



南方医科大学

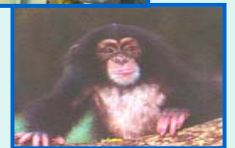
Southern Medical University

# CELL & TISSUE ENGINEERING

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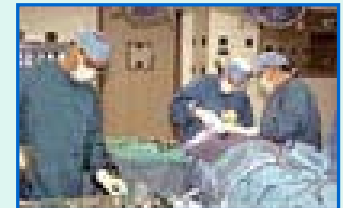
Joint Research Center of Biomedical Engineering and Material



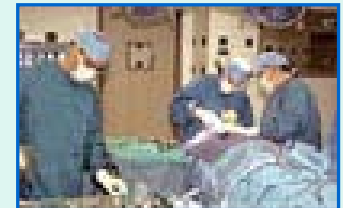
- What if lost limbs or organs could be regenerated?
- What if drug development and chemical testing were more efficient and less costly?
- Can deadly biological and/or chemical agents be rapidly identified?
- Are there ways to reduce our reliance on animal testing?

# What's Tissue Engineering?

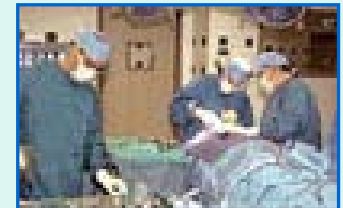
- TE is an emerging **multidisciplinary field** involving **biology, medicine, and engineering** that is likely to revolutionize the ways we improve the health and quality of life for millions of people worldwide by restoring, maintaining, or enhancing tissue and organ function. (**NIH**)



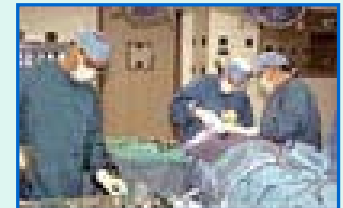
- **Tissue engineering research includes:**
  - **biomaterials,**
  - **cells,**
  - **biomolecules,**
  - **engineering design aspects,**
  - **biomechanics,**
  - **informatics to support tissue engineering**
  - **stem cell research.**



- **A brief history of TE**
  - **First put forward in 1985**
  - **Pre-clinical research during 1990s**
  - **TEMP (Tissue Engineering Medical Product) in late 1990s**



- **A brief history of TE**
  - **The Multi-Agency Tissue Engineering Science (MATES) Interagency Working Group (IWG), 2000**



# The Multi-Agency Tissue Engineering Science (MATES) Interagency Working Group (IWG)

[Department of Agriculture](#) (USDA)

[Department of Commerce](#)

[National Institute of Standards and Technology](#) (NIST)

[Department of Defense](#) (DoD)

[Defense Advanced Research Projects Agency](#) (DARPA)

[Centers for Medicare & Medicaid Services](#) (CMS)

[Department of Health and Human Services](#) (HHS)

[Food and Drug Administration](#) (FDA)

[National Institutes of Health](#) (NIH)

[Department of Energy](#) (DOE)

[National Aeronautics and Space Administration](#) (NASA)

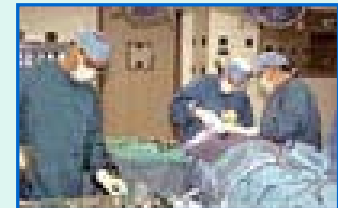
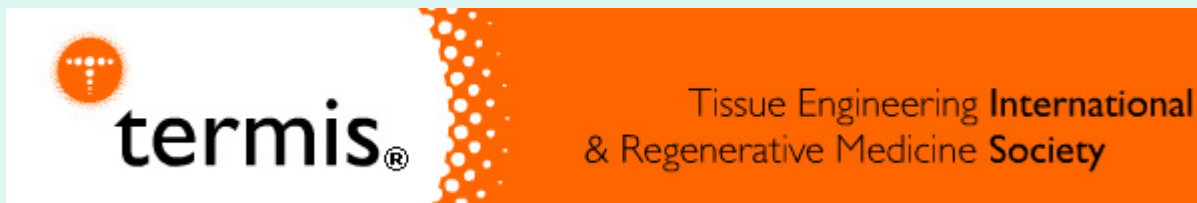
[National Science Foundation](#) (NSF)

[Naval Research Laboratory](#) (NSF)

[Veterans Administration](#) (VA)



