SOCIETAT CATALANA DE TRASPLANTAMENT

BANFF FOUNDATION FOR ALLOGRAFT PATHOLOGY

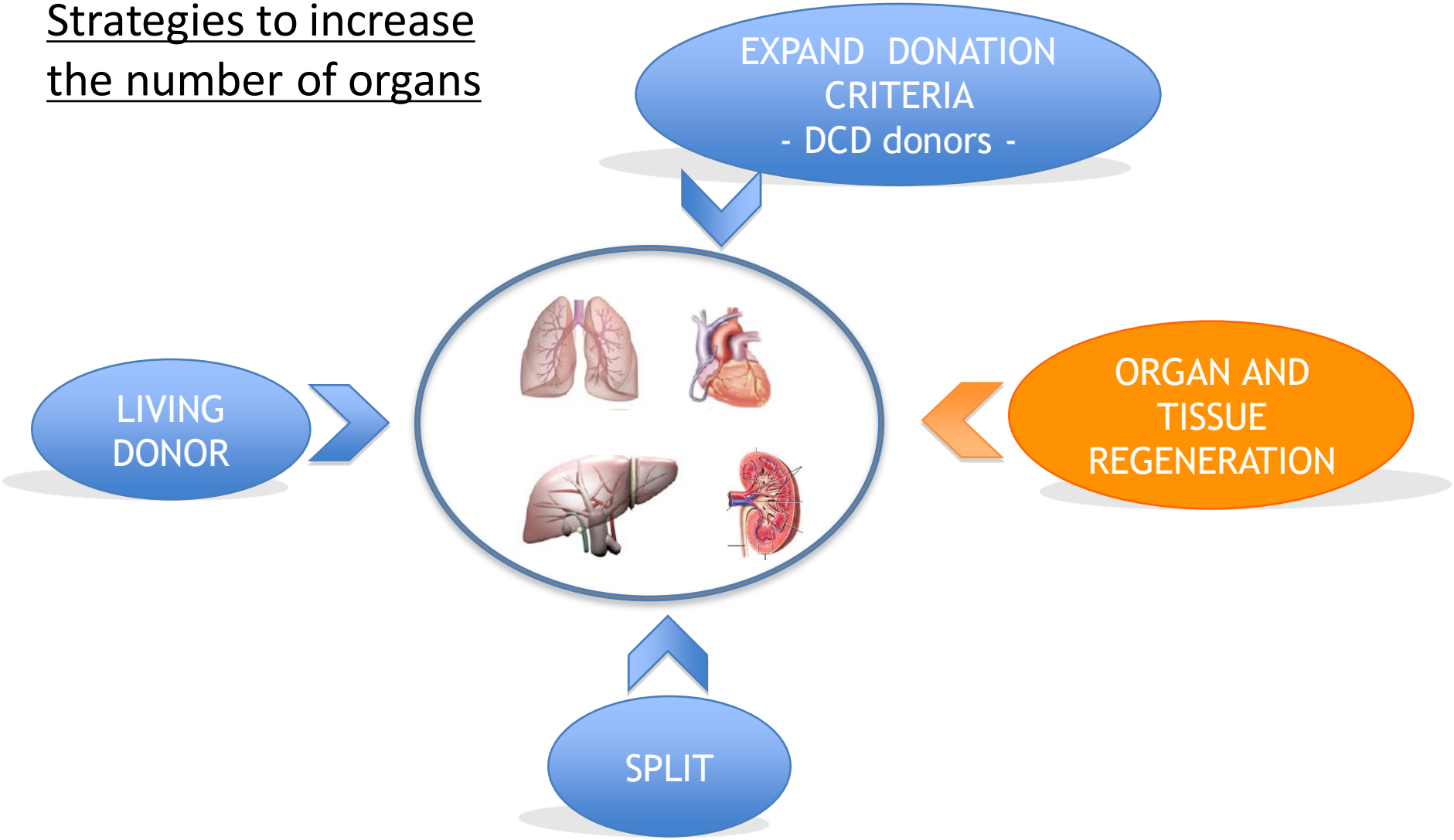
Cellular repair of damaged organs

Repopulating scaffoldings in kidney and liver

Vall d'Hebron Barcelona Hospital Campus

Mireia Caralt, MD PhD
Servei Cirurgia HBP i Trasplantaments
March 29, 2017

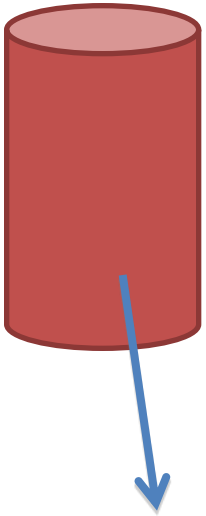
Strategies to increase the number of organs

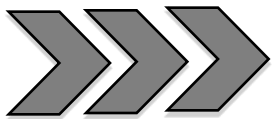


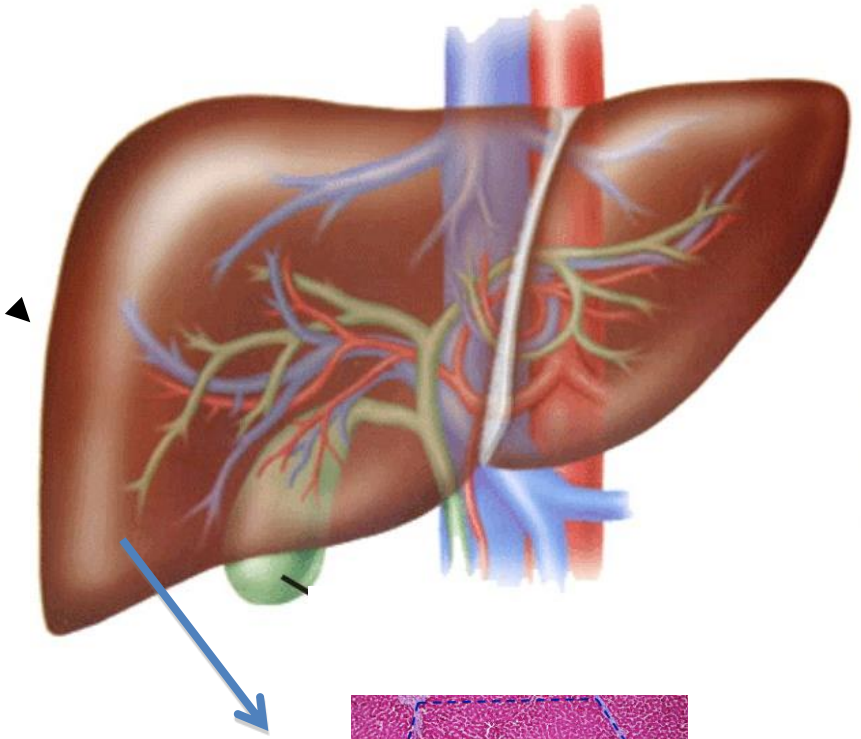
Introduction

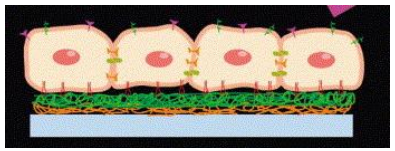
Solid organs: structural complexity!!

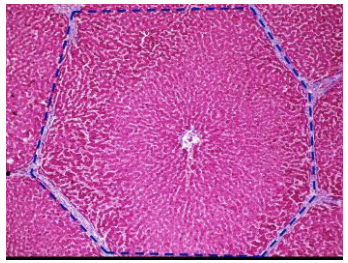
Traquea
Urine Bladder







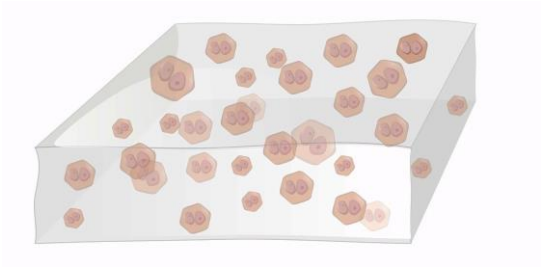




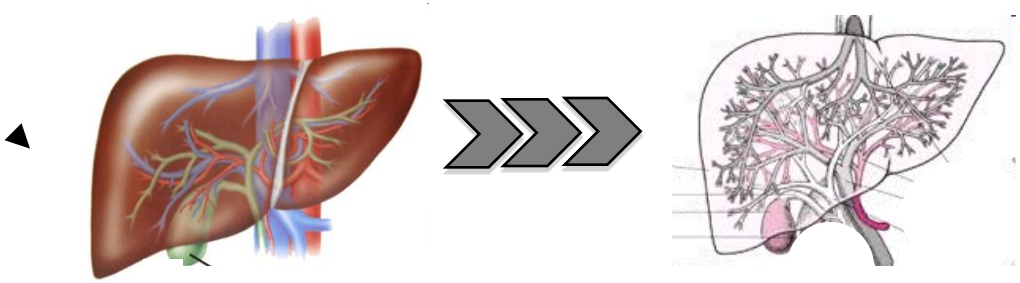
Introduction

Ideal scaffold : biocompatibility, biodegradability, porosity, structural support

Artificial Scaffold



Natural Scaffold



Organ

Extracellular Matrix

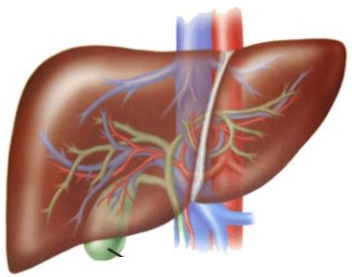
- Difficult control of size, microarchitecture and interconnectivity of pores

- Physical, chemical and molecular stimuli that enable cell engraftment

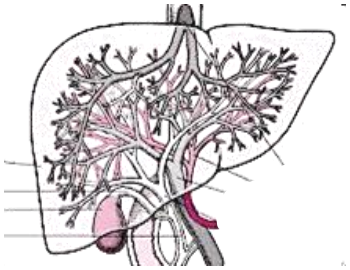
- Preservation of vascular network

- Low inmunogenicity

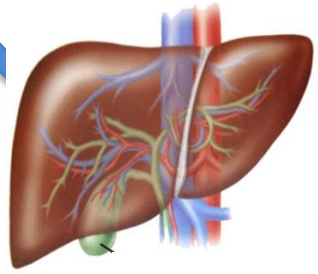
Introduction



Decellularization



Recellularization
Maturation



Transplantation

Introduction

Table 1. Studies in the literature

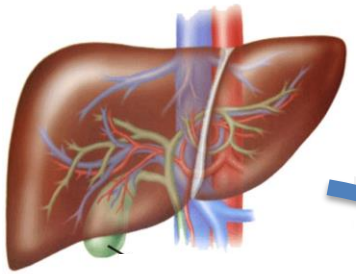
Author	Type Cells	Infusion Method	Via	Number Cells	Flow Rate	Time
Baptista 2010	hUVEC hFLC MS1	Continuous	PV IVC, PV, IVC+PV	$30x10^6$ $70 x10^6$ $100 x10^6$	3ml/min → 0.5ml/min 5ml/min	7d 3d
Uygun 2010	Rat MH Endothelial cells	Multistep	PV	$200 x10^6$	15ml/min	5d 5d
Soto 2011	Rat MH	Direct injection Continuous Multistep	PV	$10–50 x10^6$	2ml/min	7d
Yagi 2013	Pig MH	Multistep	PV	$100 x10^6$	4ml/min	7d

Table 2. Culture media used by different authors

Author	Media
Baptista 2010[1]	RPMI 1640, FBS, dexamethasone, penicillin-streptomycin, prolactin, glucagon, niacinamide, lipoic acid, triiodothyronine, hEGF, hHDL, hHGF, hGH, insulin, transferrin
Uygun 2010[3]	William's E, FBS, insulin, EGF, glucagon, hydrocortisone, penicillin-streptomycin
Soto 2011[63]	EMEM, EGF, HGF, dexamethasone, insulin, human transferring, selenous acid supplement, penicillin-streptomycin
Yagi 2013[63]	DMEM, EGF, hidrocortisone, insulin, glucagon, penicillin-streptomycin

