Study protocol for a matter of heart: a qualitative study of patient factors driving overuse of cardiac catheterisation

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ABSTRACT

Introduction
Overuse of cardiac catheterisation (CC) for stable coronary artery disease (CAD) is documented in Germany and other regions, although percutaneous coronary interventions do not provide a benefit over medical therapy for stable patients. Various studies investigated health system, physician and patient factors driving non-adherence to guidelines which recommend a stepwise approach with invasive procedures only in case of signs of ischaemia in non-invasive testing. In a large-scale project, we aim to better understand the patients’ perspective in order to develop an intervention that enhances patient’s acceptance of this stepwise diagnostic approach for stable CAD. As a first step, this qualitative study aims to identify patient factors that prevent and promote the described overuse.

Methods and analysis
The explorative qualitative interview study will include about 20 patients with stable CAD and a history of acute coronary syndrome from two German teaching practices. Narrative, structured interviews designed to last 30 to 90 min will be conducted. The interviews will be analysed using qualitative content analysis by Mayring. The analysis will address the following questions: (1) What are reasons for stable patients to undergo CC? (2) How do patients deal with their heart disease (secondary prevention)? (3) Which processes do patients describe regarding decision-making for non-invasive and invasive coronary procedures? (4) What information needs exist on behalf of patients to better understand the stepwise diagnostic approach outlined in guidelines and thereby avoid low-appropriate CCs? Based on these data, empirical typification will be conducted.

Ethics and dissemination
Ethical approval for the study was obtained. All participants will provide written informed consent. Data will be pseudonymised for analysis. The findings will contribute to the development of an appropriate intervention. Results will be disseminated by conference presentations and journal publications.

INTRODUCTION
Overuse of cardiac catheterisation (CC) with and without interventions for asymptomatic patients with stable coronary artery disease (CAD) is documented for Germany, the USA and Canada.1 2 It is estimated that 4%–18% of coronary angiography procedures in the USA and Canada are inappropriate according to guidelines.3 Although conflicting data of the scope of overuse in Germany are reported, country comparisons show markedly higher rates compared with nations such as Sweden and the Netherlands, without any difference in cardiovascular mortality.4 5

Evidence shows that percutaneous coronary interventions (PCI) do not provide a benefit in quality of life compared with medical therapy in stable CAD.5 6 Therefore, guidelines recommend a stepwise approach with invasive testing for patients with signs of ischaemia in non-invasive testing.2 7–10 A 2015 Swiss study from Chmiel et al11 revealed that 37.5% (n=1018) of 2714 stable patients had not received non-invasive diagnostics prior to coronary angiography. In a retrospective study of 147 individual patient careers spanning back up to 28 years, we showed that patients with more than 10 CCs in a lifetime had a higher rate of procedures classified as ‘low appropriateness’ (72%) compared with those with fewer procedures (20% in 1–5 procedures and 52% in 6–9 procedures).12

Analysing factors that drive such overuse of CC studies describe three aspects, that is, healthcare system, physician and patient...
factors. Health system factors comprise, for example, the availability of and access to CC laboratories as well as economic incentives within health systems.\textsuperscript{13,14} A 1994 study in three UK cardiac centres showed that the use of CC varied significantly, with a higher rate of CC in the two centres with in-house cardiothoracic surgery. The authors assumed that the physicians’ philosophy as well as the availability of surgery play a role.\textsuperscript{15} A national cohort study of 158 891 elderly patients followed for up to 7 years after a hospital stay for acute myocardial infarction showed marked differences by region of residency regarding the medical management and the intensity of invasive procedures: patients in regions with more CC laboratory capacities were more likely to receive interventional procedures, regardless of their age, clinical indication or risk profile.\textsuperscript{16} Similarly, a 2003 Canadian study examining payment claims for physician services in 47 036 inpatients with acute myocardial infarction found that hospitals with on-site catheterisation laboratories, those with university affiliations and those closer to tertiary institutions showed a higher 90-day angiography use.\textsuperscript{17,19}

