

BMJ Open Coronary calcium scoring as first-line test to detect and exclude coronary artery disease in patients presenting to the general practitioner with stable chest pain: protocol of the cluster-randomised CONCRETE trial

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ABSTRACT

Introduction Identifying and excluding coronary artery disease (CAD) in patients with atypical angina pectoris (AP) and non-specific thoracic complaints is a challenge for general practitioners (GPs). A diagnostic and prognostic tool could help GPs in determining the likelihood of CAD and guide patient management. Studies in outpatient settings have shown that the CT-based coronary calcium score (CCS) has high accuracy for diagnosis and exclusion of CAD. However, the CT CCS test has not been tested in a primary care setting. In the COroNary Calcium scoring as fiRst-line Test to dEct and exclude coronary artery disease in GPs patients with stable chest pain (CONCRETE) study, the impact of direct access of GPs to CT CCS will be investigated. We hypothesise that this will allow for early diagnosis of CAD and treatment, more efficient referral to the cardiologist and a reduction of healthcare-related costs.

Methods and analysis CONCRETE is a pragmatic multicentre trial with a cluster randomised design, in which direct GP access to the CT CCS test is compared with standard of care. In both arms, at least 40 GP offices, and circa 800 patients with atypical AP and non-specific thoracic complaints will be included. To determine the increase in detection and treatment rate of CAD in GP offices, the CVRM registration rate is derived from the GPs electronic registration system. Individual patients' data regarding cardiovascular risk factors, expressed chest pain complaints, quality of life, downstream testing and CAD diagnosis will be collected through questionnaires and the electronic GP dossier.

Ethics and dissemination CONCRETE has been approved by the Medical Ethical Committee of the University Medical Center of Groningen.

Trial registration number NTR 7475; Pre-results.

INTRODUCTION

Cardiovascular diseases (CVD) have a large impact on mortality, with 17.9 million annual deaths worldwide,¹ and coronary artery

Strengths and limitations of this study

- COroNary Calcium scoring as fiRst-line Test to dEct and exclude coronary artery disease in GPs patients with stable chest pain (CONCRETE) determines the efficiency of CT coronary calcium scoring for diagnosing and excluding coronary artery disease in primary care patients with stable chest pain.
- CONCRETE gives insight into downstream testing and (unnecessary) referral rates of both strategies.
- CONCRETE gives insight into the cost-effectiveness of the CT CCS-based strategy and the standard of care in primary care.
- CONCRETE may initiate a change in primary healthcare policy for patients with stable chest pain.
- CONCRETE is an implementation study based on the Dutch Healthcare situation in which patients with stable chest pain first visit their general practitioner.

disease (CAD) as a leading cause.² From 2015 to 2040, the number of adults with CAD is expected to increase with about 50% in the Netherlands.³ CAD is often expressed by chest discomfort⁴; for diagnostic purposes, physicians use three core symptoms to describe typicality of chest pain: (1) retrosternal complaints; (2) complaints provoked by exertion, cold, emotional stress or heavy meals and (3) complaints that are relieved with the rest and/or within 2–15 min after using sublingual nitroglycerine.⁴ Presence of all three symptoms indicates typical angina pectoris (AP). If two out of three symptoms are present, the chest pain is called atypical AP, and patients presenting with none or one

of the three symptoms are determined as having non-specific thoracic complaints.

