

Numerical Simulations for the Planning of Surgical Procedures

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Why pediatric cardiac surgery?

How many people in the United States and in Europe have a congenital heart defect?

Estimates suggest that about 1,000,000 Americans have a congenital heart defect (American Heart Association). Approximately 36,000 babies are live born with a heart defect each year in the European Union (EUROCAT Central Registry).

How serious is the problem?

Congenital heart defects are the most common birth defect and are the number one cause of death from birth defects during the first year of life. Nearly twice as many children die from congenital heart disease in the United States each year as die from all forms of childhood cancers combined. Over 91,000 life years are lost each year in the US due to congenital heart disease. Charges for care exceed 2.2 billion dollars, for inpatient surgery alone (American Heart Association).

Are things improving?

Definitely. Overall mortality has significantly declined over the past few decades. For example, in the 1960s and 1970s the risk of dying following congenital heart surgery was about 30% and today it is around 5%.

Pediatric cardiac surgery

How the story started: back to Great Ormond Street Hospital for Children, London, 1993, and to the Total CavoPulmonary Connection (TCPC)







LaBS

Prof. Marc R. de Leval Laurea ad honorem in Biomedical Engineering, Politecnico di Milano, 2007

Aim:

To investigate the effects of geometric/anatomic features (e.g. *offsetting* and shape of the anastomosis between the inferior vena cava and the pulmonary artery)

de Leval et al. Use of computational fluid dynamics in the design of surgical procedures: application to the study of competitive flows in cavo-pulmonary connections, J Thorac Cardiovasc Surg. **1996**

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TCPC: Total CavoPulmonary Connection





(de Leval et al., J Thorac Cardiovasc Surg, 96:682-695, 1988)

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(de Leval et al., J Thorac Cardiovasc Surg, 113:502-13, 1996)



Numerical simulation for pediatric cardiac surgery:

- 1. To investigate phenomena hardly measurable in the clinical setting
 - energy dissipation (recirculation areas, jets, etc.)
 - blood and oxygen flow distribution to organs

- 2. To optimize the surgery technique
- 3. To plan surgery

Multi-domain modelling approaches - 1





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Multi-domain modelling approaches - 2





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accuracy

CPU time

Computational Fluid Dynamics (CFD) 3D modelling



local haemodynamics information

- rigid or deformable walls
- from idealized to patient-specific

1D modelling

• pulse wave propagation phenomena (curvature, branching, stenoses)

• rigid or elastic

0D modelling (lumped parameter, LPM)

- more extensive circulatory networks
- fast-computing

Multi-domain modelling

- coupling of 3D domain reproducing specific vascular portions (e.g. surgery site) with the downstream circulation described by LPM or 1D model
- closed-loop approach to account for the mutual interaction between the two domains.

Multi-domain modelling approaches - 4



CFD allows an approximation of a real situation, dividing the physical problem domain (object) in smaller (elements parts or volumes) and solving each single part. If the division (mesh) is the results good, are an approximation, but very close to the reality.



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Multi-domain models





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[from http://xpsolutions.com/Software/XP2D/]

[from http://www.ricardo.com/en-GB/What-we-do/Software/Products/VECTIS/Coupled-1D3D-Analysis/]

Multi-domain model approach already in use in the automotive industry (e.g.intake and exhaust runners), environmental planning (e.g. impact of surface flooding on an existing sewer network), power generation industry (e.g. wind plant design), etc.



[from http://www.mentor.com/products/mechanical/flowmaster/flowmaster-process-power-energy/]

Multi-domain models





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Multi-domain models



Partial differential equations (Navier-Stokes)



Ordinary differential equations (ODE)

Quarteroni et al. "Coupling between lumped and distributed models for blood flow problems." Computing and Visual Science 4, 111-124, 2001.

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Introduction: single-ventricle congenital heart diseases







www.childrenshospital.org

Steps of a virtual surgical planning:

- implementation of a **pre-operative computational model** to reproduce the hemodynamics of the patient
- comparison of **different surgical options** through CFD analysis.



WORKFLOW





WORKFLOW





STEP 1: Pre-operative clinical data

Doppler acquisition



Catheterization



MR imaging and flows







STEP 1: Pre-operative clinical data

- ✓ 6 months-old infant (weight 6.9 kg, BSA 0.34 m²)
- Hypoplastic right heart syndrome + pulmonary atresia
- ✓ 3.5 mm central shunt (from AoA to MPA) surgically placed on day of life #5

Doppler acquisition (5 weeks prior to surgery, under sedation) Echocardiograpy system (iE33, Philips, Best, Netherlands)

Catheterization (under sedation, 3 days before the surgery)

Biplane fluoroscopy suite (Toshiba America Med. Syst., Inc., CA, or Siemens Med. Solutions, USA, Inc. PA).

MR imaging (under general anesthesia immediately prior to surgery) 1.5 T scanner (Philips Intera Achieva, Best, Netherlands). Imaging parameters: 4.4/2.3 (repetition time msec/echo time msec), 12° flip angle, 1.3–1.5-mm section thickness, 256 × 512 matrix, 400–500-mm field of view, one breath hold, one signal acquired, no electrocardiographic gating.

Three-dimensional MR data reconstruction

Mimics software (Materialise NV, Leuven, Belgium). Image elaboration for each patient's MR data took 2–3 hours.









