Optical coherence tomography to guide stent placement during percutaneous coronary intervention

Submission date	Recruitment status Stopped	[X] Prospectively registered		
11/03/2015		☐ Protocol		
Registration date 11/03/2015 Last Edited 04/06/2019	Overall study status Stopped Condition category Circulatory System	Statistical analysis plan		
		☐ Results		
		☐ Individual participant data		
		Record updated in last year		

Plain English summary of protocol

Background and study aims

Coronary artery disease (CAD) and its consequences are the leading cause of death and disease in the Western world, leading to major healthcare and economic burdens. CAD is the thickening of the walls of the coronary arteries that supply blood to the heart, caused by the build-up of fatty deposits in these blood vessels. Percutaneous coronary intervention (PCI) is a routine and widely used treatment to reopen narrowed or blocked coronary arteries. PCI is usually performed by feeding a hollow wire (catheter) through the blood vessels to the site of the blockage, inflating a balloon at the tip of the catheter to stretch the artery wall vessel and then placing an expandable metal tube (stent) at the site of the blockage. The stent stays in the wall of the artery in order to keep the artery wall stretched and allow blood to flow freely through the artery. Tools that can take pictures of the inside of the coronary arteries, have contributed substantially to our understanding of the how best to place a stent. Optical coherence technology (OCT) is a method for taking pictures of the inside of the coronary arteries, which provides an image with much greater resolution than current alternatives (e.g. intravascular ultrasound). The definition of the OCT images is at least ten times greater than with ultrasound, which enables very detailed imaging of the artery wall, the fatty deposits and, once placed, the stent. To date, there have been no studies of the usefulness of OCT for improving stent placement and the effectiveness of PCI. In this study we wish to assess whether routine use of OCT to guide stent placement will lead to improved stent placement.

Who can participate?

Patients aged 30 to 90 who are undergoing PCI for the treatment of CAD or acute coronary syndrome

What does the study involve?

Participants are randomly allocated to undergo either angiographically guided or OCT-guided stent placement. Angiographically guided stent placement is the standard care pathway, where a type of X-ray is used to examine the blood vessels. OCT-guided stent placement involves

taking pictures of the inside of the coronary arteries to guide decisions about the size and type of stent to use. All patients receive standard hospital follow up. 12 months after their procedure all patients complete a questionnaire.

What are the possible benefits and risks of participating? Not provided at time of registration

Where is the study run from? University of Bristol (UK)

When is the study starting and how long is it expected to run for? March 2014 to May 2018

Who is funding the study? National Institute for Health Research (UK)

Who is the main contact? Lucy Culliford lucy.culliford@bristol.ac.uk

Contact information

Type(s)

Scientific

Contact name

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Additional identifiers

Protocol serial number 18505

Study information

Scientific Title

Optical Coherence Tomography to guide stent placeMent during percutaneouS coronary intErvention: a randomised controlled trial

Acronym

OCTIMISE

Study objectives

The aim of this study is to assess whether routine use of optical coherence technology (OCT) to guide stent placement will lead to improved stent placement and improved clinical outcomes at one year for people with coronary heart disease (CHD).

Ethics approval required

Old ethics approval format

Ethics approval(s)

West Midlands -Coventry and Warwick NRES committee, 25/02/2015, ref: 15/WM/0075E

Study design

Randomised; Interventional; Design type: Treatment

Primary study design

Interventional

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Topic: Cardiovascular disease; Subtopic: Cardiovascular (all Subtopics); Disease: Cardiovascular

Interventions

Following the consenting process the intervention begins in the Catheter laboratory:

1. Angiographically guided stent placement: This is the standard care pathway, the patient has their stent placed using standard treatment plan, once the clinician has stated that they have completed deployment the OCT catheter is taken past the stent(s,) this is known as a 'pullback', to visually record the inside of the lumen. The procedure ends. The patient has standard care and is discharged.

Duration is approximately 5 minutes longer than standard care.

2. OCT guided stent placement: After the standard angiogram and review of the lesions, the OCT catheter is used to visualise the inside of the lumen, the clinician uses the images to guide their decisions about size and type of stent used. The catheter is used again to check stent placement, the clinician makes any adjustments or corrections to the stent placement and if necessary another pullback is made to check the stent. Once the clinician is content, that the deployment is complete, a reference pullback occurs.

The procedure ends. The patient has standard care and is discharged. Duration is approximately 10 minutes longer than standard care. All patients receive standard hospital follow up. 12 months after their procedure all patients will receive a trial questionnaire.

Intervention Type

Procedure/Surgery

Primary outcome(s)

To estimate the difference in minimal luminal area (MLA) immediately after the completion of PCI between the OCT-guided and the angiographically guided PCI groups. This outcome will be assessed by processing the digitally stored OCT images.

Key secondary outcome(s))

In the OCT-guided and angiographically guided groups:

- 1. To quantify and compare the prevalence of strut apposition, edge dissection and plaque prolapse. This will be determined from the images taken to assess the primary endpoint
- 2. To compare the nature and extent of additional interventions indicated from OCT images of the deployed stent, such as additional balloon dilations or further stenting
- 3. To quantify and compare the PCI procedure duration, the volume of contrast used and the X-ray dosage
- 4. To quantify and compare the number of major clinical events at 12 months follow-up

In OCT-guided group only:

Information about how the pre-deployment OCT images informed the choice of treatment strategy and stent

Completion date

01/05/2018

Reason abandoned (if study stopped)

Objectives no longer viable

Eligibility

Key inclusion criteria

A participant may enter study if ALL of the following apply:

- 1. Adults of either sex aged 30 to 90 years who are undergoing EITHER,
- 1.1. Elective PCI for the treatment of CAD OR
- 1.2. Urgent PCI for the treatment of ACS
- 2. Patients who are willing and able to give written informed consent for participation in the study
- 3. Patients who are anticipated to have at least a 20 mm stent length in at least one lesion

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

All

Key exclusion criteria

A participant may not enter study if ANY of the following apply:

- 1. Patients unable to provide written informed consent.
- 2. Patient undergoing emergency PCI
- 3. Patients that have had previous PCI within the target vessel.
- 4. Women who are pregnant or breast feeding
- 5. Haemodynamic instability
- 6. Renal impairment (eGFR =50 ml/min)
- 7. Multivessel PCI (>2 vessels) during a single procedure
- 8. Left main stem disease
- 9. Estimated angiographic vessel calibre <2.5mm or >4.5mm
- 10. Additional significant disease (>50% stenosis) in same vessel proximal or distal to culprit lesion
- 11. Target vessels / lesions of excessive tortuosity
- 12. Patients participating in another interventional study

Date of first enrolment

01/05/2015

Date of final enrolment

01/05/2017

Locations

Countries of recruitment

United Kingdom

England

Study participating centre Bristol Heart Institute

Bristol United Kingdom BS2 8HW

Sponsor information

Organisation

University Hospitals Bristol NHS Foundation Trust

ROR

https://ror.org/04nm1cv11

Funder(s)

Funder type

Government

Funder Name

National Institute for Health Research

Alternative Name(s)

National Institute for Health Research, NIHR Research, NIHRresearch, NIHR - National Institute for Health Research, NIHR (The National Institute for Health and Care Research), NIHR

Funding Body Type

Government organisation

Funding Body Subtype

National government

Location

United Kingdom

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Available on request

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
HRA research summary			28/06/2023	No	No
Participant information sheet	Participant information sheet	11/11/2025	11/11/2025	No	Yes