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Gender differences in patients with acute chest discomfort: a cross-sectional study

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Loes TCM Wouters¹, Dorien LM Zwart¹, Daphne CA Erkelens¹, Esther De Groot¹, Arno W Hoes², Roger AMJ Damoiseaux¹, Frans H Rutten¹

- Dept. General Practice, Julius Centre for Health Sciences and Primary Care, University Medical Centre, Utrecht University, Utrecht, the Netherlands
- 2. Dean University Medical Center, Utrecht University, Utrecht, the Netherlands

All authors take responsibility for all aspects of the reliability and freedom from bias of the data presented and their discussed interpretation

Corresponding author: L.T. Wouters, MD, Dept. General Practice, Julius Centre for Health Sciences and

Primary Care, University Medical Centre Utrecht, Utrecht University

STR 6.131, PO Box 85500, 3508 GA Utrecht, The Netherlands

Phone number: +31 (0)88 75 51470

Email: L.T.C.Wouters-2@umcutrecht.nl

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All authors have completed the <u>Unified Competing Interest form</u> and declare no support from any organization for the submitted work; no financial relationships with any organizations that might have an interest in the submitted work in the previous three years, no other relationships or activities that could appear to have influenced the submitted work.

Transparency declaration

The lead author (the manuscript's guarantor) affirms that the manuscript is an honest, accurate, and transparent account of the study being reported; that no important aspects of the study have been omitted; and that any discrepancies from the study as planned (and, if relevant, registered) have been explained.

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Objectives To assess differences in symptoms predictive of acute coronary syndrome (ACS) between women and men presenting with chest discomfort in out-of-hours primary care (OHS-PC).

Design Cross-sectional study.

Setting Nine OHS-PC in the Netherlands.

Participants 993 women and 802 men who called OHS-PC for acute chest discomfort (pain, pressure, tightness, or discomfort) between 2014 and 2017.

Primary outcome measure Diagnosis of acute coronary syndrome (ACS). We compared patient and call characteristics of triage call recordings between women with and without ACS, and men with and without ACS. Diagnoses were retrieved from the patient's medical record in general practice, including hospital specialists' discharge letters.

Results Among 1,795 patients (mean age 58.8 (SD 19.5) years, 55.3% women), 15.0% of men and 8.6% of women had an ACS. In both sexes, retrosternal chest pain was discriminative for ACS (women with ACS vs. without 62.3% vs. 40.3%, p=0.002, men with ACS vs. without 52.5% vs. 39.7%, p=0.032), as was pressing/tightening pain (women 78.6% vs. 61.5%, p=0.011, men 82.1% vs. 57.4%, p=<0.001) and radiation to arm (women 75.6% vs. 45.9%, p<0.001, men 56.0% vs. 34.8%, p<0.001). In women, severe pain (65.4% vs. 38.1%, p=0.006), a pale face (50.0% vs. 22.9%, p=0.007) and radiation to jaw (50.0% vs. 29.8%, p=0.017) were also discriminative for ACS, while in men this was sweating (52.4% vs. 38.1%, p=0.015), and against ACS stabbing pain (8.4% vs. 26.5%, p<0.001).

Ambulances were dispatched equally in women (72.9%) and men with ACS (70.0%).

Conclusion Symptoms predictive of ACS were rather similar for women and men with chest discomfort, with some important exceptions, such as severity, type, and radiation of pain, and some autonomous nervous system related symptoms.

Trial number: NTR7331

Keywords (5): acute coronary syndrome, gender, primary care, telephone triage, chest discomfort

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- We could analyse the very initial conversation with symptom presentation of a large sample of patients calling for acute chest discomfort.
- We analysed the conversations without knowledge of the eventual diagnosis and therefore no hindsight bias of the researcher or recall bias of the patient could occur.
- Results are generalizable to comparable primary care settings in European countries, and our results may even be generalizable to emergency medical service (EMS, '112' or '911') settings.
- We had missing values on some clinical variables, a phenomenon common in routine care data.



Adequate triage and early diagnosis is key in patients with acute chest discomfort because they might have an acute coronary syndrome (ACS) for which lifesaving early interventions are available. ACS is an umbrella term including ST-elevated myocardial infarction (STEMI), non-ST-elevated myocardial infarction (NSTEMI) and unstable angina pectoris (UAP). ¹ Increased preventive measures and development of (timely) effective therapeutic interventions ('time is muscle') have resulted in improved outcomes and prognosis in ACS. ² Telephone triage of patients with chest discomfort, as done in out-of-hours service primary care (OHS-PC) and emergency medical services (EMS or ambulance dispatch centres) is, however, challenging because it is difficult to differentiate ACS from other causes of chest discomfort based on symptoms only. ^{3,4} Importantly, the majority of patients with chest discomfort (80%) in the Netherlands first approach the general practitioner (GP) or OHS-PC, and 20% directly calls the ambulance (112) or are self-referrals to the emergency department (ED). ⁵

Previous hospital-based studies reported a delayed recognition of ACS in women compared to men. ^{6,7} It was suggested that this delayed recognition was related to a less specific presentation in women. ^{8,9} This caused an ongoing debate on whether women with ACS compared to men present with less specific symptoms, and how this affects diagnosis, but also treatment, and prognosis. ^{10,11} A recent meta-analysis of 27 studies showed that women with ACS compared to men with ACS had higher odds of presenting with pain between the shoulder blades (OR 2.15, 95% CI 1.95-2.37), nausea or vomiting (OR 1.64, 95% CI 1.48-1.82) and shortness of breath (OR 1.34 (95% CI 1.21-1.48). ¹² Women with ACS had lower odds of sweating (OR 0.84, 95% CI 0.76-0.94) and presenting with chest pain (OR 0.70, 95% CI 0.63-0.78), but in both sexes chest pain remained the most common symptom (pooled prevalence men 79% and women 74%). ¹² Importantly, researchers suggested standardization in methods of symptoms assessment is needed, because of the difficulties to formulate any definitive statements about symptom presentation, as studies assessed symptoms in different ways (questionnaires or abstracting from medical records). ^{9, 13-15} Abstracting symptom presentation from medical records may dilute symptom presentation, as they are translated by the clinician in medical terminology. ¹⁶ Moreover, many studies suffer from recall or hindsight bias of both patient and researcher as they know the outcome (ACS).

For the clinician or telephone triage nurse it is crucial to differentiate ACS from other causes of chest discomfort. For that, studies are needed that include female and male patients presenting with chest discomfort, in which women and men who turn out to have ACS are compared to those who do not

have ACS. Such studies are scarce. In a study performed among patients with chest discomfort seen at the ED in the USA (77 women and 244 men with ACS, and 195 women and 240 men without ACS were compared. Women with ACS more often reported arm pain than women without ACS (47% vs. 32%, p=0.021), while men with ACS reported pressing feeling (63% vs. 54%, p=0.035) and chest pain (72% vs 60%, p=0.005) more often than men without ACS. ¹³ In a recent Dutch OHS-PC study among 23 women and 34 men with ACS, and 253 women and 208 men without ACS, differences in symptom presentation between women and men were small. ¹⁷ In men, radiation of pain (89.3% vs. 64.9%, p-value=0.011) was discriminative for ACS, while stabbing chest pain (3.7% vs. 24.0%, p-value=0.014) was discriminative for the absence of ACS. Both these symptoms were not discriminative in women (90.0% vs 78.6%, p=0.227, and 15.8% vs. 18.8%, p=0.743), respectively. ¹⁷

We aimed to assess symptoms predictive of ACS in women and men separately, among patients presenting with acute chest discomfort to OHS-PC based on analyses of recorded telephone triage conversations.

Methods

We performed a cross-sectional study with a random sample of 1,795 OHS-PC calls for chest discomfort (chest pain, pressure, tightness, or discomfort) between 2014 and 2016. ¹⁸ We first selected calls on the basis of International Code for Primary Care (ICPC; a WHO world-wide code system for primary care) codes (K01, K02, K03, K24, K74, K75, K76, K77, K93, L04, P74, R02, R98) and keywords (thoracic pain, chest pain, myocardial infarction, heart attack and their common abbreviations). ^{19, 20} On purpose, we sampled a broad variety of symptoms to capture the entire domain of patients that could be suspected of ACS. We drew a random sample of all available calls of these patients with the Random Number Generator (RAND) function in Microsoft Excel. Calls were excluded before re-listening when the patients' age was below 18 years or when the patient did not live in the surrounding area of the OHS-PCs (because then we could not retrieve a diagnosis from the general practitioner of these patients). Calls were excluded during re-listening when it not concerned a triage call (e.g. inter-collegial consultation) or when the recording was of poor quality.

We re-listened the telephone triage recordings and collected information about patient and conversation characteristics, on symptom presentation, medical history, urgency allocation, and involvement of a supervising general practitioner (GP) in the triage. Nine OHS-PC in the Netherlands

Context

In the Netherlands, OHS-PC covers primary care during 73% of the week hours, and the initial contact is by telephone. In most OHS-PC and EMS, the 'Netherlands Triage Standard' (NTS) is used as a decision support to classify the urgency of the patients' conditions. ²¹ Based on the patient's symptom presentation, the triage nurse needs to choose the most appropriate complaint out of 56 'main complaints'. Each NTS 'main complaint' incorporates a decision tree with hierarchically ordered questions, which are similar for men and women. Triage nurses fill out the caller's responses in the semi-automatic NTS system, which then generates urgency allocations (U0 (reanimation) to U5 (self-care advice, see also appendix-table1). The triage nurse can overrule this recommendation and up- or downscale the urgency allocation, often after consulting the supervising GP. ²² Since its introduction in 2011, the NTS system has, however, never been formally validated by correlating the generated urgencies to clinical endpoints. ²¹

Data analyses

We compared patient and call characteristics between women with and without ACS, and men with and without ACS. We analysed the association between urgency allocation and the final diagnosis ACS (with or without other life threatening events (LTEs)). We considered pulmonary embolism, thoracic aortic dissection and acute abdominal aneurysm as LTEs; patients with LTEs as well as those with ACS should receive an U1-level urgency (full list in table 2).

For comparison of dichotomous variables we used the Chi² test or Fisher exact test and for continuous variables the independent sample t-test or Mann-Whitney U test. Data analysis was performed using SPSS, IBM version 25.

Patient and public involvement

No patients were involved in setting the research question or the outcome measures, or in developing plans for design, however, they were involved in the implementation of the study. In addition, they

were asked to advise on interpretation and writing up of results. Results will be shared and discussed with the national patient community of cardiovascular diseases ('Harteraad').

Ethics

The study (National Trial Register identification number: NTR7331) was approved by the Medical Ethics Committee Utrecht, the Netherlands (reference number WAG/mb/16/003208) and complied with the Declaration of Helsinki. A waiver of informed consent was given because our study had minimal risk to subjects and could otherwise not be carried out logistically. Personal and research data were handled and stored according to the European General Data Protection Regulation.

Results

Among the 1,795 callers with chest discomfort (mean age 58.8 (SD 19.5) years, 55.3% women), 8.6% of women and 15.0% of men had an ACS. In women with an ACS, 18.8% had a STEMI, 48.2% a NSTEMI, 20.0% an UAP, and 13.0% non-classified ACS. In men with ACS, 32.5% had a STEMI, 36.7% a NSTEMI, 27.5% an UAP, and 3.3% non-classified ACS.

A total of 22 (2.2%) women and 23 men (2.9%) had another LTE than ACS (e.g. pulmonary embolism, thoracic aortic dissection, acute abdominal aneurysm, acute heart failure).

Patient and call characteristics

In table 1, patient and call characteristics are presented for women and men with and without ACS. Men and women with ACS were older than those without ACS (mean age of women 73.6 vs. 57.8 years, p<0.001, men 67.2 vs. 56.9 years, p<0.001), and the mean duration of the telephone calls shorter (women 6:47 vs. 7:47 minutes, p=0.021, men 6:31 vs. 7:33 minutes, p=0.004).

The GP was consulted by the triage nurse in the majority of cases (52.2% in women and 55.5% in men, p=0.161). However, in women with ACS, the GP was less often consulted than in women without ACS (41.2% vs. 53.2%, p=0.034). In men such a difference was not observed (53.3% vs. 55.9%, p=0.607). In around half of the calls, someone else called on behalf of the patient; somewhat less often in women than in men (49.5% vs. 54.7%, p=0.029). In cases with ACS, for both sexes more often someone else called than in those without ACS (in women 69.4% vs. 47.7%, p<0.001, in men 65.8% vs. 52.8%, p=0.008). In men with ACS, most often their female partner (53.3%) called, while in women with ACS, it was either their male partner (17.6%), their daughter (20.0%) or a nurse (17.6%). See appendix-table 2.

Both women and men with ACS more often had a history of cardiovascular disease or cardiovascular risk factors (e.g. hypertension and diabetes) than those without (women 78.3% vs. 57.5%, p=0.001, and in men these proportions were 72.1% vs. 61.9% (p=0.046), respectively.

Symptom presentation

Chest pain was the most common complaint, both in those with and without an ACS; in women with and without ACS 98.8% and 93.1% (p=0.055), and in men 92.4% and 94.5%, respectively (p=0.364). Retrosternal located chest pain was more common in women and men with ACS than in those without ACS (women 62.3% vs. 40.3%, p=0.002, and men 52.5% vs. 39.7%, p=0.032). Also, radiation of pain to the arms seemed indicative of ACS in both sexes (women with vs. without ACS 75.6% vs. 45.9%, p<0.001, and men 56.0% vs. 34.8%, p<0.001), as was pressing/heavy/tightening chest pain (women with vs. without ACS 78.6% vs. 61.5%, p=0.011 and men 82.1% vs. 57.4%, p=<0.001). Only in women radiation to the jaw (50.0% vs. 22.9%, p=0.007) and severe pain (8 or more on a Numeric Rating Scale 0-10) was indicative for ACS (65.4% vs. 38.1, p=0.006). Only in men, stabbing pain was discriminative in that it was very rare in those with ACS (8.4% vs. 26.5%, p<0.001).

Of the autonomous nervous system (ANS)-related symptoms, nausea/vomiting and dizziness/near fainting were not indicative for ACS in either sex, with the exception of a pale face that was discriminative in women (50.0% vs. 29.8%, p=0.017), and sweating in men (52.4% vs. 38.1%, p=0.015). Recognition of symptoms being similar to a previous cardiac event was discriminative for ACS in men (52.9% vs. 32.1%, p=0.004), but not clearly for ACS in women (32.5% vs. 21.4%, p=0.108).

Diagnoses

The most common non-ACS diagnoses in both sexes were (i) non-urgent cardiovascular diseases such as stable angina pectoris, stable heart failure and arrhythmias (19.5% of all female callers with chest discomfort vs. 21.2% of male callers, p=0.384) and (ii) non-cardiac unspecified chest pain (women 16.4% vs. 19.8% men, p=0.061). Women more often than men were diagnosed with musculoskeletal problems (women 20.8% vs. men 14.1%, p=0.001) and psychogenic conditions (women 14.0% vs. men 8.4%, p<0.001).

Women and men were equally sent an ambulance; overall (43.6% vs. 46.6%, p=0.200). These proportions were in women compared to men with ACS or other LTEs 66.4% vs. 67.1%, p=0.897). Women and men in whom ACS was diagnosed had more often ambulances dispatched than in those showed not to have an ACS (women 72.9% vs. 40.9%, p<0.001, and men 70.0% vs. 42.5%, p<0.001). See table 3.

Discussion

For both sexes, retrosternal pain, pain described as pressing, heavy or tightening, and radiation to the arm were indicative of ACS in patients presenting with chest discomfort to out-of-hours primary care (OHS-PC). Radiation of pain to the jaw, severe pain, and a pale face was indicative for ACS in women. Sweating was positive related to ACS and stabbing pain negatively in men with chest discomfort. Women and men were equally often sent an ambulance.

Our finding that radiation of pain to the arm and chest pressure were discriminative for ACS in both sexes was not reported in the two previous studies that assessed ACS symptoms using the same methodology. The US study in ED-setting found that radiation to the arm was indicative for ACS in women but not for men, and chest pressure was distinctive for men but not for women. ¹³ The OHS-PC study reported opposite that radiation to the arm was indicative for ACS in men, but not in women, and reported similar to our study that stabbing pain is very uncommon in men with ACS. ¹⁷ Regarding dispatching the ambulance, the aforementioned OHS-PC study and two EMS studies showed, similar to our findings, that there was no difference in dispatch priorities between men and women with ACS. ^{4, 17, 23} This is in contrast with studies that show delay in hospital presentation of women with ACS. ^{9, 24}

We need to realize that focusing on differences may blur the large overlap in symptoms. Moreover, comparing *selectively* women with ACS to men with ACS as many previous studies did, is clinically irrelevant. ^{9, 14, 25} Clinicians, including GPs, and triage nurses need to know whether and how women with ACS differ from women without ACS, with the same question for men. Nevertheless, even guidelines stick to comparing those with established disease, and express the view that women with ACS more likely present with less specific symptoms than men with ACS. ^{1, 26} Unfortunately, public awareness campaigns follow this reasoning, and over-emphasize sex differences in women awareness campaigns

A likely reason behind the predominant message that women present with other ACS symptoms than men is the difference in pathophysiology of coronary artery disease. Women compared to men more often have elongated plaques, located on bifurcations in epicardial coronaries, coronary spasm, microvascular dysfunction, and spontaneous coronary dissection. ^{27, 29} These pathophysiological differences have an effect on interventional treatment and prognosis. ^{2, 29} However, these differences do not necessarily imply an effect on symptom presentation because the pain pathway is equal in women and men: i.e. triggered by myocardial ischemia. ^{30, 31} A supply and demand mismatch of the myocardial oxygen consumption triggers sensory nerve endings in the myocardium and cause ischemia symptoms, and this is irrespective of the fact whether the ischemia is caused by a plaque rupture in an epicardial artery or spasm, or any other cause. ³¹ The sex differences in pathophysiology of ACS do therefore not support the belief in differences in ACS symptoms between women and men.

