

# “Synthetic Biology: ENGINEERING LIFE”

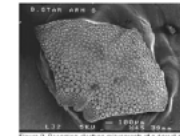
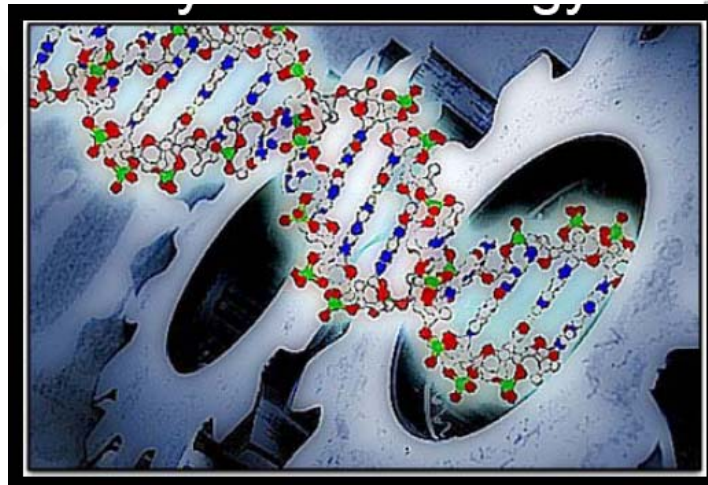
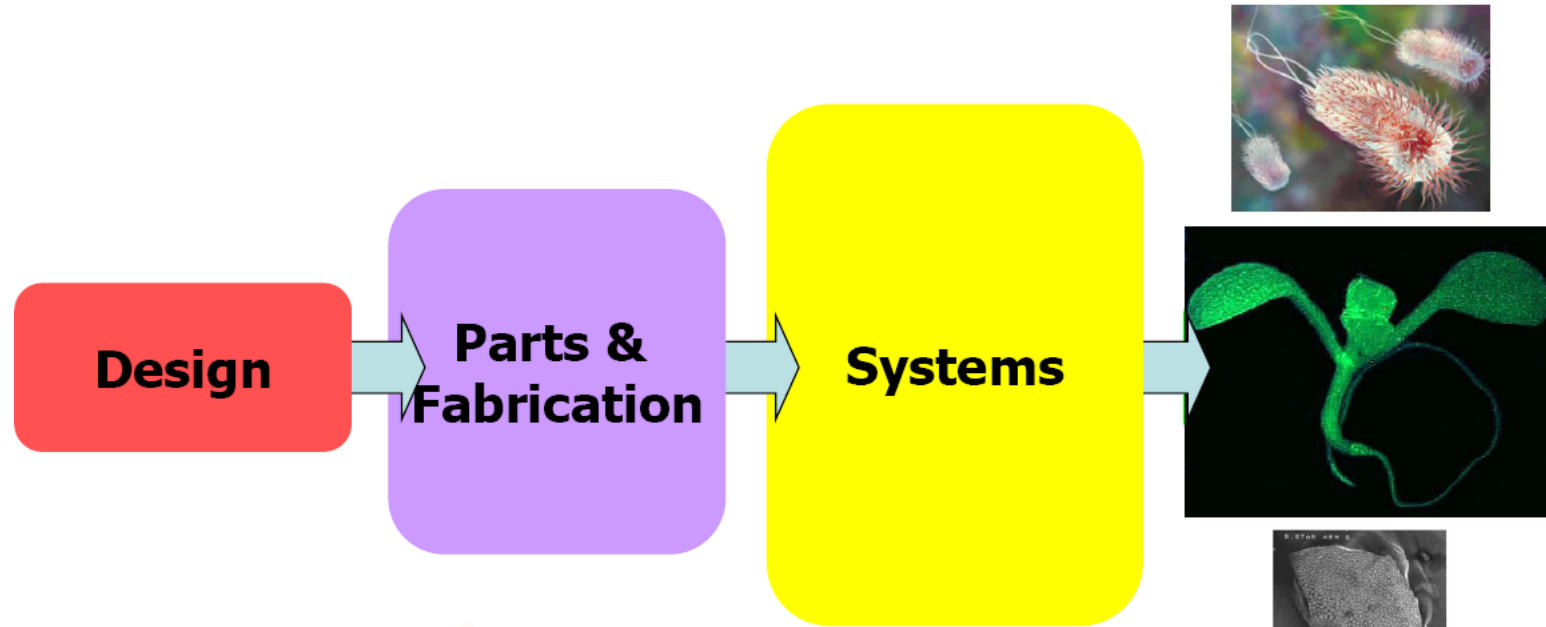


Figure 2. Scanning electron micrograph of a starfish arm. The arm is a highly porous, brittle structure. The structure is made up of single crystal cables. It is decorated with microchannels that concentrate light on the myoepithelial bundles. Courtesy of L. Arbib.



- There are many definitions of synthetic biology and each have varying ethical, legal, social and economic implications
- One general definition of synthetic biology is; the design and construction of new biological parts, devices, and systems and the re-design of existing, natural biological systems for useful purposes

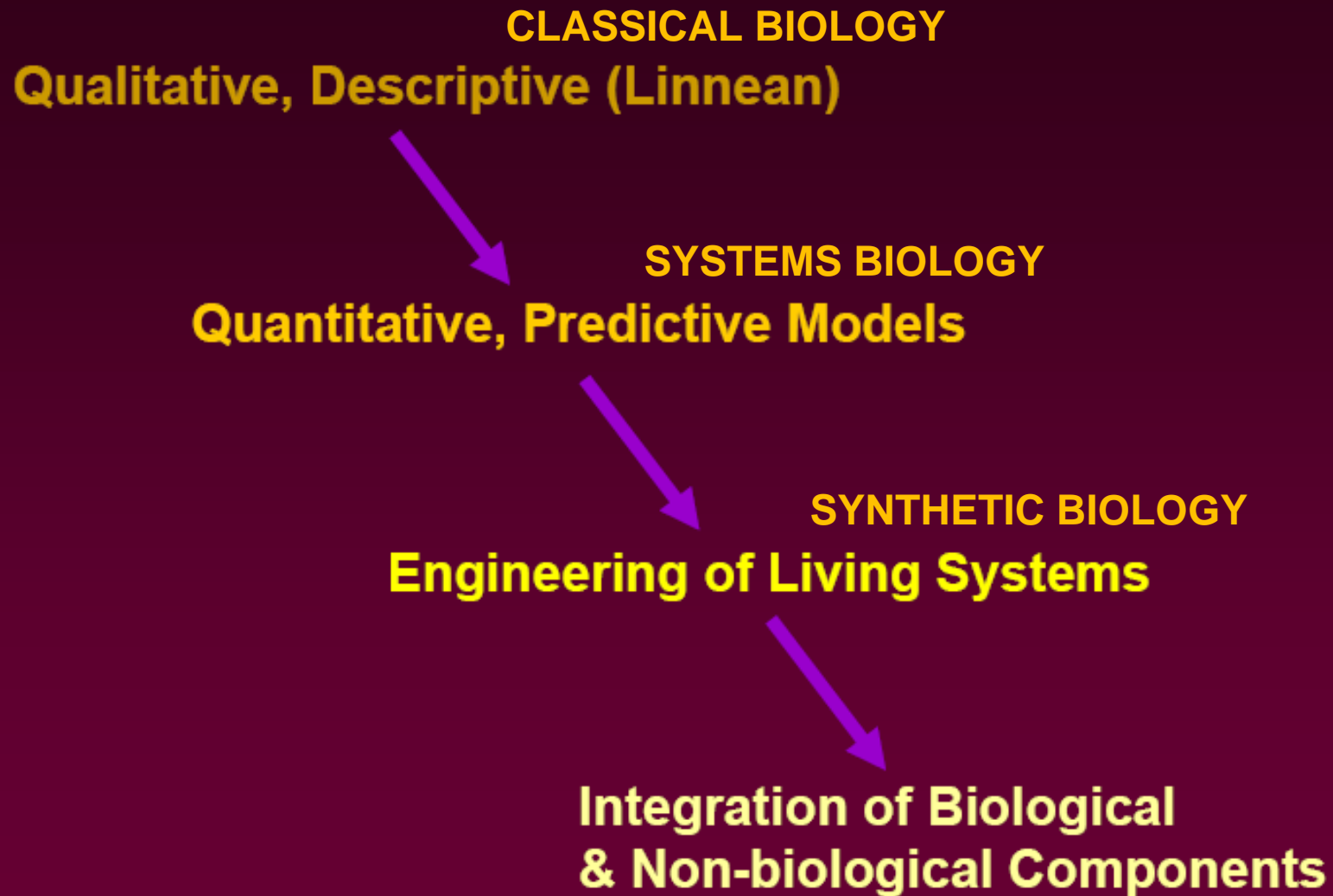
- Synthetic biology contains elements of nanotechnology and biotechnology
- Synthetic biology can be a technology based on the traditional genetic code
- Or it can be based on artificial genetic code, artificial amino acids and proteins.
- It can also be the creation of biological system with non-biological material or making of non-biological system with biological material.

# **“Synthetic Biology: ENGINEERING LIFE”**

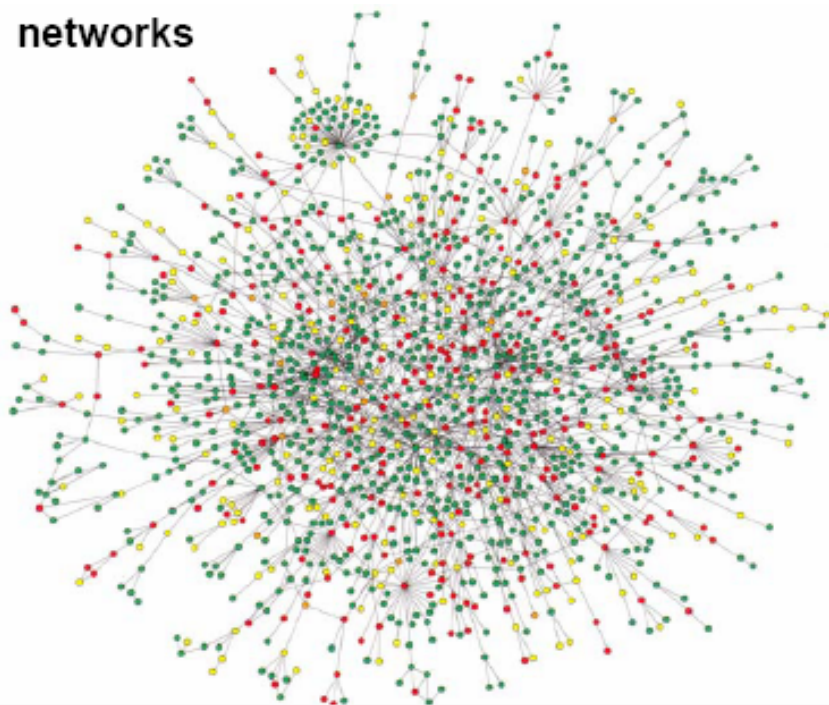
Mainly developed at the USA , with some minor disperse efforts In Europe



# Bioscience: From Discovery to Creation



## networks



## pathways & circuits



## complexes



## molecules



## contributing disciplines

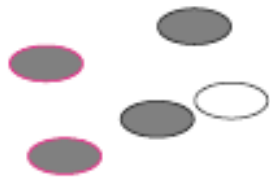
**Maths and Computing**  
Simulation-based systems  
analysis  
Modelling of complex states

**BIOSYSTEMS**

**Physical Sciences**  
Photonics/Imaging  
Spectroscopy  
Reaction  
mechanisms/kinetics  
Structural methods

**Engineering**  
Nanoscale  
technology/metrology  
Electronics  
Process control theory  
Systems modelling

# Biology poses unique challenges for engineering



Cell growth/death



Crosstalk (no insulation)

CAT → ACT  
Mutation



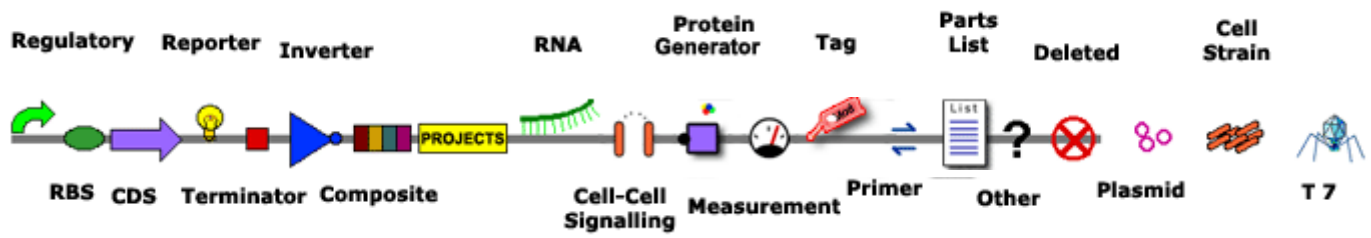
Evolution



Chemical diffusion

We need new engineering rules to control or utilize these properties.

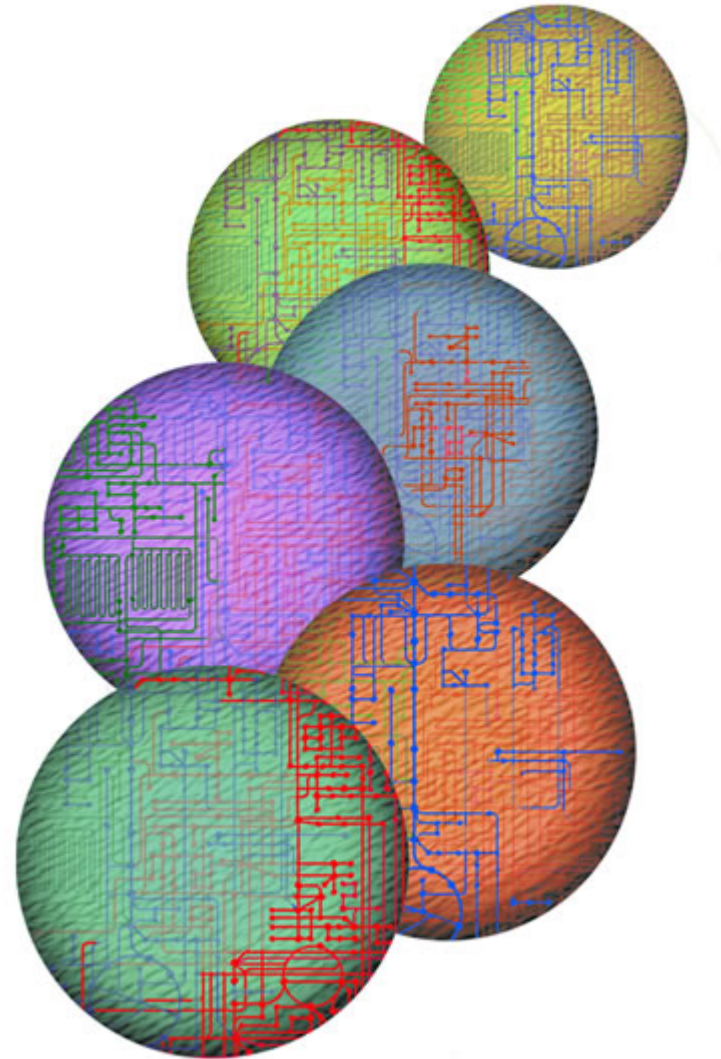
# SYNTHETIC BIOLOGY: ENGINEERING STANDARDIZATION



**Why now Synthetic Biology?**

# Why now?

- Advances in computing power
- Genomic sequencing
- Crystal structures of proteins
- High through-put technologies
- Biological databases
- Diverse biological sampling/collection



How do advances in DNA synthesis affect synthetic biology?

Our capacity to assemble long stretches of DNA that approach the length of genomes has increased.

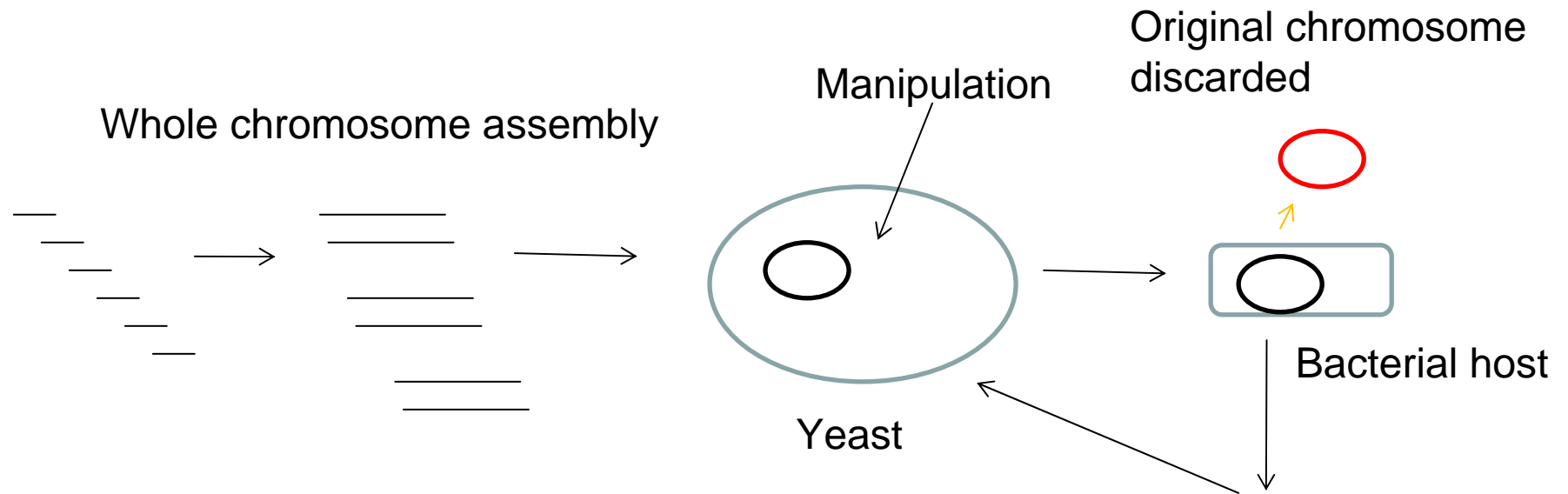
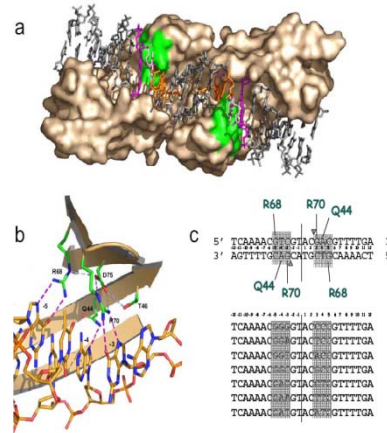
DNA synthesis is an enabling tool for construction of a genome of known sequence, in the absence of a **physical** template.