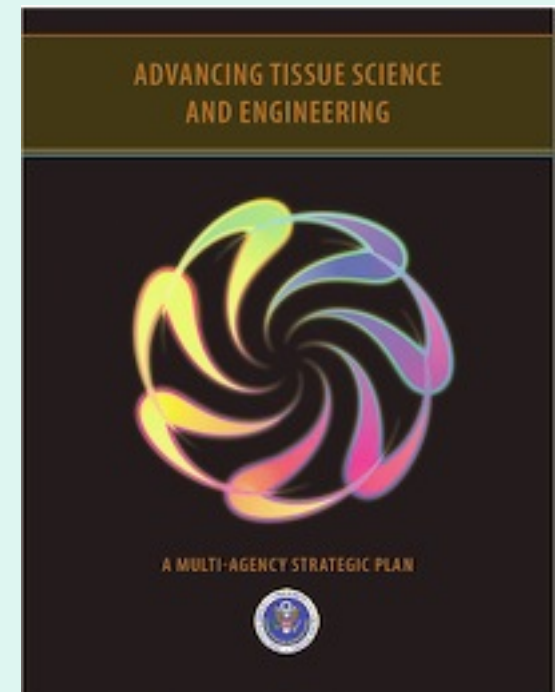
- **A brief history of TE**
  - **TERMIS (Tissue Engineering & Regenerative Medicine International Society, 2005)**





# Advancing Tissue Science And Engineering

- A Multi-agency Strategic Plan ( 2007 )
- From earliest diagnostic testing To the advanced stages of therapy.

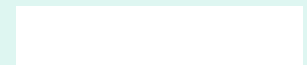


# Four overarching goals

- Understanding and controlling the cellular response;
- Formulating biomaterial scaffolds and the tissue matrix environment;
- Developing enabling tools;
- Promoting scale-up, translation and commercialization;

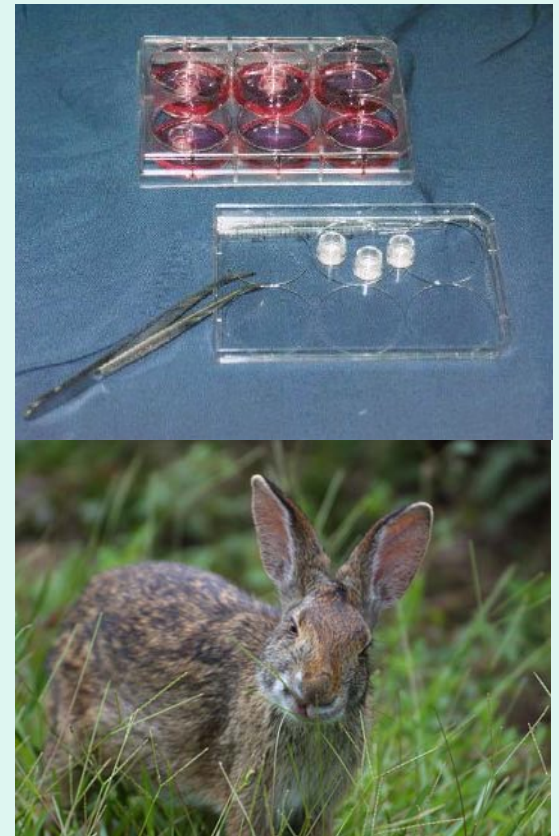
# **Understanding and Controlling The Cellular Response**

- **Tissue-Engineered Disease Models:**
- **3-D tissue models was established to study the growth of tumors in tissue beds and for use in drug screening.**



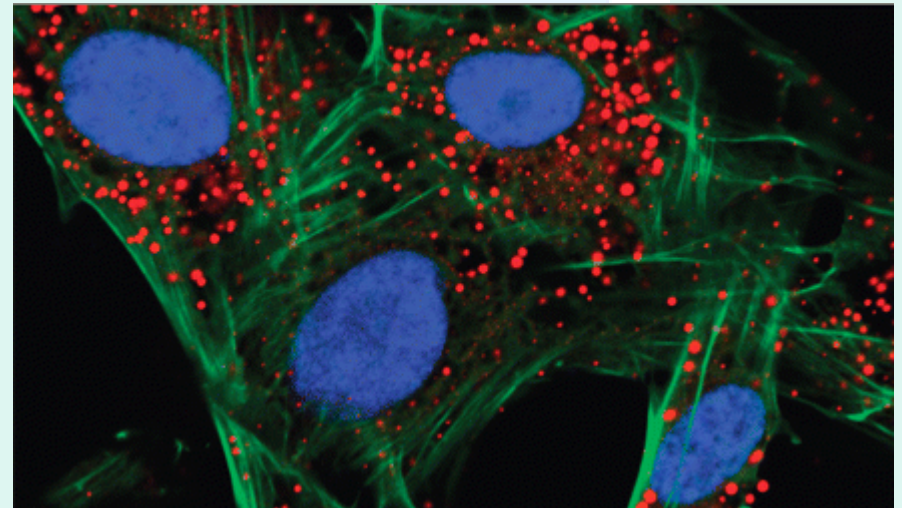
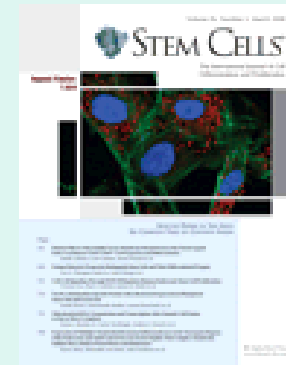
# Understanding and Controlling The Cellular Response

