - strict for non-recombinant enzymes
- microorganisms used for screening for „new“ catalytic enzymes
- screen for enzymes active at desired process conditions  
(e.g., pH, temperature)

## ❑ recombinant microorganisms

- most technical enzymes produced using **recombinant technology**
- when yield in wild type organism is low or desired enzyme is not in class I organism
- bacteria, fungi and yeasts  
(e.g., *E.coli*, *Bacillus*, *Aspergillus*, *Saccharomyces*)

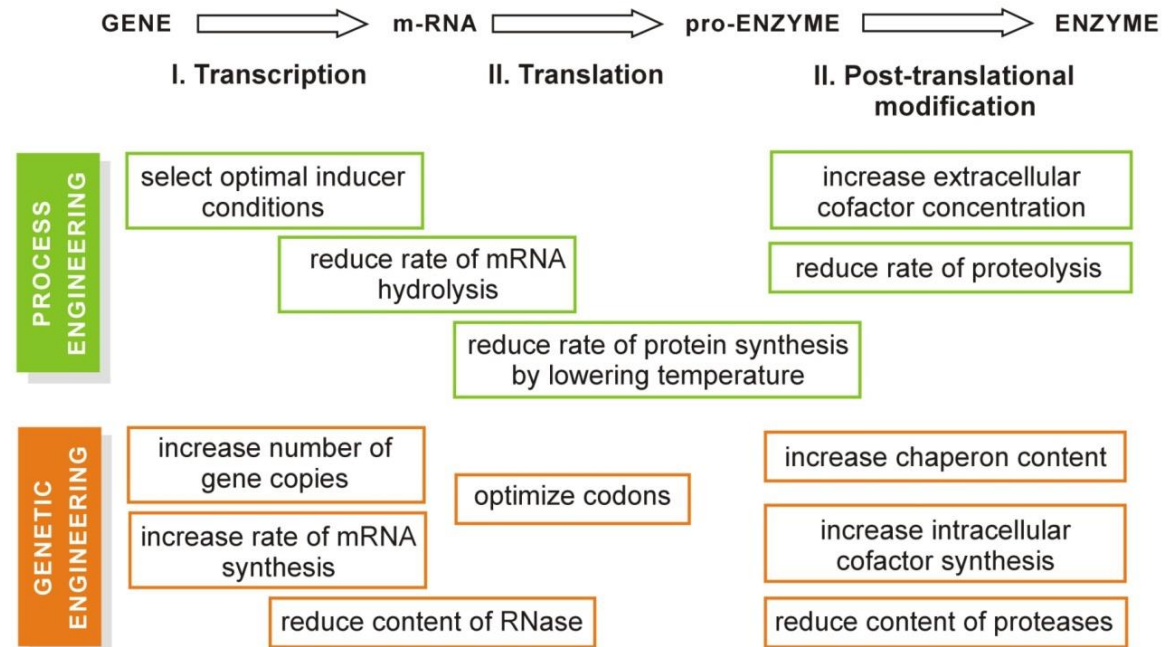




# Industrial production of proteins

## □ fermentation

- non-recombinant and recombinant organisms
- steady and safe (class I or GRAS) organisms
- up-scale and optimization
- **high cell density** fermentation (50 g cell dry weight per liter)
- upper limit of **protein concentration** ( $10 \text{ g.L}^{-1}$ ; 40% of total cell protein)