The genomes of known viruses have been synthesized.

Improved DNA synthesis does not address the essential issue of how to synthesize an organism with predefined, useful altered properties.

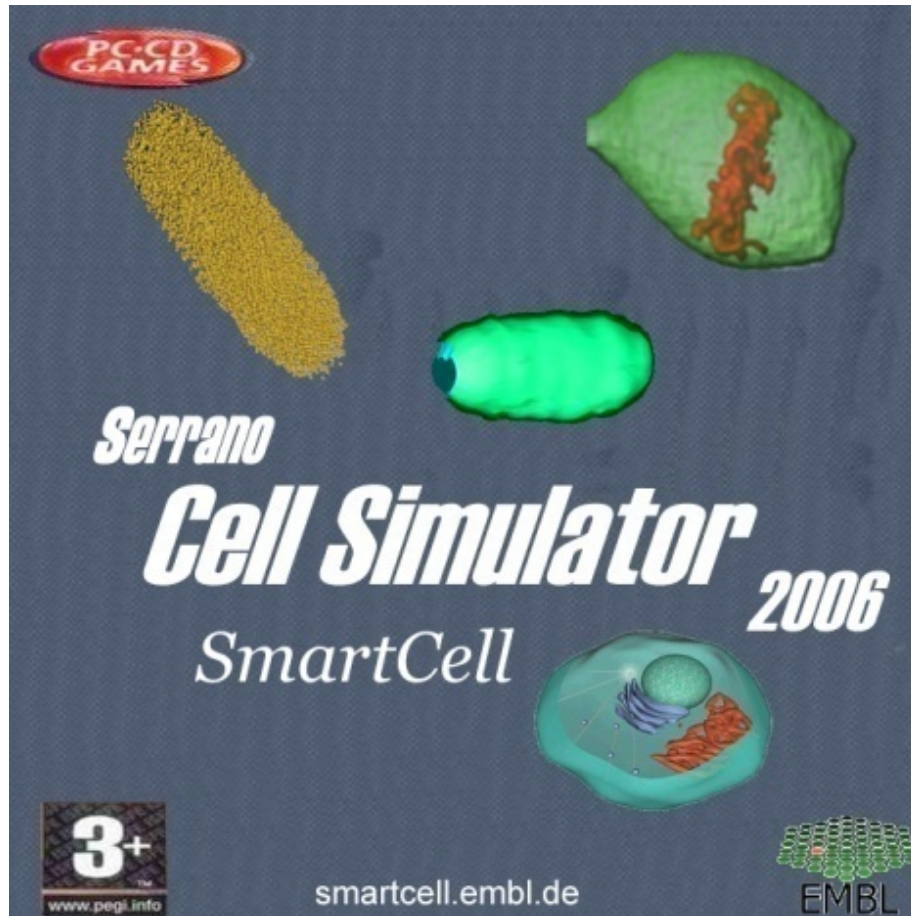
# NEW TOOLS FOR GENOME ENGINEERING

Precision scalpels  
for genome engineering

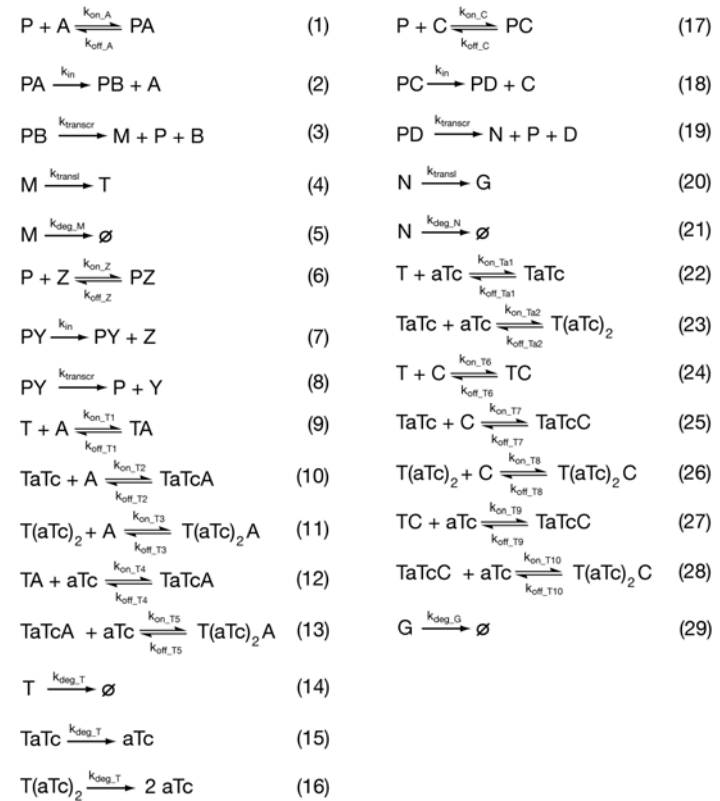




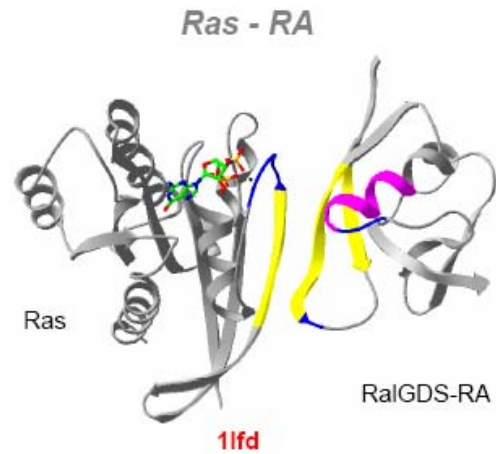
www.smartcell.crg.es



## COMPUTER TOOLS TO SIMULATE BIOLOGICAL SYSTEMS

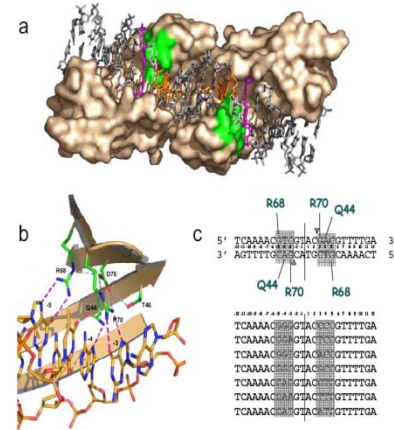


## Successful examples for structure-based predictions and protein design



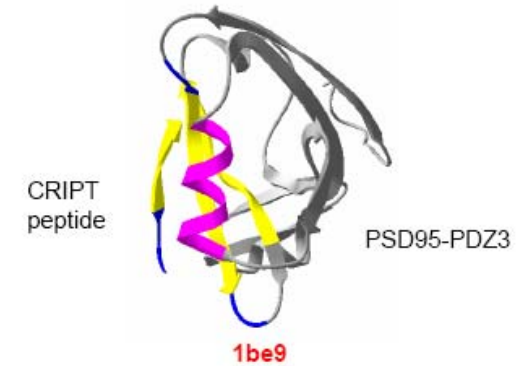
Kiel et al., J Mol Biol, **2005**  
 Kiel et al., J Mol Biol, revised, **2007**

### *Meganuclease-DNA*



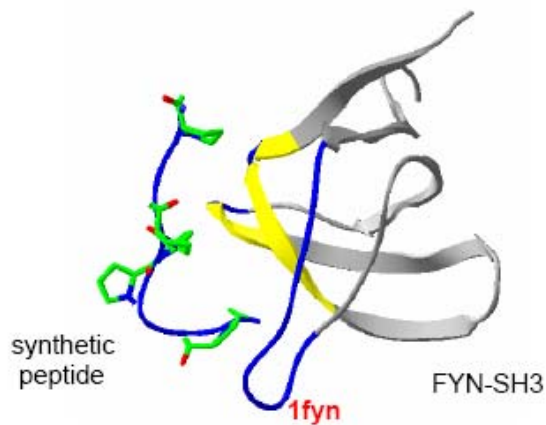
Arnould et al., J. Mol. Biol, **2006**

### *Peptide - PDZ*

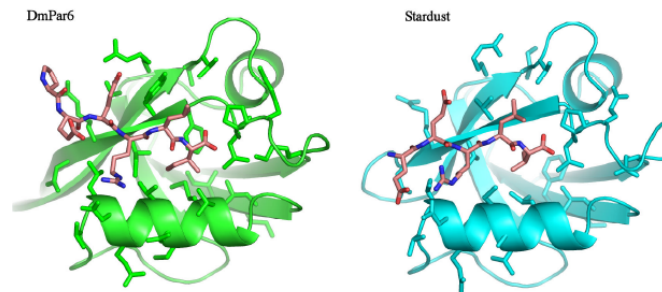


Kolsch et al., Science, **2007**

### *Peptide - SH3*

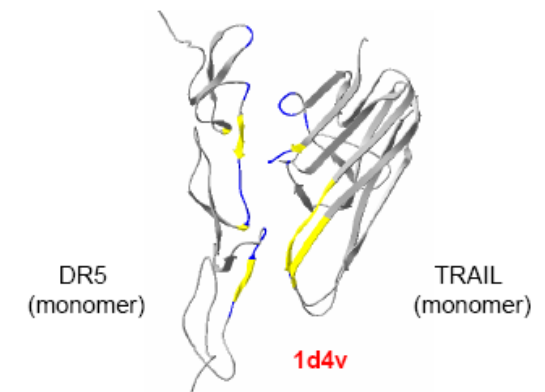


Musi et al., Protein Sci, **2006**



Kempkens, O., E. Medina, et al. (2006)

### *TNFR - TNF*



Van der Sloot, et al, PNAS, **2006**

## Defining the Rule Sets for Biological Design, Assembly and Function

- common genetic (digital) code in all life forms
- genomes encode a limited series of structural building blocks (protein motifs and programmed assembly)
- combinatorial assembly of protein building blocks generates extravagant structural and functional diversity

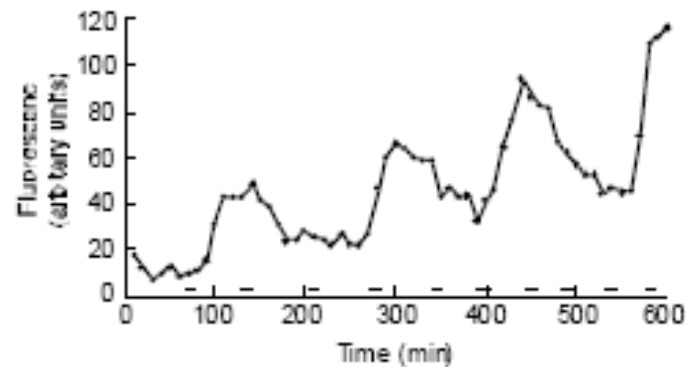
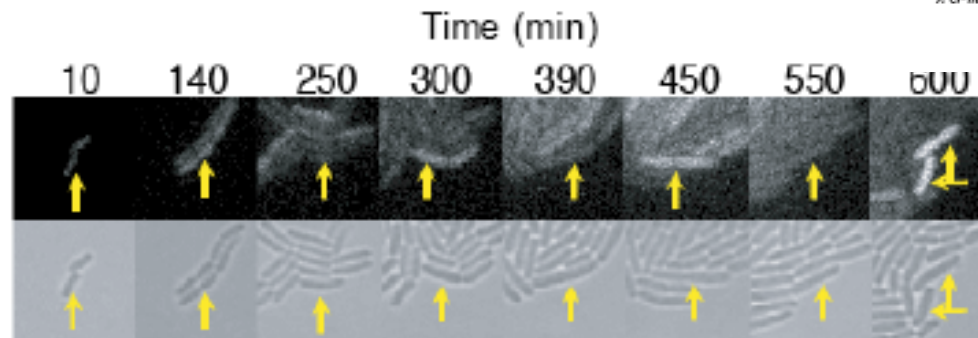
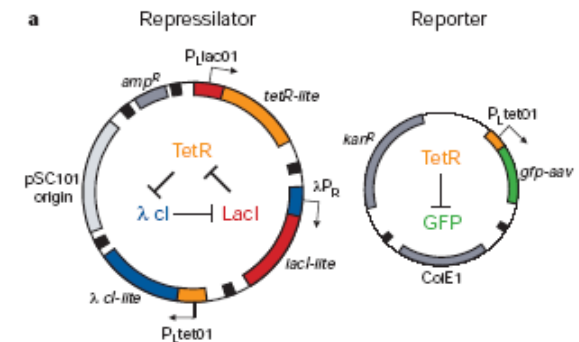


# **Playing with Synthetic Biology**

# Engineering of Circuits

What sorts of modules have been made by synthetic biologists?

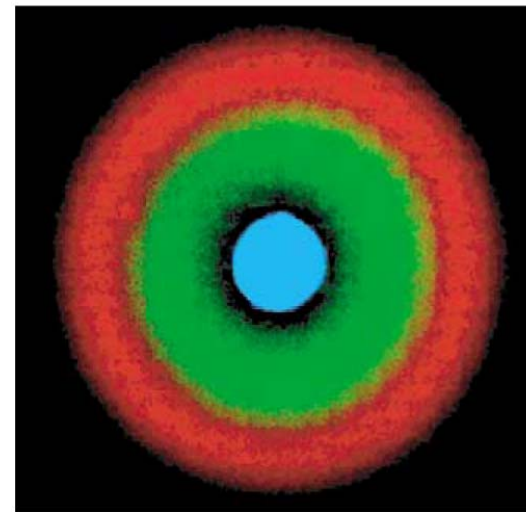
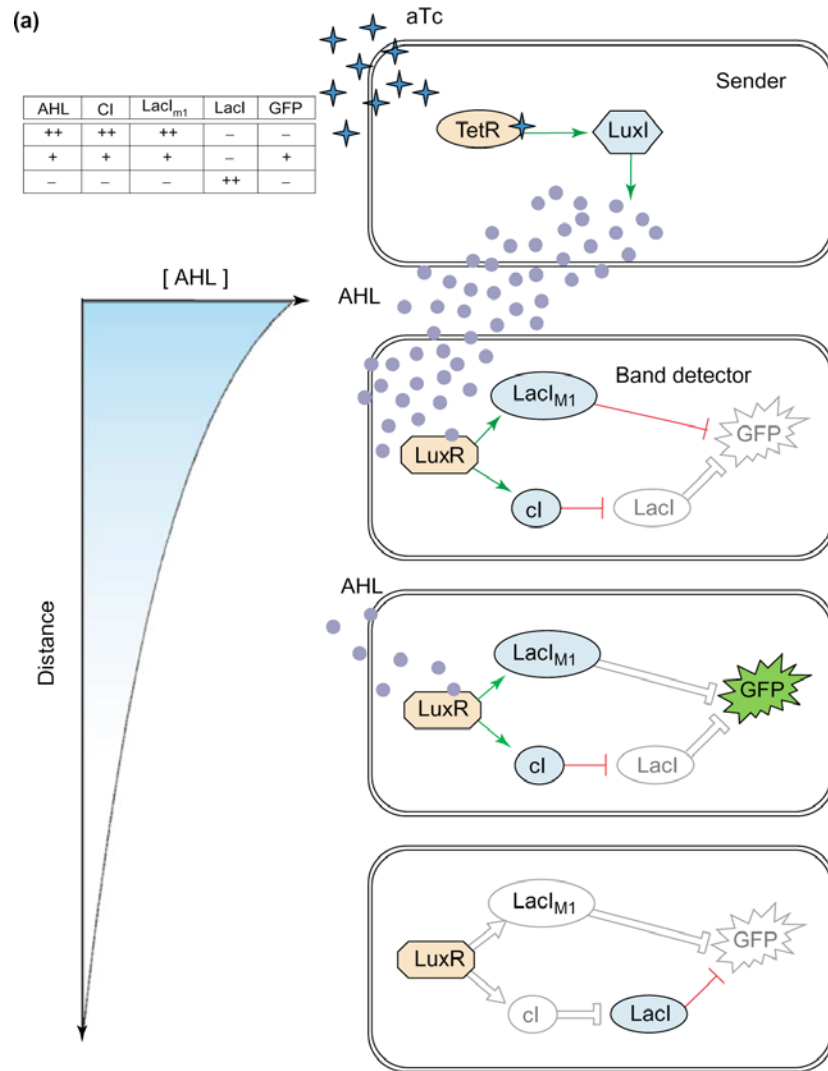
## Oscillators





