



***Executive Director's Presentation
Annual General Meeting
30 November 2009***



Mesoblast Capital Overview

Fund Raisings

\$m

IPO @ 50 cents	21.0
<i>Equity Placements:</i>	
Jul-06	17.4
Dec-07	13.4
Apr-09	10.8
Option funds	3.2
Total Funds Raised	\$65.8

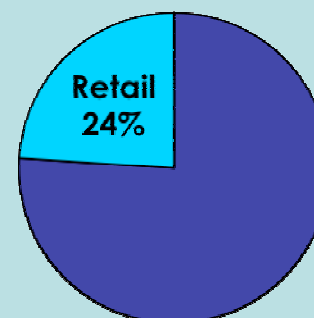


Use of capital:

- Mesoblast Preclinical and Clinical Programs
- Investment in Angioblast, 38.4%

Shares on Issue	138.3m
Current Share Price	\$1.43
Current Market Cap (circa)	\$200m

Mesoblast Ownership



**Institutions &
Sophisticated
Investors
76%**

MSB Financials

Cash Balance at 30 Sep 09*

\$15.8m

Cash Burn (annual)

\$m

G&A & Clinical Support

5.5

Capital Expenditure

0.1

Intellectual Property

0.2

Interest Revenue

(0.6)

Total Net Overheads, IP & Clinical Support

5.2

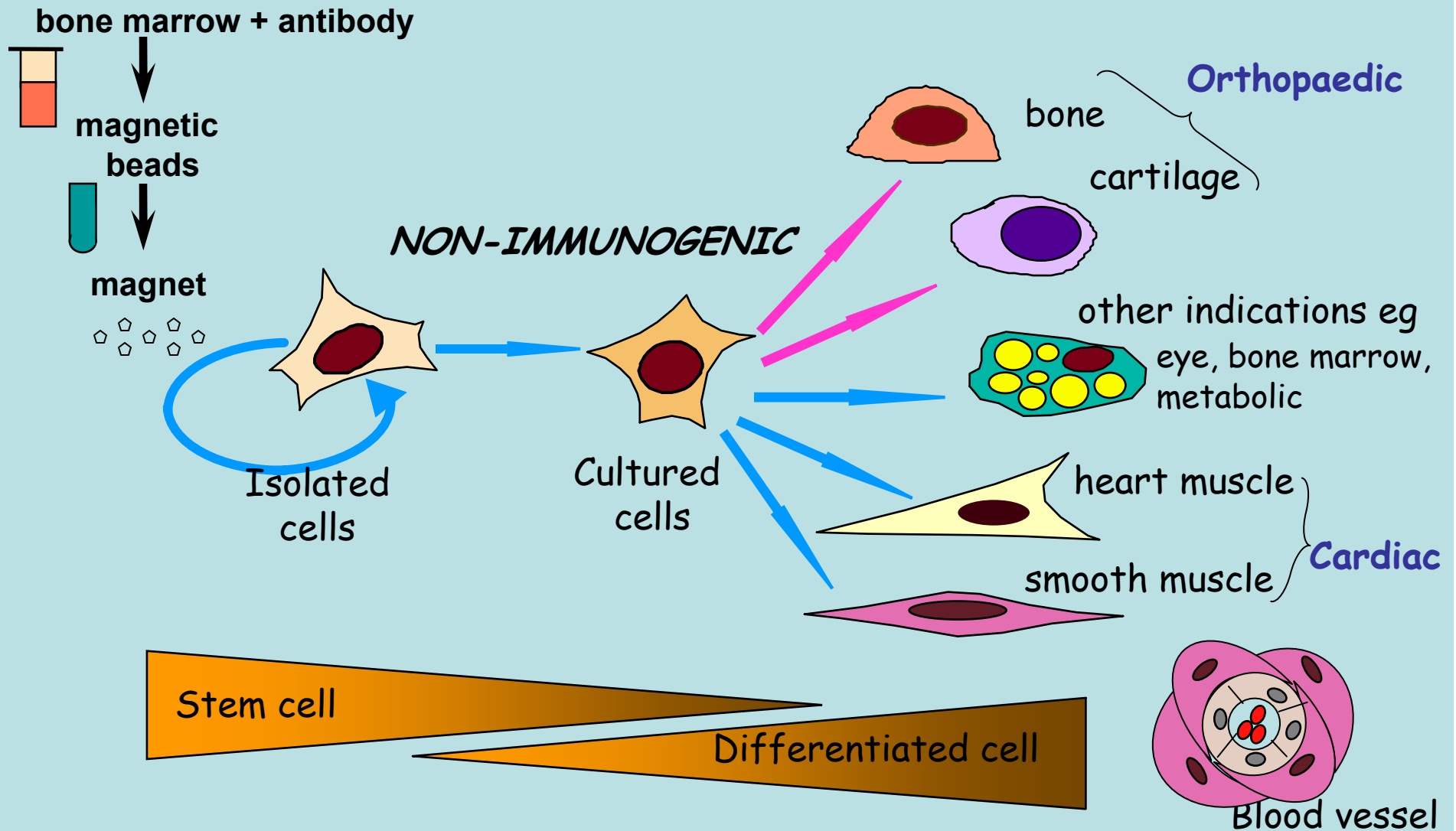
Clinical Development/R&D (external)**

3.0

Total Annual Spend

\$8.2m

Our Cells are Highly Expandable and Non-Immunogenic; Potential for “Off-the-Shelf” Allogeneic Products

