STUDY PROTOCOL

A study protocol for a randomized controlled trial on the prevention of atrial fibrillation after coronary artery bypass grafting surgery using Tocotrienol, an isomer of Vitamin E derived from palm oil [version 1; referees: awaiting peer review]

Ahmad Farouk Musa[^1], Jeswant Dillon[^2], Mohamed Ezani Md Taib[^2], Alwi Mohamed Yunus[^2], Rusli Nordin[^1], Yuen Kah Hay[^3]

[^1]: Jeffrey Cheah School of Medicine and Health Sciences, Monash University Malaysia, Bandar Sunway, Selangor, Malaysia
[^2]: Department of Cardiothoracic Surgery, National Heart Institute, Kuala Lumpur, Malaysia
[^3]: School of Pharmaceutical Sciences, Universiti Sains Malaysia, Penang, Malaysia

Abstract

**Background:** One of the most common complications following coronary artery bypass grafting (CABG) surgery is atrial fibrillation (AF), which contributes towards increasing morbidity and mortality, length of hospital stay (LoHS) and reduced quality of life (QoL) of patients.

**Objectives:** To determine whether the intake of Tocotrienol, a Vitamin E isomer derived from palm oil, before and immediately following CABG prevents AF, reduces LoHS, and improves the QoL of patients.

**Protocol:** The study is registered with the National Medical Research Register Protocol: with a trial number NMRR-17-1994-34963 and designed as a prospective, randomized controlled trial (RCT) with parallel groups. The experimental group will receive two 200mg Tocotrienol capsules each day, while the control group will receive two identical placebo (palm Super Olein) capsules per day. ECG readings will be used to detect AF post operatively, LoHS will be measured by checking the records from the National Heart Institute Hospital register, and the health-related Quality of Life (HRQoL) analysis (the Malay version of the Short Form 36 Questionnaire) will be used to analyse QoL. The sample size was calculated to be 140 in each arm of the RCT for a power of 0.8 and a significance level of 0.05.

**Funding:** HOVID Berhad funds this research project.

**Expected outcomes:** The primary endpoint is the development of postoperative AF, whilst the secondary endpoints are the LoHS and HRQoL of patients post CABG.

**Future implications:** Prevention of AF and its complications such as cardiovascular or cerebrovascular events, especially stroke, is an important output. Malaysia is one of the biggest producers and exporters of palm oil and palm oil products. Thus, the possibility of marketing Tocotrienol, in reducing AF post CABG surgery, is a very important proposition indeed.

**Trial number:** NMRR-17-1994-34963
Corresponding author: Ahmad Farouk Musa (farouk@monash.edu)

Author roles: Musa AF: Conceptualization, Data Curation, Formal Analysis, Funding Acquisition, Investigation, Methodology, Project Administration, Resources, Software, Supervision, Validation, Visualization, Writing – Original Draft Preparation, Writing – Review & Editing; Dillon J: Project Administration, Resources, Supervision, Validation, Visualization; Md Taib ME: Project Administration, Resources, Validation, Visualization; Mohamed Yunus A: Project Administration, Resources, Validation, Visualization; Nordin R: Project Administration, Supervision, Validation, Visualization, Writing – Original Draft Preparation, Writing – Review & Editing; Kah Hay Y: Funding Acquisition, Project Administration, Resources, Supervision, Visualization

Competing interests: No competing interests were disclosed.

How to cite this article: Musa AF, Dillon J, Md Taib ME et al. A study protocol for a randomized controlled trial on the prevention of atrial fibrillation after coronary artery bypass grafting surgery using Tocotrienol, an isomer of Vitamin E derived from palm oil [version 1; referees: awaiting peer review] F1000Research 2018, 7:215 (doi: 10.12688/f1000research.12912.1)

Copyright: © 2018 Musa AF et al. This is an open access article distributed under the terms of the Creative Commons Attribution Licence, which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

Grant information: HOVID Berhad (contact information: 121, Jalan Tunku Abdul Rahman, 30010 Ipoh, Perak, Malaysia) sponsored this study (grant number, MMRD-MS-1801, awarded to the principle investigator, Ahmad Farouk Musa). The study design is solely developed by the main investigator. The sponsor, HOVID Berhad, supplies the materials for the research and the financial support. The funders had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript.

