

BIOLOGY, BIOTECHNOLOGY

in English


2 hour lecture/week, 3 credits

2 midterm tests, no final examination

12 lectures, 3 lecturers

Handouts, slide shows and readings:


http://oktatas.ch.bme.hu/oktatas/konyvek/abet/Biology-biotechnology_in_English/



1

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Date	Lecture	Topic	Lecturer	tests	room
1-Mar	1	Cells	M. Pécs		
8-Mar	2	Industrial microbiology	Á. Németh		
15-Mar		National Holiday			
22-Mar	3	Enzymes	M. Pécs		
29-Mar	4	Enzymes	M. Pécs		
05-Apr	5	Microbial growth	Á. Németh		
12-Apr		Spring Holiday			
19-Apr	6	Aeration, agitation	Á. Németh		
26-Apr	7	Sterilization	Á. Németh	midterm test 1	
3-May	8	Downstream processing	M. Pécs		
10-May	9	Technologies, case studies	M. Pécs		
17-May	10	Wastewater treatment	V. Bakos		
24-May	11	Wastewater treatment	V. Bakos		
31-May	12			midterm test 2	
07-Jun				makeup tests	



2

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Lecturers:

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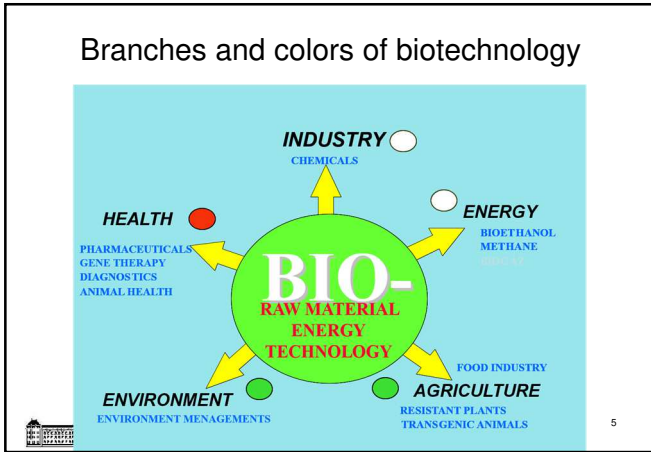
3

BIOLOGY, BIOTECHNOLOGY

Biology: everybody knows - a natural science dealing with living beings.
 But what is Biotechnology?
 ... is an integrated application of
 biochemistry,
 microbiology and
 engineering sciences
 ... principles in order to the technological use of
 microorganisms
 animal and plant cells/tissues
 or parts of these (e.g. enzymes)
 ...to produce something.