- Engineered Tissues Provide Alternatives to Animal Testing.
- Currently, engineered skin products are the most widely used.



# Stem Cells for Tissue Science and Engineering

- Various tissues of the adult body have been used as a source of stem cells ( e.g., fat, muscle, skin).
- To control cell phenotype and direct tissue formation is the ultimate goal.

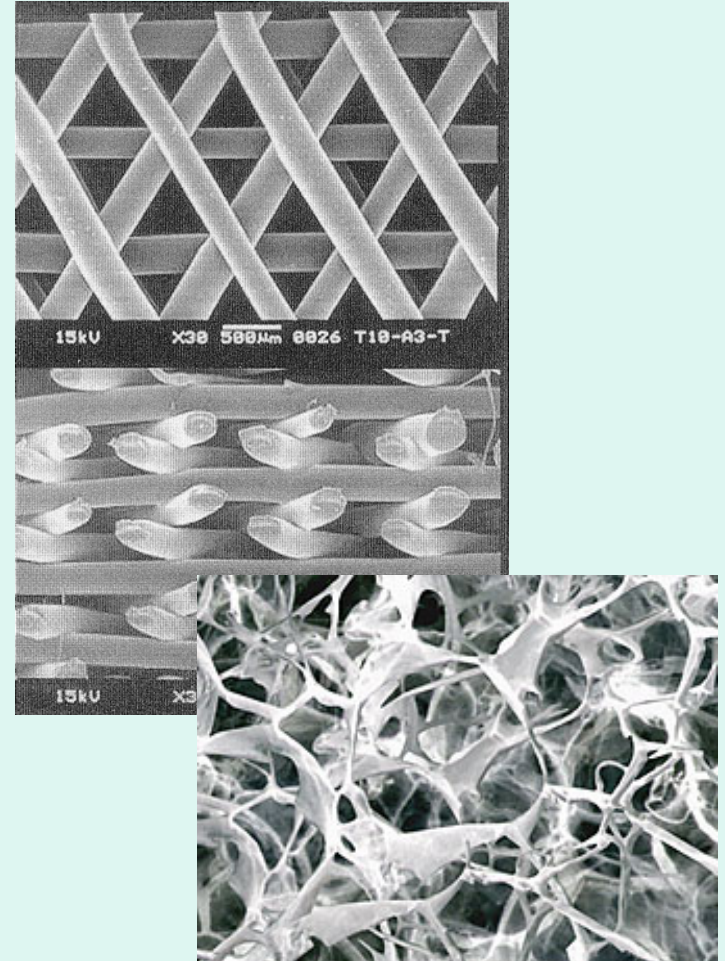


# Formulating biomaterial scaffolds and the tissue matrix environment

- Guide the growth, differentiation and organization of cells;
- Give support, strength and for to tissue and organs.

# Formulating biomaterial scaffolds and the tissue matrix environment

- Hight molecular compound
  - Polylactic acid (PLA),
  - Polyglycolic acid (PGA),
  - Polycaprolactone (PCL)



# Formulating biomaterial scaffolds and the tissue matrix environment

- Matrix:

collagen, elastin, proteoglycans and wide variety of other proteins and signaling molecules.



# Facing difficulties

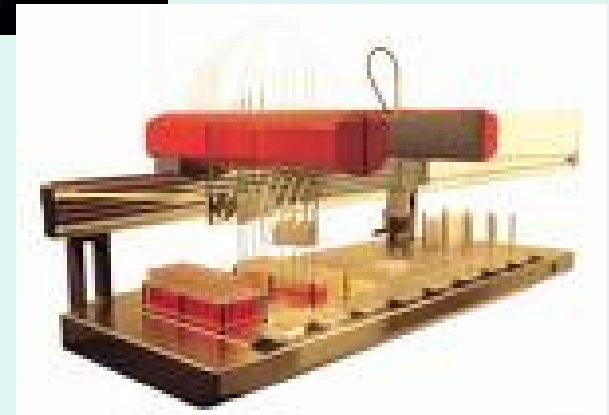
- Understanding and quantifying the relationship between biomaterial scaffolding characteristics and physiological responses remain among the most pressing materials challenges.

