International Symposium on 3D Imaging for Interventional Catheterization in CHD

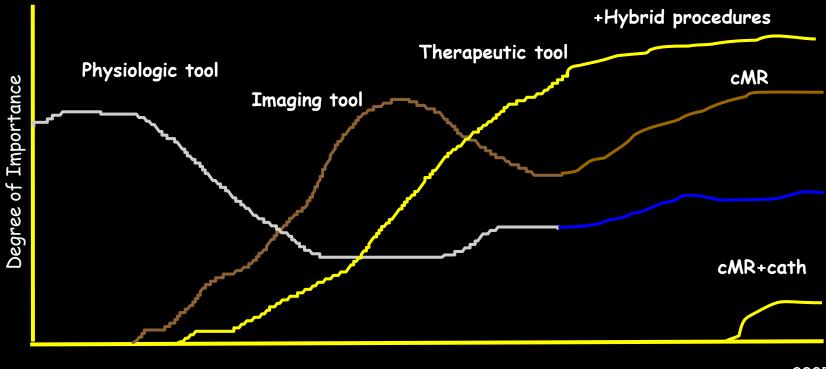
I Like My MR Scanner Adjacent to My Angio Suite Because.....

Lee Benson MD The Hospital for Sick Children, Toronto





The Changing Role of the Cardiac Catheterization Laboratory



1940 thru the 1950's 1960 thru the 1970's 1980 thru the 1990's 2000 thru 2005 >2005

The question to ask is why have a combined unit?

- The 2 imaging technologies are complimentary
- Allows acquisition of pressure & flow variables under consistent physiological conditions
- cMR imaging 2D/3D may facilitate catheter procedures
- Avoid multiple sedation/GA episodes
- Reduces radiation exposure
 - Many diagnostic angiograms can be replaced by cMRAs
 - IMR instead of x-ray guided interventions (?)

Combined UNIT System Requirements

cMR unit

Biplane/monoplane catheterization unit Sliding door with RF & X-ray shielding Transportation mechanism

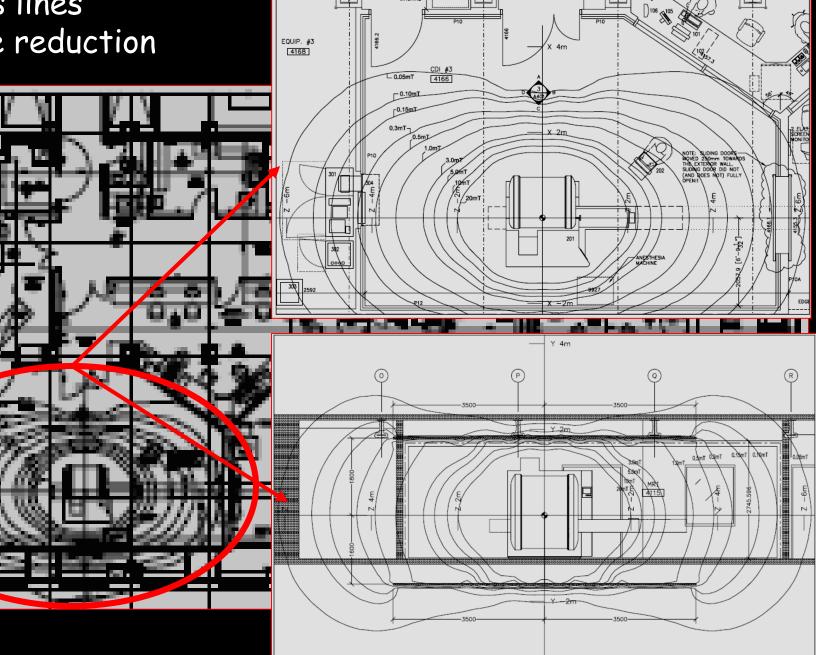
- Separate trolley
- Extended tables with rollers