Introduction

Atrial fibrillation (AF) after coronary artery bypass grafting (CABG) surgery is common in clinical practice, with an incidence of up to 70%. It has the potential to double the risk of mortality and is associated with a 6-fold increase in the risk of stroke.

In a recent local study by Musa et al. on post-CABG patients, it was demonstrated that patients who developed postoperative AF had a prolonged Intensive Care Unit (ICU) stay and High Dependency Unit (HDU) stay, and a prolonged total hospital stay. This leads to an increase in resource utilization. With an increased rate of morbidity and mortality, preoperative prophylactic strategies are necessary to reduce the incidence of AF post-operatively and improve the Quality of Life (QoL) of patients.

The financial burden in managing AF is huge, and many new innovative ideas and materials have been tested in order to reduce the occurrence of post-operative AF. One of the very promising but not widely studied treatment alternatives is the use of Tocotrienol, an isomer of vitamin E.

AF is thought to be initiated by rapid electrical activity arising from the muscular sleeves of pulmonary veins. Multiple re-entrant wavelets, shorter refractory periods, conduction delays and fragmented electrograms maintain the arrhythmia. If this arrhythmia continues, electrical remodeling occurs, which will further facilitate AF maintenance. The pathogenesis of post-operative AF is believed to be multifactorial. Previously it was thought to be caused by inflammatory pathways, but now it is known that oxidative stress plays a significant role. Reactive oxygen species (ROS) are the cause of oxidative stress, and generate numerous pathological processes such as DNA damage, apoptosis, and cellular hypertrophy, as well as signal pathway modulation.

Cardiac surgery is characterized by ischaemia and reperfusion injury, which leads to the release of ROS. This causes oxidative stress and initiates a systemic inflammatory response. The extracorporeal circulation during on-pump surgery also activates inflammatory cells, increasing the oxidative stress. Therefore it was always thought that an off-pump surgery would reduce the incidence of AF.

Oxidative stress causes the depletion of endogenous antioxidants, and this results in oxidative damage. This oxidative stress is mainly the result of on-pump surgery that requires aortic cross-clamping and the bypass machine.

Almost all cardiac centers have guidelines on the pharmacological management of postoperative AF. But the challenge has always been how to prevent AF from happening. Efforts have been made to develop alternative preventive strategies. With the recent understanding of the pathogenesis of AF via the oxidative pathway, perhaps antioxidant vitamins would be a promising treatment.

Previous data have shown that both Vitamin E and C are known to have antioxidant properties, and have the ability to counter the action of free radicals. Vitamin E is able to maintain membrane stability. It could also prevent the process of lipid peroxidation. Vitamin C removes water-soluble free radicals and acts synergistically with Vitamin E.

There have been numerous mechanistic studies on the role of antioxidant vitamins in the prevention of postoperative AF. However, the therapeutic potential is uncertain and there is no universally accepted protocol. It is interesting to note that Carnes et al. reported a 16.3% incidence of postoperative AF in Vitamin C-treated patients compared to 34.9% in controls in their prospective study. In their meta-analysis, Harling et al. demonstrated a reduction in the incidence of postoperative AF and other forms of arrhythmia following antioxidant vitamin therapy, which remained significant when only randomized controlled trials (RCTs) were analyzed. They also showed that this reduction in incidence was not related to the bypass or cross-clamp time, where there was no significant difference between the two-groups. Lastly, the results of the meta-analysis comparing the groups of anti-oxidants and control also demonstrated a reduced stay at the ICU and at the hospital. While this finding cannot be attributed solely to a reduction in postoperative AF, it is possible that this reduction in both ICU and hospital stay was a result of the reduced incidence of AF in the treatment group.

Rasoli et al. came to a similar conclusion that the use of antioxidant vitamins plays a role in the reduction of postoperative AF. However, they argued that the effect was somewhat variable, because both vitamins, C and E, were weak anti-oxidants. They were limited in their capacity to cross the cell membrane in order to counter the superoxides.