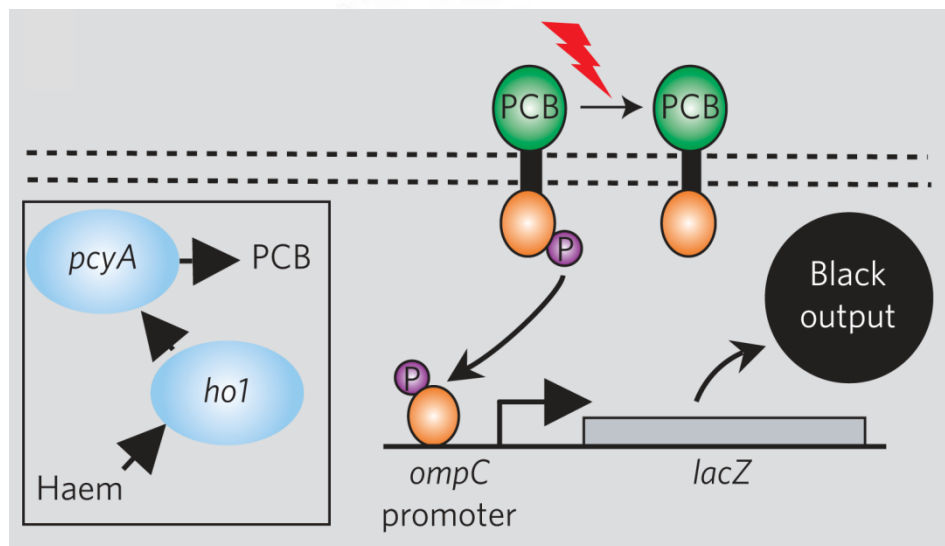
# Engineering of Circuits

## Building a sender & receiver



# Engineering of Circuits

## Bacterial photography



*Synechocystis*: Photoreceptor, Phytochrome synthesis genes (*ho1*, *pcyA*)  
*E. coli*: osmotic shock response circuit (*ompC*),  $\beta$ -galactosidase

Levskaya et al (2005) *Nature*,

**We need more Parts**



**Biological Design: "Endless Forms Most Beautiful":  
Limitless Diversity From Combinatorial Assemblies of Limited Building Blocks**

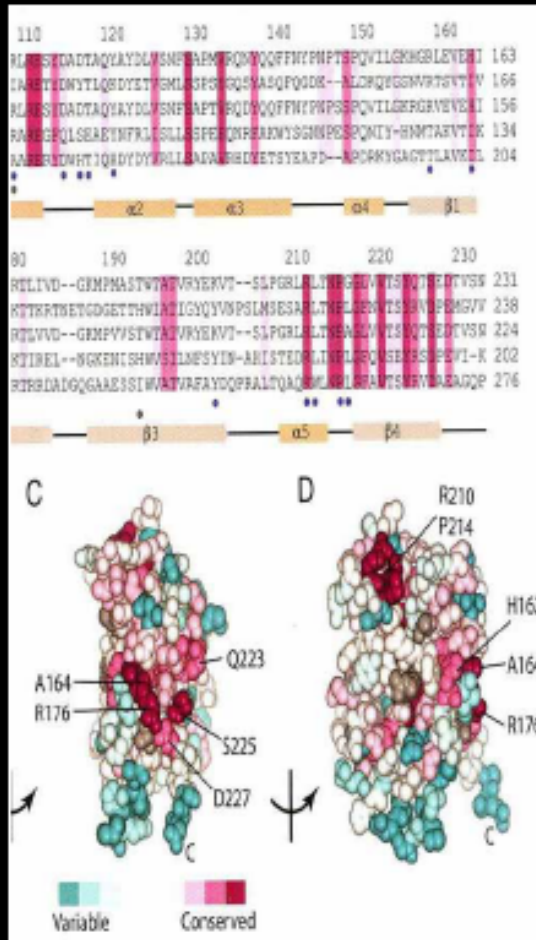


## **Metagenomics: Sampling the Extravagant Functional Diversity of Microorganisms**

- **estimated 100 billion microbial species**
- **only 6000 species cultivated and characterized**
- **massive repertoire of uncharacterized genes/proteins/metabolomes**
- **metagenomic sampling**
  - **mass screening of complete genomes of unknown/unculturable organisms**
  - **high throughput profiling to identify transfer of gene(s) with desired function into 'universal acceptor' organisms**



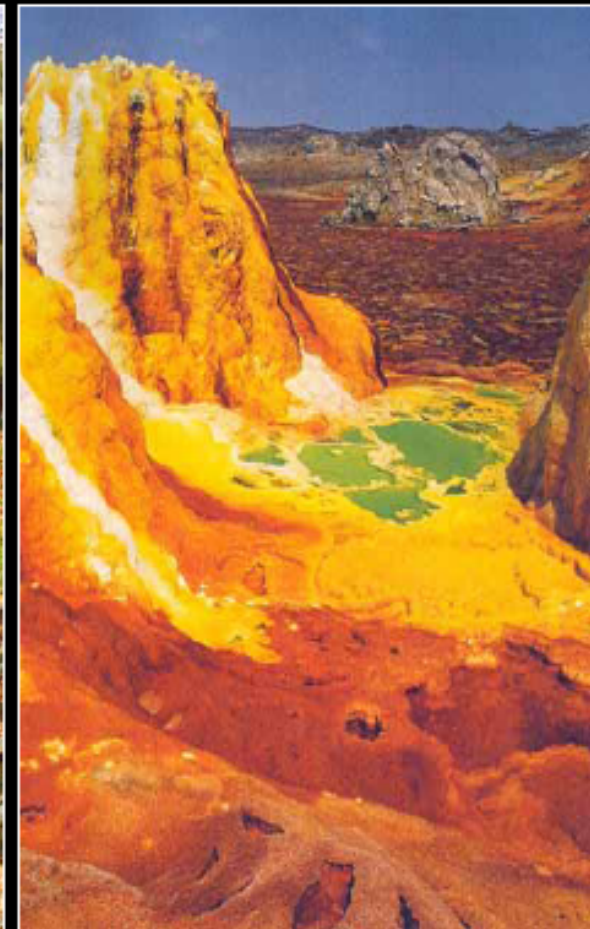
# Ecogenomics: Mapping the Extraordinary Genomic Diversity and Biosynthetic Capabilities of Microbial Life



genomics

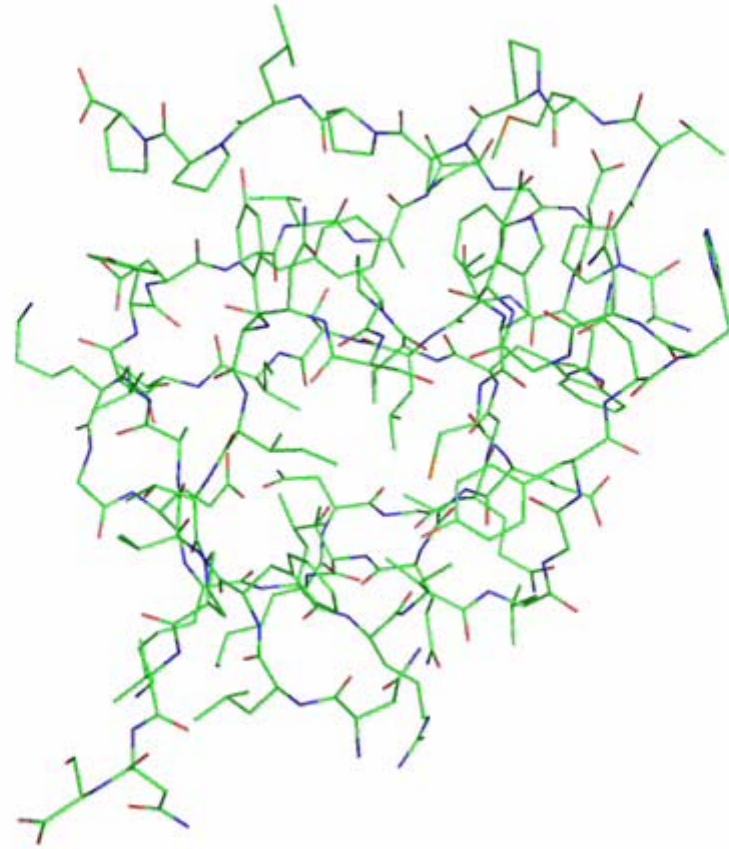


eco-niches



extremophiles

# PROTEIN DESIGN



# Exploring Biospace: Iterative Selection of Novel Variants for Substantial Functional Performance Improvements Versus Naturally Occurring "Parent System"



Activity Versus Starting Material

	Reduced	Same	Increased
cycle 1	● ●	●	●
cycle 2	●	●	●
cycle 3	●	●	●
cycle 4	●	●	●

# **Possible applications of Synthetic Biology**



# Synthetic Biology: An Emerging Technology with Diverse Industrial Applications

Healthcare



Public Health



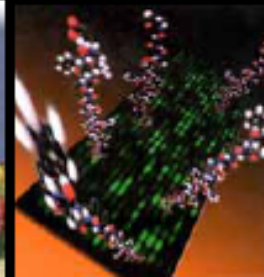
Agriculture



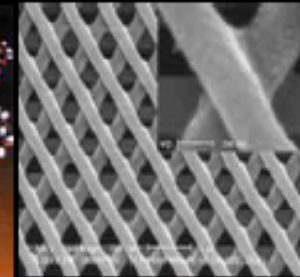
Functional Foods



Novel Materials



Textiles



Bioenergy  
and  
Biofuels



Industrial  
Enzymes



'Green'  
Mfg



Bio-  
remediation



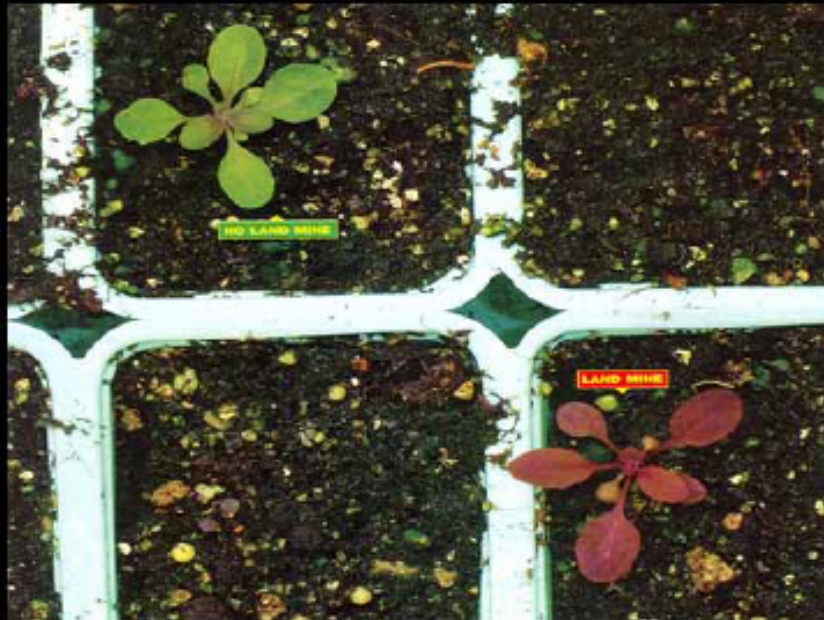
Clean  
Water



Ubiquitous  
Sensors

# Biological Sensor Systems for Environmental Monitoring and Ecosystem Status

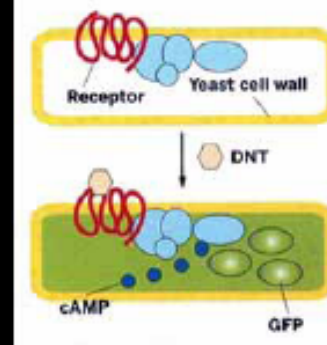
Genetically-Engineered *Arabidopsis thaliana*  
Change-Color when Exposed to Landmine  
Degradation Products



Aresa Biotection, Demark

## Engineering Sentinel Organisms as Environmental Sensors

**SNIFFER** DNT induces yeast cells with transplanted rat olfactory machinery to produce cAMP, which then triggers production of green fluorescent protein (GFP).



D. N. Dhanasekaran et al (2007) Nature Chem. Biol.

DOI: 10.1038/nchembio882

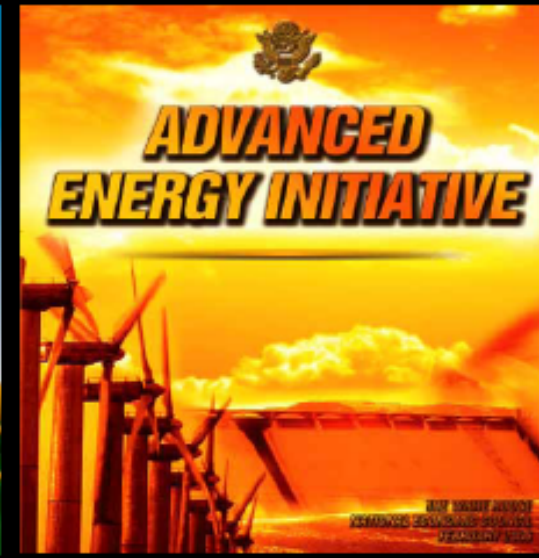
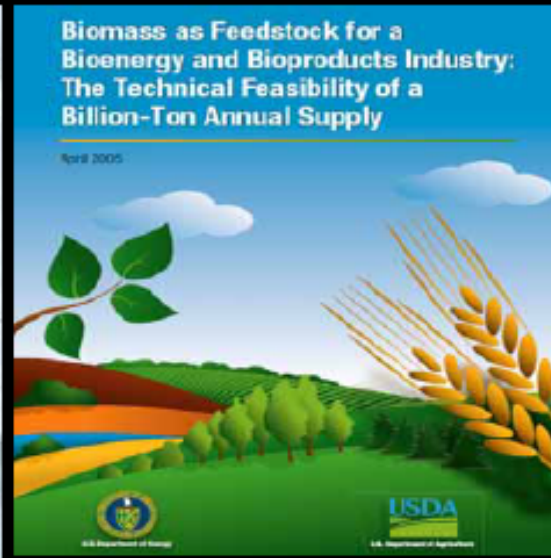


# Directed Evolution and Design of Novel Enzymes for Catalysis of Lignocellulosic Biomass



- cellulosic energy crops
- agricultural, forestry and mill residues
- food processing residues
- municipal solid waste
- non-recycled paper
- construction and demolition wood

# Bio-Inspired Systems for Energy Production



## Production of Biodiesel from Biomass: A Widely Differing Yield Spectrum\*

Source	Yield	
	US gal/acre	L/Km <sup>2</sup>
● Soybean	40-50	35-45,000
● Rapeseed	110-145	100-130,000
● Mustard	140	130,000
● <i>Jatropha</i>	175	160,000
● Palm oil	650	580,000
● Photosynthetic Microbes	10-20,000	9-18,000,000

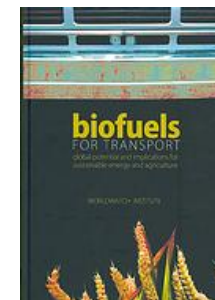
\* <http://en.wikipedia.org/wiki/Biodiesel>



# Biofuels



- First generation
  - Food feedstock: sugar, starch, vegetable oil or animal fats using conventional technology (food for fuel debate)
  - Fuel types: vegetable oil, biodiesel, butanol, ethanol, syngas
- Second generation **(Needs Synthetic Biology)**
  - Non food crop feedstock: cellulose, waste biomass: wheat, corn, wood
  - Fuel types: biohydrogen, biomethanol, DMF, bio-DME, Fischer-Tropsch diesel, biohydrogen diesel, mixed alcohols and wood diesel
- Third generation **(Needs Synthetic Biology)**
  - Algae feedstock
- Fourth generation **(Needs Synthetic Biology)**
  - CO<sub>2</sub> feedstock: CO<sub>2</sub> converted to methane by bacteria