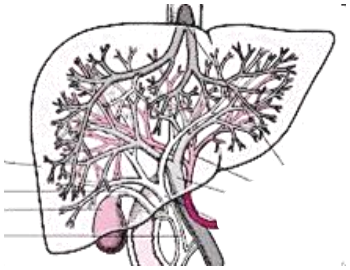
Caralt M. Organogenesis 2014;10(2):250

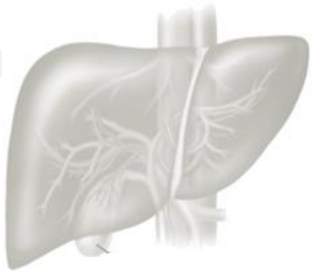
Decellularization



Decellularization

Transplantation





Recellularization
Maturation

Decellularization

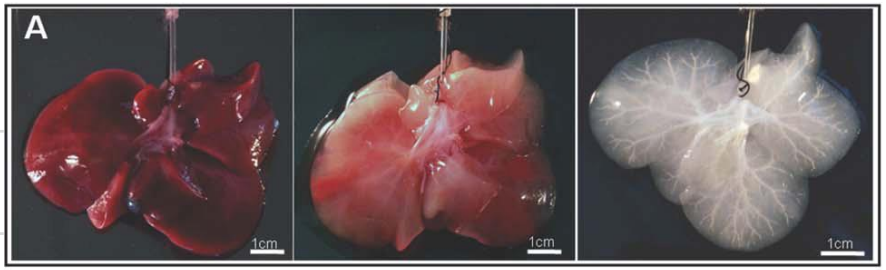
Different decellularization "recipes" in liver

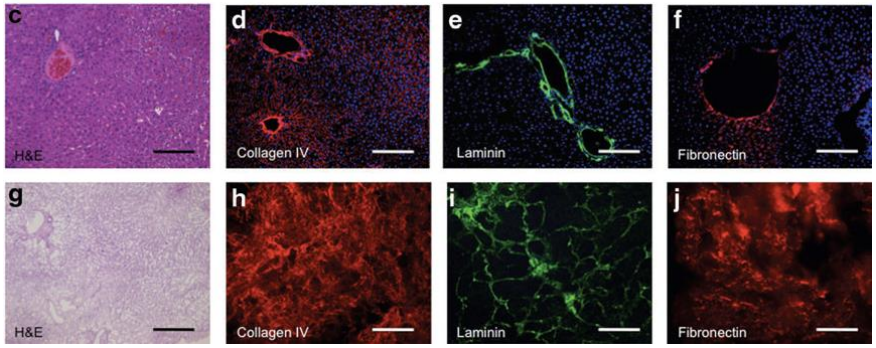
Uygun	rat	SDS Triton X-100	0.01% 24h, 0.1% 24h, 1% 24h 1% 30min	1ml/min
Baptista	rat	Triton X-100 + NH4 OH	1%+3%, 3h	5ml/min
Soto	rat	Trypsin + EGTA Triton X-100 + EGTA	0.02%+0.05%, 2h 3%+0.05%, 24h	8ml/min
Bao	rat	Adenosine SDS	10mM 1% 4h, 0.5% 4h, 0.25% 4h	25ml/min
Yagi	pig	SDS Triton X-100	0.01% 24h, 0.1% 24h, 1% 48h 1% 30min	30ml/min
Ko	pig	Triton X-100 + NH4 OH	1%+3%, 3h	0.5ml/min

Decellularization

Different decellularization "recipes" in liver

Uygun	rat	SDS Triton X-100
Baptista	rat	Triton X-100 + NH4 OH
Soto	rat	Trypsin + EGTA Triton X-100 + EGTA
Bao	rat	Adenosine SDS
Yagi	pig	SDS Triton X-100
Ko	pig	Triton X-100 + NH4 OH





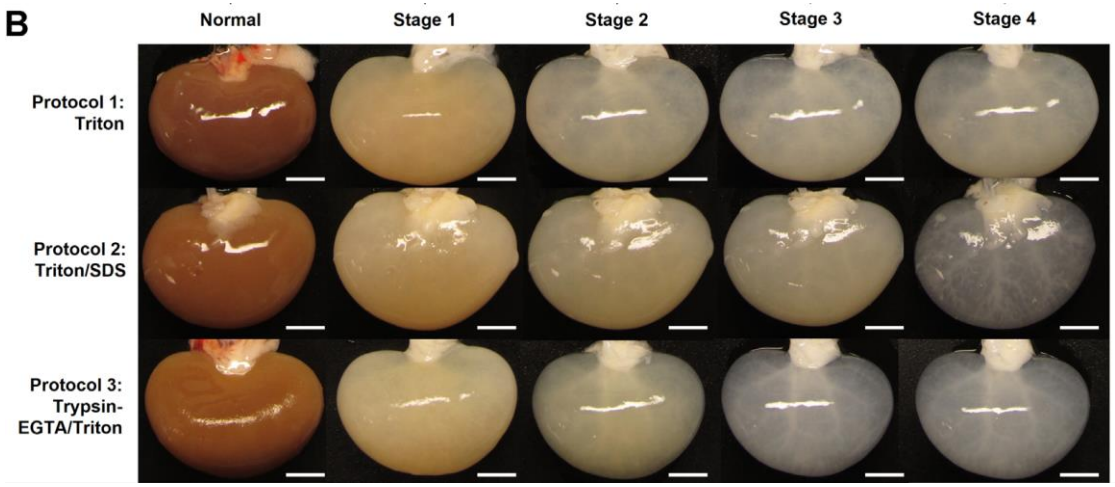
Decellularization

… best decellularization protocol in kidneys

A

	1 h, 40 min.	3 h, 20 min.	16 h, 40 min.	3 h, 20 min.	1 h, 40 min.
Protocol 1: Triton	dH_2O 500 mL	1% Triton X-100 1000 mL	1% Triton X-100 1000 mL	1% Triton X-100 1000 mL	dH_2O 500 mL
Protocol 2: Triton/SDS	dH_2O 500 mL	1% Triton X-100 1000 mL	1% Triton X-100 1000 mL	0.1% SDS 1000 mL	dH_2O 500 mL
Protocol 3: Trypsin-EGTA/Triton	dH_2O 500 mL	2 h: 0.02% trypsin-0.05% EGTA	1% Triton X-100 1000 mL	1 % Triton X-100 1000 mL	dH_2O 500 mL

B



Caralt et al. Am J Transplant 2015; 15:64-75

Decellularization

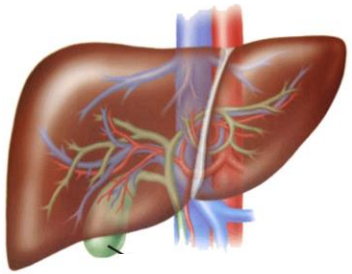
Table 1: Summary evaluation of decellularization protocols

Selection criteria	Protocol 1: Triton	Protocol 2: Triton/SDS	Protocol 3: Trypsin-EGTA/Triton
Macroscopic appearance (transparency of gross tissue is the goal)	–	+	+
Microscopic appearance (H&E, SEM, cell removal, maintenance of architecture is goal)	–	+	+/–
DNA reduction (goal: >95%)	–	+	–
Nuclear basophilia score (goal score is 1.0–2.0)	– Glomeruli: 2.1 ± 0.4; Tubules: 1.9 ± 0.9; Vessels: 3.4 ± 0.8	+ Glomeruli: 1.0 ± 0.0; Tubules: 1.0 ± 0.0; Vessels: 1.0 ± 0.0	+ Glomeruli: 1.1 ± 0.4; Tubules: 1.1 ± 0.4; Vessels: 1.3 ± 0.8
Architectural score (goal score is 4.0–5.0)	+ Glomeruli: 4.3 ± 0.5; Tubules: 4.4 ± 0.8; Vessels: 5.0 ± 0.0	+ Glomeruli: 4.0 ± 0.6; Tubules: 4.2 ± 0.8; Vessels: 5.0 ± 0.0	+ Glomeruli: 3.9 ± 0.7; Tubules: 4.1 ± 0.4; Vessels: 5.0 ± 0.0
ECM (collagen, laminin) retention (goal is retention of proteins)	+	+/–	+/–
Growth factor (bFGF, VEGF) retention (goal: >30% retention)	+	+	–

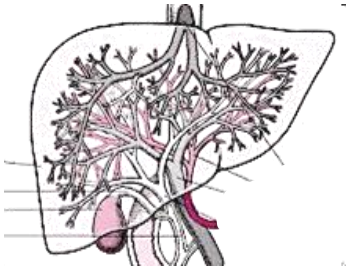
bFGF, basic fibroblast growth factor; ECM, extracellular matrix; H&E, hematoxylin & eosin; SEM, scanning electron microscopy; SDS, sodium dodecyl sulfate; VEGF, vascular endothelial growth factor.
For each criterion, protocols were evaluated and assigned one of three values: good (+), fair (+/–) or poor (–) at reaching a target goal. Each protocol was evaluated independently of the other two protocols, and is compared to normal kidneys.

Caralt et al. Am J Transplant 2015; 15:64-75

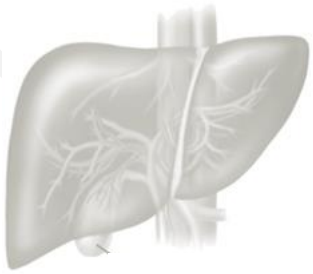
Recellularization



Decellularization



Recellularization
Maturation



Transplantation

Recellularization

Organ reengineering through development of a transplantable recellularized liver graft using decellularized liver matrix

Basak E Uygun[1]

NATURE MEDICINE VOLUME 16
NUMBER 7 | JULY 2010

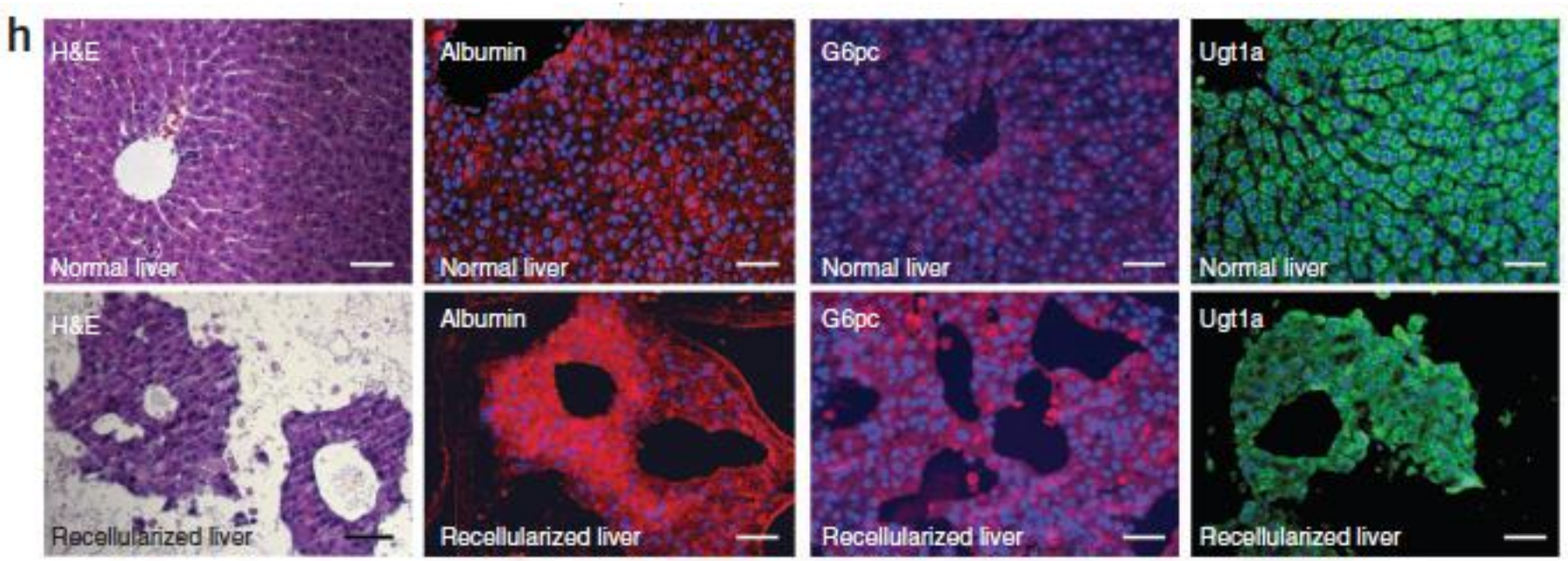
Primary rat hepatocytes

The Use of Whole Organ Decellularization for the Generation of a Vascularized Liver Organoid

