

Collection of Information on the long-term results of treatment with the Supraflex™ Cruz drug-releasing blood vessel scaffold

Submission date	Recruitment status	<input type="checkbox"/> Prospectively registered
01/07/2020	No longer recruiting	<input type="checkbox"/> Protocol
Registration date	Overall study status	<input type="checkbox"/> Statistical analysis plan
01/07/2020	Completed	<input checked="" type="checkbox"/> Results
Last Edited	Condition category	<input type="checkbox"/> Individual participant data
29/10/2024	Circulatory System	

Plain English summary of protocol

Background and study aims

Coronary heart disease (CHD), also known as ischemic heart disease, is one of the leading causes of death worldwide. CHD develops because of the build-up of fatty deposits (plaque) on the walls of the coronary arteries (the arteries that supply the heart with oxygen-rich blood). When arteries are blocked or narrowed, the heart does not receive enough blood to function properly, which can cause pain and tightness in the chest (angina), or a heart attack. An angioplasty is a common procedure where a thin tube (catheter) is placed inside the narrowed blood vessel. A small balloon on the tip of the catheter is gradually inflated to reopen the artery and flatten the blockage against the artery wall. In some cases, the surgeon also places a mesh-like tube (stent) into the artery to keep it open. One of the main problems following this type of surgery is that the artery may re-narrow and become blocked again (restenosis), as the cells which make up the obstruction multiply (proliferate). New techniques have been developed where the stent placed in the artery is coated in a drug which prevents cell proliferation (drug-eluting stent). The Supraflex™ Cruz Stent is a type of drug-eluting stent which uses the drug sirolimus to prevent restenosis. The aim of this study is to look at patients who have had a Supraflex™ Cruz Stent to find out if it is a safe and efficient way of preventing restenosis and future heart problems.

Who can participate?

Adults who have had a Supraflex™ Cruz Stent implanted as part of their clinical care.

What does the study involve?

Participants who are having a Supraflex™ Cruz Stent implanted are asked for their consent to take part in the study. The patients are asked to attend a follow-up appointment 12 months after their operation so that any blockages in their stent (stent thrombosis) can be measured, using an angiogram (a scan of the coronary arteries). The patients are also interviewed in order to find out whether there have been any problems since their operation.

What are the possible benefits and risks of participating?

There are no direct benefits or risks to patients taking part in this study

Where is the study run from?

Freeman Hospital (lead centre) and around 30 other hospitals in the UK

When is the study starting and how long is it expected to run for?

April 2018 to January 2023

Who is funding the study?

Sahajanand Medical Technologies Pvt. Ltd (India)

Who is the main contact?

Manoj Virupil, manoj@smt.in

Contact information

Type(s)

Public

Contact name

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Contact details

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Additional identifiers

Clinical Trials Information System (CTIS)

Nil known

Integrated Research Application System (IRAS)

246372

ClinicalTrials.gov (NCT)

Nil known

Protocol serial number

IRAS 246372

Study information

Scientific Title

Prospective evaluation of thin-strut biodegradable polymer-coated Supraflex sirolimus-eluting stents in an all-comers patient population (S-FLEX UK-II)

Acronym

S-FLEX UK-II

Study objectives

To evaluate the clinical safety and performance of thin-strut biodegradable polymer-coated Supraflex Cruz sirolimus-eluting stents in an all-comers patient population requiring stent implantation.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Approved 22/08/2019, South West - Cornwall & Plymouth REC (Level 3, Block B, Whitefriars, Lewins Mead, Bristol, BS1 2NT, UK; +44 (0)207 104 8214; cornwallandplymouth.rec@hra.nhs.uk), ref: 18/SW/0130

Study design

Multi-centre observational cohort study

Primary study design

Observational

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Atherothrombosis

Interventions

Once a patient has been identified as potentially meeting the inclusion and exclusion criteria written informed consent is obtained prior to the patient's enrollment. Potential participants will have up to 24 hours to consider the information presented to them, after which signed consent will be sought. It should be noted that the patient would have already provided consent for the DES implantation procedure as per normal standard of care at the investigational site.