CDIU physical plant integration



Gauss lines Noise reduction

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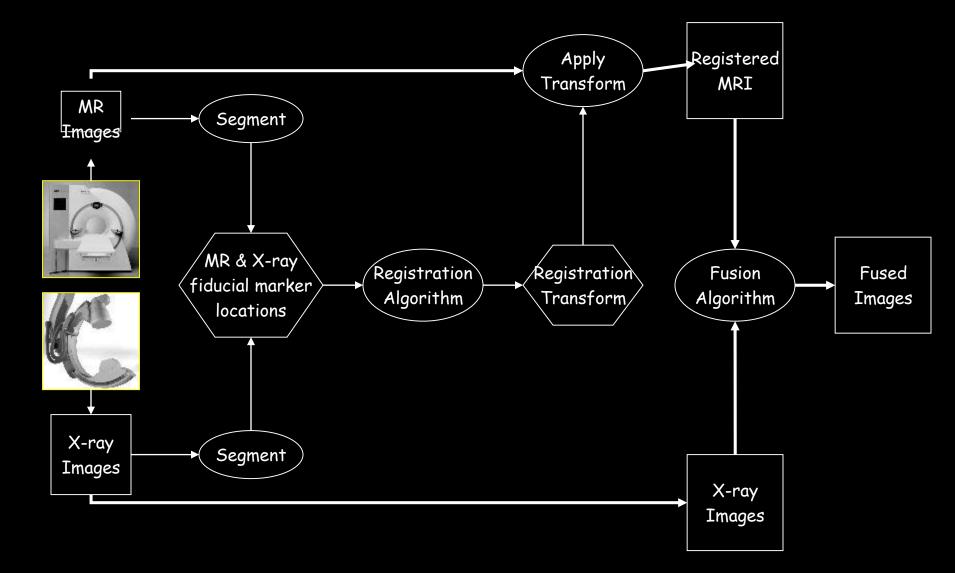




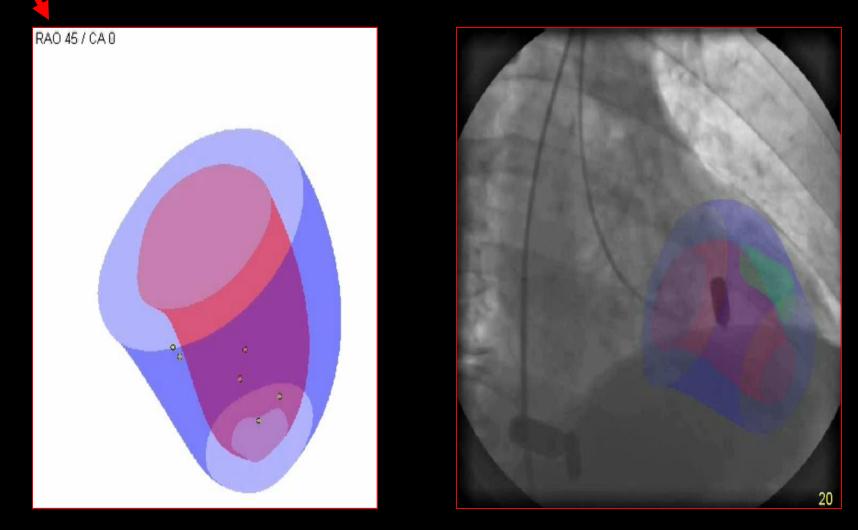




cMR data may help facilitate catheter procedures Real-time X-ray FUSED with cMRI



Real-time X-ray FUSED with MRI





Combined procedures (36 in 2015, 62 so far 2016)

Before diagnostic catheterization

prior to a Fontan operation assessment of APC's CA anomaly after re-implantation -ALCAPA; ARCAPA after a Fontan operation PAH study (multiple sources PBF) anatomical assessment

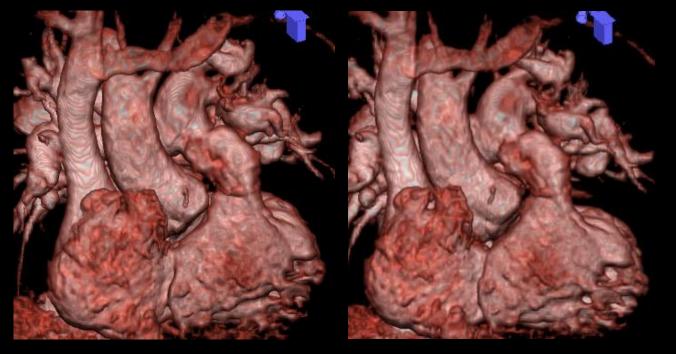
Before &/or after catheter interventions

