



LOSCHMIDT  
LABORATORIES

## 2. Introduction to Molecular Biotechnology

Bi7430 Molecular Biotechnology

### Outline

- ❑ Definition of biotechnology
- ❑ History of biotechnology
- ❑ Fundamentals of molecular biotechnology
- ❑ Basic concept of rDNA technology
- ❑ Methods of gene transfer
- ❑ Main fields of biotech applications
- ❑ Risks and positives

### Definition of biotechnology

- ❑ **biotechnology** („biotech“)  
*bios – techné – logos*
- ❑ **Kalr Ereky, 1917** – „biotechnology is a process by which raw materials could be biologically upgraded into socially useful products“
- ❑ „any technological application that uses biological systems, living organisms, or derivatives thereof, to make or modify products or processes for specific use“  
(The United Nations Convention on Biological Diversity, 1992)

### History of biotechnology

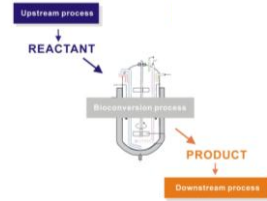
- ❑ a story that began long time ago
- ❑ 10,000 B.C. neolithic revolution  
**cultivation and domestications**
- ❑ 8,000 B.C. **fermented bread**  
(ancient Egypt)
- ❑ 8,000 B.C. **cheese making**  
(the Middle East)
- ❑ 6,000 B.C. **wine production**  
(Egypt and the Middle East)
- ❑ 5,000 B.C. **brewing**  
(ancient Egypt)
- ❑ developed without any knowledge about  
existence of cells, enzymes, genes



## History of biotechnology

## Traditional biotechnology

- 1970s biotechnology recognized as **scientific discipline** (interlink of chemical engineering, microbiology nad biochemistry)
- traditional biotechnology **based on fermentation**
- development focused on **process technology** (bioreactor design, upstream, downstream)



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- biotransformation component**
  - natural strains - far from optimum
  - difficult to optimise
  - induced mutagenesis and selection (chemical mutagens, UV radiation)
  - limited by inherited properties of the strain



## Revolution in biotechnology

- 1973 Stanley Cohen and Herbert Boyer development of **recombinant DNA technology**



genetic engineering provided the means to create, rather than merely isolate, highly productive strains

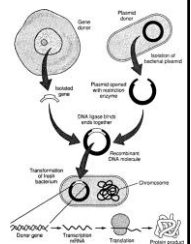
*Proc. Natl. Acad. Sci. USA*  
Vol. 70, No. 11, pp. 3240-3244, November 1973

### Construction of Biologically Functional Bacterial Plasmids *In Vitro*

(*Orbital restriction enzymes/transformation/inducible tetracycline resistance*)

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## Molecular biotechnology

- 1976 Herbert Boyer and Robert Swanson



- 1978 production of **human insulin in *E. coli*** by Genentec (recombinant "human" insulin approved by FDA 1982)
- 1981 production of **recombinant growth hormone**
- 1987 production of **recombinant tissue plasminogen activator** used to dissolve blood clots during myocardial infarction
- 1980-83 about 200 **small biotechnological companies** founded in US

## Molecular biotechnology

- 1974 Rudolf Jaenisch - **first transgenic mammal** (a mouse)

### ANIMALS AND PLANTS ACT AS NATURAL BIOREACTORS

- 1982 first recombinant animal vaccine approved
- 1983 engineered Ti plasmid – **plant transformation**
- 1988 Kary Mullis - **PCR method (Nobel Prize in 1993)**
- 1994 **first genetically engineered food** approved by FDA (tomato)



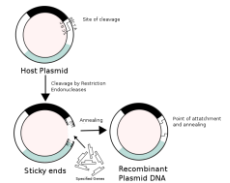
## Molecular biotechnology

- 1995 **first genome** sequenced (bacterium *Haemophilus influenzae*)
- 1996 complete **eukaryotic DNA sequence**
- 1996 commercial planting of **GMO crops begins**
- 1997 Ian Wilmut – **nuclear cloning of a mammal**
- 1998 first **antisense drug** approved by FDA
- 1999 **Drosophila genome** sequenced
- 2000 **Arabidopsis genome** sequenced
- 2000 development of „**golden rice**“
- 2001 **human genome** sequenced
- 2009 **first drug** produced in genetically engineered animal (a goat)



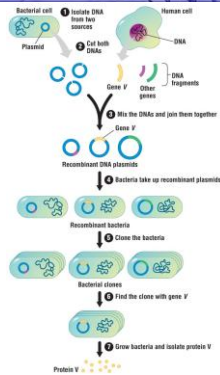
## Molecular biotechnology

- classical biotechnology** based on selective breeding
- molecular biotechnology** (modern; „mol biotech“) revolutionary scientific discipline based on methods of gene manipulation (*Lecture 3*)
- the ability to transfer specific units of genetic information from one organism to another
- recombinant DNA technology (rDNA)**
- genetic engineering** enable create rather than isolate highly productive organisms



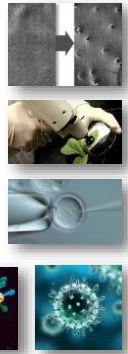
## Concept of rDNA technology

- ❑ isolate gene(s) of interest
- ❑ modify gene(s)
- ✓ protein engineering (Lecture 4)
- ❑ ligate gene(s) into a vector
- ❑ transform host organism
- ❑ select transformed cells
- ❑ culture host organism
- ❑ application of gene product



## Techniques of DNA transfer

- ❑ transformation and transfection
- ❑ direct methods
  - electroporation (2.5 kV, 5 ms)
  - chemical transformation ( $\text{CaCl}_2$ )
  - heat shock ( $42^\circ\text{C}$ )
  - micro-injection
  - biolistic delivery - „gene gun“
  - liposomal transfection
- ❑ indirect methods
  - transduction (bacteriophage)
  - viral and bacterial vectors



## Mol. biotech applications

- ❑ **white** - industrial biotechnology (Lecture 8)
  - production of fine chemicals
  - production of proteins/enzymes
- ❑ **green** - agricultural biotechnology (Lecture 9)
  - transgenic plants and animals
  - biofertilizers and biopesticides
- ❑ **red** - medical biotechnology (Lecture 10-11)
  - developing new vaccines and drugs
  - tissue engineering and regenerative therapies
  - molecular diagnostics and pharmacogenomics
  - cell and gene therapy
- ❑ **grey** - environmental biotechnology (Lecture 12)
  - biosensing and bioremediation



## Pros and cons

- ❑ **safety and ethical concerns of molecular biotechnology**
  - do we have a right to move genes, creating new life forms, „playing God“?
  - will transgenic organisms be harmful to other organisms or environment?
  - should humans be genetically engineered?
- ❑ **positive aspects of molecular biotechnology**
  - opportunities to accurately diagnose, prevent and cure a wide range of infectious and genetic diseases
  - increase crop yield and resistance to insects and diseases, environmental stress (e.g., drought, heat, cold)
  - develop microorganisms that produce chemicals in a sustainable manner
  - facilitate removal of pollutants and waste materials from environment