



Assessing the Impact of Colchicine on Coronary Plaque Phenotype After Myocardial Infarction with Optical Coherence Tomography: Rationale and Design of the COCOMO-ACS Study

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Abstract

Introduction Recurrent event rates after myocardial infarction (MI) remain unacceptably high, in part because of the continued growth and destabilization of residual coronary atherosclerotic plaques, which may occur despite lipid-lowering therapy. Inflammation is an important contributor to this ongoing risk. Recent studies have shown that the broad-acting anti-inflammatory agent, colchicine, may reduce adverse cardiovascular events in patients post-MI, although the mechanistic basis for this remains unclear. Advances in endovascular arterial wall imaging have allowed detailed characterization of the burden and compositional phenotype of coronary plaque, along with its natural history and responsiveness to treatment. One such example has been the use of optical coherence tomography (OCT) to demonstrate the plaque-stabilizing effects of statins on both fibrous cap thickness and the size of lipid pools within plaque.

Methods The Phase 2, multi-centre, double-blind colchicine for coronary plaque modification in acute coronary syndrome (COCOMO-ACS) study will evaluate the effect of colchicine 0.5 mg daily on coronary plaque features using serial OCT imaging in patients following MI. Recruitment for the trial has been completed with 64 participants with non-ST elevation MI randomized 1:1 to colchicine or placebo in addition to guideline recommended therapies, including high-intensity statins. The primary endpoint is the effect of colchicine on the minimal fibrous cap thickness of non-culprit plaque over an 18-month period.

Summary The COCOMO-ACS study will determine whether addition of colchicine 0.5 mg daily to standard post-MI treatment has incremental benefits on high-risk features of coronary artery plaques. If confirmed, this will provide new mechanistic insights into how colchicine may confer clinical benefits in patients with atherosclerotic cardiovascular disease.

Trial Registration ANZCTR trial registration number: ACTRN12618000809235. Date of trial registration: 11th of May 2018.

Keywords Colchicine · Optical coherence tomography · Coronary plaque · Inflammation · Fibrous cap

Introduction

Atherosclerotic coronary artery disease (CAD) is the principal source of cardiovascular mortality, morbidity and health economic burden worldwide [1, 2]. While aggressive modification of traditional atherosclerosis risk factors has led to a substantial reduction in the incidence of myocardial

infarction (MI) [3], the recurrence of major adverse cardiovascular events (MACE) post-MI remains a major concern [4]. Inflammation is independently and strongly associated with future MACE in both primary and secondary prevention settings [5–8]. Current therapies for CAD, such as lipid-lowering statins, do not adequately control plaque inflammation, meaning that new anti-atherosclerotic agents targeting inflammation are needed [9]. The positive results for the anti-interleukin-1-beta (IL-1 β) monoclonal antibody, canakinumab, in the CANTOS clinical trial [10], highlighted that not only is inflammation important from a pathogenic and risk prediction perspective in CAD, but that reducing

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inflammation can be beneficial. More recently, low-dose colchicine, a broad acting anti-inflammatory agent, has been repurposed for the treatment of CAD. Results from the LoDoCo [11], LoDoCo 2 [12], COPS [13] and COLCOT [14] studies have demonstrated potential clinical benefit of colchicine use in patients with both stable and unstable CAD, with other Phase 3 clinical studies still ongoing [9]. Colchicine's potential mechanism of effect in atherosclerosis is protean, and although it may primarily act via its known action on innate immune pathways, we hypothesize that it may also promote favourable plaque healing following MI, leading to the formation of a thicker plaque cap which is more resilient to rupture and thrombosis.

Coronary Atherosclerosis: an Inflammatory Process Requiring Treatment Beyond Statins

Considered a chronic inflammatory disease of blood vessels, atherosclerosis is characterized by a maladaptive immune response, triggered by subintimal lipoprotein accumulation, and resulting in the formation of lipid-rich, inflamed plaques [9]. Ultimately, systemic and local activation of both the adaptive and innate immune systems can result in either rupture or erosion of an atherosclerotic plaque, leading to thrombosis, which may manifest as unstable angina, acute MI or sudden cardiac death [15, 16]. Despite no established association between genetically elevated C-reactive protein (CRP) and coronary heart disease in genome-wide association studies [17, 18], measurement of the inflammatory biomarker, high-sensitivity CRP (hs-CRP), has been shown to independently predict future cardiovascular events in both patients with stable coronary disease [19] and those with acute coronary syndrome (ACS) [20]. After adjusting for traditional risk factors, a meta-analysis of 54 long-term prospective studies involving 160,000 participants revealed that each standard deviation increase in log-normalized hs-CRP was associated with an increase in relative risk of 1.37 (95% CI, 1.27 to 1.48) for CAD and 1.55 (95% CI, 1.37 to 1.76) for cardiovascular mortality [21]. Statins reduce hs-CRP in a partly low-density lipoprotein cholesterol (LDL-C) independent manner, and in the JUPITER study, the most favourable clinical outcomes were observed in participants who achieved both a reduction in LDL-C and hs-CRP [22]. However, the concerning frequency of recurrent cardiovascular events despite highly effective LDL-C reduction has led to concerns that residual inflammatory risk is also causal. The PROVE-IT trial showed that of the 2,099 patients with prior ACS prescribed atorvastatin 80 mg daily, 44% achieved target reductions of both LDL-C (< 1.8 mmol/L) and hs-CRP (< 2 mg/L) [23, 24]. However, 29% were left with a residual inflammatory burden (hs-CRP > 2 mg/L), 13% a residual cholesterol risk (LDL-C > 1.8 mmol/L) and 14% with both,