RV-PA conduits branch PAS BCPS a prior hybrid procedure - HLHS

Ventricular performance, anatomy (volumes)

endomyocardial biopsy (tumor) combined with brain MRI prior to a hybrid stage II

Research studies



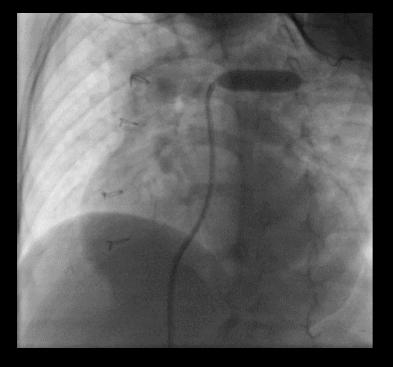


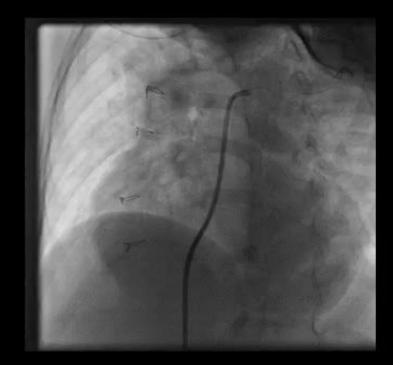




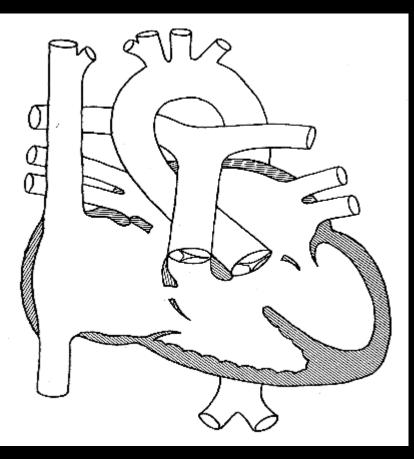


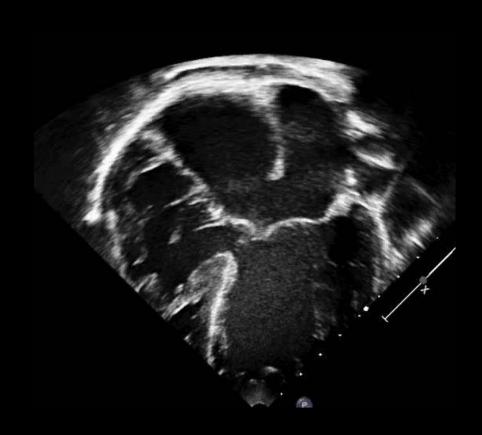






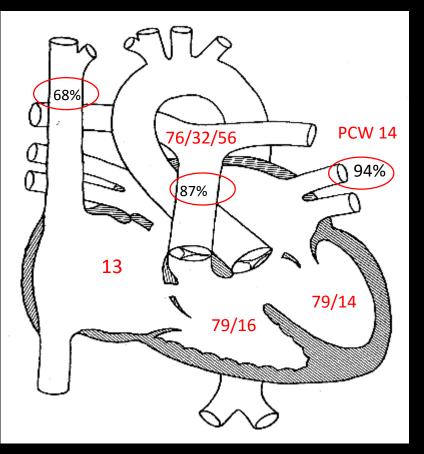
Can cMR help with hemodynamic assessments? Cardiac catheterization in patients with known PAH

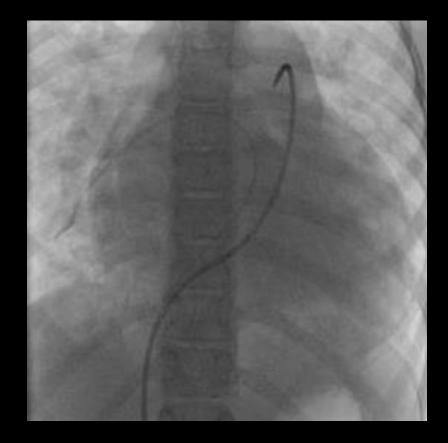




12 year old AVSD..is she operable?

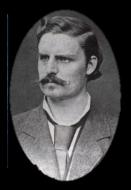
Can cMR help with hemodynamic assessments? Cardiac catheterization in patients with known PAH





12 year old AVSD..is she operable?

Based on Fick principle



Pulmonary blood flow =

Pulmonary AV-O₂ content difference

VO

Adolf Gaston Fick

Pulmonary vascular resistance:



<u>mean PAP - mean LAP (or mean PCWP)</u> PBF Wood units (mmHg/l/min)

Not Tiger Woods

That's pretty easy



Any invasive catheterization has risks, especially in patients with PVD.