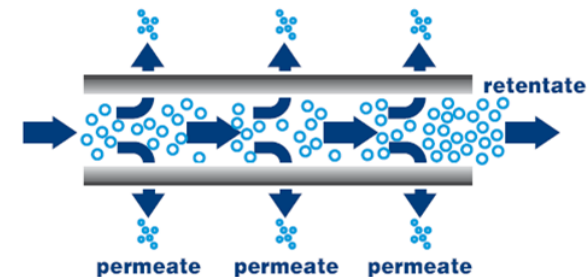
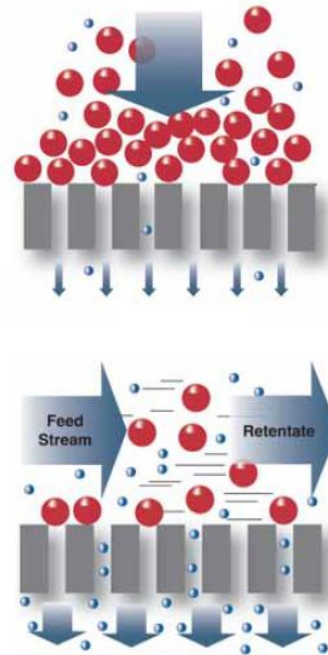
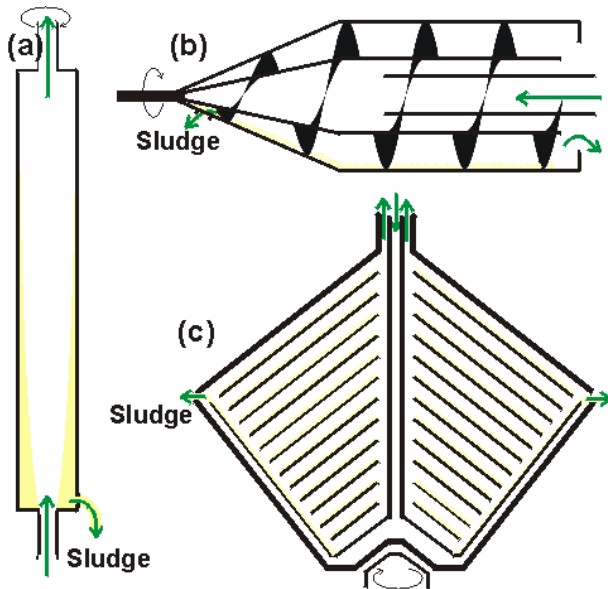
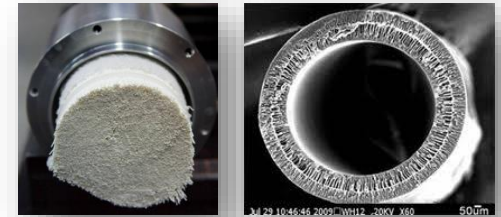
# Downstream process

