Study to Improve Cardiovascular Outcomes in high-risk older patients (ICON1) with acute coronary syndrome: study design and protocol of a prospective observational study

Vijay Kunadian,1,2 R Dermot G Neely,3 Hannah Sinclair,1,2 Jonathan A Batty,1,2 Murugapathy Veerasamy,1,2 Gary A Ford,4 Weiliang Qiu5

To cite: Kunadian V, Neely RDG, Sinclair H, et al. Study to Improve Cardiovascular Outcomes in high-risk older patients (ICON1) with acute coronary syndrome: study design and protocol of a prospective observational study. BMJ Open 2016;6:e012091. doi:10.1136/bmjopen-2016-012091

ABSTRACT

Introduction: The ICON1 study (a study to Improve Cardiovascular Outcomes in high-risk older patients with acute coronary syndrome) is a prospective observational study of older patients (≥75 years old) with non-ST-elevation acute coronary syndrome managed by contemporary treatment (pharmacological and invasive). The aim of the study was to determine the predictors of poor cardiovascular outcomes in this age group and to generate a risk prediction tool.

Methods and analysis: Participants are recruited from 2 tertiary hospitals in the UK. Baseline evaluation includes frailty, comorbidity, cognition and quality-of-life measures, inflammatory status assessed by a biomarker panel, including microRNAs, senescence assessed by telomere length and telomerase activity, cardiovascular status assessed by arterial stiffness, endothelial function, carotid intima media thickness and left ventricular systolic and diastolic function, and coronary plaque assessed by virtual histology intravascular ultrasound and optical coherence tomography. The patients are followed-up at 30 days and at 1 year for primary outcome measures of death, myocardial infarction, stroke, unplanned revascularisation, bleeding and rehospitalisation.

Ethics and dissemination: The study has been approved by the regional ethics committee (REC 12/NE/016). Findings of the study will be presented in scientific sessions and will be published in peer-reviewed journals.

Trial registration number: NCT01933581: Pre-results.

Strengths and limitations of this study

- Older patients with non-ST-elevation acute coronary syndrome represent a high-risk population, who remain understudied in contemporary cardiovascular research.
- This prospective cohort study is designed and powered to identify risk factors for adverse outcomes, at 30 days and 1 year, in patients aged ≥75 years undergoing invasive management of non-ST-elevation acute coronary syndrome.
- This study will evaluate the role of frailty, using a well-defined frailty index, and invasive imaging modalities (including optical coherence tomography and virtual histology intravascular ultrasound) as determinants of clinical outcome and also evaluate the quality of life in this age group.
- Limitations include (1) the non-randomised character of this study, which is not able to derive definitive insights regarding the causality of factors associated with clinical outcomes, and (2) that intracoronary imaging will be performed in only a subset of patients recruited, owing to anatomical contraindications and patient wishes.
- The results of this study will enable improved risk stratification for older patients presenting with non-ST-elevation acute coronary syndrome and will have implications for the design of future clinical trials in this high-risk population.

INTRODUCTION

In the general population, ischaemic heart disease (IHD) is the leading cause of death worldwide.1 Mortality due to IHD increases steeply among those aged >70 years.2 In 2010, in the UK, more than twice as many individuals >75 years of age (n=55 028) died from IHD, compared to younger individuals <75 years (n=25 540).3 According to the Myocardial Ischaemia National Audit Project Database annual public report 2012–2013, there were 80 974 admissions with a final diagnosis of myocardial infarction (MI). Of these, 60% had non-ST-elevation myocardial infarction (NSTEMI). Of the patients with NSTEMI, 59% were >70 years of age (26% were aged 70–79 years, 26% were aged 80–89 years and 7% were aged ≥90 years).4

For numbered affiliations see end of article.

CrossMark
Mortality benefit from advances in the management of acute coronary syndrome (ACS) has largely been realised in patients aged <65 years. There has been an increase in IHD burden in older patients, who are at risk of poorer outcomes due to frailty and comorbidity.

Until recent years, there has persisted a paucity of evidence from clinical trials and studies to inform the management of ACS in older patients. More than half of all randomised controlled trials for ACS failed to enrol participants >75 years of age and, even in those that did, only 9% were >75 years of age. Notable studies, recruiting patients >75 years of age, have been reported in recent years, in the context of invasive and non-invasive management of ST-elevation MI and non-ST-elevation ACS. Evidence-based recommendations from trials do not account for age-related differences in physiology, disease and comorbidities, which may alter the risk–benefit profile of cardiovascular treatments and interventions. The age mismatch between trial and community populations begins at 75 years and widens with age. Furthermore, older people who are included in trials have lower than expected rates of traditional cardiovascular risk factors, fewer comorbidities and better renal function than the community population. Risks and benefits derived from trials cannot always be extrapolated to older patients in daily clinical practice due to the differences between the patient groups and their baseline characteristics.