BAPTISTA ET AL. HEPATOLOGY, Vol. 53, No. 2, 2011

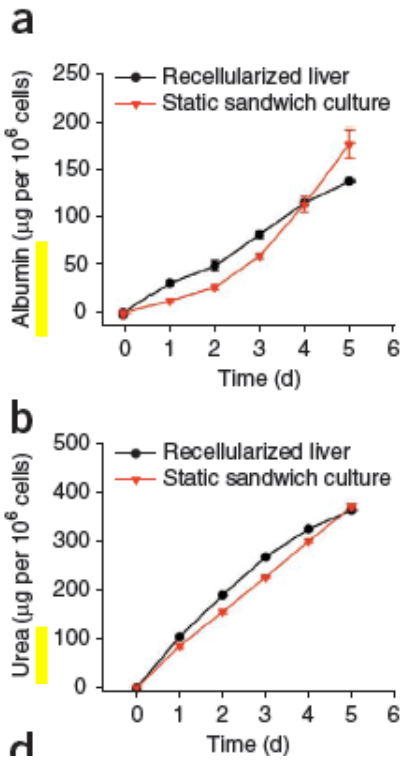
hFLCs + hUVEC

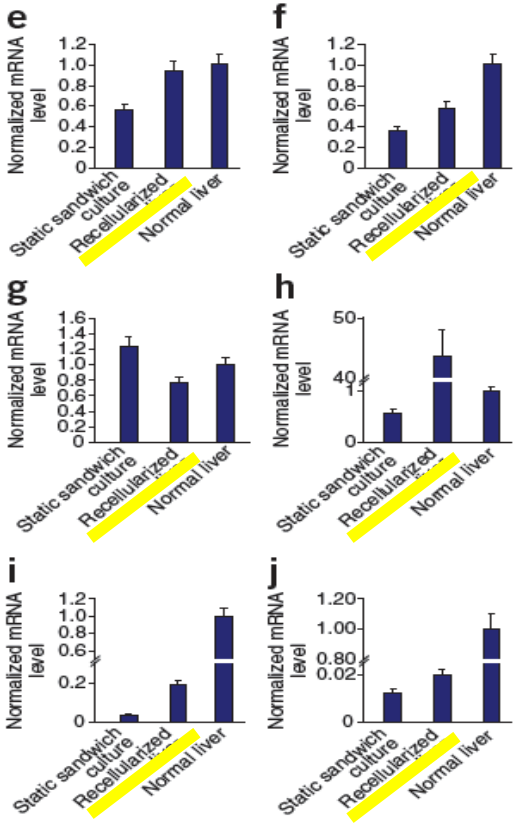
Recellularization



Uygun. Nature 2010;16(7):814-821

Recellularization





(e–j) Normalized gene expression of *Cyp2c11* (e), *Gstm2* (f), *Ugt1a1* (g), *Cyp1a1* (h), *Adh1* (i) and *Cyp3a18* (j). All error bars represent s.e.m. ($n = 3$).

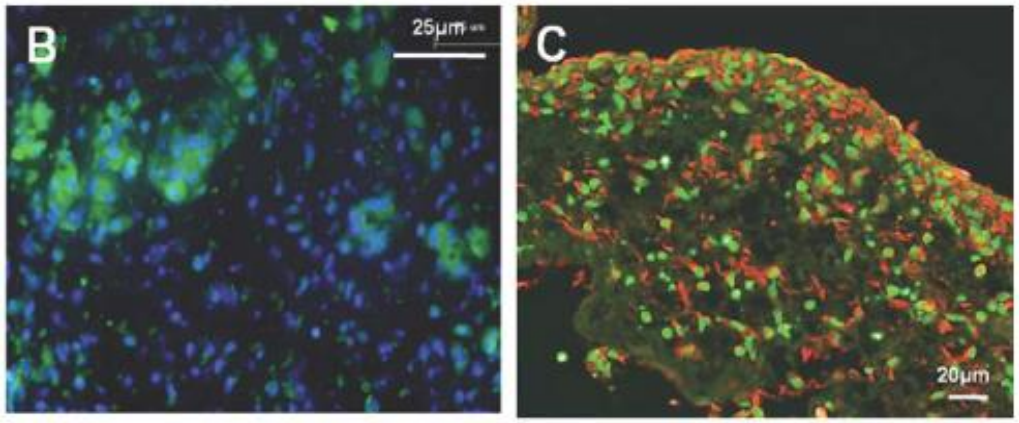
20% albumin production of *in vivo* levels

30% drug metabolism gene expression of *in vivo* levels

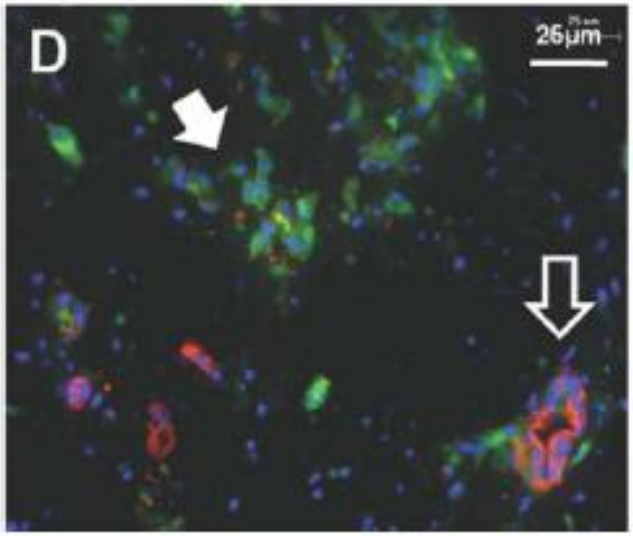
Uygun. Nature 2010;16(7):814-821

Recellularization

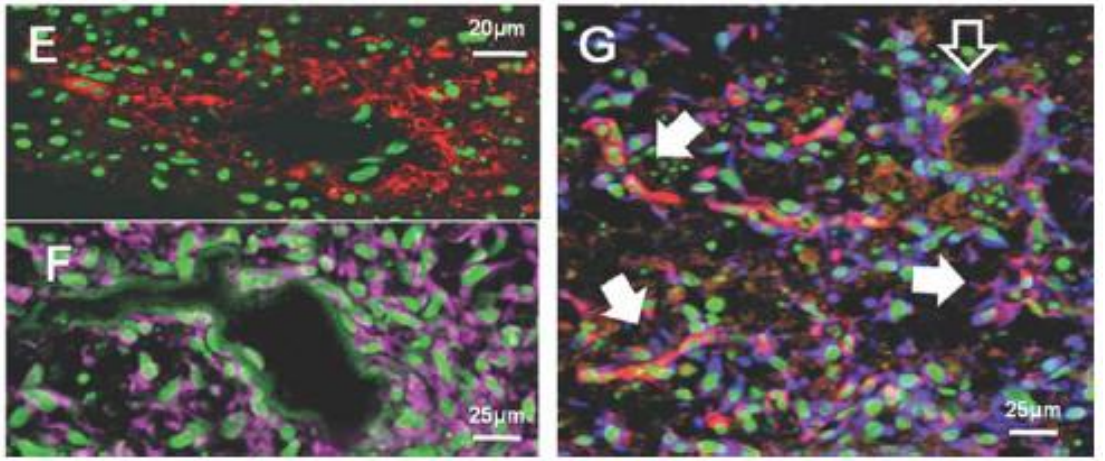
hFLC
hUVEC



p450 CYP2A (green) and p450 CYP3A (red)



CK 19 (red) and albumin (green)



Von Willebrand (red) and endothelial oxid nitric sinthetase (pink)

Baptista et al. Hepatology 2010 Nov;12:604-617

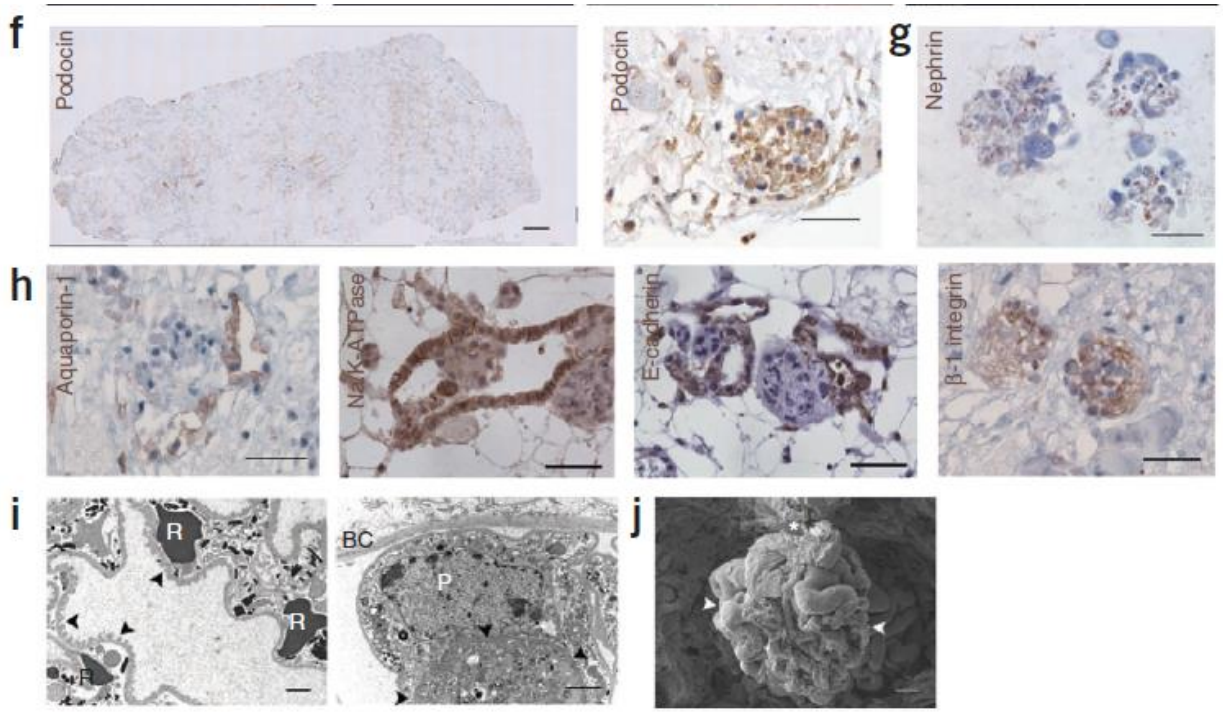
Recellularization

Regeneration and experimental orthotopic transplantation of a bioengineered kidney

Jeremy J Song[1,2], Jacques P Guyette[1,2], Sarah E Gilpin[1,2], Gabriel Gonzalez[1,2], Joseph P Vacanti[1–3] & Harald C Ott[1,2,4]