## □ separation and homogenization

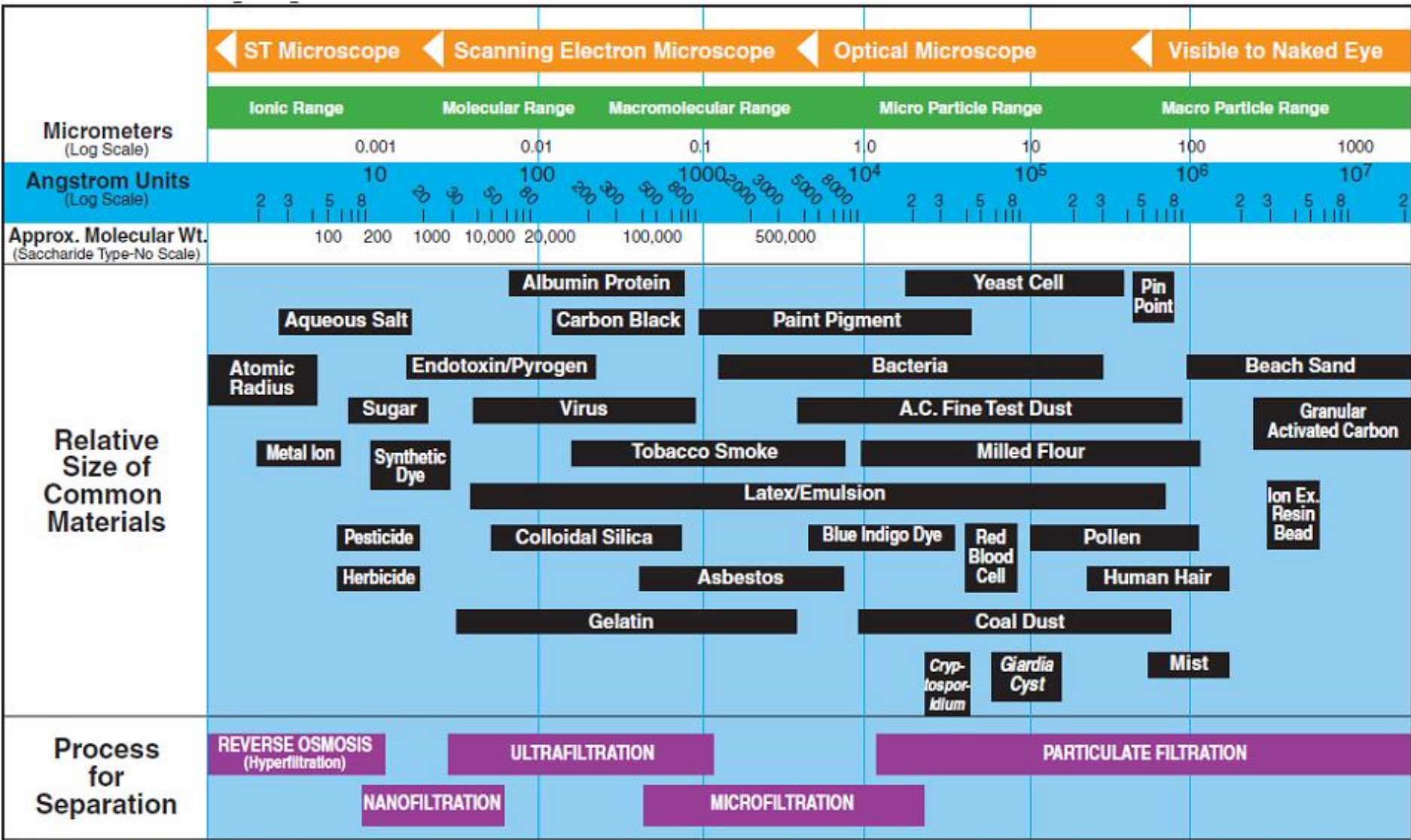
- dependent on application and required purity
- **technical enzymes** - low to moderate purity
- proteins for **therapy** and **diagnostics** - high purity

Intracellular                      Periplasmatic                      Extracellular

separation of cells from medim by centrifugation or microfiltration



# Downstream process



Note: 1 Micron (1 x 10<sup>-6</sup> Meters) = 4 x 10<sup>-5</sup> Inches (0.00004 Inches)  
 1 Angstrom Unit = 10<sup>-10</sup>Meters = 10<sup>-4</sup> Micrometers (Microns)

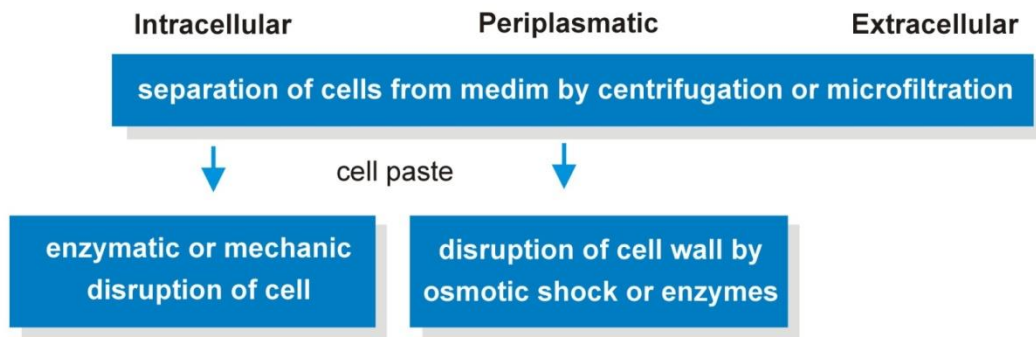
© Copyright 1998, 1996, 1993, 1990, 1984 Osmonics, Inc., Minnetonka, MN, USA