In the ageing population, there is increasing evidence for the association of cardiovascular disease (CVD) and frailty. Depending on the frailty scale used and the population studied, almost half of the patients with CVD can be identified as frail. There is an increased risk of mortality and major adverse cardiovascular events in frail patients with CVD, especially those undergoing invasive procedures or suffering from coronary artery disease and heart failure. In patients aged >75 years, frailty was strongly and independently associated with inhospital mortality (OR 4.6; 95% CI 1.3 to 16.8) and 1 month mortality (OR 4.7; 95% CI 1.7 to 13.0). At 1 year, there was a significant increase in mortality among frail patients compared with non-frail patients (HR 4.3, 95% CI 2.4 to 7.8). Similarly, in >65-year-old patients, frailty was associated with increased long-term mortality and MI among patients undergoing percutaneous coronary intervention (PCI).

No studies have been performed in older patients undergoing an invasive treatment strategy to evaluate predictors of poor outcomes or to develop strategies to improve outcomes following ACS. The ACS and PCI risk models that are currently available were mainly derived from patients <65 years of age and, hence, cannot be applied to the increasing proportion of older patients (aged >75 years) with ACS managed by contemporary treatment. The goal of Improve Cardiovascular Outcomes in high-risk older patients with acute coronary syndrome (ICON1) study is to determine the predictors of adverse outcomes (death, MI, stroke, unplanned revascularisation, bleeding and rehospitalisation for any reason) at 1 month and at 1 year following invasive management of non-ST-elevation acute coronary syndrome (NSTEACS) in older patients and to develop an integrated risk score to predict adverse outcomes at 1 year that will inform clinical decision-making. In addition, the impact of contemporary NSTEACS management on the quality of life will be assessed.

**HYPOTHESIS**

Frailty and comorbid status in older patients are associated with worse outcomes following invasive treatment for NSTEACS.

**TRIAL DESIGN**

The study has been designed as a multicentre, prospective, observational study of patients aged ≥75 years undergoing invasive management (coronary angiography with a view to revascularisation) for NSTEACS.

**METHODS**

**Study setting**

This ongoing, multicentre, observational study is being conducted in two tertiary cardiac care hospitals in the North-East of England. The Freeman Hospital, in Newcastle upon Tyne, is a tertiary cardiac centre with a catchment population of 2 million. Approximately 300 PCI procedures are performed each year. The James Cook University Hospital, in Middlesbrough, performs ~1750 PCI procedures every year. The study participants are recruited from patients referred to these hospitals from the neighbouring district general hospitals for invasive treatment of NSTEACS. Patients are diagnosed on the basis of clinical symptoms, electrocardiography criteria and high-sensitivity troponin testing, in line with guidelines transferred the day before or on the day of procedure to the tertiary hospitals. Prospective ICON1 patients are identified from an electronic referral system and, on arrival to the tertiary hospitals, are approached for recruitment into the study. The research team explains the study to the patients and a patient information sheet is provided. If a patient agrees to participate in the study, a written informed consent is obtained. All patients screened for the study are entered in a screening log, with details regarding the patients consented, declined and consented but not recruited (due to alternative diagnosis following coronary angiography). The inclusion and exclusion criteria are shown in box 1. Recruitment to the study started in October 2012 with the 1-year follow-up is projected to reach completion in December 2016.

**Treatment protocol**

During the course of the study, the patients were treated according to contemporary evidenced-based guidelines, as directed by an interventional cardiologist, at the time...
of study enrolment.\textsuperscript{20, 21} According to standard practice, the patients are revascularised by PCI or coronary artery bypass graft surgery. The patients may also be managed medically, if deemed not appropriate for either of the revascularisation strategies at the discretion of the operating cardiologist.

**Data collection**

Data are collected on standardised case report forms by members of the research team. The data collected include demographics, baseline characteristics, and details of coronary angiography and/or PCI. Periprocedural complications and in-hospital complications are recorded. Further data are collected on the cardiovascular status, Canadian Cardiovascular Society (CCS) angina grade, New York Heart Association (NYHA) dyspnoea grade, frailty category, functional health status, quality of life and cognitive status. These are listed in box 2. The assessments and techniques used for the above data collection are discussed in the following sections. The study flow chart is displayed in figure 1. All questionnaires were administered verbally, in person and by a trained, clinical researcher. Appropriate training was provided to researchers, ensuring that these scripted questionnaires were performed, and results recorded, in an unbiased fashion.