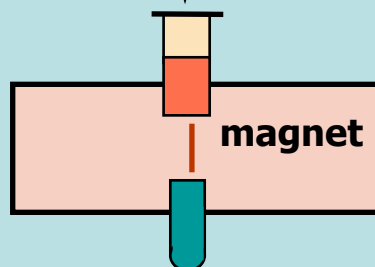


Industrial-Scale Manufacturing Process

Bone Marrow (BM)

BM + stem cell-binding antibody

magnetic beads



fresh,
quiescent

master
cell bank

MPC

final
product



Competitive Advantages:

*Precise identification,
ease of isolation and scale-up*

- purer initial stem cell pool
- homogeneous population
- efficient large-scale expansion
- lower costs of cell culture process
- batch-to-batch consistency
- stringent release criteria
- greater potency of expanded product



Lead Orthopaedic Programs

1. Spinal Fusion

- ***Lumbar fusion: Phase 2 trial (minimally-invasive) ongoing***
- ***Cervical fusion: Phase 2 trial ongoing***

2. Intervertebral disc repair/regeneration

- ***Phase 2 clinical trial to commence 2010***

3. Long bone fracture repair

- ✓ ***Phase 1b trial long bone non-union completed***

4. Knee Osteoarthritis

- ***Phase 2 trial post-knee trauma ongoing***



Lead Non-Orthopaedic Programs, In Association With Angioblast Systems

5. Congestive Heart Failure (CHF)

- ***Phase 2 trial ongoing***

6. Acute Myocardial Infarction (AMI)

- ***Phase 2 trial ongoing***

7. Bone Marrow Transplantation, Expansion of Umbilical Cord Blood

- ***Orphan Drug Designation***
- ***Phase 1/2 trial ongoing***

8. Age-Related Macular Degeneration (AMD) and Diabetic Retinopathy

- ***Phase 2 trial to commence early 2010***



Commercialisation and Revenue Generation Strategies

- 1. Broad-based Partnering of Platform Technology**
e.g. For Pharma Licensing
- 2. Partnering Specific Applications**
e.g. For Medtech Applications
- 3. Taking Individual Applications to Market on Our Own**
e.g. For Orphan Drug Applications
- 4. Product Manufacturing and Supply Agreements**
e.g. Transfer Price on Finished Product



Commercial Strategies For MPC-Based Biologicals

1. Broad-Based Partnering Of Platform Technology e.g. For Pharma Indications

- ***partner(s) for lead programs***
- ***partner(s) for non-core/second generation programs***
- ***extensive resources applied to programs***
- ***enable simultaneous development of multiple core and new applications***
- ***de-risk funding requirements for more costly Phase 3 programs***
- ***de-risk need for sales and marketing force***