indicating that residual inflammatory risk is frequent after high-intensity statin therapy. Together, these results highlight that recurrent cardiovascular events are determined by multiple biological processes that require a tailored therapeutic approach and that future treatment for CAD must tackle residual inflammation not completely addressed by statin therapy alone.

Repurposing Potential of Colchicine in CAD

Colchicine, an orally administered, inexpensive anti-inflammatory medication used to treat gout, pericarditis and familial Mediterranean fever (FMF), has been repurposed over recent years to treat CAD. Colchicine binds to microtubule ends and inhibits cytoskeletal microtubule processes [25]. At low concentrations, it inhibits their formation, and at higher concentrations, it promotes their depolymerization [26]. Neutrophil chemotaxis, phagocytosis and protein excretion are all microtubule-dependent processes. In vitro experiments have identified that colchicine lessens endothelial selectin family-dependent adhesiveness, affecting both endothelial E-selectin and neutrophil L-selectin surface expression [27]. Additionally, it has been found to increase leukocyte cyclic adenosine monophosphate levels, inhibit IL-1 production by activated neutrophils and to downregulate tumour necrosis factor-alpha receptors in macrophages [28]. In the setting of atherosclerotic plaque containing cholesterol crystals, NLRP3 inflammasome activation is dampened in colchicine-treated neutrophils and macrophages [29]. Colchicine also suppresses neutrophil extracellular traps (NETs) by restoring cytoskeletal dynamics in patients treated with percutaneous coronary intervention (PCI) following MI [30]. Comprising strands of DNA in the form of decondensed chromatin, and liberated by activated or dying neutrophils, NETs localize at the interface of blood and the intimal surface of diseased arteries [31]. Here they form a fibrin-like base for platelet adhesion, activation and aggregation, promote the accumulation of prothrombotic molecules, including von Willebrand factor and fibrinogen, and facilitate erythrocyte adhesion, all of which contribute to thrombus formation [32].

Retrospective observational studies identified lower rates of MI and other vascular events in patients with gout and FMF treated with colchicine [33, 34]. The first prospective study to examine the impact of colchicine in cardiovascular disease was the LoDoCo study, which evaluated the effect of colchicine 0.5 mg daily in addition to antiplatelet and statin therapy in 532 patients with stable CAD [11]. After a median follow-up of 3 years, participants who had received colchicine demonstrated a significant reduction in a composite endpoint consisting of MI, cardiac arrest or non-cardioembolic stroke when

compared to the no colchicine control group (hazard ratio (HR), 0.33). The LoDoCo2 study, which randomized 5,522 patients with stable CAD to colchicine 0.5 mg daily or matching placebo revealed a similar therapeutic benefit at a median follow-up of 28.6 months [12]. The colchicine-treated group had a significant reduction in the primary composite endpoint of cardiovascular death, non-procedural MI, ischemic stroke, or ischemia-driven coronary revascularization compared to placebo (HR, 0.69). The secondary endpoint composite of cardiovascular death, spontaneous MI and ischemic stroke was also significantly lowered (HR, 0.72), as were the individual endpoints of MI and revascularization.

Inflammation is especially accentuated in the coronary vascular bed following acute MI [35]. The COLCOT trial demonstrated the potential benefits of colchicine in 4,755 patients randomized within 30 days following MI to receive colchicine 0.5 mg daily or placebo in addition to standard guideline-recommended therapy [14]. The primary efficacy endpoint was a composite of cardiovascular death, resuscitated cardiac arrest, MI, stroke or urgent hospitalization for angina requiring coronary revascularization and occurred in 5.5% of the participants who received colchicine compared to 7.1% in the placebo arm (HR, 0.77). The difference in the primary endpoint was driven by a significant risk reduction in urgent hospitalization for angina requiring revascularization (HR, 0.50) and for stroke (HR, 0.26). Trends towards reduced cardiovascular death (HR, 0.83) and MI (HR, 0.91) in colchicine-treated participants did not reach statistical significance. A recent substudy of COLCOT established that there was a significant reduction in the incidence of the primary endpoint for patients in whom colchicine was initiated between days 0 and 3 compared with placebo (HR, 0.52), in contrast to patients who either started colchicine between days 4 and 7 (HR, 0.96), or from day 8 onwards (HR, 0.82) [36]. This intimated that early initiation of low-dose colchicine after MI provides maximal clinical benefit with a 48% reduction in ischemic cardiovascular events when compared to placebo and supports the strategy of early in-patient initiation of therapy.