# Building a Super H<sub>2</sub> Producer

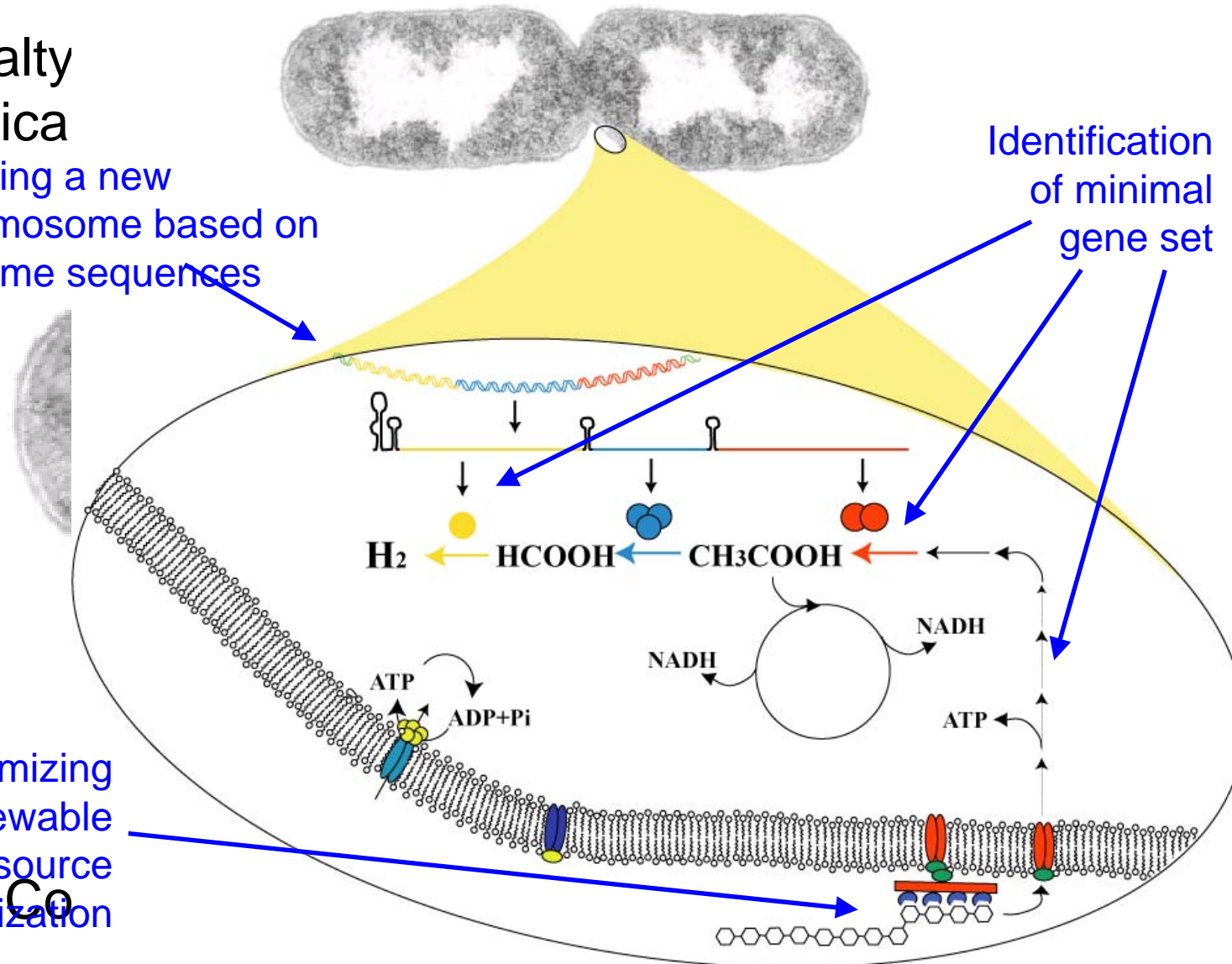
Specialty  
Chemical

Building a new  
chromosome based on  
genome sequences

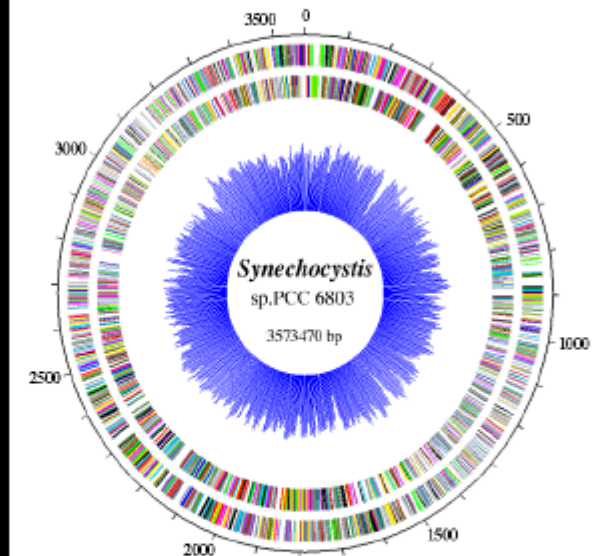
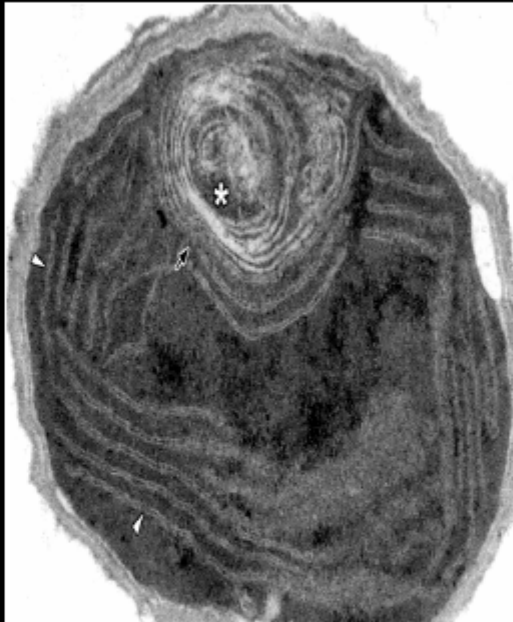
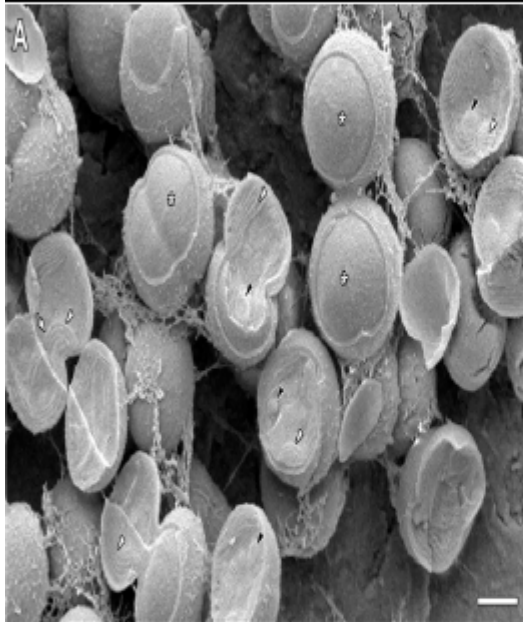
Identification  
of minimal  
gene set

Maximizing  
renewable  
resource  
utilization

Co



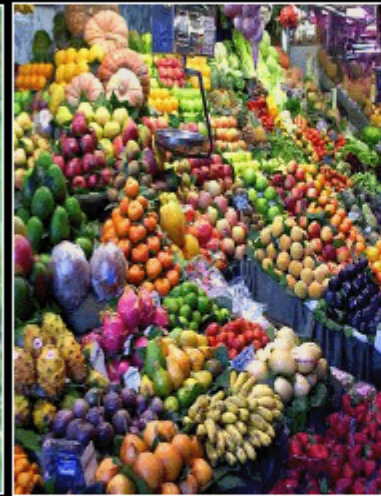
## Biomass to Energy: The Merits of Cyanobacteria as a Feedstock



- well characterized, tractable genome for trait enhancement
- no cellulose
- compositional homogeneity
- rapid assay of genetic modifications
- siting flexibility
- no or limited transport costs
- CO<sub>2</sub> fixation



## Synthetic Biology and Engineering Enhanced Traits in Food, Feed and Fiber Products



# Synthetic Biology and Novel Industrial Process Chemistry

**The Natural World  
(Evolutionary Biospace)**

**The Synthetic World  
(Exploring Unexplored  
Biospace)**

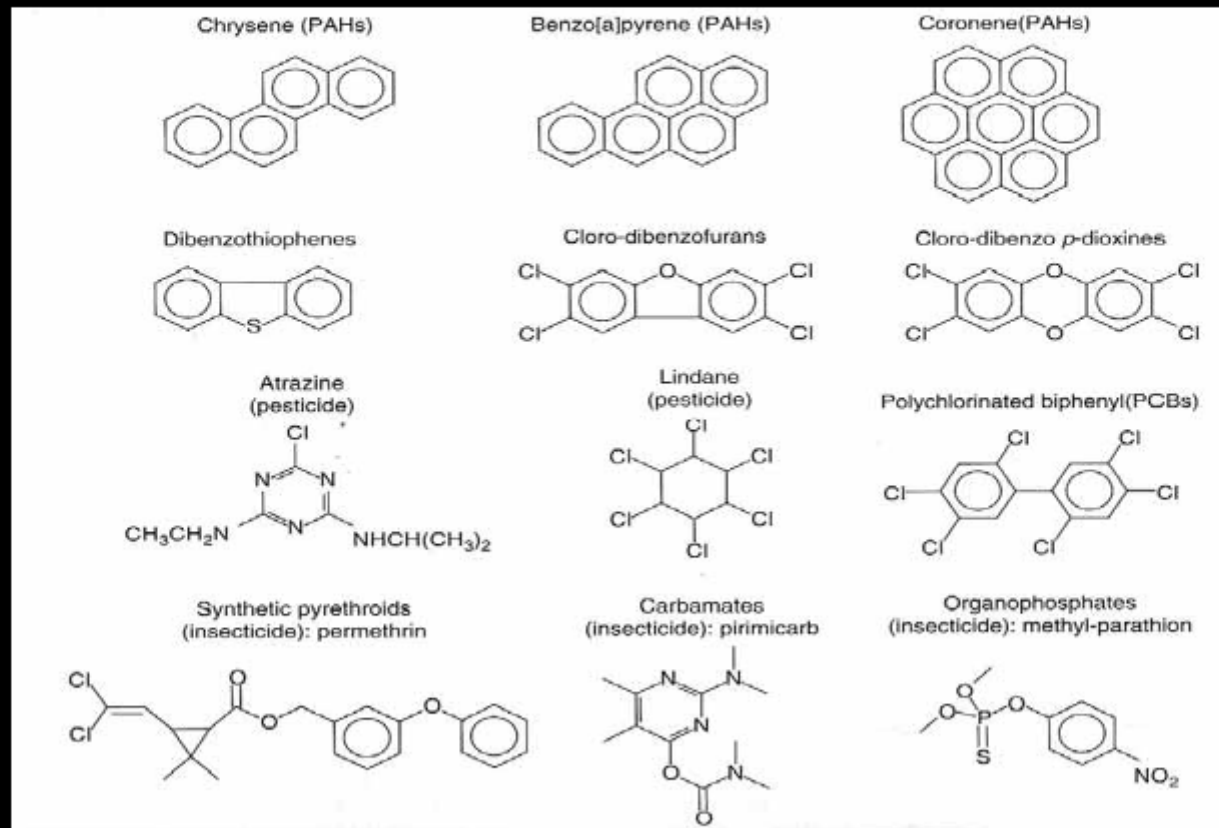
**mapping  
protein  
families  
and  
functional  
design**

**metagenomics  
and  
expanded  
knowledge  
of  
evolved  
'biospace'**

**directed  
(accelerated)  
evolution  
and  
expansion of  
novel  
'biospace'**



# Microbial Genomics and Synthetic Biology: New Technology Platforms for Bioremediation and Improved Efficiency of Wastestream Management



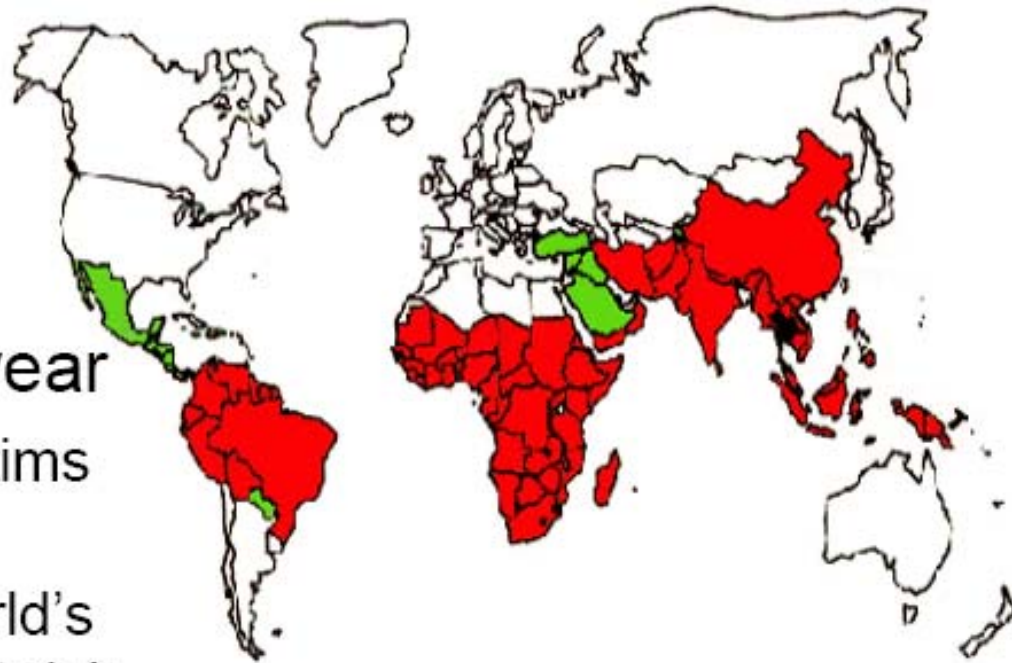
## **Bio-Inspired Engineering: Removal of Perchlorate Contamination by Chemical Reduction by Biofilms of Hydrogen Producing Bacteria**

- ✓ pilot scale project at La Puente, CA (400 L/min)
- ✓ influent: 25 mg/L of  $\text{NO}_3^-$  and 60  $\mu\text{g/L}$   $\text{ClO}_4^-$
- ✓  $\approx 95\%$  removal of  $\text{ClO}_4^-$ , from 60  $\mu\text{g/L}$  to below the CA action level of 4  $\mu\text{g/L}$
- ✓  $\approx 98\%$   $\text{NO}_3^-$  removal of 25 mgN/L to  $\approx 0.5\text{mgNO}_3^-/\text{L}$
- ✓ essentially 100%  $\text{H}_2$  usage



# Malaria

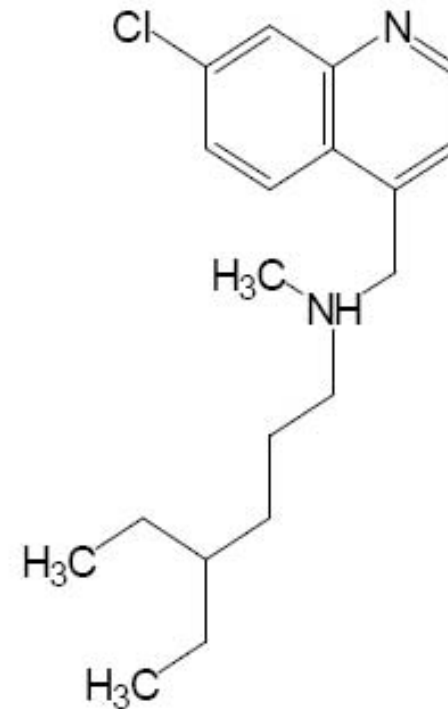
- 1.5-2.7 million people die of malaria every year
  - 90% of the victims are children
  - 40% of the world's population is at risk



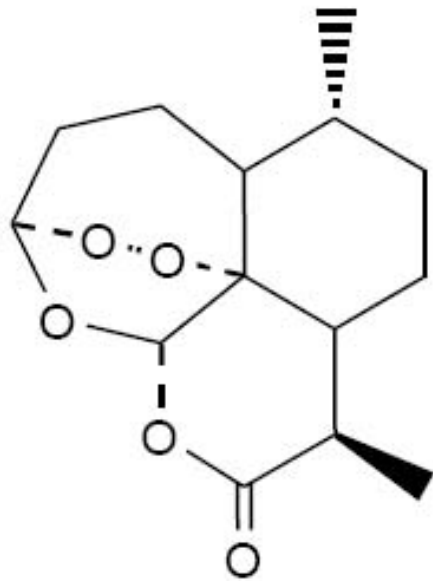
- Economists have proposed that malaria decreases the GDP of affected countries by as much as 50%.