In the Dutch healthcare system, the general practitioner (GP) is usually the first physician a patient consults with chest discomfort. Chest pain is the primary reason to contact the GP in about 4% of the consultations.⁵ In only 10%–15%, obstructive CAD will eventually be diagnosed as cause of the symptoms.⁵ The challenge for the GP is to diagnose CAD based on symptoms, age and sex.⁵ Distinguishing life-threatening and non-life-threatening diseases is essential for the treatment of patients, but may be challenging in particular in case of atypical AP or non-specific thoracic complaints.^{5,6} The Dutch College of General Practitioners (*Nederlands Huisartsen Genootschap (NHG)*) clinical Standard for Stable AP serves as the guideline for GPs regarding, among others, referral for additional testing and treatment strategies.^{5,7} According to the prior clinical standard, the GP could order exercise electrocardiogram (ECG).⁵ However, exercise ECG is known to have suboptimal sensitivity and specificity for CAD. Furthermore, in clinical practice an important part of this heterogeneous group of patients was sent directly to the cardiologist for evaluation.⁸ The most recent standard (as of January 2020) has been adapted to this practice, and recommends that patients with typical and atypical AP be referred to the cardiologist (without first having exercise ECG ordered by the GP).⁷ In non-specific thoracic complaints, the GP should consider diseases other than CAD, unless there is reason to regard CAD as a possible cause. A recent Dutch nationwide analysis based on data from 2012 shows that 61% of patients who are referred to the cardiologist undergo an exercise ECG, 22% receive no testing and 19% undergo (also) a non-invasive CAD imaging test, most often an ischaemia test.⁸ Presently, in the Netherlands, about 105 000 patients are referred to the cardiologist (52% women) each year.⁸ Ultimately, only 5% of men and 1% of women have obstructive CAD requiring invasive treatment.⁸ There is a clinical need in patients with chest pain to optimise diagnostic management and referral to the cardiologist. An accurate diagnostic and prognostic tool could help GPs in determining the likelihood of CAD and guide patient management. At this moment, the Dutch GP does not have access to advanced imaging tests for CAD, such as computed tomography (CT). Furthermore, the most commonly performed first test, exercise ECG, apart from suboptimal accuracy for obstructive CAD, cannot detect early stages of CAD. A sensitive test for early diagnosis of CAD, including non-obstructive stages, will allow earlier treatment based on the Dutch guideline for cardiovascular risk management (CVRM).⁹ Early treatment could potentially allow a reduction in the incidence of major acute cardiac events (MACE).

In this paper, we present the rationale, objectives and study design of the COroNary Calcium scoring as fiRst-line Test to dETect and exclude coronary artery disease in GPs patients with stable chest pain (CONCRETE) trial. In CONCRETE, we investigate the impact of giving GPs

direct access to CT coronary calcium scoring for the diagnosis and exclusion of CAD in a pragmatic randomised trial. The coronary calcium score (CCS) is a robust, quantitative measure of coronary calcification based on non-contrast, low-dose ECG-triggered CT with a standardised protocol for scanning and postprocessing, with virtually no contraindications. In outpatient cardiology clinic setting, the CCS has proven to have better diagnostic and prognostic power than exercise ECG.^{10–12} The CCS has a negative predictive value of 93%–99% for obstructive CAD in symptomatic patients,^{13–17} and 99% for MACE, with similar prognostic results in men and women.^{14,15} The sensitivity of CCS for obstructive CAD is 95%–99%.^{18–20} The severity of coronary calcification is strongly related to CAD burden,^{13,21,22} ischaemia²² and MACE,^{23–25} and allows for early detection of patients with non-obstructive CAD, who can then receive early treatment.^{26–29} Research in symptomatic patients undergoing exercise ECG and calcium scoring in an outpatient cardiology setting showed that the CCS can be safely used for patient stratification, with no events after 1 year in patients with CCS <10 (52% of all patients), and better identification of patients with obstructive CAD and/or subsequent coronary events.¹¹ Other Dutch studies in outpatient cardiology setting have confirmed these findings, and indicate that CCS improves diagnostic and prognostic stratification compared with exercise ECG testing.^{10,12} A recent Dutch study in 1551 cardiology outpatients with chest discomfort and low or intermediate CAD probability showed a CCS 0 in 48%, CCS 1–100 in 32%, CCS 101–400 in 14% and CCS >400 in 6%.²⁴ Only 3% of patients with CCS 0 had obstructive CAD on CT coronary angiography. During a follow-up of nearly 2 years, the MACE rate was 0.3% in CCS 0, 1% in CCS 1–100, 4% in CCS 101–400 and 7% in CCS >400.²⁴ A CCS of 0 was found to have a safe and efficient approach to exclude CAD in patients with low-to-intermediate pretest probability, while probability of obstructive CAD increased particularly from a CCS of 100.^{10,20,23,25} The CAD consortium recently reported that inclusion of calcium scoring in risk stratification tools improves prediction of CAD probability in patients with chest pain undergoing invasive or CT coronary angiography.³⁰ In contrast to exercise ECG, the CT CCS test detects also early stages of CAD. Early detection of CAD combined with early treatment can potentially prevent adverse cardiac events, lower the burden of disease and increase the patient's life expectancy.^{3,31} It remains uncertain whether the diagnostic accuracy of CT CCS is the same in primary care, although prior outpatient studies included mainly low and intermediate probability patients, similar to the risk profile of GP patients. Also, the impact of implementation of calcium scoring in a GP setting on CAD diagnosis and treatment rate is unknown. CONCRETE is an implementation study focusing on the Dutch healthcare system in which the GP is usually the first physician a patient consults with chest discomfort. We hypothesise that GP access to the CT CCS test allows for early diagnosis of CAD and treatment, more efficient