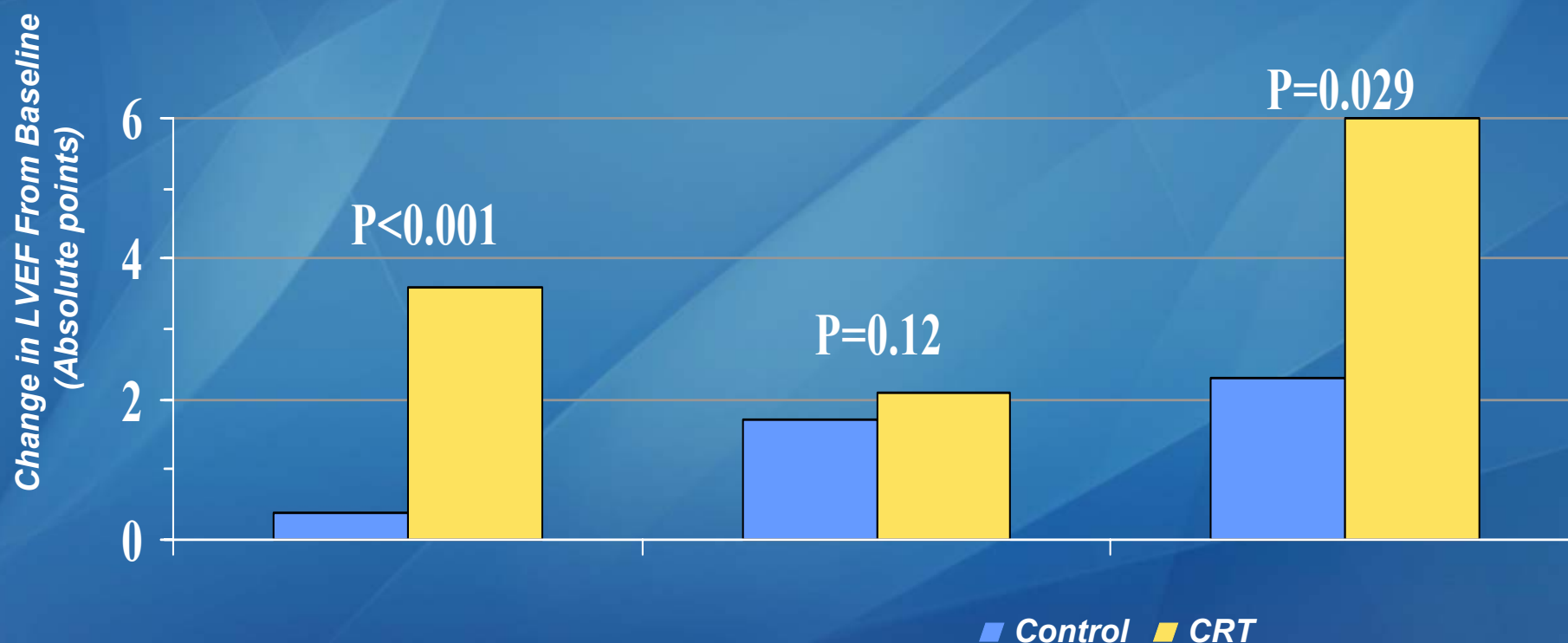


Congestive Heart Failure

- *in United States alone, existing 5.5M plus 0.55M new-onset/year*
- *Class II-IV patients with EF < 35%*
 - (a) have highest mortality at 6 and 12 months*
 - (b) account for 40% of all CHF patients*
- *product effective in this group will generate premium pricing*
- *predicted 45% annual growth rate, peak penetration year 8*

Cardiac Resynchronization Therapy (CRT): A Good Model For Product Success In CHF Patients With EF < 35%.

At 6 Months, CRT Increases EF In Patients With Conduction Defects (15% Of Total CHF Pool)...
4 point difference from controls impacts on mortality!