Another reason behind the belief of 'vague' symptom presentation in women with ACS might be that they seem to present a larger number of symptoms than men with ACS, and this may be interpreted as 'vague' by physicians. ^{15, 32} The presentation of multiple symptoms may influence the prompt recognition of heart disease and initial actions on the part of providers. ^{32, 33} In a study from 2018, 2009 women and 976 men hospitalized for myocardial infarction, healthcare providers initially thought symptoms of women (53.4%) were less often heart-related than in men (36.7%). ³² Women and men had the same chest pain symptoms, but women reported more additional symptoms. ³² In our study, the call duration and the number of GP consultations by the triage nurse were similar among women and men, suggesting that triage nurses seem not to experience more difficulties in interpreting symptoms in women than men. This is in line with a prospective study with 2,795 patients with chest discomfort in ED-setting that showed the physicians' diagnostic uncertainty for the presence of ACS in women was not more common as compared to men.³⁴ Interestingly, in the majority of calls in our study someone else than the patient called the OHS-PC (women with ACS 69.4%, without ACS 47.7%, and men with ACS 65.8%, and without ACS 52.8%). This was also highlighted in an Australian study among 1,681 patients with an acute myocardial infarction; in

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90.5% of the women with AMI someone else called on behalf of the patient and in 87.8% of the men with AMI. 35

Strengths and limitations

We could analyse the very initial conversation with symptom presentation of a large sample of patients calling the OHS-PC because of chest discomfort. We analysed the conversation without knowledge of the eventual diagnosis (no hindsight bias of the researcher or recall bias of the patient). Another strength is that our results are generalizable to comparable primary care settings, e.g. UK and Scandinavian countries, and some other European countries. Our results may even be generalizable to EMS settings, since the prior probability of having an ACS is comparable in EMS setting as in OHS-PC settings. ⁴

We had missing values on some clinical variables, a phenomenon common in routine care data.

Conclusions

Symptoms predictive of ACS were rather similar for women and men with chest discomfort, with some important exceptions, such as severity, type, and radiation of pain, and some autonomous nervous system related symptoms.

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Data sharing statement

The data can be made available for researchers whose proposed use of the data has been approved at request of the corresponding author, with a signed data access agreement.

Author contribution

FR and DZ are the lead investigators who conceived the research idea and methodology. Funding acquisition was done by FR, DZ and RD. LW and DC conducted data acquisition. LW performed the analyses and wrote the first draft of the manuscript. She was supervised by FR and DZ, who critically revised the manuscript. DC, EdG, AH and RD contributed with and approved the final version of the manuscript.

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Table 1. Baseline characteristics of 993 (55.3%) women and 802 (44.7%) men patients contacting the OHS-PC with chest discomfort (pain, pressure, tightness, or discomfort)

	993 women			802 men		
3	ACS	No ACS	p-value	ACS	No ACS	p-value
0 Characteristics	N = 85 (8.6%)	N = 908 (91.4%)		N = 120 (15.0%)	N = 682 (85.0%)	
and the states	14 - 85 (8.0%)	14 - 308 (31.470)		14 - 120 (13.0%)	14 - 082 (83.070)	
Mean age in years (SD)	73.6 (±14.1)	57.8 (±20.1)	<0.001	67.2 (±13.0)	56.9 (±19.2)	<0.001
	75.0 (±14.1)	37.8 (±20.1)	\0.001	07.2 (±13.0)	30.9 (±19.2)	\0.001
4 Çall characteristics						
6						
Mean call duration in min (SD)	6:47 (±5:16)	7:47 (±3:48)	0.021	6:31 (±3:13)	7:33 (±3:42)	0.004
8	0.47 (25.10)	7.47 (23.40)	0.021	0.51 (±5.15)	7.55 (±5.42)	0.004
Mean patient's introduction duration in	0:14 (± 0:08)	0:19 (±0:13)	<0.001	0:16 (±0:11)	0:19 (±0:14)	0.060
20	3.11 (2 0.00)	(20.13)	10.001	(_0.11)	0.13 (_0.17)	0.500
2finin (SD)						
22						
 2₃riage nurse consulted the GP	35 (41.2)	483 (53.2)	0.034	64 (53.3)	381 (55.9)	0.607
24	(.=,	(33.2)				
25omeone else called on behalf of patient	59 (69.4)	433 (47.7)	<0.001	79 (65.8)	360 (52.8)	0.008
26				- ()		
The patient or person who called	36 (97.3)	409 (88.9)	0.109	52 (96.3)	321 (86.5)	0.041
28 expressed concerned (n=497;425)	32 (2135)			- ()		
expressed concerned (n=497;425)						
30	0.1 (0.0.0)	0.17 (00.1)		110 (00 1)	601 (01 =)	0.001
Chest pain (n=960;779)	81 (98.8)	817 (93.1)	0.055	110 (92.4)	624 (94.5)	0.364
32	1= (0= 1)	(00.1)	2000	10 (20 0)	07 (00 1)	
Severe Pain (>7 on a scale 0-10) (412;341)	17 (65.4)	147 (38.1)	0.006	13 (23.6)	87 (30.4)	0.312
34						
guration						
36 	66 (400)	720 (05.0)	0.402	00 (07.4)	F44 (04 C)	0.202
3≯ 15 min (n=827;674)	66 (100)	729 (95.8)	0.102	99 (97.1)	541 (94.6)	0.292
38	60 (05.7)	F75 (72 7)	0.040	00 (04 5)	424 (72.6)	0.052
39 12 hrs (n=861;702)	60 (85.7)	575 (72.7)	0.018	88 (81.5)	431 (72.6)	0.052
10						
լկocation (n=706;561)*						
12	22 (62 2)	262 (40.2)	0.003	42 (52 5)	101 (20.7)	0.022
18etrosternal	33 (62.3)	263 (40.3)	0.002	42 (52.5)	191 (39.7)	0.032
14 1Pight or left side thoray	14 (26.4)	260 (39.8)	0.054	26 (22 5)	211 (42 0)	0.057
Bight or left side thorax	14 (26.4)	200 (33.8)	0.054	26 (32.5)	211 (43.9)	0.057
16 1 7 ype of pain (n=744;590)**						
₽sing/heavy/tightening	11 (79 6)	422 (61 E)	0.011	70 (02 1)	294 (57.4)	<0.001
60	44 (78.6)	423 (61.5)	0.011	78 (82.1)	284 (57.4)	<0.001
5\$tabbing	0 (14.2)	155 (22.5)	0.153	0 (0 4)	121 (26 5)	<0.001
52	8 (14.3)	155 (22.5)	0.152	8 (8.4)	131 (26.5)	<0.001
Radiation of chest pain (n=778;613) ***						
44						
5						

2						
Any location	61 (85.9)	485 (68.6)	0.002	65 (63.7)	292 (57.1)	0.218
5 Arm 6	31 (75.6)	188 (45.9)	<0.001	47 (56.0)	117 (34.8)	<0.001
7 Between the shoulder blades	11 (52.4)	159 (41.7)	0.336	14 (27.5)	91 (29.4)	0.781
9 Jaws 10	10 (50.0)	66 (22.9)	0.007	1 (2.6)	28 (11.6)	0.146
11 Shortness of breath (n=751;614) 12	47 (72.3)	455 (66.3)	0.328	53 (60.9)	329 (62.4)	0.788
13 13 13 14 13 13 13 13 13 13 13 13 13 13 13 13 13	13 (32.5)	79 (21.4)	0.108	27 (52.9)	92 (32.1)	0.004
1 gvents (n=410;338)						
16						
1 ANS-related symptoms						
18						
1 9 weating (n=607;523)	28 (49.1)	234 (42.5)	0.340	43 (52.4)	168 (38.1)	0.015
20						
2 Nausea or vomiting (n=463;345) 22	16 (44.4)	240 (56.2)	0.173	27 (45.8)	122 (42.7)	0.661
2Pallor (n=339;295)	17 (50.0)	91 (29.8)	0.017	20 (40.8)	82 (33.)	0.315
24						
2 5 izziness or near fainting (879;720) 26	13 (18.6)	197 (24.4)	0.277	17 (15.9)	127 (20.2)	0.249
² Medical history						
28						
$\frac{28}{8}$ V disease or risk factors (n=777;684)	54 (78.3)	407 (57.5)	0.001	75 (72.1)	359 (61.9)	0.046
31 Coronary artery disease (n=494;472) 32	17 (42.5)	114 (25.1)	0.017	45 (57.0)	151 (38.4)	0.002
33 Hypertension (n=435;321) 34	22 (71.0)	142 (35.1)	<0.001	17 (47.2)	94 (33.0)	0.091
35 Diabetes mellitus (n=412;333) 36	12 (41.4)	56 (14.6)	<0.001	12 (28.6)	59 (20.3)	0.220
Hypercholesterolemia or use of statins 38			0.161	19 (45.2)	62 (23.3)	0.003
39 40 (n=371;308)						
41						

P-value comparing retrosternal or left/right side thorax vs. others locations of pain together (restrosternal, left/right side thorax,

56 57 58

59

⁴gack/shoulder, epigastric region)

^{45*} Pressing/ heavy/tightening pain vs. other types of pain (stabbing, burning, cramping, tearing). Stabbing pain: stabbing vs. other types of 46pain (pressing/heavy/tightening, burning, cramping, tearing) 47

 $^{^{44}}$ ** P-value comparing radiation arm or back/shoulder or jaws vs. no radiation 49

⁵⁰ NRS: Numeric Rating Scale 51

⁵² ANS-related symptoms: Autonomous nervous system related symptoms 53

ÇV disease or risk factors; a history of previous coronary artery disease, heart failure, stroke, cardiac arrhythmia, hypertension, and/or

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	Women n= 993 (55.3%)	Men n=802 (44.7%)	p-value
Acute coronary syndrome	85 (8.6)	120 (15.0)	<0.001
STEMI	16 (18.8)	39 (32.5)	0.037
NSTEMI	41 (48.2)	44 (36.7)	0.114
UAP	17 (20.0)	33 (27.5)	0.250
Non-classified ACS	11 (13.0)	4 (3.3)	0.013
Life threatening events (LTE's)	22 (2.2)	23 (2.9)	0.448
Pulmonary embolism	6 (27.3)	7 (30.4)	0.815
Thoracic aortic dissection	4 (18.2)	2 (8.7)	0.349
Acute abdominal aneurysm	3 (13.6)	2 (8.7)	0.598
Other*	9 (40.9)	12 (52.2)	0.449
Non-urgent cardiovascular diseases**	194 (19.5)	170 (21.2)	0.384
Non-cardiac chest pain, not further	163 (16.4)	159 (19.8)	0.061
specified ***			
Musculoskeletal pain	199 (20.0)	113 (14.1)	0.001
Psychogenic disorders	139 (14.0)	67 (8.4)	<0.001
Gastrointestinal tract disorders	76 (7.7)	62 (7.7)	0.951
Respiratory tract disorders	52 (5.2)	45 (5.6)	0.727
Other non-urgent diagnoses***	63 (6.3)	43 (5.4)	0.380

^{*}Acute heart failure, stroke, severe COPD exacerbation, sepsis, coronary spasm probably caused by hypokalaemia, diabetic ketoacidosis, epileptic insult, bleeding from oesophageal varices, ovarian torsion, ventricular fibrillation.

^{**} Stable angina pectoris (including atypical chest pain), stable heart failure, arrhythmias, hypertension

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**** Amongst others: anemia, malignancy, vasovagal collapse, side effects medication, dermatologic diseases



Table 3. Association between urgency allocation, diagnose ACS and other life threatening events (LTE)

Women	ACS n=85 (8.6%)	No ACS n=908 (91.4%)	p-value 1*	p-value 2**
U1	62 (72.9)	371 (40.9)	<0.001	<0.001
U2	12 (14.1)	231 (25.4)	-	
U3-U5	11 (13.0)	306 (33.6)	_	
Men	ACS n=120 (15.0%)	No ACS n=682 (85.0%)		
U1	84 (70.0)	290 (42.5)	<0.001	<0.001
U2	19 (15.8)	142 (20.8)	-	
U3-U5	17 (14.2)	250 (36.7)	-	
Women	ACS or LTE n=107 (10.8%)	No ACS or LTE n=886 (89.2%)		
U1	71 (66.4)	362 (40.9)	<0.001	<0.001
U2	21 (19.6)	222 (25.1)	=	
U3-U5	15 (14.0)	302 (34.0)		
Men	ACS or LTE n=143 (17.8%)	No ACS or LTE n=659 (82.2%)		
U1	96 (67.1)	278 (42.2)	<0.001	<0.001
U2	24 (16.8)	137 (20.8)		
U3-U5	23 (16.1)	244 (37.0)		

^{*} P-value 1: U1 vs. U2, U3, U4 and U5

^{**} P-value 2: U1,U2 vs. U3,U4,U5

^{***}LTE = life threatening event. Life threatening events consist of: ACS; pulmonary embolism; thoracic aortic dissection; acute heart failure; stroke; abdominal aortic aneurysm; severe COPD exacerbation; diabetic ketoacidosis; coronary spasm probably caused by hypokalaemia; epileptic insult; bleeding from oesophageal varices; ovarian torsion; ventricular fibrillation.

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Appendix-Table 1: Urgency levels

U0	
	Reanimation
J1	Life-threatening, GP/ ambulance should arrive within 15 minutes
U2	Emergency, GP should arrive within 60 minutes
U3	Urgent, consultation by GP within three hours
U4	Routine, consultation by GP the same day
U5	Advice given by triage nurse

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Appendix-Table 2: Relation of caller to patient in women and men with ACS

	Women	Men	P-value
	n= 85 (8.6%)	n = 120 (15.0%)	
Someone else calls on behalf of patient	59 (69.4)	79 (65.8)	0.590
Partner	16 (18.8)	64 (53.3)	<0.001
Son, daughter or other family member	20 (23.5)	8 (6.7)	0.001
Nurse	15 (17.6)	4 (3.3)	<0.001
Other (neighbor, friend, colleague)	8 (9.4)	3 (2.5)	0.030

STRODE Statement		eklist of items that should be included in reports of <i>cross-sectional studies</i>	_
	Item No	Recommendation	Page No
Title and abstract	1	(a) Indicate the study's design with a commonly used term in the title or the	1
		abstract	
		(b) Provide in the abstract an informative and balanced summary of what was	3
		done and what was found	
Introduction			
Background/rationale	2	Explain the scientific background and rationale for the investigation being	5,6
8		reported	- , -
Objectives	3	State specific objectives, including any prespecified hypotheses	5,6
Methods			•
Study design	4	Present key elements of study design early in the paper	6
Setting	5	Describe the setting, locations, and relevant dates, including periods of	6,7
· ·		recruitment, exposure, follow-up, and data collection	
Participants	6	(a) Give the eligibility criteria, and the sources and methods of selection of	6,7
1		participants	
Variables	7	Clearly define all outcomes, exposures, predictors, potential confounders, and	6,7
		effect modifiers. Give diagnostic criteria, if applicable	
Data sources/	8*	For each variable of interest, give sources of data and details of methods of	6,7
measurement		assessment (measurement). Describe comparability of assessment methods if	
		there is more than one group	
Bias	9	Describe any efforts to address potential sources of bias	6
Study size	10	Explain how the study size was arrived at	6
/Quantitative	11	Explain how quantitative variables were handled in the analyses. If applicable,	6
variables		describe which groupings were chosen and why	
Statistical methods	12	(a) Describe all statistical methods, including those used to control for	7
		confounding	
		(b) Describe any methods used to examine subgroups and interactions	7
		(c) Explain how missing data were addressed	7
		(d) If applicable, describe analytical methods taking account of sampling	
		strategy	
		(e) Describe any sensitivity analyses	7
Results			
Participants	13*	(a) Report numbers of individuals at each stage of study—eg numbers	7
· wivio ip wiito	10	potentially eligible, examined for eligibility, confirmed eligible, included in	'
		the study, completing follow-up, and analysed	
		(b) Give reasons for non-participation at each stage	7
		(c) Consider use of a flow diagram	
Descriptive data	14*	(a) Give characteristics of study participants (eg demographic, clinical, social)	7,8
2 courper v uniu		and information on exposures and potential confounders	,,,
		(b) Indicate number of participants with missing data for each variable of	15
		interest	
Outcome data	15*	Report numbers of outcome events or summary measures	7
Main results	16	(a) Give unadjusted estimates and, if applicable, confounder-adjusted	7,8
	10	estimates and their precision (eg, 95% confidence interval). Make clear which	,,0
		confounders were adjusted for and why they were included	
		- comounders were adjusted for and why they were included	

		(b) Report category boundaries when continuous variables were categorized	7
		(c) If relevant, consider translating estimates of relative risk into absolute risk	NA
		for a meaningful time period	
Other analyses	17	Report other analyses done—eg analyses of subgroups and interactions, and	9
		sensitivity analyses	
Discussion			
Key results	18	Summarise key results with reference to study objectives	9
Limitations	19	Discuss limitations of the study, taking into account sources of potential bias	9,10
		or imprecision. Discuss both direction and magnitude of any potential bias	
Interpretation	20	Give a cautious overall interpretation of results considering objectives,	9,
		limitations, multiplicity of analyses, results from similar studies, and other	10,11
		relevant evidence	
Generalisability	21	Discuss the generalisability (external validity) of the study results	11
Other information			
Funding	22	Give the source of funding and the role of the funders for the present study	2
		and, if applicable, for the original study on which the present article is based	

^{*}Give information separately for exposed and unexposed groups.

Note: An Explanation and Elaboration article discusses each checklist item and gives methodological background and published examples of transparent reporting. The STROBE checklist is best used in conjunction with this article (freely available on the Web sites of PLoS Medicine at http://www.plosmedicine.org/, Annals of Internal Medicine at http://www.annals.org/, and Epidemiology at http://www.epidem.com/). Information on the STROBE Initiative is available at www.strobe-statement.org.

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Loes TCM Wouters¹, Dorien LM Zwart¹, Daphne CA Erkelens¹, Esther De Groot¹, Arno W Hoes², Roger AMJ Damoiseaux¹, Frans H Rutten¹

- Dept. General Practice, Julius Centre for Health Sciences and Primary Care, University Medical Centre, Utrecht University, Utrecht, the Netherlands
- 2. Dean University Medical Centre, Utrecht University, Utrecht, the Netherlands

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Corresponding author: L.T. Wouters, MD, Dept. General Practice, Julius Centre for Health Sciences and

Primary Care, University Medical Centre Utrecht, Utrecht University

STR 6.131, PO Box 85500, 3508 GA Utrecht, The Netherlands

Phone number: +31 (0)88 75 51470

Email: L.T.C.Wouters-2@umcutrecht.nl

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All authors have completed the <u>Unified Competing Interest form</u> and declare no support from any organization for the submitted work; no financial relationships with any organizations that might have an interest in the submitted work in the previous three years, no other relationships or activities that could appear to have influenced the submitted work.

Transparency declaration

The lead author (the manuscript's guarantor) affirms that the manuscript is an honest, accurate, and transparent account of the study being reported; that no important aspects of the study have been omitted; and that any discrepancies from the study as planned (and, if relevant, registered) have been explained.

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Objectives To assess differences in symptom presentation of women with and without an acute coronary syndrome (ACS), and the same in men, in patients with chest discomfort who contact out-of-hours primary care (OHS-PC).

Design Cross-sectional study.

Setting Nine OHS-PC in the Netherlands.

Participants 993 women and 802 men who called OHS-PC for acute chest discomfort (pain, pressure, tightness, or discomfort) between 2014 and 2016.

Primary outcome measure Diagnosis of ACS. We compared patient and call characteristics of triage call recordings between women with and without ACS, and men with and without ACS. Diagnoses were retrieved from the patient's medical record in general practice, including hospital specialists' discharge letters.

