

Standards in Scientific Communities II; Cell Viability

Module 3, Lecture 4

20.109 Spring 2010

Lecture 3 review

- What can you learn from a confidence interval? A t-test?
- What three general engineering principles might help make biology more “engineerable”?

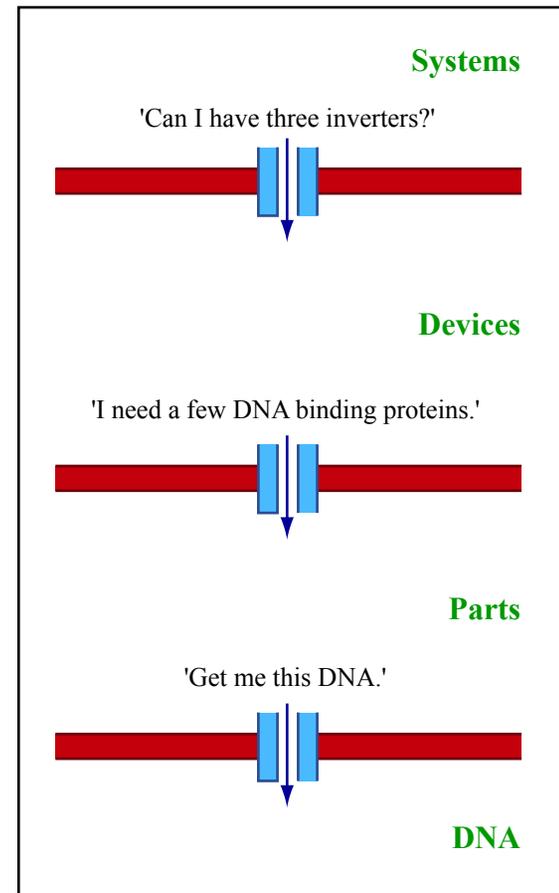


Image by MIT OpenCourseWare.

See D. Endy, *Nature* **438**:449

Topics for Lecture 4

- Standards in tissue engineering
 - introduction
 - writing exercise
 - discussion
- Cell viability
 - your data
 - relation to diffusion

How valued are TE standards?

- 2007 strategic plan for TE clinical success by 2021
 - 24 int'l leaders in TE listed high-priority areas
 - 1/3 named standards

- Analysis
 - concept dominance
 - progress so far
 - standards 7th of 14

P.C. Johnson et al., *Tissue Eng* 13:2827 (2007)

TABLE 6. NORMALIZED CONCEPT DOMINANCE
(I.E., TAKING PRESENT PROGRESS INTO CONSIDERATION)

	O/P
Angiogenic control	3.3
Stem cell science	3.2
4. Cell sourcing/characterization.	2.2
Clinical understanding/interaction	2.2
Immunologic understanding and control	2.0
Manufacturing/scale-up	1.1
Regulatory transparency	1.1
7 (tie). Standardized models.	0.8
Multidisciplinary understanding/cooperation	0.8
Expectation management/communication	0.4
Pharmacoeconomic/commercial pathway	0.3
Multilevel funding	0.0

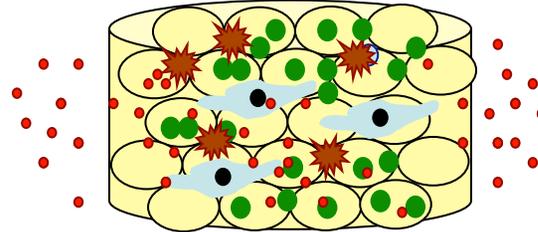


- 2007 US govt. strategic plan
 - standards listed as part of “implementation strategy”

Courtesy of Mary Ann Liebert, Inc. Used with permission.
Source: Johnson, P. C., et al. "Strategic Directions in Tissue Engineering." *Tissue Eng* 13, no. 12 (2007): 2827-2837. doi:10.1089/ten.2007.0335.

How useful are TE standards?

- See 2005 editorial by A. Russell
 - proposes need for standards
 - in data collection and sharing
- Choose and respond to a student excerpt (~10')
- Pros/cons/etc... ?