Data sources: MIRACLE: *Circulation* 2003;107:1985-1990

MIRACLE ICD: *JAMA* 2003;289:2685-2694

Contak CD: *J Am Coll Cardiol* 2003;2003;42:1454-1459

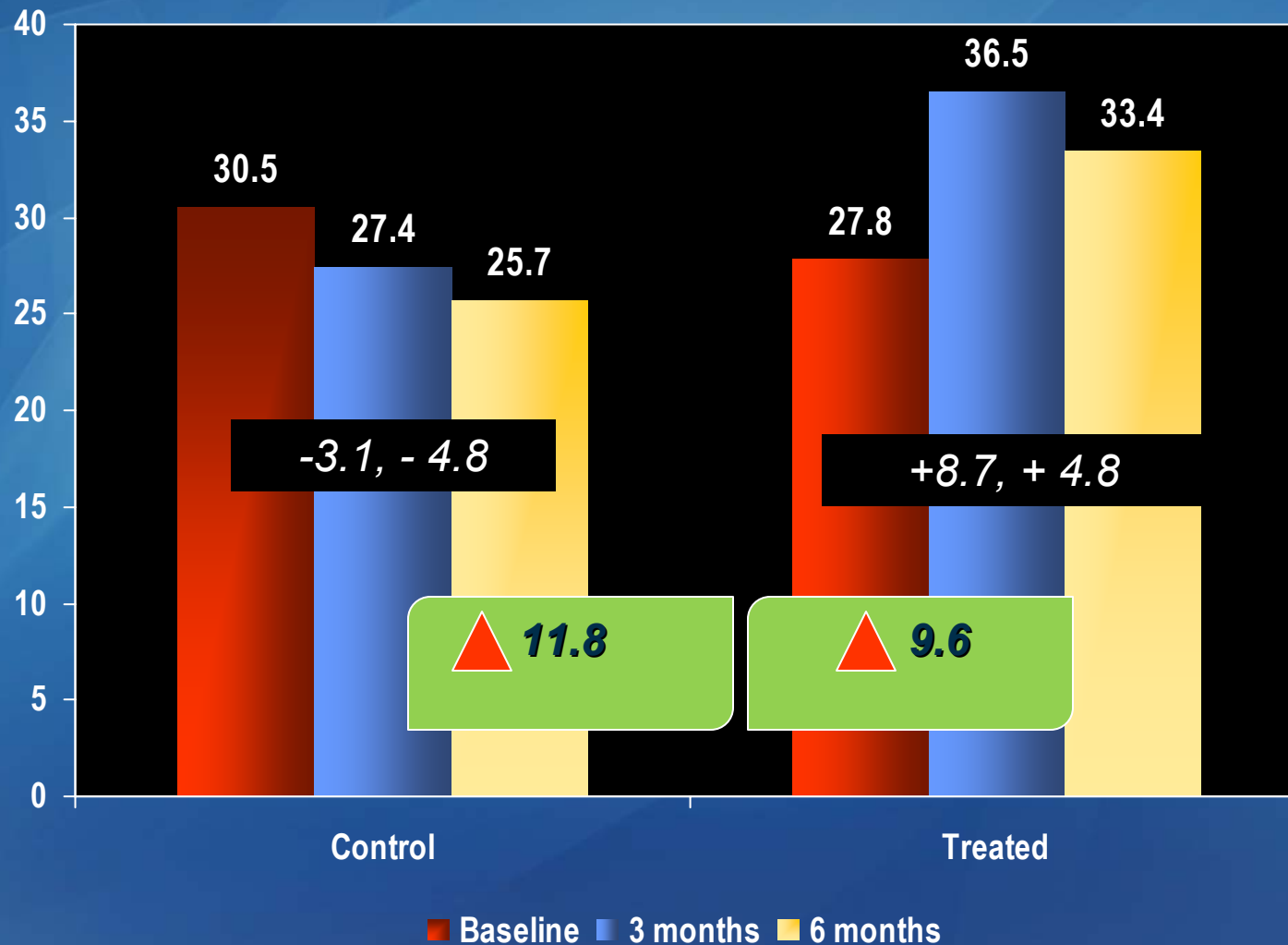


Phase 2a Trial Of Allogeneic MPC In CHF Patients EF<40%

- ***60 patient multi-center, randomized, controlled trial***
- ***class II-IV CHF, EF < 40%, high 6- and 12-month mortality***
- ***three cohorts of 20 patients, each cohort testing progressively increasing allogeneic MPC dose***
- ***cells injected by JNJ NOGA Myostar catheter***
- ***40 patients first two doses completed, no adverse events***
- ***evaluating cardiac functional recovery at 3, 6 and 12 months***