Box 1 Primary and secondary objectives of CONCRETE

Primary objective (cluster-based)

1. To determine the increase in detection/treatment rate of CAD in GP offices with the CT CCS-based strategy, compared with GP offices with the standard of care strategy, in patients with atypical AP and non-specific thoracic complaints.

Secondary objectives (patient-based)

1. To establish the diagnostic yield to diagnose obstructive CAD, for both strategies.
2. To establish the effectiveness in terms of CAD diagnosis and exclusion of GP referral to the cardiologist for the calcium score cluster as well as the time to (exclusion of) CAD diagnosis.
3. To compare downstream diagnostic testing and treatment for both strategies.
4. To evaluate whether diagnostic stratification, in particular cut-offs for referral to the cardiologist, can be optimised for the calcium score.
5. To estimate the effect of calcium scoring versus the standard of care on quality of life and cardiac complaints after 6, 12 and 24 months.
6. To estimate the effect of calcium scoring on reduction of MACE (after 2 years).
7. To derive data on the costs per diagnosis of obstructive and diagnosis of non-obstructive CAD in the setting of calcium score testing versus the standard of care.
8. To estimate the cost-utility of implementing the calcium score test in GP setting.
9. To develop machine learning tools to evaluate big data on (combinations of) symptoms and family history/risk factors, in relationship to CAD.
10. To establish and visualise relationships between (combinations of) symptoms and family history/risk factors and probability of CAD, using innovative techniques for big data analysis; these results will form the input for a risk assessment tool to be developed.

All analyses will be analysed by sex.

AP, angina pectoris; CAD, coronary artery disease; CCS, coronary calcium score; CONCRETE, COroNary Calcium scoring as fiRst-linE Test to dEct and exclude coronary artery disease in GPs patients with stable chest pain; GP, general practitioner; MACE, major acute cardiac event.

referral to the cardiologist and a reduction in healthcare-related costs.

Objectives

The primary objective is to determine the increase in detection and treatment rate of CAD in GP offices with the CT CCS-based strategy, compared with GP offices with the standard of care strategy, in patients presenting with atypical AP and non-specific thoracic complaints. The primary end point is the (early) CAD diagnosis registration as expressed by the CVRM registration rate.⁹ The primary and secondary objectives of the trial are listed in box 1.

METHODS AND ANALYSIS

Study design and setting

CONCRETE is a pragmatic multicentre study with a cluster-randomised design. This type of design is

increasingly used in primary care for the evaluation of healthcare interventions.^{32–35} An overview of the study design is provided in figure 1. From January 2019 onward, GPs are recruited from multipractice GP organisations in urban and rural regions from the provinces Gelderland, Groningen, Limburg and Overijssel in the Netherlands. The initial aim was to include 80 GP offices. Through a permutated block randomisation scheme (1:1), GP offices are divided into two equally large strategy groups; one group of GP offices refers patients for CT CCS testing and another group of GP offices provides the standard of care (figure 1). The primary analysis will be performed in the two clusters. After GP office randomisation, approximately 800 patients will be included over a period of at least two years in both diagnostic strategies (figure 1).