Results Among 1,795 patients (mean age 58.8 (SD 19.5) years, 55.3% women), 15.0% of men and 8.6% of women had an ACS. In both sexes, retrosternal chest pain was associated with ACS (women with ACS vs. without 62.3% vs. 40.3%, p=0.002, men with ACS vs. without 52.5% vs. 39.7%, p=0.032), as was pressing/tightening pain (women 78.6% vs. 61.5%, p=0.011, men 82.1% vs. 57.4%, p=<0.001) and radiation to arm (women 75.6% vs. 45.9%, p<0.001, men 56.0% vs. 34.8%, p<0.001). In women, severe pain (65.4% vs. 38.1%, p=0.006), a pale face (50.0% vs. 22.9%, p=0.007) and radiation to jaw (50.0% vs. 29.8%, p=0.017) were also associated with ACS, while in men this was sweating (52.4% vs. 38.1%, p=0.015), and against ACS stabbing pain (8.4% vs. 26.5%, p<0.001).

Ambulances were dispatched equally in women (72.9%) and men with ACS (70.0%).

Conclusion There were more similarities than differences in symptoms associated with the diagnosis ACS for women and men. Important exceptions were severity, type, and radiation of pain, and in women a pale face, and in men sweating.

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Keywords (5): acute coronary syndrome, gender, primary care, telephone triage, chest discomfort

Strengths and limitations of this study

- We could analyse the very initial conversation with symptom presentation of a large sample of patients calling for acute chest discomfort.
- We analysed the conversations without knowledge of the eventual diagnosis and therefore no hindsight bias of the researcher or recall bias of the patient could occur.
- Results are generalizable to comparable primary care settings in the United Kingdom and Scandinavian Countries, and our results may even be generalizable to emergency medical service (EMS, '112' or '911') settings.
- We had missing values on some clinical variables, a phenomenon common in routine care data.



Adequate triage and early diagnosis is key in patients with acute chest discomfort because they might have an acute coronary syndrome (ACS) for which lifesaving early interventions are available. ACS is an umbrella term including ST-elevated myocardial infarction (STEMI), non-ST-elevated myocardial infarction (NSTEMI) and unstable angina pectoris (UAP). ¹ Increased preventive measures and development of (timely) effective therapeutic interventions ('time is muscle') have resulted in improved outcomes and prognosis in ACS. ² Telephone triage of patients with chest discomfort, as done in out-of-hours service primary care (OHS-PC) and emergency medical services (EMS or ambulance dispatch centres) is, however, challenging because it is difficult to differentiate ACS from other causes of chest discomfort based on symptoms only. ^{3, 4} Importantly, the majority of patients with chest discomfort (80%) in the Netherlands first approach the general practitioner (GP) or OHS-PC, and 20% directly calls the ambulance (112) or are self-referrals to the emergency department (ED). ⁵

Previous hospital-based studies reported a delayed recognition of ACS in women compared to men. ^{6,7} It was suggested that this delayed recognition was related to a less specific presentation in women. ^{8,9} This caused an ongoing debate on whether women with ACS compared to men present with less specific symptoms, and how this affects diagnosis, but also treatment, and prognosis. ^{10,11} A recent meta-analysis of 27 studies showed that women with ACS compared to men with ACS had higher odds of presenting with pain between the shoulder blades (OR 2.15, 95% CI 1.95-2.37), nausea or vomiting (OR 1.64, 95% CI 1.48-1.82) and shortness of breath (OR 1.34 (95% CI 1.21-1.48). ¹² Women with ACS had lower odds of sweating (OR 0.84, 95% CI 0.76-0.94) and presenting with chest pain (OR 0.70, 95% CI 0.63-0.78), but in both sexes chest pain remained the most common symptom (pooled prevalence men 79% and women 74%). ¹² Importantly, researchers suggested standardization in methods of symptoms assessment is needed, because of the difficulties to formulate any definitive statements about symptom presentation, as studies assessed symptoms in different ways (questionnaires or abstracting from medical records). ^{9, 13-15} Abstracting symptom presentation from medical records may dilute symptom presentation, as they are translated by the clinician in medical terminology. ¹⁶ Moreover, many studies suffer from recall or hindsight bias of both patient and researcher as they know the outcome (ACS).

For the clinician or telephone triage nurse it is crucial to differentiate ACS from other causes of chest discomfort. For that, studies are needed that include female and male patients presenting with chest discomfort, in which women and men who turn out to have ACS are compared to those who do not

have ACS. Such studies are scarce. In a study performed among patients with chest discomfort seen at the ED in the USA (77 women and 244 men with ACS, and 195 women and 240 men without ACS were compared. Women with ACS more often reported arm pain than women without ACS (47% vs. 32%, p=0.021), while men with ACS reported pressing feeling (63% vs. 54%, p=0.035) and chest pain (72% vs 60%, p=0.005) more often than men without ACS. ¹³ In a recent Dutch OHS-PC study among 23 women and 34 men with ACS, and 253 women and 208 men without ACS, differences in symptom presentation between women and men were small. ¹⁷ In men, radiation of pain (89.3% vs. 64.9%, p-value=0.011) was discriminative for ACS, while stabbing chest pain (3.7% vs. 24.0%, p-value=0.014) was discriminative for the absence of ACS. Both these symptoms were not discriminative in women (90.0% vs 78.6%, p=0.227, and 15.8% vs. 18.8%, p=0.743), respectively. ¹⁷

We aimed to assess whether symptoms were different in patients with ACS from patients without ACS in women and men separately. This, in the domain patients presenting with acute chest discomfort who contact the OHS-PC. For analyses we used the very initial symptom presentation as available from the recorded telephone triage conversations.

Methods

We performed a cross-sectional study with a random sample of 1,795 OHS-PC calls for chest discomfort (chest pain, pressure, tightness, or discomfort) between 2014 and 2016. ¹⁸ We first selected calls on the basis of International Code for Primary Care (ICPC; a WHO world-wide code system for primary care) codes (K01, K02, K03, K24, K74, K75, K76, K77, K93, L04, P74, R02, R98) and keywords thoracic pain, chest pain, myocardial infarction, heart attack and their common abbreviations mentioned by the triage nurse in the electronic medical file at the OHS-PC. ^{19, 20} General practitioners who work at the OHS-PC assign the ICPC codes to the call (see also appendix-table1). We combined ICPC-codes and keywords to achieve a sample with a broad variety of symptoms to capture the entire domain of patients suspected of ACS. We drew a random sample of all available calls of these patients with the Random Number Generator (RAND) function in Microsoft Excel. Calls were excluded before re-listening when the patients' age was below 18 years or when the patient did not live in the surrounding area of the OHS-PCs (because then we could not retrieve a diagnosis from the general practitioner of these patients). Calls were excluded during re-listening when it not concerned a triage call (e.g. inter-collegial consultation) or when the recording was of poor quality (Figure 1). For a descriptive observational study,

We re-listened the telephone triage recordings and collected information about patient and conversation characteristics, on symptom presentation, medical history, urgency allocation, and involvement of a supervising general practitioner (GP) in the triage. Nine OHS-PC in the Netherlands participated, serving a total population of 1.5 million people. The diagnosis was made after the phone call, which was in the case of ACS nearly always done by the cardiologist (97.1%) in the hospital based on (i) symptom presentation, (i) levels of (high-sensitivity) troponin and (iii) electrocardiography results. The final diagnoses were provided by the patients' GP, based on the electronic medical file including ED and cardiologist discharge letters, and also the notes from the OHS-PC. We used medical information up to 30-days following the contact with the OHS-PC, to allow us to include diagnoses of ACS that was initially missed because the patient was not referred to the cardiologist the same day of the OHS-PC contact. In none of the patients in the study we had evidence of a missed diagnosis of ACS.

Context

In the Netherlands, OHS-PC covers primary care during 73% of the week hours, and the initial contact is by telephone. In most OHS-PC and EMS, the 'Netherlands Triage Standard' (NTS) is used as a decision support to classify the urgency of the patients' conditions. ²¹ Based on the patient's symptom presentation, the triage nurse needs to choose the most appropriate complaint out of 56 'main complaints'. Each NTS 'main complaint' incorporates a decision tree with hierarchically ordered questions, which are similar for men and women. Triage nurses fill out the caller's responses in the semi-automatic NTS system, which then generates urgency allocations linked to a timeframe within which the patient should be seen by a physician or ambulance personnel (U0 (reanimation) to U5 (selfcare advice, see also appendix-table 2. The triage nurse can overrule this recommendation and up- or downscale the urgency allocation, often after consulting the supervising GP. ²² Since its introduction in 2011, the NTS system has, however, never been formally validated by correlating the generated urgencies to clinical endpoints. ²¹ All telephone calls to the OHS-PC are routinely recorded and archived for five years for training and quality control purposes.

Data analyses

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We compared patient and call characteristics between women with and without ACS, and men with and without ACS. We analysed the association between urgency allocation and the final diagnosis ACS (with or without other life threatening events (LTEs)). We considered pulmonary embolism, thoracic aortic dissection and acute abdominal aneurysm as LTEs; patients with LTEs as well as those with ACS should receive an U1-level urgency.

For comparison of dichotomous variables we used the Chi² test or Fisher exact test and for continuous variables the independent sample t-test or Mann-Whitney U test. Data analysis was performed using SPSS, IBM version 25.

Patient and public involvement

No patients were involved in setting the research question or the outcome measures, or in developing plans for design, however, they were involved in the implementation of the study. In addition, they were asked to advise on interpretation and writing up of results. Results will be shared and discussed with the national patient community of cardiovascular diseases ('Harteraad').

Ethics

The study (National Trial Register identification number: NTR7331) was approved by the Medical Ethics Committee Utrecht, the Netherlands (reference number WAG/mb/16/003208) and complied with the Declaration of Helsinki. A waiver of informed consent was given because our study had minimal risk to subjects and could otherwise not be carried out logistically. Personal and research data were handled and stored according to the European General Data Protection Regulation.

Results

Among the 1,795 callers with chest discomfort (mean age 58.8 (SD 19.5) years, 55.3% women), 8.6% of women and 15.0% of men had an ACS. In women with an ACS, 18.8% had a STEMI, 48.2% a NSTEMI, 20.0% an UAP, and 13.0% non-classified ACS. In men with ACS, 32.5% had a STEMI, 36.7% a NSTEMI, 27.5% an UAP, and 3.3% non-classified ACS.

A total of 22 (2.2%) women and 23 men (2.9%) had another LTE than ACS (e.g. pulmonary embolism, thoracic aortic dissection, acute abdominal aneurysm, acute heart failure).

Patient and call characteristics

In table 1, patient and call characteristics are presented for women and men with and without ACS. Men and women with ACS were older than those without ACS (mean age of women 73.6 vs. 57.8 years, p<0.001, men 67.2 vs. 56.9 years, p<0.001), and the mean duration of the telephone calls shorter (women 6:47 vs. 7:47 minutes, p=0.021, men 6:31 vs. 7:33 minutes, p=0.004). The GP was consulted for supervision by the triage nurse in the majority of cases (52.2% in women and 55.5% in men, p=0.161). However, in women with ACS, the GP was less often consulted than in women without ACS (41.2% vs. 53.2%, p=0.034). In men such a difference was not observed (53.3% vs. 55.9%, p=0.607). In around half of the calls, someone else called initially on behalf of the patient; somewhat less often in women than in men (49.5% vs. 54.7%, p=0.029). In cases with ACS, for both sexes more often someone else called than in those without ACS (in women 69.4% vs. 47.7%, p<0.001, in men 65.8% vs. 52.8%, p=0.008). In men with ACS, most often their female partner (53.3%) called, while in women with ACS, it was either their male partner (17.6%), their daughter (20.0%) or a nurse (17.6%). See appendix-table 3. Callers expressed concerns in nearly all calls, also in those without an ACS; women with ACS vs. women without ACS 97.3% vs. 88.9% (p=0.109), and men with ACS vs. men without ACS 96.3% vs. 86.5% (p=0.041).

Both women and men with ACS had more often a history of coronary artery disease (women 42.5% vs. 25.1%, p=0.017, and men 57.0% vs. 38.4%, p=0.002), but women with ACS had more often a history of diabetes (41.4% vs. 14.6%, p<0.001).

Symptom presentation

Chest pain was the most common complaint, both in those with and without an ACS; in women with and without ACS 98.8% and 93.1% (p=0.055), and in men 92.4% and 94.5%, respectively (p=0.364). Retrosternal located chest pain was more common in women and men with ACS than in those without ACS (women 62.3% vs. 40.3%, p=0.002, and men 52.5% vs. 39.7%, p=0.032). Also, radiation of pain to the arms seemed indicative of ACS in both sexes (women with vs. without ACS 75.6% vs. 45.9%, p<0.001, and men 56.0% vs. 34.8%, p<0.001), as was pressing/heavy/tightening chest pain (women with vs. without ACS 78.6% vs. 61.5%, p=0.011 and men 82.1% vs. 57.4%, p=<0.001). Only in women radiation to the jaw (50.0% vs. 22.9%, p=0.007) and severe pain (8 or more on a Numeric Rating Scale 0-10) was associated with ACS (65.4% vs. 38.1, p=0.006). Only in men, stabbing pain was very rare in those with ACS (8.4% vs. 26.5%, p<0.001).

Of the autonomous nervous system (ANS)-related symptoms, nausea/vomiting and dizziness/near fainting were not associated with ACS in either sex, with the exception of a pale face that was in women (50.0% vs. 29.8%, p=0.017), and sweating in men (52.4% vs. 38.1%, p=0.015).

Recognition of symptoms being similar to a previous cardiac event was associated with ACS in men (52.9% vs. 32.1%, p=0.004), but not clearly for ACS in women (32.5% vs. 21.4%, p=0.108). Subgroup analyses in 56 women and 58 men with diabetes showed that both women (85.7% vs. 58.3%, p<0.001) and men with diabetes (67.2% vs. 51.5%, p=0.033) more often had shortness of breath than those without diabetes, but as often chest discomfort (women 90.9% vs. 95.0%, p=0.193, men 89.2% vs

94.1%, p=0.162). Shortness of breath in patients with diabetes was not related to ACS diagnosis (women

81.8% vs. 86.7%, p=0.680, men 75.0% vs. 66.0%, p=0.615).

Diagnoses

Of the 205 patients with an ACS (85 women, 120 men), 55 (26.8%) patients had a STEMI (women 18.8%, men 32.5%), 85 (41.5%) a NSTEMI (women 48.2%, men 36.7%), 50 (24.4%) unstable angina pectoris (UAP) (women 20.0%, men 27.5%) and 15 (7.3%) unspecified ACS (women 13.0%, men 3.3%), the latter also including two sudden cardiac deaths in women and one in men (Table 2). In nearly all cases (97.1%) the ACS diagnosis was made by a cardiologist based on symptom presentation, troponin levels and electrocardiography. Three patients died before arrival of the ambulance (they were classified as acute cardiac death) and one patient died after resuscitation at the ED. Two patients were classified as ACS by the GP; they were not referred to the hospital because of short life expectancy due to cancer. There were 45 patients with other LTEs (2.5%) and the majority of patients had non-urgent medical conditions (86.1%). The most common non-urgent diagnoses in both sexes were (i) non-urgent cardiovascular diseases such as stable angina pectoris, stable heart failure and arrhythmias (19.5% of all female callers with chest discomfort vs. 21.2% of male callers, p=0.384) and (ii) non-cardiac unspecified chest pain (women 16.4% vs. 19.8% men, p=0.061). Women more often than men were diagnosed with musculoskeletal problems (women 20.8% vs. men 14.1%, p=0.001) and psychogenic conditions (women 14.0% vs. men 8.4%, p<0.001). Of the patients who were diagnosed with a non-ACS diagnoses, 45.4% were classified by a cardiologist, 5.5% by another hospital specialist (e.g. pulmonologist or internal medicine specialist) and the remaining patients were diagnosed by a GP.

Urgencies

Women and men with chest discomfort were equally sent an ambulance (43.6% vs. 46.6%, p=0.200). This was also in women and men who had an ACS (72.9% vs. 70.0%, p=0.647), and in those with either ACS or other LTEs (66.4% vs. 67.1%, p=0.897). See table 3.

Discussion

For both sexes, retrosternal pain, pain described as pressing, heavy or tightening, and radiation to the arm were associated with ACS in patients who contacted the OHS-PC for chest discomfort. Radiation to the jaw, severe pain, and a pale face was indicative for ACS in women. Sweating was positively associated to ACS, and stabbing pain negatively associated to ACS in men with chest discomfort. Women and men with chest discomfort as also those with ACS were equally often sent an ambulance.

Our finding that radiation of pain to the arm and retrosternal chest pain was discriminative for diagnosing ACS in both sexes was also reported in a study among 2,475 patients with acute chest pain in a multicentre ED-study. ²³ Another ED-study among 1,334 patients with ACS showed that regardless of ethnics status the most common presenting symptom was retrosternal pain/discomfort of any intensity. ²⁴ The aforementioned US study in the ED-setting reported that radiation to the arm was indicative for ACS in women but not for men, and chest pressure was distinctive for men but not for women. ¹³ The only previously published OHS-PC study reported the opposite; radiation to the arm was associated with ACS in men, but not in women. However, similar to our study, this study reported that stabbing pain was very uncommon in men with ACS. ¹⁷

In our study, women with ACS had more often a history of diabetes and were older than men with ACS, which is in line with other studies. ^{8, 12} Some studies claim that patients with diabetes more often have atypical symptoms of ACS, however a review of eight studies concluded the evidence of these studies was conflicting. ²⁵ We showed that both women and men with diabetes had more often shortness of breath than those without diabetes, but shortness of breath in patients with diabetes was not helpful to diagnose ACS. Regarding dispatching the ambulance, the aforementioned OHS-PC study and two EMS studies showed, similar to our findings, that there was no difference in dispatch priorities between men and women with ACS. ^{4, 17, 26} This is in contrast with studies that show delay in hospital presentation of women with ACS. ^{9, 27}

We need to realize that focusing on differences may blur the large overlap in symptoms. Moreover, comparing *selectively* women with ACS to men with ACS as many previous studies did, is clinically

irrelevant. ^{9, 14, 28} Clinicians, including GPs, and triage nurses need to know whether and how women with ACS differ from women without ACS, with the same question for men. Nevertheless, even guidelines stick to comparing those with established disease, and express the view that women with ACS more likely present with less specific symptoms than men with ACS. ^{1, 29} Unfortunately, public awareness campaigns follow this reasoning, and over-emphasize sex differences in women awareness campaigns ('Go Red for Women' in the United States and 'Invisible me' in Australia). ^{7, 28, 30} Unbalanced attention to differences, neglecting the much larger overlap may even introduce new blind spots in recognizing ACS in women. ³¹

A likely reason behind the predominant message that women present with other ACS symptoms than men is the difference in pathophysiology of coronary artery disease. Women compared to men more often have elongated plaques, located on bifurcations in epicardial coronaries, coronary spasm, microvascular dysfunction, and spontaneous coronary dissection. ^{30, 32} These pathophysiological differences have an effect on interventional treatment and prognosis. ^{2, 32} However, these differences do not necessarily imply an effect on symptom presentation because the pain pathway is equal in women and men: i.e. triggered by myocardial ischemia. ^{33, 34} A supply and demand mismatch of the myocardial oxygen consumption triggers sensory nerve endings in the myocardium and cause ischemia symptoms, and this is irrespective of the fact whether the ischemia is caused by a plaque rupture in an epicardial artery or spasm, or any other cause. ³⁴ The sex differences in pathophysiology of ACS do therefore not support the belief in differences in ACS symptoms between women and men.