MPC Treatment Results In Sustained 10-Point Increase In EF Relative To Controls...

?Impact On Mortality



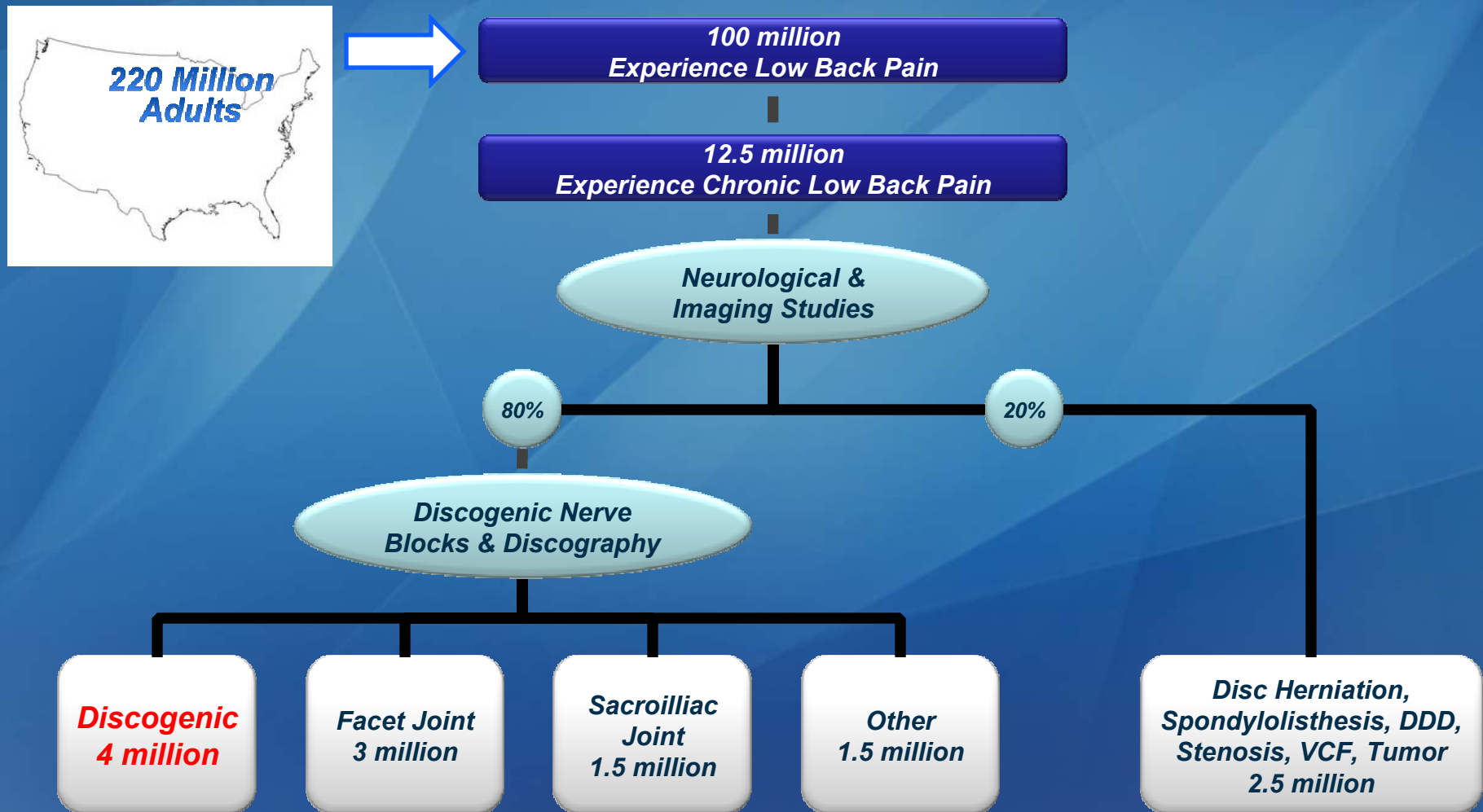


Commercial Strategies For MPC-Based Biologicals

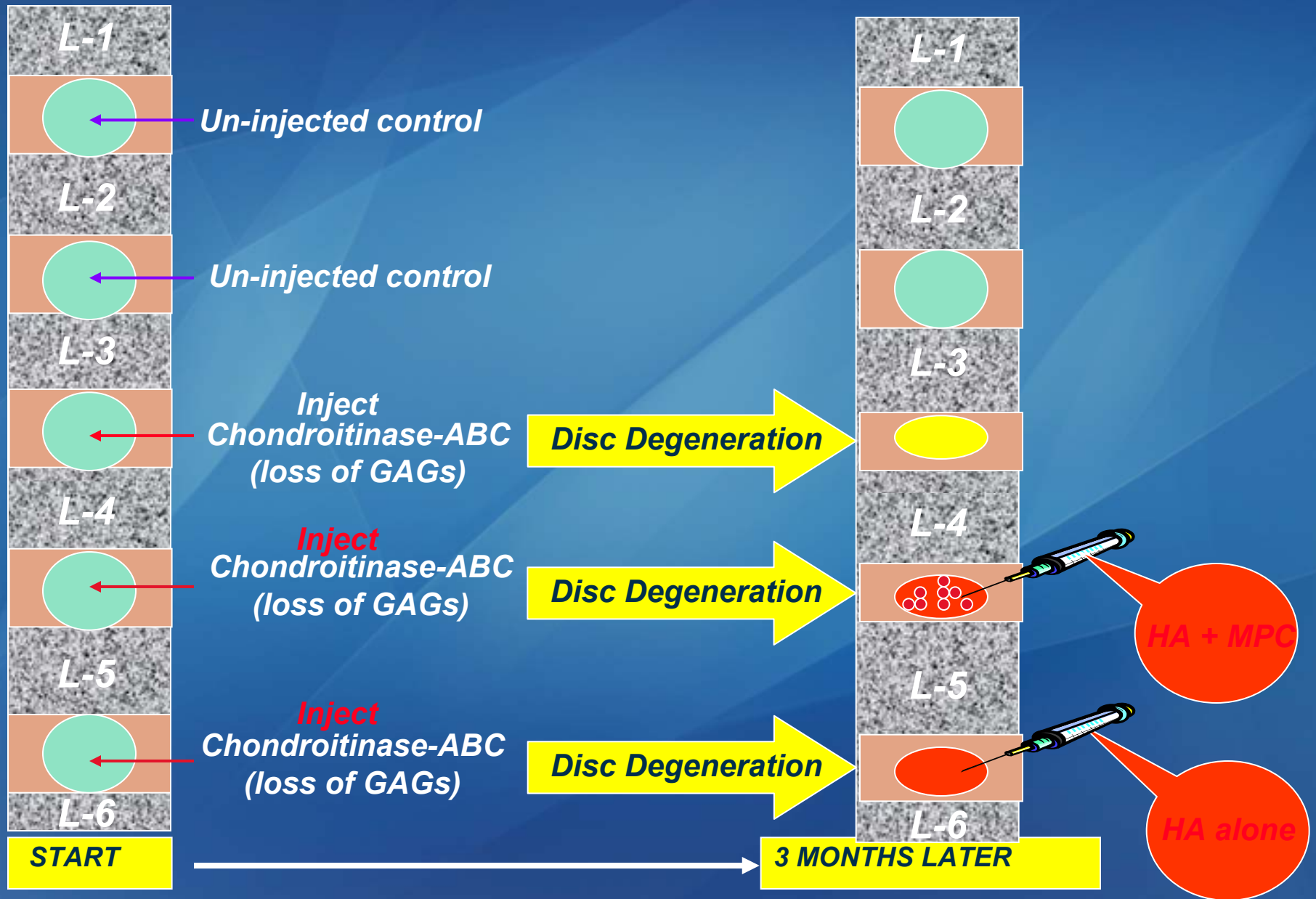
2. Partnering Specific Applications, e.g. For Spine Indications

- *partner(s) when technology/clinical risk reduced*
- *may drive more favorable distribution deal as Phase 3 costs lower than Pharma indications*
- *share the risk, share the reward*
- *de-risk need for sales/marketing force*

