

A Sphere Model for Atrial Fibrillation

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Introduction

Atrial Fibrillation (AF) is a type of cardiac arrhythmia and one of the major causes of stroke and heart failure. AF is also the most widespread cardiac condition, with over 30 millions of people worldwide suffering from it. In the UK, expenses related to this disease account for more than **1%** of the total budget of the National Health Service, which is more than **800 million pounds**. Atrial fibrillation usually manifests itself in patients affected through alterations of the normal cardiac rhythm lasting short periods of time.

Phenomenology of the Heart

► **Heartbeat Propagation:** Each heartbeat originates as an electric signal in the sinoatrial (SA) node that propagates first into the atria, then through the atrioventricular (AV) node, through Purkinje fibres, and finally from the ventricular endocardium to the ventricular epicardium (Figure 1). The electric signal is conducted in the cardiac muscle cells thanks to polarisation (change in voltage) of the cell membrane [1].

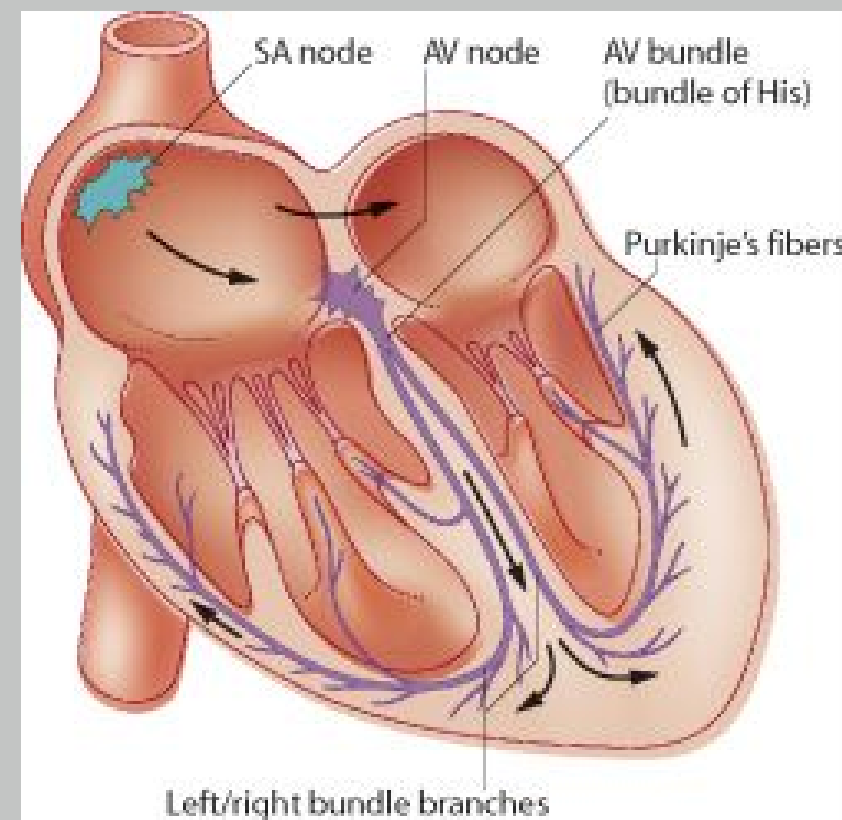


Figure 1: This is the structure of the heart.

► **Cells Excitation Rules:** As the electric impulse propagates, each cardiac single muscle cell can be in one of three stages:

- *Excitable state:* the muscle cell is at rest with a negative built up potential.
- *Excited state:* the muscle cell is excited by the excited cells it is connected to and its voltage is at its peak.
- *Refractory state:* the muscle cell goes through a phase during which it can't be excited again, a decrease in voltage—this period is called *refractory period*. The refractory period duration is related to the heartbeat rate and this relationship is called *restitution*.