More recently, studies focused on physician factors driving overuse. A 2007 focus group study asked 20 US cardiologists about their intentions to use PCI for stable CAD. The authors showed that physicians had a firm belief in the benefits of the new technologies, and they feared bad outcomes and/or being sued if they did not use PCI. Some even reported negative experiences like a young patient dying of CAD. Also, physicians argued that it is better to visualise the coronary arteries than to rely on non-invasive results (so-called ‘oculostenotic reflex’). Also, they aimed to reduce patients’ anxieties and wish to match patients’ requests.\textsuperscript{18} In a qualitative study, 40 interactions between US cardiologists and patients with stable angina from the Verilogue Point-of-Practice Database were analysed in detail. In general, cardiologists overstated the benefits of coronary interventions and understated the associated risks and alternatives.\textsuperscript{19,20}

Other studies addressed patient factors that drive an overuse of CC. A questionnaire study from Rothberg \textit{et al}\textsuperscript{20} showed a marked discrepancy between US cardiologists’ and patients’ beliefs about the effects of PCI. While physicians aimed at symptom relief, three quarters of the 153 patients with elective CC believed in a prognostic procedure, 79\% (n=45) were convinced of suffering from heart disease; interestingly, 16 months after CC, 44\% (n=25) were still convinced of an undiagnosed heart disease. In agreement with others, this study concluded that uncertainties and fears play a major role, and that these should be addressed by adequate patient information. However, interventions with proven effectiveness to address the various factors on behalf of patients with stable CC, for example, lack of information, inadequate risk perception and emotional factors, are missing.\textsuperscript{20} Also, it is unclear whether there are different types of patients which may require different information strategies.

To better prevent overuse of CC, we assume that it is important to obtain a more in-depth understanding which factors influence stable patients’ decisions for invasive procedures. Therefore, this qualitative study uses a transdisciplinary approach well established in sociology and social sciences. Based on the reasons of stable patients to opt for CC, we will conduct an empirical typification to describe the relationship between a patient’s attitude towards medical information and their trust in suggested treatments and decisions for CC with low appropriateness. We assume that patients have different attitudes towards the healthcare environment which result in different decisions and actions even in comparable situations. To show that, we will draw on and integrate findings of a 2007 German study which addressed patients’ normative values and behavioural patterns in the healthcare system. Based on about 1500 randomly selected participants from the general population, four types of patients were characterised: (1) the indifferent patient, who has little interest in gathering medical information and little trust in medical treatment, (2) the accepting patient, who has little interest in gathering medical information and high trust in medical treatment, (3) the sceptical patient, who has high interest in gathering medical information and little trust in medical treatment and (4) the cophysician, who has high interest in gathering information and high trust in medical treatment.\textsuperscript{25} We expect that these four different types of patients can also be found among patients with stable CAD, and that these deal differently with their heart disease, describe different processes in decision-making, have different reasons for undergoing a CC and need different information strategies to avoid CCs of low appropriateness.

\textbf{METHODS AND ANALYSIS}

\textbf{Study setting and design}

The interviews will be conducted in two German teaching practices which are affiliated with our Institute for General Medicine. Teaching physicians will select patients with a history of acute coronary syndrome (ACS) who underwent at least one or better yet multiple coronary catheterisation procedures with or without intervention.
The interviews will be conducted by a female researcher who is qualified and experienced in qualitative research methods and interview techniques. Each interview will last about 30 to 90 min.

**Sample size**
About 20–25 patients (ideally a 70/30 male to female ratio) will be interviewed.

**Inclusion criteria**
Patients with history of ACS or known CAD who have undergone at least one or better yet multiple coronary procedures will be asked to participate. Patients must be able to communicate in German in order to understand the study information sheet, to provide informed consent and to answer questions of the interviewer.

**Exclusion criteria**
Patients with known anxiety disorders and those who do not match the inclusion criteria will be excluded.

**Recruitment and sampling**
The teaching physicians will ask patients who meet the study criteria as they come to the practice for routine care. Patients will be informed that participation is voluntary, that they can contribute to a better scientific understanding in the field and that non-participation has no adverse effect on their medical care. According to the patients’ wishes, the practice will arrange an appointment for the interview, or the contact data will be passed on to the interviewer who will then contact the patient by phone. The recruitment will end after 20–25 interviews are conducted and saturation is reached.