**Frailty and comorbidity assessments**

Frailty is assessed by Fried Frailty Index, derived from Cardiovascular Health Study\textsuperscript{22} and Rockwood Frailty Index, derived from Canadian Study of Health and Aging.\textsuperscript{23} The Fried Frailty Index is based on assessing five criteria, comprising subjective answers provided by the patient (regarding weight loss, physical energy, physical activity) and objective assessment (hand grip strength). A score of 0 is categorised as robust, 1 or 2 as intermediate or pre-frail and 3 or more as frail (see online supplementary appendix 1). The Rockwood Frailty Index is, based on the assessment by the researchers, grouped into categories 1–7, from very fit to severely frail, depending on functional status and independence/dependence on others for activities of daily living (see online supplementary appendix 2).

In addition, the Charlson Comorbidity Index,\textsuperscript{24} a method of predicting mortality based on a weighted index of the number and seriousness of comorbid conditions, is evaluated for each patient. The Charlson
Comorbidity Index has been demonstrated to be an appropriate indicator of in-hospital and 1-year outcomes in the setting of ACS.  

### Functional status and quality-of-life measures

The Short Form-36 Standard (SF-36 Standard) health survey is completed by each patient prior to discharge from the hospital and at 1-year follow-up to assess functional health and quality of life. The responses will be used to obtain physical component summary score and mental component summary score. In addition, the EQ-5D-3L questionnaire is used to assess the health outcome of each patient at discharge and 1-year follow-up.

### Cognitive status assessment

Atherosclerosis is associated with increased risk of cognitive impairment in older patients. To assess the cognitive status of patients during admission, the Montreal Cognitive Assessment (MoCA) test is used. The MoCA has been shown to have high sensitivity in screening patients with known CVD for mild cognitive impairment, even in a non-memory clinic setting. This test is repeated at 1-year follow-up.

### Biomarker sampling

Blood samples are collected at the time of coronary angiography and/or PCI for biomarker analysis. Serum for biomarkers is stored for analysis in batches. Peripheral blood mononuclear cells are separated by centrifugation techniques for storage at −80°C for analysis of telomeres and telomerase activity. High-sensitivity C reactive protein (hsCRP), parathyroid hormone and total vitamin D are analysed. Full blood count, renal function, blood glucose, cholesterol and high-sensitivity cardiac troponin T levels are measured in patients as part of our routine care.

Inflammation plays a central role in acute thrombotic complications of unstable atherosclerotic coronary plaque. Increased levels of markers of inflammation predict CV outcomes following ACS. Inflammatory markers including myeloperoxidase, hsCRP and soluble CD40 ligand have been associated with ACS and have been shown to predict the outcome. The patients with ACS have decreased levels of anti-inflammatory ω-3 fatty acids (eicosapentaenoic acid and docosahexaenoic acid). Increased lipoprotein-associated phospholipase A2 activity has been associated with increased cardiovascular event rates. An elevated level of asymmetric dimethyl arginine is a strong and independent predictor of adverse outcomes following ACS. Interleukin-6 (IL-6) levels in the serum were increased in patients with ACS. IL-6 expressed in atherosclerotic plaques may increase plaque instability. Elevated IL-6 was a predictor of 6-month and 12-month mortality in patients with unstable coronary artery disease. Tumour necrosis factor-α (TNF-α) is a proinflammatory cytokine associated with myocardial dysfunction and remodelling following ACS. In patients with recent MI, increased levels of TNF-α were associated with adverse cardiovascular outcomes (recurrent MI and cardiac death). Vitamin D deficiency has been associated with elevated CAD burden and worse cardiovascular outcomes. These biomarkers will be analysed in this group of ≥75-year-old patients to enable the determination of predictors of adverse CV outcomes at 1 year. Telomere shortening has been associated with ageing and senescence, and shorter telomere length is associated with increased cardiovascular risk and mortality. Shorter telomere length predicted high-risk plaque morphology on virtual histology intravascular ultrasound (VH-IVUS). Whether shorter telomere length is a predictor of adverse events among older patients undergoing PCI is not known and will be evaluated in this study.

### MicroRNA analysis

MicroRNAs (miRNAs) are small non-coding RNAs that post-transcriptionally inhibit gene expression. In the...
past few years, miRNAs have emerged as key tools for the understanding of IHD pathophysiology, with great potential to be used as new biomarkers and therapeutic targets. miRNAs seem to possess ideal characteristics to be used as disease biomarkers, as they are detectable in biofluids in a reproducible and stable fashion, even after years of sample storage and freeze–thaw cycles. In the blood, circulating miRNAs are found mainly within extracellular vesicles, such as exosomes, microvesicles and apoptotic bodies, and, to a lesser extent, associated with vesicles, such as exosomes, microvesicles and apoptotic bodies, and to a lesser extent, associated with vesicles, such as exosomes, microvesicles and apoptotic bodies. Several studies have demonstrated elevated or decreased levels of specific circulating miRNAs in patients with ACS. However, few have addressed their prognostic value with regards to major cardiovascular events or death, especially among older cohorts of patients presenting with NSTEACS.