Enrolment of clusters, GP offices (for primary analysis)

GPs and medical staff of the GP offices, from the collaborating multipractice GP organisations, are informed of the trial through written information (eg, information brochure, newsletters and website) and verbal information (eg, presentations during mandatory educational courses and local policy meetings) by the (local) researchers of the trial. Then, every GP office is contacted by the researchers to schedule a face-to-face appointment in order to discuss the trial and their potential participation. GP offices are included cluster-wise if they are willing to take part in the trial and provide written informed consent. Thereafter, the GP office is randomised into one of the two strategy groups using a computerised randomisation scheme and is informed by the researcher by telephone or per email of the randomisation outcome. If a GP experiences difficulties in fulfilling the study tasks, a meeting will take place with the researcher to find solutions in order to sustain participation. In case GPs wish to discontinue their participation, collaboration is ended and data of patients included up to that moment are used in the study, since written consent of the patients was obtained.

Enrolment of patients (for secondary analysis)

For the individual-based analyses, patients with chest discomfort, either atypical AP or non-specific thoracic complaints (figure 1), with indication for further diagnostic evaluation as determined by the GP, will be informed about the trial and asked to participate by the GP. Men of 40 years and older, and women of 45 years and older will be included. Exclusion criteria for individual patients are pregnancy, unwillingness to provide written informed consent for the individual level (secondary) outcomes and prior diagnosis of CAD (percutaneous coronary intervention, coronary artery bypass surgery, myocardial infarct, stable CAD).

Patients receive an information brochure and informed consent form. The signed informed consent form is returned to the GP and sent to the researcher. For a patient, participation comprises agreement to share clinical data from the GP system with the researchers, and