Another reason behind the belief of 'vague' symptom presentation in women with ACS might be that they seem to present a larger number of symptoms than men with ACS, and this may be interpreted as 'vague' by physicians. ^{15, 35} Presentation of multiple symptoms may influence the prompt recognition of heart disease and initial actions on the part of health care providers. ^{35, 36} In a study from 2018, 2009 women and 976 men hospitalized for myocardial infarction, healthcare providers initially thought symptoms of women (53.4%) were less often heart-related than in men (36.7%). ³⁵ Women and men had the same chest pain symptoms, but women reported more additional symptoms. ³⁵ In our study, the call duration and the number of GP consultations by the triage nurse were similar among women and men, suggesting that triage nurses seem not to experience more difficulties in interpreting symptoms in women than men. This is in line with a prospective study with 2,795 patients

with chest discomfort in ED-setting that showed the physicians' diagnostic uncertainty for the presence of ACS in women was not more common as compared to men.³⁷

Interestingly, in the majority of calls in our study someone else than the patient called the OHS-PC (women with ACS 69.4%, without ACS 47.7%, and men with ACS 65.8%, and without ACS 52.8%). This was also highlighted in an Australian study among 1,681 patients with an acute myocardial infarction; in 90.5% of the women with AMI someone else called on behalf of the patient and in 87.8% of the men with AMI. ³⁸ According to protocol in OHS-PC, triage nurses ask the patient to the phone, this to prevent loss of (paralinguistic) information from the patient him/herself. In our study, in about 50% of the conversations the patient took over the phone call.

Strengths and limitations

We could analyse the very initial conversation with symptom presentation of a large sample of patients calling the OHS-PC because of chest discomfort. We analysed the conversations without knowledge of the eventual diagnosis (no hindsight bias of the researcher or recall bias of the patient). Another strength is that our results are generalizable to comparable primary care settings, e.g. UK and Scandinavian countries, and possibly some other European countries.³ Our results may even be generalizable to EMS settings, since the prior probability of having an ACS is comparable in EMS setting as in OHS-PC settings. ^{4 39}

We had missing values on some clinical variables, a phenomenon common in routine care data. Future research could focus on developing a multivariable prediction model useful with telephone triage to estimate the risk of ACS in men and women suspected of ACS.

Conclusions

There were more similarities than differences in symptoms associated with the diagnosis ACS for women and men. Important exceptions were severity, type, and radiation of pain, and in women a pale face, and in men sweating.

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Data sharing statement

The data can be made available for researchers whose proposed use of the data has been approved at request of the corresponding author, with a signed data access agreement.

Author contribution

FR and DZ are the lead investigators who conceived the research idea and methodology. Funding acquisition was done by FR, DZ and RD. LW and DC conducted data acquisition. LW performed the analyses and wrote the first draft of the manuscript. She was supervised by FR and DZ, who critically revised the manuscript. DC, EdG, AH and RD contributed with and approved the final version of the manuscript.

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Table 1. Baseline characteristics of 993 (55.3%) women and 802 (44.7%) men patients contacting the OHS-PC with chest discomfort (pain, pressure, tightness, or discomfort)

<u>0</u> 1	993 women			802 men		
2	333 Women			302 men		
3	ACS	No ACS	p-value	ACS	No ACS	p-value
	ACS	NO ACS	p-value	ACS	NO ACS	p-value
4 Characteristics	N = 85 (8.6%)	N = 908 (91.4%)		N = 120 (15.0%)	N = 682 (85.0%)	
6	14 - 85 (8.0%)	14 - 308 (31.470)		N - 120 (13.0%)	N - 082 (83.0%)	
Mean age in years (SD)	73.6 (±14.1)	57.8 (±20.1)	<0.001	67.2 (±13.0)	56.9 (±19.2)	<0.001
8	73.0 (±14.1)	37.8 (120.1)	<0.001	07.2 (±13.0)	30.9 (±19.2)	<0.001
Pall characteristics						
0						
Mean call duration in min (SD)	6:47 (±5:16)	7:47 (±3:48)	0.021	6:31 (±3:13)	7:33 (±3:42)	0.004
22	0.47 (±3.10)	7.47 (±3.46)	0.021	0.31 (±3.13)	7.33 (±3.42)	0.004
Mean patient's introduction duration in	0:14 (± 0:08)	0:19 (±0:13)	<0.001	0:16 (±0:11)	0:19 (±0:14)	0.060
4	0.14 (± 0.06)	0.19 (±0.15)	<0.001	0.16 (±0.11)	0.19 (±0.14)	0.060
ក្សាin (SD)						
11111 (3D) 16						
Triage nurse consulted the GP	35 (41.2)	483 (53.2)	0.034	64 (53.3)	381 (55.9)	0.607
18	33 (41.2)	463 (33.2)	0.034	04 (33.3)	361 (33.9)	0.007
Someone else called on behalf of patient	59 (69.4)	433 (47.7)	<0.001	79 (65.8)	360 (52.8)	0.008
0	39 (09.4)	455 (47.7)	0.001	79 (03.8)	300 (32.8)	0.008
The nationt or person who called	36 (97.3)	409 (88.9)	0.109	52 (96.3)	321 (86.5)	0.041
The patient or person who called	30 (37.3)	409 (88.9)	0.103	32 (90.3)	321 (80.3)	0.041
gxpressed concerned (n=922)						
4						
Shest pain (n=1739)	81 (98.8)	817 (93.1)	0.055	110 (92.4)	624 (94.5)	0.364
6			•			
Severe Pain (>7 on a scale 0-10) (n=753)	17 (65.4)	147 (38.1)	0.006	13 (23.6)	87 (30.4)	0.312
8						
9 uration						
-0						
-1> 15 min (n=1501)	66 (100)	729 (95.8)	0.102	99 (97.1)	541 (94.6)	0.292
2						
-3× 12 hrs (n=1563)	60 (85.7)	575 (72.7)	0.018	88 (81.5)	431 (72.6)	0.052
4						
Σ̄ocation (n=1267)*						
.6						
Retrosternal	33 (62.3)	263 (40.3)	0.002	42 (52.5)	191 (39.7)	0.032
.8						
Right or left side thorax	14 (26.4)	260 (39.8)	0.054	26 (32.5)	211 (43.9)	0.057
0						
Type of pain (n=1334)**						
Pressing/heavy/tightening	44 (78.6)	423 (61.5)	0.011	78 (82.1)	284 (57.4)	<0.001
4						

2						
³ Stabbing 4	8 (14.3)	155 (22.5)	0.152	8 (8.4)	131 (26.5)	<0.001
5 Radiation of chest pain (n=1391) ***						
7 Any location	61 (85.9)	485 (68.6)	0.002	65 (63.7)	292 (57.1)	0.218
9 Arm 10	31 (75.6)	188 (45.9)	<0.001	47 (56.0)	117 (34.8)	<0.001
11 Between the shoulder blades	11 (52.4)	159 (41.7)	0.336	14 (27.5)	91 (29.4)	0.781
13 14 Jaws	10 (50.0)	66 (22.9)	0.007	1 (2.6)	28 (11.6)	0.146
15 Shortness of breath (n=1365)	47 (72.3)	455 (66.3)	0.328	53 (60.9)	329 (62.4)	0.788
17 Symptoms similar to previous cardiac	13 (32.5)	79 (21.4)	0.108	27 (52.9)	92 (32.1)	0.004
1 9 vents (n=748) 20						
	· ·					
2ANS-related symptoms 22						
2 3 weating (n=1130) 24	28 (49.1)	234 (42.5)	0.340	43 (52.4)	168 (38.1)	0.015
2¶ausea or vomiting (n=808) 26	16 (44.4)	240 (56.2)	0.173	27 (45.8)	122 (42.7)	0.661
2₱allor (n=634) 28	17 (50.0)	91 (29.8)	0.017	20 (40.8)	82 (33.)	0.315
29 Dizziness or near fainting n=1599) 30	13 (18.6)	197 (24.4)	0.277	17 (15.9)	127 (20.2)	0.249
³ Medical history 32						
3 eV disease or risk factors (n=1461) 34	54 (78.3)	407 (57.5)	0.001	75 (72.1)	359 (61.9)	0.046
35 Coronary artery disease (n=966) 36	17 (42.5)	114 (25.1)	0.017	45 (57.0)	151 (38.4)	0.002
37 38 Hypertension (n=756)	22 (71.0)	142 (35.1)	<0.001	17 (47.2)	94 (33.0)	0.091
39 40 Diabetes mellitus (n=745)	12 (41.4)	56 (14.6)	<0.001	12 (28.6)	59 (20.3)	0.220
41 Hypercholesterolemia or use of statins 42	8 (36.4)	81 (23.2)	0.161	19 (45.2)	62 (23.3)	0.003
43 44 ⁽ⁿ⁼⁶⁷⁹⁾						
45						

*P-value comparing retrosternal or left/right side thorax vs. others locations of pain together (restrosternal, left/right side thorax,

48

57 58

59

⁴Back/shoulder, epigastric region)

^{49*} Pressing/ heavy/tightening pain vs. other types of pain (stabbing, burning, cramping, tearing). Stabbing pain: stabbing vs. other types of 50 pain (pressing/heavy/tightening, burning, cramping, tearing) 51

P-value comparing radiation arm or back/shoulder or jaws vs. no radiation

⁵⁴ NRS: Numeric Rating Scale 55

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BANS-related symptoms: Autonomous nervous system related symptoms 5CV disease or risk factors; a history of previous coronary artery disease, heart failure, stroke, cardiac arrhythmia, hypertension, and/or able or unstable a. diabetes (patient reported Coronary artery disease: History of prior MI, PCI, CABG, stable or unstable angina pectoris (patient reported)

	Women n= 993 (55.3%)	Men n=802 (44.7%)	p-value
Acute coronary syndrome	85 (8.6)	120 (15.0)	<0.001
STEMI	16 (18.8)	39 (32.5)	0.037
NSTEMI	41 (48.2)	44 (36.7)	0.114
UAP	17 (20.0)	33 (27.5)	0.250
Non-classified ACS	11 (13.0)	4 (3.3)	0.013
Life threatening events (LTE's)	22 (2.2)	23 (2.9)	0.448
Pulmonary embolism	6 (27.3)	7 (30.4)	0.815
Thoracic aortic dissection	4 (18.2)	2 (8.7)	0.349
Acute abdominal aneurysm	3 (13.6)	2 (8.7)	0.598
Other*	9 (40.9)	12 (52.2)	0.449
Non-urgent cardiovascular diseases**	194 (19.5)	170 (21.2)	0.384
Non-cardiac chest pain, not further	163 (16.4)	159 (19.8)	0.061
specified ***			
Musculoskeletal pain	199 (20.0)	113 (14.1)	0.001
Psychogenic disorders	139 (14.0)	67 (8.4)	<0.001
Gastrointestinal tract disorders	76 (7.7)	62 (7.7)	0.951
Respiratory tract disorders	52 (5.2)	45 (5.6)	0.727
Other non-urgent diagnoses***	63 (6.3)	43 (5.4)	0.380

^{*}Acute heart failure, stroke, severe COPD exacerbation, sepsis, coronary spasm probably caused by hypokalaemia, diabetic ketoacidosis, epileptic insult, bleeding from oesophageal varices, ovarian torsion, ventricular fibrillation.

^{**} Stable angina pectoris (including atypical chest pain), stable heart failure, arrhythmias, hypertension

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**** Amongst others: anaemia, malignancy, vasovagal collapse, side effects medication, dermatologic diseases



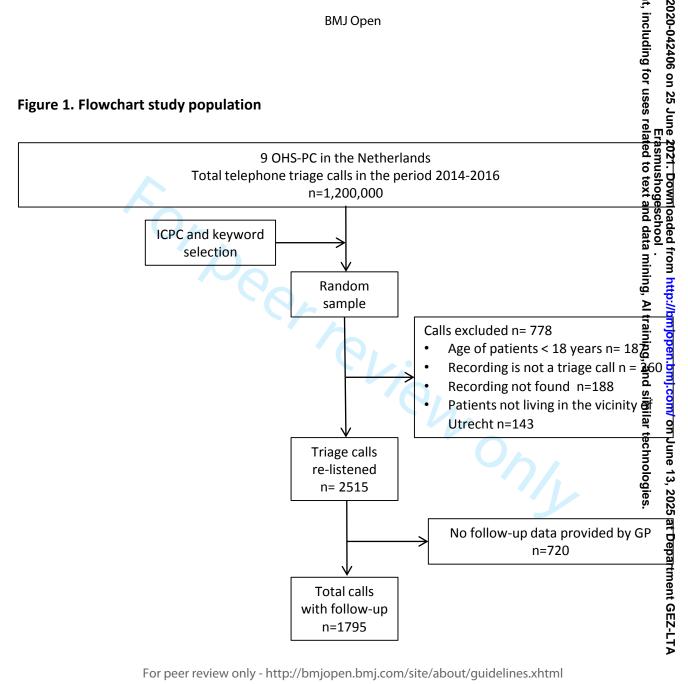
Women	ACS n=85 (8.6%)	No ACS n=908 (91.4%)	p-value 1*	p-value 2**
U1	62 (72.9)	371 (40.9)	<0.001	<0.001
U2	12 (14.1)	231 (25.4)	_	
U3-U5	11 (13.0)	306 (33.6)	_	
Men	ACS n=120 (15.0%)	No ACS n=682 (85.0%)		
U1	84 (70.0)	290 (42.5)	<0.001	<0.001
U2	19 (15.8)	142 (20.8)	_	
U3-U5	17 (14.2)	250 (36.7)		
Women	ACS or LTE n=107 (10.8%)	No ACS or LTE n=886 (89.2%)		
U1	71 (66.4)	362 (40.9)	<0.001	<0.001
U2	21 (19.6)	222 (25.1)		
U3-U5	15 (14.0)	302 (34.0)		
Men	ACS or LTE n=143 (17.8%)	No ACS or LTE n=659 (82.2%)		
U1	96 (67.1)	278 (42.2)	<0.001	<0.001
U2	24 (16.8)	137 (20.8)		
U3-U5	23 (16.1)	244 (37.0)		

^{*} P-value 1: U1 vs. U2, U3, U4 and U5

^{**} P-value 2: U1,U2 vs. U3,U4,U5

^{***}LTE = life threatening event. Life threatening events consist of: ACS; pulmonary embolism; thoracic aortic dissection; acute heart failure; stroke; abdominal aortic aneurysm; severe COPD exacerbation; diabetic ketoacidosis; coronary spasm probably caused by hypokalaemia; epileptic insult; bleeding from oesophageal varices; ovarian torsion; ventricular fibrillation.

Figure 1. Flowchart study population



Supplementary file

Appendix-Table 1: Overview of ICPC-codes used to select calls

ICPC code	Calls of patients with ACS	Calls of patients without ACS
	N=205 (%)	N=1,590 (%)
K01 Pain attributed to the heart	112 (54.6)	485 (30.5)
K02 Pressure/tightness attributed to the heart	39 (19.0)	184 (11.6)
K03 Other cardiovascular pain	1 (0.5)	6 (0.4)
K24 Fear of heart attack	1 (0.5)	8 (0.5)
K74 Angina pectoris	19 (9.3)	101 (6.4)
K75 Acute myocardial infarction	7 (3.4)	22 (1.4)
K76 Other/chronic ischaemic heart disease	0 (0.0)	2 (0.1)
K77 Heart failure	0 (0.0)	11 (0.7)
K93 Pulmonary embolism	0 (0.0)	15 (0.9)
L04 Chest discomfort	25 (12.2)	689 (43.9)
P74 Anxiety disorder	0 (0.0)	12 (0.8)
R02 Shortness of breath	1 (0.5)	36 (2.3)
R98 Hyperventilation	0 (0.0)	10 (0.6)

Appendix-Table 2: Urgency levels

Urgency level	Implication
U0	Reanimation
U1	Life-threatening, GP/ ambulance should arrive within 15 minutes
U2	Emergency, GP should arrive within 60 minutes
U3	Urgent, consultation by GP within three hours
U4	Routine, consultation by GP the same day
U5	Advice given by triage nurse

	Women	Men	P-value
	n= 85 (8.6%)	n = 120 (15.0%)	
Someone else calls on behalf of patient	59 (69.4)	79 (65.8)	0.590
Partner	16 (18.8)	64 (53.3)	<0.001
Son, daughter or other family member	20 (23.5)	8 (6.7)	0.001
Nurse	15 (17.6)	4 (3.3)	<0.001
Other (neighbour, friend, colleague)	8 (9.4)	3 (2.5)	0.030
Patient takes over the phone call on request	27 (45.8)	58 (73.4)	<0.001
of the triage nurse			

STROBE Statement—Checklist of items that should be included in reports of *cross-sectional studies*

STROBE Statement-	—Chec	eklist of items that should be included in reports of <i>cross-sectional studies</i>	1
	Item No	Recommendation	Page No
Title and abstract	1	(a) Indicate the study's design with a commonly used term in the title or the abstract	1
		(b) Provide in the abstract an informative and balanced summary of what was done and what was found	3
Introduction			
Background/rationale	2	Explain the scientific background and rationale for the investigation being reported	5,6
Objectives	3	State specific objectives, including any prespecified hypotheses	5,6
Methods		A	•
Study design	4	Present key elements of study design early in the paper	6
Setting	5	Describe the setting, locations, and relevant dates, including periods of	6,7
		recruitment, exposure, follow-up, and data collection	
Participants	6	(a) Give the eligibility criteria, and the sources and methods of selection of participants	6,7
Variables	7	Clearly define all outcomes, exposures, predictors, potential confounders, and	6,7
Data saumass/	0*	effect modifiers. Give diagnostic criteria, if applicable	6.7
Data sources/	8*	For each variable of interest, give sources of data and details of methods of assessment (measurement). Describe comparability of assessment methods if	6,7
measurement		there is more than one group	
Bias	9	Describe any efforts to address potential sources of bias	6
Study size	10	Explain how the study size was arrived at	6
/Quantitative	11	Explain how due study size was arrived at: Explain how quantitative variables were handled in the analyses. If applicable,	6
variables	11	describe which groupings were chosen and why	
Statistical methods	12	(a) Describe all statistical methods, including those used to control for confounding	7
		(b) Describe any methods used to examine subgroups and interactions	7
		(c) Explain how missing data were addressed	7
		(d) If applicable, describe analytical methods taking account of sampling	,
		strategy	7
		(\underline{e}) Describe any sensitivity analyses	7
Results			Τ_
Participants	13*	(a) Report numbers of individuals at each stage of study—eg numbers	7
		potentially eligible, examined for eligibility, confirmed eligible, included in	
		the study, completing follow-up, and analysed (b) Give reasons for non-participation at each stage.	7
		(b) Give reasons for non-participation at each stage (c) Consider use of a flow diagram	/
Descriptive data	14*	(a) Give characteristics of study participants (eg demographic, clinical, social)	7,8
Descriptive data	14	and information on exposures and potential confounders	7,0
		(b) Indicate number of participants with missing data for each variable of	15
		interest	13
Outcome data	15*	Report numbers of outcome events or summary measures	7
	16	(a) Give unadjusted estimates and, if applicable, confounder-adjusted	7,8
Main results	10	(, anagueta termate ana, ii appiitatie, contounat aajastea	1 ,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,
Main results		estimates and their precision (eg, 95% confidence interval). Make clear which	

			_
		(b) Report category boundaries when continuous variables were categorized	7
		(c) If relevant, consider translating estimates of relative risk into absolute risk	NA
		for a meaningful time period	
Other analyses	17	Report other analyses done—eg analyses of subgroups and interactions, and	9
		sensitivity analyses	
Discussion			
Key results	18	Summarise key results with reference to study objectives	9
Limitations	19	Discuss limitations of the study, taking into account sources of potential bias	9,10
		or imprecision. Discuss both direction and magnitude of any potential bias	
Interpretation	20	Give a cautious overall interpretation of results considering objectives,	9,
		limitations, multiplicity of analyses, results from similar studies, and other	10,11
		relevant evidence	
Generalisability	21	Discuss the generalisability (external validity) of the study results	11
Other information			
Funding	22	Give the source of funding and the role of the funders for the present study	2
		and, if applicable, for the original study on which the present article is based	

^{*}Give information separately for exposed and unexposed groups.

