



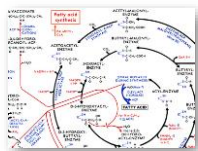
7. Molecular Biotechnology in Industry

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

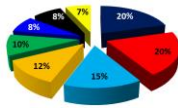
Enzyme applications

<ul style="list-style-type: none"> restrictases DNA ligases polymerases 		<ul style="list-style-type: none"> phosphatases peroxidases
<ul style="list-style-type: none"> amylases proteases cellulases phytases lipases 		<ul style="list-style-type: none"> lipases nitrilases peptidases amidases aldolases
<ul style="list-style-type: none"> asparaginase DNase urokinases proteases 		<ul style="list-style-type: none"> cellulases ligninase lipases
<ul style="list-style-type: none"> amylase cellulases catalase 		<ul style="list-style-type: none"> 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

■ Pharmaceuticals ■ Food and Beverages ■ Household ■ Biotech R&D
 ■ Biofuels ■ Diagnostics ■ Animal feed ■ Biocatalysis



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

Source	Enzyme	Application
Animal tissues		
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	
Plant tissues		
Pineapple stem	bromelain (mixture of proteases)	anti-inflammatory agents, meat tenderizer
Papaya latex	papain (protease)	anti-inflammatory agents
Aspergillus	proteases, lipases, amylases, cellulases	natural food supplements, digestive enzymes

Enzyme sources

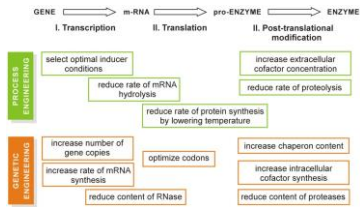
- ❑ **wild-type microorganisms**
 - enzymes from microorganisms long been safely used in food industry
 - food processing regulations - strict for non-recombinant enzymes
 - microorganisms used for screening for „new“ catalytic enzymes
 - screen for enzymes active at desired process conditions (e.g.,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

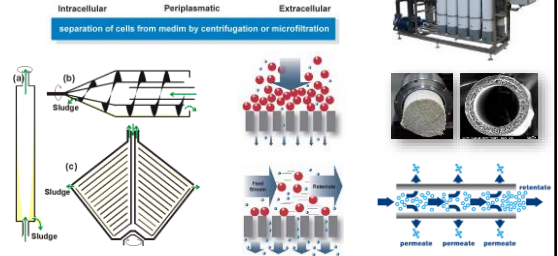
- non-recombinant and recombinant organisms
- steady and safe (class I or GRAS) organisms
- up-scale and optimization
- high cell density fermentation (50 g cell dry weight per liter)
- upper limit of protein concentration (10 g.L⁻¹; 40% of total cell protein)



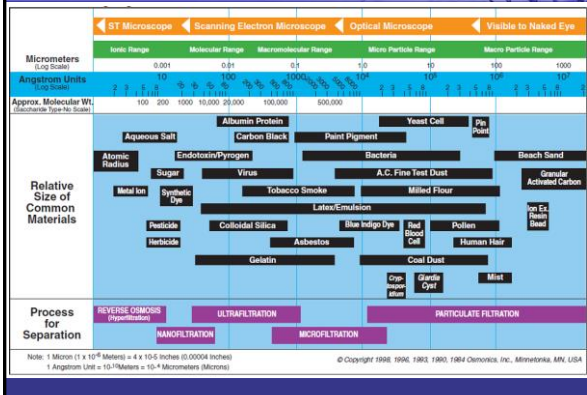
Downstream process

separation and homogenization

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



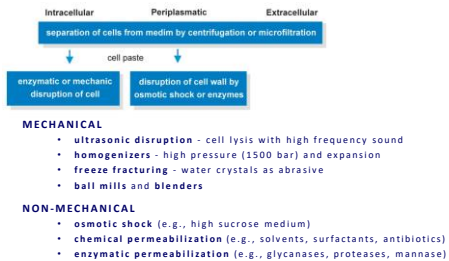
Downstream process



Downstream process

separation and homogenization

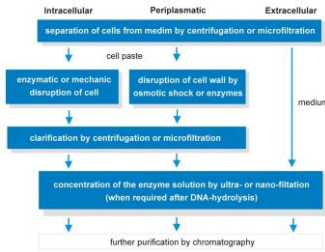
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Downstream process

