Novel Imaging of the heart for new structural and metabolic diagnosis



# Novel Imaging of the heart for new structural and metabolic diagnosis

## **Fact Sheet**

**Project Information** 

#### **ANATOMY-FOUND**

Grant agreement ID: 707663

#### DOI

10.3030/707663

## Project closed

EC signature date

4 April 2016

## Start date

1 February 2017

## End date

31 January 2019

#### **Funded under**

EXCELLENT SCIENCE - Marie Skłodowska-Curie Actions

#### **Total cost**

€ 200 194.80

#### **EU** contribution

€ 200 194,80

## Coordinated by

**AARHUS UNIVERSITET** 

**Denmark** 

# This project is featured in...



# **Objective**

Each year cardiovascular disease (CVD) causes over 4 million deaths in Europe. Predictions suggest that 80% of premature heart disease and stroke is preventable, but ~20% of CVDs are misdiagnosed, and in elderly patients up to 60% of CVDs remain undiagnosed. We need to develop new technologies and strategies for early and accurate diagnosis of the biggest killer in western society.

Currently, diagnosis of CVD is based on assessment of macro-anatomy, whole heart function, and whole body metabolic markers. Early CVD processes begin in heart cells, before evolving to the 'macro' pathology recognisable by traditional techniques. Clinicians desire a non-invasive technique that can provide high-resolution 3D micro-anatomy, and organ specific regional metabolic assessment. However the intricate micro-anatomy of the heart and its dynamic relationship with cardiac function is still fiercely debated.

To elucidate the true micro-anatomy of the heart, and understand its correlation with both pump- and metabolic function in health and disease, I will investigate the following questions:

- 1) What is the true micro-anatomy of the heart, and how is it remodelled in disease?
- 2) What is the role of 3D micro-anatomy in 4D pump function?
- 3) Does micro-anatomical remodelling correlate with regional changes in metabolism?

Using novel state-of-the-art non-invasive 3D and 4D ex-vivo and in-vivo imaging methodologies, I will answer these questions. We will finally understand the microanatomy of the heart in 3D, and its relationship with cardiac contraction and metabolism in disease. I hypothesise detecting and correlating novel morphological and metabolic changes upstream of the macro-anatomical and non-specific metabolic changes identifiable by traditional techniques, will offer the foundation for a step-wise change in diagnosis of CVD.

Working with world-renowned researchers, I will develop interdisciplinary skills in functional cardiac imaging to support a productive career in academia.

## Fields of science (EuroSciVoc) 6

medical and health sciences > basic medicine > anatomy and morphology

medical and health sciences > clinical medicine > cardiology > cardiovascular diseases

medical and health sciences > basic medicine > pathology

medical and health sciences > basic medicine > neurology > stroke

engineering and technology > medical engineering > diagnostic imaging > magnetic resonance <u>imaging</u>



## Programme(s)

H2020-EU.1.3. - EXCELLENT SCIENCE - Marie Skłodowska-Curie Actions (MAIN PROGRAMME

H2020-EU.1.3.2. - Nurturing excellence by means of cross-border and cross-sector mobility

# Topic(s)

MSCA-IF-2015-EF - Marie Skłodowska-Curie Individual Fellowships (IF-EF)

## Call for proposal

H2020-MSCA-IF-2015

See other projects for this call

## **Funding Scheme**

MSCA-IF-EF-ST - Standard EF

## Coordinator



## **AARHUS UNIVERSITET**

Net EU contribution

€ 200 194,80

Total cost

€ 200 194,80

Address

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Region

## Danmark > Midtjylland > Østjylland

Activity type

**Higher or Secondary Education Establishments** 

Links

Contact the organisation Website Medicipation in EU R&I programmes Medicipation network

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European Union, 2025