# Downstream process

## □ separation and homogenization

- dependent on application and required purity
- **technical** enzymes - low to moderate purity
- proteins for **therapy** and **diagnostics** - high purity



### MECHANICAL

- **ultrasonic disruption** - cell lysis with high frequency sound
- **homogenizers** - high pressure (1500 bar) and expansion
- **freeze fracturing** - water crystals as abrasive
- **ball mills** and **blenders**

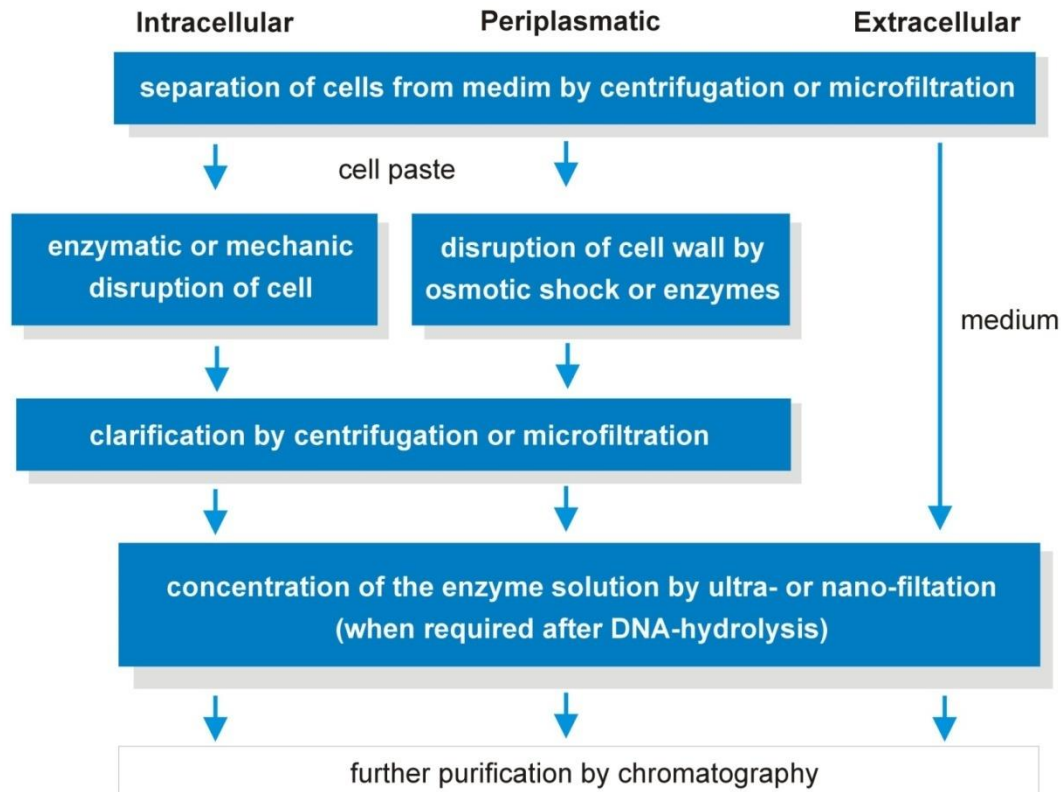
### NON-MECHANICAL

- **osmotic shock** (e.g., high sucrose medium)
- **chemical permeabilization** (e.g., solvents, surfactants, antibiotics)
- **enzymatic permeabilization** (e.g., glycanases, proteases, mannanase)

# Downstream process

## □ separation and homogenization

- dependent on application and required purity
- **technical** enzymes - low to moderate purity
- proteins for **therapy** and **diagnostics** - high purity