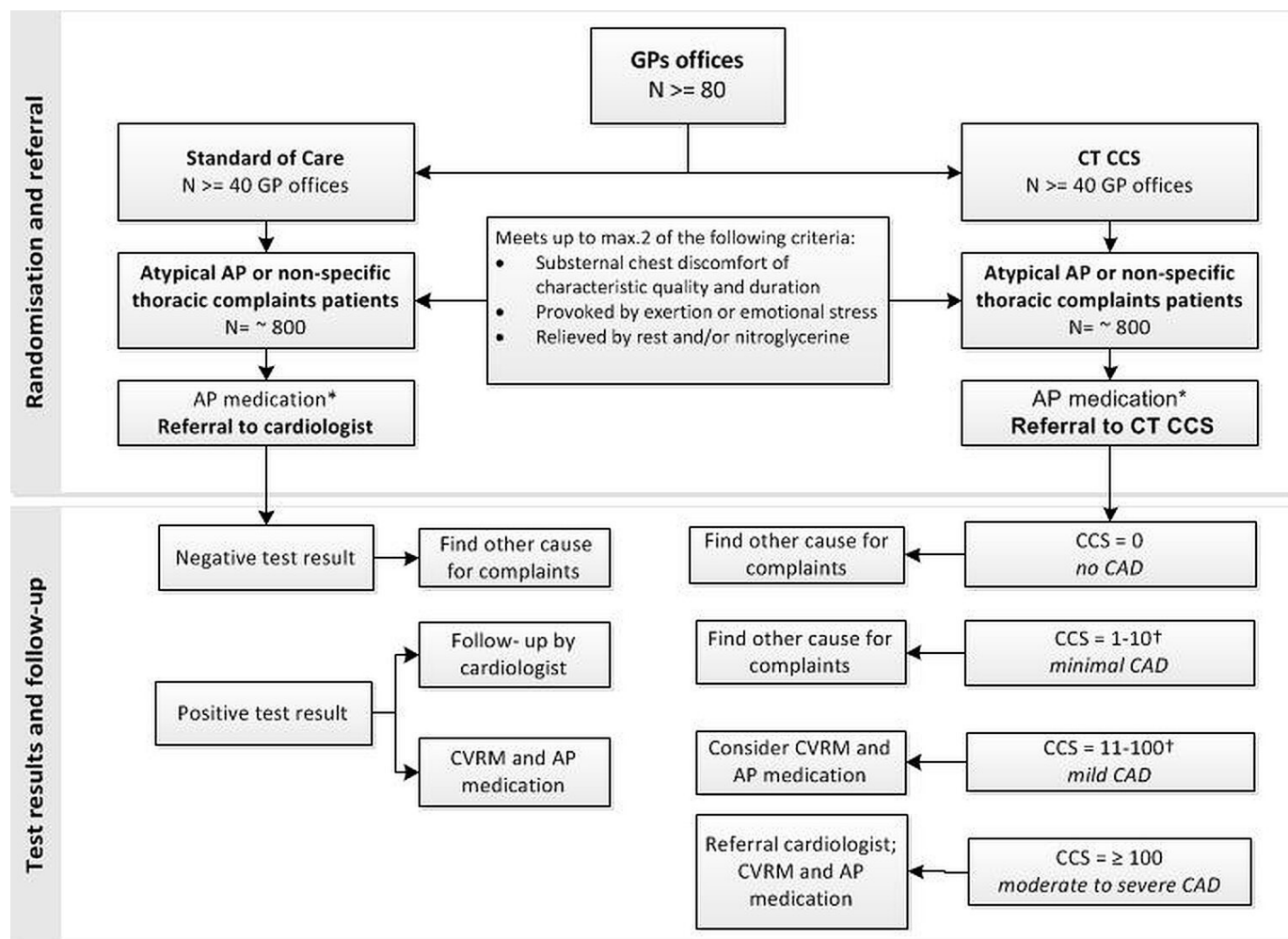


Figure 1 Flow chart of randomisation, referral and test results. AP, angina pectoris; CAD, coronary artery disease; CCS, coronary calcium scoring; CVRM, cardiovascular risk management. *Guidelines indicate patients with stable chest pain have to start with medication for AP before undergoing diagnostic testing.⁵ †With CCS ≥75th percentile for age and sex, the GP is advised to consider classifying the patient one CCS category higher.

filling in questionnaires. The first questionnaire is sent to the patient either digitally or by letter, to be completed prior to the visit to the cardiologist or the CT scan. Patients will receive digital reminders to fill in the questionnaires. The diagnostic management executed by the GP (CT CCS or standard of care) does not depend on the signing of the informed consent by the patient; this depends on the cluster to which the GP office is randomised.

CT coronary calcium scoring strategy

The CT CSS is carried out in accordance with the routine procedure of the participating radiology departments, and performed on the routine (single or dual source) CT scanner used for cardiac imaging in these departments. In this way, adherence to common clinical practice and generalisability of results are optimised. In practice, the scan and reconstruction protocol for CCS is rather standardised. In all centres, the CT scan consists of an ECG-synchronised acquisition without intravenous iodine contrast, during breath hold. The entire heart is included in the scan range. Images are commonly acquired around 60% of the cardiac cycle. Tube voltage is 120 kVp and tube

current is generally set at around 80 mAs as reference. Images are reconstructed with a slice thickness of 3.0 mm and slice increment of 1.5 or 2.5 mm. The radiation dose is approximately 0.5–1 mSv, which is <50% of the annual background radiation dose in the Netherlands.^{36 37} The CCS is calculated according to the method by Agatston *et al.*³⁸ Patients receive AP medication subscribed by the GP while awaiting the results of the CT CCS test, the preventive or pre-emptive treatment for stable CAD can be stopped if the diagnosis CAD has been ruled out, as recommended by the NHG Standard for Stable AP.⁷ The GP will be informed of the CT CCS result based on the radiologist report (figure 1). The report of the radiologist consists of the total CCS and the CCS per coronary artery (right coronary artery, left coronary artery, left anterior descending artery and left circumflex artery) and the Multi-Ethnic Study of Atherosclerosis (MESA) percentile.³⁹ CCS results are categorised as CCS of 0 (no CAD), 1–10 (minimal CAD), 11–100 (mild CAD), 101–399 (moderate CAD) and ≥400 (severe CAD), respectively. When CCS is 0 or 1–10, GPs are advised to stop AP