□ 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

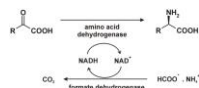


□ advantages

- smaller reactors
- less side reactions
- higher productivity

□ disadvantages

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



Immobilisation methods

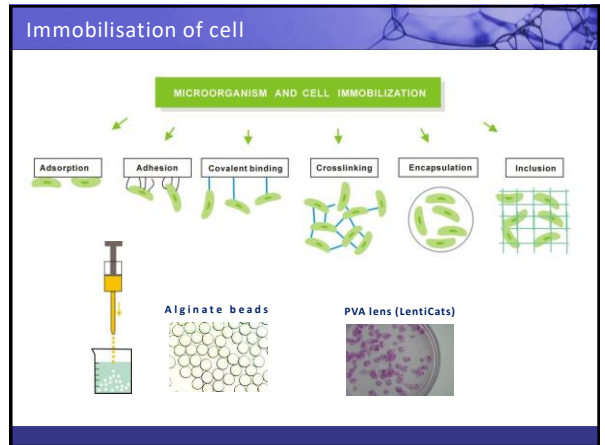
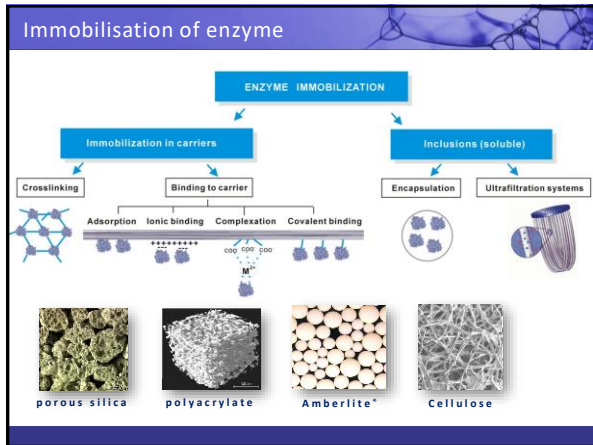
- biocatalysts (enzyme or cell) **limited in moving** due to **chemical or physical treatment**

□ benefits

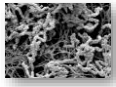
- **stabilization** by immobilization
- **easy separation** of product
- **repeated use** of biocatalyst
- **continuous bioprocessing**

□ limitations


- **expenses** of carriers and immobilization
- **activity loss** during immobilization
- **changes in properties** of biocatalyst
- **mass transfer limitations**




Examples of whole cell biocatalysis

- 

synthesis of **agrochemical intermediates** by microbial hydroxylation of heteroatoms (Lonza)

Cc1ccc(C(=O)O)cc1.O=O>>Oc1ccc(C(=O)O)cc1
Achromobacter xylooxidans
- 

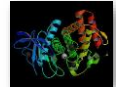
mandelic acid - **urinary antiseptic, skin care cosmetics** (du Pont, Nitto Chemicals, etc.)

Oc1ccc(C#N)cc1.O.O>>Oc1ccc(C(=O)O)cc1.N
Alcaligenes faecalis (R)-mandelic acid 100% e.e., 91%
- 


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

C#CC=C.N>>NC(=O)C=C
Rhodococcus rhodochrous II

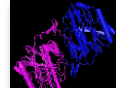
Examples of enzyme biocatalysis

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

COC(=O)N[C@@H](Cc1ccc(C(=O)O)cc1)C(=O)O.NC(=O)C(=O)O>>COC(=O)N[C@@H](Cc1ccc(C(=O)O)cc1)C(=O)N[C@@H](Cc2ccc(C(=O)O)cc2)C(=O)O
thermolysin
- 

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

OC[C@H]1O[C@H](O)[C@H](O)[C@@H](O)[C@H]1O>>OC[C@H]1O[C@H](O)[C@H](O)[C@@H](O)[C@H]1O
Mg
- 

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

C1=CC=C(C=C1)C(=O)Cl.NaCN>>NC(=O)C(=O)C1=CC=C(C=C1)C(=O)O
nitrilase (R)

Reading

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