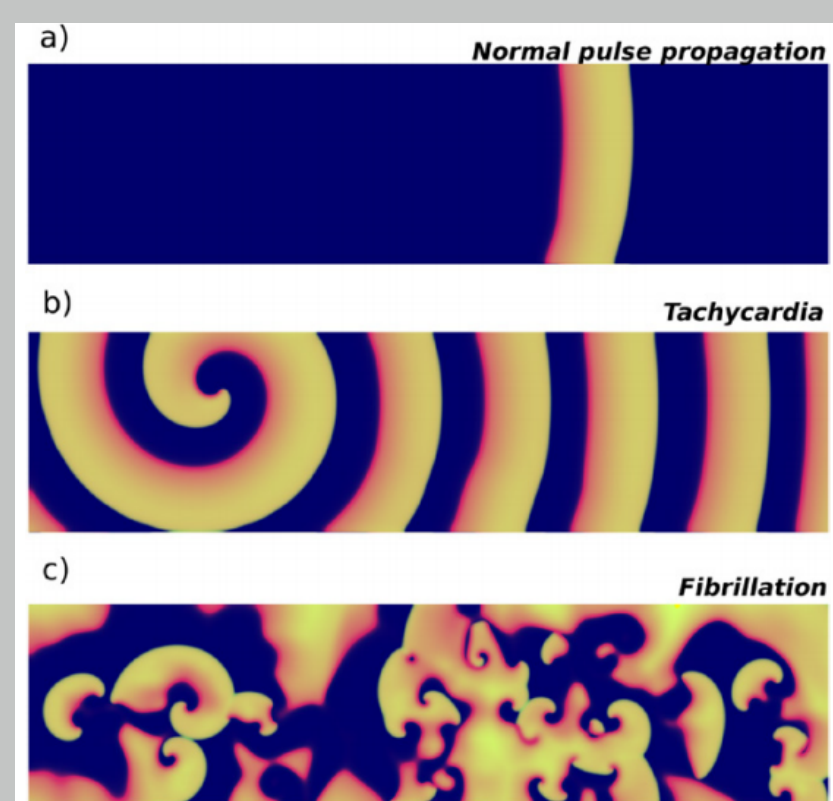


Figure 2: Idealised diagram of: (a), normal heartbeat propagation; (b), tachycardia and (c), fibrillation.

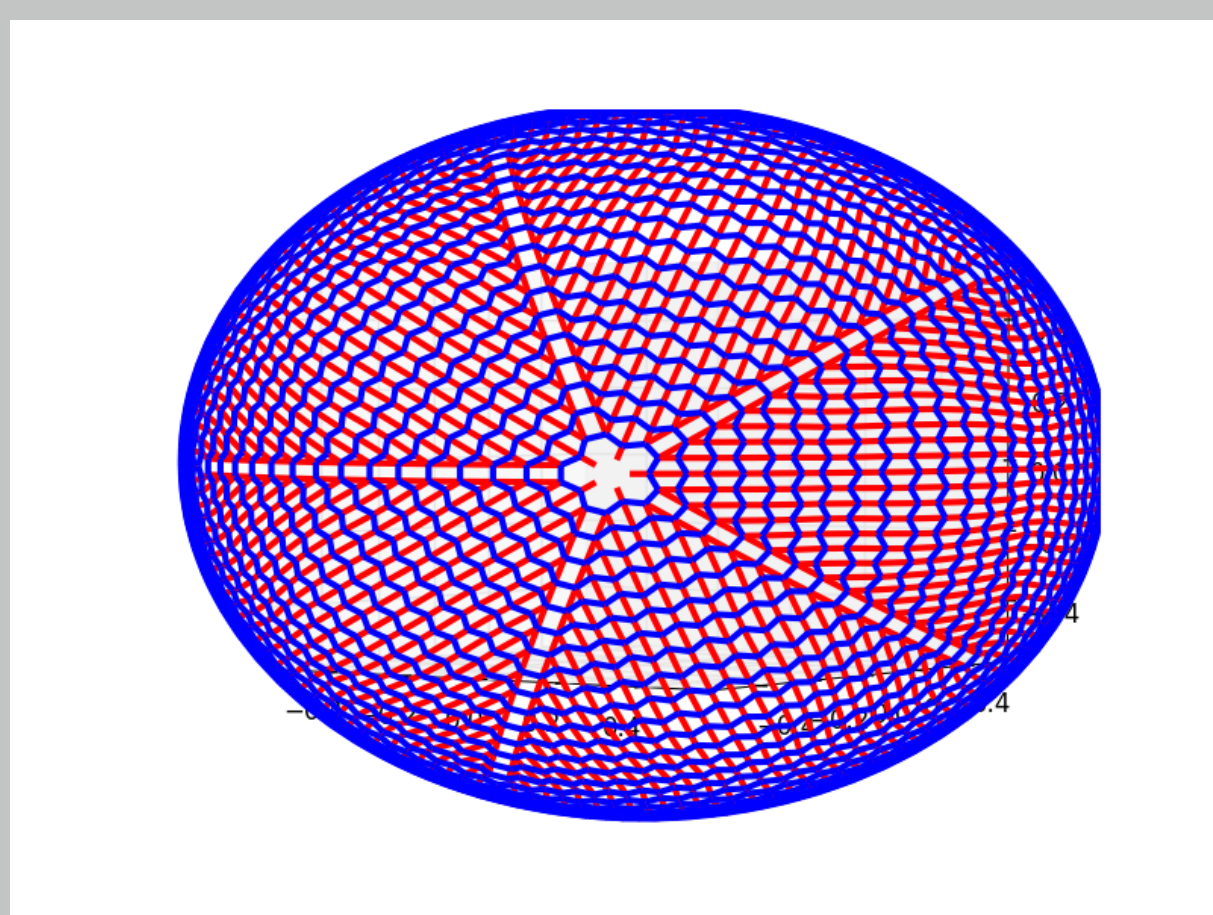
► **Base for our Research:** Our research is based on expanding on the work done by Christensen *et al.* which models the left atrium on a 2D surface [3].