The levels of nine circulating miRNAs, known to be differentially expressed in patients with ACS (miR-21-5p, miR-126-5p, miR-132-3p, miR-133a-3p, miR-142-3p, miR-150-5p, miR-208-3p, miR-223-3p and miR-320a), will be quantified by reverse transcription quantitative PCR, in serum and circulating microvesicles (isolated from an additional 200 µL of serum) and correlated with clinical variables with a view to assess their value as a prognostic biomarker in older patients with NSTEACS.

Invasive coronary artery imaging
Postmortem studies have identified that vulnerable plaques, with specific morphological characteristics, are implicated in the pathophysiology of ACS. These plaques, which are prone to erosion and rupture, have inflamed fibrous caps, rich in macrophages, overlying a lipid pool. Burke et al examined the hearts of 113 men who had died suddenly and found that 95% of ruptured plaques had fibrous caps <65 µm thick (mean thickness 23±19 µm) with an infiltrate of macrophages. ICON1 aims to identify whether the increased mortality in the older population with ACS is due to an increased prevalence of these vulnerable thin-capped fibroatheromas (TCFAs). Following diagnostic coronary angiography, the patients undergo VH-IVUS imaging and optical coherence tomography (OCT) imaging in all three coronary arteries prior to PCI, where feasible and not contraindicated, and VH-IVUS imaging post-PCI in the culprit vessel at the discretion of the operating cardiologist.

Virtual histology intravascular ultrasound
The greyscale IVUS image uses only the amplitude of the reflected ultrasound wave. VH-IVUS uses spectral analysis of the frequency and power of the reflected wave to generate a more accurate reflection of the tissue subtypes present within the vessel wall. This can then be used to differentiate plaque components (fibrous, fibro-fatty, dense calcium and necrotic core) and identify high-risk vulnerable plaques. Although VH-IVUS lacks the resolution to identify the thin fibrous cap of the TCFA, it is well placed to accurately identify the necrotic core of these plaques. A 20 MHz, phased-array Eagle Eye Platinum catheter is mounted on an R-100 pullback device and connected to either an integrated S5i system or a mobile S5 tower. Image acquisition is performed at a pullback speed of 0.5 mm/s and is ECG-gated to ensure one frame is acquired per cardiac cycle. The maximum length of all three coronary arteries is imaged, where feasible and not contraindicated. The data are anonymised and transferred to DVD for off-line data analysis. The operator is blinded to these data.

VH-IVUS data analysis is performed using the Medis QIVUS software v2.0 (Leiden, the Netherlands). Contours are drawn manually around the external elastic membrane and lumen of the vessel for each greyscale IVUS frame, excluding any ring-down artefact or previously stented segments. The software then calculates several parameters such as minimum lumen area and diameter, per cent stenosis, and absolute volume and percentage of each plaque component. The image reader can also calculate the remodelling index and classify the lesion type from these data. Lesion classification in ICON1 is based on previously published recommendations for tissue characterisation by radiofrequency data analysis (figure 2).

Optical coherence tomography
OCT generates an image analogous to IVUS using a low coherence, near-infrared (wavelength 1.3 µm) light source, instead of sound. A bloodless field inside the coronary artery is vital, as red blood cells strongly backscatter the near-infrared light. This is obtained by using a flush of contrast during image acquisition. OCT has a greater resolution than IVUS (20–40 vs 100–200 µm) and is thus able to delineate the thin fibrous cap present in a TCFA. However, its poorer penetration (1–2.5 mm) can limit its capacity to identify deep lipid pools and quantify plaque volume.

OCT images are obtained using a Dragonfly catheter (St Jude Medical, Minnesota, USA) connected to the Ilumien PCI Optimization System. Just before image acquisition, a short flush of iso-osmolar contrast is administered to ensure that the guide catheter is well engaged with the coronary artery and the catheter is clear of blood. The system is calibrated and OCT pull-back is initiated with a further flush of iso-osmolar contrast (10 mL in the right coronary artery and 15 mL in the left coronary artery). OCT images are obtained in 54 mm segments at a pullback rate of 20 mm/s in all three coronary arteries, where feasible. Data are transferred anonymously to a DVD for off-line analysis; the operator is blinded to these data during the procedure.

OCT data are analysed using the Medis QIVUS software. Contours are drawn around the lumen to generate data on the minimum lumen area and diameter. The whole vessel is then analysed to identify plaque subtypes. An atherosclerotic lesion is seen on OCT as a mass lesion within the arterial wall, with focal intimal thickening or loss of the normal vessel architecture. Fibrous plaque is...
homogenous and highly backscattering, calcified plaques are signal-poor areas with sharply delineated borders and lipid pools are signal-poor regions with poorly defined borders and a fast OCT signal drop-off. Using side branches and areas of calcification as landmarks, it is possible to compare the accuracy of lesion subtypes identified by VH-IVUS and OCT.