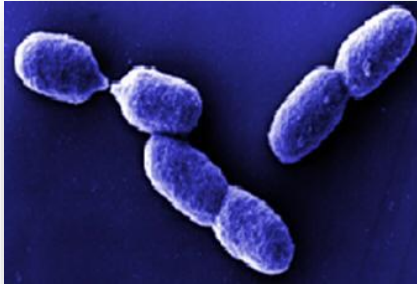
# Downstream process

## ❑ enzyme purification

- **impurities** (e.g., proteins, DNA and others)
- further purification when **safety** (e.g., recombinant DNA, viruses) or **function** reasons (impurities disturbing catalytic function)
- basic knowledge of **protein properties** necessary
  - molecular weight (MW)
  - isoelectric point (pI)
  - cofactors
  - pH range
  - temperature stability
- **methods** of protein purification
  - precipitation and differential solubilization (e.g., ammonium sulfate, pH, solvents)
  - membrane filtration
  - chromatographic methods (e.g., size exclusion, ion exchange, hydrophobic, metal affinity, biospecific)
- **more steps -> higher purity** (each step loss >10% of enzyme)



# Whole cell vs. isolated enzyme

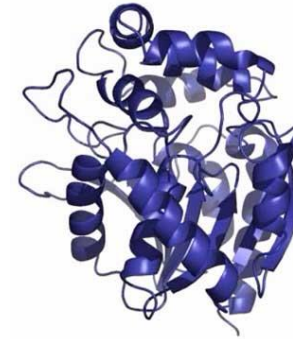
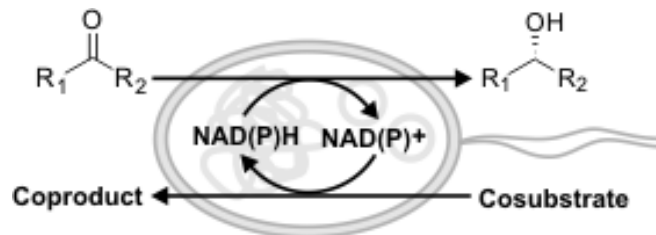