# Chloroquine-based drugs

- Most widely-used drugs to treat malaria
- *Plasmodium* in South America and Southeast Asia is largely resistant to chloroquine



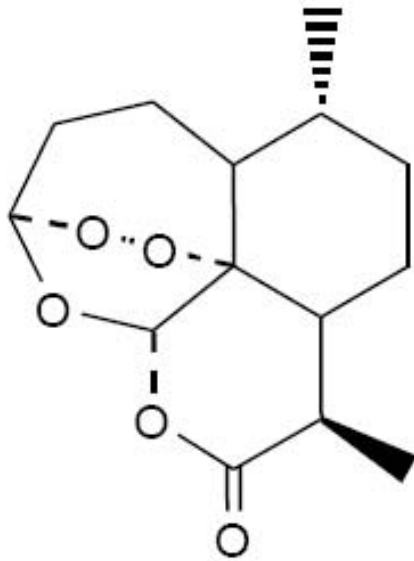
# Artemisinin-based drugs



- The current cost is approximately \$2.40.
  - Artemisinin adds \$1.00-1.50 to the cost for drugs
  - Most developing countries spend less than \$4/person/year on health care
- As many as 10-12 treatments are needed for each person annually
- World Health Organization estimates that 700 tons will be needed annually

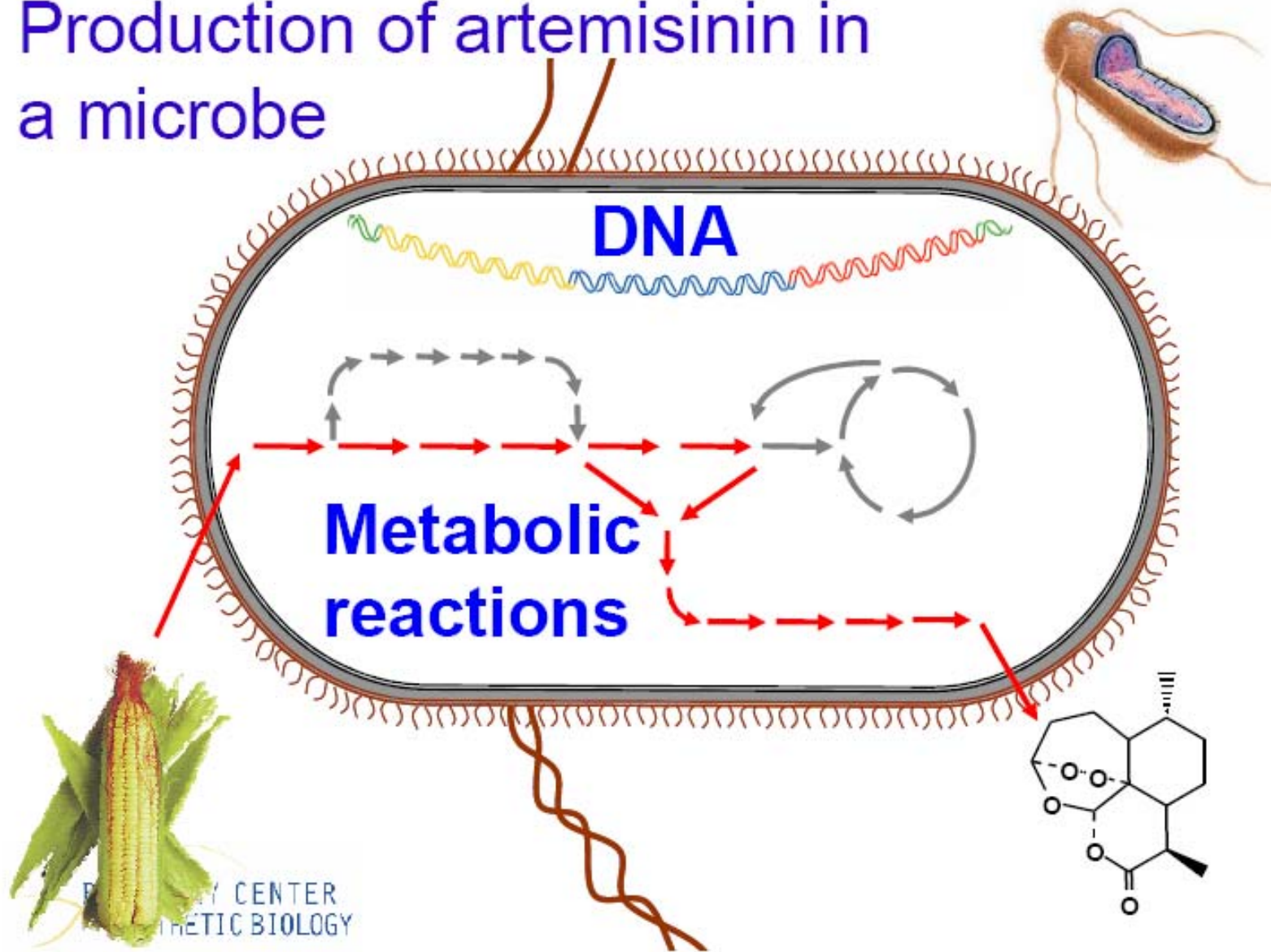


# Potential sources for artemisinin

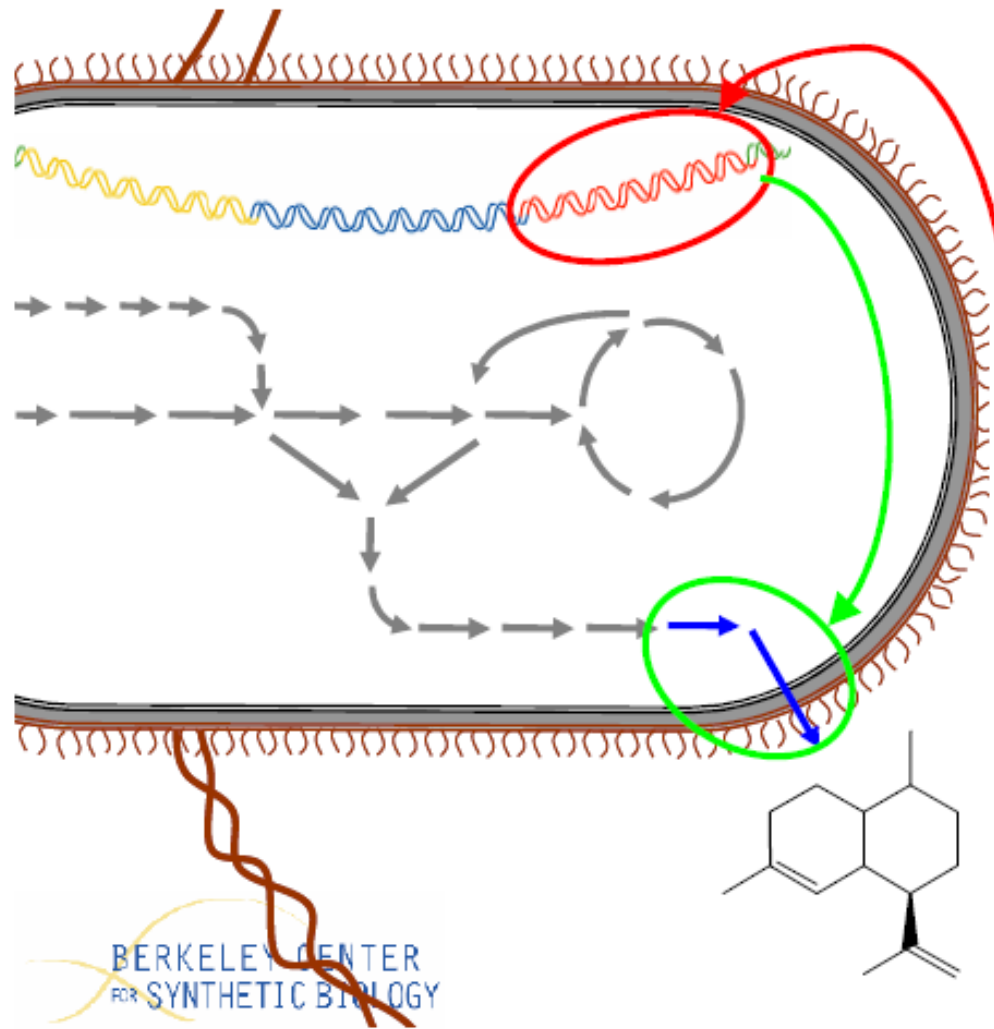


- **Agriculture**
  - Efforts are under way to plant *Artemisia annua* around the world
- **Chemical synthesis**
  - A synthesis route is known but it is too complicated for economical production
- **Microbial**
  - Need all of the genes from the plant

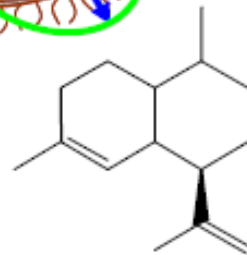
# Production of artemisinin in a microbe



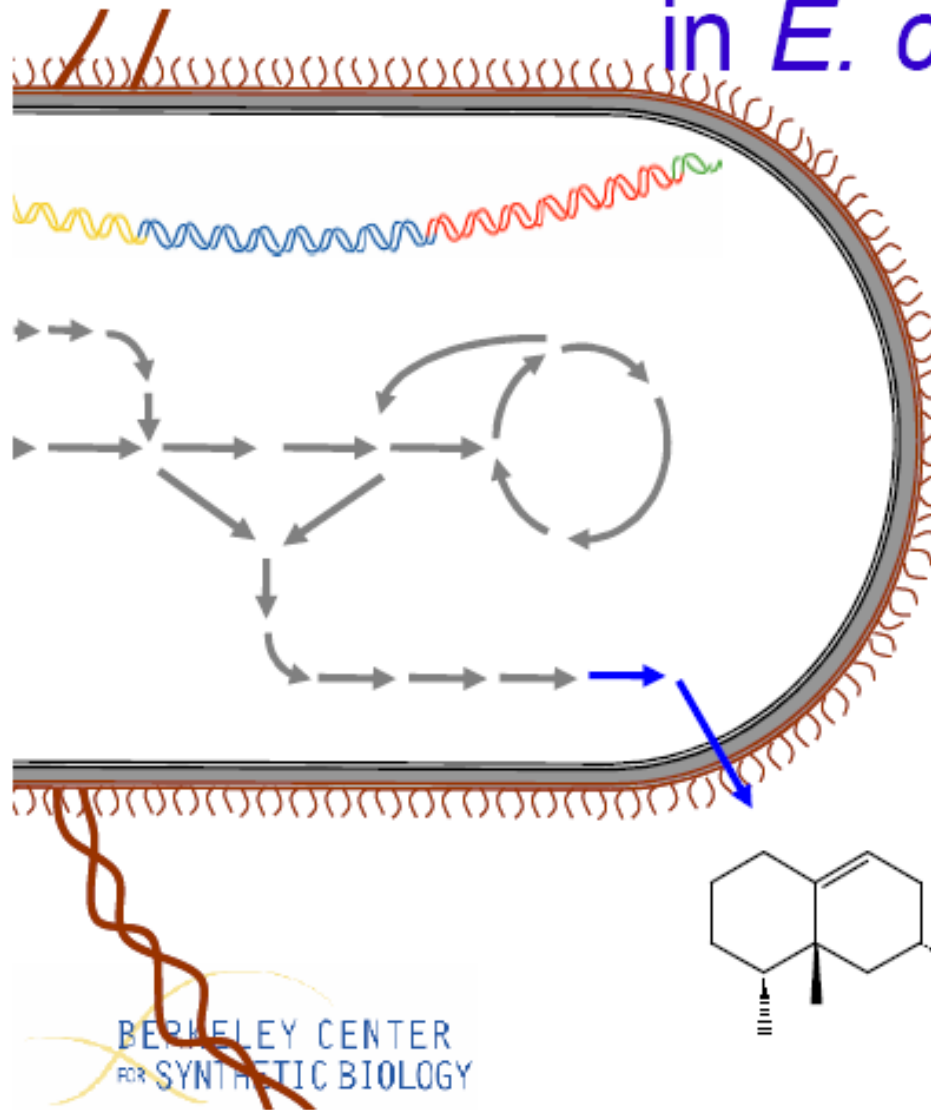
# Transplanting artemisinin pathways



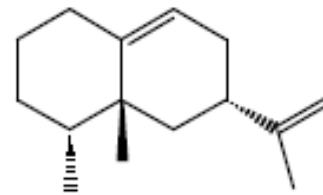
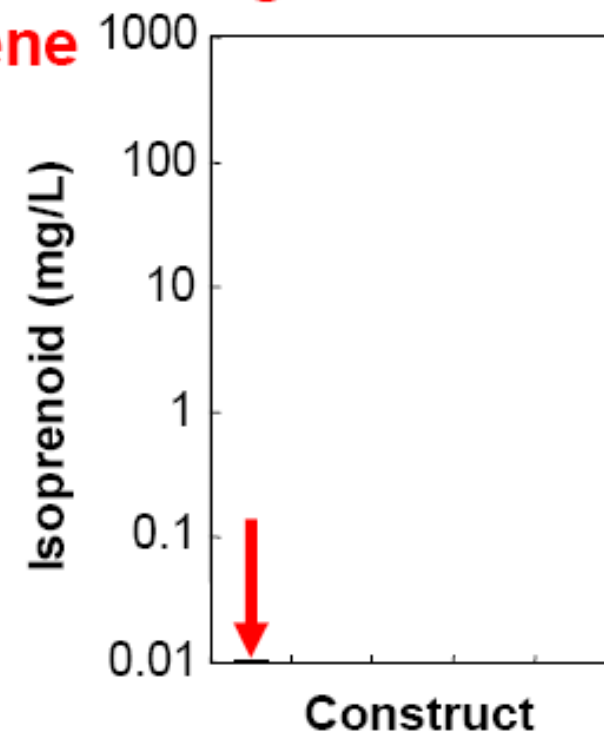
Clone the genes



# Production of a model isoprenoid in *E. coli*



**Very low production  
of isoprenoid resulted  
when using the native  
gene**





atcttgat catccaaga caaaaccaga gaaaagacc tgtctgttt ttaagaagt  
ctttatatta tttttttgt cggagaatct tataagcatg gcttcaggag gatcaaagtc  
ggcagcttc atgcttctga tgctgaatct tggctctat ttcgtcatca ccatcatcgc  
ttcttgggct gttaatcacg gcatcgagag aactcgcgag tctggtaact aacaaagata  
acaactgatt aagtaacaat taatccaacg ttagaaaatg tcatcatcaa tcttctttt  
gtggtatfff gcagcgtcga cactgtcact tccggcgaag atattcccga tataactccc  
ggtggggaac atggcgaccg gtttttcgt aatattcacg ttaatcgccg gcgtcgtcgg



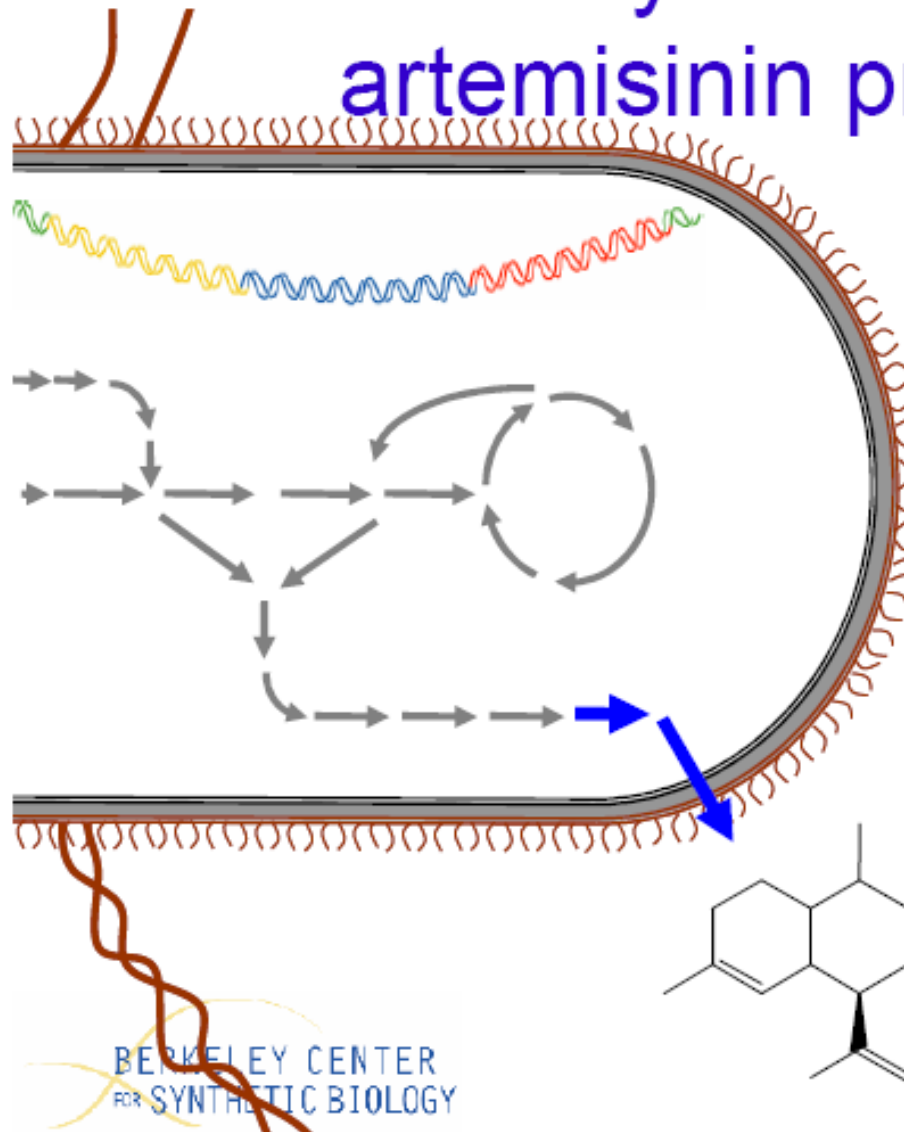
## Making a plant gene look like a microbial gene



