

The background of the slide is a grayscale photograph of three surgeons in an operating room. They are wearing surgical masks, hairnets, and scrubs, and are focused on a patient. Large surgical lamps are visible at the top of the frame.

# Closed-Loop Computational Modelling of the Heart with Applications in Xenotransplantation

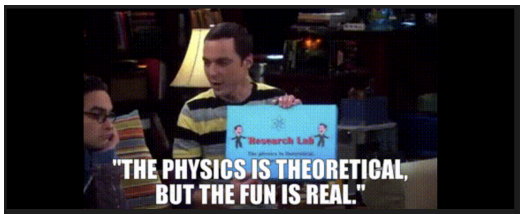
Research Studies Panel Meeting

David Kelly, Prof. Philip Cardiff

UCD School of Mechanical and Materials Engineering

December 15, 2025

## A bit about me...



- Graduated Sept 2025 from Theoretical Physics in UCD
- Started PhD supervised by Prof. Cardiff in Sept. 2025
- Working on ERC-funded XenoSim project.

# Background & Motivation

## Big Picture

- Donor heart shortage continues to limit cardiac transplantation.
- Xenotransplantation is a promising alternative, but haemodynamic performance remains poorly understood.
- Porcine-to-human implantation introduces distinct haemodynamic and electromechanical challenges.



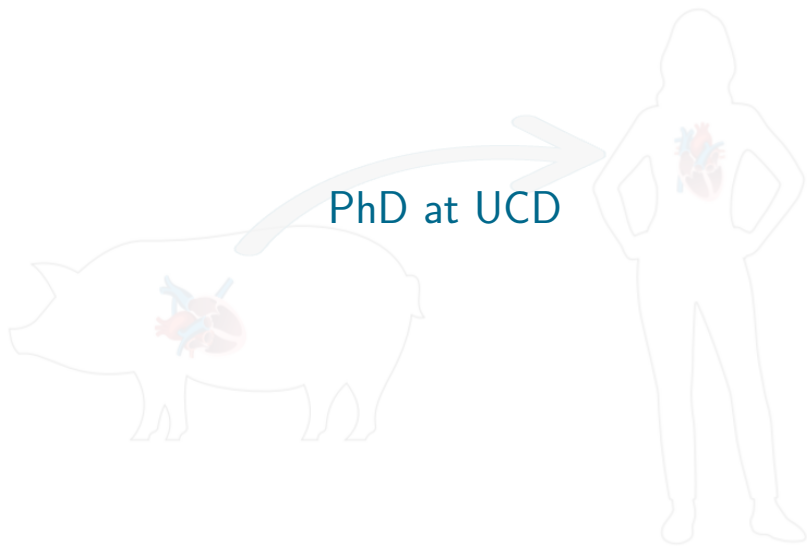
XenoSim

## Key Drivers

- Fundamental pig-human differences in anatomy, compliance, and vascular impedance.
- Limited in-vivo haemodynamic access motivates high-fidelity *in-silico* modelling.
- Need for predictive tools to assess pig-heart performance under human circulatory loading.



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# The Credits (all 30 of them!)



## Completed (Hopefully)

- *ACM40660* – Scientific Programming Concepts (ICHEC) (5 Credits)
- *MEEN40150* – Computational Continuum Mechanics II (5 Credits)

## Potential Future Modules

- *ACM40640* – High Performance Computing (ICHEC) (5 Credits – Spring)
- *ACM41000* – Uncertainty Quantification (5 Credits – Spring)
- *ACM40290* – Numerical Algorithms (5 Credits – next Autumn)
- *MEEN40910* – Teaching Assistant in MME (5 Credits – next Autumn)