However, with the currently available techniques & materials, including non-ionic contrast media, the risks are relatively low.

• What are the pitfalls in the calculation of PVR?

Common sources of errors

- Use of inappropriate value/sample for "mixed venous" blood
- Failure to calculate dissolved O₂ when using enriched gas (for example, 100% O₂)

Common sources of errors

- Assumed O₂ consumption is notoriously unreliable
- Unfamiliarity with O₂ consumption measurement technique—leads to unpredictable/unreliable results

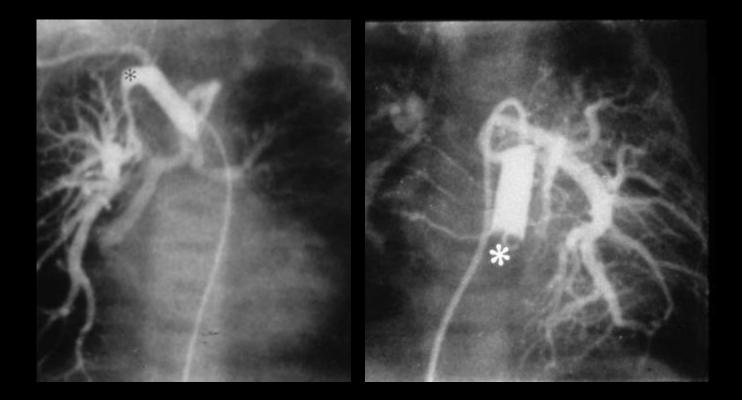
The LaFarge & Miettinen table has been largely used for the assumed O_2 consumption, but in young (infants) it tends to give higher O_2 consumption values resulting in optimistic results (lower values) for PVR.

Calculations based on assumed O_2 consumption is a real problem in cyanotic patients (measured values >assumed ones).

If assumed, use the range of values.

infants: 2 -5 years: adolescents: adult females: adult males: 130 (<3mon) - 170 ml/min/m² 150 - 200 ml/min/m² 120 - 180 ml/min/m² 100 ml/min/m² 110 - 120 ml/min/m²

Patients with multiple sources of PBF (i.e. MAPCAs) or discontinuous PA's pose a very special problem.



Novel Method of Quantifying Pulmonary Vascular Resistance by Use of Simultaneous Invasive Pressure Monitoring and Phase-Contrast Magnetic Resonance Flow Muthurangu et al Circ 110:2004

Phase-Contrast Magnetic Resonance Quantification of Normal Pulmonary Venous Return

Goo et al JMRI 29:2009

Cardiovascular Magnetic Resonance catheterization derived pulmonary vascular resistance and medium-term outcomes in congenital heart disease Pushparajah et al JMRI 2015



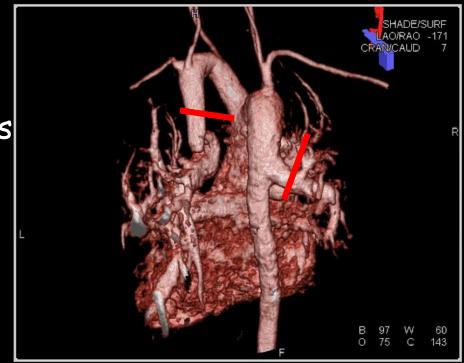




Flow volumes in ml/beat

```
Q_{P} = Q_{PV} = Q_{AA}
Q_{S} = Q_{SVC} + Q_{IVC} = Q_{PA}
Q_{APC} = Q_{PV} - Q_{PA}
= Q_{AA} - (Q_{SVC} + Q_{IVC})
= Q_{AA} - Q_{PA}
```

14 year boy unrepaired PA/VSD, MAPCAs unrestrictive VSD O₂ saturation 88% moderately limited