The COPS trial randomized 795 patients with ACS, managed with either PCI or medical therapy, to receive either colchicine 0.5 mg twice daily for 1 month, then 0.5 mg daily for 11 months, or placebo [13]. At 12-month follow-up, there was no significant difference in the primary composite endpoint of all-cause mortality, ACS, ischemia-driven urgent revascularization and non-cardioembolic ischemic stroke in the colchicine group compared to placebo (6.1 versus 9.5%; $P=0.09$). However, a trend towards clinical benefit following the use of colchicine was observed and may not have reached statistical significance due to the small sample size. Noteworthy, in a post hoc analysis of the composite endpoint using only cardiovascular death rather than total death, a

significant reduction in events in favour of colchicine was demonstrated at 12 months (HR, 0.51).

The propensity of low-dose colchicine to cause gastrointestinal intolerance in up to 10–15% of patients [37], the increased risk of pneumonia seen in COLCOT (0.9% compared with 0.3% in placebo arm; $P=0.03$) [14] and a non-significant trend toward higher mortality from non-cardiovascular causes in some studies, including a recent meta-analysis [12, 38], provide some cause for caution in colchicine's routine use. Additional randomized controlled trial data is needed to help clarify its safety and efficacy.

Mechanistic Actions of Colchicine on Atherosclerotic Plaque

The successful repurposing of colchicine for atherosclerotic cardiovascular disease will also benefit from a better understanding of how it works. However, few studies have directly examined colchicine's effects on atherosclerotic plaque. Published preclinical data are mostly limited to studies performed more than 20 years ago and provide conflicting evidence that colchicine may either exacerbate [39] or inhibit [40] plaque formation or have no effect [41]. Most recently, in a rabbit model of atherosclerosis, in vivo multimodality imaging using magnetic resonance imaging, ^{18}F -Fluorodeoxyglucose-positron emission tomography, optical coherence tomography (OCT) and immunohistology demonstrated that colchicine may stabilize atherosclerotic plaque by reducing plaque inflammation and burden, without altering macrophage infiltration or atherosclerotic plaque microstructure. In this study, no difference was observed in terms of lipid plaque proportion or mean minimum fibrous cap thickness measured by OCT in either colchicine- or placebo-treated cohorts [42].

So far the only clinical evaluation of colchicine's ability to modify coronary plaque comes from a non-randomized, single centre observational study of 80 patients post-ACS, in whom colchicine significantly reduced low attenuation plaque volume (treatment: mean 15.9 [−40.9%] ± 17.3 mm³; control mean 6.61 [−17.0%] ± 12.8 mm³; $P=0.008$), as measured by serial coronary computed tomography angiography over 12 months [43]. This was accompanied by a reduction in hs-CRP with colchicine compared to the non-placebo control group and supports the notion that colchicine may help to stabilize plaques. Although the predominant focus on the mechanism of action of colchicine in atherosclerosis has been on its purported ability to attenuate cholesterol crystal-induced inflammasome activation [44], there is increasing evidence that other pleiotropic mechanisms may also be instrumental in providing atheroprotection and plaque stabilization [45]. This is supported by a recent proteomic substudy of LoDoCo 2 [46], in which

colchicine was associated with a reduction in serum levels of 11 proteins not directly involved in the NLRP3 inflammasome pathway, including strong attenuation in proteins involving neutrophil degranulation (myeloblastin, carcinoembryonic antigen-related cell adhesion molecule 8, azurocidin and myeloperoxidase). Colchicine also resulted in an increase in another 23 proteins, including intestinal fatty acid-binding protein, a biomarker of intestinal barrier dysfunction, protein-glutamine γ -glutamyltransferase 2, related to tissue repair, and both fibroblast growth factor 21 and insulin-like growth factor-binding protein 1, hypothesized to protect against atherosclerosis.

Assessment of Therapeutic Modulation of Plaque Using Endovascular Imaging

Intracoronary OCT is a catheter-based imaging technique which utilizes coherent near-infrared light, typically of a wavelength of approximately 1,300 nm, to create images of plaque atheroma in the coronary arteries from optical backscatter. Its greatest advantage is its high spatial resolution (approximately ten times higher than that of intravascular ultrasound), with an axial resolution of up to 10 μ m and lateral resolution of up to 20 μ m [47, 48]. This enables qualitative and quantitative analysis of the atheroma below the intimal endothelial surface, providing the ability to discriminate between fibrous, lipid-rich and calcified plaques [47, 49, 50], quantitate lipid content and macrophage burden [49, 51] and accurately measure fibrous cap thickness [50].