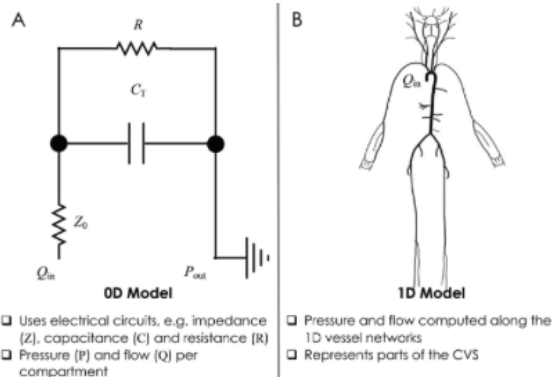
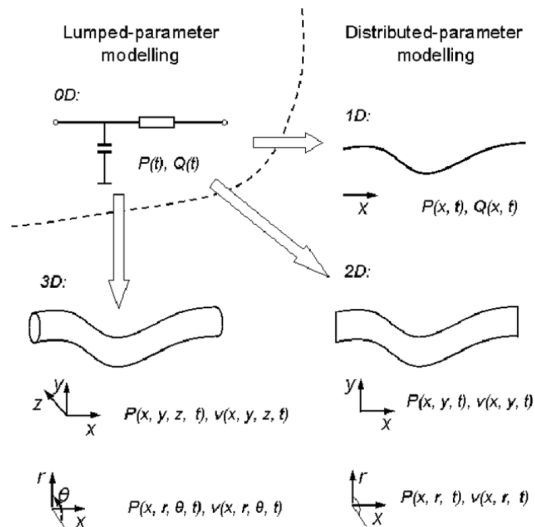


- Completed online Research Integrity Training on Brightspace
- Abstract accepted to Bioengineering in Ireland Conference (BINI) 2026

## Current Knowledge, Research Gaps & Objectives



# 0D & 1D Cardiovascular Models





# Why Closed-Loop 0D/1D Coupling?

## Motivation

- Xenotransplantation is fundamentally an **interaction problem**: a porcine heart operating under **human vascular loading**.
- Closed-loop circulation models provide a physiologically consistent way to impose **afterload, preload, and feedback** on the 3D heart.

## What 0D vs 1D adds

- **3D-0D**: robust, efficient closed-loop boundary conditions (mean pressure/flow and PV behaviour).
- **3D-1D**: captures spatial wave propagation and reflections  $\Rightarrow$  time-varying, phase-dependent afterload.
- Both support: **“pig heart + human circulation”** experiments in-silico.

# 3D–1D Coupling: Current Knowledge & Research Gaps

## Current Knowledge

- Coupling of 3D vessels or stenosis (*Hilhorst et al.*, *Formaggia et al.*) with 1D circulatory tree.
- Limited studies couple with 3D heart chambers, primarily focus only on **left ventricle only**.
- Caforio *et al.* couple a 3D human LV electromechanical model to a 1D arterial system.
- Found only one CFD-based studied (*Chen et al.*) couple 3D left ventricle to 1D circulation models.

## Research Gaps

- Extension of 3D–1D coupling beyond the LV to **multi-chamber heart models** is largely unexplored.
- Existing 3D–1D frameworks are restricted to **human physiology**.
- CFD-based (*Chen et al.*) approaches do not incorporate full cardiac electromechanics.
- Electromechanical approaches (e.g. Caforio *et al.*) do not resolve intraventricular flow or fluid–structure interaction.
- **No existing framework combines multi-chamber 3D modelling, closed-loop 1D circulation, and xenogeneic (pig–human) configurations.**

# 3D-0D Coupling: Current Knowledge & Research Gaps

## Current Knowledge

- Closed-loop coupling of 3D cardiac models with lumped-parameter (0D) circulation models is well established.
- Several studies couple multi-chamber **human** heart models to full systemic LPMs.
- Zingaro *et al.* demonstrate coupling of the full left heart (LV, LA, valves) to a closed-loop 0D circulation.
- 3D-0D coupling provides robust and computationally efficient haemodynamic prediction.