**Interview guide**
We will use narrative, structured interviews as a survey method which is known for high methodological standards such as comparability, objectivity and transparency. The interview guide (see table 1) was prepared by a multidisciplinary research team which includes a sociologist (AH) (qualified in qualitative research methods and interview techniques) and a Board-certified primary care physician and epidemiologist (BW). The interview guide was reviewed by an external Board-certified primary care physician specialised in psychotherapy and supervision. To familiarise herself with the field, the sociologist performed a participatory observation in a CC laboratory and interviewed cardiac patients in a general practice. The interview guide was based on these observations and conversations with patients as well as theoretical preparatory work from the field of qualitative research, our prior retrospective study and a systematic literature review.

**Compensation**
The participants will receive compensation in the form of a small gift worth 10€.

**Data analysis**
The interviews will be tape-recorded and transcribed according to a simplified transcription system by the research team prior to analysis. The transcripts will be anonymised so that no names or other identifying features will appear in any form of data reporting. Data will be analysed by means of qualitative content analysis according to Mayring using the summarising approach with a combination of inductive category development and deductive category application. Two researchers will work independently on a system of categories with a subsequent discussion of the systems until a consensus is reached. The interpretation will focus on emotional and factual arguments that motivate patients with stable CAD to undergo CC. In addition, the patients’ needs and wishes will be analysed with regard to strategies for a future intervention. The following questions will guide our analysis:

1. What are reasons for stable patients to undergo CC?
2. How do patients deal with their heart disease (secondary prevention)?
3. Which processes do patients describe regarding decision-making for non-invasive and invasive coronary procedures?
4. What information needs exist on behalf of patients to better understand a stepwise diagnostic approach and thereby avoid low-appropriate CCs?

In order to compare different types of patients and how they deal with their heart disease, an empirical typification of patients will be conducted. Objectivity and reliability will be ensured by a second coder. Validity will be verified by construct validity, the verification of the results on the basis of proven theories and/or results from prior studies on plausibility. Qualitative data analysis will be performed using the software ATLAS.ti.