The total duration of observation is 12 months with a follow up at 12 months.

The total duration of the follow-up at 12 months is no longer than 1 hour.

Intervention Type

Other

Primary outcome(s)

The safety and efficacy of the Supraflex Cruz stent measured through patient interviews and the review of medical notes at the 12 month follow up

Key secondary outcome(s)

Measured using a review of patient records:

1. Procedural endpoints [at the end of the index procedure (12 months)]
 - 1.1. Device success, lesion success, procedural success
2. Safety and efficacy endpoints [30 days and 12 months]
 - 2.1. Overall stent thrombosis

- 2.2. All deaths (cardiac, vascular and non-cardiovascular)
- 2.3. Any myocardial infarction (Q wave and non-Q wave MI)
- 2.4. Any repeat revascularization (target lesion and target vessel revascularization)
- 2.5. Target Lesion Failure (TLF)
- 2.6. Target Vessel Failure (TVF): a composite endpoint of cardiac death, target vessel myocardial infarction and clinically driven target vessel revascularization

Completion date

10/01/2023

Eligibility

Key inclusion criteria

- 1. Aged 18 years or older
- 2. Symptomatic coronary disease
- 3. Clinical indication for PCI and stenting of at least one coronary lesion visually confirmed on coronary angiography
- 4. Only Supraflex Cruz stent(s) is/are to be implanted into the coronary vasculature during the index procedure
- 5. Give informed consent to participate in this registry and sign the informed consent form approved by the institutional review board of each registry site before the index PCI
- 6. Agree to undergo all clinical follow-up procedures specified in the study protocol (S-FLEX UK-II) for this registry

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Lower age limit

18 years

Sex

All

Total final enrolment

1904

Key exclusion criteria

- 1. Cardiogenic shock.
- 2. Pregnant or possibly pregnant
- 3. Known hypersensitivity or contraindication to aspirin, clopidogrel, ticlopidine, heparin or any other anticoagulation / antiplatelet therapy required for PCI, cobalt chromium, sirolimus or contrast media
- 4. Scheduled to undergo elective surgery within 12 months post-index PCI
- 5. Currently participating in a clinical study of another drug or medical device and in whom

assessment of the primary endpoint of that study has not been completed or clinically interferes with the endpoints of this registry

6. Mental incapacity, unwillingness or language barrier precluding understanding of the registry procedure or cooperation with registry site personnel

Date of first enrolment

10/03/2020

Date of final enrolment

30/09/2021

Locations

Countries of recruitment

United Kingdom

England

Wales

Study participating centre

Freeman Hospital

Freeman Road

High Heaton

Newcastle Upon Tyne

United Kingdom

NE7 7DN

Study participating centre

University Hospital of Wales - Cardiff

Heath Park

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United Kingdom

CF14 4XW

Study participating centre

Royal Berkshire Hospital

London Road

Reading

United Kingdom

RG1 5AN

Study participating centre

Bedford Hostpital

King's Place
Britannia Road
Bedford
United Kingdom
MK42 9DJ

Study participating centre

Wigan Hospital
Brick Kiln Lane
Wigan
United Kingdom
WN1 1XX

Sponsor information

Organisation

Sahajanand Medical Technologies Pvt. Ltd (SMT)

Funder(s)

Funder type

Industry

Funder Name

Sahajanand Medical Technologies Pvt. Ltd (SMT)

Results and Publications

Individual participant data (IPD) sharing plan

The current data sharing plans for this study are unknown and will be available at a later date.

IPD sharing plan summary

Data sharing statement to be made available at a later date

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	safety and performance	22/10/2024	29/10/2024	Yes	No
HRA research summary			28/06/2023	No	No