4



1st lecture: Composition and structure of cells

1. Prokaryotes and eukaryotes


Karyon = nucleus pro- = before/first eu- = true/good

Basic difference: they don't have/have real, isolated nucleus

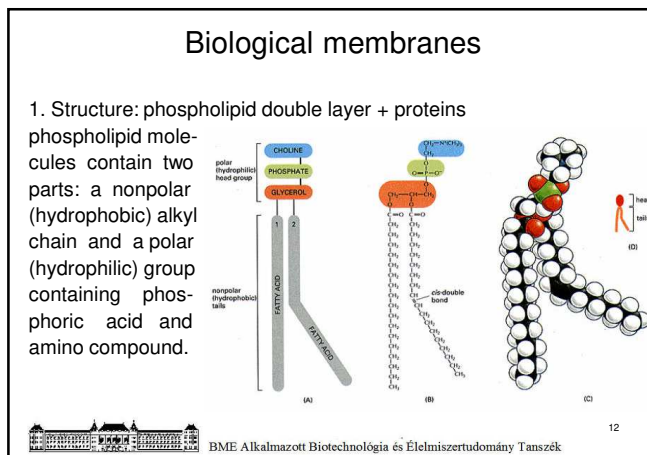
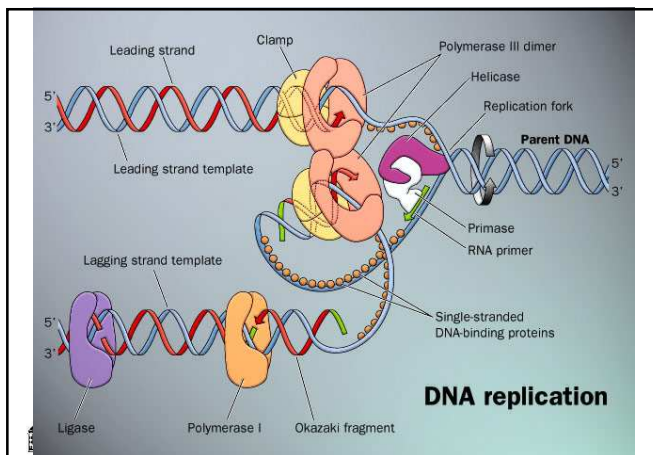
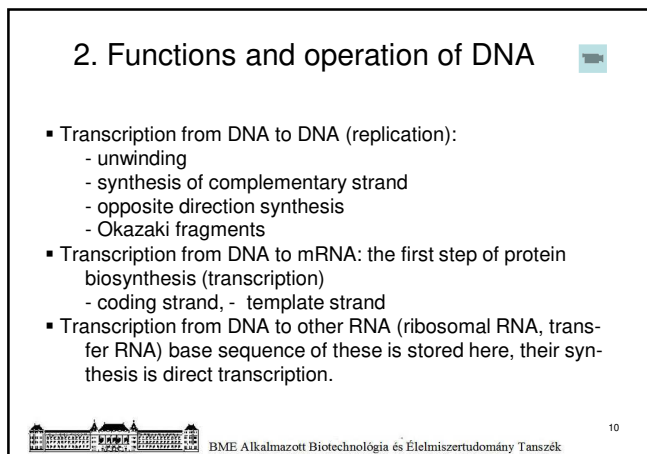
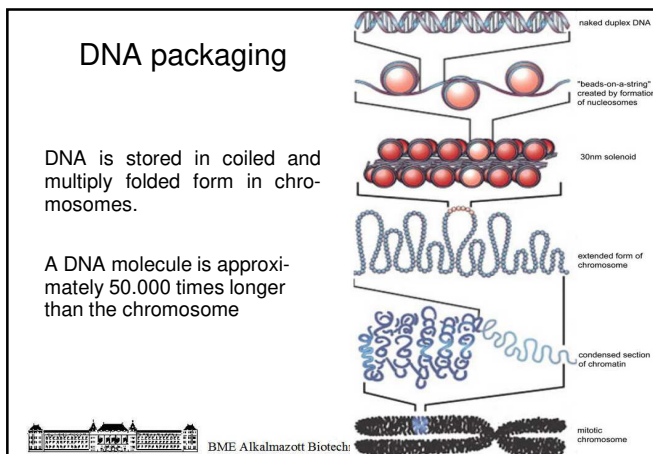
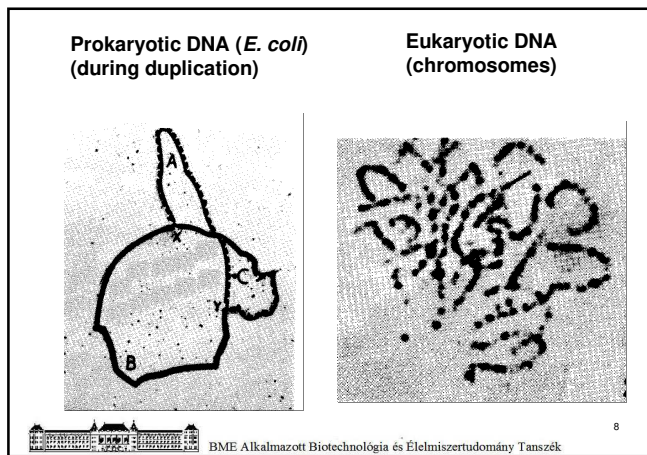
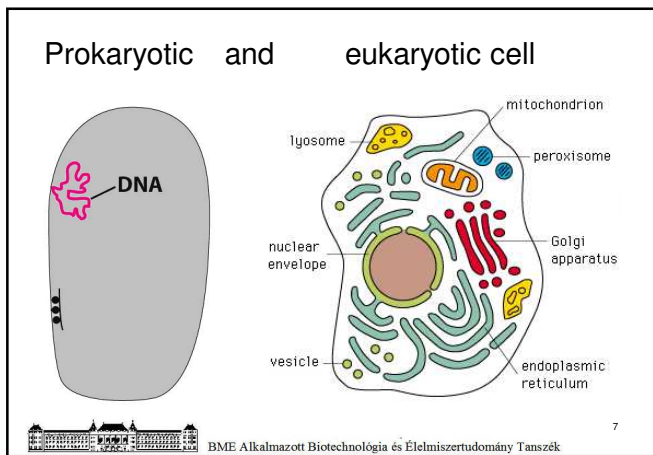
In the evolution: the prokaryotes are ancient, simple forms, the eukaryotes are more complex and evolved later

Prokaryotes: all bacteria, included the filiform Actinomycetales and blue algae (Cyanobacteriales)

Eukaryotes: yeasts, moulds, protozoa, green algae, and all multicellular living being.