Note: An Explanation and Elaboration article discusses each checklist item and gives methodological background and published examples of transparent reporting. The STROBE checklist is best used in conjunction with this article (freely available on the Web sites of PLoS Medicine at http://www.plosmedicine.org/, Annals of Internal Medicine at http://www.annals.org/, and Epidemiology at http://www.epidem.com/). Information on the STROBE Initiative is available at www.strobe-statement.org.

BMJ Open

Gender-stratified analyses of symptoms associated with acute coronary syndrome in telephone triage: a cross-sectional study

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Loes TCM Wouters¹, Dorien LM Zwart¹, Daphne CA Erkelens¹, Esther De Groot¹, Maarten van Smeden², Arno W Hoes³, Roger AMJ Damoiseaux¹, Frans H Rutten¹

- Dept. General Practice, Julius Centre for Health Sciences and Primary Care, University Medical Centre, Utrecht University, Utrecht, the Netherlands
- 2. Dept. Epidemiology, Julius Centre for Health Sciences and Primary Care, University Medical Centre, Utrecht University, Utrecht, the Netherlands
- 3. Dean University Medical Centre, Utrecht University, Utrecht, the Netherlands

All authors take responsibility for all aspects of the reliability and freedom from bias of the data presented and their discussed interpretation

Corresponding author: L.T. Wouters, MD, Dept. General Practice, Julius Centre for Health Sciences and Primary Care, University Medical Centre Utrecht, Utrecht University

STR 6.131, PO Box 85500, 3508 GA Utrecht, The Netherlands

Phone number: +31 (0)88 75 51470

Email: L.T.C.Wouters-2@umcutrecht.nl

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Transparency declaration

The lead author (the manuscript's guarantor) affirms that the manuscript is an honest, accurate, and transparent account of the study being reported; that no important aspects of the study have been omitted; and that any discrepancies from the study as planned (and, if relevant, registered) have been explained.

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Objectives To identify clinical variables that are associated with the diagnosis acute coronary syndrome (ACS) in women and men with chest discomfort who contact out-of-hours primary care (OHS-PC) by telephone, and to explore whether there are indications whether these variables differ among women and men.

Design Cross-sectional study in which we compared patient and call characteristics of triage call recordings between women with and without ACS, and men with and without ACS.

Setting Nine OHS-PC in the Netherlands.

Participants 993 women and 802 men who called OHS-PC for acute chest discomfort (pain, pressure, tightness, or discomfort) between 2014 and 2016.

Primary outcome measure Diagnosis of ACS retrieved from the patient's medical record in general practice, including hospital specialists' discharge letters.

Results Among 1,795 patients (mean age 58.8 (SD 19.5) years, 55.3% women), 15.0% of men and 8.6% of women had an ACS. In both sexes, retrosternal chest pain was associated with ACS (women with ACS vs. without 62.3% vs. 40.3%, p=0.002, men with ACS vs. without 52.5% vs. 39.7%, p=0.032, gender interaction p=0.323), as was pressing/heavy/tightening pain (women 78.6% vs. 61.5%, p=0.011, men 82.1% vs. 57.4%, p=<0.001, gender interaction p=0.368) and radiation to the arm (women 75.6% vs. 45.9%, p<0.001, men 56.0% vs. 34.8%, p<0.001, gender interaction p=0.339). Results indicate that only in women, severe pain (65.4% vs. 38.1%, p=0.006, gender interaction p=0.007) and radiation to jaw (50.0% vs. 22.9%, p=0.007, gender interaction p=0.015) were associated with ACS. Ambulances were dispatched equally in women (72.9%) and men with ACS (70.0%).

Conclusion Our results indicate there were more similarities than differences in symptoms associated with the diagnosis ACS for women and men. Important exceptions were severity and radiation of pain in women. Whether these differences have an impact on predicting ACS needs to be further investigated with multivariable analyses.

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Strengths and limitations of this study

- We could analyse the very initial conversation with symptom presentation of a large sample of
 patients calling for acute chest discomfort, without the risk of hindsight bias of the researcher or
 recall bias of the patient.
- We analysed clinical variables associated with ACS in gender subgroup analyses and across gender with statistical interaction terms.
- Results are generalizable to comparable primary care settings in the United Kingdom and Scandinavian Countries, and our results may even be generalizable to emergency medical service (EMS, '112' or '911') settings.
- For the purpose of improving telephone triage interviewing, prediction rule development with multivariable regression analysis is needed.

Adequate triage and early diagnosis is key in patients with acute chest discomfort because they might have an acute coronary syndrome (ACS) for which lifesaving early interventions are available. ACS is an umbrella term including ST-elevated myocardial infarction (STEMI), non-ST-elevated myocardial infarction (NSTEMI) and unstable angina pectoris (UAP). ¹ For the diagnosis of ACS an abnormal electrocardiogram (ST and/or T wave abnormalities) and/or elevated blood levels of troponin I or T are needed. ACS may than be further subdivided into ST-elevation myocardial infarction (STEMI) and non-ST-elevation myocardial infarction (NSTEMI) if the troponin levels are elevated.¹ If troponin levels are not elevated (or increased over time), it is called unstable angina pectoris (UAP). ¹ Increased preventive measures and development of (timely) effective therapeutic interventions ('time is muscle') have resulted in improved outcomes and prognosis in ACS. ² Telephone triage of patients with chest discomfort, as done in out-of-hours service primary care (OHS-PC) and emergency medical services (EMS or ambulance dispatch centres) is, however, challenging because it is difficult to differentiate ACS from other causes of chest discomfort based on symptoms only. ³, ⁴ Importantly, the majority of patients with chest discomfort (80%) in the Netherlands first approach the general practitioner (GP) or OHS-PC, and 20% directly calls the ambulance (112) or are self-referrals to the emergency department (ED). ⁵

Previous hospital-based studies reported a delayed recognition of ACS in women compared to men. ^{6,7} It was suggested that this delayed recognition was related to a less specific presentation in women. ^{8,9} This caused an ongoing debate on whether women with ACS compared to men present with less specific symptoms, and how this affects diagnosis, but also treatment, and prognosis. ^{10,11} A recent meta-analysis of 27 studies showed that women with ACS compared to men with ACS had higher odds of presenting with pain between the shoulder blades (OR 2.15, 95% CI 1.95-2.37), nausea or vomiting (OR 1.64, 95% CI 1.48-1.82) and shortness of breath (OR 1.34 (95% CI 1.21-1.48). ¹² Women with ACS had lower odds of sweating (OR 0.84, 95% CI 0.76-0.94) and presenting with chest pain (OR 0.70, 95% CI 0.63-0.78), but in both sexes chest pain remained the most common symptom (pooled prevalence men 79% and women 74%). ¹² Importantly, researchers suggested standardization in methods of symptoms assessment is needed, because of the difficulties to formulate any definitive statements about symptom presentation, as studies assessed symptoms in different ways (questionnaires or abstracting from medical records). ^{9, 13-15} Abstracting symptom presentation from medical records may dilute symptom presentation, as they are translated by the clinician in medical terminology. ¹⁶ Moreover, many studies suffer from recall or hindsight bias of both patient and researcher as they know the outcome (ACS).

For the clinician or telephone triage nurse it is crucial to differentiate ACS from other causes of chest discomfort. For that, studies are needed that include female and male patients presenting with chest discomfort, in which women and men who turn out to have ACS are compared to those who do not have ACS. Such studies are scarce. In a study performed among patients with chest discomfort seen at the ED in the USA (77 women and 244 men with ACS, and 195 women and 240 men without ACS were compared. ¹³ Women with ACS more often reported arm pain than women without ACS (47% vs. 32%, p=0.021), while men with ACS reported pressing feeling (63% vs. 54%, p=0.035) and chest pain (72% vs 60%, p=0.005) more often than men without ACS. ¹³ In a recent Dutch OHS-PC study among 23 women and 34 men with ACS, and 253 women and 208 men without ACS, symptoms associated with ACS in women and men seemed quite similar and the authors conclude that discriminating ACS in patients with chest discomfort who contacted primary care OHS is difficult in both women and men. ¹⁷

We aimed to identify clinical variables that are associated with the diagnosis ACS in women and men with chest discomfort who contact out-of-hours primary care (OHS-PC) by telephone, and to explore whether there are indications these variables differ among women and men. For analyses we used the very initial symptom presentation as available from the recorded telephone triage conversations.

Methods

We performed a cross-sectional diagnostic factor study with a random sample of 1,795 OHS-PC calls for chest discomfort (chest pain, pressure, tightness, or discomfort) between 2014 and 2016. ^{18, 19} We first selected calls on the basis of International Code for Primary Care (ICPC; a WHO world-wide code system for primary care) codes (K01, K02, K03, K24, K74, K75, K76, K77, K93, L04, P74, R02, R98, appendix-table 1) and keywords thoracic pain, chest pain, myocardial infarction, heart attack and their common abbreviations mentioned by the triage nurse in the electronic medical file (EMF) at the OHS-PC. ^{20, 21} General practitioners who work at the OHS-PC assign the ICPC codes to the call. We combined ICPC-codes and keywords to achieve a sample with a broad variety of symptoms to capture the entire domain of patients suspected of ACS. We listed all available calls of these patients and assigned random numbers with the Random Number Generator (RAND) function in Microsoft Excel to retrieve a random sample. Calls were excluded before re-listening when the patients' age was below 18 years or when the patient did not live in the surrounding area of the OHS-PCs (because then we could not retrieve a diagnosis from the general practitioner of these patients). Calls were excluded during re-listening when

We re-listened the telephone triage recordings to collect information about patient and conversation characteristics, on symptom presentation, medical history, urgency allocation, and involvement of a supervising general practitioner (GP) in the triage conversation. Gender considered the self-identified gender of the patient. Call duration and age were retrieved from the electronic EMF of the OHS-PC. Nine OHS-PC in the Netherlands participated, serving a total population of 1.5 million people. The diagnosis was made after the phone call, which was in the case of ACS nearly always done by the cardiologist (97.1%) in the hospital based on (i) symptom presentation, (i) levels of (high-sensitivity) troponin and (iii) electrocardiography results. The final diagnoses were provided by the patients' GP, based on the electronic medical file including ED and cardiologist discharge letters, and also the notes from the OHS-PC. We used medical information up to 30-days following the contact with the OHS-PC, to allow us to include diagnoses of ACS that was initially missed because the patient was not referred to the cardiologist the same day of the OHS-PC contact. In none of the patients in the study we had evidence of a missed diagnosis of ACS.

Context

In the Netherlands, OHS-PC covers primary care during 73% of the week hours, and the initial contact is by telephone. In most OHS-PC and EMS, the 'Netherlands Triage Standard' (NTS) is used as a decision support to classify the urgency of the patients' conditions. ²² Based on the patient's symptom presentation, the triage nurse needs to choose the most appropriate complaint out of 56 'main complaints'. Each NTS 'main complaint' incorporates a decision tree with hierarchically ordered questions, which are similar for men and women. Triage nurses fill out the caller's responses in the semi-automatic NTS system, which then generates urgency allocations linked to a timeframe within which the patient should be seen by a physician or ambulance personnel (U0 (reanimation) to U5 (self-care advice) appendix-table 2). The triage nurse can overrule this recommendation and up- or downscale the urgency allocation, often after consulting the supervising GP. ²³ A recent validation study showed that the diagnostic accuracy of the NTS for patients with chest discomfort is poor (sensitivity 0.73 (95% CI 0.68-0.78) and specificity 0.43 (95% CI 0.40-0.45)), as calculated on the outcome ACS or

other life-threatening events (LTEs). ²⁴ All telephone calls to the OHS-PC are routinely recorded and archived for five years for training and quality control purposes.

Data analyses

We compared patient and call characteristics between women with and without ACS, and men with and without ACS. For comparison of dichotomous variables we used the Chi² test or Fisher exact test and for continuous variables the independent sample t-test or Mann-Whitney U test. We performed interaction analysis across gender separately for each clinical variable with logistic regression analyses, to explore whether there are indications that these variables are differently associated with the diagnosis ACS among men and women. We analysed the association between urgency allocation and the final diagnosis ACS (alone or including other LTEs) with the Chi² test. We considered pulmonary embolism, thoracic aortic dissection and acute abdominal aneurysm as LTEs; patients with LTEs as well as those with ACS should receive an U1-level urgency.

Data analysis was performed using SPSS, IBM version 25.

Patient and public involvement

No patients were involved in setting the research question or the outcome measures, or in developing plans for design, however, they were involved in the implementation of the study. In addition, they were asked to advise on interpretation and writing up of results. Results will be shared and discussed with the national patient community of cardiovascular diseases ('Harteraad').

Ethics

The study (National Trial Register identification number: NTR7331) was approved by the Medical Ethics Committee Utrecht, the Netherlands (reference number WAG/mb/16/003208) and complied with the Declaration of Helsinki. A waiver of informed consent was given because our study had minimal risk to subjects and could otherwise not be carried out logistically. Personal and research data were handled and stored according to the European General Data Protection Regulation.

Results

Among the 1,795 callers with chest discomfort (mean age 58.8 (SD 19.5) years, 55.3% women), 8.6% of women and 15.0% of men had an ACS. In women with an ACS, 18.8% had a STEMI, 48.2% a NSTEMI,

A total of 22 (2.2%) women and 23 men (2.9%) had another LTE than ACS (e.g. pulmonary embolism, thoracic aortic dissection, acute abdominal aneurysm, acute heart failure).

Patient and call characteristics

Men and women with ACS were older than those without ACS (mean age of women with ACS 73.6 vs. without ACS 57.8 years, p<0.001, men 67.2 vs. 56.9 years, p<0.001, gender interaction p=0.094), and the mean duration of the telephone calls was shorter (women 6:47 vs. 7:47 minutes, p=0.021, men 6:31 vs. 7:33 minutes, p=0.004, gender interaction p=0.803) (table 1). The GP was consulted for supervision by the triage nurse in the majority of cases (52.2% in women and 55.5% in men). However, in women with ACS, the GP was less often consulted than in women without ACS (41.2% vs. 53.2%, p=0.034), but in men such a difference was not observed (53.3% vs. 55.9%, p=0.607, gender interaction p=0.208). In around half of the calls, someone else called initially on behalf of the patient (49.5% in women vs. 54.7% in men). In cases with ACS, for both sexes more often someone else called than in those without ACS (in women 69.4% vs. 47.7%, p<0.001, in men 65.8% vs. 52.8%, p=0.008, gender interaction p=0.251). In men with ACS, most often their female partner (53.3%) called, while in women with ACS, it was either their male partner (17.6%), their daughter (20.0%) or a nurse (17.6%) (appendix-table 3). Callers expressed concerns in nearly all calls, also in those without an ACS; women with ACS vs. women without ACS 97.3% vs. 88.9% (p=0.109), and men with ACS vs. men without ACS 96.3% vs. 86.5% (p=0.041) (gender interaction p=0.935).

Both women and men with ACS had more often a history of coronary artery disease (women 42.5% vs. 25.1%, p=0.017, and men 57.0% vs. 38.4%, p=0.002, gender interaction p=0.927), but women with ACS had more often a history of diabetes (41.4% vs. 14.6%, p<0.001, gender interaction p=0.079).

Symptom presentation

Chest pain was the most common complaint, both in patients with and without an ACS (women with ACS and without 98.8% and 93.1%, p=0.055, in men 92.4% and 94.5%, p=0.364, gender interaction p=0.048). Retrosternal located chest pain was more common in women and men with ACS than in those without ACS (women 62.3% vs. 40.3%, p=0.002, and men 52.5% vs. 39.7%, p=0.032, gender interaction p=0.323). Also, radiation of pain to the arms was associated with ACS in both sexes (women with vs.

without ACS 75.6% vs. 45.9%, p<0.001, and men 56.0% vs. 34.8%, p<0.001, gender interaction p=0.339), as was pressing/heavy/tightening chest pain (women with vs. without ACS 78.6% vs. 61.5%, p=0.011 and men 82.1% vs. 57.4%, p=<0.001, gender interaction p=0.368). Only in women radiation to the jaw (50.0% vs. 22.9%, p=0.007, gender interaction p=0.015) and severe pain (8 or more on a Numeric Rating Scale 0-10) was associated with ACS (65.4% vs. 38.1, p=0.006, gender interaction p=0.007). Only in men, stabbing pain was very rare in those with ACS (8.4% vs. 26.5%, p<0.001, gender interaction p=0.141). Of the autonomous nervous system (ANS)-related symptoms, nausea/vomiting and dizziness/near fainting were not associated with ACS in either sex, with the exception of a pale or ashen face that was in women (55.6% vs. 35.5%, p=0.019, gender interaction p=0.545), and sweating in men (52.4% vs. 38.1%, p=0.015, gender interaction p=0.418). Recognition of symptoms being similar to a previous cardiac event was associated with ACS in men (52.9% vs. 32.1%, p=0.004), but not clearly for ACS in women (32.5% vs. 21.4%, p=0.108, gender interaction p=0.532).