VOLUME 19 | NUMBER 5 | MAY 2013 NATURE MEDICINE

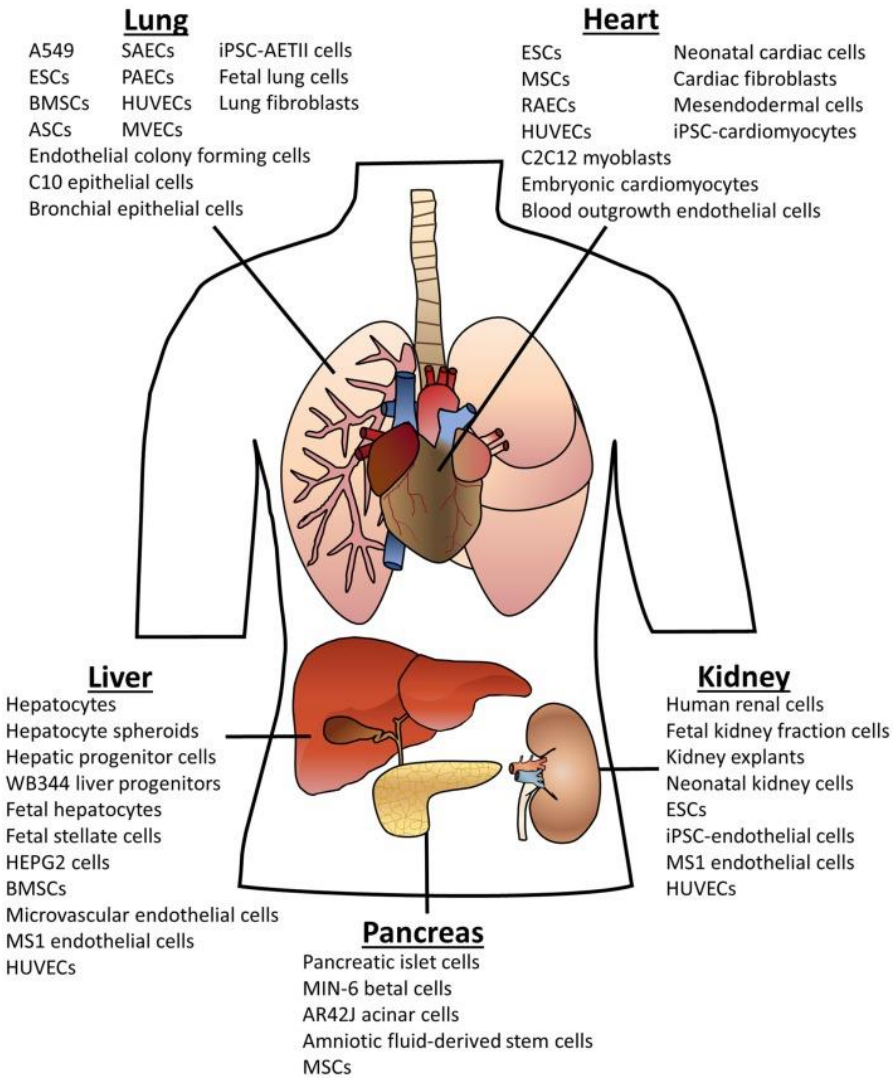
Neonatal kidney cells



Recellularization

Function		Dcell	ReCell	Native
Albumin retention		46.9%	23.3%	89.9%
Glucose absortion		2.8%	47.38%	91.7%
Filtration	Volume	4.9 ±0.1ul/min	1.2 ±0.1ul/min	3.2 ±0.1ul/min
	Glucose	249±62.9mg/dl	160±20mg/dl	29±8.5mg/dl
	Albumin	26.85±4.03g/dl	4.67±2.51g/dl	0.6±0.4g/dl
	Urea	18 ±42.2mg/dl	28.3±8.5mg/dl	617.3±34.8mg/dl
	Creatinine	0.5±0.3mg/dl	1.3±0.2mg/dl	24.6±5.8mg/dl
	ClCr		0.01±0.002ml/min	0.36±0.09ml/min
	Urea Excretion		0.003±0.002mg/min	0.19±0.01mg/min

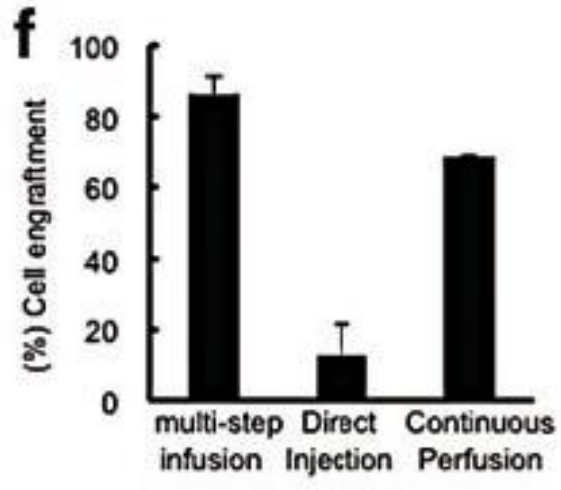
Recellularization

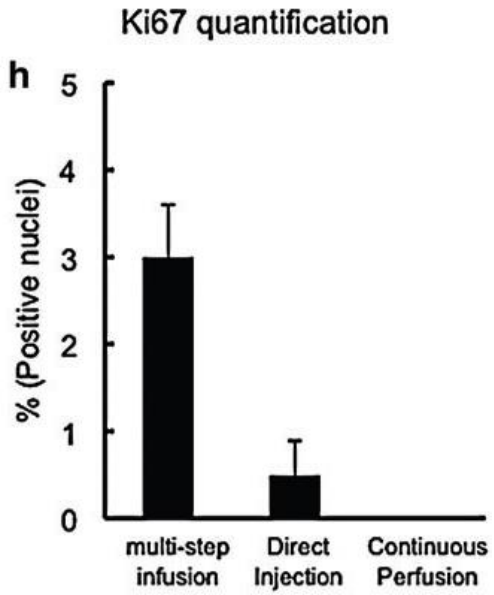


Scarritt et al. Front Bioeng Biotechnol 2015;3(43):1-17

Recellularization

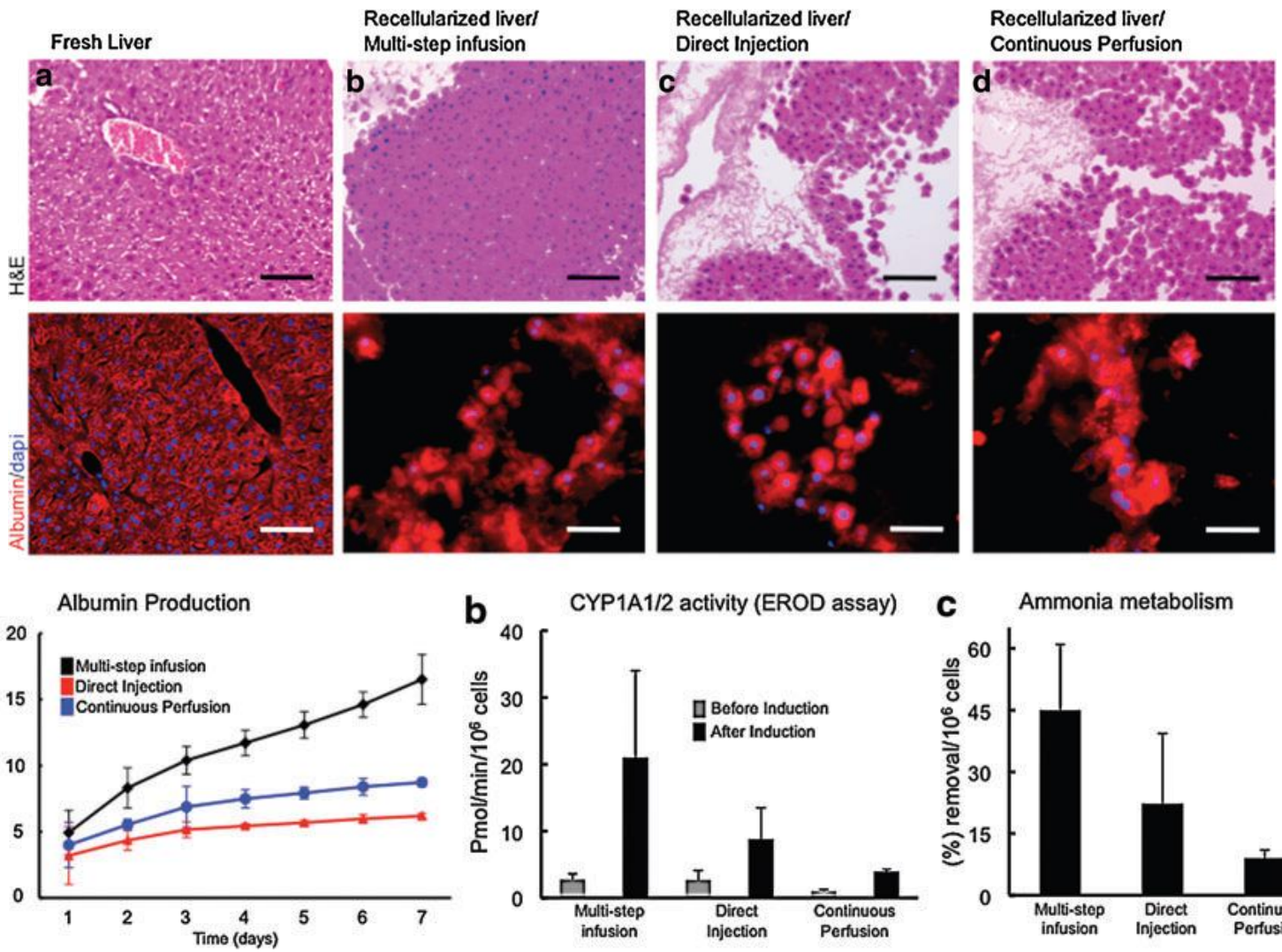
Direct parenchymal injection	Continuous perfusion	Multistep infusion
Direct injections into different lobes	Cells suspendended in culture media	Cells through Porta Vein 4 steps at 10-15min interval





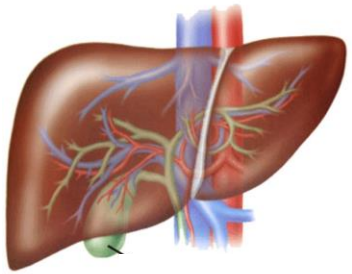
Soto A. Tissue Eng Part C Methods 2011;17(6):677-86