NON-INVASIVE ASSESSMENT OF CARDIOVASCULAR STATUS

Arterial stiffness

Arterial stiffness is now increasingly recognised as a surrogate end point for the assessment of CVD status. It can lead to angina in the presence of even minor coronary artery disease and to the development of diastolic dysfunction, the commonest form of heart failure in the elderly. Arterial stiffness is determined by carotid-femoral pulse-wave velocity (PWV), which is a simple, non-invasive, robust and reproducible investigation method that can be performed at the bedside. In older patients, arterial stiffness assessed by increased PWV is associated with poor cardiovascular outcomes. In the ICON1 study, carotid-femoral PWV is assessed by the Vicorder device (Skidmore Medical Limited, Bristol, UK). In addition, brachiofemoral PWV, pulse-wave analysis (includes pulse pressure, augmentation pressure and augmentation index) and ankle brachial pressure index are also assessed.

Endothelial function

Endothelial dysfunction is considered one of the earliest markers of atherosclerosis, contributing to lesion development and its later clinical manifestations. It is associated with increased risk of cardiovascular events and has been proposed as a marker of poor CV outcomes. Peripheral arterial tonometry (PAT) by finger plethysmography (EndoPAT; Itamar Medical, Caesarea, Israel) is a novel method of measuring the peripheral vasodilator response. Hyperaemic response measured by PAT signal amplitude gives a measure of nitric oxide-mediated endothelial function. In patients with low-risk findings during stress testing and/or the absence of new obstructive lesions on angiography, lower natural logarithmic-scaled reactive hyperaemia index (<0.40) is associated with increased cardiovascular death over 6 years. In the ICON1 study, endothelial function is measured by EndoPAT. PAT signals are recorded from the index fingers with pneumatic probes at baseline, during cuff occlusion and during hyperaemia. A measure of endothelial function is calculated from the ratio of PAT signal amplitude at baseline and postocclusion. Reactive hyperaemia index data from the study will be used in the prediction of adverse CV outcomes and will be incorporated in the risk model.

Carotid intima media thickness

Carotid intima media thickness (CIMT) is a significant predictor of incident adverse cardiovascular events.
Increased CIMT was associated with severity of coronary atherosclerosis in ACS. CIMT and its association with predicting CV events in older patients with NSTEMACS are not known. In a meta-analysis, addition of CIMT to Framingham risk score in general population did not improve 10-year prediction of first MI or stroke. However, CIMT and arterial stiffness together increase the cardiovascular risk in patients with known vascular disease or cardiovascular risk factors. In the ICON1 study, CIMT is assessed using a Vivid I GE machine, with a vascular probe. CIMT measurement is obtained via semiautomated software, which uses an edge detection technique. CIMT values will be analysed for the prediction of adverse outcomes and will be incorporated in the risk model.

**Transthoracic echocardiogram**

In hospitalised elderly patients with known CVD, left ventricular diastolic dysfunction was similar in prevalence to systolic dysfunction and was associated with similar cardiovascular and all-cause mortality. Transthoracic echocardiography will be performed using a Vivid I GE echo machine, according to the British Society of Echocardiography guidelines, to assess systolic function, diastolic function and valvular heart disease. Systolic and diastolic function will be analysed for the prediction of adverse CV outcomes.

**Follow-up**

One-month outcomes are recorded using general practitioner summary documents, obtained from the patients’ general practitioner. The patients are followed-up in a study outpatient clinic at 1 year. During this follow-up visit, repeat blood samples are collected for biomarker analysis. In addition, NYHA class, CCS angina class, SF-36, EQ-5D and MoCA assessments are completed. Frailty status is reassessed using Fried and Rockwood Frailty Criteria.

**Primary outcome measures**

The primary outcome measure is a composite of death, MI, stroke, repeat, unplanned revascularisation and BARC (Bleeding Academic Research Consortium)-defined bleeding (type 2 or greater) at 1 year (see online supplementary appendices 3 and 4). We also intend to analyse 1-year mortality as an independent outcome measure. All-cause hospitalisation comprises a secondary outcome measure.

**Sample size**

For the primary outcome, Hsieh and Lavori’s method was used to calculate the power for testing the association of the risk score with adverse outcomes, based on 300 participants with a type I error rate of 0.05. From the national-level registry data, the 1-year mortality rate for NSTEMI in all patients undergoing invasive strategy is ~2–5%. Estimates of the SD and HR of the risk score are unknown. An assumption was made on the HRs being an increment of 1 SD of the risk score (see figure 3).