► **How AF Occurs:** AF is correlated to the amount of **fibrosis**—which is the scarring of cells. This creates dysfunctional/inactive cells, decreasing the number of connections between the cells in the cardiac tissue. This promotes a process called *reentry*, during which, the electric signal wavefront propagation breaks [2]. Even though many treatments have been developed, the reentry mechanism is not completely understood and, as such, AF remains a major topic in medical research.

Our Objectives

1. Replicate the model of a 2D cardiac atrium from Christensen *et al.*
2. Translating the model onto a sphere in order to simulate the effects of a more realistic morphology.
3. Incorporate restitution of heart cells into the original model (where the *refractory period* is instead set to a constant).
4. Observe the risk of AF in the new models and compare it with the original model.

Our Sphere model's Morphology



Implementing Our Sphere Model

In research by Fedotov, a model of the atrium was constructed on the surface of a triangulated sphere to AF on a closed, heterogeneous surface [4]. Inspired by this research, we built a new model which translates Kim Christensen model of a two dimensional atrium onto a sphere.

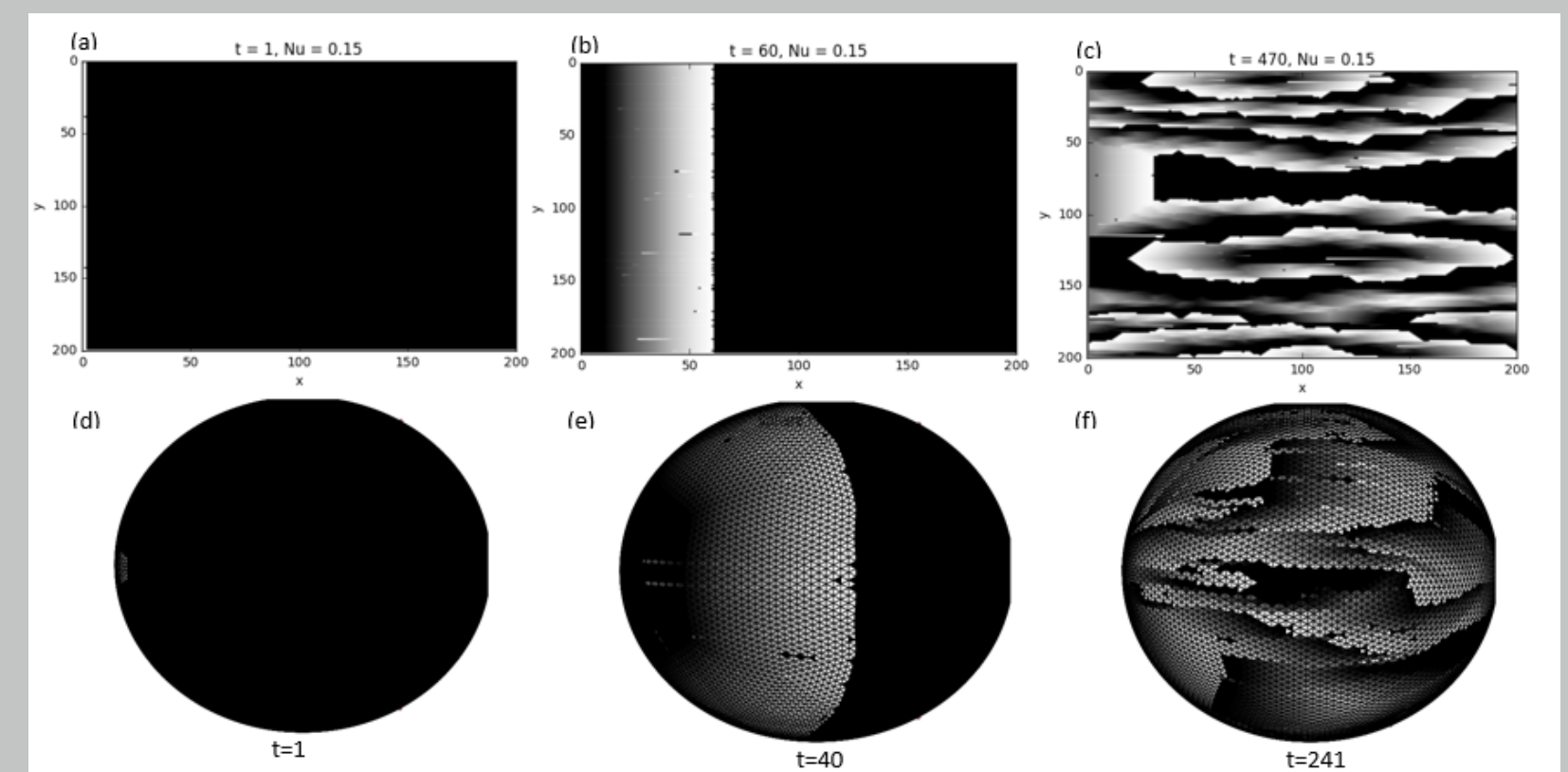


Figure 4: A comparison between Christensen *et al.*'s model (a, b and c) and the sphere model (for d, e and f) evolving in time in a not-healthy atrium. Normal wavefront propagation is seen starting in the leftmost figures. In figures b,e the excitation wavefront propagates unevenly due to fibrosis, starting reentry. On the right AF has fully arisen from both models generating chaotic fibrillatory activity.

Implementing the Restitution Model

In Christensen *et al.*'s model [3], the refractory period of each muscle cell was always fixed to a constant. In a real heart, the refractory period depends on the heart beat rate and on the last time each cell it was excited. The relationship between refractory period and heart beat rate is called restitution. We introduced a linear restitution function in Christensen *et al.*'s model in order to study its effects.