Recellularization

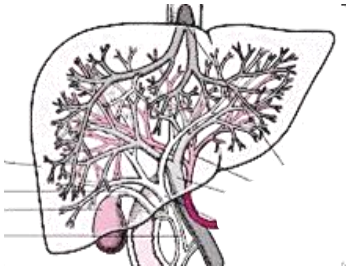


Soto A. Tissue Eng Part C Methods 2011;17(6):677-86

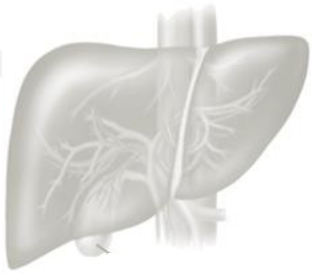
Transplantation



Decellularization

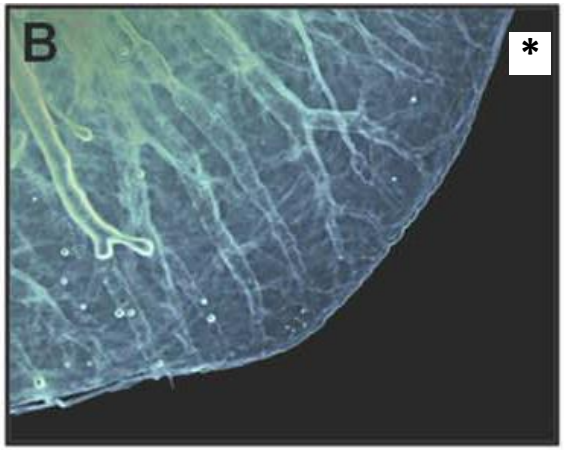


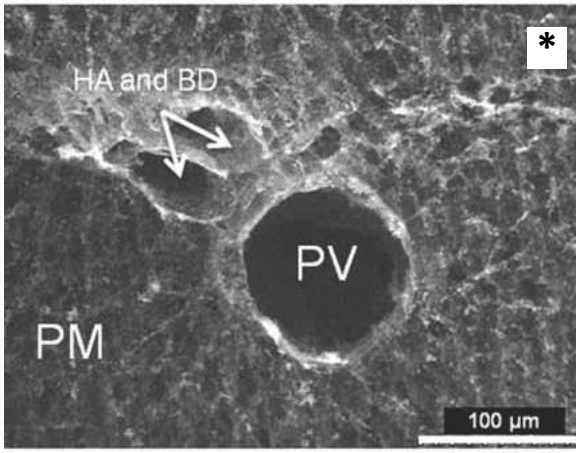
Recellularization
Maturation

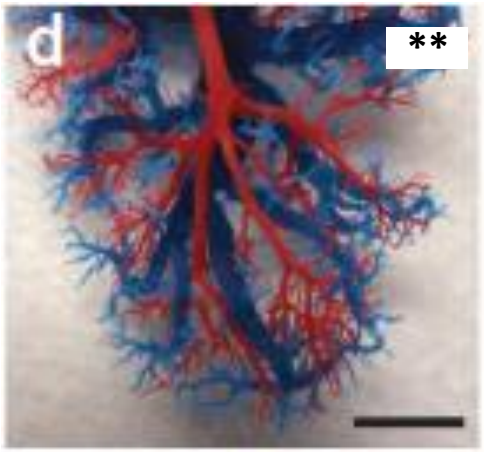


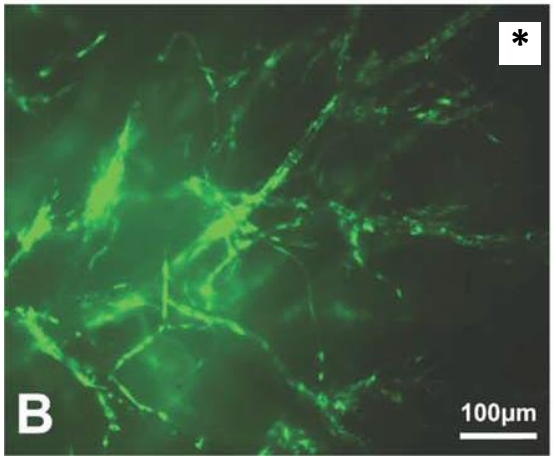
Transplantation

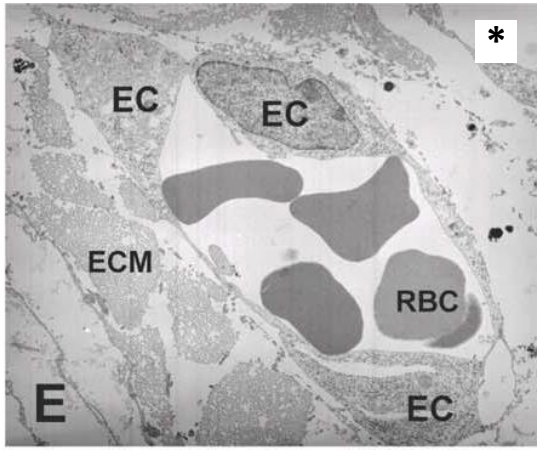
Transplantation

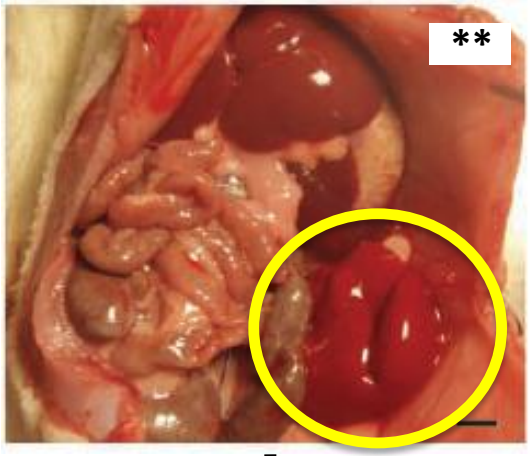






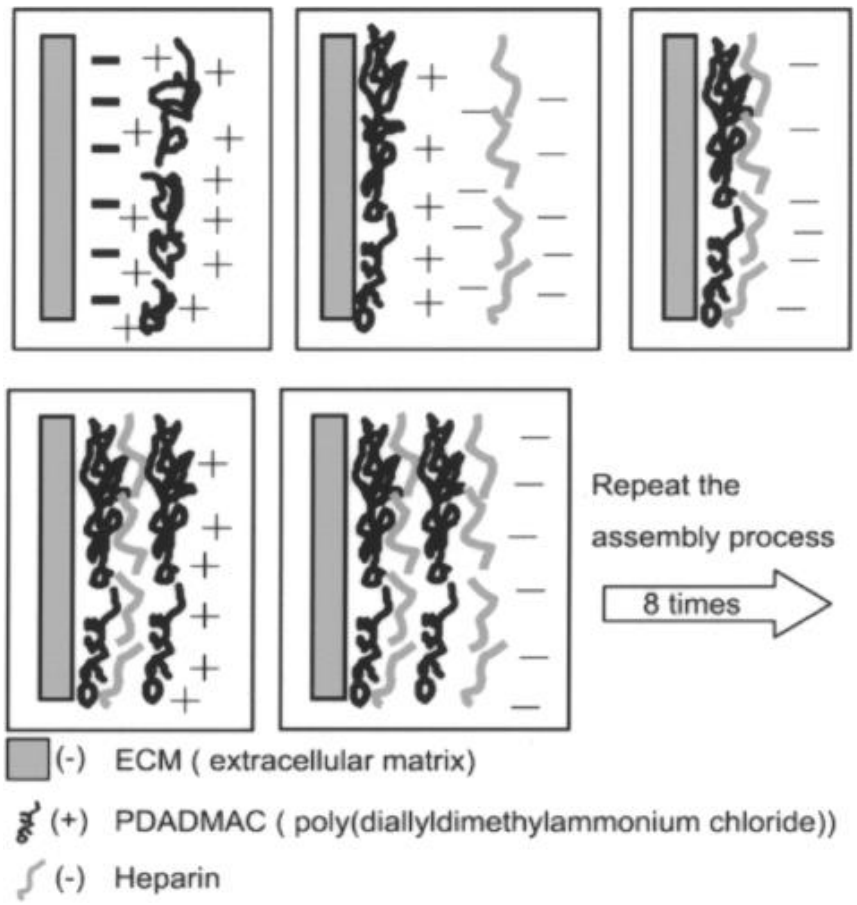






8h viability
thrombosis

* Baptista et al. Hepatology 2010;12:604-617
** Uygun. Nature 2010;16(7):814-821



LbL self-assembly technique

Polyelectrolyte polydiallydimethylammoniuum chloride (PDADMAC) positively charged

Heparine negatively charged

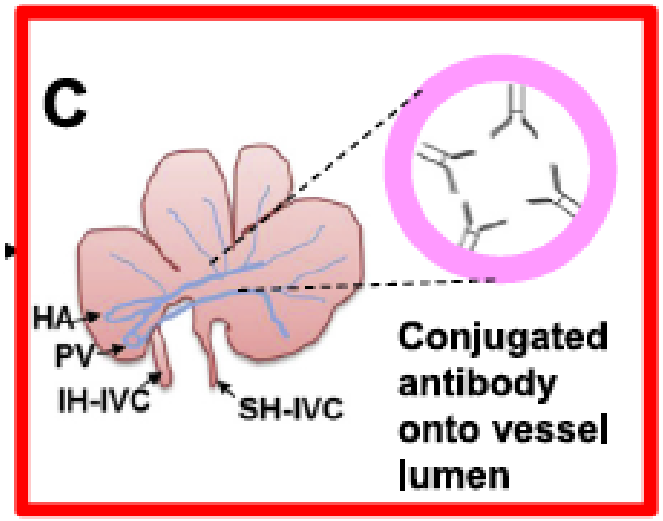
Thromboresistant after 3h of blood perfusion

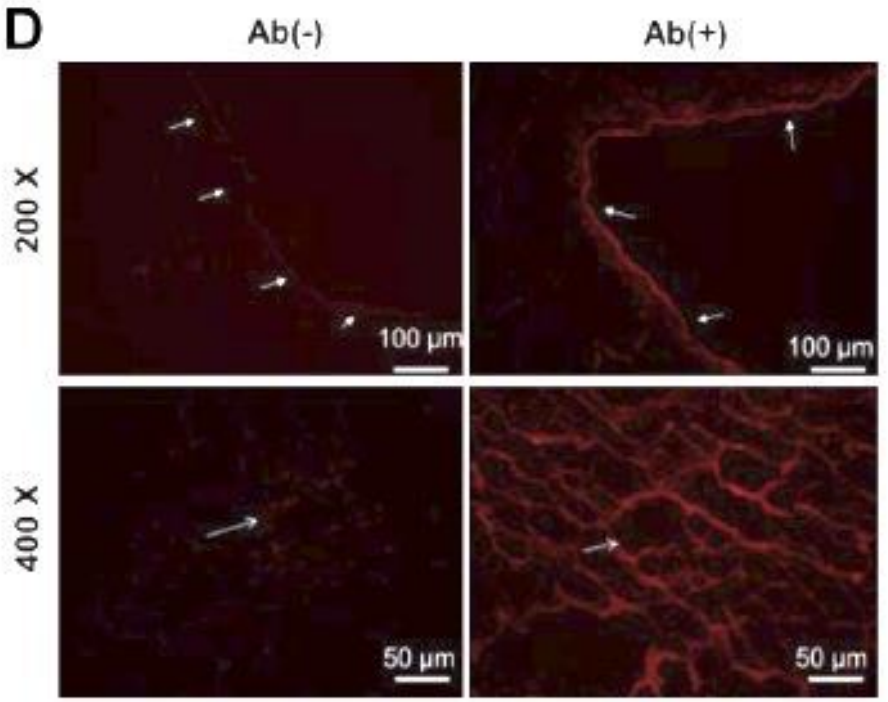
After 72h, hepatocytes maintained normal morphology

Bao et al. Cell Transplantation 2011;20:753-766

Antibody conjugation method

1. Anti endothelial cell antibodies to stabilize seeded cells on the vessel walls.
   Rat anti-mouse CD31 antibody was conjuated to the acellular liver scaffold