STEP 1: Pre-operative clinical MRI and Doppler flows













WORKFLOW





STEP 2: Pre-operative 3D reconstruction

Whole geometry reconstruction

Selection of the sub-model of the pulmonary arterial vasculature











Whole geometry reconstruction



Selection of the sub-model of the pulmonary arterial vasculature



MESHSIM (Simmetrix, Inc., NY)



Volume meshing



STEP 2: Pre-operative pulmonary LPM - morphometric approach





 - R_p, C, R_d are given by least-squares fit of the impedance spectrum

> Troianowski et al., 2011 J. Biomech. Eng. Spilker et al., 2007 Ann. Biomed. Eng.



- ΔP between 3D inlet and single atrium SA
- size of each outlet branch $Q_i \propto S_i^{\alpha}$

Iterative simulations until convergence



STEP 2: Pre-operative closed-loop LPM



Setting of the parameters

- <u>HR</u> = 120 bpm
- <u>BSA</u> = 0.34 m²
- Vascular resistances [mmHg*s/ml]

$$UBSVR = \frac{P_{Ao} - P_{SA}}{Q_{UBA}} = \frac{43 - 5}{11.2} = 3.39$$

$$LBSVR = \frac{P_{Ao} - P_{SA}}{Q_{THAO}} = \frac{43 - 5}{5.7} = 6.67$$

- <u>Pulmonary RCRs</u> provided by morphometric approach
- <u>Shunt</u> Operating point: dP= 43-13 = 30 mmHg, Q_{SH} = 7.5ml/s Relation: $dP = R_{SH}Q_{SH} + K_{SH}Q_{SH}^2$

Migliavacca et al., 2001, Am J Physiol Heart Circ Physiol

- Respiratory effects were not included
- Single values of resistance and compliance vary depending on the BSA

Snyder, Rideout, 1969, IEEE Trans Bio-Med Eng





STEP 2: Pre-operative closed-loop LPM - Results



[ml/s]	Clinical mean values	LPM
Q _{UB}	11.2	11.3
Q _{LB}	5.7	5.7
Q _{LPA}	2.7	2.8
Q _{RPA}	4.8	4.8
СО	24.4	24.4
[mmHg]	Reference clinical data	LPM
_		
P _{SV}	6 / 103 / 54	0 / 72 / 25
P _{sv} P _{sA}	6 / 103 / 54 5	0 / 72 / 25 6.8
P _{sv} P _{sa} P _{pa}	6 / 103 / 54 5 13	0 / 72 / 25 6.8 14.7
P _{SV} P _{SA} P _{PA} P _{SVC}	6 / 103 / 54 5 13 9	0 / 72 / 25 6.8 14.7 7.9
P _{SV} P _{SA} P _{PA} P _{SVC} P _{DAo}	6 / 103 / 54 5 13 9 43	0 / 72 / 25 6.8 14.7 7.9 43.8
P _{SV} P _{SA} P _{PA} P _{SVC} P _{DAo} [ml]	6 / 103 / 54 5 13 9 43 Clinical mean values	0 / 72 / 25 6.8 14.7 7.9 43.8 LPM
P _{SV} P _{SA} P _{PA} P _{SVC} P _{DA0} [<i>ml</i>] EDV	6 / 103 / 54 5 13 9 43 Clinical mean values 22.6	0 / 72 / 25 6.8 14.7 7.9 43.8 LPM 21.8
P _{sv} P _{SA} P _{PA} P _{Svc} P _{DAo} [<i>ml</i>] EDV ESV	6 / 103 / 54 5 13 9 43 Clinical mean values 22.6 9.6	0 / 72 / 25 6.8 14.7 7.9 43.8 LPM 21.8 9.6



STEP 2: Pre-operative closed-loop LPM - Results

S/A IVC



-1,15

-0,95



STEP 2: Pre-operative closed-loop LPM - Results







WORKFLOW







Different options:





Glenn

hemi-Fontan



STEP 3: Post-operative model – closed-loop 3D-0D models





Hemi-Fontan



Pulmonary RCR blocks





WORKFLOW





STEP 3: Post-operative model – Simulation results



STEP 3: Post-operative model – Simulation results







Investigated fluid dynamic quantities	Glenn	Hemi-Fontan
Q _{svc} (ml/s)	11.4	11.3
Q _{ıvc} (ml/s)	7.87	7.89
Q _{LPA} (ml/s)	4.17	4.21
Q _{RPA} (ml/s)	7.21	7.07
P _{svc} (mmHg)	15.1	15.4
P _{Ao} (mmHg)	53.1	53.1
P _{SA} (mmHg)	2.48	2.43
Power Loss Surgical Junction (mW)	0.61	1.26
Efficiency Surgical Junction	0.97	0.95
Ventricular power (mW)	166	166



Conclusions



The workflow

- constant interactions among clinicians and engineers to:
 - collect the patient's clinical data
 - select the data to be used in the modeling process
 - build a preoperative patient-specific model that reproduced the patient
 - simulate different postoperative surgical options
 - compare the local and global hemodynamic results

Not shown

• Different postoperative scenarios (e.g. agitation, exercise) have been simulated



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"Multi-scale Modeling of Single Ventricle Hearts for Clinical Decision Support" Transatlantic Project

http://modelingventricle.clemson.edu



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