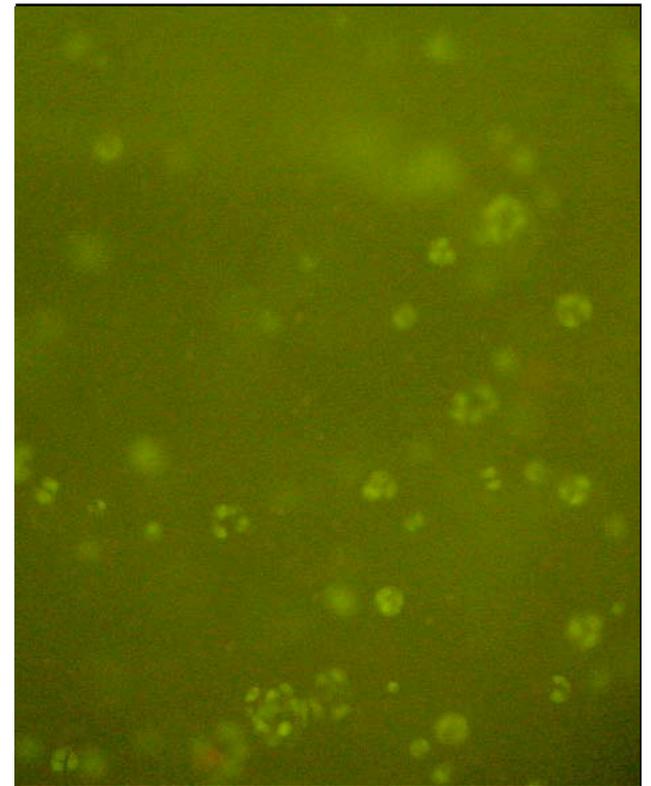


Can we standardize this TE construct?

Module progress: week 2

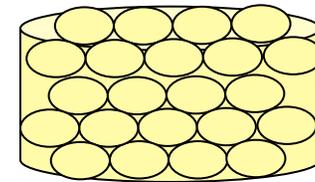
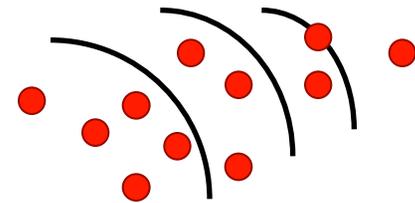
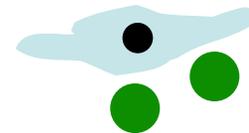
- Day 3: viability/cytotoxicity testing
- Groups generally found
 - mostly live
 - mostly round
 - some clustering
- How do we explain the results?
- How can we improve the assay?
- What conditions killed cells?

Image from W/F Yellow Group



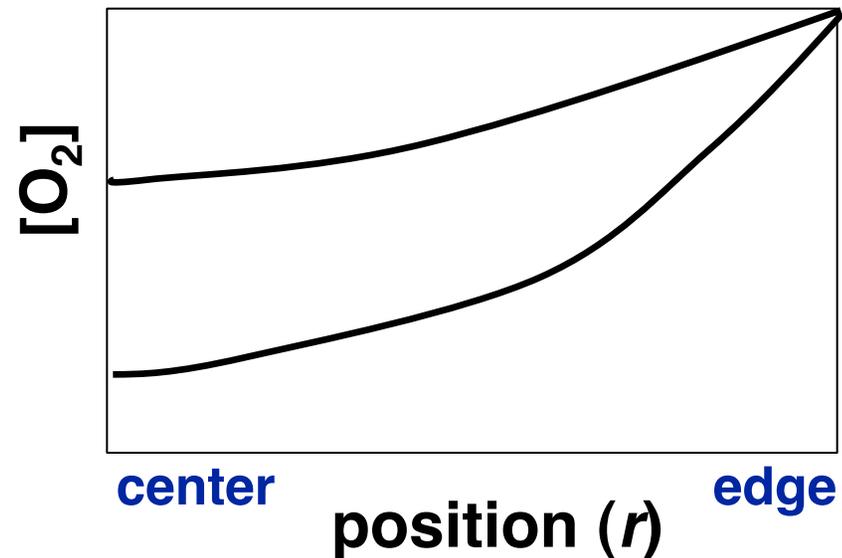
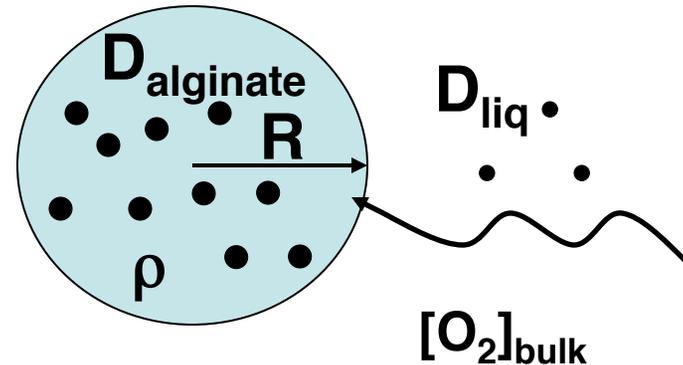
Factors affecting cell viability