**STATISTICAL METHODS**

**Risk factor selection**

Cox proportional hazards regression analysis will be performed to estimate HRs of the risk factors and associated p values for the primary outcome. Multiple logistic regression analysis will be performed to estimate ORs of the risk factors and associated p values for the secondary outcome. The bootstrap method will be used to avoid overfitting the data. One thousand bootstrapping will be performed. For each bootstrapping, we will sample with replacement 300 patients from the original 300 patients. Backward selection with a p value of <0.05 for statistical significance will be used to remove variables in each sample. Variables selected ≥800 times (80%) in the overall sample will be included in the final model. All missing values will be reported, and appropriate statistical methods will be used to handle missing values.

**Risk score construction**

To construct the risk score, risk factors identified through the multivariable model will be assigned a weight. Weights are the estimated regression coefficients from the Cox proportional hazards regression or logistic regression model. The risk score is thus the weighted average of the identified risk factors. Another Cox proportional hazards regression or logistic regression model will be applied to detect the association of the proposed risk score to the outcomes.

**Risk score evaluation**

Harrell’s C-index will be used to assess the discriminatory capacity of the integrated risk score for primary and secondary outcomes. The Jackknife method will be used to estimate the SE of the estimated Harrell’s C-index or area under the curve. The difference between model-predicted and observed event rates (goodness-of-fit) will be evaluated using the Hosmer-Lemeshow test (p value of >0.10 will be considered to indicate the lack of
deviation between the model-predicted and observed event rates). Reclassification calibration measures (eg, net reclassification improvement and integrated discrimination improvement) will be used to evaluate the improvement of new predictors (relative to existing predictors) on the agreement between observed and predicted outcomes. A cross-validation technique will be used to assess how the results of statistical analysis generalise to an independent data set. Finally, a prediction nomogram will be developed to facilitate calculating the risk scores and the corresponding survival probability at 1 year.

CONCLUSION

The ICON1 study will identify predictors of poor cardiovascular outcomes among older patients (aged ≥75 years) presenting with NSTEACS managed by contemporary pharmacotherapy and invasive revascularisation strategy. Based on clinical characteristics, frailty status, comorbidities and cardiovascular status, an integrated risk stratification tool to help decision-making in the management of older patients will be developed. The variables that we hypothesise may be relevant to such a model would be either (1) routinely collected in clinical practice as part of current evidence-based practice or (2) should not be unduly burdensome to collect during routine clinical assessment.

Author affiliations

1. Institute of Cellular Medicine, Newcastle University, Newcastle upon Tyne, UK
2. Cardiothoracic Centre, Freeman Hospital, Newcastle upon Tyne, UK
3. Department of Biochemistry, Newcastle upon Tyne Hospitals NHS Foundation Trust, Newcastle upon Tyne, UK
4. Institute for Ageing and Health, Newcastle University, Newcastle upon Tyne, UK
5. Channing Division of Network Medicine, Brigham and Women’s Hospital and Harvard Medical School, Boston, Massachusetts, USA

Twitter

Follow Jonathan Batty at @jonnybatty

Acknowledgements

The authors thank Dr D Ahmed, Dr A Bagnall, Dr R Das, Dr R Edwards, Dr M Eged, Dr I Purcell, Professor I Spyridopoulos and Professor A Zaman of Freeman Hospital, Newcastle upon Tyne, and Dr Mark de Belder and Mrs Bev Atkinson of the James Cook University Hospital, South Tynes Hospitals NHS Foundation Trust, Middlesbrough, UK for their help with data collection. They also thank the Cardiology CRN research team at Freeman Hospital, Mrs Kathryn Proctor and Mrs Jennifer Adams-Hall, for their support with the follow-up of study patients. Dr Carmen Martin-Ruiz and Dr Gabriele Saretzki of Institute for Ageing and Health, Newcastle University, Newcastle upon Tyne, UK on behalf of the investigators. Invasive versus conservative strategy in patients aged 80 years or older with non-ST-segment-elevation acute myocardial infarction: TRITIUM (TRatamiento del Infarto Agudo de miocardio en Ancianos) randomized trial and pooled analysis with previous studies. Eur Heart J 2011;32:51–60.