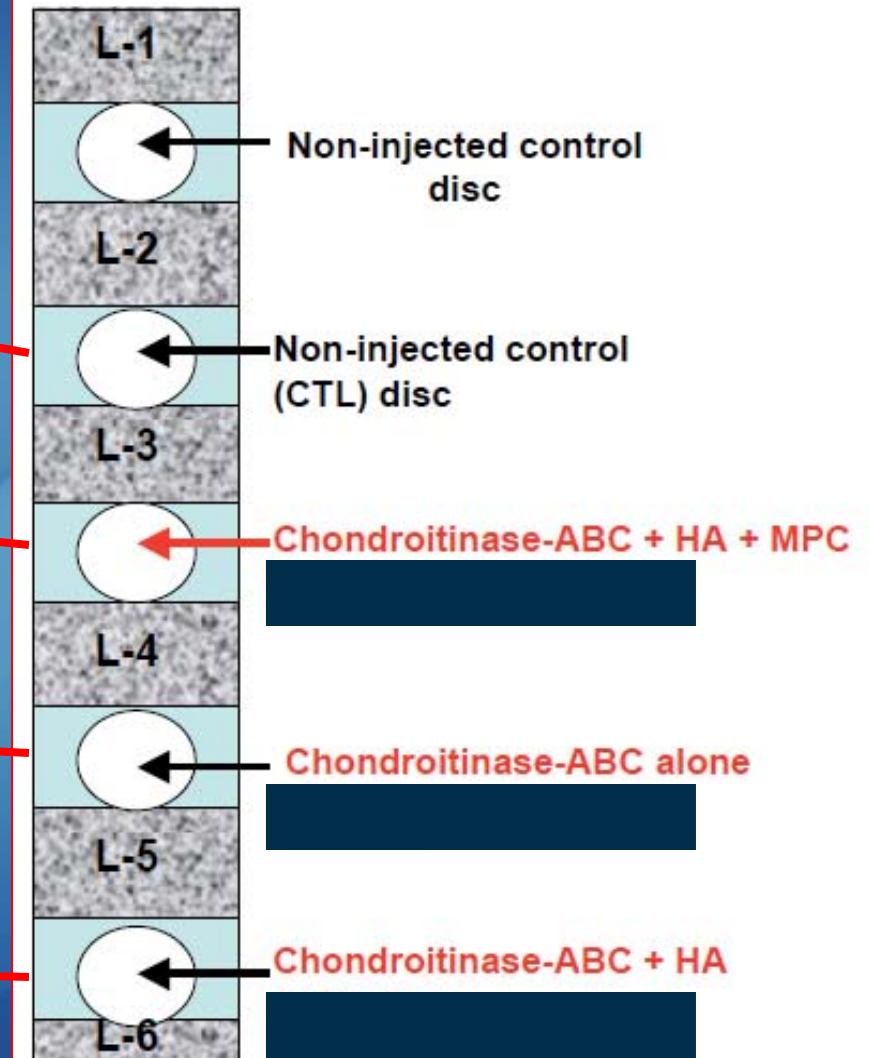
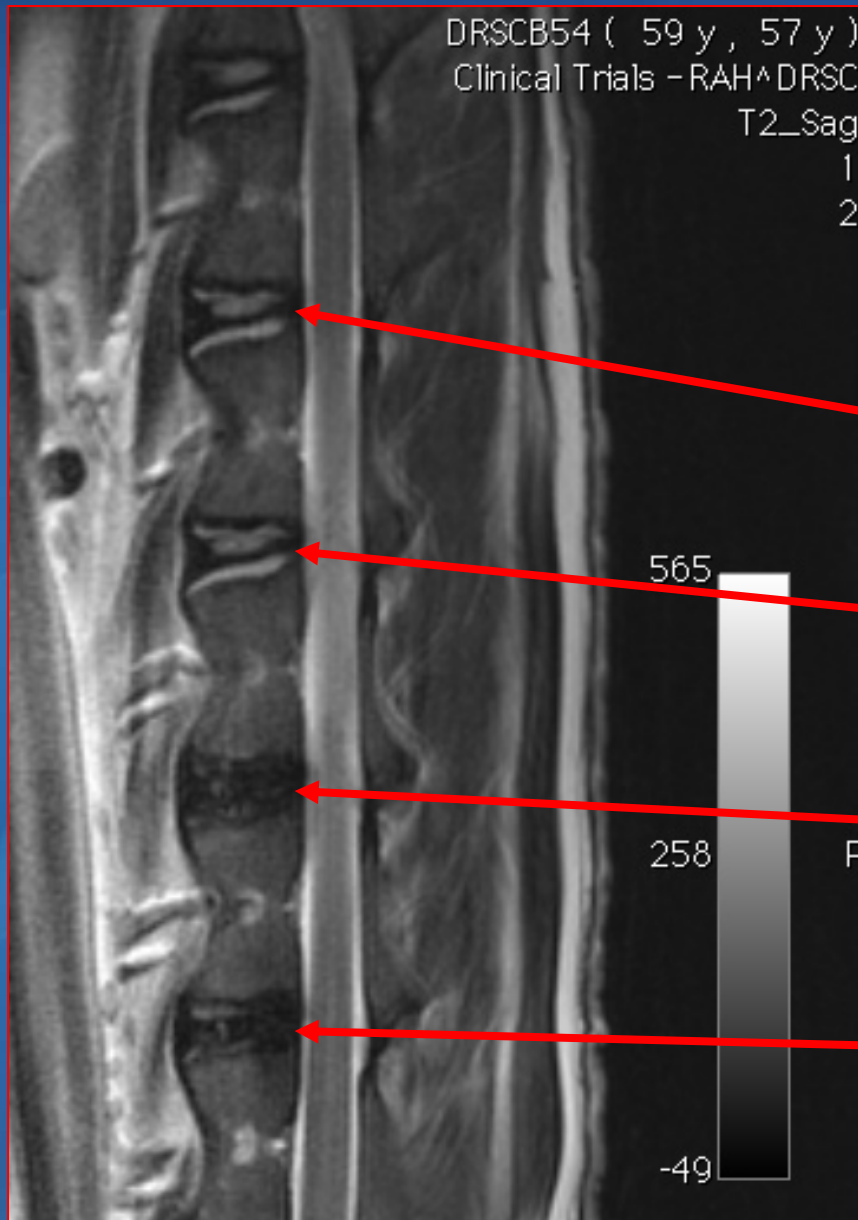
Disc Degeneration *Number One* Cause of Chronic Back Pain



Sheep model to evaluate the re-constitution of degenerate discs by injections of MPC + hyaluronan (HA)