medication and consider other causes for the complaints. In case of CCS 11–100, continuation of AP medication and inclusion of these patients into CVRM is to be considered.⁹ In patients with CCS 101–400 and CCS ≥ 400 , AP medication is continued and in addition, the patient will be included into CVRM⁹ and referred to the cardiologist. In case of a CCS result above the 75th percentile for age and sex based on MESA percentile,³⁹ the GP is recommended to classify the patient one CCS category higher than the category matching the absolute score, in view of the premature atherosclerosis and the associated long-term cardiovascular risk (figure 1).^{40 41} The categories are based on recent literature from Dutch studies,^{9 11 22 41} and on the experience of physicians who have been applying the CCS for years in practice. The management advice for the CCS categories is based on discussions with cardiologists, GPs and radiologists, and is not obligatory but meant as guidance. The GP discusses the results with the patient. Decisions regarding patient management remain at the discretion of the GP, who takes all available patient information into account.

Standard of care strategy

In this arm, all steps are in agreement with routine standard of care, as determined by the NHG Standard for Stable AP. Thus, patients with atypical AP or with non-specific thoracic complaints in whom the GP wants to exclude CAD as underlying cause, will be referred to the outpatient cardiology clinic. There, evaluation of the patient takes place with additional testing at the discretion of the attending cardiologist in accordance with the guidelines of the European Society of Cardiology (ESC).⁴² GPs will be informed of the findings and cardiac diagnosis based on the letter from the cardiologist (figure 1).

Data collection

The primary end point is an increase in CAD diagnosis and treatment in patients in the CT CCS strategy compared with the standard care strategy. The detection and treatment rate of CAD in GP offices is based on the CVRM registration rate. According to the CVRM guideline, patients are registered into the CVRM registry when they have high risk of developing CVD and/or are diagnosed with CVD.⁹ CVRM registry data are derived from the GP electronic registration system with registration date 1 year before baseline, at baseline and 1 and 2 years after baseline. Individual patient data are collected over a period of 2 years, using four questionnaires (table 1) containing questions with regard to experienced chest pain complaints,⁴³ quality of life (QoL) (EQ-5D-5L)^{44 45} and heart-related QoL (HeartQoL).^{46 47} In addition, the cardiovascular risk profile of the patient, and information on downstream testing and CAD diagnosis will be collected through the electronic patient dossier (table 2).

Sample size

We used sample size calculations for cluster randomised trials.⁴⁸ In order to detect a 7.5% difference in CVRM registrations between the two clusters, with a power of 80%, a significance level of 5% and an intracluster correlation of 0.01, inclusion of 36 GP offices in each cluster would be necessary, with an estimated total of 20 patients per GP office (10 patients per year per GP office). To anticipate unforeseen circumstances, such as drop out, we initially decided to include 40 GP offices in each strategy group. The calculations are based on the assumption that yearly about 50 patients will consult their GP with atypical AP and non-specific thoracic complaints,⁵ with an estimated 10 patients who will be referred for additional evaluation

Table 1 SPIRIT schedule of enrolment, test strategies and assessments

	Study period					
	Enrolment	Baseline	Execution of strategy	Postexecution strategy		Close-out
Time points	0	0	1–2 weeks	6 months	12 months	24 months
Enrolment						
Eligibility screening by GPs	x					
Informed consent	x					
Test strategies						
Standard care strategy			x			
CT CCS strategy			x			
Assessment						
CVRM GPs offices		x			x	x
Risk factors	x					
Chest pain complaints		x				
Quality of life		x		x	x	x
Heart-related quality of life		x		x	x	x

CCS, coronary calcium score; CVRM, cardiovascular risk management; GP, general practitioner; SPIRIT, Standard Protocol Items: Recommendations for Interventional Trials.



Table 2 Cardiovascular risk assessment items

General health assessment

Smoking habits	Tobacco use per day. Total number of smoking pack years.
Alcohol use	Mean number of glasses per day. General behaviour with regard to alcohol use using the 5SH1 test.
Family history	Fatal or non-fatal cardiovascular diseases in a first-degree family member?
Diabetes mellitus	Does the patient have diabetes mellitus?
Rheumatoid arthritis	Does the patient have rheumatoid arthritis?