REFERENCES

1. Murray CJ, Lopez AD. Mortality by cause for eight regions of the world: Global Burden of Disease Study. Lancet 1997;349:1269–76.
2. Finegold JA, Asaria P, Francis DP. Mortality from ischaemic heart disease by country, region, and age: statistics from World Health Organisation and United Nations. Int J Cardiol 2013;168:934–45.
3. Townsend N, Wickramasinghe K, Bhatnagar P, et al. Coronary heart disease statistics 2012 edition. London: British Heart Foundation, 2012.
4. Gavala L, Weston C. Myocardial Ischaemia National Audit Project Annual Public Report April 2012–March 2013. London: National Institute for Cardiovascular Outcomes Research, 2013.
5. Veerasamy M, Edwards R, Ford G, et al. Acute coronary syndrome among older patients: a review. Cardiol Rev 2015;23:26–32.
6. Lee PY, Alexander KP, Hammill BG, et al. Representation of elderly persons and women in published randomized trials of acute coronary syndromes. JAMA 2001;286:708–13.
7. Bueno H, Betriu A, Heras M, et al. Acute coronary care in elderly patients: a review. Heart 2001;86:176–81.
8. Savonitto S, Cavallini C, Petronio AS, et al. Early aggressive versus initially conservative treatment in elderly patients with non-ST-segment elevation acute coronary syndrome: a randomized controlled trial. JACC Cardiovasc Interv 2012;5:906–16.
9. Tegn N, Abdelnoor M, Aaberge L, et al. After Eighty study investigators. Invasive versus conservative strategy in patients aged 80 years or older with non-ST-elevation myocardial infarction or unstable angina pectoris (After Eighty study): an open-label randomised controlled trial. Lancet 2016;387:1057–65.
10. Roe MT, Goodman SG, Ohman EM, et al. Elderly patients with acute coronary syndromes managed without revascularization: insights into the safety of long-term dual antiplatelet therapy with reduced-dose prasugrel versus standard-dose clopidogrel. Circulation 2013;128:823–33.
11. Alexander KP, Newby LK, Cannon CP, et al. Acute coronary care in the elderly, part I: non-ST-segment-elevation acute coronary syndromes: a scientific statement for healthcare professionals from the American Heart Association Council on Clinical Cardiology; in collaboration with the Society of Geriatric Cardiology. Circulation 2007;115:2549–69.
12. Kandzari DE, Roe MT, Chen AY, et al. Influence of clinical trial enrollment on the quality of care and outcomes for patients with non-ST-segment elevation acute coronary syndromes. Am Heart J 2005;149:74–81.
Potential pitfalls of fitness and frailty in elderly people.

A global clinical measure of fitness and frailty in elderly people. Arch Intern Med 2001;56:1146–56.

Rockwood K, Song X, MacKnight C, et al. A global clinical measure of fitness and frailty in elderly people. CMAJ 2005;173:489–95.

Charlson ME, Pompei P, Ales KL, et al. A new method of classifying performance of acute coronary syndrome in patients presenting without persistent ST-segment elevation: Task Force for the Management of Acute Coronary Syndromes. European Society of Cardiology (ESC). Eur Heart J 2007;28:2595–611.

Valadi H, Ekstrom K, Bossios A, et al. Exosome-mediated transfer of mRNAs and microRNAs is a novel mechanism of genetic exchange between cells. Nat Cell Biol 2004;6:658–69.

Moldovan L, Batte KE, Trgovich J, et al. Methodological challenges in utilizing microRNAs as circulating biomarkers. J Cell Mol Med 2014;18:371–90.

Vickers KC, Palimaso BT, Shoucri BM, et al. MicroRNAs are transported in plasma and delivered to recipient cells by high-density lipoproteins. Nat Cell Biol 2011;13:423–33.

Wagner J, Riwanto M, Bessler C, et al. Characterization of levels and cellular transfer of circulating lipoprotein-bound microRNAs. Arterioscler Thromb Vasc Biol 2013;33:1392–400.

Turchinovich A, Burwinkel B, Distinct AGO1 and AGO2 associated with 1-year mortality for elderly patients with non-ST-segment elevation myocardial infarction. Eur J Prev Cardiol 2014;21:1216–24.

Singh M, Rihal CS, Lennon RJ, et al. Frailty in older adults: a global clinical measure of fitness and frailty in elderly people. Arch Intern Med 2001;56:1146–56.
recommendations for acquisition, analysis, interpretation and reporting. **EuroIntervention** 2009;5:177–89.

63. Mintz GS, Kent KM, Pichard AD, et al. Contribution of inadequate arterial remodeling to the development of focal coronary artery stenoses. An intravascular ultrasound study. **Circulation** 1997;95:1791–8.

64. Huang D, Swanson EA, Lin CP, et al. Optical coherence tomography. **Science** 1991;254:1178–81.

65. Sinclair H, Bourantas C, Bagnall A, et al. OCT for the identification of vulnerable plaque in acute coronary syndrome. **JACC Cardiovasc Imaging** 2015;8:198–209.

66. Prati F, Guagnioli G, Mintz GS, et al. Expert’s OCT Review Document. Expert review document part 2: methodology, terminology and clinical applications of optical coherence tomography for the assessment of interventional procedures. **Eur Heart J** 2012;33:2513–20.

67. Tearney GJ, Regar E, Akasaka T, et al. Expert consensus document on arterial stiffness: methodological issues and clinical applications. **Eur Heart J** 2006;27:2588–605.