Lipid-rich thin-cap fibroatheromas (TCFAs) that are believed to be more prone to rupture can be identified and quantitated by OCT. The relationship between culprit site lipid-rich plaques (LRP) and cardiovascular events has been documented in patients with ACS utilizing OCT [52, 53]. A retrospective study demonstrated that OCT-detected non-culprit LRP led to a two-fold increase in non-culprit MACE (*HR*, 2.06), primarily driven by revascularization for recurrent ischemia [54]. In addition, OCT has been used to assess modulation of coronary atherosclerosis in response to different established therapies. Serial OCT assessment of non-culprit lesions was used in 70 patients with unstable angina and dyslipidemia in the Japanese EASY-FIT study to investigate plaque stability. Patients were randomized to either 20 mg or 5 mg of atorvastatin daily and imaging was performed at baseline and 12-month follow-up [55]. Lower LDL-C levels were obtained in the higher dose group (69 versus 78 mg/dL, $P=0.039$) with OCT suggesting a more stable plaque in this group, characterized by a significant increase in fibrous cap thickness of 69% compared to 17% ($P<0.001$). The increase in fibrous cap correlated with the decrease in LDL-C levels and grade of OCT measured macrophages. Similar results were seen in a single centre OCT

study that examined the effects of the addition of ezetimibe 10 mg daily to treatment with fluvastatin 30 mg daily. In this study, the reduction in LDL-C was significantly larger in the group receiving combination therapy (-34.0 ± 32.0 versus -8.8 ± 17.4 mg/dL, $P<0.001$), and while OCT demonstrated a significant increase in fibrous cap thickness after 9 months of therapy in both groups, this was greater in those receiving combination therapy [56].

These findings contribute to our understanding of the mechanistic effects of statins on plaque. As new therapies evolve in the treatment of CAD, intracoronary imaging studies will continue to complement Phase 3 outcome trials by evaluating their effectiveness at modifying the burden and composition of coronary plaques [57].

Design of the COCOMO-ACS Study

Objectives

The COCOMO-ACS study will employ serial OCT imaging to evaluate the impact of colchicine on coronary atheroma phenotype. The primary objective is to evaluate the effect of colchicine on changes in fibrous cap thickness on non-culprit plaques in patients with non-ST-elevation myocardial infarction (NSTEMI), taking maximally tolerated oral lipid-lowering therapy. Additional objectives are to evaluate the impact of colchicine on other measures of plaque phenotype and the safety and tolerability of colchicine when administered to patients in the post-ACS setting. The primary hypothesis of the study is that the addition of colchicine to high-intensity statin therapy will have a favourable effect on coronary plaque composition, with a greater increase in the minimum fibrous cap thickness compared to lipid-lowering therapy alone.

Study Design

COCOMO-ACS is an investigator-initiated, multicentre, randomized, double-blind, placebo-controlled, Phase 2 clinical trial in patients hospitalized due to NSTEMI to evaluate the effect of colchicine on non-culprit plaque phenotype as assessed by serial OCT. The COCOMO-ACS study includes patients from nine sites across Australia with no maximum limit on recruitment at each centre.

Ethical Aspects

All procedures in this study are conducted in accordance with the Declaration of Helsinki, International Conference on Harmonization, Good Clinical Practice guidelines and applicable regulatory requirements. The final protocol and informed consent forms were approved by the Royal

Adelaide Hospital Human Research Ethics Committee (HREC) under the National Mutual Acceptance (NMA) system with local Research Governance Officer (RGO) approval at each site (HREC/17/RAH/366; Central Adelaide Local Health Network Reference Number R20170904). This clinical trial was registered with the Australian New Zealand Clinical Trials Registry (ACTRN12618000809235) on the 11th of May 2018.

Study Population

Randomization has now been completed for this study. Eligible participants were those aged between 18 and 82 years of age, who had a clinically indicated coronary angiogram

within 72 h of presenting with NSTEMI and gave written informed consent for the study before angiography. A culprit atherosclerotic lesion was identified and managed as clinically indicated. There must have been at least one intermediate lesion in a non-culprit coronary artery, determined visually to be 20–50% stenotic, with no intention to revascularize it in the next 18 months. The non-culprit lesion was imaged by OCT at baseline angiography. OCT images were assessed within 48 h at the Atherosclerosis Imaging Core Laboratory at the South Australian Health and Medical Research Institute (SAHMRI), to ensure that they were of suitable quality and that the target vessel contained a lipid-rich plaque (fibrous cap thickness $\leq 120 \mu\text{m}$ and lipid arc $> 90^\circ$). A full list of inclusion and exclusion criteria is provided in Table 1.