## Research Gaps

- Existing 3D-0D frameworks are restricted to **human cardiac anatomy and physiology**.
- No models investigate porcine heart behaviour under **human circulatory loading**, even at the 0D level.
- Limited reuse or extension of existing frameworks to support xenogeneic configurations.
- **A predictive 3D-0D framework for porcine hearts operating within a human circulatory system is absent.**

# Aim & Objectives

## Overall Aim

**To develop a multiscale computational framework for cardiac haemodynamics, and to apply it to the study of porcine hearts operating within a human circulatory system.**

## Specific Objectives

- ① Develop a stable closed-loop coupling between reduced-order circulation models and 3D cardiac chambers.
- ② Extend existing 1D–3D coupling approaches beyond the left ventricle to multi-chamber heart models incorporating electromechanics and fluid–structure interaction.
- ③ Apply the framework to xenogeneic configurations by coupling a 3D porcine heart with a human circulatory model.

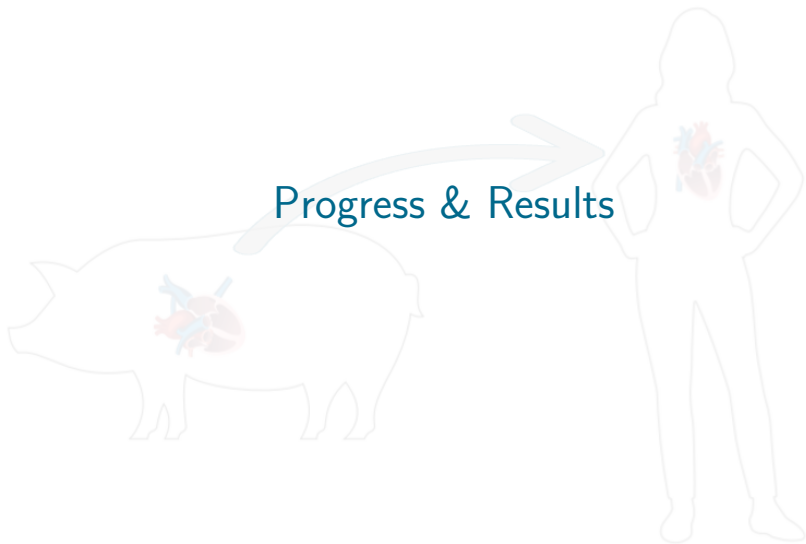
## Modelling Framework

- Lumped-parameter (0D) and one-dimensional (1D) circulation models.
- Three-dimensional cardiac chamber models within OpenFOAM.
- Closed-loop pressure–flow coupling at physiological interfaces.

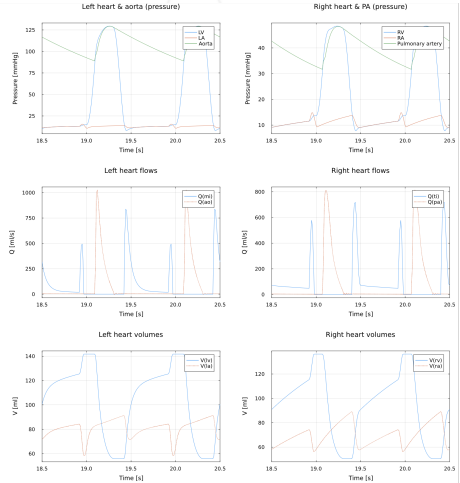
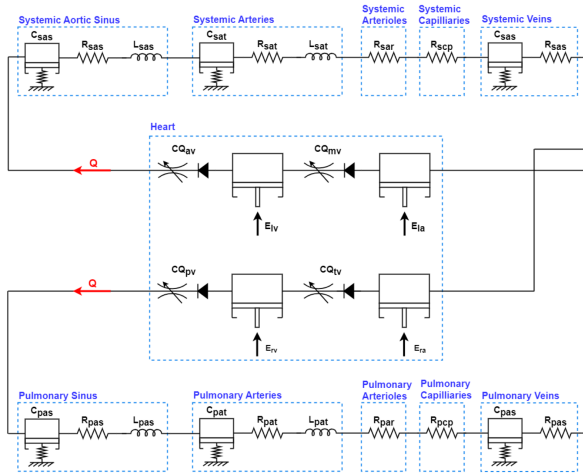
## Verification & Assessment

- Verification via volume conservation and flow balance.
- Comparison against published human physiological benchmarks.

