

Plate 1. Primary explant and outgrowth. A, 4× objective, B, 10× objective. (See Fig. 1.2 for details.)

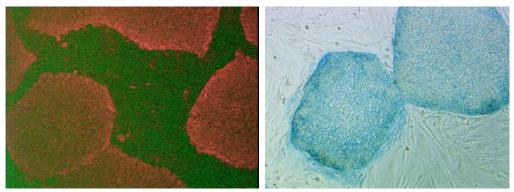


Plate 2. hES cell colonies on mouse embryonic fibroblasts. SSEA-4, red, left and ALP, blue, right. (See Fig. 3.1 for details.)

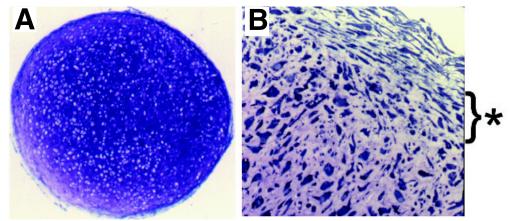


Plate 3. The Toluidine Blue metachromatic matrix of cartilaginous aggregates of human marrow-derived cells after 14 days in chondrogenic medium. A, section of paraffin-embedded whole aggregate; B, higher magnification of edge of a methyl methacrylate-embedded section with the region of flattened cells indicated by asterisk. (See Fig. 4.5 for details.)

Culture of Cells for Tissue Engineering, edited by Gordana Vunjak-Novakovic and R. Ian Freshney Copyright © 2006 John Wiley & Sons, Inc.

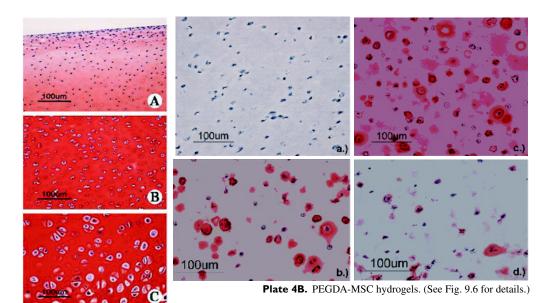
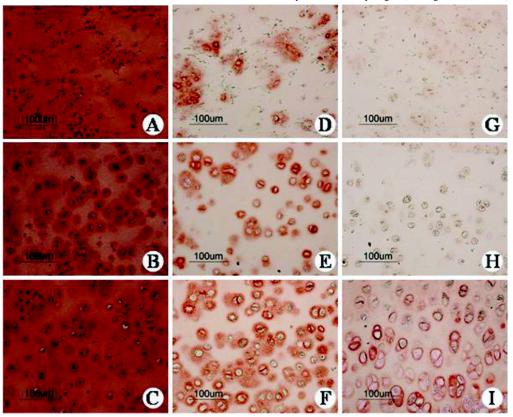


Plate 4A. Juvenile bovine cartilage. (See Fig. 9.2 for details.)

Plate 4C. Multilayered PEGDA hydrogel. (See Fig. 9.5 for details.)



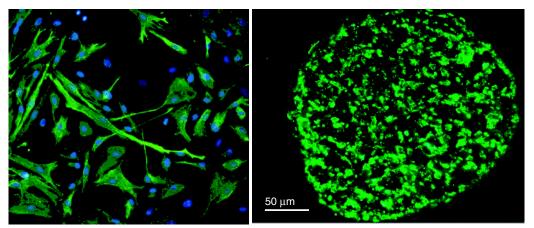


Plate 5A. Human skeletal muscle cells. (See Fig. 10.2.) Plate 5B. Cross section of 10-day in vitro HBAM. (See Fig. 10.5.)