- Cell-related
 - density
 - interactions
- Cytokine-related
 - proliferative
 - apoptotic
- Materials-related
 - bulk permeability
 - macro-porosity
 - toxicity



Diffusion in 3D constructs

- Nutrients, O_2
- Affected by
 - construct size R
 - cell density ρ
 - diffusivity D
 - conc. in medium $[O_2]_{\text{bulk}}$
- Concentration profile
 - can be solved (DE)
 - $[O_2] \downarrow$ toward center
 - steepness = $f(D, \rho, \dots)$



Significance of diffusion in TE

- Characteristic limit $\sim 100 \mu\text{m}$
- Diffusion and viability profiles correlated
- How can we make thick tissues?
 - *in vitro*: dynamic/perfusion culture
 - *in vivo*: promote rapid angiogenesis

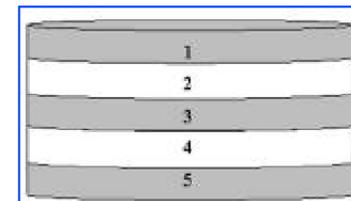
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perfusion system
zeiss.com.sg

Modeling cell viability in TE constructs

- Porous PLGA scaffolds
- Seeded cells as in (A) or (B)
- Observed after 10 days
- Model includes
 - Diffusion
 - O₂ use
 - Cell growth
- Model assumes
 - [O₂]_{bulk} is constant
 - Quasi-steady state

A Cells in odd layers



B Cells in all layers



Dunn, et al. *Tissue Eng* 12:705 (2006)