## □ advantages

- allow more enzymes
- cofactor regeneration
- cheap

## □ disadvantages

- side-reactions
- low tolerance to solvents
- low productivity

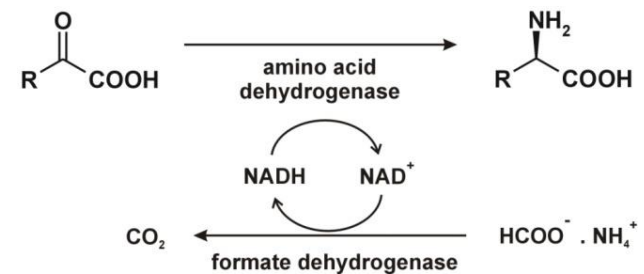


## □ advantages

- smaller reactors
- less side reactions
- higher productivity

## □ disadvantages

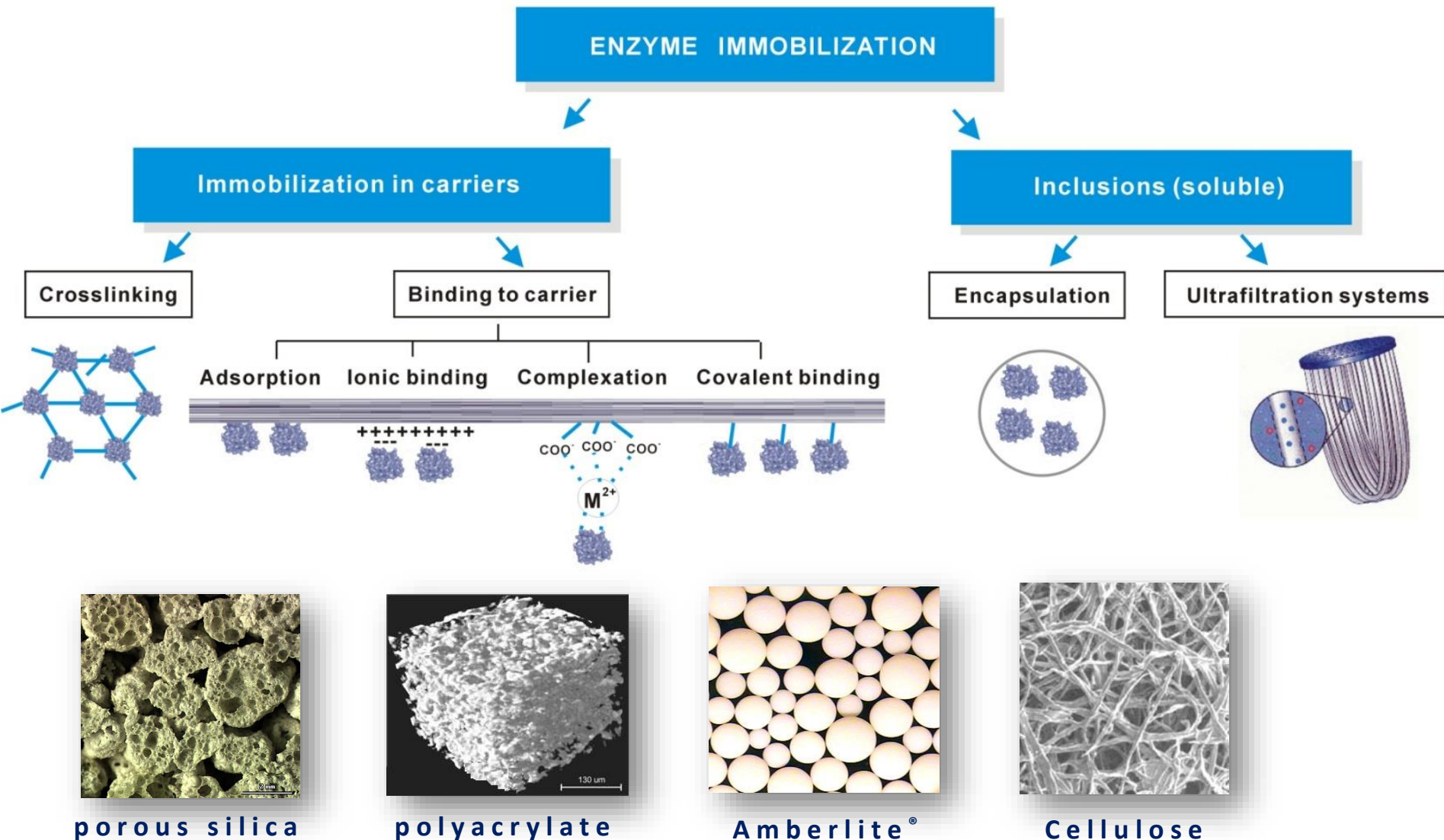
- more expensive
- addition of cofactors
- less stable outside cell