Subgroup analyses in 56 women and 58 men with diabetes showed that both women (85.7% vs. 58.3%, p<0.001) and men with diabetes (67.2% vs. 51.5%, p=0.033, gender interaction p=0.119) more often had shortness of breath than those without diabetes, but as often chest discomfort (women 90.9% vs. 95.0%, p=0.193, men 89.2% vs 94.1%, p=0.162, gender interaction p=0.969). Shortness of breath in patients with diabetes was not related to ACS diagnosis (women 81.8% vs. 86.7%, p=0.680, men 75.0% vs. 66.0%, p=0.615, gender interaction p=0.520).

Diagnoses

Of the 205 patients with an ACS (85 women, 120 men), 55 (26.8%) patients had a STEMI (women 18.8%, men 32.5%), 85 (41.5%) a NSTEMI (women 48.2%, men 36.7%), 50 (24.4%) unstable angina pectoris (UAP) (women 20.0%, men 27.5%) and 15 (7.3%) unspecified ACS (women 13.0%, men 3.3%), the latter also including two sudden cardiac deaths in women and one in men (Table 2). In nearly all cases (97.1%) the ACS diagnosis was made by a cardiologist based on symptom presentation, troponin levels and electrocardiography. Three patients died before arrival of the ambulance (they were classified as acute cardiac death) and one patient died after resuscitation at the ED. Two patients were classified as ACS by the GP; they were not referred to the hospital because of short life expectancy due to cancer. There were 45 patients with other LTEs (2.5%) and the majority of patients had non-urgent medical conditions (86.1%). The most common non-urgent diagnoses in both sexes were (i) non-urgent cardiovascular diseases such as stable angina pectoris, stable heart failure and arrhythmias (19.5% of all

female callers with chest discomfort vs. 21.2% of male callers, p=0.384) and (ii) non-cardiac unspecified chest pain (women 16.4% vs. 19.8% men, p=0.061). Women more often than men were diagnosed with musculoskeletal problems (women 20.8% vs. men 14.1%, p=0.001) and psychogenic conditions (women 14.0% vs. men 8.4%, p<0.001). Of the patients who were diagnosed with a non-ACS diagnoses, 45.4% were classified by a cardiologist, 5.5% by another hospital specialist (e.g. pulmonologist or internal medicine specialist) and the remaining patients were diagnosed by a GP.

Urgencies

Women and men with chest discomfort were equally sent an ambulance (43.6% vs. 46.6%, p=0.200). This was also in women and men who had an ACS (72.9% vs. 70.0%, p=0.647), and in those with either ACS or other LTEs (66.4% vs. 67.1%, p=0.897). See table 3.

Discussion

For both sexes, retrosternal pain, pain described as pressing, heavy or tightening, and radiation to the arm were associated with ACS in patients who contacted the OHS-PC for chest discomfort. Radiation to the jaw and severe pain were related to ACS in women. Our results indicate there were more similarities than differences in symptoms associated with the diagnosis ACS for women and men. However, whether these differences have an impact on predicting ACS needs to be further investigated. Women and men with chest discomfort as also those with ACS were equally often sent an ambulance.

Our finding that radiation of pain to the arm and retrosternal ('mid') chest pain were associated with the diagnosis ACS in both sexes was also reported in a study among 2,475 patients with acute chest pain in a multicentre ED-study. ²⁵ Another ED-study among 1,334 patients with ACS showed that regardless of ethnics status the most common presenting symptom was retrosternal pain/discomfort of any intensity. ²⁶ The aforementioned US study in the ED-setting reported that radiation to the arm was associated with ACS in women but not for men, and chest pressure was associated with ACS for men but not in women. ¹³ The only previously published OHS-PC study reported the opposite; radiation to the arm was associated with ACS in men, but not in women. ¹⁷ In our study, women with ACS had more often a history of diabetes and were older than men with ACS,

which is in line with other studies. ^{8, 12} Some studies claim that patients with diabetes more often have atypical symptoms of ACS, however a review of eight studies concluded the evidence of these studies was conflicting. ²⁷ We showed that both women and men with diabetes had more often shortness of

breath than those without diabetes, but shortness of breath in patients with diabetes was not associated with ACS. Regarding dispatching the ambulance, the aforementioned OHS-PC study and two EMS studies showed, similar to our findings, that there was no difference in dispatch priorities between men and women with ACS. ^{4, 17, 28} This is in contrast with studies that show delay in hospital presentation of women with ACS. ^{9, 29}

We need to realize that focusing on gender differences may blur the large overlap in symptoms in women and men. Moreover, comparing *selectively* women with ACS to men with ACS as many previous studies did, is clinically irrelevant. ^{9, 14, 30} Clinicians, including GPs, and triage nurses need to know whether and how women with ACS differ from women without ACS, with the same question for men. Nevertheless, even guidelines stick to comparing those with established disease, and express the view that women with ACS more likely present with less specific symptoms than men with ACS. ^{1, 31} Unfortunately, public awareness campaigns follow this reasoning, and over-emphasize sex differences in women awareness campaigns ('Go Red for Women' in the United States and 'Invisible me' in Australia). ^{7, 30, 32} Unbalanced attention to symptom differences, neglecting the much larger overlap may even introduce new blind spots in recognizing ACS in women. ³³

A likely reason behind the predominant message that women present with other ACS symptoms than men is the difference in pathophysiology of coronary artery disease. Women compared to men more often have elongated plaques, located on bifurcations in epicardial coronaries, coronary spasm, microvascular dysfunction, and spontaneous coronary dissection. ^{32, 34} These pathophysiological differences have an effect on interventional treatment and prognosis. ^{2, 34} However, these differences do not necessarily imply an effect on symptom presentation because the pain pathway is equal in women and men: i.e. triggered by myocardial ischemia. ^{35, 36} A supply and demand mismatch of the myocardial oxygen consumption triggers sensory nerve endings in the myocardium and cause ischemia symptoms, and this is irrespective of the fact whether the ischemia is caused by a plaque rupture in an epicardial artery or spasm, or any other cause. ³⁶ The sex differences in pathophysiology of ACS do therefore not support the belief in differences in ACS symptoms between women and men.

Another reason behind the belief of 'vague' symptom presentation in women with ACS might be that they seem to present a larger number of symptoms than men with ACS, and this may be interpreted as 'vague' by physicians. ^{15, 37} Presentation of multiple symptoms may influence the prompt recognition of

Interestingly, in the majority of calls in our study someone else than the patient called the OHS-PC (women with ACS 69.4%, without ACS 47.7%, and men with ACS 65.8%, and without ACS 52.8%). This was also highlighted in an Australian study among 1,681 patients with an acute myocardial infarction; in 90.5% of the women with AMI someone else called on behalf of the patient and in 87.8% of the men with AMI. ⁴⁰ According to protocol in OHS-PC, triage nurses ask the patient to the phone, this to prevent loss of (paralinguistic) information from the patient him/herself. In our study, in about 50% of the conversations the patient took over the phone call.

Strengths and limitations

A major strength is that we could analyse the very initial conversation with symptom presentation of a large sample of patients calling the OHS-PC because of chest discomfort. We analysed the conversations without knowledge of the eventual diagnosis and have prevented risk of hindsight bias of the researcher or recall bias of the patient. Another strength is that we performed gender subgroup analyses combined with interaction analyses across gender, to investigate whether there are indications that ACS related symptoms differ among men and women. Furthermore, our results are generalizable to comparable primary care settings, e.g. UK and Scandinavian countries, and possibly some other European countries.³ Our results may even be generalizable to EMS settings, since the prior probability of having an ACS is comparable in EMS setting as in OHS-PC settings. ^{4 41}

As the intention of our analysis was to describe whether symptoms were different in patients with ACS from patients without ACS in women and men separately, none of our results can be used to adjust interview questions for the triage nurses. For that purpose, prediction rule development with multivariable analyses is necessary. Also, only with multivariable analysis it can be truly investigated

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whether the potential differences are clinically relevant in prediction of ACS. Another limitation is missing values on some clinical variables, a phenomenon common in routine care data, and therefore the results should be interpreted with caution.

Conclusions

Our results indicate there were more similarities than differences in symptoms associated with the diagnosis ACS for women and men. Important exceptions were severity and radiation of pain in women. However, whether these differences have an impact on predicting ACS needs to be further investigated yses. with multivariable analyses.

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Data sharing statement

The data can be made available for researchers whose proposed use of the data has been approved at request of the corresponding author, with a signed data access agreement.

Author contribution

FR and DZ are the lead investigators who conceived the research idea and methodology. Funding acquisition was done by FR, DZ and RD. LW and DC conducted data acquisition. LW performed the analyses and wrote the first draft of the manuscript. She was supervised by FR and DZ, who critically revised the manuscript. DC, EdG, AH and RD contributed with and approved the final version of the manuscript. MvS was involved in adjusting the analyses and revising of the manuscript.

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Figure 1. Flowchart study population.

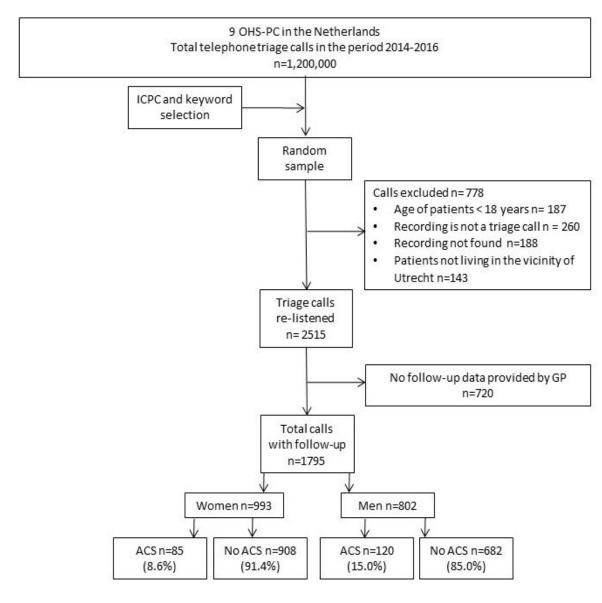


Table 1. Baseline characteristics of 993 (55.3%) women and 802 (44.7%) men patients contacting the OHS-PC with open pressure, tightness, or discomfort)

Characteristics	993 women			Difference (95% CI)	802 men		© 122 © 96 Difference (95% CI)		P-value
8 9	ACS	No ACS	p-value		ACS		r ja Ug p-Majlue	-	interaction gender
10 11	N = 85 (8.6%)	N = 908 (91.4%)			N = 120 (15.0%)	N = 682 (85.0%)	iune 20 Era:		
13 Mean age in years (SD)	73.6 (±14.1)	57.8 (±20.1)	<0.001	15.8 (11.4-20.2)	67.2 (±13.0)	56.9 (±19.2)	202001	10.3 (6.7-13.9)	0.094
14		Uh					Dowr Ishog		
15 Call characteristics		/ /					Žige ≸		
16							loa Sc		
17 Mean call duration in min (SD) 18	6:47 (±5:16)	7:47 (±3:48)	0.021	1.00 (0.19-1.80)	6:31 (±3:13)	7:33 (±3:42)	Downloa∰ed i ushogesch∞ol	1.02 (0.37-1.68)	0.803
19 Mean patient's introduction in min (SD) 20	0:14 (± 0:08)	0:19 (±0:13)	<0.001	0.05 (0.02-0.09)	0:16 (±0:11)	0:19 (±0:14)	¬. =~	0.03 (0.00-0.06)	0.042
21 Triage nurse consulted the GP	35 (41.2)	483 (53.2)	0.034	12.0 (0.3-22.9)	64 (53.3)	381 (55.9)	ing. Alt. 0. \$ 37	2.5 (-7.3-12.6)	0.208
22 23 Someone else called on behalf of patient 24	59 (69.4)	433 (47.7)	<0.001	21.7 (10.1-31.6)	79 (65.8)	360 (52.8)	0. 0 .8	13.0 (3.0-22.2)	0.251
25 The patient or person who called	36 (97.3)	409 (88.9)	0.109	8.4 (-5.0-12.6)	52 (96.3)	321 (86.5)	0.041	9.8 (-0.9-14.8)	0.935
26 27 expressed concerned (n=922)					100		o. and sin		
29 Chest pain (n=1739)	81 (98.8)	817 (93.1)	0.055	5.7 (-0.8-8.0)	110 (92.4)	624 (94.5)	0.364 On	2.1 (-2.3-9.0)	0.048
30 3 Severe Pain (>7 on a scale 0-10) (n=753)	17 (65.4)	147 (38.1)	0.006	27.3 (5.7-44.7)	13 (23.6)	87 (30.4)	n પ્રાne 13, 20 ા	6.8 (-7.9-18.3)	0.007
32 3 ₃ Duration) 13, 1		
34 35 > 15 min (n=1501)	66 (100)	729 (95.8)	0.102	4.2 (-2.8-6.0)	99 (97.1)	541 (94.6)	7. 20292 0. 292	2.5 (-3.8-5.6)	0.998
36 37 < 12 hrs (n=1563)	60 (85.7)	575 (72.7)	0.018	13.0 (1.7-20.6)	88 (81.5)	431 (72.6)	_	8.9 (-0.6-16.5)	0.490
38 39 Location (n=1267)*							: D@partment GE		
40							ä		

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							جَ ي		
Retrosternal	33 (62.3)	263 (40.3)	0.002	22.0 (7.1-35.2)	42 (52.5)	191 (39.7)	, 0. 65 2	12.8 (0.5-24.8)	0.323
Right or left side thorax	14 (26.4)	260 (39.8)	0.054	13.4 (-1.2-24.8)	26 (32.5)	211 (43.9)	2020-0#2406 (0.0.0.0.1 ht, including for	11.4 (-1.0-22.2)	0.757
Type of pain (n=1334)**							9 9		
Pressing/heavy/tightening	44 (78.6)	423 (61.5)	0.011	17.1 (3.2-27.2)	78 (82.1)	284 (57.4)	1 250un	24.7 (14.3-32.9)	0.368
1 Stabbing 2	8 (14.3)	155 (22.5)	0.152	8.2 (-4.6-16.4)	8 (8.4)	131 (26.5)		18.1 (9.2-24.1)	0.141
Radiation of chest pain (n=1391) ***		04					ા e ² 021. Dow¤toaded Erasmushogeschool Hated to text and data		
5 Any location 6	61 (85.9)	485 (68.6)	0.002	17.3 (6.0-25.0)	65 (63.7)	292 (57.1)	wi્રીoa ogesc xt and	6.6 (-4.5-16.7)	0.071
7 8 Arm	31 (75.6)	188 (45.9)	<0.001	29.7 (12.8-42.2)	47 (56.0)	117 (34.8)	data	21.1 (8.7-32.9)	0.339
9 Between the shoulder blades	11 (52.4)	159 (41.7)	0.336	10.7 (-12.0-32.5)	14 (27.5)	91 (29.4)	noga 0.281 mining,	1.9 (-13.4-14.3)	0.352
1 2 ^{Jaws}	10 (50.0)	66 (22.9)	0.007	27.1 (4.3-49.7)	1 (2.6)	28 (11.3)	ttp 88 0. 8/br	8.7 (-4.6-14.1)	0.015
3 4 Shortness of breath (n=1365)	47 (72.3)	455 (66.3)	0.328	6.0 (-7.2-16.7)	53 (60.9)	329 (62.4)	0.288	1.5 (-9.5-13.3)	0.355
5 Symptoms similar to previous cardiac	13 (32.5)	79 (21.4)	0.108	11.1 (-3.0-28.4)	27 (52.9)	92 (32.1)	an 0.0004	20.8 (5.4-35.8)	0.532
7 <mark>event (n=748)</mark> 8							0.@mj.com/ and simi		
9 ANS-related symptoms						7/.	욕으		
1 Sweating (n=1130) 2	28 (49.1)	234 (42.5)	0.340	6.6 (-7.4-20.7)	43 (52.4)	170 (38.5)	Juge 1	13.9 (16.7-25.8)	0.418
3 Nausea or vomiting (n=808) 4	16 (44.4)	240 (56.2)	0.173	11.8 (-6.2-28.6)	27 (45.8)	122 (42.7)	13,602 0.602 hogies.	3.1 (-11.1-17.7)	0.186
5 Pallor or ashen skin (n=652) 6	20 (55.6)	110 (35.5)	0.019	20.1 (1.9-37.0)	28 (53.8)	103 (40.6)	0. 01	13.2 (-2.3-28.3)	0.545
7 Dizziness or near fainting n=1599) 8	13 (18.6)	197 (24.4)	0.277	5.8 (-6.0-14.3)	17 (15.9)	127 (20.7)	0. 23 19	4.8 (-4.3-11.9)	0.963
9 Medical history							ment (
1 2 3		1		1	1	1	025 at Degartment GEZ-LTA	I	21

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3 4	CV disease or risk factors (n=1461)	54 (78.3)	407 (57.5)	0.001	20.8 (8.4-30.2)	75 (72.1)	359 (61.9)	7, 0. 22 6	10.2 (-0.3-19.3)	0.179	
5	Coronary artery disease (n=966)	17 (42.5)	114 (25.1)	0.017	17.4 (1.7-34.3)	45 (57.0)	151 (38.4)	n-2820-0824063n ;	18.6 (5.9-30.5)	0.927	
7	Hypertension (n=756)	ion (n=756) 22 (71.0) 142 (35.1) <0.001 35.9 (16.0-50.7) 17 (47.2) 94 (33.0) 7 0.091 14.2 (-3.0-32.1) 0.094									
9 10	Diabetes mellitus (n=745)	12 (41.4)	56 (14.6)	<0.001	26.8 (9.0-46.5)	12 (28.6)	59 (20.3)	25QJun o. uses re	8.3 (-5.1-25.1)	0.079	
11 12	Hypercholesterolemia (n=679)	8 (36.4)	81 (23.2)	0.161	13.1 (-5.8-36.4)	19 (45.2)	62 (23.3)	ne∰02 Ecasr elated	21.9 (5.8-38.6)	0.527	
13 14	18 *P-value comparing retrosternal or left/right side thorax vs. others locations of pain together (restrosternal, left/right side thorax, back/shoulder, epigas (right)										
15 16	** Pressing/ heavy/tightening pain vs. oth	er types of pain	stabbing, burning,	cramping, tea	aring). Stabbing pain: s	stabbing vs. other	types of pain (press	ặc g sin g∯hg a y /tig	htening, burning, cram	ping, tearing)	
17 18	ዕር አል ር አልር 2 *** P-value comparing radiation arm or back/shoulder or laws vs. no radiation										
19 20	NRS: Numeric Rating Scale							rom h			
21 22	ANS-related symptoms: Autonomous nerv	ous system relat	ed symptoms					ig, Al			
23 24	CV disease or risk factors; a history of prev	rious coronary ar	tery disease, heart	failure, strok	e, cardiac arrhythmia,	hypertension, and	l/or diabetes (patie	— • • • • • • • • • • • • • • • • • • •			
25								ng,			
26	Coronary artery disease: History of prior N	II, PCI, CABG, sta	ble or unstable ang	gina pectoris	(patient reported)			<mark>bmj</mark> and			
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28 29								om/ on June 13, 202 imilar technologies			
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	202 (55 204)	200 (44 70/)	
	Women n= 993 (55.3%)	Men n=802 (44.7%)	p-value
Acute coronary syndrome	85 (8.6)	120 (15.0)	<0.001
STEMI	16 (18.8)	39 (32.5)	0.037
NSTEMI	41 (48.2)	44 (36.7)	0.114
UAP	17 (20.0)	33 (27.5)	0.250
Non-classified ACS	11 (13.0)	4 (3.3)	0.013
Life threatening events (LTE's)	22 (2.2)	23 (2.9)	0.448
Pulmonary embolism	6 (27.3)	7 (30.4)	0.815
Thoracic aortic dissection	4 (18.2)	2 (8.7)	0.349
Acute abdominal aneurysm	3 (13.6)	2 (8.7)	0.598
Other*	9 (40.9)	12 (52.2)	0.449
Non-urgent cardiovascular	194 (19.5)	170 (21.2)	0.384
diseases**			
Non-cardiac chest pain, not further	163 (16.4)	159 (19.8)	0.061
specified ***		0,	
Musculoskeletal pain	199 (20.0)	113 (14.1)	0.001
Psychogenic disorders	139 (14.0)	67 (8.4)	<0.001
Gastrointestinal tract disorders	76 (7.7)	62 (7.7)	0.951
Respiratory tract disorders	52 (5.2)	45 (5.6)	0.727
Other non-urgent diagnoses***	63 (6.3)	43 (5.4)	0.380

^{*}Acute heart failure, stroke, severe COPD exacerbation, sepsis, coronary spasm probably caused by hypokalaemia, diabetic ketoacidosis, epileptic insult, bleeding from oesophageal varices, ovarian torsion, ventricular fibrillation.