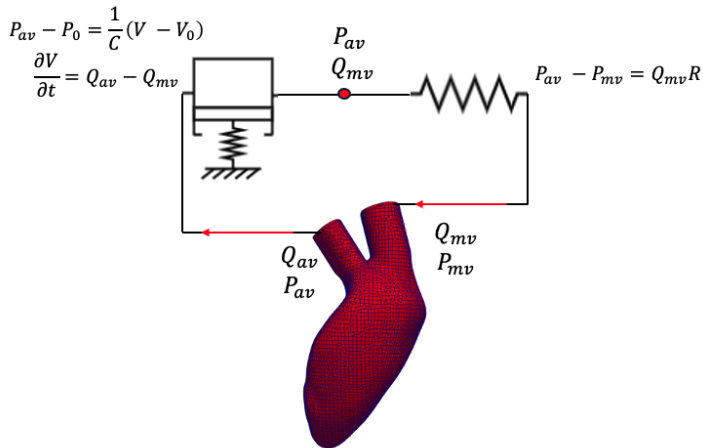
## Progress & Results



# Completely Lumped System



# Incorporating a 3D Left Ventricle in OpenFOAM



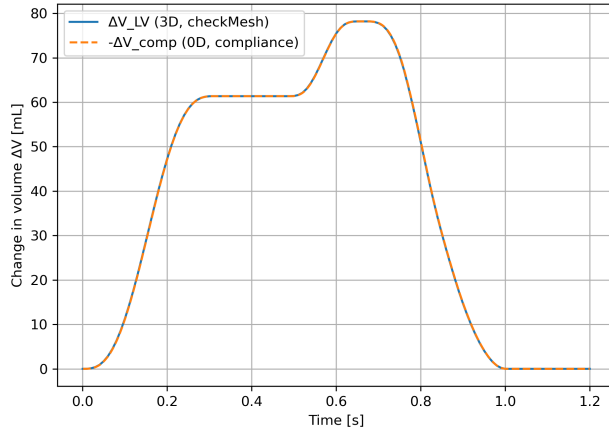
## OpenFOAM BCs

- Prescribe  $P_{av}$  and  $Q_{mv}$  at the OpenFOAM boundaries
- Compute  $Q_{av}$  and  $P_{mv}$  from the 3D model and use them to update the LPM
- $\frac{\partial P_{av}}{\partial t} = \frac{1}{C}(Q_{av} - Q_{mv})$   
(setting  $P_0 = V_0 = 0$ )
- $\frac{\partial P_{av}}{\partial t} = \frac{1}{C}(Q_{av} - \frac{1}{R}(P_{av} - P_{mv}))$
- $Q_{mv} = \frac{1}{R}(P_{av} - P_{mv})$