68. Laurent S, Cockcroft J, Van Bortel L, et al. Expert consensus document on arterial stiffness: methodological issues and clinical applications. **Eur Heart J** 2006;27:2588–605.

69. Weber T, Auer J, O. el al. Prolonged mechanical systole and increased arterial wave reflections in diastolic dysfunction. **Heart** 2006;92:1616–22.

70. Veerasamy M, Ford GA, Neely D, et al. Association of aging, arterial stiffness and coronary artery disease: a review. **Cardiol Rev** 2014;22:22–32.

71. Lüscher TF, Barton M. Biology of the endothelium. **Circ Cardiol** 1997;20(Suppl 2):II-3–20.

72. Laurent S, Crockcroft J, Van Bortel L, et al. Expert consensus document on arterial stiffness: methodological issues and clinical applications. **Eur Heart J** 2012;59:1058–72.

73. Ross R. The pathogenesis of atherosclerosis: a perspective for the 1990s. **Nature** 1993;362:801–9.

74. Halcox JP, Schenke WH, Zalos G, et al. Prognostic value of coronary vascular endothelial function in the assessment of interventional procedures. **Eur Heart J** 2012;33:1058–72.

75. Schächinger V, Britten MB, Zeiher AM. Prognostic impact of coronary vasodilator dysfunction on adverse long-term outcome of coronary heart disease. **Circulation** 2000;101:1899–906.

76. Suwaidi JA, Hamasaki S, Higano ST, et al. Long-term follow-up of patients with mild coronary artery disease and endothelial dysfunction. **Circulation** 2000;101:948–54.

77. Kuvín JT, Patel AR, Slény KA, et al. Assessment of peripheral vascular endothelial function with finger arterial pulse wave amplitude. **Am J Cardiol** 2003;14:168–74.

78. Pavie L, Schnall RP, Sheffy J, et al. Peripheral vasoconstriction during REM sleep detected by a new plethysmographic method. **Nat Med** 2000;6:606.

79. Nohria A, Gerhard-Herman M, Creager MA, et al. Role of nitric oxide in the regulation of digital pulse volume amplitude in humans. **J Appl Physiol** 2006;101:545–8.

80. Noon JP, Haynes WG, Webb DJ, et al. Local inhibition of nitric oxide generation in man reduces blood flow in finger pulp but not in hand dorsum skin. **J Physiol** 1996;490(Pt 2):501–8.

81. Rubinstein R, Kuvín JT, Soffler M, et al. Assessment of endothelial function by non-invasive peripheral arterial tonometry predicts late cardiovascular adverse events. **Eur Heart J** 2010;31:1142–8.

82. Lorenz MW, Markus HS, Bots ML, et al. Prediction of clinical cardiovascular events with carotid intima-media thickness: a systematic review and meta-analysis. **Circulation** 2007;115:659–67.

83. Carpenter M, Sinclair H, Kunadian V. Carotid intimal media thickness and its utility as a predictor of cardiovascular disease: a review of evidence. **Cardiol Rev** 2016;24:70–5.

84. Karayiorgou M, Aricò I, et al. The assessment of atherosclerosis on transthoracic echocardiogram: a guideline protocol from the British Society of Echocardiography. **Echo Res Pract** 2015;2:99–24.

85. Thaygesen K, Alpert JS, White HD, et al. Universal definition of myocardial infarction. **Eur Heart J** 2007;28:2525–38.

86. Mehran R, Yao SV, Bhat DL, et al. Standardized bleeding definitions for cardiovascular clinical trials: a consensus report from the Bleeding Academic Research Consortium. **Circulation** 2011;123:2736–47.

87. Hsieh FY, Lavori PW. Sample-size calculations for the Cox proportional hazards regression model with nonbinary covariates. **Control Clin Trials** 2002;23:552–60.

88. British Cardiovascular Intervention Society. **BCIS audit result 2011**. London (UK): British Cardiovascular Intervention Society, 2011.

89. Harrell FE Jr, Lee KL, Pollock BG. Regression models in clinical studies: determining relationships between predictors and response. **J Natl Cancer Inst** 1988;80:1198–202.

90. Cook NR, Ridker PM. Advances in measuring the effect of individual predictors of cardiovascular risk: the role of reclassification measures. **Ann Intern Med** 2009;150:795–802.

91. Picard RD, Cook RD. Cross-validation of regression-models. **J Am Stat Assoc** 1984;79:575–83.

92. Iasonos A, Schrag D, Raj GV, et al. How to build and interpret a nomogram for cancer prognosis. **J Clin Oncol** 2008;26:1364–70.

93. World Medical Association. World Medical Association Declaration of Helsinki: ethical principles for medical research involving human subjects. **JAMA** 2013;310:2191–4.