Physical assessment

Systolic blood pressure	Mean of three blood pressure measurements by general practitioner.
Body mass index (BMI)	Determination of BMI (kg/m ²)

Laboratory assessment

Blood lipids spectrum	▶ HDL ratio ▶ LDL ratio ▶ Total cholesterol ratio ▶ Total cholesterol-HDL ratio ▶ Triglycerides ratio
Blood glucose level	Fasting glucose ratio
Estimated glomerular filtration rate (eGFR)	eGFR ratio

Measurements to assess cardiovascular risk of the patients.

HDL, high-density lipoprotein; LDL, low-density lipoprotein; 5SH1, FiveShot1 questionnaire, that is used to determine alcohol use or possible alcohol abuse.

to diagnose or exclude CAD.⁶ In the CT CCS strategy, a CCS of 0 are expected in 45% of patients, a CCS of 1–10 in 10%, a CCS of 11–100 in 20% and a CCS >100 in 10%, actual percentages will only be known during this study.^{24 30 49 50} In the standard care strategy, 27.5% of the patients are expected to be included in CVRM registry for CAD diagnosis/treatment, and 35% of the patients in the CT CCS test strategy. These percentages are based on similar populations from previously published Dutch research.^{10 51 52}

COVID-19 pandemic

The trial was stopped due to the COVID-19 pandemic from March until June 2020. However, even after the re-start in July 2020, the inclusion of GP offices and patients was so far (June 2021) severely slowed due to effects of the COVID-19 pandemic (among others, lower GP consultation rates, procedures to restart clinical practice in adherence to COVID-19 regulations and vaccination procedures). The COVID-19 pandemic urged us to realign the patient inclusion rate, sample size and the interdependent calculations of the trial. To reduce the impact of COVID-19 on the progress of CONCRETE, we aim to increase the number of participating GP offices from 80 to 130.

Data analysis

Baseline characteristics (including baseline rate of CVRM registration) will be summarised by mean (SD), median (IQR) and percentage. The primary outcome is the (early) CAD diagnosis/treatment registration as

expressed by the CVRM registration rate at GP office level. The increase for each individual GP office will be calculated by subtracting the percentage of registrations at baseline from the percentages of CAD diagnosis/treatment registrations at follow-up. The difference in increase of CVRM registration rate of the two clusters will be compared with an independent t-test or non-parametric test, depending on distribution. A multiple multilevel linear regression will be performed to adjust for potential confounders (differences between GP practices), such as GP practice size and the ratio of men/women per practice (characteristics of each practice). In case of missing data, due to loss to follow-up, multiple imputations will be used.

In the CT CCS strategy, receiver operating characteristic analyses will be used to find the optimal CCS and CCS percentile for referral to the cardiologist, by sex and age.

Follow-up

Patients will be followed for acute myocardial infarction and sudden cardiac death as well as CAD diagnostic procedures, cardiac interventions and diagnostic/treatment costs, for up to 5 years; source data about CAD diagnosis, diagnostic procedures and follow-up cardiovascular events will periodically be obtained from the GP electronic dossier. As part of the current grant, follow-up procedures up to 2 years after the inclusion period are covered. A follow-up up to 5 years will be performed if additional funding is secured.

Quality of life

QoL and HeartQoL will be compared between the strategies.^{45 53} Patients fill in both questionnaires prior to the CT CCS or cardiologist referral, and after 6, 12 and 24 months. Multiple linear regression will be performed, to adjust for potential confounders. Analyses will be performed on an intention-to-treat basis.

Cost-utility

Costs of diagnosis are recorded in this clinical trial and used to calculate the costs per CAD diagnosis (obstructive and non-obstructive) for each strategy. Additionally, costs for events during follow-up are recorded for each strategy. Cost-utility analysis will be performed to reflect the balance between incremental monetary cost and incremental quality-adjusted life years resulting in an incremental cost-utility ratio (ICUR). A societal perspective will be used and discounting will be applied for costs (4%) and health outcomes (1.5%) according to Dutch guidelines.⁵⁴