# Immobilisation methods

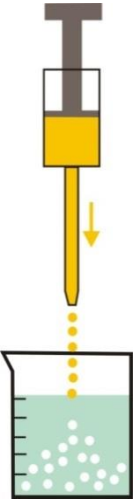
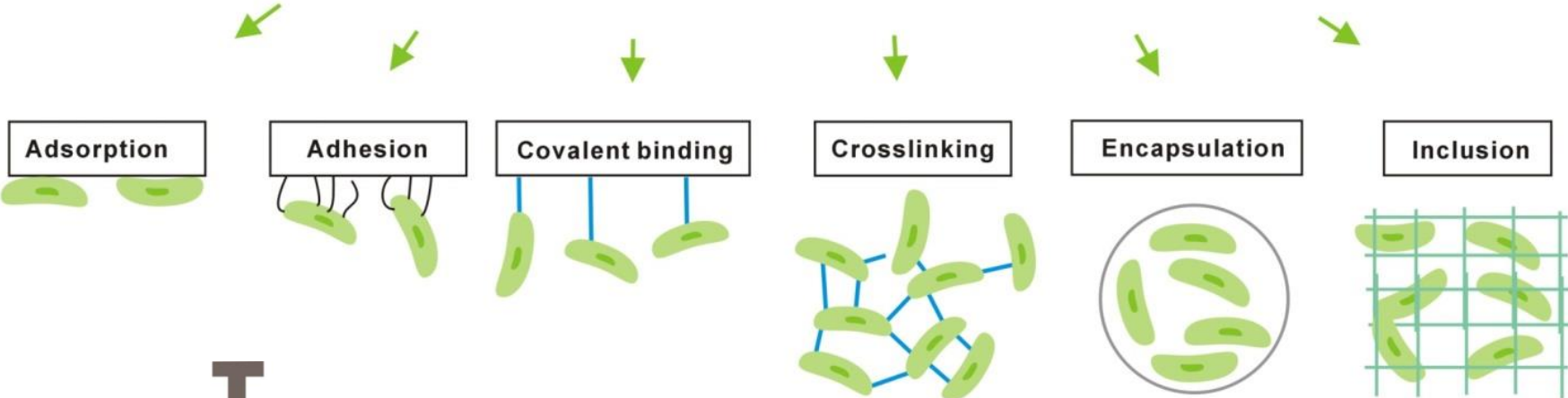
- ❑ biocatalysts (enzyme or cell) **limited in moving** due to **chemical** or **physical treatment**
  
- ❑ **benefits**
  - **stabilization** by immobilization
  - **easy separation** of product
  - **repeated use** of biocatalyst
  - **continuous** bioprocessing
  
- ❑ **limitations**
  - **expenses** of carriers and immobilization
  - **activity loss** during immobilization
  - **changes in properties** of biocatalyst
  - **mass transfer** limitations

# Immobilisation of enzyme

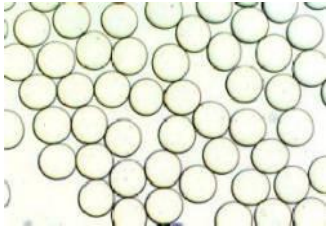


# Immobilisation of cell

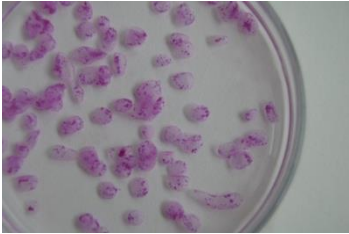
## MICROORGANISM AND CELL IMMOBILIZATION