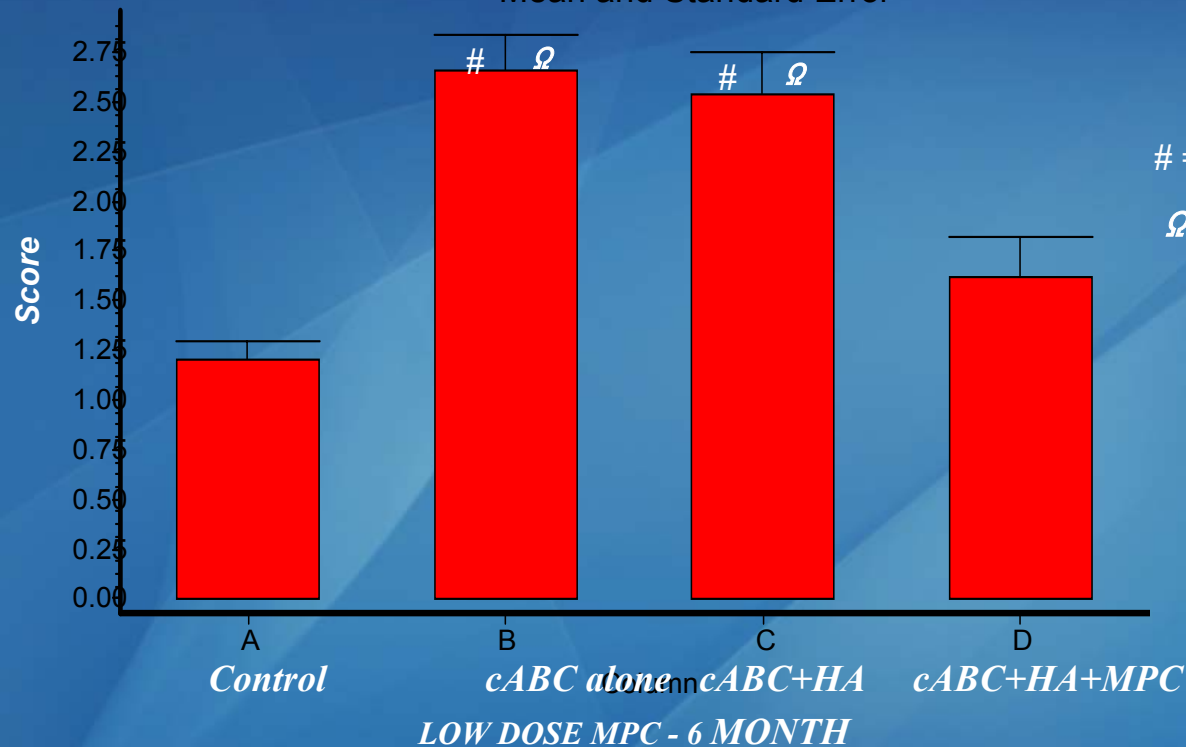


MRI of an ovine spine 6 months after injecting 0.5 million MPC showing similar H₂O signal (white) to normal control ovine disc. Note: L4L5 & L5L6 low H₂O signal



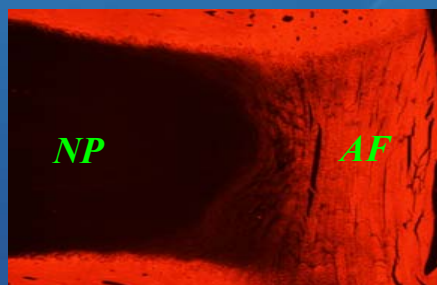
Aggregate Histopathology Disc Degeneration Scores Show MPC Return Disc Pathology To Normal

Means of Aggregate Histopathology Disc Degeneration Scores at 6 Months
Mean and Standard Error



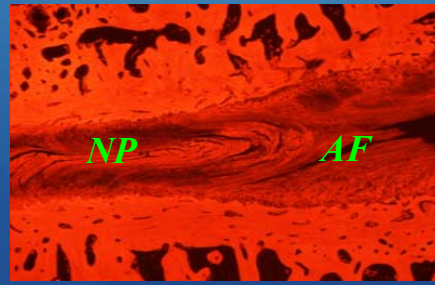
= significantly different from control $p < 0.001$;

Ω = from MPC $p < 0.01$



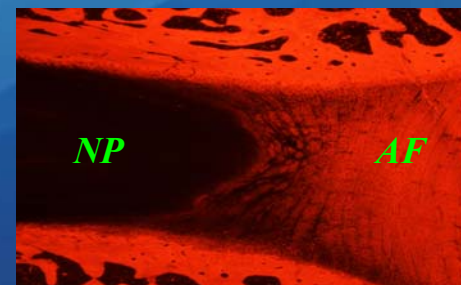
Normal Control Disc

(AF histopath score = 1).



Degenerate Disc

(AF histopath score = 4).