# Discretisation and Verifying Closed Loop

Volume change: 3D LV (checkMesh) vs 0D compliance

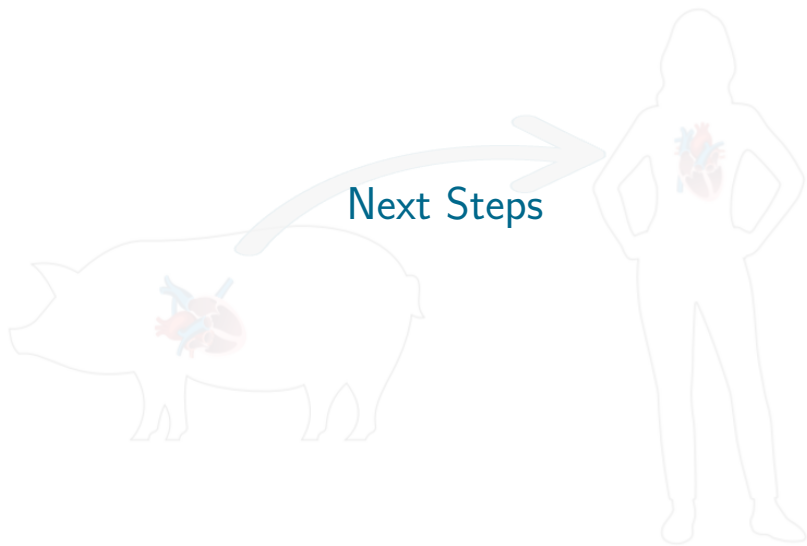


## Discretisation

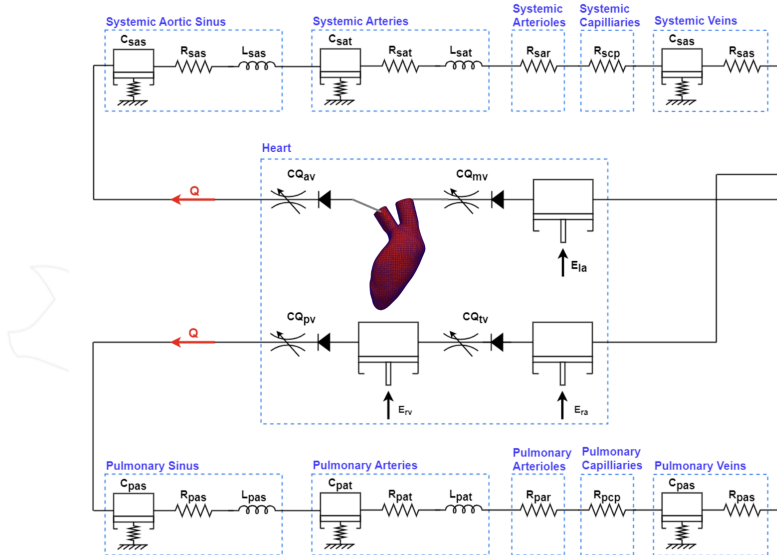
$$P_{av}^{[1]} = \frac{P_{av}^{[0]} + \frac{\Delta t}{C} \left( Q_{av}^{[1]} + \frac{P_{mv}^{[1]}}{R} \right)}{1 + \frac{\Delta t}{RC}}, \text{ initial } n = 0$$

$$P_{av}^{[n]} = \frac{4 P_{av}^{[n-1]} - P_{av}^{[n-2]} + \frac{2\Delta t}{C} \left( Q_{av}^{[n]} + \frac{P_{mv}^{[n]}}{R} \right)}{3 + \frac{2\Delta t}{RC}}$$

$$Q_{mv}^{[n]} = \frac{P_{av}^{[n]} - P_{mv}^{[n]}}{R}$$



# WIP: Fully Coupled 0D – 3D System



## Planned Work (Next 6–12 Months)

### Timeline

- **0–3 months:** complete integration of the full lumped-parameter circulation with a 3D left ventricle in closed loop; verify numerical stability and conservation properties.
- **3–6 months:** incorporate additional cardiac chambers and develop a modular OpenFOAM framework enabling interchangeable 0D and 3D heart components within the closed loop.
- **6–12 months:** introduce 3D porcine heart representations and perform initial xenogeneic simulation studies under human circulatory loading.

# References

- Hilhorst P.L.J., et al. *A Comparison of Coupling Strategies for 1D–3D Simulations of Coronary Hemodynamics* International Journal for Numerical Methods in Biomedical Engineering, 2025.
- Formaggia L., et al. *On the physical consistency between three-dimensional and one-dimensional models in haemodynamics* Journal of Computational Physics, 2013.
- Caforio, F., et al. *A coupling strategy for a first 3D-1D model of the cardiovascular system to study the effects of pulse wave propagation on cardiac function.* Computational Mechanics 2022.
- Chen, W.W., et al. *Study of cardiovascular function using a coupled left ventricle and systemic circulation model.* Journal of Biomechanics, 2016.
- Zingaro, A., et al. *A multiscale CFD model of blood flow in the human left heart coupled with a lumped-parameter model of the cardiovascular system.* Discrete and Continuous Dynamical Systems , 2022.



**Thank you!**

Happy to take questions.