We are proposing a more potent isomer of vitamin E, namely Tocotrienol, which is derived from palm oil, instead of the usual isomer of vitamin E, namely tocopherol, which was being used in all previous research. Tocotrienol has been proven to possess more potent antioxidant activity than tocopherols. Laboratory studies have proven that Tocotrienol has the ability to scavenge peroxyl radicals in liposomes. Tocotrienol is seen to be more evenly distributed in the phospholipid bilayer of the plasma membrane. It is also shown to have more efficient collision with radicals. These are some of the reasons why Tocotrienol is thought to have a more potent antioxidant properties.

All these recent discoveries suggest the promising antioxidant properties for Tocotrienol. We hypothesize that it would be able to reduce the incidence of postoperative AF and subsequently reduce the complications associated with postoperative AF, along with the economic burden of prolonged intensive care stay and prolonged hospital stay. Overall, we also hypothesized that Tocotrienol will increase the health related quality of life (HRQoL) of patients. The potential value of marketing a national product that can be produced locally from palm oil and its potential uses, not only for patients, who are undergoing CABG, but also for patients with standalone AF, makes this an even more attractive product.
Protocol

Ethical statement

The National Heart Institute Ethics Committee has approved the study protocol (IJNREC/20112017). The ethics committee will also serve as the data safety committee. Similar approval has been obtained from MUHREC - Monash University Human Research Ethics Committee – (9277) and from MREC – the Malaysian Research Ethics Committee. The study is registered with the National Medical Research Registry (NMRR), Ministry of Health, Malaysia, (NMRR-17-1994-34963) and will be carried out in accordance with the ethical principles outlined in the Malaysian Good Clinical Practice Guidelines.

Written informed consent for participation in the trial will be obtained from each participant on recruitment (Supplementary File 1).

The clinical data of patients enrolled in this study will be de-identified. No personal data will be disclosed to anyone outside of the National Heart Institute Ethics Committee.

This is Protocol Version 2.0 dated on 15 June 2017.

Trial registration: NMRR-17-1994-34963

Trial registry: National Medical Research Register, Malaysia

Trial registration date: 13th March 2017

Study status: The study has not yet started at the time of submission of this manuscript since we are still waiting for the Clinical Trial Exemption (CTX) from the National Pharmaceutical Regulatory Agency (NPRA).

Hypothesis

Tocotrienol is a commercial product produced by Hovid Berhad, and is marketed as Tocovid Suprabio, containing 61.52mg alpha-Tocotrienol, 112.80mg gamma-Tocotrienol, 25.68mg delta-Tocotrienol and 91.60IU alpha-tocopherol. Its availability in Malaysia has allowed this innovative and promising study to be conducted. We hypothesize that:

1. Intake of Tocotrienol capsules before and immediately following CABG prevents AF post CABG;
2. Intake of Tocotrienol capsules before and immediately following CABG improves the QoL of patients post CABG;
3. Intake of Tocotrienol capsules before and immediately following CABG shortens the length of hospital stay (LoHS) of patients post CABG.

Objectives

To determine whether the intake of Tocotrienol capsules before and immediately following CABG prevents AF post CABG.

To determine whether the intake of Tocotrienol capsules before and immediately following CABG improves the QoL of patients post CABG.

To determine whether the intake of Tocotrienol capsules before and immediately following CABG shortens the LoHS of patients post CABG.

Study design

Study site: National Heart Institute, Kuala Lumpur, Malaysia

The study is designed as a prospective, randomized controlled trial, with two parallel groups of the same demographics and comorbidities. The main aim is to look at the effect of Tocotrienol in reducing AF post CABG.

Patients who are scheduled for CABG under the co-researchers at the National Heart Institute will be approached for their consent to be enrolled in this study once they are admitted to the wards. We will assign the patients enrolled in the study to one of two groups:

1. The control group: Routine CABG surgery procedure plus placebo containing palm Super Olein capsules (supplier: HOVID Berhad, Malaysia). The placebo is an identical-looking capsule that contains tocotrienols-stripped palm oil (termed palm Super Olein), which is also the non-active excipient in the treatment capsule. The choice of control is to mimic the treatment capsule as much as possible, without the active component of tocotrienols.