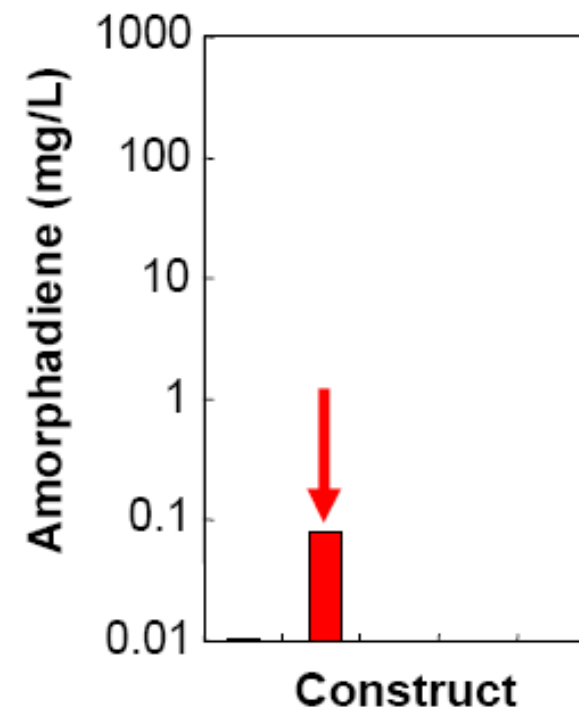
atcttgat catccaaga caaaaccaga gaaaagacc tgtctgttt ttaagaagt  
ctttatatta tttttttgt cggagaatct tataagcatg gcttcaggag gatcaaagtc  
ggcagcttc atgcttctga tgctgaatct tggctctat ttcgtcatca ccatcatcgc  
ttcttgggct gttaatcacg gcatcgagag aactcgcgag tctggtaact aacaaagata  
acaactgatt aagtaacaat taatccaacg ttagaaaatg tcatcatcaa tcttctttt  
gtggtatfff gcagcgtcga cactgtcact tccggcgaag atattcccga tataactccc  
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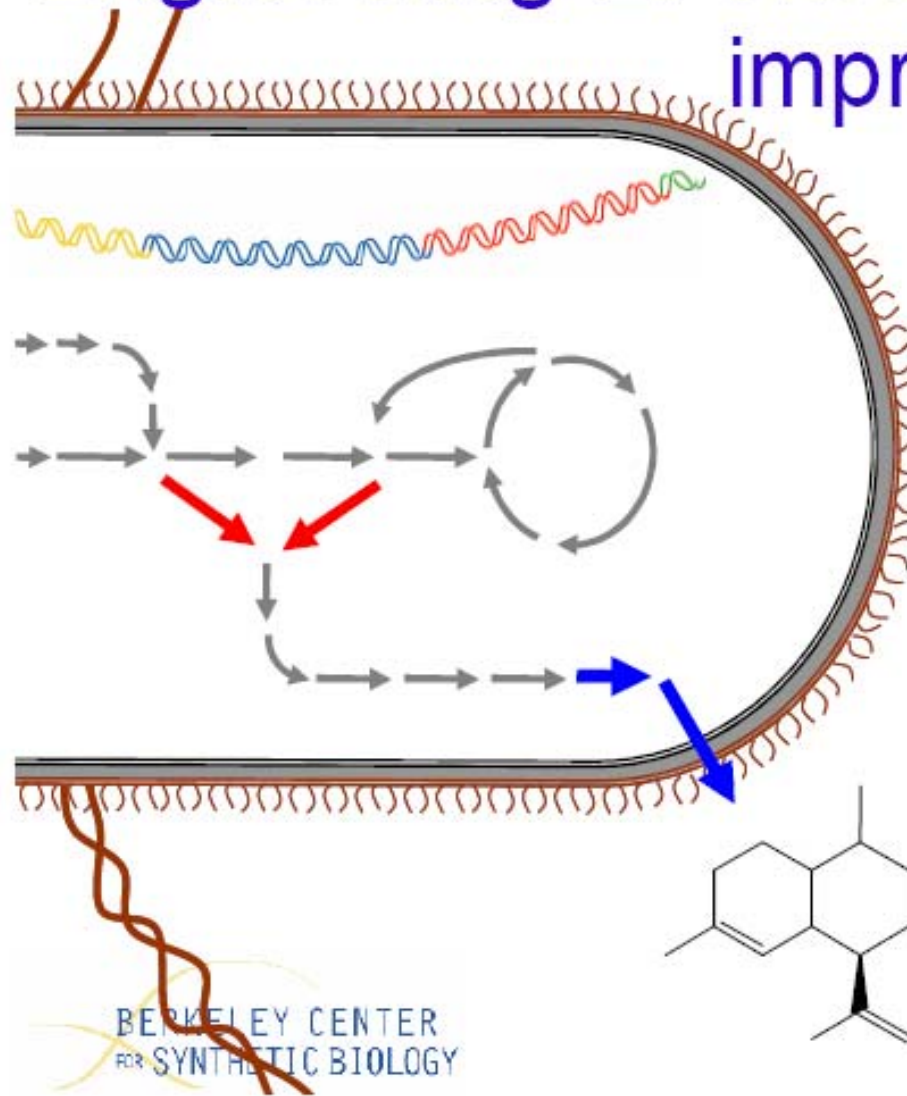
# Gene resynthesis improves artemisinin production



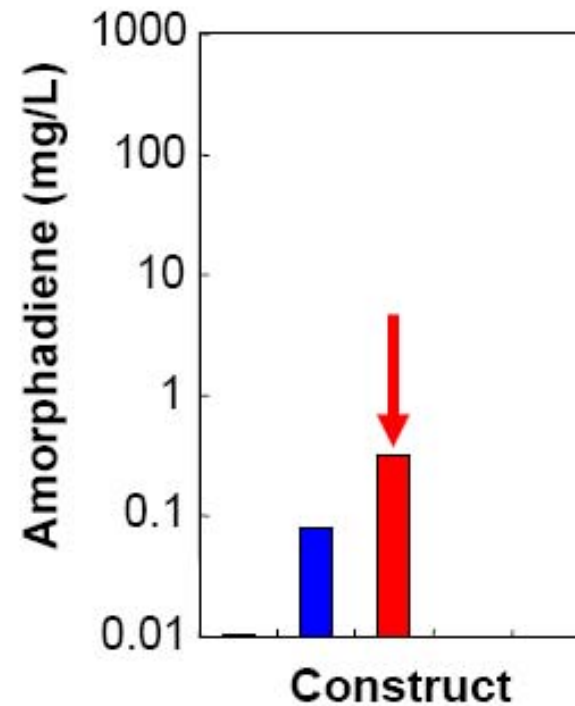
**142-fold improved production!**



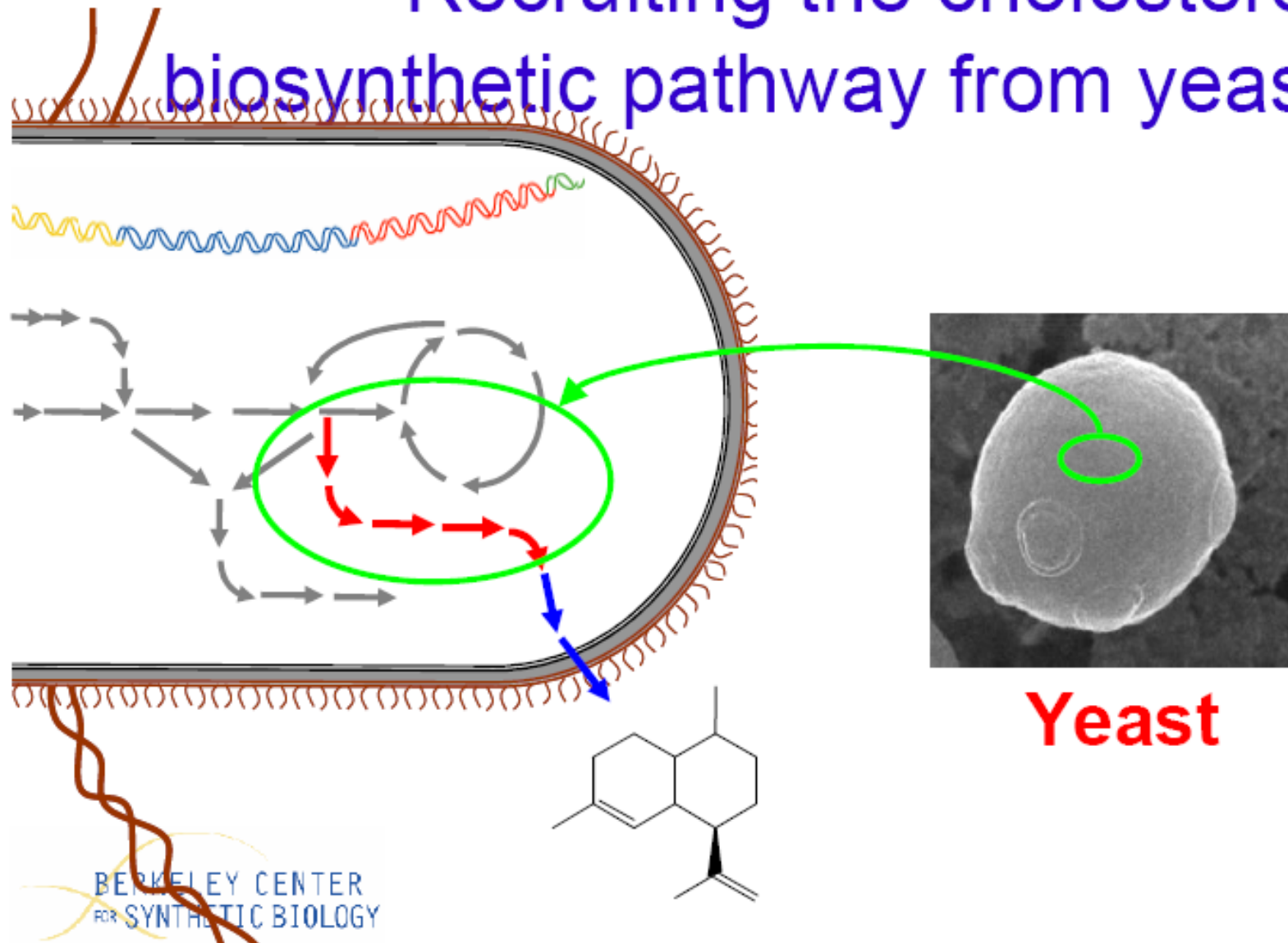
# Engineering *E. coli*'s native pathway improves production



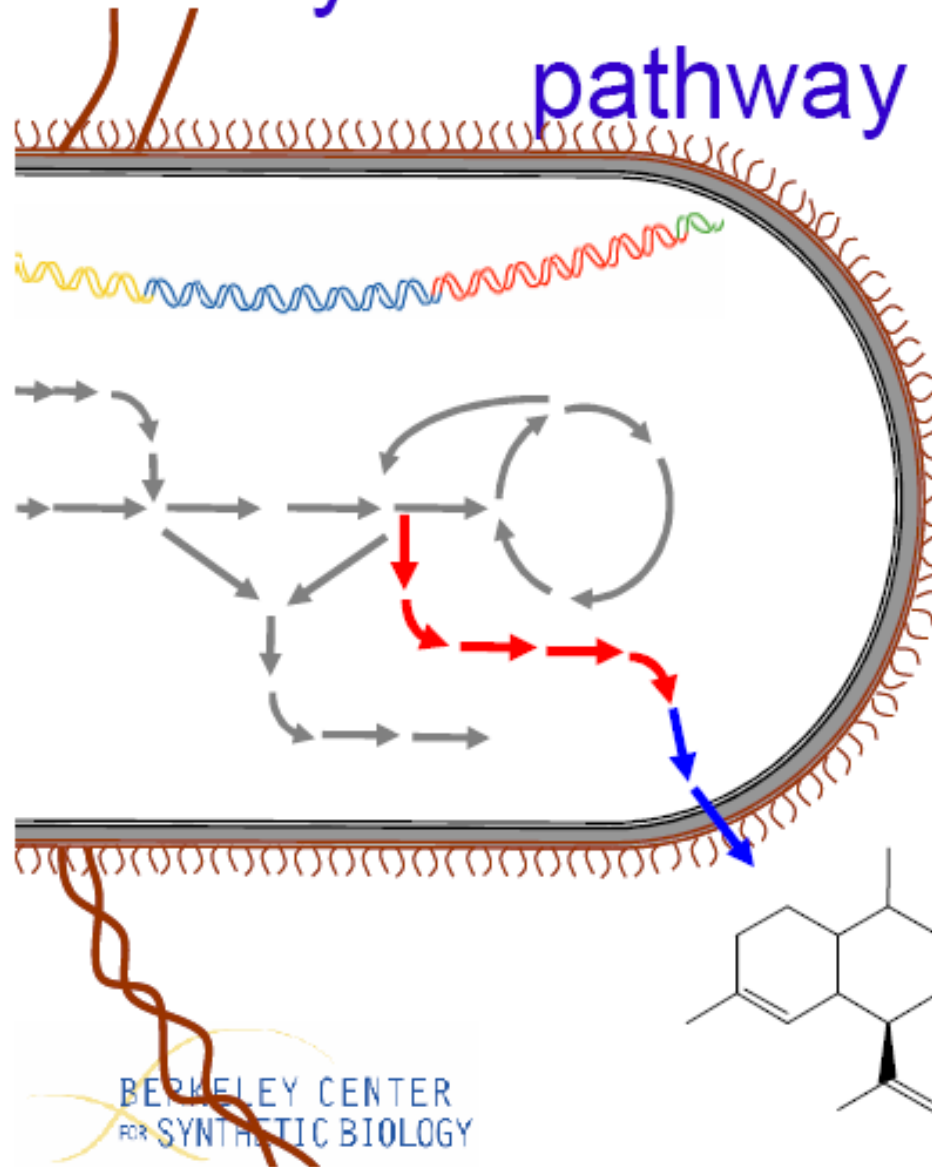
Three-fold improvement in production



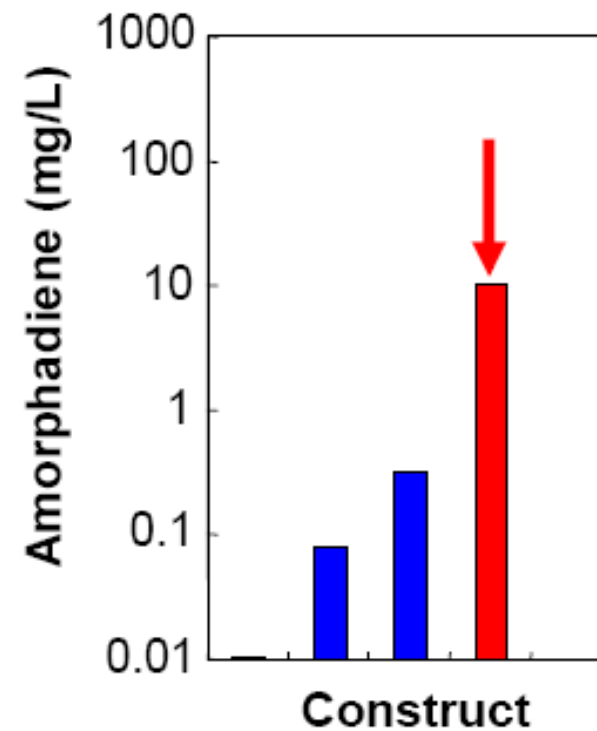
# Recruiting the cholesterol biosynthetic pathway from yeast

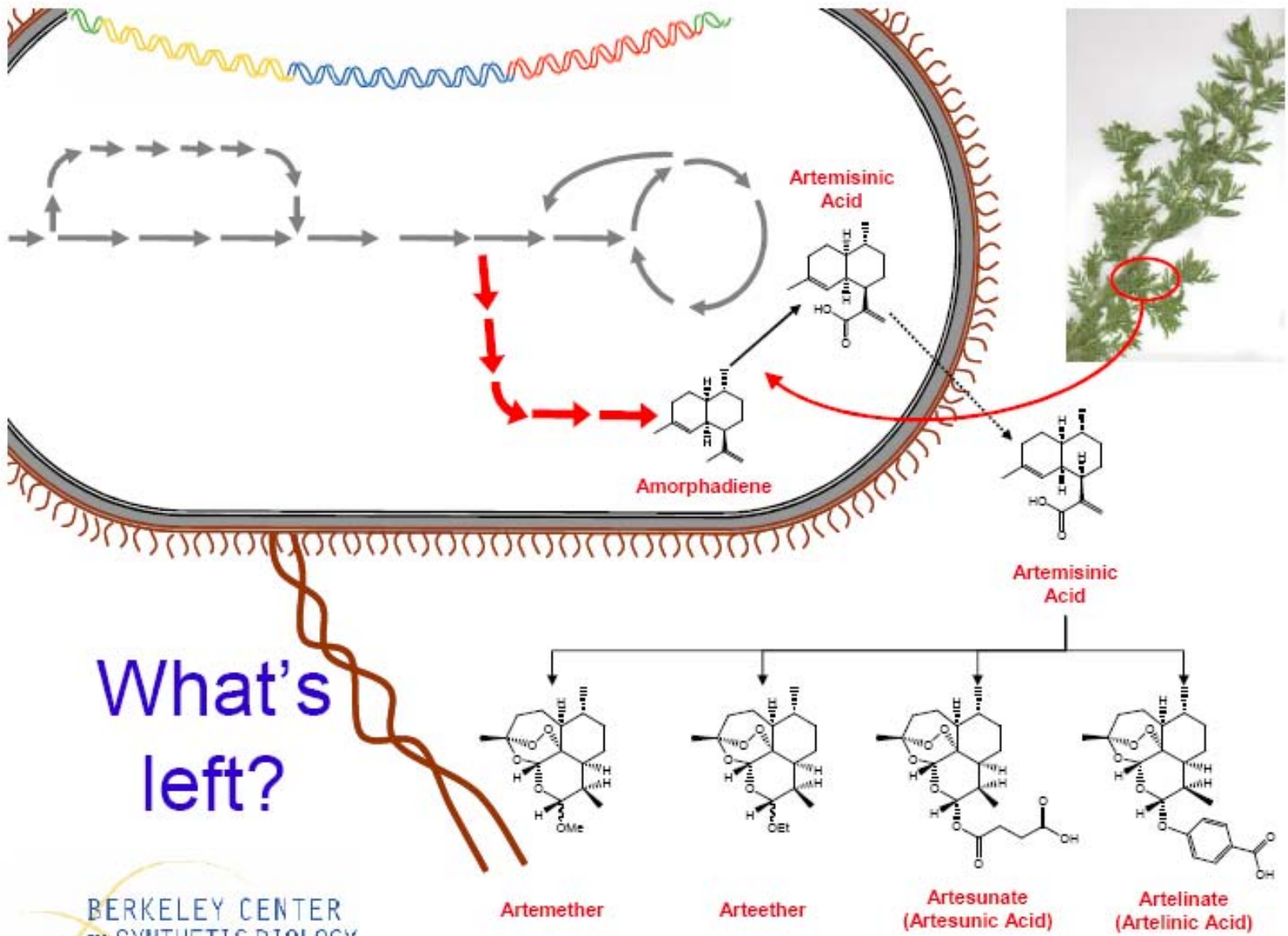


# The yeast cholesterol biosynthetic pathway improves yields



**~90-fold improvement in production**





What's left?