MRI Qs =SVC+DAO flows Qp = pulmonary vein flow Qp/Qs = 2.58

r-MAPCAs: inaccurate

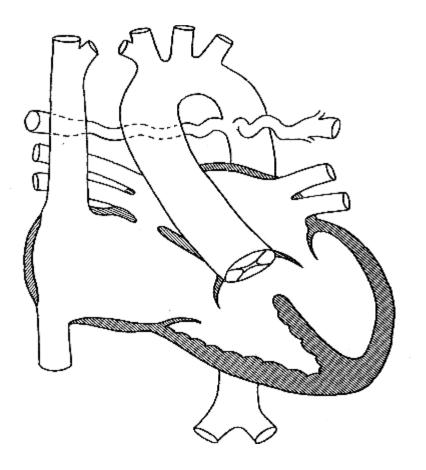
right pulmonary veins: I-MAPCA

Cath r-mAP wedge I-mAP = 2.40 l/min/m² = 6.20 l/min/m²

= 2.16 l/min/m² = 3.61 l/min/m²

= 61mmHg = 15mmHg = 17mmHg

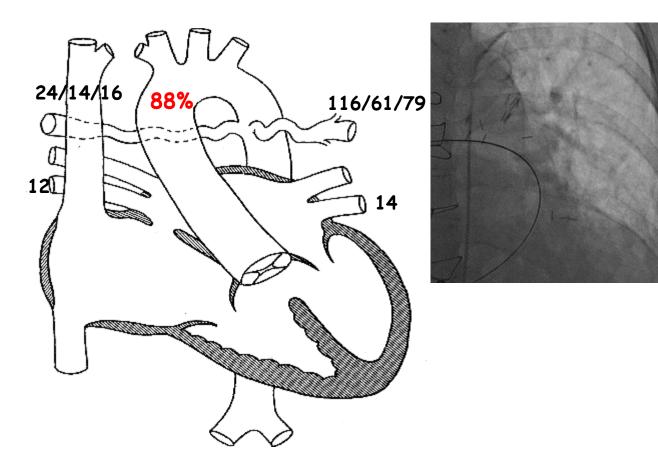
Total resistance= $(R_1 * R_2)/R_1 + R_2$

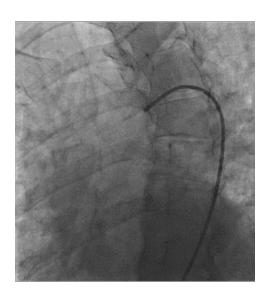


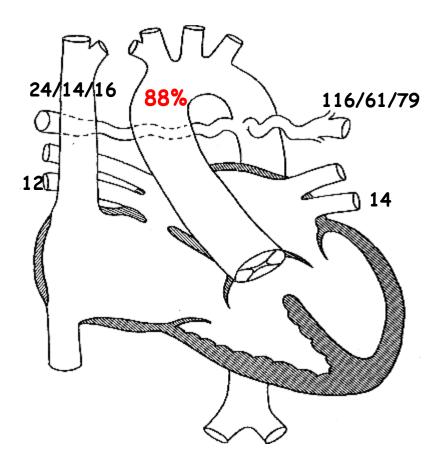
41 year old Increasing cyanosis with effort RA O_2 saturation 80% in 15 | O_2

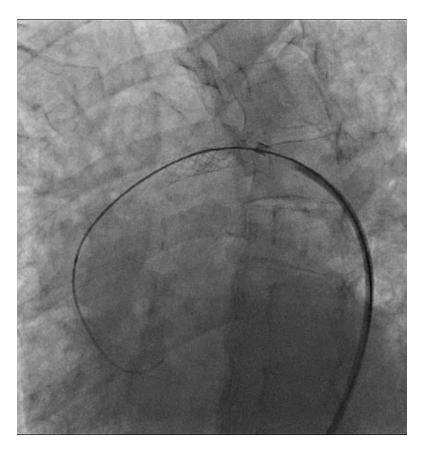
Is there something we can do to improve her clinical status, can she be repaired?

Pulmonary atresia, VSD, ductal origin of RPA, APC to LPA









Data from cMR (includes arterial collateral flow)