6



The structure of double layer

(A) (B)

water
lipid bilayer
water

1 nm

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Membrane proteins

Integral and peripheral membrane proteins. Fluid mosaic model

Peripheral membrane protein
Carbohydrate
Integral membrane protein
Peripheral membrane protein

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Membrane functions

Separates and connects the two spaces.

- > Diffusion barrier – osmotic barrier
- > Selective transports
- > Types of transports:
 - passive transport - uniport
 - active transport - symport
 - antiport

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Passive transport

Driving force: concentration gradient (→ diffusion)
No energy demand.

It may be:

- Membrane diffusion
- Pore diffusion
- Carrier diffusion

Transported molecules
Uniport
Channel
Carrier molecule
Simple diffusion
Facilitated diffusion
Passive transport

Uniport: the molecular transport is independent from other transports

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Active transport

Against concentration gradient → energy is required
An active (energy-transforming) protein is necessary.

Symport: two molecules transport together, to the same direction.

Antiport: two molecules transport together, to opposite direction

Carrier molecule
Transported ion
Symport
Antiport
Energy
Coupled transport
Active transport

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Biological membranes in cells

Cytoplasmic/cell membrane
Nuclear membrane
Other membranes:

- Mitochondrion
- Endoplasmic reticulum
- Golgi complex
- Chloroplast
- Vesicles
- Special (retina, neuron)

lysosome
mitochondrion
peroxisome
Golgi apparatus
endoplasmic reticulum
vesicle
nuclear envelope

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Nuclear envelop

Nuclear pores for transporting mRNA out into cytoplasm

Outer membrane
Inner membrane
Nucleoplasm
Nucleolus
Chromatin
Nuclear envelope
Pore in nuclear envelope

19

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Endoplasmic reticulum and Golgi complex

Endoplasmic reticulum: flat, closed membrane sacks, covering the nucleus in few layers.
 RER: rough endoplasmic reticulum, it has small particles on the surface = ribosomes (→ protein synthesis)
Golgi apparatus: flat, closed membrane sacks surrounding ER in more layers.
 The synthesized proteins are let into ER lumen and during the maturation process they are moved through the layers of Golgi and transported to proper place. This transport is carried out in small transport vesicles covered with double lipid membrane, too.

20

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Nucleus
Nuclear pore
Rough endoplasmic reticulum
Ribosome
Smooth endoplasmic reticulum
Secretory vesicle
Cisterna
Trans face
Golgi apparatus
Protein expelled
Cell membrane
Proteins
Transport vesicle

21

http://www.fredonia.edu/bio241/images6_19_ER_and_Golgi.jpg

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MITOCHONDRIA – structure

Elongated particles, observable with microscope
 Number: ~10 – 1000 /cell
 They only occur in eukaryotes

Matrix
Cristae
Inner membrane
Outer membrane

22

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MITOCHONDRIA – biochemical functions

Located in the matrix space:

- The citrate cycle = Krebs cycle
- β-oxidation of fatty acids

Located in the inner membrane:

- Terminal oxidation

23

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Terminal oxidation

The substrate hydrogens arrive in the form of NADH or FADH. These are oxidized in three steps with oxygen. H⁺ ions accumulate in the intermembrane space. This Δc is converted to ATP.

1 NADH₂ → 3 ATP 1 FADH₂ → 2 ATP

Outer mitochondrial membrane
Intermembrane space
Inner mitochondrial membrane
Mitochondrial matrix

Electron transport (the respiratory chain) ATP production

24

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Protein biosynthesis

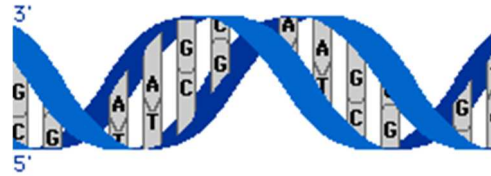
All proteins have a fixed sequence of amino acids. This must be exactly (re)produced in the biosynthesis.

The sequence is stored in the DNA encoded (genetic code, 64 different base triplets). This information is transcribed to mRNA in the nucleus.

The mRNA moves out of nucleus and the assembly of amino acids is going on the surface of ribosomes (translation).