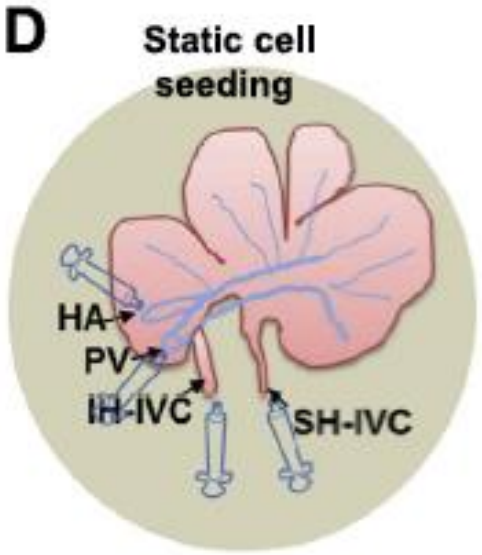


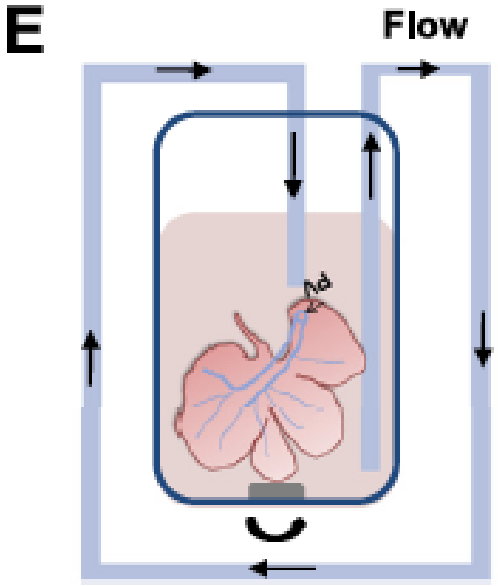


Ko et al. Biomaterials 2015;40:72-79

Antibody conjugation method

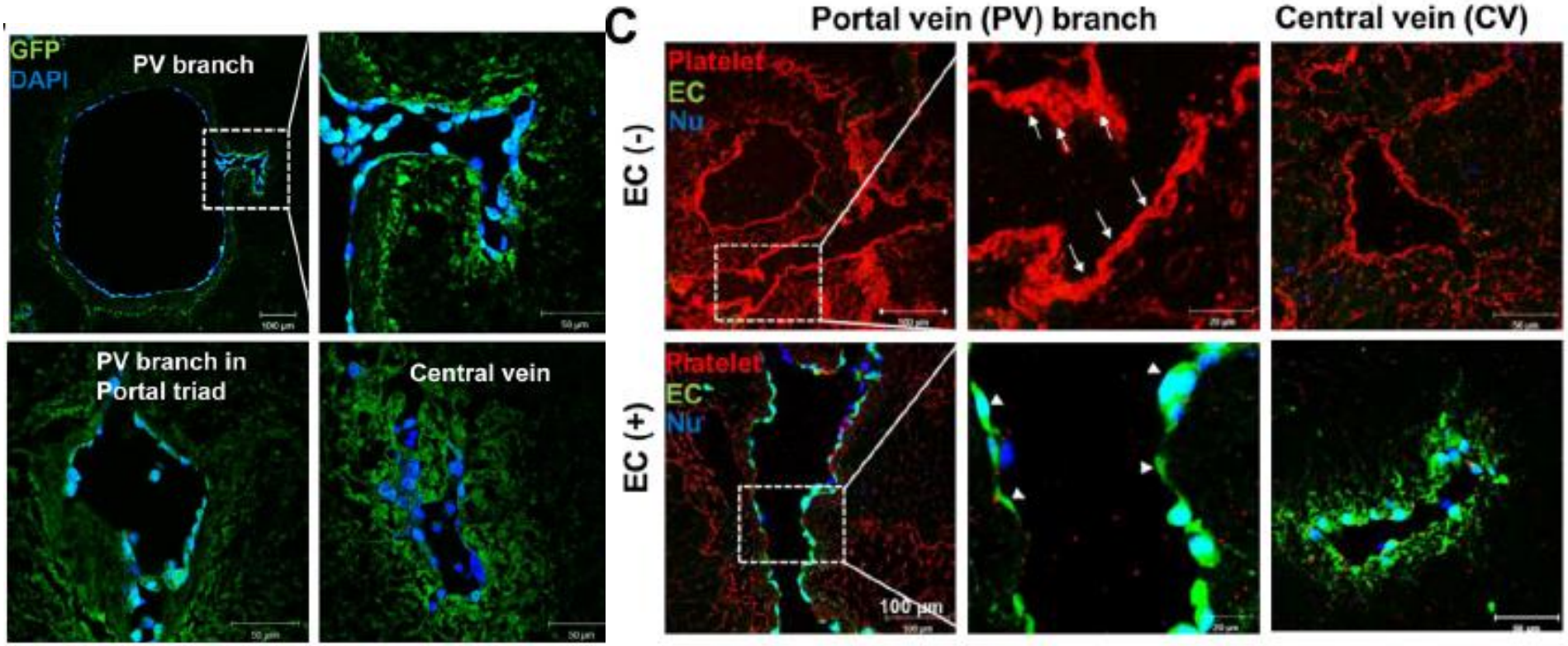
1. Anti endothelial cell antibodies to stabilize seeded cells on the vessel walls: Rat anti-mouse CD31 antibody was conjuated to the acellular liver scaffold
2. ReEndothelization with endothelial cells (MS1) expressing GFP protein





Ko et al. Biomaterials 2015;40:72-79

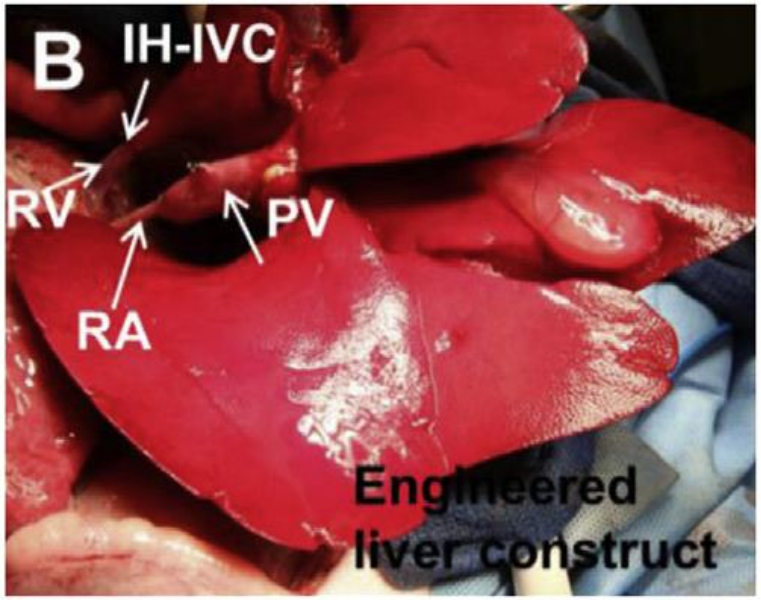
Re-endothelialization characterization



Ko et al. Biomaterials 2015;40:72-79

Antibody conjugation method

1. Anti endothelial cell antibodies to stabilize seeded cells on the vessel walls: Rat anti-mouse CD31 antibody was conjuated to the acellular liver scaffold
2. ReEndothelization with endothelial cells (MS1) expressing GFP protein
3. Implantation of engineered porcine liver construct

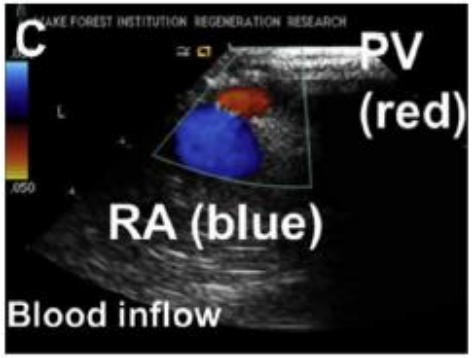


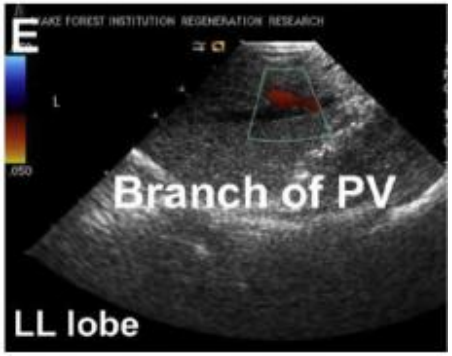
Heterotopically implantation into pig
Left renal artery – Portal vein
Left renal vein - Inferior vena cava