Table 1 Eligibility criteria for COCOMO-ACS study

Inclusion criteria (if all of the following met)	Exclusion criteria (if any of the following met)
1. Participants who undergo clinically indicated coronary angiography within 72 h of presentation with a NSTEMI	1. Left main coronary disease ($> 50\%$ reduction in lumen diameter by angiographic visual estimation)
2. Able to provide written informed consent before baseline angiography	2. Cardiogenic shock
3. Male or female ≥ 18 and ≤ 82 years of age	3. Heart failure (NYHA class IV) or LVEF $\leq 35\%$
4. Must meet all of the following criteria at the qualifying coronary catheterization procedure:	4. Episode of acute gout within the last 5 years
a. Angiographic evidence of coronary artery disease, with a culprit lesion identifiable for the NSTEMI and managed as clinically indicated	5. Currently prescribed colchicine for other indication, presence of contraindications to colchicine, known prior intolerance to colchicine. Concomitant therapy with drugs that could interact with colchicine
b. Target coronary artery for OCT:	6. Dialysis or eGFR $< 30 \text{ ml/min/1.73 m}^2$
i. At least one non-culprit lesion in a non-culprit artery, determined angiographically to be 20–50% stenotic	7. Thyroid stimulating hormone (TSH) $<$ lower limit of normal (LLN) or $> 1.5 \times$ upper limit of normal (ULN)
ii. When multiple non-culprit lesions of 20–50% are present, it is preferable that the most angiographically severe one is imaged	8. Active liver disease or hepatic dysfunction
iii. Vessel for interrogation must be accessible to the OCT catheter	9. Known major active infection, or major hematologic, renal, metabolic, gastrointestinal or endocrine dysfunction
iv. Target vessel has not undergone prior PCI or CABG surgery and is not a bypass graft	10. Significant haematological abnormalities on assessment of complete blood picture: Hb $< 100 \text{ g/L}$, Plt $< 150 \times 10^3 /\mu\text{L}$, white cell count $< 3.5 \times 10^3 /\mu\text{L}$
v. Target vessel is not currently a candidate for intervention or a likely candidate for intervention over the next 18 months	11. Active malignancy (except non-melanoma skin cancers, cervical in situ carcinoma, breast ductal carcinoma in situ, or stage 1 prostate carcinoma)
5. Able to be randomized within 10 days of catheterization	12. Female participants cannot be pregnant or breast feeding and premenopausal participants must be willing to use at least two highly effective methods of birth control during treatment and for an additional 12 weeks after the end of treatment
6. Baseline OCT interrogation determined to be of acceptable quality, and target vessel contains a lipid-rich plaque with a FCT $\leq 120 \mu\text{m}$ and lipid arc $> 90^\circ$, at core laboratory review	13. Unable to give informed consent
	14. Not willing or able to attend follow-up visits or follow-up OCT procedure
	15. Any other information that the investigator considers will limit the ability of the participant to complete all study associated procedures

CABG coronary artery bypass graft, eGFR estimated glomerular filtration rate, FCT fibrous cap thickness, LVEF left ventricular ejection fraction, NSTEMI non-ST-elevation myocardial infarction, NYHA New York Heart Association, OCT optical coherence tomography, PCI percutaneous coronary intervention

Randomization and Allocation Concealment

Participants who passed angiographic and OCT screening were randomized 1:1 within 10 days of their angiogram and OCT study, to colchicine 0.5 mg or placebo, administered orally daily for 18 months. Randomization was by a computer generation Web response system, and to ensure balance, patients were stratified into two groups according to prior treatment with statins. Group 1 includes patients with no statin use in the preceding 4 weeks prior to admission. Group 2 includes patients who had been on a statin at any dose for at least 4 weeks prior to the admission. The randomization schedule included blocking within each stratum with variable block size of two to four. Treatments were randomly allocated within blocks with an equal number of each treatment within each block.

Once the participants were randomized, the randomization number was given to the pharmacist, and kit numbers containing the appropriate matching allocation were dispensed to participants. All treatment packs are identical and contain either active tablets (Colgout® manufactured by Aspen Pharmacare, NSW, Australia) or matching placebo tablets consisting of PROSOLV® EASYtab SP (composition: 95.0–98.0% microcrystalline cellulose, 1.5–2.5% colloidal silicon dioxide, 0.5–2.0% sodium starch glycolate and 0.3–1.0% sodium stearyl fumarate) (manufactured by Pharmaceutical Packaging Professionals Pty Ltd trading as PCI Pharma Services, Victoria, Australia). Researchers have no access to the randomization list. All researchers, study participants and adjudicators remain blinded to the treatment allocation. Only the pharmacist and a small unblinded team at SAHMRI have access to the randomization list to ensure allocation concealment throughout the study.

Standard of Care

All participants also receive currently recommended therapies after MI at the discretion of their treating clinician: dual antiplatelets, cholesterol-lowering medication including maximally tolerated dose of a statin to achieve LDL-C of < 1.8 mmol/L, with or without an ACE inhibitor/angiotensin II-receptor blocker or beta-blocker.

Study Procedures

Initial background assessments included demographics, cardiovascular risk factors, relevant medical and surgical history and laboratory data. Follow-up study visits are performed in the clinic at months 3, 6 and 12 and by telephone at months 1, 9 and 15. Pill counts from study medication bottles returned by participants are used to assess adherence. Imaging of the target artery by OCT was performed at the time of screening visit coronary angiography and will

then be repeated at month 18. A final end of study visit will be performed at 18 months to complete the safety evaluation. The final OCT originally planned to be performed at 12 months has been extended to 18 months in the updated protocol due to restrictions on elective surgery schedules due to the COVID-19 pandemic. The study flowchart is presented in Fig. 1.