A patient-level simulation model using a life-long time horizon will be created to assess the long-term impact of CT CCS access by GPs regarding costs and health outcomes. Due to the late onset of symptoms for CAD and the major impact of cardiovascular events caused by CAD, extrapolating the results to a lifelong time horizon is desired to determine the true impact of CT CCS. Clinical data will be used as input for the simulation model where possible. Bootstrapping is used to determine uncertainty surrounding these input parameters.⁵⁵ Model parameters beyond the scope of the clinical trial are supplemented by literature. Expert opinion will be used in case literature yielded inconclusive results. To reflect outcome (ICUR) uncertainty, a probabilistic analysis will be performed using the Monte Carlo approach with 5000 iterations.⁵⁶ Results of the probabilistic analysis will be visualised in a cost-effectiveness plane and cost-effectiveness acceptability curve. Finally, subgroup-specific cost-utility outcomes will be determined for age and sex.

Patients and public involvement

Patients were involved in the design and reporting of the study. Cardiovascular patients gave input on the information and questionnaires to be provided to patients. Patients are also included in the user committee of the study, which gives input on every aspect of the trial including recruitment and conduct. There is a specific part of the CONCRETE website dedicated to informing patients about the trial and its results.

ETHICS AND DISSEMINATION

CONCRETE has been approved by the Medical Ethical Committees of the University Medical Center of Groningen (UMCG) on 12 November 2018, V.2.0, under number 2018/404. The study will be conducted according to the principles of the Declaration of Helsinki, Brazil, October 2013,⁵⁷ and in accordance with

the Dutch Medical Research Involving Human Subjects Act (WMO).⁵⁸ Study data will be collected in a pseudonymised fashion and managed using REDCap electronic data capture tools hosted by the UMCG. Pseudonymised CT image data will be stored in the XNAT imaging data archive. We will handle personal data in compliance with the Dutch General Data Protection Regulation (AVG).⁵⁹ The trial has a Data Management Plan not accessible to the public, consisting—among other things—of information regarding metadata, the used software for data management and a platform for storing and sharing data. Monitoring of the data will be performed independently of the sponsor, by a monitor of the monitoring committee of the UMCG using a preconceived plan. The participating (University) Medical Centers are subject to internal audit programmes, randomly evaluating scientific studies, independently of the sponsor and investigators. The accredited METc will receive a summary of the progress of the trial once a year and will make the final decision to continue or (prematurely) terminate the trial. In case of (serious) events continuation of the trial will be discussed within the consortium and the accredited METc. However, due to the negligible risk associated with the CT CCS, we do not anticipate adverse events due to the implementation of CT calcium scoring in GP setting, nor a premature termination of the study. There is liability insurance for ancillary and post-trial care for those who could have suffered harm from trial participation. The study is registered in the Netherlands National Trial Register on <https://www.trialregister.nl/>. Information on CONCRETE for patients, physicians and researchers is available at the website www.concrete-project.nl. Results from the trial will be published in peer-reviewed journals with the consortium members as coauthors and presented at (inter)national conferences. We do not intend to use professional writers. Following Findability, Accessibility, Interoperability and Reuse of digital assets principles, data will be made available to third parties, after completion of the trial. Procedures and criteria for sharing of the data will be designed.

DISCUSSION

The CONCRETE study aims to determine whether direct GP access to CT CCS leads to earlier and more cost-effective diagnosis and treatment of CAD in patients with atypical AP or non-specific thoracic complaints, compared with standard of care. To our knowledge, CONCRETE is the first study that will test the implementation of CT CCS in primary care, as previous studies have been performed in secondary care settings.^{4 12 60 61} Furthermore, the study will assess and optimise sex-specific diagnostic stratification based on the CCS. In addition, CONCRETE will give insight into the QoL of these patients, downstream testing, (unnecessary) referral rates and the cost-effectiveness of both strategies. Implementation of CONCRETE findings could initiate a change in healthcare policy

early CAD may prevent future MACE or sudden cardiac death, although at this moment this has not yet been proven in randomised clinical trials.

CONCLUSION

CONCRETE is a pragmatic implementation study to determine the effectiveness of direct GP access to CT CCS to diagnose and exclude CAD in patients with atypical AP and non-thoracic complaints, compared with the standard of care. The CONCRETE results are expected to lead to change of GP referral indications for patients with atypical AP and non-specific thoracic complaints and change of reimbursement policies in healthcare.

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