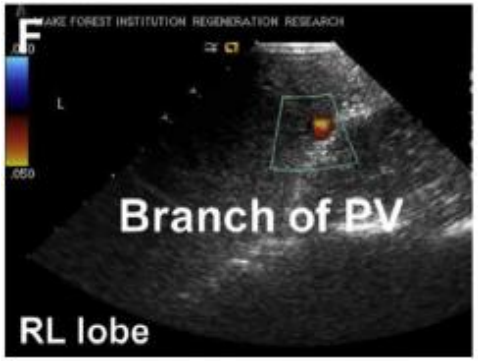
Ko et al. Biomaterials 2015;40:72-79

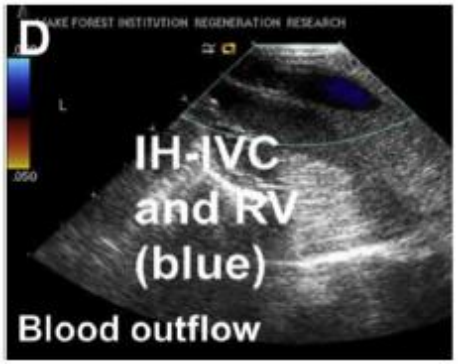
In vivo functional testing

Intraoperative

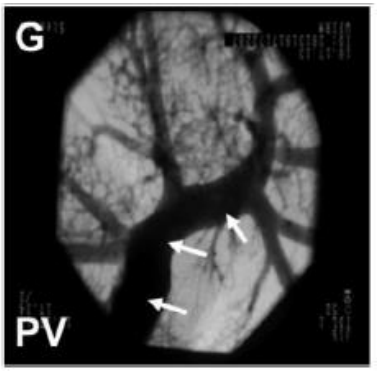


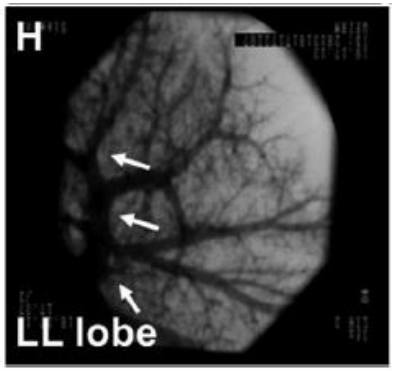


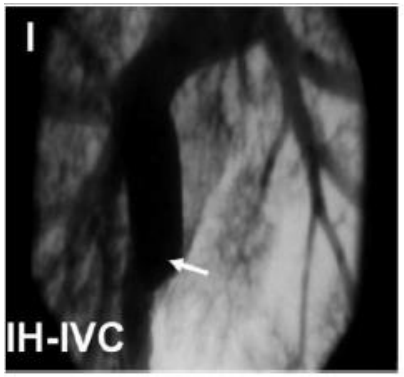




4h after implantation

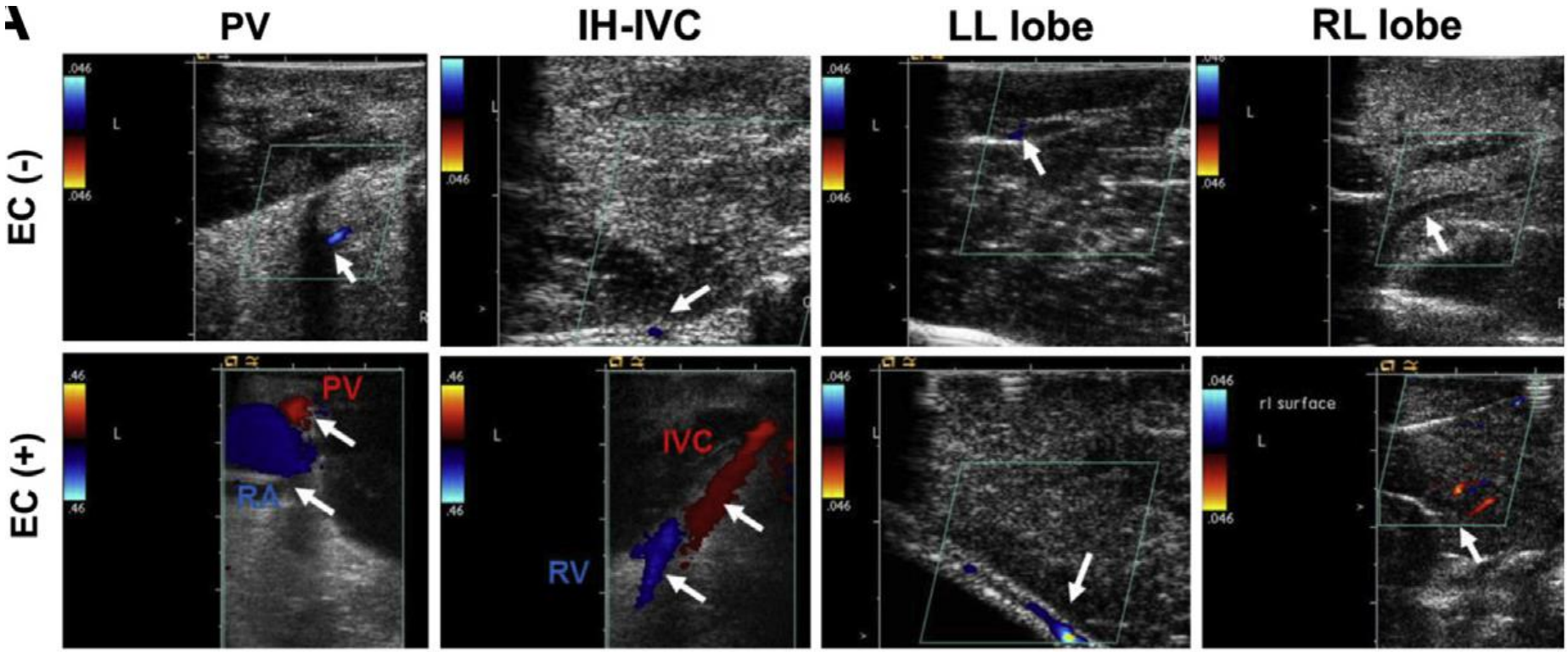






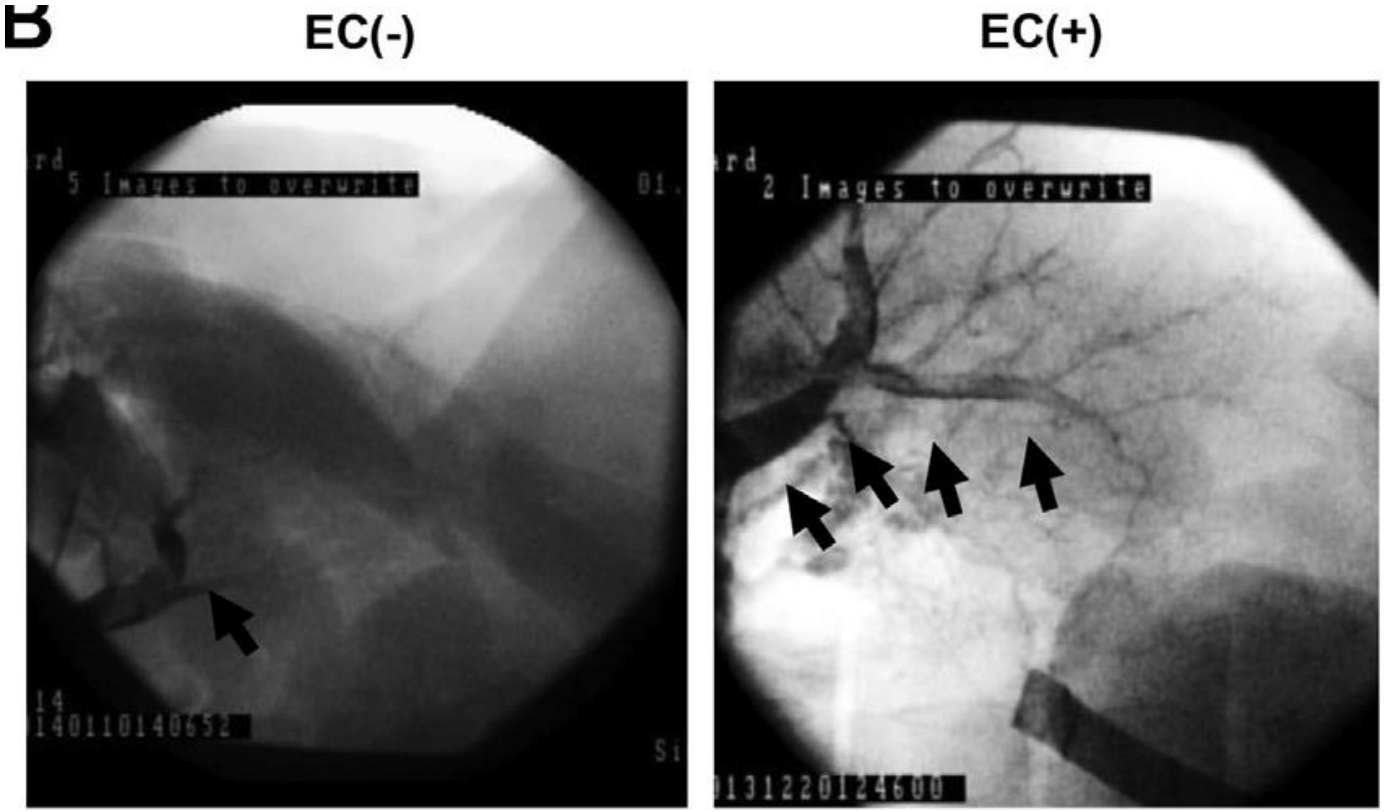
Ko et al. Biomaterials 2015;40:72-79

In vivo functional testing: POD 1



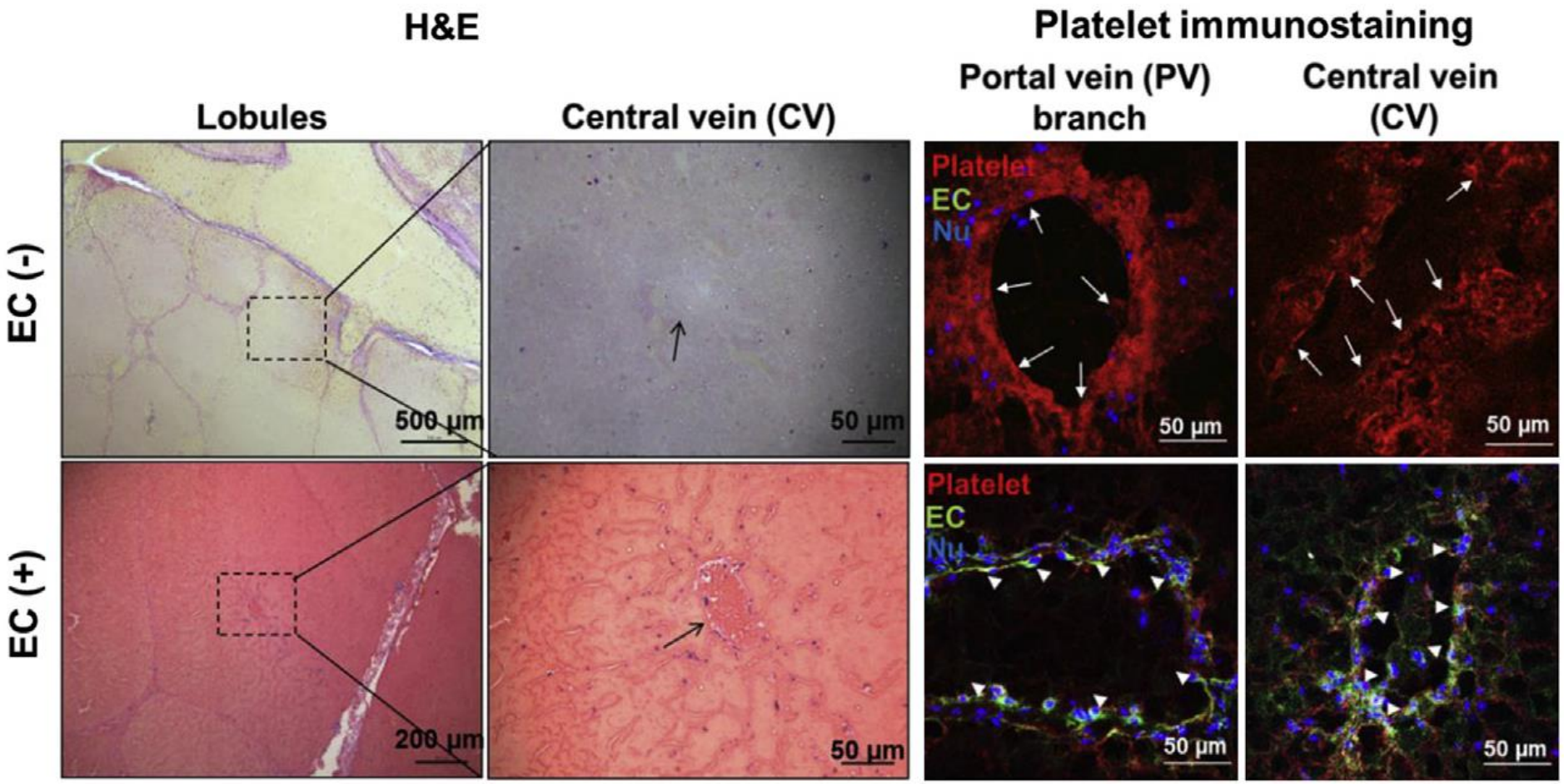
Ko et al. Biomaterials 2015;40:72-79

In vivo functional testing: POD 1



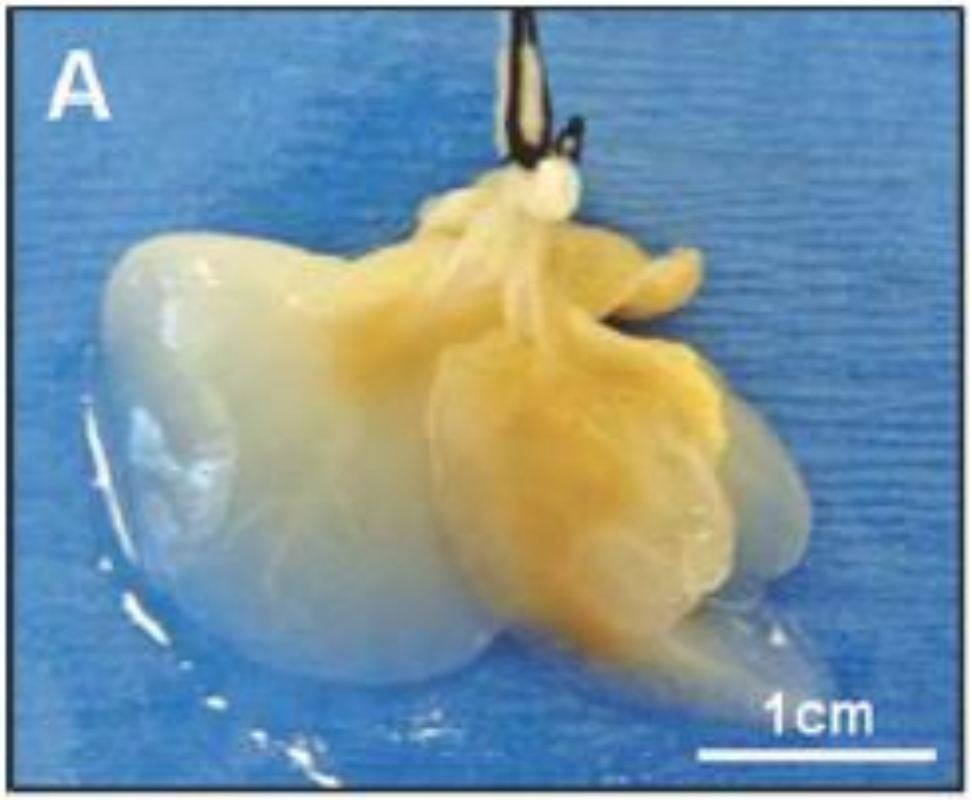
Ko et al. Biomaterials 2015;40:72-79

In vitro functional testing: POD 1



Ko et al. Biomaterials 2015;40:72-79

Main concerns



Baptista et al. Hepatology 2010 Nov;12:604-617

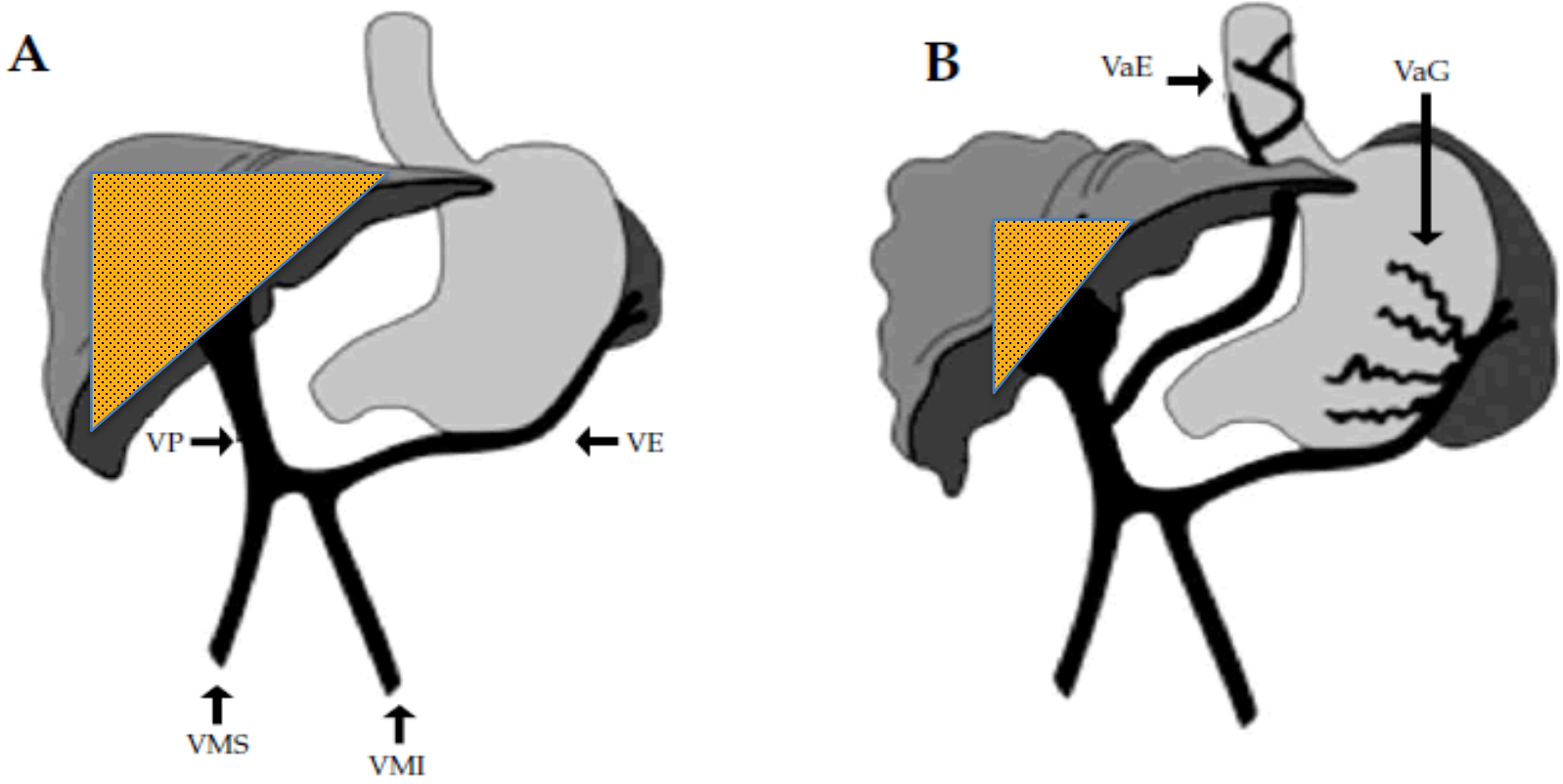
70M human fetal hepatocytes

=

4 livers
(17-21 weeks gestation)