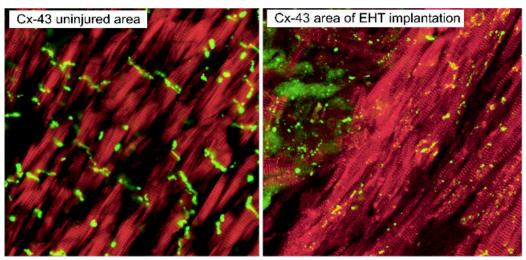


Plate 6A. Effect of EHT implantation of the spatial organization of connexin 43 (Cx-43) in rat hearts. (See Fig. 11.3)

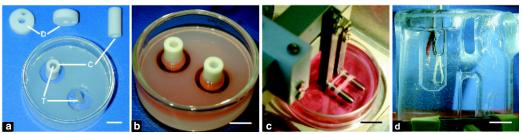
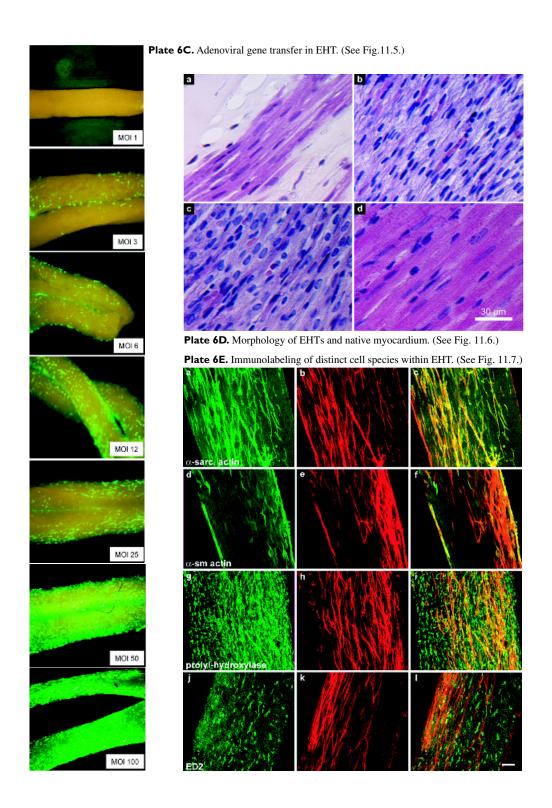


Plate 6B. Experimental setup for EHT preparation, culture, phasic stretch and analysis of contractile function in the organ bath. (See Fig. 11.4 for details.)



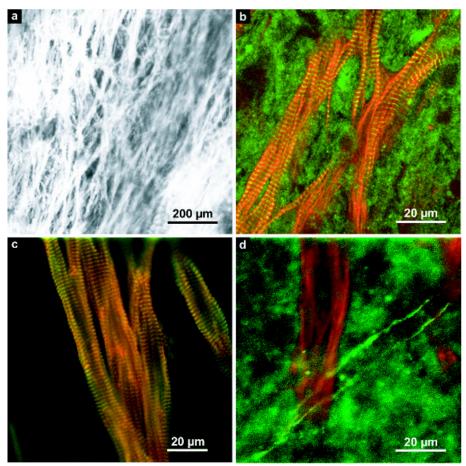


Plate 6F. High-power CLSM of EHT. (See Fig. 11.8 for details.)

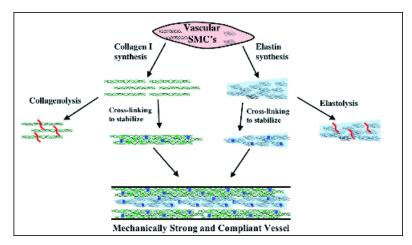


Plate 7A. Secretion of collagen and elastin by smooth muscle cells. (See Figure 12.3.)

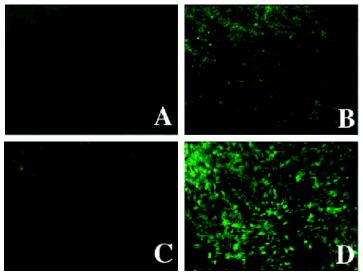


Plate 7B. Enhanced green fluorescent protein (EGFP) expression in cultured ECs. (See Fig. 12.8.)

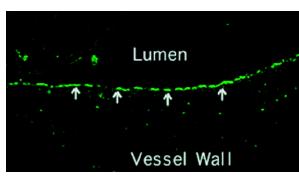
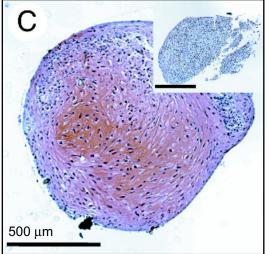
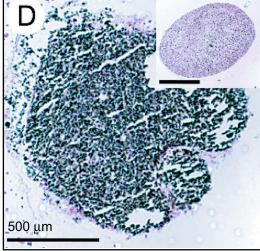


Plate 7C. EGFP expressed on engineered vessel lumen. (See Fig.12.9.)