2. The study group: Routine CABG surgery procedure plus Tocotrienol capsules (supplier: HOVID Berhad, Malaysia)

For patients in the study group, Tocotrienol will be administered as two capsules of 200mg per day in two divided doses, and administration will start immediately after randomization and will continue until the first follow-up, which is normally six weeks after discharge.

Since there is no available data for preferred dose of treatment, we decided on two divided doses of Tocotrienol of one capsule twice daily of 200mg Tocovid. This dosage was estimated based on the regime used by Olaf Stanger et al.17 at the Department of Cardiac Surgery, Paraceles Medical University Salzburg, Austria, where three ampoules of 45IE Vitamin E were used, with 30mg per ampoule or 90mg in total. Because in our study we would be using an oral preparation instead of an IV preparation as in the Austrian study, and absorption of Tocotrienol has been shown to be low and incomplete via the oral route, we estimated a higher dosage18. Considering the bioavailability of Tocotrienol can be as low 10–30% if administered orally18; and taking into consideration that this is a pivotal study, it is reasonable to use the highest dose possible that is safe without any adverse effects. Many clinical studies with Tocotrienol have used 400mg daily in two divided doses and have been proven to be safe19,20.

Tocovid will be administered right after randomization, and will continue after the patient is discharged until the first follow-up, which is normally about six-weeks later. If the patient is on prolonged ventilation in the ICU, Tocotrienol will be administered through a nasogastric tube. The intensive care nurses and cardiothoracic ward nurses will monitor compliance.

All patients undergoing CABG surgery, either or with valve surgery, will be included. Similarly, this study will include...
participants undergoing both on-pump surgery using cold potassium cardioplegia, and off-pump beating heart surgery.

All patients will be admitted to the intensive care unit (ICU) after CABG surgery, with close monitoring on one-patient-one-nurse basis. They will then be transferred to a monitored high dependency unit (HDU) if their condition is stable, while some might be transferred straight back to the ward. Continuous rhythm monitoring will be performed using the 12-lead ECG for all patients in ICU and HDU. The monitoring will be continued for the first four to five postoperative days on the normal cardiothoracic wards using Holter monitors until the patients are discharged.

We will review the electrocardiographic (ECG) data on a daily basis. We will also review all the printouts of all abnormal rhythms, which will then be included in the clinical records. Additionally, an ECG will also be recorded when patients are symptomatic or when there is a suspicion of arrhythmia clinically. We will treat all AF episodes, in both the study arm and control arm, according to the protocols of the Cardiothoracic Department. The patients will be managed by the cardiothoracic surgeon and their team. The first-line drug used for treatment of new onset AF post-operatively by the Cardiothoracic team in the National Heart Institute (IJN) Hospital is Amiodarone, unless contraindicated, as a 300mg infusion over one hour to be followed by 900mg over the next 23 hours.

Patients of both the study and control arms will come back for follow-up at the Cardiotoracic Clinic at the IJN usually six weeks after discharge. They will also be advised to report to the Outpatient Department at the IJN if they develop any symptoms. During follow-up, all patients will be assessed routinely with blood work and a 12-lead ECG.

For the study flowchart, see Figure 1.

Hovid Berhad follow strict GLP Guidelines in the manufacturing process of Tocotrienols and matching placebo capsules. Each 200mg capsule of Tocotrienol will contain 61.52mg alpha-Tocotrienol, 112.80mg gamma-Tocotrienol, 25.68mg delta-Tocotrienol and 91.60IU alpha-tocopherol. The placebo capsules will contain palm Super Olein oil.

Randomization

The study is designed as a randomized controlled trial (RCT), where patients with coronary artery disease requiring CABG surgery at the IJN will be prospectively and randomly divided into two parallel groups by means of computer generated numbers using Excel 2016 (Microsoft, Redmont, WA, USA) in a block of 10 experimental to 10 matching controls at enrollment. The experimental group will receive Tocovid capsules whilst the control group will receive identical placebo capsules containing palm Super Olein oil.