RPA/LPA flow ratio 92%/8%

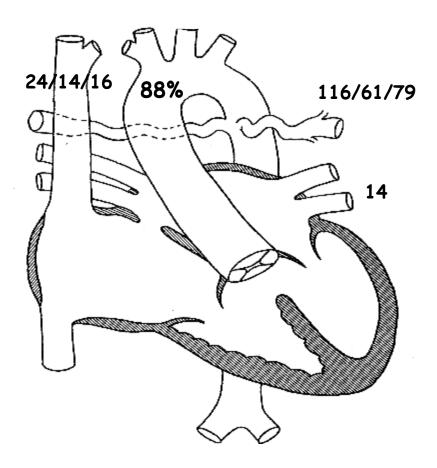
HR 57beats/min

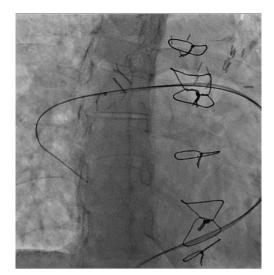
LPV flow 24ml/beat TPG 65mmHg

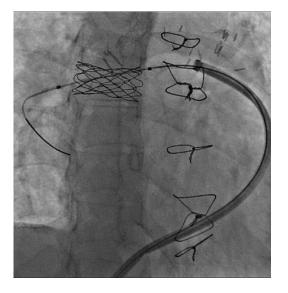
RPV flow 35ml/beat TPG 4mmHg



R-PVR = 4/1.995 = 2 Wood Units







rPAP systolic ~40mm Hg

A Combined Unit SickKids Lessons Learned

*Expensive investment

*Impacts on lab resources/scheduling

Excellent for research purposes

Benefits outweigh risks

* Allows for enhanced understanding of pathophysiology not possible with 2D invasive studies alone

Do I like the cMRI right next to the cath lab?

Yes



Jose Bautista Game 5 3-run home run AL Div series 2015

When it comes to improving our understanding of physiology & anatomy

Do I like the cMRI right next to the cath lab?

No

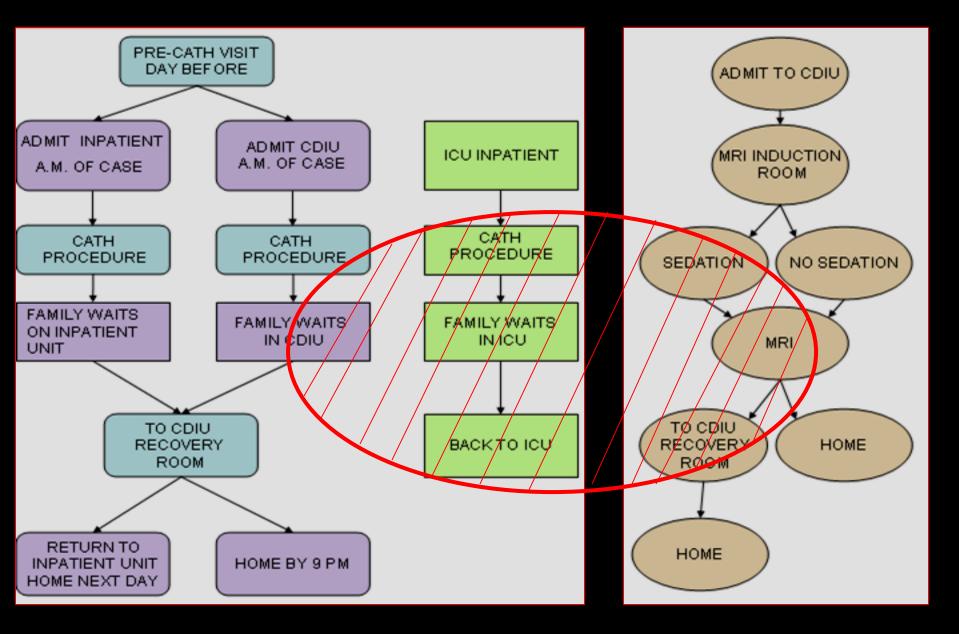


Jose Bautista taking a hard shot from Texas Ranger 2nd basemen Rougned Odor

When it comes room utilization and scheduleing

Cath Lab Patient Flow

cMRI Patient Flow



The Cardiac Diagnostic and Interventional Unit (CDIU)

Dr. Lee Benson Dr. Jin Lee Dr. Rajiv Chatervurdi Dr. Andrews Wan Interventional Dr. Shi-Joon Yoo Dr. Mike Seed Dr. Lars Grosse-Wortmann * cMRI/Diagnostic Dr. Robert Hamilton Dr. Joel Kirsch Dr. Gil Gross

Dr. Beth Stephenson

Electrophysiology

Susan Johnston Sandra Skrt-Martin Janine Barclay Jacqueline Viegas Martine Dubreuil *Nurse technicians

Darius Mroczek Biomedical techn. Rachel Enkin Cath technician Kelly Paredes *****Inform. Co-ordinator **Omar Thabit** Vinay Kainthla Jothi Thind Sundar Devadas *cMR/MRT

"The future is not some place we are going to, but one we are creating. The paths are not to be found, but made, & the activity of making them changes both the maker & the destination"

John Schaar, Letgitimacy in the Modern State. 1989

THANK YOU