Degenerate Disc + MPC

(AF histopath score = 1)



Commercial Strategies For MPC-Based Biologicals

3. Taking Individual Applications to Market On Own e.g. for Orphan Drug Indications

- ***relatively inexpensive Phase 3 trials***
- ***can build small sales/marketing team for niche indications***
- ***easy access to key hospital centres of excellence, e.g. bone marrow transplant sites***
- ***fastest route to market***
- ***special reimbursement opportunities***
- ***greatest shareholder return***



Bone Marrow Transplantation

- ***139,000 new cases hematologic cancers each year in US alone***
- ***17,500 BMTs in USA in 2008***
- ***39,000 BMTs in EU in 2008***
- ***Allogeneic BMT could increase BMT market by > three-fold if an effective alternative to adult bone marrow existed, without risk of graft-versus-host disease (GVHD)***
- ***Umbilical Cord Blood lower risk of GVHD, but less effective than adult bone marrow because insufficient cell numbers***



MPC-Expanded Cord Blood For Bone Marrow Transplantation

- ***Our proprietary MPC induce 40-fold expansion of Hematopoietic Stem Cells (HSC) in co-culture***
- ***Orphan Drug designation received for increasing HSC in cancer patients needing allogeneic BMT***
- ***NIH-funded FDA-cleared trial in up to 30 patients transplanted with MPC-expanded cord blood HSC***
- ***Pilot trial to show earlier bone marrow engraftment, reduce in-hospital stay, reduce GVHD***
- ***Plan for Phase 3 trial to obtain early FDA approval***

Clinical Trial Results

- ***18 patients transplanted to date, mean age 40 yrs***
- ***median time to neutrophil engraftment 16.5 days
(historic controls approx. 30 days)***
- ***median time to platelet engraftment 38.5 days
(historic controls >90 days)***
- ***one patient has Grade III GVHD
(historic controls 40% Grade III/IV GVHD)***

Conclusions:

MPC-expanded cord blood comparable to allogeneic adult marrow for bone marrow reconstitution, with significantly reduced risk of GVHD

Results support accelerated Phase 3 program



Timelines For Lead Programs



angioblast
systems

IND
Clearance

FDA
Approval

Mesoblast Programs

Preclinical

Phase I

Phase II

Phase III

Spinal Fusion

Intervertebral Disc Repair

Long Bone Fracture Repair

Knee Osteoarthritis

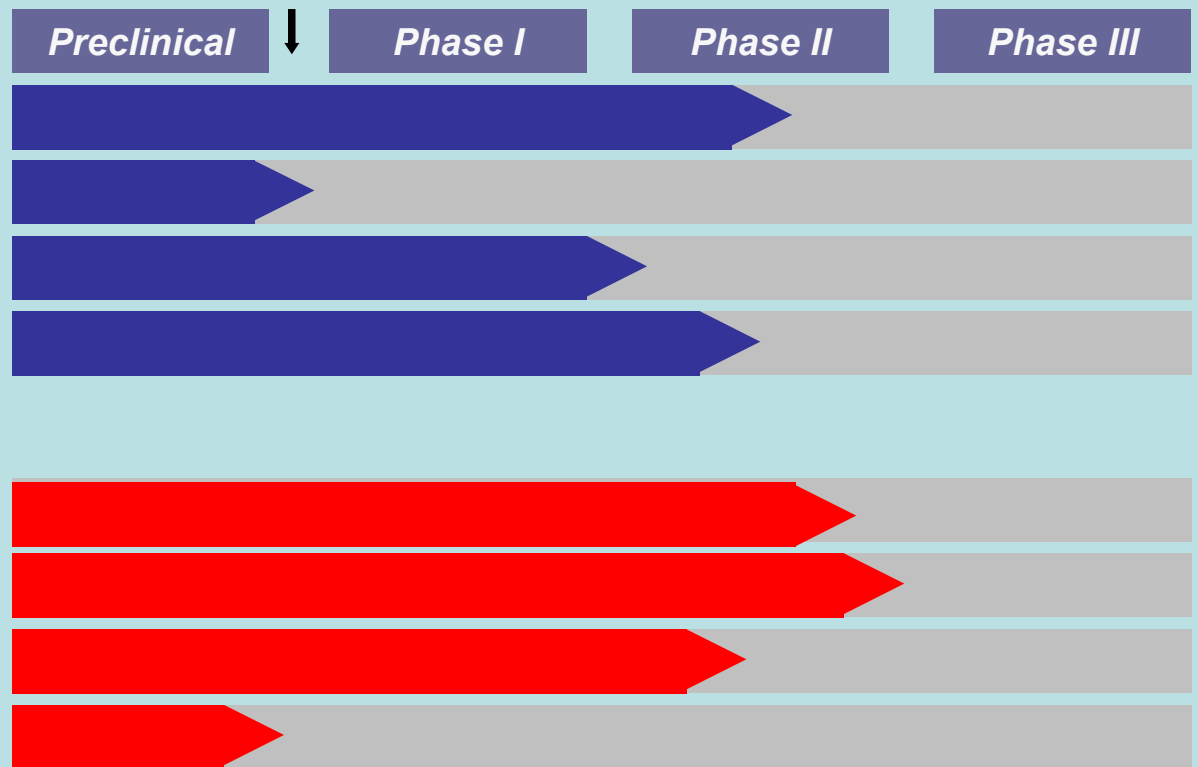
Angioblast Programs

Congestive Heart Failure

Bone Marrow Transplantation

Acute Myocardial Infarction

Eye Disease (AMD)



Australian Regulatory Approval May Mean Earlier Revenues

Approval Process Underway

- **Manufacturing process contracted with TGA-approved manufacturer**
- **Documents for product manufacturing license submitted to TGA**
- **Decision expected within months**

Commercial Advantages

- **Early revenues via Special Access Scheme**
- **Accelerated entry into fracture and other markets with high-end pricing points**
- **Registration template for Asian and other jurisdictions**
- **Early clinical data for US registration**