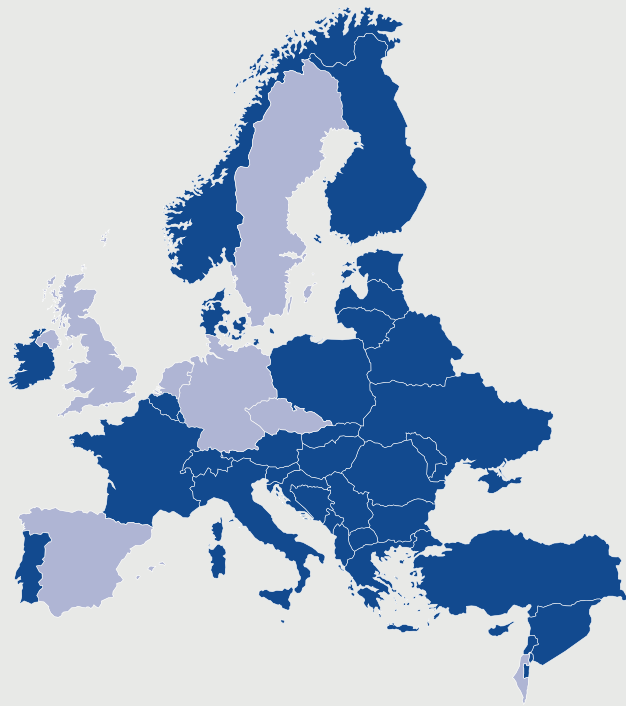


Partners & regions



Learn more



Find out more about this groundbreaking study with its extraordinary consortium of international teams, online at :



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RITA-MI2



A PHASE 2 CLINICAL STUDY

Toward an innovative and cost effective treatment for major cardiovascular events



Given the role of inflammation in heart disease, can an existing drug, CD20 mAb rituximab, be repurposed to target inflammation-based heart disease after heart attack?

At a glance :

5
YEARS
DURATION

13
PARTNERS

8
COUNTRIES

€6.5m
EUROPEAN COMMISSION FUNDING

550
PATIENTS RECRUITED

The RITA-MI 2 project is a European-funded research initiative coordinating over a dozen teams of researchers from across Europe. Its aim is to improve heart function recovery and reduce progression to heart failure after heart attack by the selective targeting of mature B lymphocytes, part of the immune system.



The problem

Cardiovascular diseases are a major cause of illness and death worldwide. Despite advances in treatment of myocardial infarction, up to 50% of patients still develop dysfunction in the heart's left ventricle, indicating 3 to 4-times higher mortality risk.



The hypothesis

The study's central idea is that using rituximab to deplete B lymphocyte cells will limit the infarct size and swelling of the heart muscle after a serious heart attack (a 'STEMI'). This should improve heart recovery and function. Additionally, the therapy will hopefully tame excessive inflammatory response and accelerated thickening of artery walls after such events.



Trial objectives

This phase 2b, placebo-controlled, doubleblind study has three main aims :



Prove that rituximab improves heart function after STEMI (a type of heart attack)

Assess impact of the drug on vascular inflammation patients



Understand immuno-pharmacology and cardio-protective mechanisms of rituximab



Anticipated impacts



Unique
no equivalent exists



Synergies
can combine existing therapies



30% reduction
in cardiovascular events



Duration
may give long term protection



Compliance
no medication to remember

Month 0
Kick-off

Month 3
Website online

Month 5
First patient, first visit

2021

Month 2
Statistical plan approved
Operating procedures setup

Month 4
Regulatory approval

Month 18
Asses single cell transcriptions
Asses mass cytometry

Month 20
140 patients reached (25%)

Month 32
280 patients reached (50%)

Month 57
Final patient, final visit (100%)

2026

Month 60
study ends