Transcription - translation

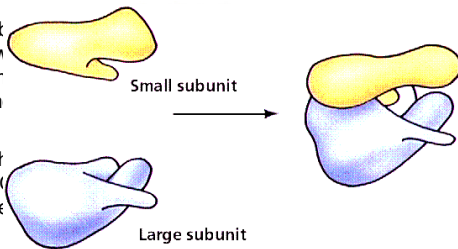


Ribosome

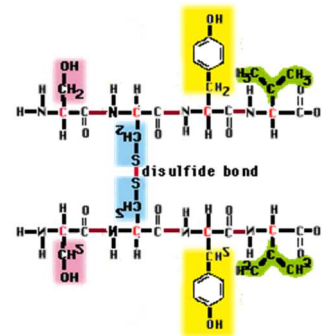
Ribosomes consist of two subunits, containing rRNA and protein. The two parts are coupled with a Mg^{2+} ion.

The size of subunit characterized by Swedberg sedimentation number (30 S and 50 S).

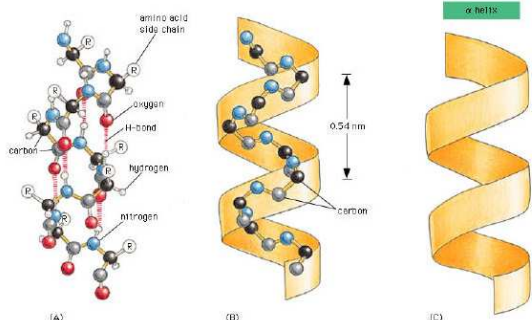
The ribosome has binding sites for mRNA, and three tRNA.



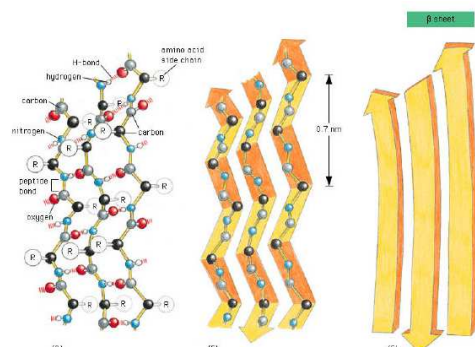
Primary structure: the amino acid sequence



SECONDARY STRUCTURE: α -helix



SECONDARY STRUCTURE: β -pleating



TERTIARY STRUCTURE

3D structure of the whole chain

12. előadás: a fehérjék szerkezete
Az előadás témakörébe tartoznak az A-, B- és C-területű előadások. A B-területű előadások az alapvető biológiai folyamatok (metabolizmus, sejtszintézis, sejtszervek) és a C-területű előadások a biotechnológiai alkalmazások (enzimek, mikroorganizmusok, sejtszervek) témakörébe tartoznak.

31

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QUATERNARY STRUCTURE

Quaternary structure: 3D structure of a protein complex consisting of more chain.
Example: hemoglobine, build up of two α and two β chain: $\alpha_2\beta_2$

32

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Levels of protein structure

33

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Cytoplasm

It is not a simple liquid, it has an inner structure, slightly elastic and deformable like *gels*.

(Gels: some macromolecules in solutions – like proteins or carbohydrates – form a crosslinked structure holding the liquid in form. This shows a quasi-solid properties – like jelly or jam.)

34

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The most important biochemical process in cytoplasm is:

GLYCOLYSIS

It is an energy producing process, it works both under aerobic and anaerobic conditions.

The energy balance of process:
-2 ATP + 4 ATP =
+2 ATP /molecule of glucose

35

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Cell wall

The microbial cell wall is a shield against mechanical stress and osmotic pressure. (Animal cells don't have cell wall, they don't need such protection.)

The two basic types of bacterial cell wall: Gram-positive, and Gram-negative.

The Gram-staining

is a staining method for microscopic prepares. Cells are stained with crystal violet and iodine, decolorized with alcohol and investigated under microscope. Cell walls colored violet-blue are identified as Gram-positive, Gram-negative cells remain pink.

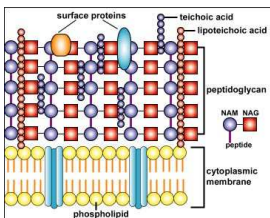
36

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Differences of cell wall structure

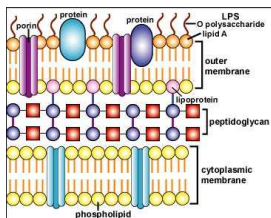
Gram positive

Cell membrane + a thick peptidoglycan layer



Gram negative

a thin peptidoglycan layer between two lipid membranes



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