# Developing enabling tools

- The development of tissues is the result of molecular and supramolecular interactions that occur on a continuum of time and spatial scales in response to the many chemical and physical parameters within cells, between cells and within organisms.
- Achieving sufficient insight into these complicated relationships needs knowledge and computational model.

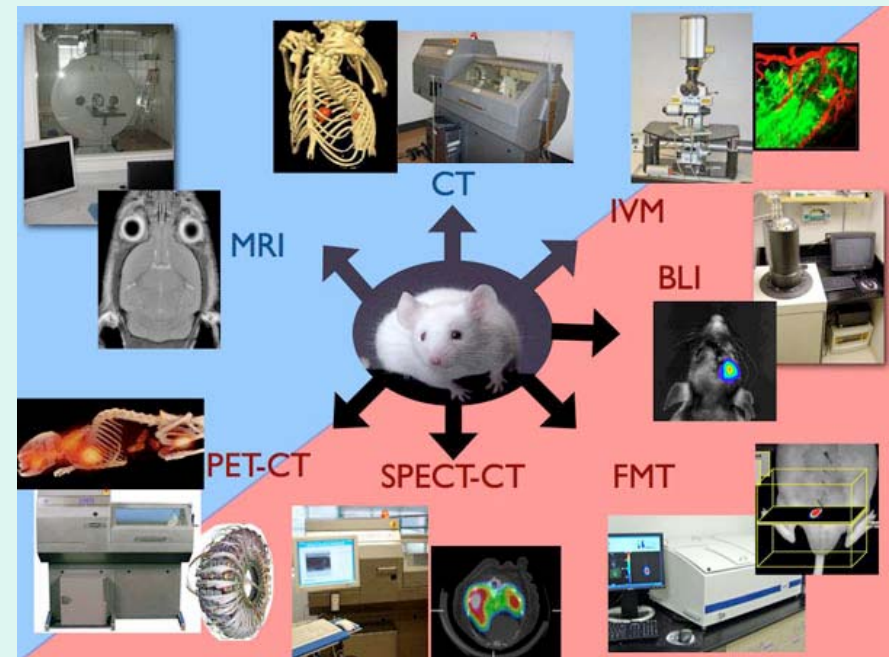
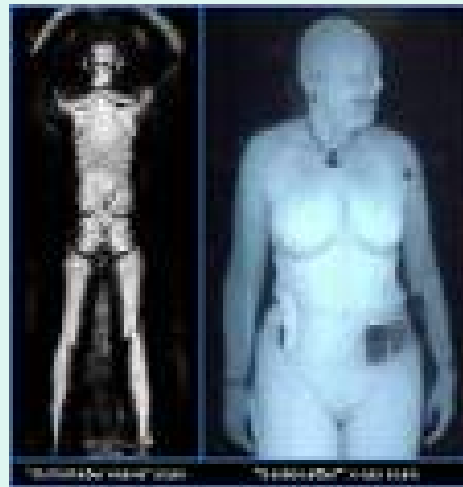
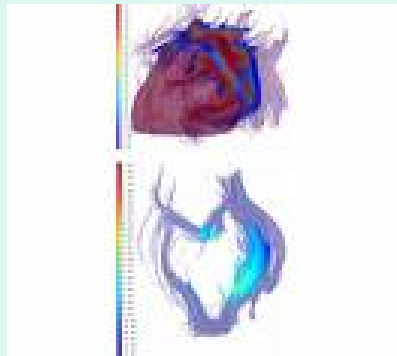
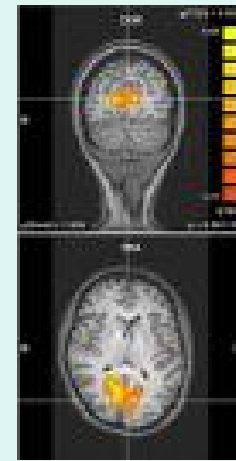
# Biomarkers, High-throughput/ Content Assays and Instrumentation

- Assessing the physiological state or condition of a cell or tissue
- Rapid screening of novel materials as candidates for scaffolding.



# Imaging Technologies

- Imaging challenges include development, optimization, and integration of technologies at the many levels-from molecular to tissue to whole body.