Results: Risk of AF for the three models

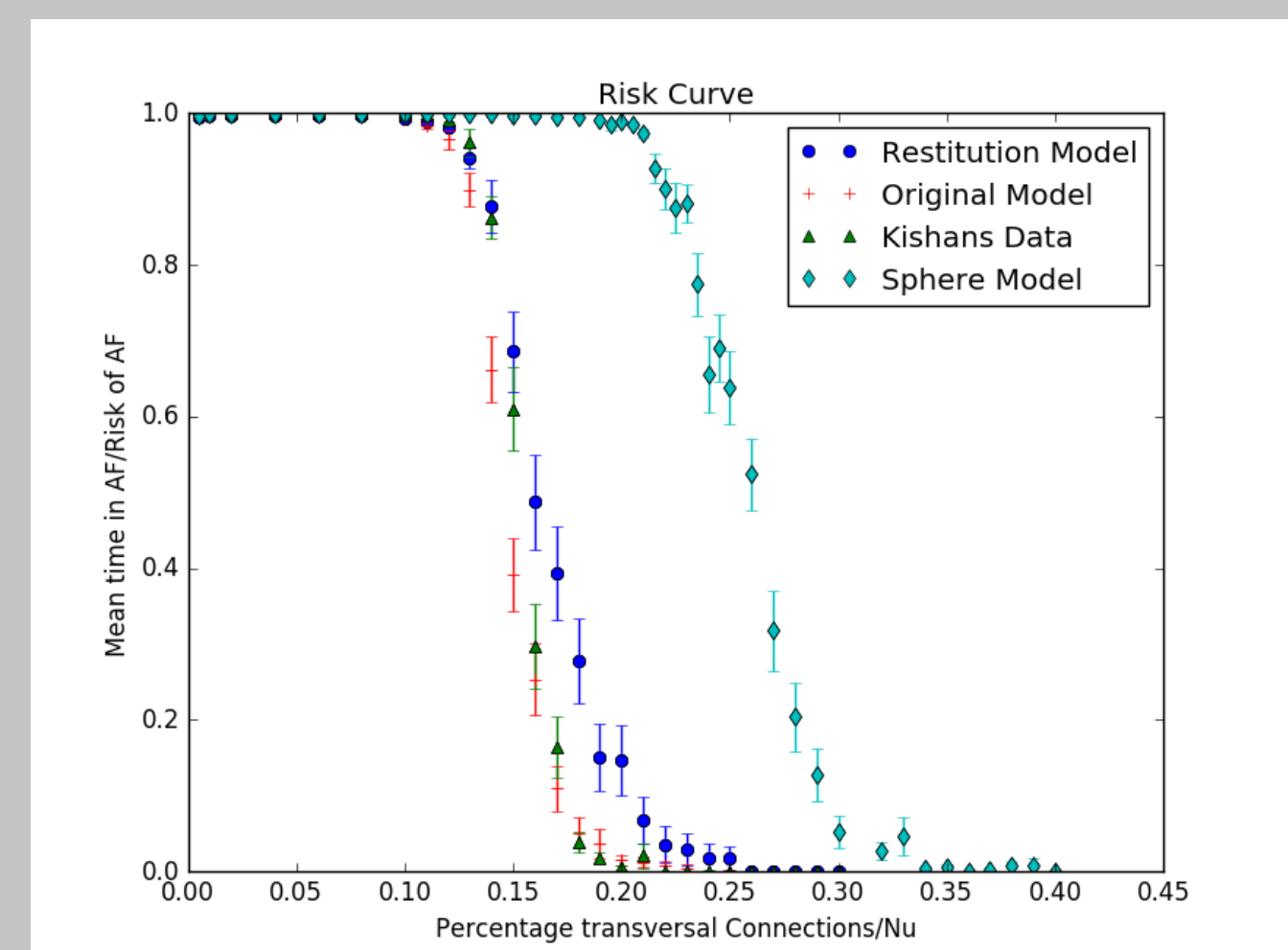


Figure 5: Risk of AF (defined as the fraction of time spent in fibrillation) for the three models studied (Prof Christensen's model, restitution curve model, sphere model) averaged over 100 simulations against percentage of transversal connections (ν) in the model. A healthy heart is expected to have ν close to 1 while a heart with fibrosis low ν .

Conclusion

A simplified mechanism of AF had been achieved by Christensen *et al.*'s model through the modelling of the atrium muscle tissue while incorporating the effects of fibrosis. We implemented this model by adding more realistic features by using a more realistic heart morphology and restitution. Our results show a higher risk of AF for the new models created.

References

- [1] P. Comtois, J. Kneller, S. Nattel, *Of circles and spirals: Bridging the gap between the leading circle and spiral wave concepts of cardiac reentry* Europace. 2005 Sep;7 Suppl 2:10-20. DOI: <http://dx.doi.org/10.1016/j.eupc.2005.05.011>
- [2] S. Alonso, M. Bar and B. Echebarria *Nonlinear physics of electrical wave propagation in the heart: a review*, Rep. Prog. Phys. 79 (2016) 096601, doi:10.1088/0034-4885/79/9/096601
- [3] Kim Christensen, Kishan A. Manani, and Nicholas S. Peters *Simple Model for Identifying Critical Regions in Atrial Fibrillation* Phys. Rev. Lett. 114, 028104 Published 16 January 2015
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