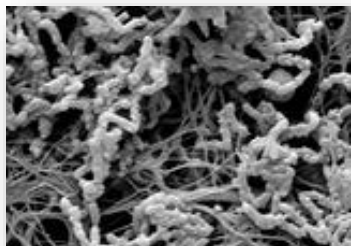
**Alginate beads**



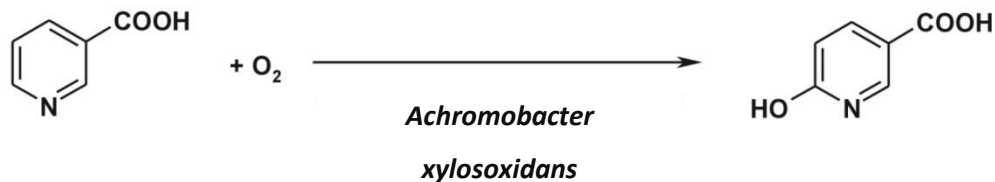
**PVA lens (LentiCats)**



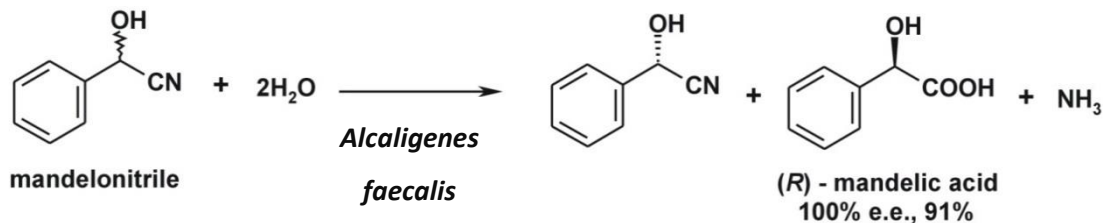
# Examples of whole cell biocatalysis



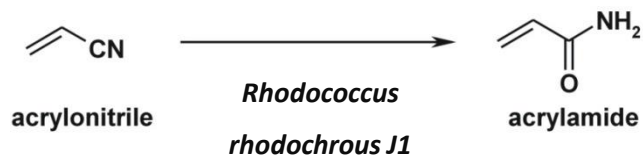
- synthesis of **agrochemical intermediates** by microbial hydroxylation of heteroatomics (Lonza)



- mandelic acid - **urinary antiseptic, skin care cosmetics** (du Pont, Nitto Chemicals, etc.)



- large-scale production of **commodity chemical** - acrylamide (Mitsubishi, Nitto Chemicals)

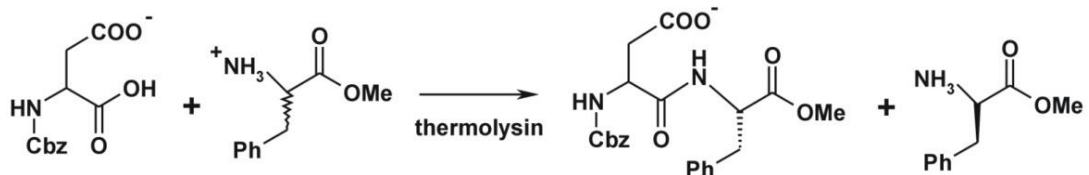




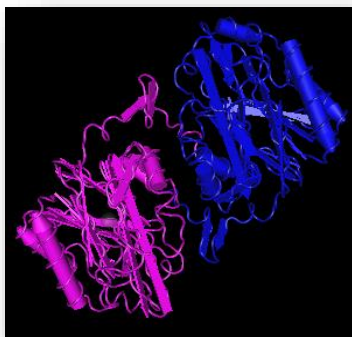
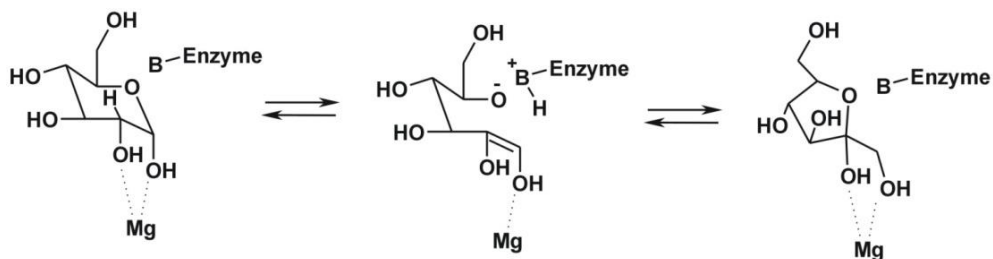
# Examples of enzyme biocatalysis



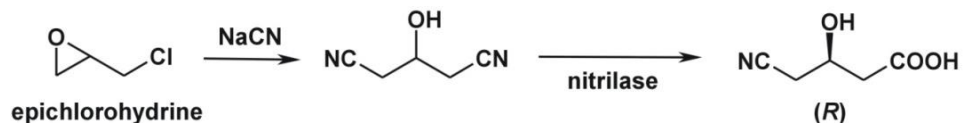
- large scale production of **Aspartame**, low-calorie sweetener (DSM, NutraSweet)



- synthesis of **high fructose syrup** from corn starch (10 million tons per year)



- synthesis of **atorvastatin, Lipitor<sup>®</sup>**, intermediate (Pfizer - sales since 1996 exceed US\$ 150 billion)





- ❑ **Enzymes at work** (Novozymes, Denmark)
  - 1. Why use enzymes for industrial processes?
  - 2. The nature of enzymes
  - 3. Industrial enzyme production