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**** Amongst others: anaemia, malignancy, vasovagal collapse, side effects medication, dermatologic diseases



Women	ACS n=85 (8.6%)	No ACS n=908 (91.4%)	p-value 1*	p-value 2**
U1	62 (72.9)	371 (40.9)	<0.001	<0.001
U2	12 (14.1)	231 (25.4)	-	
U3-U5	11 (13.0)	306 (33.6)	_	
Men	ACS n=120 (15.0%)	No ACS n=682 (85.0%)		
U1	84 (70.0)	290 (42.5)	<0.001	<0.001
U2	19 (15.8)	142 (20.8)	-	
U3-U5	17 (14.2)	250 (36.7)	-	
Women	ACS or LTE n=107 (10.8%)	No ACS or LTE n=886 (89.2%)		
U1	71 (66.4)	362 (40.9)	<0.001	<0.001
U2	21 (19.6)	222 (25.1)		
U3-U5	15 (14.0)	302 (34.0)		
Men	ACS or LTE n=143 (17.8%)	No ACS or LTE n=659 (82.2%)		
U1	96 (67.1)	278 (42.2)	<0.001	<0.001
U2	24 (16.8)	137 (20.8)		
U3-U5	23 (16.1)	244 (37.0)	2/	

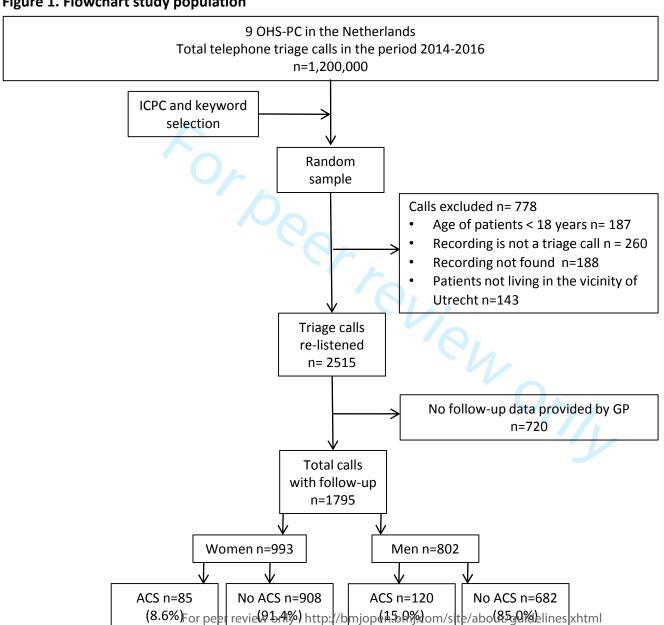
^{*} P-value 1: U1 vs. U2, U3, U4 and U5

^{**} P-value 2: U1,U2 vs. U3,U4,U5

^{***}LTE = life threatening event. Life threatening events consist of: ACS; pulmonary embolism; thoracic aortic dissection; acute heart failure; stroke; abdominal aortic aneurysm; severe COPD exacerbation; diabetic ketoacidosis; coronary spasm probably caused by hypokalaemia; epileptic insult; bleeding from oesophageal varices; ovarian torsion; ventricular fibrillation.

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Figure 1. Flowchart study population



Supplementary file

Appendix-Table 1: Overview of ICPC-codes used to select calls

ICPC code	Calls of patients with ACS	Calls of patients without ACS
	N=205 (%)	N=1,590 (%)
K01 Pain attributed to the heart	112 (54.6)	485 (30.5)
K02 Pressure/tightness attributed to the heart	39 (19.0)	184 (11.6)
K03 Other cardiovascular pain	1 (0.5)	6 (0.4)
K24 Fear of heart attack	1 (0.5)	8 (0.5)
K74 Angina pectoris	19 (9.3)	101 (6.4)
K75 Acute myocardial infarction	7 (3.4)	22 (1.4)
K76 Other/chronic ischaemic heart disease	0 (0.0)	2 (0.1)
K77 Heart failure	0 (0.0)	11 (0.7)
K93 Pulmonary embolism	0 (0.0)	15 (0.9)
L04 Chest discomfort	25 (12.2)	689 (43.9)
P74 Anxiety disorder	0 (0.0)	12 (0.8)
R02 Shortness of breath	1 (0.5)	36 (2.3)
R98 Hyperventilation	0 (0.0)	10 (0.6)

Appendix-Table 2: Urgency levels

Urgency level	Implication
U0	Reanimation
U1	Life-threatening, GP/ ambulance should arrive within 15 minutes
U2	Emergency, GP should arrive within 60 minutes
U3	Urgent, consultation by GP within three hours
U4	Routine, consultation by GP the same day
U5	Advice given by triage nurse

Appendix-Table 3: Relation of caller to patient in women and men with ACS

	Women	Men	P-value
	n= 85 (8.6%)	n = 120 (15.0%)	
Someone else calls on behalf of patient	59 (69.4)	79 (65.8)	0.590
Partner	16 (18.8)	64 (53.3)	<0.001
Son, daughter or other family member	20 (23.5)	8 (6.7)	0.001
Nurse	15 (17.6)	4 (3.3)	<0.001
Other (neighbour, friend, colleague)	8 (9.4)	3 (2.5)	0.030
Patient takes over the phone call on request	27 (45.8)	58 (73.4)	<0.001
of the triage nurse			

STROBE Statement—Checklist of items that should be included in reports of cross-sectional studies

	Item No	Recommendation	Page No
Title and abstract	1	(a) Indicate the study's design with a commonly used term in the	1,3
		title or the abstract	
		(b) Provide in the abstract an informative and balanced summary of	3
		what was done and what was found	
Introduction			
Background/rationale	2	Explain the scientific background and rationale for the investigation being reported	5,6
Objectives	3	State specific objectives, including any prespecified hypotheses	6
Methods			•
Study design	4	Present key elements of study design early in the paper	6
Setting	5	Describe the setting, locations, and relevant dates, including periods	6,7
S		of recruitment, exposure, follow-up, and data collection	
Participants	6	(a) Give the eligibility criteria, and the sources and methods of	6,7
•		selection of participants	
Variables	7	Clearly define all outcomes, exposures, predictors, potential	6,7
		confounders, and effect modifiers. Give diagnostic criteria, if	
		applicable	
Data sources/	8*	For each variable of interest, give sources of data and details of	6,7
measurement		methods of assessment (measurement). Describe comparability of	
		assessment methods if there is more than one group	
Bias	9	Describe any efforts to address potential sources of bias	8
Study size	10	Explain how the study size was arrived at	7
/Quantitative	11	Explain how quantitative variables were handled in the analyses. If	8
variables		applicable, describe which groupings were chosen and why	
Statistical methods	12	(a) Describe all statistical methods, including those used to control	8
		for confounding	
		(b) Describe any methods used to examine subgroups and	8
		interactions	
		(c) Explain how missing data were addressed	14
		(d) If applicable, describe analytical methods taking account of	
		sampling strategy	
		(g) Describe any sensitivity analyses	8
Results			•
Participants	13*	(a) Report numbers of individuals at each stage of study—eg	19 (figure 1)
1 articipants	15	numbers potentially eligible, examined for eligibility, confirmed	(ligare 1)
		eligible, included in the study, completing follow-up, and analysed	
		(b) Give reasons for non-participation at each stage	19 (figure 1)
		(c) Consider use of a flow diagram	19
Descriptive data	14*	(a) Give characteristics of study participants (eg demographic,	7
200011ptive data	17	clinical, social) and information on exposures and potential	,
		confounders	
		(b) Indicate number of participants with missing data for each	20,21 (table 1)
		variable of interest	20,21 (more 1)
Outcome data	15*	Report numbers of outcome events or summary measures	9
Outcome data	13	report numbers of outcome events of summary measures	

Main manulta	1.6	(a) Circa and directed actions and if applicable confounder	
Main results	16	(a) Give unadjusted estimates and, if applicable, confounder-	n.a.
		adjusted estimates and their precision (eg, 95% confidence interval).	
		Make clear which confounders were adjusted for and why they were	
		included	
		(b) Report category boundaries when continuous variables were	20,21 (table 1)
		categorized	
		(c) If relevant, consider translating estimates of relative risk into	NA
		absolute risk for a meaningful time period	
Other analyses	17	Report other analyses done—eg analyses of subgroups and	9,10 (subgroup
		interactions, and sensitivity analyses	diabetes)
Discussion			
Key results	18	Summarise key results with reference to study objectives	11
Limitations	19	Discuss limitations of the study, taking into account sources of	14
		potential bias or imprecision. Discuss both direction and magnitude	
		of any potential bias	
Interpretation	20	Give a cautious overall interpretation of results considering	13,14
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Funding	22	Give the source of funding and the role of the funders for the present	2
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^{*}Give information separately for exposed and unexposed groups.

Note: An Explanation and Elaboration article discusses each checklist item and gives methodological background and published examples of transparent reporting. The STROBE checklist is best used in conjunction with this article (freely available on the Web sites of PLoS Medicine at http://www.plosmedicine.org/, Annals of Internal Medicine at http://www.annals.org/, and Epidemiology at http://www.epidem.com/). Information on the STROBE Initiative is available at www.strobe-statement.org.

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Gender-stratified analyses of symptoms associated with acute coronary syndrome in telephone triage: a cross-sectional study

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Loes TCM Wouters¹, Dorien LM Zwart¹, Daphne CA Erkelens¹, Esther De Groot¹, Maarten van Smeden², Arno W Hoes³, Roger AMJ Damoiseaux¹, Frans H Rutten¹

- Dept. General Practice, Julius Centre for Health Sciences and Primary Care, University Medical Centre, Utrecht University, Utrecht, the Netherlands
- 2. Dept. Epidemiology, Julius Centre for Health Sciences and Primary Care, University Medical Centre, Utrecht University, Utrecht, the Netherlands
- 3. Dean University Medical Centre, Utrecht University, Utrecht, the Netherlands

All authors take responsibility for all aspects of the reliability and freedom from bias of the data presented and their discussed interpretation

Corresponding author: L.T. Wouters, MD, Dept. General Practice, Julius Centre for Health Sciences and Primary Care, University Medical Centre Utrecht, Utrecht University

STR 6.131, PO Box 85500, 3508 GA Utrecht, The Netherlands

Phone number: +31 (0)88 75 51470

Email: L.T.C.Wouters-2@umcutrecht.nl

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All authors have completed the <u>Unified Competing Interest form</u> and declare no support from any organization for the submitted work; no financial relationships with any organizations that might have an interest in the submitted work in the previous three years, no other relationships or activities that could appear to have influenced the submitted work.

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The lead author (the manuscript's guarantor) affirms that the manuscript is an honest, accurate, and transparent account of the study being reported; that no important aspects of the study have been omitted; and that any discrepancies from the study as planned (and, if relevant, registered) have been explained.

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Objectives To identify clinical variables that are associated with the diagnosis acute coronary syndrome (ACS) in women and men with chest discomfort who contact out-of-hours primary care (OHS-PC) by telephone, and to explore whether there are indications whether these variables differ among women and men.

Design Cross-sectional study in which we compared patient and call characteristics of triage call recordings between women with and without ACS, and men with and without ACS.

Setting Nine OHS-PC in the Netherlands.

Participants 993 women and 802 men who called OHS-PC for acute chest discomfort (pain, pressure, tightness, or discomfort) between 2014 and 2016.

Primary outcome measure Diagnosis of ACS retrieved from the patient's medical record in general practice, including hospital specialists' discharge letters.

Results Among 1,795 patients (mean age 58.8 (SD 19.5) years, 55.3% women), 15.0% of men and 8.6% of women had an ACS. In both sexes, retrosternal chest pain was associated with ACS (women with ACS vs. without 62.3% vs. 40.3%, p=0.002, men with ACS vs. without 52.5% vs. 39.7%, p=0.032, gender interaction p=0.323), as was pressing/heavy/tightening pain (women 78.6% vs. 61.5%, p=0.011, men 82.1% vs. 57.4%, p=<0.001, gender interaction p=0.368) and radiation to the arm (women 75.6% vs. 45.9%, p<0.001, men 56.0% vs. 34.8%, p<0.001, gender interaction p=0.339). Results indicate that only in women, severe pain (65.4% vs. 38.1%, p=0.006, gender interaction p=0.007) and radiation to jaw (50.0% vs. 22.9%, p=0.007, gender interaction p=0.015) were associated with ACS. Ambulances were dispatched equally in women (72.9%) and men with ACS (70.0%).

Conclusion Our results indicate there were more similarities than differences in symptoms associated with the diagnosis ACS for women and men. Important exceptions were pain severity and radiation of pain in women. Whether these differences have an impact on predicting ACS needs to be further investigated with multivariable analyses.

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Keywords (5): acute coronary syndrome, gender, primary care, telephone triage, chest discomfort

Strengths and limitations of this study

- We could analyse the very initial conversation with symptom presentation of a large sample of
 patients calling for acute chest discomfort, without the risk of hindsight bias of the researcher or
 recall bias of the patient.
- We analysed clinical variables associated with ACS in gender subgroup analyses and across gender with statistical interaction terms.
- Results are generalizable to comparable primary care settings in the United Kingdom and Scandinavian Countries, and our results may even be generalizable to emergency medical service (EMS, '112' or '911') settings.
- For the purpose of improving telephone triage interviewing, prediction rule development with multivariable regression analysis is needed.

Adequate triage and early diagnosis is key in patients with acute chest discomfort because they might have an acute coronary syndrome (ACS) for which lifesaving early interventions are available. ACS is an umbrella term including ST-elevated myocardial infarction (STEMI), non-ST-elevated myocardial infarction (NSTEMI) and unstable angina pectoris (UAP). ¹ For the diagnosis of ACS an abnormal electrocardiogram (ST and/or T wave abnormalities) and/or elevated blood levels of troponin I or T are needed. ACS may than be further subdivided into ST-elevation myocardial infarction (STEMI) and non-ST-elevation myocardial infarction (NSTEMI) if the troponin levels are elevated.¹ If troponin levels are not elevated (or increased over time), it is called unstable angina pectoris (UAP). ¹ Increased preventive measures and development of (timely) effective therapeutic interventions ('time is muscle') have resulted in improved outcomes and prognosis in ACS. ² Telephone triage of patients with chest discomfort, as done in out-of-hours service primary care (OHS-PC) and emergency medical services (EMS or ambulance dispatch centres) is, however, challenging because it is difficult to differentiate ACS from other causes of chest discomfort based on symptoms only. ³, ⁴ Importantly, the majority of patients with chest discomfort (80%) in the Netherlands first approach the general practitioner (GP) or OHS-PC, and 20% directly calls the ambulance (112) or are self-referrals to the emergency department (ED). ⁵

Previous hospital-based studies reported a delayed recognition of ACS in women compared to men. ^{6,7} It was suggested that this delayed recognition was related to a less specific presentation in women. ^{8,9} This caused an ongoing debate on whether women with ACS compared to men present with less specific symptoms, and how this affects diagnosis, but also treatment, and prognosis. ^{10,11} A recent meta-analysis of 27 studies showed that women with ACS compared to men with ACS had higher odds of presenting with pain between the shoulder blades (OR 2.15, 95% CI 1.95-2.37), nausea or vomiting (OR 1.64, 95% CI 1.48-1.82) and shortness of breath (OR 1.34 (95% CI 1.21-1.48). ¹² Women with ACS had lower odds of sweating (OR 0.84, 95% CI 0.76-0.94) and presenting with chest pain (OR 0.70, 95% CI 0.63-0.78), but in both sexes chest pain remained the most common symptom (pooled prevalence men 79% and women 74%). ¹² Importantly, researchers suggested standardization in methods of symptoms assessment is needed, because of the difficulties to formulate any definitive statements about symptom presentation, as studies assessed symptoms in different ways (questionnaires or abstracting from medical records). ^{9, 13-15} Abstracting symptom presentation from medical records may dilute symptom presentation, as they are translated by the clinician in medical terminology. ¹⁶ Moreover, many studies suffer from recall or hindsight bias of both patient and researcher as they know the outcome (ACS).

For the clinician or telephone triage nurse it is crucial to differentiate ACS from other causes of chest discomfort. For that, studies are needed that include female and male patients presenting with chest discomfort, in which women and men who turn out to have ACS are compared to those who do not have ACS. Such studies are scarce. In a study performed among patients with chest discomfort seen at the ED in the USA (77 women and 244 men with ACS, and 195 women and 240 men without ACS were compared. ¹³ Women with ACS more often reported arm pain than women without ACS (47% vs. 32%, p=0.021), while men with ACS reported pressing feeling (63% vs. 54%, p=0.035) and chest pain (72% vs 60%, p=0.005) more often than men without ACS. ¹³ In a recent Dutch OHS-PC study among 23 women and 34 men with ACS, and 253 women and 208 men without ACS, symptoms associated with ACS in women and men seemed quite similar and the authors conclude that discriminating ACS in patients with chest discomfort who contacted primary care OHS is difficult in both women and men. ¹⁷

We aimed to identify clinical variables that are associated with the diagnosis ACS in women and men with chest discomfort who contact out-of-hours primary care (OHS-PC) by telephone, and to explore whether there are indications these variables differ among women and men. For analyses we used the very initial symptom presentation as available from the recorded telephone triage conversations.