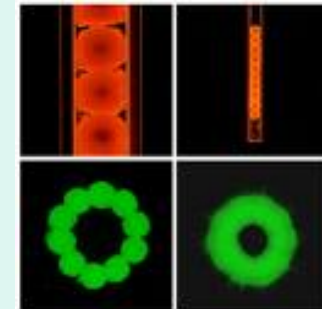
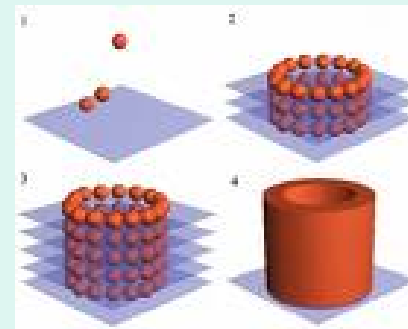
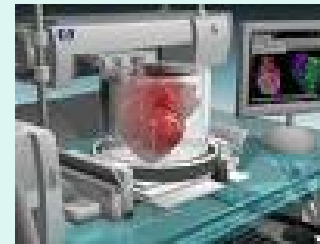


# Computational Modeling and Bioinformatics

- Understanding the intracellular machinery, will require assimilating vast amounts of data into computational models;
- Such models will be critical for allowing prediction and subsequent testing of the large number of parameters that influence cells, tissues, and organs.

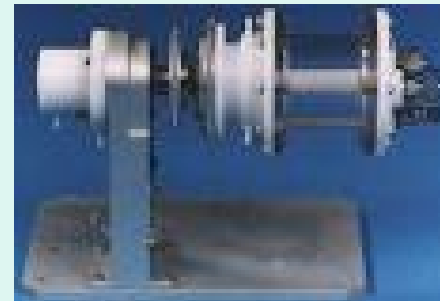
# Fabrication Technologies

- Rapid prototyping (RP) technology was used in TE field;
- Majority of effort should on deposit cells in the layers during the fabrication process.



# Bioreactors

- Producing 3D tissues in vitro requires;
- **NEED:** Design devices that can support self-assembly of multiple cell types into complex tissue structures.



# Tissue Preservation and Storage methodologies

- It is very important for basic research, device development and clinical applications.
- Wood frogs
- Preservation techniques are far from perfect.





# Promoting scale-up, translation and commercialization

- Much work to be done on the technical side in translating lab-scale fabrication techniques into large-scale production of high-quality products that meet GMP standards.
- TEMP is “combination products” and need new standards and rules.

# **TEMPs Industrialization**

# Tremendous market potential

- **Organ replacement >65 ys, 1/5 (USA)**
- **8% medical spending, \$350 Billion per year**
- **Operation \$8 Million/year worldwide**
- **40-90 million days/year in hospital**
- **Bringing a new drug estimated \$800 Million over 12 years**

# Enterprise of TEMPs

- In 2002, 89 biotechnology organizations were involved in TE R&D worldwide.
- Theragnostics TEMP is anticipated to be \$3.7 Billion by 2009.

# TEMP Classification

## ➤ Structure and function

❖ **Simple** (Cartilage,tendon,bone,skin et al.)

❖ **Complex** (Liver、 Kidney et al)

❖ **Cell transplantation and capsule** (Pancreas)

➤ **Composition**

❖ **Cells only( Epicel, Carticel )**

❖ **Cell and biomaterial( Apligraf )**

❖ **Biomaterial only( Alloderm )**

# Total spending



— Lysaght, Tis. Eng. 7,2001

# Enterprise of TEMPs

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	<b>Structural</b>	<b>Metabolic</b>	<b>Cellular</b>
<b>Examples</b>	<b>Skin, bone, heart valves</b>	<b>Bioartificial organs</b>	<b>Cell transplantation</b>
<b>Employee</b>	<b>1980(60%)</b>	<b>570(11%)</b>	<b>890(27%)</b>
<b>R&amp;D spending</b>	<b>\$3.63亿</b>	<b>\$0.68亿</b>	<b>\$1.74亿</b>

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— *Lysaght, Tis. Eng. 7, 2001*



# Comparison of USA & European

	<b>USA</b>	<b>European</b>
<b>Funding Sources</b>	Industry	Government
<b>Research focus</b>	Application	Basic
<b>Cellular sources</b>	auto/allograft	autograft
<b>Biomaterials</b>	new material	modification
<b>Bioreactor</b>	On top	
<b>Standards</b>	On top	

*WTEC, Jan, 2002*

Thank you fro your attention!