The unblinded pharmacist will generate the allocation sequence to assign the participants to the interventions. Surgeon investigators will enroll the participants. The participants will then be assign randomly to either the study arm or the control arm based on the sequence assigned to them.

Inclusion/exclusion criteria

Inclusion criteria:
1. Males or females
2. More than 18 years of age
3. Elective, on-pump or off-pump CABG surgery of coronary artery revascularization, single or double procedure

Exclusion criteria:
1. Less than 18 years of age
2. Refusal to have surgery
3. Urgent or emergency surgery
4. Inability to give informed consent
5. Documented allergy to palm oil

Patients will be recruited from those who are scheduled electively for surgery under the Consultant Surgeons, JD, MEMT, and AMY, who are co-researchers in this project. All public and private patients are eligible to be enrolled in this study. Consent will be taken upon admission to the wards.

Patients can choose to withdraw from the study at any time. Subjects may be withdrawn if the investigator deems that it is detrimental or risky for the subject to continue. The patients who withdraw will not be replaced.

Study endpoints

As the primary study endpoint, we will look at the development of AF post-surgery. This will be documented with ECG. We take a 30 second duration as a cut-off point and define AF as when there is a loss of p-waves and irregular ventricular rate or a confirmed atrial flutter. The secondary endpoint would include the LoHS after surgery, which will be obtained from the IJN registry, and the HRQoL. Three measurements will be used to determine the LoHS:

1. Total number of days that the patients stay in the Intensive care unit (ICU)
2. Total number of days in the high-dependency unit (HDU); and
3. Total number of days in the hospital, overall.

We will measure the HRQoL of patients using the validated Malay and English versions of Short-Form 36 Questionnaires (SF-36). Since the researchers are blinded to the trial, we will be evaluating all endpoints, primary and secondary, independently. We will also review the patients’ clinical records and all ECG tracings. In fact, all the trial participants, care-givers, outcome assessors and data analysts will be blinded to the study treatment allocation. An assigned pharmacist maintaining the randomisation list
Figure 1. Study flowchart.
will be unblinded to the study treatment allocation. In the event of serious adverse events or emergency clinical treatment requiring knowledge of the study treatment, a request for unblinding shall be submitted to the pharmacist with reasonable justification for unblinding. The unblinding details and justification will be documented in the case record form. If unblinded, the participant will be withdrawn from the study in the event of unblinding, and post-treatment follow-up conducted as per protocol or as applicable. Where possible, the outcome assessor will remain blinded to all treatment allocation until end of study, even after participants are withdrawn from the study.

Study procedures

Preoperative

The two randomized groups, based in the IJN, will be matched according to sex, age, NYHA criteria, ejection fraction, and diabetic status. Operative and peri-operative conditions will also be similar for both groups. All subjects will receive an identical prophylactic antibiotic regime consisting of Cefazolin 2gm at induction and 1gm 12 hourly for 48 hours. Gentamicin 2mg/kg will also be given at induction.

Intra-operative

- On-pump CABG

A standard procedure of performing CABG will be done under general anaesthesia. Veins will be harvested from the legs and the left internal mammary will be taken down once the chest is opened. Titanium clips will be used to secure all side branches of saphenous veins and also the branches of the internal mammary artery. The patient will then be cannulated and placed on a heart-lung machine. Once the ascending aorta is cross-clamped, cold cardioplegic solution will be perfused in an ante-grade manner through the aortic root into the heart and the pericardial sac will be buried with ice sludge to create myocardial hypothermia.

The diseased coronary arteries will then be identified and arteriotomies will be performed beyond the level of blockages. The open ends of the saphenous veins and the internal mammary artery are sewn to the openings artertomies using Prolene 7/0 sutures. Once the distal anastomoses are constructed, proximal anastomoses will commence.

Once all the anastomoses are completed, the cross-clamp will be taken off and the heart-lung machine will then be gradually weaned off. Subsequently, the patient will be decannulated. Drainage catheters will be placed around the heart and temporary pacing will be used to the surface of the heart. Sternum will then be closed with steel wire and subcutaneous tissue and skin in the usual manner.