Number is important…

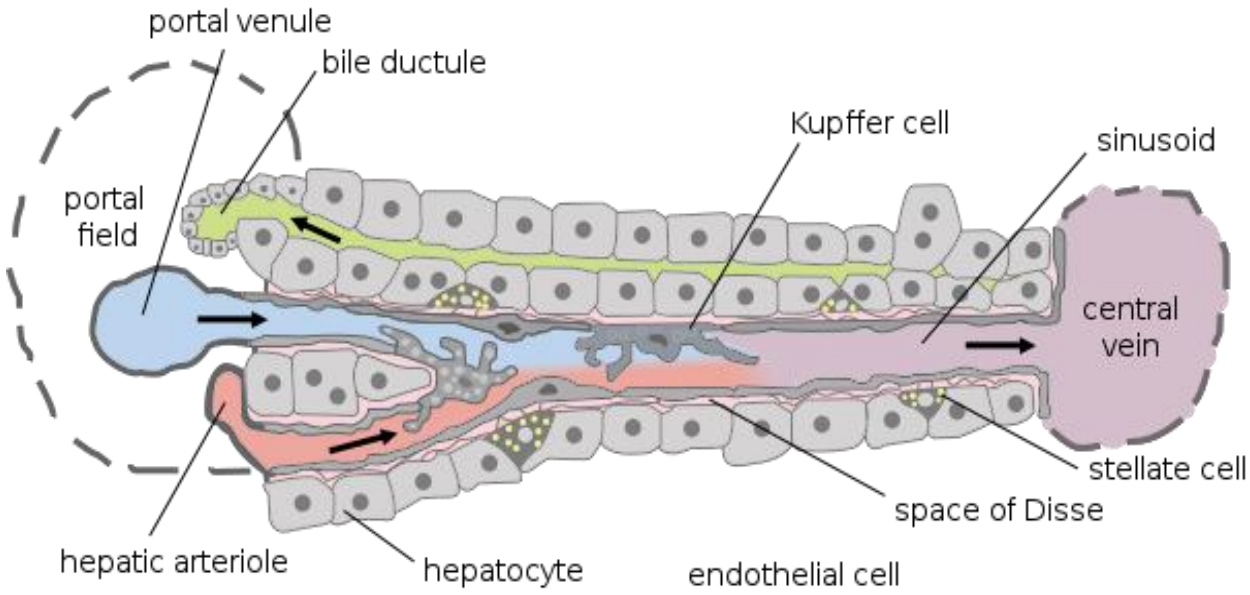
Main concerns



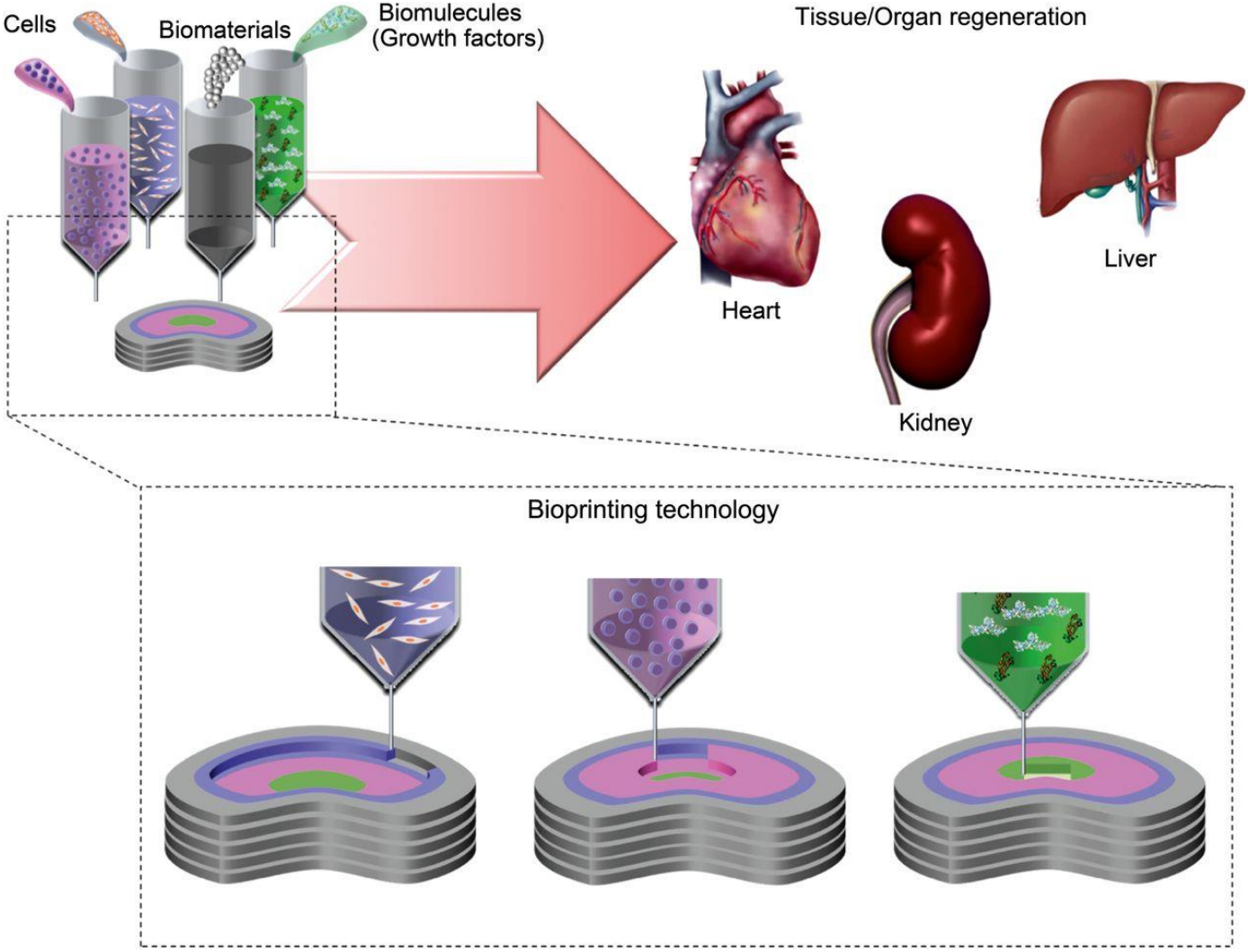
… size is important too

Future directions

- Co-Culture with other non-parenchymal cells. iPSC?
- Cholangiocytes - Biliary tree
- Bioprinting

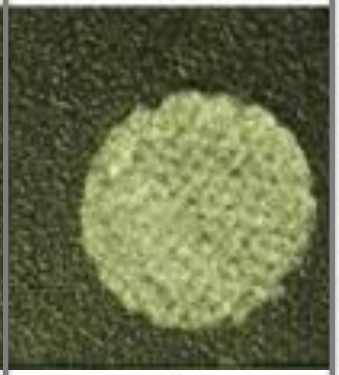


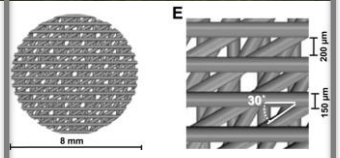
Future directions



Seol et al. Eur J Cardiothorac Surg 2014;46:342-348

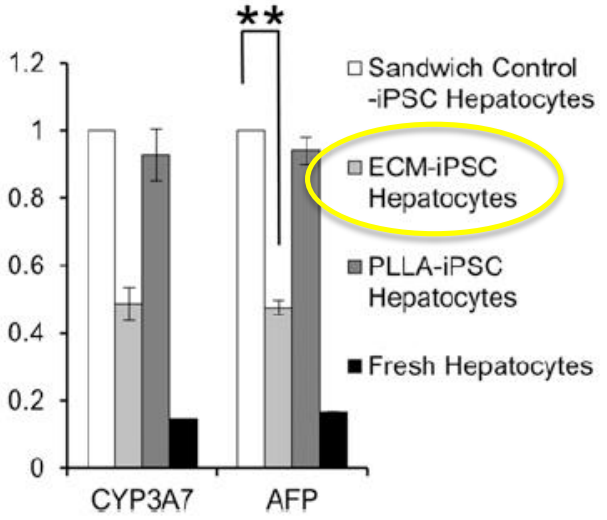
Future directions

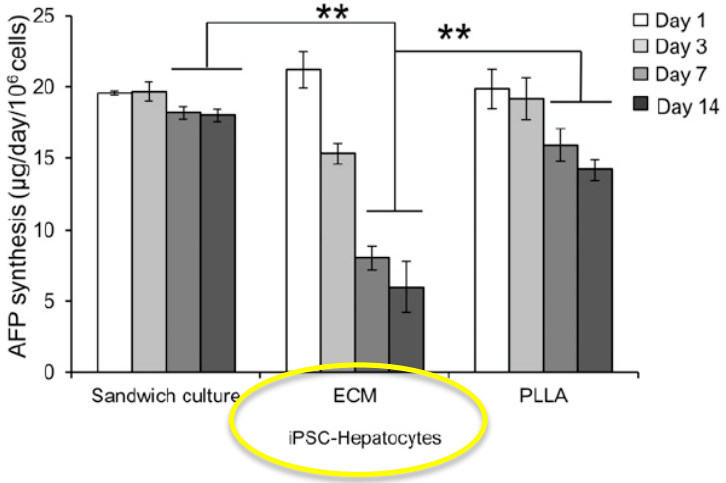
ECM scaffold	PLLA scaffold
	
type I collagen, fibronectin, laminin HGF, bFGF	type I collagen



Biochemical milieu in the ECM leads to the enhanced maturation of iPSC hepatocytes

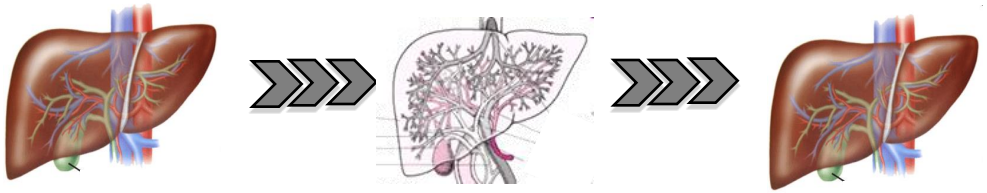
Hepatocyte maturation





Wang et al. Stem Cells Transl Med 2016;5(9):1257-1267

Summary



- Decellularized organs are a good option to obtain scaffolds because architecture and vasculature are well preserved
- Cells are “happy” in the scaffolds

  mature hepatocytes: viable and functional

  immature cells: differentiate into cells present in the liver
- Multistep recellularization
- Endothelization needed for transplantation
- Main concerns: number of cells and size of organoid
- Future directions:

  Coculture with other non-parenchymal cells

  Cholangiocytes and Bile Duct