Image Analysis

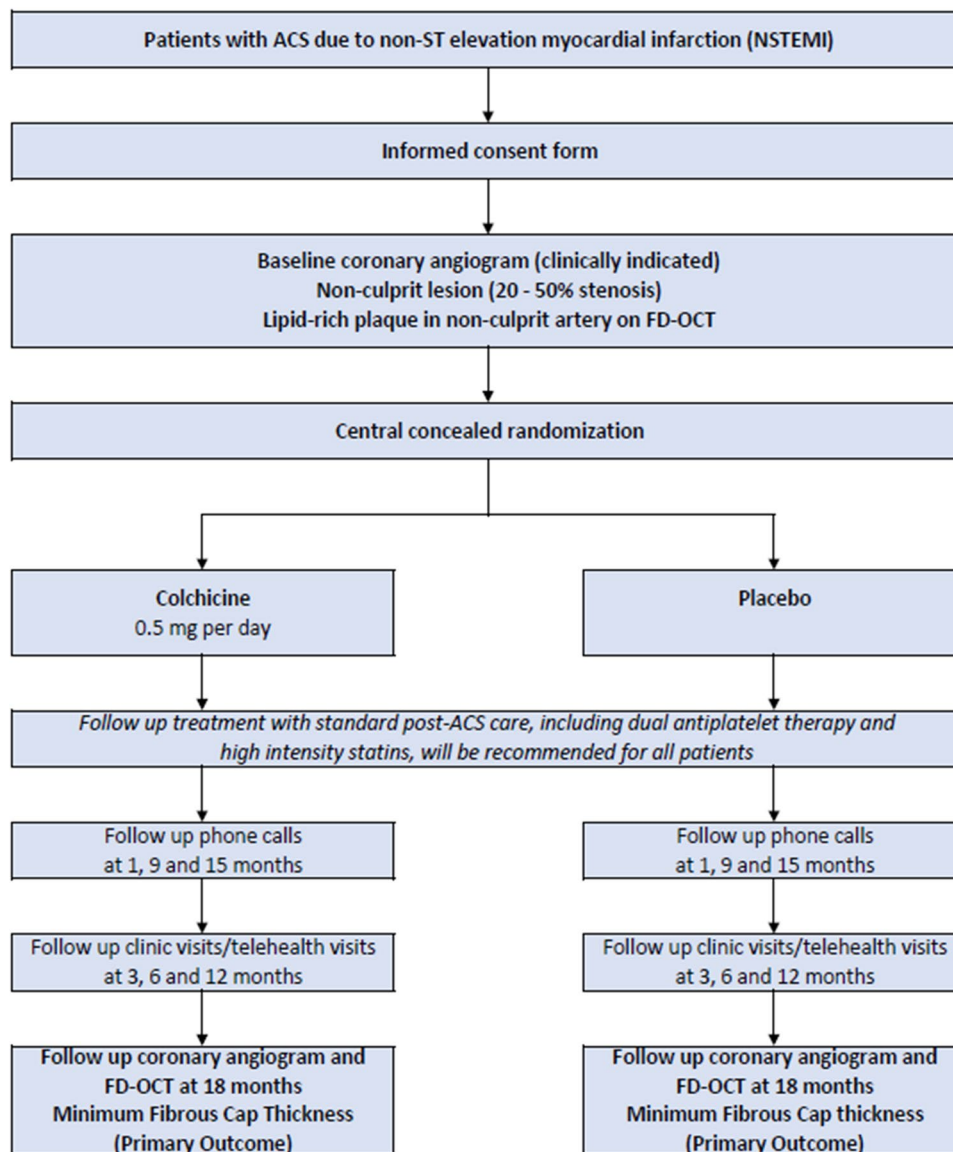
All imaging performed is anonymized, digitally stored and transferred to the SAHMRI Atherosclerosis Imaging Core Laboratory for analysis. Screening OCT imaging was assessed within 48 h for image quality and to determine that the patient met all eligible imaging inclusion criteria. Paired baseline and final OCT images will be analysed at the end of the study, with measurements of plaque phenotype. The segment selected for analysis will be defined by proximal and distal side branches and by the presence of at least one image containing a fibrous cap thickness $\leq 120 \mu\text{m}$ and one image with a lipid arc $> 90^\circ$. The same segment will undergo measurements by analysts who are blinded to the treatment status of the patient. Cross-sectional images spaced 0.2 mm apart will be selected for analysis. For each image, where plaque is present, measurement of minimum fibrous cap thickness, mean fibrous cap thickness, lipid arc and the lipid length will be performed (Fig. 2). Each image will be graded for the presence of macrophage infiltration, microchannels and calcium accumulation [58]. The presence of TCFA, plaque rupture and thrombus will also be recorded.