- Off-pump CABG

In this case, the bypass surgery will be done without the use of heart-lung machine. Surgery will be done on a beating heart. The procedure is similar to the on-pump surgery. Chest is opened and the left internal mammary artery is taken down. A stabilizer is placed on the heart to limit the motion of the heart. The anastomosis is done by sewing the open end of the internal mammary to the coronary artery, namely the left anterior descending artery. Chest is closed in the usual manner after placement of the drains and the temporary pacing wire.

Post-operative

Both groups will continue taking either two capsules of Tocotrienol or two capsules of placebo (palm Super Olein) daily for the entire hospitalization period until follow-up at six weeks after discharge. The capsules will be taken on a bd – twice-daily dosing. If the patient is still ventilated in the cardiac ICU, the capsules will be broken and the content administered via a Ryle’s tube.

After surgery, patients will be admitted to the ICU with close monitoring on one-patient-one-nurse basis and will be subsequently transferred to a monitored high dependency unit if their condition is stable. Continuous heart rhythm monitoring will be carried out using the 12-lead ECG. The monitoring will be continued for the first four to five postoperative days on the normal cardiothoracicwards using Holter monitors until the patients are discharged.

The electrocardiographic data will be stored for 24 h and reviewed on a daily basis by the cardiothoracic team involved in the research. The printouts of all abnormal rhythms will also be reviewed for any episodes of arrhythmia. All printouts will be included in the clinical records. Additionally, an ECG will also be recorded in case of symptoms or when arrhythmia is suspected on clinical grounds; AF episodes will always be treated under the direction of the attending cardiothoracic surgeon.

After discharge, all patients will be asked to report to the outpatient department of our institution in case of any relevant symptom. Additionally, all patients will be followed up six weeks after discharge; this will include physical examination and a 12-lead ECG measurement.

Measurements

ECG readings to detect AF post operatively

Diagnostic criteria:
1. Absent P waves
2. Evidence of fibrillation (f) waves instead of P waves. These are irregular undulations of the base line in ECG.
3. Irregular R-R intervals

Length of hospital stay (LoHS)

This information will be obtained from the IJN registry of patients. Three measurements will be used to determine the LoHS:

1. Total Cardiac ICU length of stay;
2. Total HDU length of stay; and
3. Total hospital length of stay.

Health related quality of life (HRQoL)

The analysis of HRQoL will be performed using the validated Short-Form 36 Questionnaire (SF-36). SF-36 has
demonstrated its efficiency in clinical studies and has been proven to be generally acceptable to patients, consistent and a valid measure of health outcome. We will be using the validated Malay version of SF-36, which is a thirty-six items questionnaire, which measures QoL across eight domains that encompass both physical and mental.

The questionnaire will be distributed to both groups of patients before CABG and at discharge and six weeks during clinic visit postoperatively. The questionnaires are going to be administered using the questionnaire-interview approach.

**Sample size calculations**

We will use the PS Power and Sample Size Calculation Software version 3.1.2 for sample size calculation. We plan to study a continuous response variable from independent control and experimental subjects with 1 control per experimental subject.

Calculation of sample size in the present study (randomized controlled trial: RCT to rule out selection bias) requires precise specification of:

- The primary hypothesis of the study (Tocotrienol consumption reduces the incidence of post-operative atrial fibrillation in subjects that had undergone CABG)
- The method of analysis (using Relative Risk: RR).

We will also take into account the possibility of ‘loss to follow-up’ or attrition bias of subjects by analysing all subjects from the start to the completion of the study according to the groups that they were originally randomized. This is called an Intention-To-Treat (ITT) analysis. To calculate the desired sample size based on the above consideration, we use the PS Power and Sample Size Calculation Software for sample size calculation.

In the present RCT, estimated sample size for the primary endpoint (incidence of POAF) was computed on the basis of findings from Calo et al. In that study, the incidence of post-operative AF was 15.2% in the treatment group. Whereas in the placebo group, the incidence was 33.3% (RR=15.2/33.3 = 0.46).