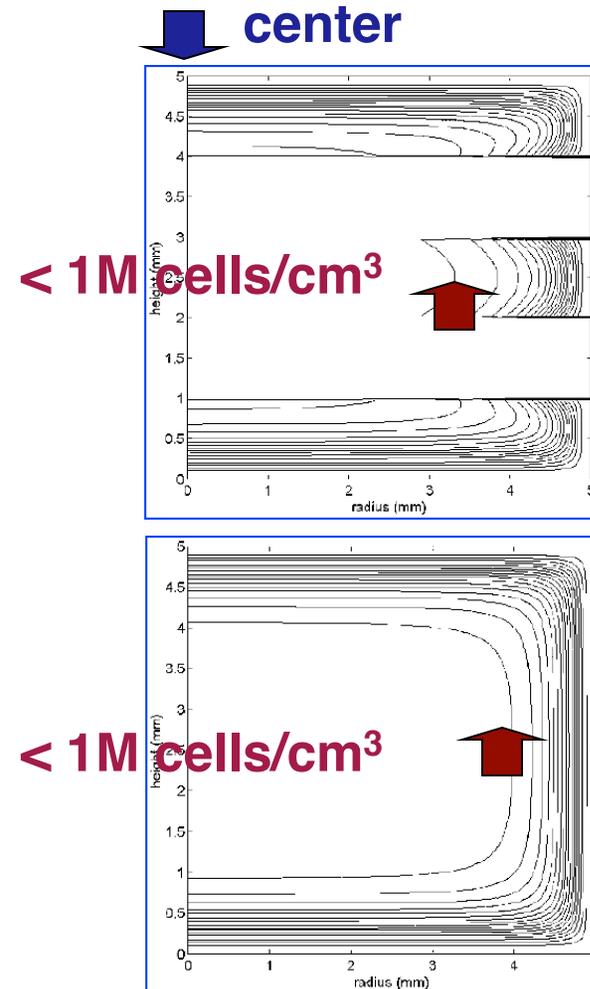
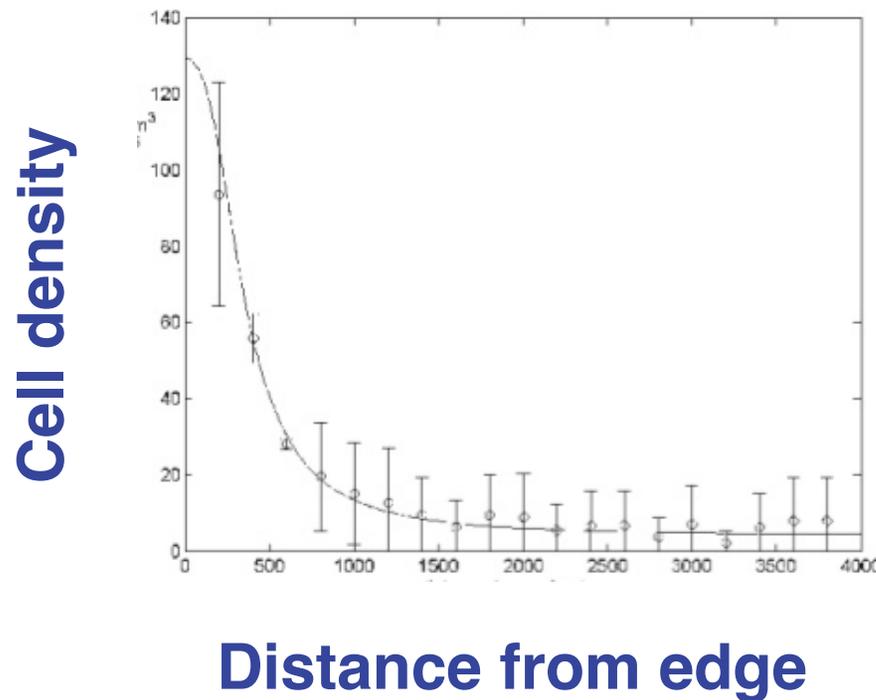
Courtesy of Mary Ann Liebert, Inc. Used with permission.

Source: Dunn, J. C. Y., et al. "Analysis of Cell Growth in Three-Dimensional Scaffolds."

Tissue Eng 12, no. 4 (2006): 705-716.

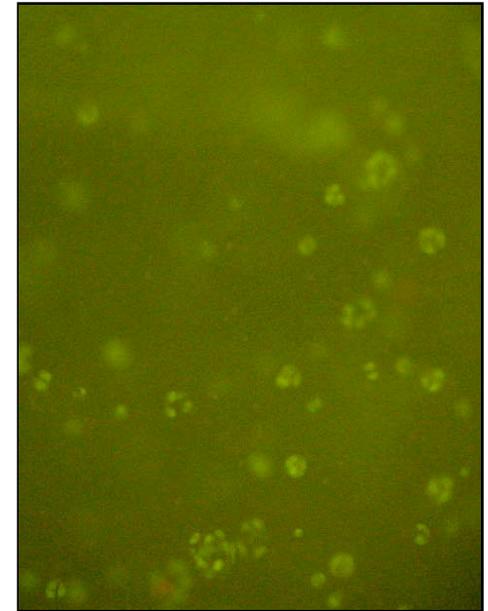
Dunn et al. results for cell viability

- A more uniform than B
- Cell growth matches O_2 tension
- Claim of predictive capability



Lecture 4: conclusions

- Strategies besides standardization may take precedence in some BE fields.
- Cell viability in TE constructs is affected by cell, material, and soluble factors.
- Modeling can elucidate nutrient diffusion and cell viability profiles.



Next time: transcript and protein assays;
advice for module 2 report revision.

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