Sample Collection

Blood samples are being collected at baseline, 3-, 6-, 12- and 18-month visits. Clinical laboratory testing is performed by the local laboratory for complete blood picture, biochemistry, creatinine and estimated glomerular filtration rate (eGFR), liver function tests, thyroid function tests, lipids (LDL-C, high-density lipoprotein cholesterol (HDL-C), triglycerides, non-HDL-C), hs-CRP, creatine kinase, troponin, HbA1c and vitamin B12 level. Plasma, serum and peripheral blood mononuclear cells are also prepared from fasting blood collected at baseline and final study visit, stored at -80°C or in liquid nitrogen and transferred to SAHMRI at the end of the study, for exploratory analysis of inflammatory cytokines and other biomarkers.

Study Endpoints

The primary endpoint of the study is the percentage change in minimum FCT in a non-culprit plaque from baseline to 18 months. The secondary endpoints include absolute change in plaque minimum FCT, absolute and percentage

Fig. 1 COCOMO-ACS study design

change in plaque mean FCT, change in lipid pool size and absolute change in plaque macrophage content. All endpoints, including exploratory endpoints, are listed in Table 2.

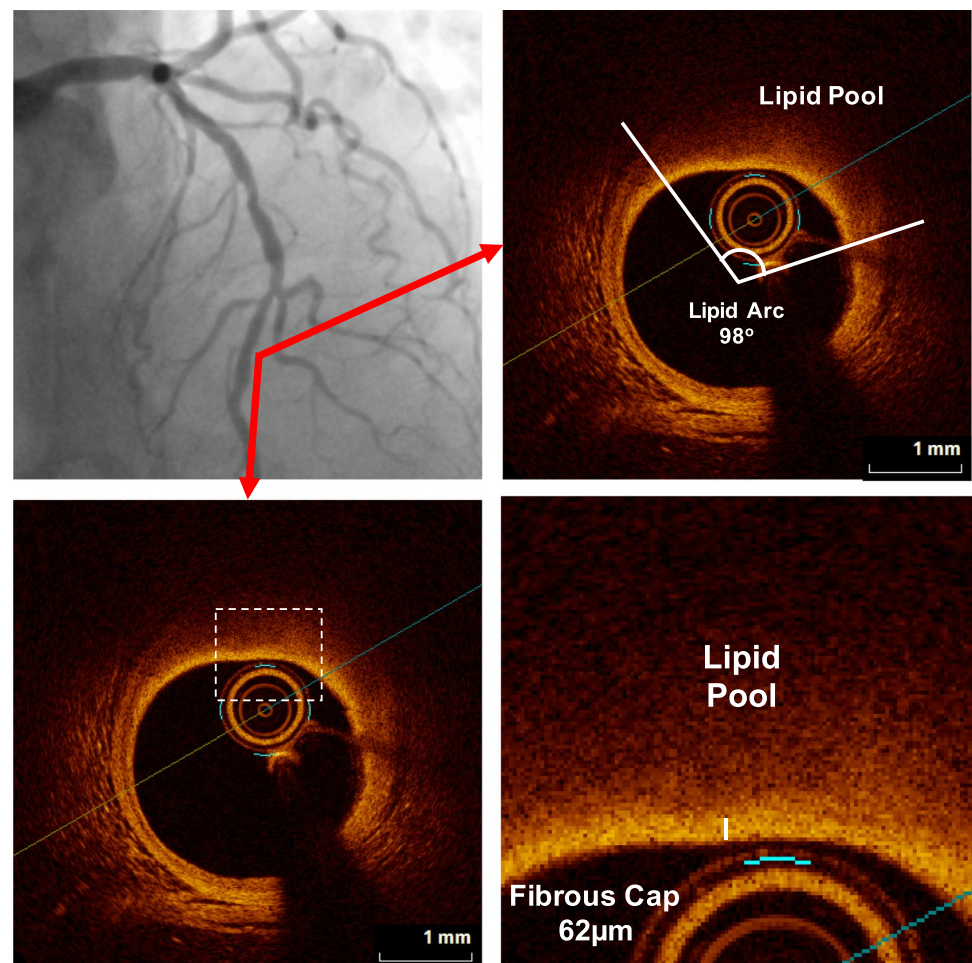
Statistical Analysis

Sample size estimation was based around studies that have demonstrated a mean % change in minimum FCT of $65 \pm 35\%$ with statins, which corresponds here to the expected change in the placebo group [55, 59–64]. We assume that colchicine will result in a segment-specific 50% relative increase of the expected change in minimum FCT ($97 \pm 35\%$). With a two-sided alpha of 0.05 and power of 90%, 52 participants (26 per group) are required to meet these expectations. Allowing for approximately 20% loss in follow-up due to missing data or patient withdrawal, we

randomized a total of 64 participants (32 per group) with recruitment completed in December of 2020.