## Artemisinin costs

### Artesunate combination treatment

**Current cost of drug**

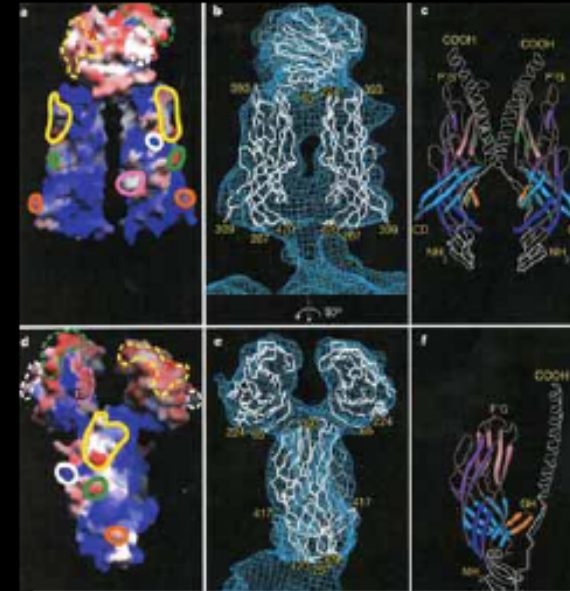
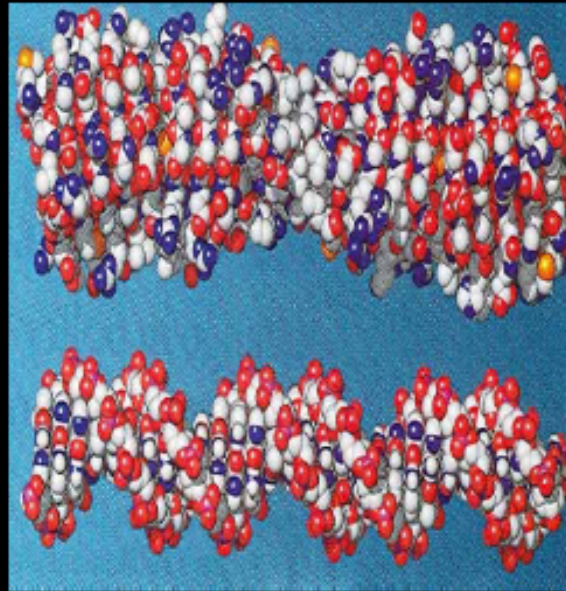
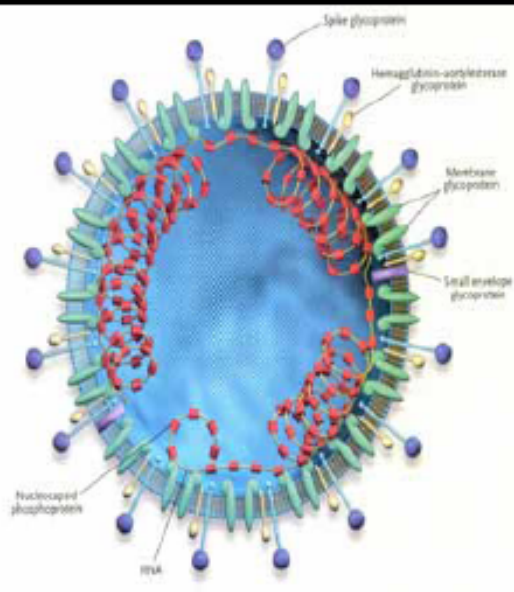
**\$2.25-2.50**

**Cost with new process**

**\$.21/.12**

# Engineered Microorganisms and Cells

## As living pills



# Coding Capacity vs Physical Limitations

A drug molecule



a protein molecule



a "minimal cell"

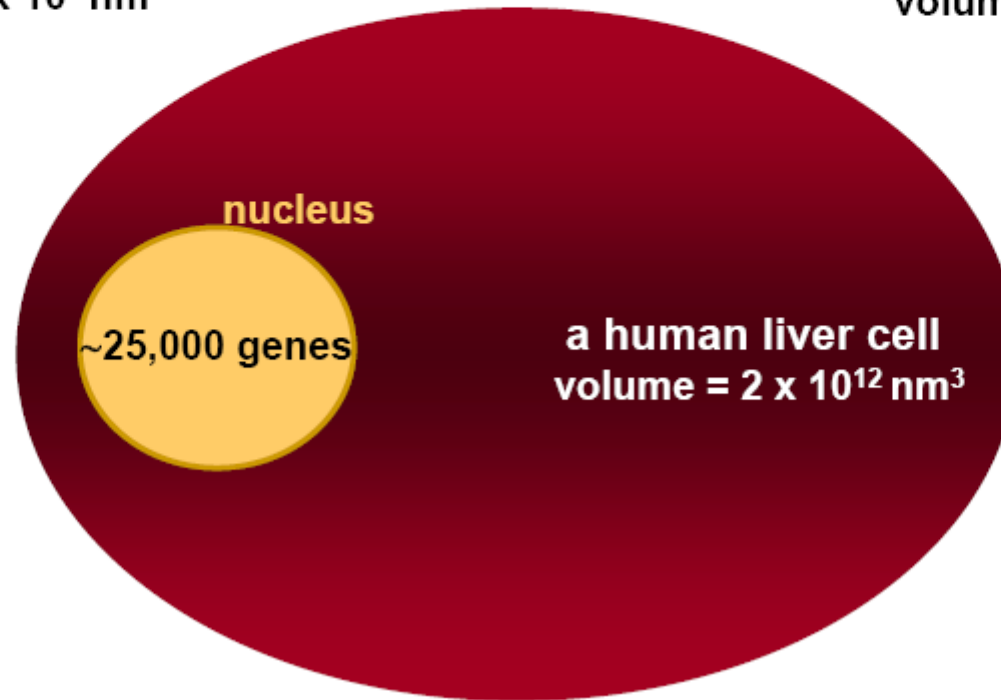


~300 genes  
volume =  $3 \times 10^6 \text{ nm}^3$

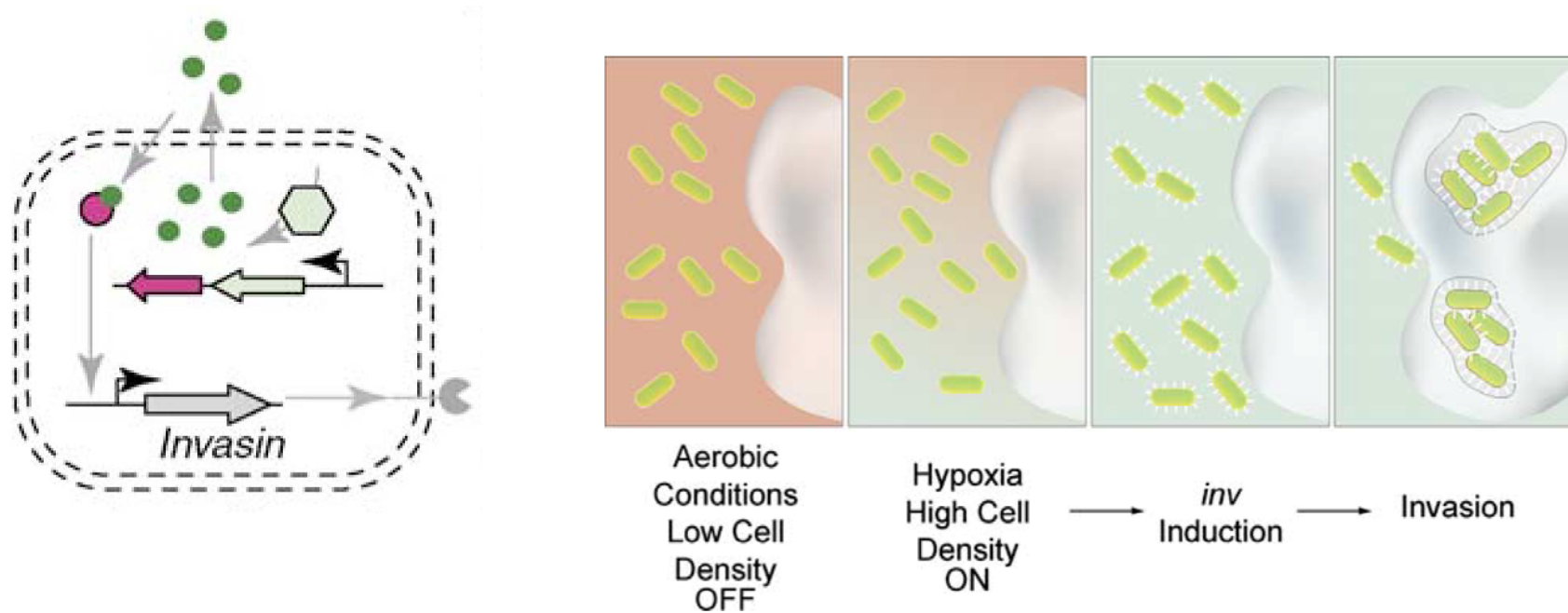
*Escherichia coli*



~4000 genes  
volume =  $5 \times 10^9 \text{ nm}^3$



# Towards “biobots”: cancer invading bacteria



*Vibrio fischeri*: Quorum sensing (cell density sensing)

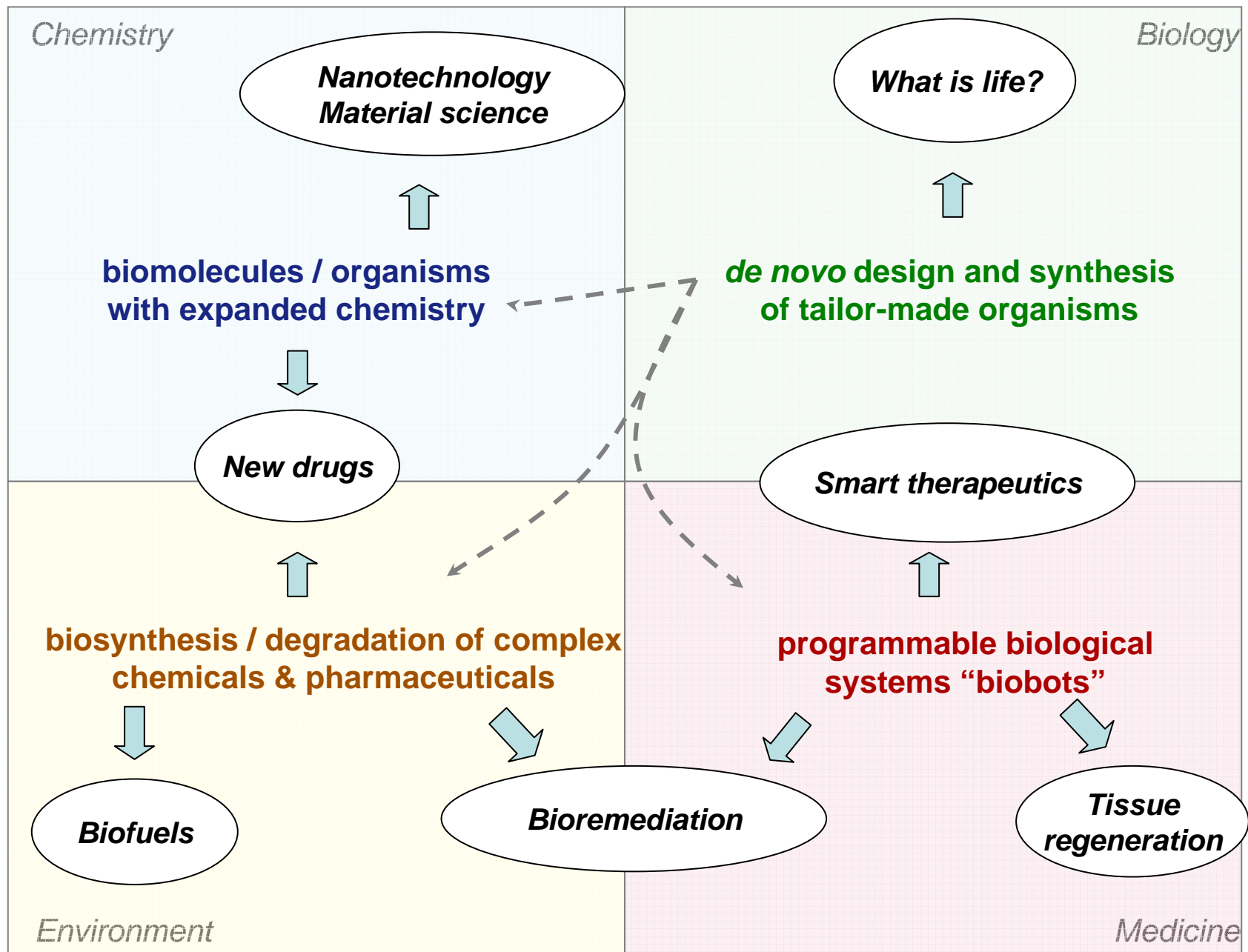
*E.coli*: hypoxia responsive promotor (*fdhF*)

*Yersinia pseudotuberculosis*: invasins (*inv*)

# Perspectives and Ethics



# What can synthetic biology deliver ( in 5-10 years time)?



## **Synthetic Biology**

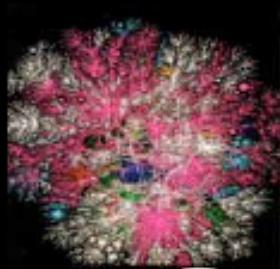
- **anticipated powerful driver of industrial innovation and market disruption**
- **new aspirants, new cross-sector relationships and new markets**
- **the next era in the evolution of human mastery of the environment**
  - **agronomic, industrial, informational, genetic, biomimetic, designed life forms**
  - **enhancement and eugenics?**

## **The End of the Darwinian Interlude**

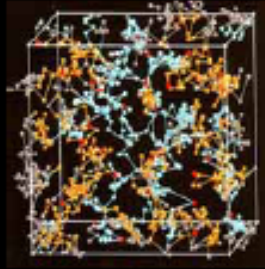
- **early 'biotic' world**
  - massive lateral gene transfer and 'loose' definition of species
  - rapid evolution as a communal affair
- **the Darwinian interlude**
  - majority period in evolution (3 billion plus) years
  - slow pace of change and species 'isolationism'
- **synthetic biology**
  - revival of pre-Darwinian era of horizontal gene transfer

# Synthetic Biology: Inter-disciplinary Convergence and Complex Policy Issues

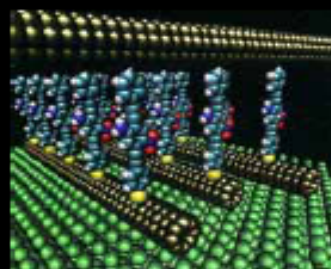
**Systems  
Biology**



**Computational  
Biology**



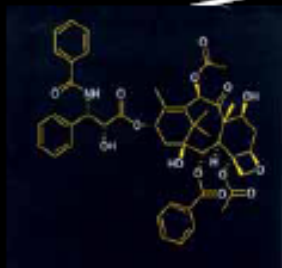
**Materials and  
Nanobiotechnology**



**Industrial Bioprocess  
Engineering**



**Specific Applications**



**Occupational  
Safety**



**BIOHAZARD**

**Dual-Use  
Applications**



**Public and Media  
Responses**



**Public Policy and  
Regulatory Oversight**

## **Synthetic Biology**

- **complex policy issues**
- **design of novel life forms and societal response**
- **dual-use applications**
- **public and media attitudes to perceived risks and benefits**
- **proactive, predictable, evidence-based regulatory frameworks**



# Key points raised in 2009

Samuel, Selgelid, Kerridge, EMBO reports

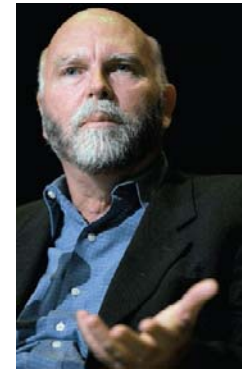
- Misuse for weapons/terrorism
  - Spelled out in greater detail
- Environmental impact
- Health impact
- Justice--patents, ownership
- Commerce and self-regulation

*Science* 15 June 2007:  
Vol. 316. no. 5831, p. 1557  
DOI: 10.1126/science.316.5831.1557

[Prev](#) | [Table of Contents](#) | [Next](#)

## News of the **Week**

### **SYNTHETIC BIOLOGY: Attempt to Patent Artificial Organism Draws a Protest**



An activist group's concern about maverick genome sequencer J. Craig Venter's intention to patent an entirely synthetic free-living organism has thrown a spotlight on the emerging intellectual-property landscape in this hot new field. The protesters claim that Venter wants his company to become the Microsoft of synthetic biology, dominating the industry.