**Possible outcome of the analysis and benefits of the study**
We assume that patients have a high level of uncertainty and fears regarding an adverse outcome. From everyday experience in general practices, we also presume that there is a lack of confidence in non-invasive diagnostic procedures as well as a lack of detailed information about the disease and the effectiveness of secondary prevention (lifestyle changes, medication) as well as stepwise diagnostic algorithms. Our study is designed to provide the basis for interventions supporting the health literacy of patients with known CAD on the role of diagnostic algorithms. To meet this objective, we hypothesise that the confidence in guidelines will need to be strengthened and the understanding of pathological processes improved as well as uncertainties and fears addressed in order to minimise them. On the basis of the study results, we are planning to develop an intervention in the form of an educational video. This intervention will empower patients to improve shared decision-making together with their physicians. Overall, we aim to promote guideline adherence, reduce the number of inappropriate CCs.
| Topic                      | Question                                                                 | Background                                                                                                                                                                                                 |
|----------------------------|--------------------------------------------------------------------------|----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| Starting the conversation  | How is your heart? <br> *Alternatively*: What happened to your heart? <br> *Alternatively*: What have you gone through with your heart? | Patients are encouraged to speak. At the same time, as much information as possible is to be collected from the patient's point of view about the heart disease. The answer to the question is meant to show how patients deal with their illness and how they handle and process the steps of their treatment. |
| Understanding the disease  | You have undergone cardiac catheterisation. I have already seen the procedure in a heart catheter laboratory, but I cannot imagine how it feels lying there. Can you please tell me how it feels? <br> *Further question*: What were your ideas/expectations regarding cardiac catheterisation? <br> *Further question*: What did you think about the catheterisation after the procedure? <br> *Further question*: What complications occurred during or after the procedure? <br> *Further question*: When did you feel most uncomfortable during the procedure? <br> Please tell me why the cardiac catheterisation had to be carried out? <br> *Further question*: What is your primary disease? <br> *Further question*: What complaints do you have? <br> *Further question*: When did you go to the physician regarding these complaints? <br> *Further question if patient had heart attack*: What did you feel when you had the heart attack? <br> *Further question if patient had heart attack*: What do you feel today when you think about that heart attack? <br> *Further question*: How did you feel when the physician told you about your disease? <br> *Further question*: How did you feel after you heard the results of your cardiac catheterisation? <br> Please tell me, how healthy do you feel today? <br> *Further question*: Why do you feel unwell? <br> *Further question*: How big is your fear of having another heart attack? | Patients are invited to share their experiences during the cardiac catheterisation. The aim is to find out how the patients were informed about the procedure before and what impressions they were left with. If it has not been explained in the narration before, the patients are asked to relate in their own words how they understood why the cardiac catheterisation had to be done, what disease they are suffering from and what affects them on an emotional and factual level. The patients should talk about their state of health after the procedure and whether they are afraid of a new heart attack. The answer to this question is to find out whether the cardiac event had an influence on everyday life. |
| Topic                                      | Question                                                                 | Background                                                                                                                                   |
|-------------------------------------------|--------------------------------------------------------------------------|-----------------------------------------------------------------------------------------------------------------------------------------------|
| Information procurement                   | What did you discuss with your family regarding your cardiac catheterisation?                                             | We want to know how family and friends support the patient.                                                                                   |
|                                            | **Further question:** How did your family react to your cardiac catheterisation?                                             |                                                                                                                                             |
|                                            | **Further question:** How did your friends react to your cardiac catheterisation?                                           |                                                                                                                                             |
|                                            | **Further question:** What did you think about **"THAT SPECIAL ADVICE/OPINION"**?                                           | We want to know if patients actively get informed about their disease and the treatment options and how the patients get this information. |
|                                            | You seem to be quite well informed about your disease and the cardiac catheterisation. How come?                           |                                                                                                                                             |
|                                            | **Further question:** What do you read about your disease?                                                                 |                                                                                                                                             |
|                                            | **Further question:** What advice does your family give you?                                                               |                                                                                                                                             |
|                                            | **Further question:** Do you talk about your disease with your family and friends a lot?                                   |                                                                                                                                             |
|                                            | **Further question:** With regard to all the information you were given, what would you have said to the physician if he had told you that you do not need cardiac catheterisation? |                                                                                                                                             |
| Enlightenment by the physicians and patient-physician relationship | Which physician did you initially contact because of your complaints?                                                        | We want to find out which physician the patients are most likely to trust and which recommendations they follow.                               |
|                                            | Which medical examinations were done before you had your cardiac catheterisation?                                         | The intention is to analyse how familiar the patient is with other diagnostic procedures.                                                   |
|                                            | How did the physician explain the need and the procedure of cardiac catheterisation to you?                               | The aim of this question is to establish the satisfaction with the ambulatory patient’s education.                                              |
|                                            | **Further question:** How did the physician explain your disease?                                                           |                                                                                                                                             |
|                                            | **Further question:** What did you feel when your physician told you that you needed cardiac catheterisation?            | The aim is to ask about the satisfaction with the patient’s education as an inpatient as well as about the patient’s needs and wants regarding information. |
|                                            | What did the physicians tell you during the procedure?                                                                    |                                                                                                                                             |
|                                            | What do you think about the information the physicians gave to you?                                                        |                                                                                                                                             |
|                                            | Do you remember a situation in which you would have liked to have more information?                                      |                                                                                                                                             |
|                                            | How satisfied are you with your treatment in general?                                                                    |                                                                                                                                             |
| Ending the conversation                    | Is there anything else you did not tell my yet but think may be of interest?                                              |                                                                                                                                             |
procedures and thereby decrease the risk of minor and major adverse effects including severe arrhythmias and death as well as radiation exposure.

**Ethics and dissemination**

All participants will receive an information sheet outlining the study, their voluntary participation and how their personal data will be protected. All participants will provide written informed consent. Procedures for pseudonymisation of the transcripts of the tape-recorded interviews will be outlined. Participants have the right to refuse answers to any question posed by the interviewer without disadvantages for the participants. The tapes will be erased on completion of the study. All electronic data will be stored in password-protected computers. Only the research team will have access to the data. Participants will not be identifiable in any publication or dissemination activity; confidentiality will be ensured by using study numbers to differentiate participant quotations. The findings will contribute to the development of an appropriate intervention. Results will be disseminated within the academic field (conference presentations, journal publications) and beyond. Our study report will adhere to the consolidated criteria for reporting qualitative studies.

Ethical approval for the study was obtained from the Ethics Committee of the Faculty of Medicine at the University of Duisburg-Essen, Germany (15-6448-BO).

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**Contributors** AH: complemented the study concept, wrote the first draft of the manuscript. BW: developed the study idea and concept, revised the manuscript. Both authors critically reviewed the first draft and provided feedback on it. Both authors read and approved the final manuscript.

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