Plate 8. Characterization of MSCs. C) Chondrocyte differentiation. D) Osteoblast differentiation. (See Fig.13.2.)





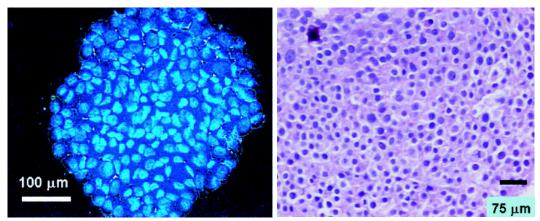


Plate 9A. 3-D assemblies of PC12. Left, static aggregate, right, dynamic aggregate in SLTV. (See Fig. 14.3 for details.)

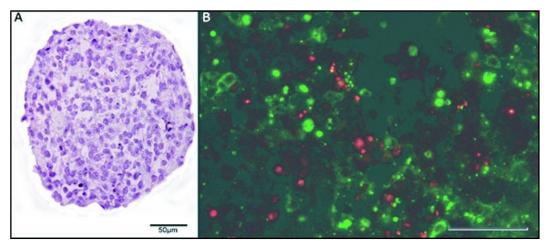
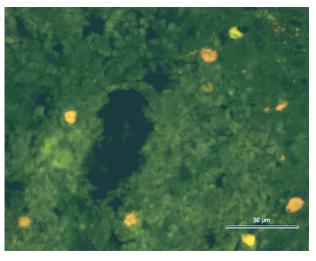


Plate 9B. Morphology and TH content of SNAC tissue constructs. A, SNAC section immumostained for human nuclei, (NT2 cells). B, double immunofluorescence; Sertoli cells, green, TH-positive NT2N neurons red. (See Fig. 14.5 for details.)

Plate 9C. Photomicrograph through a SNAC tissue construct transplant into the rat striatum 4 weeks postsurgery. Surviving TH-positive NT2N neurons (red) double immunostained with antihuman nuclei antibody (green) can be seen along the course of the penetration. These NT2N neurons contain a green nucleus and lighter green cytoplasm, which now appears yellow because of the double label. Some neurite outgrowth is seen in the TH-positive NT2N neuron near the top right of the photomicrograph.



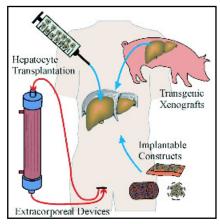


Plate 10A. Cell-based therapies for liver disease. (See Fig. 15.1 for details.)

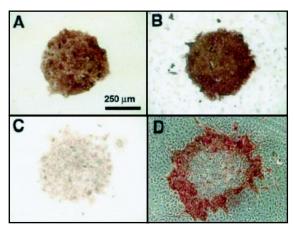


Plate 10B. Intracellular albumin in micropatterned hepatocytes. (See Fig.15.5 for details.)

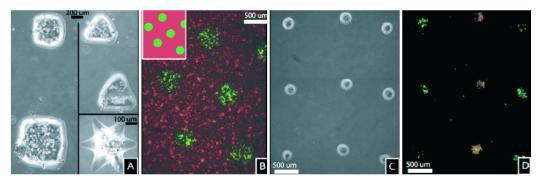


Plate 10C. Hydrogel microstructures containing living cells. (See Fig. 15.10 for details.)

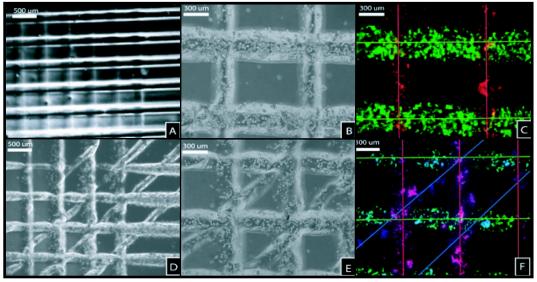


Plate 10D. Multilayer hydrogel microstructures containing living cells. (See Fig. 15.11 for details).