We will perform intention-to-treat analysis as the primary analysis. Descriptive summary will be presented as percentage frequencies for categorical variables and as *mean* \pm *SD* (or as median with interquartile range) for continuous variables by treatment group. Continuous variables will be tested for normality and appropriate non-parametric test will be used for variables not normally distributed. Group comparisons will be performed by chi-squared (or Fisher's exact) test for categorical variables and independent samples *t* test for continuous variables. The % change in minimum FCT (primary endpoint) will be compared between the two groups using ANCOVA adjusting for baseline measurements. Chi-squared test or independent *t* test will be used for the secondary endpoints to compare the difference in changes between groups. We will explore relationships

Fig. 2 OCT plaque measurements. Example of coronary angiogram image in AP Cranial projection, identifying non-culprit plaques in the left anterior descending artery, with corresponding cross-sectional OCT image from the segment denoted by red arrow, highlighting the thin fibrous cap, lipid pool and lipid arc.



between % change in minimum FCT and % change in inflammatory biomarkers in the colchicine group using simple regression model; multivariable analysis will subsequently be used to determine independent predictors of colchicine responsiveness. Per-protocol analysis will be performed on those with treatment compliance > 80%, as a sensitivity analysis. A complete case

analysis will include all patients with evaluable imaging at both time points. We will also perform a sensitivity analysis using multiple imputation to accommodate patients who were lost to follow-up. A two-sided p value of < 0.05 will be considered statistically significant. Analysis will be performed with Stata MP 14 (StataCorp, TX).

Table 2 Endpoints of COCOMO-ACS study

Primary endpoint

Percentage change in coronary plaque minimum fibrous cap thickness (FCT) as determined by OCT

Secondary endpoints

- Absolute change in plaque minimum FCT
- Absolute and percentage change in plaque mean FCT
- Change in plaque lipid pool size, as determined by OCT measurement of lipid arc and length
- Change in plaque macrophage content

Exploratory endpoints

- Changes in other high-risk features of coronary plaque morphology (e.g. plaque microchannels, cholesterol crystals, spotty calcification) as determined by OCT
 - Absolute and percentage changes in blood lipid levels and inflammatory markers (e.g. hs-CRP, cytokines)
 - Relationship between changes in blood lipids and inflammatory markers and plaque characteristics
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Discussion

There is compelling evidence attesting to the importance of local and systemic inflammation in all stages of atherosclerosis and its complications. Residual inflammation despite current treatment predominantly centred on statin therapy is a harbinger of recurrent cardiovascular risk. Results from recent studies have demonstrated the potential clinical benefits of colchicine in patients with established CAD, including when initiated early after MI. Further, the use of serial arterial wall imaging has provided the opportunity to characterize the effect of lipid-lowering interventions on atherosclerotic plaque *in vivo*.

COCOMO-ACS utilizes intracoronary OCT to extend these observations in order to determine the effect of anti-inflammatory therapy with low-dose colchicine on non-culprit, high-risk lipid-rich plaques in patients presenting with MI. Recruitment for the trial has been completed with 64 participants enrolled, and it is anticipated that statistical analysis on the entire cohort will be complete by late 2022. COCOMO-ACS findings are poised to provide unique mechanistic insights into the potential effects of colchicine on modifying coronary plaque biology. If positive, this study will contribute to the rapidly accumulating evidence that supports the use of colchicine in atherosclerotic CAD.

All Principal Investigators and Participating Centres

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Data Availability There is currently no plan to make individual participant data publicly available for this trial.

Declarations

Ethics Approval and Consent to Participate All procedures in this study will be conducted in accordance with the Declaration of Helsinki, International Conference on Harmonization, Good Clinical Practice guidelines and applicable regulatory requirements. The final protocol and informed consent forms were approved by the Royal Adelaide Hospital Human Research Ethics Committee (HREC) under the National Mutual Acceptance (NMA) system with local Research Governance Officer (RGO) approval at each site (HREC/17/RAH/366; Central Adelaide Local Health Network Reference Number R20170904). This clinical trial is registered with the Australian New Zealand Clinical Trials Registry (ACTRN12618000809235). Informed consent was obtained from all individual participants included in the study.

Consent for Publication. Not applicable.

Competing Interests K.S. has received research support and consultant fees from Abbott Vascular and Bayer. A.S. has received consultant fees and speaker honoraria from Edwards Lifesciences, Medtronic, Boston Scientific, AstraZeneca and Abbott Vascular. D.T.L.W. has received speaker honoraria from AstraZeneca, Pfizer, Bayer and Boehringer Ingelheim. P.L.T. has received research grants or honoraria from Amgen, Amgen, Aspen, AstraZeneca, Boehringer Ingelheim, Bristol-Myers Squibb, DalCor, Merck and Pfizer. S.J.N. has received research support from AstraZeneca, Amgen, Anthera, Eli Lilly, Esperion, Novartis, Cerenis, The Medicines Company, Resverlogix, InfraReDx, Roche, Sanofi-Regeneron and Liposcience and is a consultant for AstraZeneca, Akcea, Eli Lilly, Anthera, Kowa, Omthera, Merck, Takeda, Resverlogix, Sanofi-Regeneron, CSL Behring, Esperion and Boehringer Ingelheim. P.J.P. has received research support from Abbott Vascular, consulting fees from Amgen and Esperion and speaker honoraria from AstraZeneca, Bayer, Boehringer Ingelheim, Merck Schering-Plough and Pfizer.

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