To account for possible subject withdrawal/non-compliance (attrition bias), we will adopt the ITT analysis so that subjects who are ‘lost to follow-up’/non-compliant will be analyzed according to the groups that they were randomized at the beginning of the study. The formula suggested by Witters is “1/1-c” where c is the proportion that is lost to follow-up/did not comply. When c= 12%, for a total enrollment of 260 patients. However, when c=20%, the estimated sample size is increased to 278 (rounded to 280).

Using the PS Power and Sample Size Calculator with α equivalent to 0.05 and power (1-β) is 0.8, the estimated sample size based on the two proportions above is 130 intervention subjects and 130 control subjects (Fisher’s exact test) if we assume the “lost to follow up” is 12%. However, if we assume that the ‘lost to follow up’ is as high as 20%, then the two proportions will be 140 (Intervention Group) and 140 (Control Group) with a total number of 280 patients.

**Statistical analysis**

IBM SPSS Statistics version 25.0 will be used to analyze the results. The ITT analysis will be used to analyze all outcome. The RR of the two-binomial proportion analysis will be used to test the occurrence of postoperative AF in the two treatment groups. The Kaplan-Meier method will be used to look at the cumulative risk. Log-rank test will compare the survival curves of the two treatment groups. Continuous variables will use mean (±SD) while categorical variables will use frequencies (percentages). We will use unpaired Student-t test for continuous variables to look for differences between groups. And group differences will be examined by the chi-square or Fisher exact tests as appropriate for categorical variables. In case of an expected frequency of less than 5 in any cell in a 2×2 table, a Fischer Exact test will be applied. A p value of less than 0.05 will be considered as statistically significant.

In order to find the predictors of AF after surgery, a stepwise multiple logistic regression analysis will be done. To examine the mean (±SD) LoHS (number of days) differences between the two groups, we will use an unpaired Student-t test.

To examine the mean (±SD) QoL score differences between the two groups (pre-operative, six weeks, and three months), the one-way mixed-mode repeated measure ANOVA with post-hoc multiple comparison test (between and within subject) of the two groups will be performed.

In order to look for the factors influencing the change in the quality of life after CABG, we will perform the univariate simple logistic regression (SLogReg) initially, and examining the statistical significance of each independent variable such as the number of revascularization and duration of surgery on the outcome. Then, we will perform a multiple logistic regression (MLogReg) including variables with a level of significance less or equal to 0.25 in the univariate logistic regression and controlling for the effects of possible confounding variables (sex, age, NYHA Criteria, ejection fraction, and diabetic status). A p value of less than 0.05 will be taken as significant.

**Monitoring**

The Principal Investigator, AFM, will be responsible for ensuring participants’ safety on a daily basis and for reporting Serious Adverse Events and Unanticipated Problems to his IJN Review Board (IJNRB) as required. The study statistician prepares reports that list adverse events, serious adverse events, deaths, and disease-or treatment-specific events required for monitoring body review in order to ensure good clinical care and identify any emerging trends. The IJNRB will act in an advisory capacity to the IJN Ethics Committee to monitor participants’ safety, evaluate the progress of the study, to review procedures for maintaining the confidentiality of data, the quality of data collection, management, and analyses.
Data dissemination
Data dissemination will be done via three ways. Firstly, via presentation at local and international conferences either in the form of posters and/or oral presentation. Secondly via writing for publication in scientific journals. Thirdly, via participation at international exhibitions and conventions on research and innovation.

Study termination
The sponsor has the full discretion to decide on the termination of the study at any time. Patients will be informed if the study is terminated and follow-up visits will be arranged if needed.

Data Availability
No data are associated with this article.

Supplementary material
Supplementary File 1: Informed consent form and participant information sheet.
Click here to access the data.

References


The benefits of publishing with F1000Research:

- Your article is published within days, with no editorial bias
- You can publish traditional articles, null/negative results, case reports, data notes and more
- The peer review process is transparent and collaborative
- Your article is indexed in PubMed after passing peer review
- Dedicated customer support at every stage

For pre-submission enquiries, contact research@f1000.com