Venter hopes to use the artificial life form, which he says does not yet exist, as a carrier for genes that would enable the bug to crank out hydrogen or ethanol to produce cheap energy. Duke University law professor Arti Rai says the patent, if awarded, "could be problematic" only if Venter's product became the standard in the field. But Venter says this application is just the start: He plans to patent methods that would cover more than the single microbe described in the application. "We'd certainly like the freedom to operate on all synthetic organisms" that could serve as a chassis for swapping out genes, says Venter, whose research team is at the nonprofit J. Craig Venter Institute in Rockville, Maryland, but who recently started a company to commercialize the work.

Filed last October and published by the U.S. Patent and Trademark Office on 31 May, the application describes "a minimal set of protein-coding genes which provides the information required for replication of a free-living organism in a rich bacterial culture medium." The application cites work by Hamilton Smith and others on Venter's team on a simple bacterium called *Mycoplasma genitalium* that they are using to determine the minimum number of genes for life. They want to synthesize this "minimal genome" from scratch, get it working inside a cell, then add genes to produce cheap fuels (*Science*, 14 February 2003, p. [1006](#)).

In a press release, the ETC Group, a technology watchdog in Ottawa, Canada, called Venter's "monopoly claims ... the start of a high-stakes commercial race to synthesize and privatize synthetic life forms." ETC calls for the U.S. and international patent offices to reject the patent so that societal implications can be considered. ETC also cited a recent *Newsweek* interview in which the scientist says he wants to create "the first billion- or trillion-dollar organism."

Venter says this is just one of several patent applications that would give his company, Synthetic Genomics Inc., exclusive rights to methods for making synthetic organisms. The artificial *Mycoplasma* "may or may not be" the one used to generate hydrogen or ethanol, he says; his team is working on several species. "We haven't given any thought to" the licensing conditions, but in any case, they would not impede work in academic labs, says Venter, adding, "This is a problem that we hope will have hundreds of solutions."

M  
V  
S  
S  
H

# iGEM

The International Genetically Engineered Machine (iGEM) competition is a worldwide competition between teams of undergraduate, graduate and PhD students from different universities who are interested in the field of synthetic biology. When creating their projects, the teams use the BioBricks (standardized biological parts) from the iGEM registry to make new, functional biological systems. The teams come from Asia , Europe, United States and Canada to the Massachusetts Institute of Technology (MIT) in Boston to present their project.

C..

# “Synthetic Biology: ENGINEERING LIFE”

## IMAGINATION IS THE LIMIT

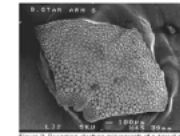
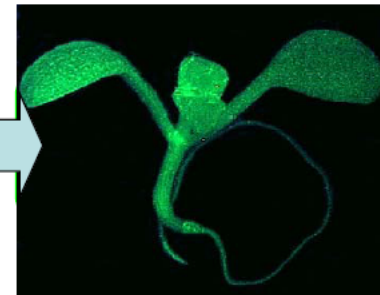
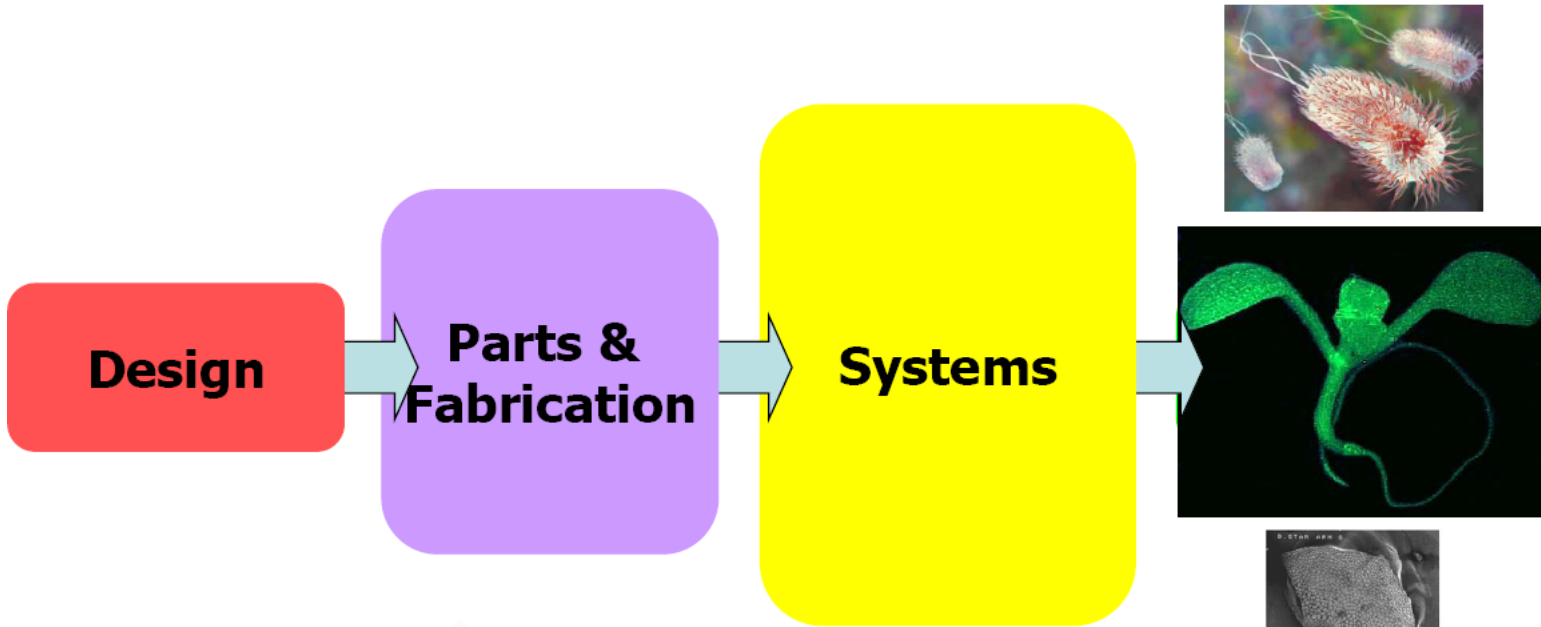
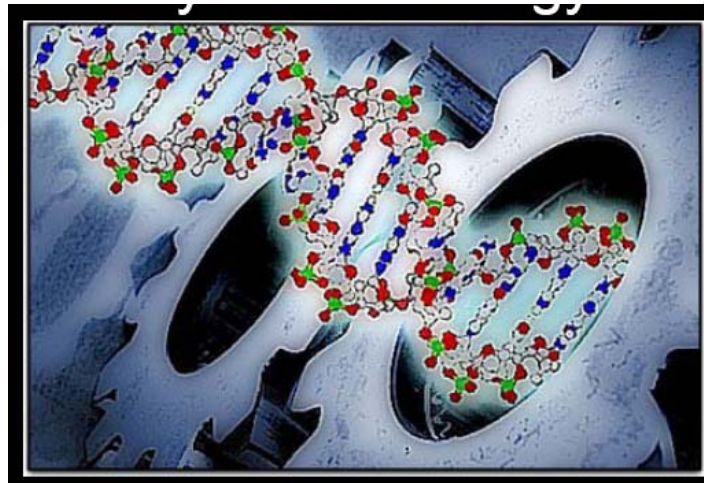


Figure 2. Scanning electron micrograph of a dorsal arm of a light-sensitive protein. The structure is made up of single crystal units, is decorated with microtubules that concentrate light on the ring nerve bundle. Courtesy of L. Araki.



# Questions



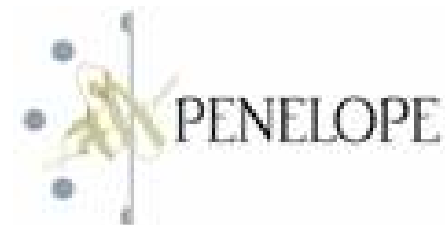


# Acknowledgements

**ZNIP**



**NetSensor**



# The Future of Synthetic Biology?

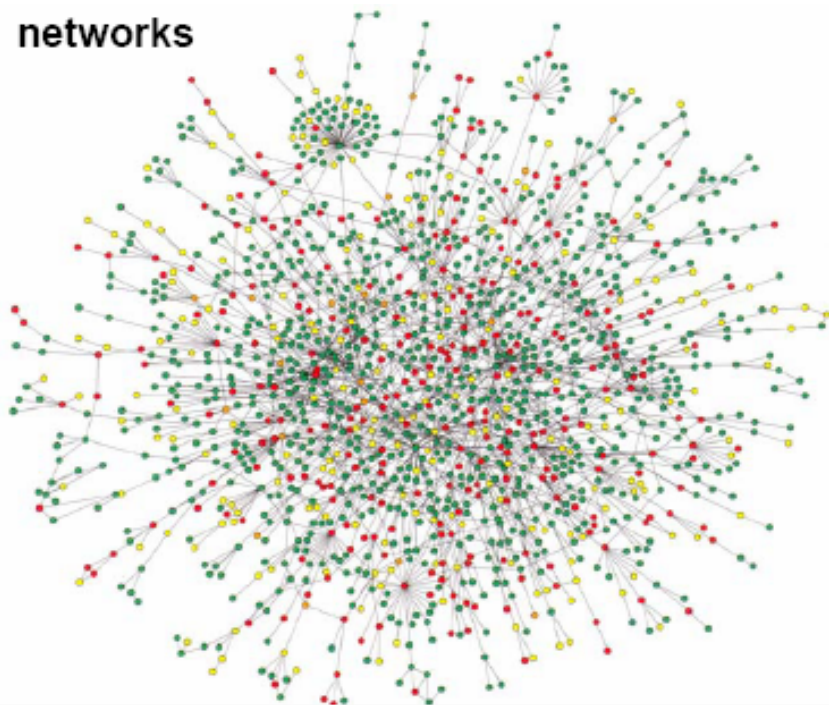


# What are the key issues? Not these

- Misuse for weapons/terrorism
  - Any technology can be misapplied/put to pernicious use
  - Issue is into whose hands does the technology fall?
- Environmental impact
  - Can be handled through regulation
- Health impact
  - Can be handled through regulation and enforcement
- Justice--patents, ownership, commerce
  - No special issue relative to synthetic biology



## networks



## pathways & circuits



## complexes



## molecules



## contributing disciplines

**Maths and Computing**  
Simulation-based systems  
analysis  
Modelling of complex states

**BIOSYSTEMS**

**Physical Sciences**  
Photonics/Imaging  
Spectroscopy  
Reaction  
mechanisms/kinetics  
Structural methods

**Engineering**  
Nanoscale  
technology/metrology  
Electronics  
Process control theory  
Systems modelling



## Defining the Rule Sets for Biological Design, Assembly and Function

- common genetic (digital) code in all life forms
- genomes encode a limited series of structural building blocks (protein motifs and programmed assembly)
- combinatorial assembly of protein building blocks generates extravagant structural and functional diversity



# Ethics of new technology: dual-use

“evil”

**Technophobic (Bill Joy)**

- Control and if necessary extinguish technology
- Top-down monitoring and control, hierarchical, few in power (surveillance)
- Philosophy of secrecy
- Licensing, monitoring, gated access, tracking, inspection
- Challenges are concentrated, government provides national security

“good”

**Technophilic (Ray Kurzweil)**

- Technology is inevitable
- Bottom-up monitoring, democratic, participatory, many in power (surveillance)
- Philosophy of openness
- Proliferation of open source projects (OpenWetWare, diybio, biopunk, biohack)
- Challenges are distributed, citizen defense, biosensors

# Biological warfare and public health

- Can these technologies be weaponized?
- Risk assessment
  - Access to existing samples
  - Creating pathogens is difficult
  - Superbugs (Staph aureus), emerging infections
- Simultaneous development of defenses
  - Sensors

## **Exploring Biospace**

### **The Design Power of Combinatorial Interactions and Assemblies**

- **30,000 genes**
- **two genes cooperate to create a function**  
**= $(30,000 \times 29,999)/2 = 449,985,000$  potential combinations**
- **100 genes generate a complex function**  
**= $10^{289}$  potential combinations**
- **if any combination of genes can generate a function**  
**=  $2 \times 10^{72,403}$  potential combinations**
- **number of theoretical possibilities for synthetic assembly (biospace) far exceed narrow molecular space sampled in evolutionary time**

# Synthetic Biology

## Identification of Relevant Genetic Source Code

- metagenomic screening for desired trait(s)

## Minimum Genomes (Universal Producer Cell)

- “plug and play” programmed expression of gene “cassettes”

## Control of Gene Expression Networks

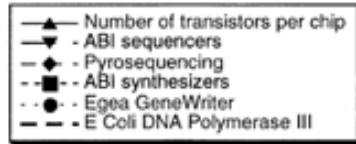
- programmed biosynthesis
- induction of novel properties in specific cells/tissues/organisms



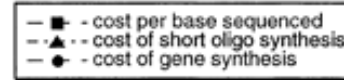
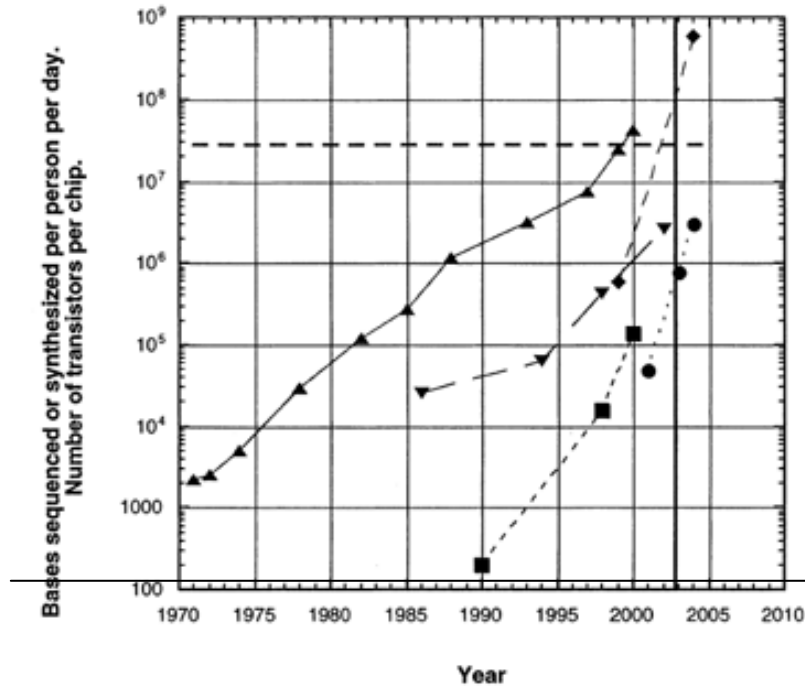
**Biological Design: "Endless Forms Most Beautiful":  
Limitless Diversity From Combinatorial Assemblies of Limited Building Blocks**



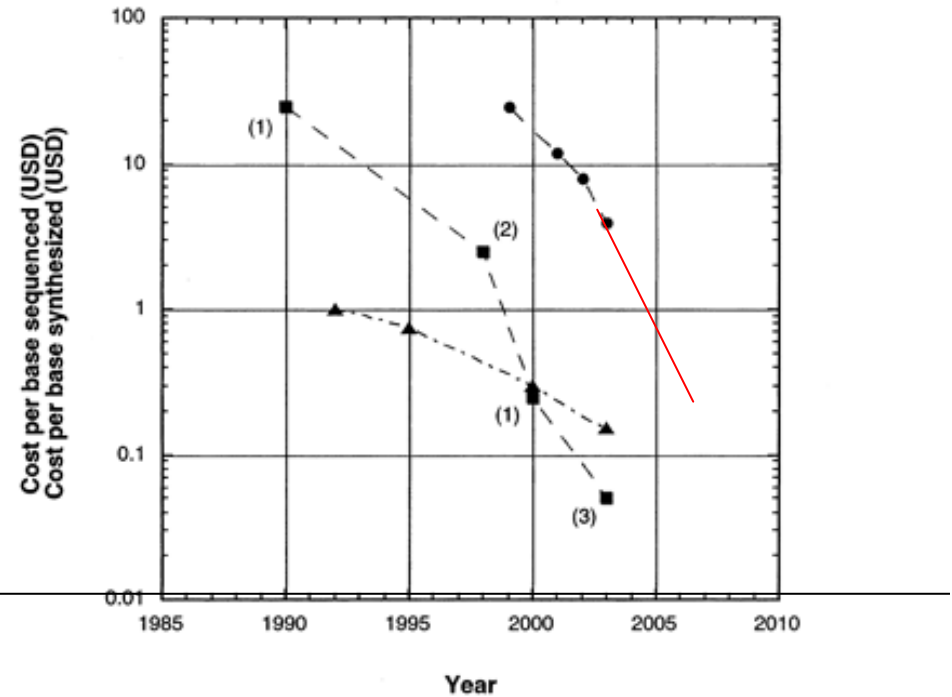
# Sequencing & synthesis follow Moore's law



**Productivity Improvements in DNA Synthesis and Sequencing**  
(as of October, 2002)



**Cost Per Base of Sequencing and Synthesis**



Human genome sequence (2001): 10 yrs, \$ 3 billion / (2007) 2 months \$ 2 million  
Total synthesis of bacterial genome (2008)

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