Methods

We performed a cross-sectional diagnostic factor study with a random sample of 1,795 OHS-PC calls for chest discomfort (chest pain, pressure, tightness, or discomfort) between 2014 and 2016. ^{18, 19} We first selected calls on the basis of International Code for Primary Care (ICPC; a WHO world-wide code system for primary care) codes (K01, K02, K03, K24, K74, K75, K76, K77, K93, L04, P74, R02, R98, appendix-table 1) and keywords thoracic pain, chest pain, myocardial infarction, heart attack and their common abbreviations mentioned by the triage nurse in the electronic medical file (EMF) at the OHS-PC. ^{20, 21} General practitioners who work at the OHS-PC assign the ICPC codes to the call. We combined ICPC-codes and keywords to achieve a sample with a broad variety of symptoms to capture the entire domain of patients suspected of ACS. We listed all available calls of these patients and assigned random numbers with the Random Number Generator (RAND) function in Microsoft Excel to retrieve a random sample. Calls were excluded before re-listening when the patients' age was below 18 years or when the patient did not live in the surrounding area of the OHS-PCs (because then we could not retrieve a diagnosis from the general practitioner of these patients). Calls were excluded during re-listening when

We re-listened the telephone triage recordings to collect information about patient and conversation characteristics, on symptom presentation, medical history, urgency allocation, and involvement of a supervising general practitioner (GP) in the triage conversation. Gender considered the self-identified gender of the patient. Call duration and age were retrieved from the electronic EMF of the OHS-PC. Nine OHS-PC in the Netherlands participated, serving a total population of 1.5 million people. The diagnosis was made after the phone call, which was in the case of ACS nearly always done by the cardiologist (97.1%) in the hospital based on (i) symptom presentation, (i) levels of (high-sensitivity) troponin and (iii) electrocardiography results. The final diagnoses were provided by the patients' GP, based on the electronic medical file including ED and cardiologist discharge letters, and also the notes from the OHS-PC. We used medical information up to 30-days following the contact with the OHS-PC, to allow us to include diagnoses of ACS that was initially missed because the patient was not referred to the cardiologist the same day of the OHS-PC contact. In none of the patients in the study we had evidence of a missed diagnosis of ACS.

Context

In the Netherlands, OHS-PC covers primary care during 73% of the week hours, and the initial contact is by telephone. In most OHS-PC and EMS, the 'Netherlands Triage Standard' (NTS) is used as a decision support to classify the urgency of the patients' conditions. ²² Based on the patient's symptom presentation, the triage nurse needs to choose the most appropriate complaint out of 56 'main complaints'. Each NTS 'main complaint' incorporates a decision tree with hierarchically ordered questions, which are similar for men and women. Triage nurses fill out the caller's responses in the semi-automatic NTS system, which then generates urgency allocations linked to a timeframe within which the patient should be seen by a physician or ambulance personnel (U0 (reanimation) to U5 (self-care advice) appendix-table 2). The triage nurse can overrule this recommendation and up- or downscale the urgency allocation, often after consulting the supervising GP. ²³ A recent validation study showed that the diagnostic accuracy of the NTS for patients with chest discomfort is poor (sensitivity 0.73 (95% CI 0.68-0.78) and specificity 0.43 (95% CI 0.40-0.45)), as calculated on the outcome ACS or

other life-threatening events (LTEs). ²⁴ All telephone calls to the OHS-PC are routinely recorded and archived for five years for training and quality control purposes.

Data analyses

We compared patient and call characteristics between women with and without ACS, and men with and without ACS. For comparison of dichotomous variables we used the Chi² test or Fisher exact test and for continuous variables the independent sample t-test or Mann-Whitney U test. We performed interaction analysis across gender separately for each clinical variable with logistic regression analyses, to explore whether there are indications that these variables are differently associated with the diagnosis ACS among men and women. We analysed the association between urgency allocation and the final diagnosis ACS (alone or including other LTEs) with the Chi² test. We considered pulmonary embolism, thoracic aortic dissection and acute abdominal aneurysm as LTEs; patients with LTEs as well as those with ACS should receive an U1-level urgency.

Data analysis was performed using SPSS, IBM version 25.

Patient and public involvement

No patients were involved in setting the research question or the outcome measures, or in developing plans for design, however, they were involved in the implementation of the study. In addition, they were asked to advise on interpretation and writing up of results. Results will be shared and discussed with the national patient community of cardiovascular diseases ('Harteraad').

Ethics

The study (National Trial Register identification number: NTR7331) was approved by the Medical Ethics Committee Utrecht, the Netherlands (reference number WAG/mb/16/003208) and complied with the Declaration of Helsinki. A waiver of informed consent was given because our study had minimal risk to subjects and could otherwise not be carried out logistically. Personal and research data were handled and stored according to the European General Data Protection Regulation.

Results

Among the 1,795 callers with chest discomfort (mean age 58.8 (SD 19.5) years, 55.3% women), 8.6% of women and 15.0% of men had an ACS. In women with an ACS, 18.8% had a STEMI, 48.2% a NSTEMI,

A total of 22 (2.2%) women and 23 men (2.9%) had another LTE than ACS (e.g. pulmonary embolism, thoracic aortic dissection, acute abdominal aneurysm, acute heart failure).

Patient and call characteristics

Men and women with ACS were older than those without ACS (mean age of women with ACS 73.6 vs. without ACS 57.8 years, p<0.001, men 67.2 vs. 56.9 years, p<0.001, gender interaction p=0.094), and the mean duration of the telephone calls was shorter (women 6:47 vs. 7:47 minutes, p=0.021, men 6:31 vs. 7:33 minutes, p=0.004, gender interaction p=0.803) (table 1). The GP was consulted for supervision by the triage nurse in the majority of cases (52.2% in women and 55.5% in men). However, in women with ACS, the GP was less often consulted than in women without ACS (41.2% vs. 53.2%, p=0.034), but in men such a difference was not observed (53.3% vs. 55.9%, p=0.607, gender interaction p=0.208). In around half of the calls, someone else called initially on behalf of the patient (49.5% in women vs. 54.7% in men). In cases with ACS, for both sexes more often someone else called than in those without ACS (in women 69.4% vs. 47.7%, p<0.001, in men 65.8% vs. 52.8%, p=0.008, gender interaction p=0.251). In men with ACS, most often their female partner (53.3%) called, while in women with ACS, it was either their male partner (17.6%), their daughter (20.0%) or a nurse (17.6%) (appendix-table 3). Callers expressed concerns in nearly all calls, also in those without an ACS; women with ACS vs. women without ACS 97.3% vs. 88.9% (p=0.109), and men with ACS vs. men without ACS 96.3% vs. 86.5% (p=0.041) (gender interaction p=0.935).

Both women and men with ACS had more often a history of coronary artery disease (women 42.5% vs. 25.1%, p=0.017, and men 57.0% vs. 38.4%, p=0.002, gender interaction p=0.927), but women with ACS had more often a history of diabetes (41.4% vs. 14.6%, p<0.001, gender interaction p=0.079).

Symptom presentation

Chest pain was the most common complaint, both in patients with and without an ACS (women with ACS and without 98.8% and 93.1%, p=0.055, in men 92.4% and 94.5%, p=0.364, gender interaction p=0.048). Retrosternal located chest pain was more common in women and men with ACS than in those without ACS (women 62.3% vs. 40.3%, p=0.002, and men 52.5% vs. 39.7%, p=0.032, gender interaction p=0.323). Also, radiation of pain to the arms was associated with ACS in both sexes (women with vs.

without ACS 75.6% vs. 45.9%, p<0.001, and men 56.0% vs. 34.8%, p<0.001, gender interaction p=0.339), as was pressing/heavy/tightening chest pain (women with vs. without ACS 78.6% vs. 61.5%, p=0.011 and men 82.1% vs. 57.4%, p=<0.001, gender interaction p=0.368). Only in women radiation to the jaw had an association with ACS (women 50.0% vs. 22.9%, p=0.007, men 23.6% vs. 30.4%, p=0.312, gender interaction p=0.015) and severe pain (8 or more on a Numeric Rating Scale 0-10) (65.4% vs. 38.1%, p=0.006, men 2.6% vs. 11.3%, p=0.098, gender interaction p=0.007), which had a differential effect towards the risk of ACS in women. Only in men, stabbing pain was very rare in those with ACS (8.4% vs. 26.5%, p<0.001,) however this had not have a differential effect on the diagnosis of ACS between men and women (gender interaction p=0.141).

Of the autonomous nervous system (ANS)-related symptoms, nausea/vomiting and dizziness/near fainting were not associated with ACS in either sex. A pale or ashen face was associated with ACS in women (55.6% vs. 35.5%, p=0.019, gender interaction p=0.545), and sweating in men (52.4% vs. 38.1%, p=0.015, gender interaction p=0.418), however without a differential effect on the risk of a diagnosis of ACS between women and men. Recognition of symptoms being similar to a previous cardiac event was associated with ACS in men (52.9% vs. 32.1%, p=0.004), but not clearly for ACS in women (32.5% vs. 21.4%, p=0.108, gender interaction p=0.532).

Subgroup analyses in 56 women and 58 men with diabetes showed that both women (85.7% vs. 58.3%, p<0.001) and men with diabetes (67.2% vs. 51.5%, p=0.033, gender interaction p=0.119) more often had shortness of breath than those without diabetes, but as often chest discomfort (women 90.9% vs. 95.0%, p=0.193, men 89.2% vs 94.1%, p=0.162, gender interaction p=0.969). Shortness of breath in patients with diabetes was not related to ACS diagnosis (women 81.8% vs. 86.7%, p=0.680, men 75.0% vs. 66.0%, p=0.615, gender interaction p=0.520).

Diagnoses

Of the 205 patients with an ACS (85 women, 120 men), 55 (26.8%) patients had a STEMI (women 18.8%, men 32.5%), 85 (41.5%) a NSTEMI (women 48.2%, men 36.7%), 50 (24.4%) unstable angina pectoris (UAP) (women 20.0%, men 27.5%) and 15 (7.3%) unspecified ACS (women 13.0%, men 3.3%), the latter also including two sudden cardiac deaths in women and one in men (Table 2). In nearly all cases (97.1%) the ACS diagnosis was made by a cardiologist based on symptom presentation, troponin levels and electrocardiography. Three patients died before arrival of the ambulance (they were classified as acute

Urgencies

Women and men with chest discomfort were equally sent an ambulance (43.6% vs. 46.6%, p=0.200). This was also in women and men who had an ACS (72.9% vs. 70.0%, p=0.647), and in those with either ACS or other LTEs (66.4% vs. 67.1%, p=0.897). See table 3.

Discussion

For both sexes, retrosternal pain, pain described as pressing, heavy or tightening, and radiation to the arm were associated with ACS in patients who contacted the OHS-PC for chest discomfort. Radiation to the jaw and severe pain were related to ACS in women. Our results indicate there were more similarities than differences in symptoms associated with the diagnosis ACS for women and men. However, whether these differences have an impact on predicting ACS needs to be further investigated. Women and men with chest discomfort as also those with ACS were equally often sent an ambulance.

Our finding that radiation of pain to the arm and retrosternal ('mid') chest pain were associated with the diagnosis ACS in both sexes was also reported in a study among 2,475 patients with acute chest pain in a multicentre ED-study. ²⁵ Another ED-study among 1,334 patients with ACS showed that regardless of ethnics status the most common presenting symptom was retrosternal pain/discomfort of any intensity. ²⁶ The aforementioned US study in the ED-setting reported that radiation to the arm was associated with ACS in women but not for men, and chest pressure was associated with ACS for men but not in women.

¹³ The only previously published OHS-PC study reported the opposite; radiation to the arm was associated with ACS in men, but not in women. ¹⁷

In our study, women with ACS had more often a history of diabetes and were older than men with ACS, which is in line with other studies. ^{8, 12} Some studies claim that patients with diabetes more often have atypical symptoms of ACS, however a review of eight studies concluded the evidence of these studies was conflicting. ²⁷ We showed that both women and men with diabetes had more often shortness of breath than those without diabetes, but shortness of breath in patients with diabetes was not associated with ACS. Regarding dispatching the ambulance, the aforementioned OHS-PC study and two EMS studies showed, similar to our findings, that there was no difference in dispatch priorities between men and women with ACS. ^{4, 17, 28} This is in contrast with studies that show delay in hospital presentation of women with ACS. ^{9, 29}

We need to realize that focusing on gender differences may blur the large overlap in symptoms in women and men. Moreover, comparing *selectively* women with ACS to men with ACS as many previous studies did, is clinically irrelevant. ^{9, 14, 30} Clinicians, including GPs, and triage nurses need to know whether and how women with ACS differ from women without ACS, with the same question for men. Nevertheless, even guidelines stick to comparing those with established disease, and express the view that women with ACS more likely present with less specific symptoms than men with ACS. ^{1, 31} Unfortunately, public awareness campaigns follow this reasoning, and over-emphasize sex differences in women awareness campaigns ('Go Red for Women' in the United States and 'Invisible me' in Australia). ^{7, 30, 32} Unbalanced attention to symptom differences, neglecting the much larger overlap may even introduce new blind spots in recognizing ACS in women. ³³

A likely reason behind the predominant message that women present with other ACS symptoms than men is the difference in pathophysiology of coronary artery disease. Women compared to men more often have elongated plaques, located on bifurcations in epicardial coronaries, coronary spasm, microvascular dysfunction, and spontaneous coronary dissection. ^{32, 34} These pathophysiological differences have an effect on interventional treatment and prognosis. ^{2, 34} However, these differences do not necessarily imply an effect on symptom presentation because the pain pathway is equal in women and men: i.e. triggered by myocardial ischemia. ^{35, 36} A supply and demand mismatch of the myocardial oxygen consumption triggers sensory nerve endings in the myocardium and cause ischemia symptoms, and this is irrespective of the fact whether the ischemia is caused by a plaque rupture in an

Another reason behind the belief of 'vague' symptom presentation in women with ACS might be that they seem to present a larger number of symptoms than men with ACS, and this may be interpreted as 'vague' by physicians. ^{15, 37} Presentation of multiple symptoms may influence the prompt recognition of heart disease and initial actions on the part of health care providers. ^{37, 38} In a study from 2018, with 2009 women and 976 men hospitalized for myocardial infarction, healthcare providers initially thought symptoms of women (53.4%) were less often heart-related than in men (36.7%). ³⁷ In that study women and men had the same chest pain symptoms, but women reported more additional symptoms. ³⁷ In our study, the call duration and the number of GP consultations by the triage nurse were similar among women and men, suggesting that triage nurses seem not to experience more difficulties in interpreting symptoms in women than men. This is in line with a prospective study with 2,795 patients with chest discomfort in ED-setting that showed the physicians' diagnostic uncertainty for the presence of ACS in women was not more common as compared to men.³⁹

Interestingly, in the majority of calls in our study someone else than the patient called the OHS-PC (women with ACS 69.4%, without ACS 47.7%, and men with ACS 65.8%, and without ACS 52.8%). This was also highlighted in an Australian study among 1,681 patients with an acute myocardial infarction; in 90.5% of the women with AMI someone else called on behalf of the patient and in 87.8% of the men with AMI. ⁴⁰ According to protocol in OHS-PC, triage nurses ask the patient to the phone, this to prevent loss of (paralinguistic) information from the patient him/herself. In our study, in about 50% of the conversations the patient took over the phone call.

Strengths and limitations

A major strength is that we could analyse the very initial conversation with symptom presentation of a large sample of patients calling the OHS-PC because of chest discomfort. We analysed the conversations without knowledge of the eventual diagnosis and have prevented risk of hindsight bias of the researcher or recall bias of the patient. Another strength is that we performed gender subgroup analyses combined with interaction analyses across gender, to investigate whether there are indications that ACS related symptoms differ among men and women. Furthermore, our results are generalizable to comparable primary care settings, e.g. UK and Scandinavian countries, and possibly some other European countries.³

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Our results may even be generalizable to EMS settings, since the prior probability of having an ACS is comparable in EMS setting as in OHS-PC settings. 4 41

As the intention of our analysis was to describe whether symptoms were different in patients with ACS from patients without ACS in women and men separately, none of our results can be used to adjust interview questions for the triage nurses. For that purpose, prediction rule development with multivariable analyses is necessary. Also, only with multivariable analysis it can be truly investigated whether the potential differences are clinically relevant in prediction of ACS. Another limitation is missing values on some clinical variables, a phenomenon common in routine care data, and therefore the results should be interpreted with caution.

Conclusions

Our results indicate there were more similarities than differences in symptoms associated with the diagnosis ACS for women and men. Important exceptions were pain severity and radiation of pain in women. However, whether these differences have an impact on predicting ACS needs to be further investigated with multivariable analyses.

The authors would like to thank the OHS-PC foundation 'Primair Huisartsenposten' and all employees of the participating locations for or their cooperation in this study, notably for providing data and technical support.

Data sharing statement

The data can be made available for researchers whose proposed use of the data has been approved at request of the corresponding author, with a signed data access agreement.

Author contribution

FR and DZ are the lead investigators who conceived the research idea and methodology. Funding acquisition was done by FR, DZ and RD. LW and DC conducted data acquisition. LW performed the analyses and wrote the first draft of the manuscript. She was supervised by FR and DZ, who critically revised the manuscript. DC, EdG, AH and RD contributed with and approved the final version of the manuscript. MvS was involved in adjusting the analyses and revising of the manuscript.

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Table 1. Baseline characteristics of 993 (55.3%) women and 802 (44.7%) men patients contacting the OHS-PC w	-	ಕ್ಷ est discomfort (pain,
pressure, tightness, or discomfort)	<u>}</u>	-042

7 Characteristics	993 women			Difference (95% CI)	802 men		6	Difference (95% CI)	P-value
8		I	T .	_		2	3	=	interaction
9	ACS	No ACS	p-value		ACS	No ACS	p- Va lue		gender
10	(2.22)						<u> </u>		gender
1 1	N = 85 (8.6%)	N = 908 (91.4%)			N = 120 (15.0%)	N = 682 (85.0%)	e me		
12							20 20 20 20 20 20 20 20 20 20 20 20 20 2		
13 Mean age in years (SD)	73.6 (±14.1)	57.8 (±20.1)	<0.001	15.8 (11.4-20.2)	67.2 (±13.0)	56.9 (±19.2)	₹ 0 .0 01	10.3 (6.7-13.9)	0.094
14		Uh				9	Sh D		
15 Call characteristics		/ h					ge ≸		
16							loa Sc		
17 Mean call duration in min (SD)	6:47 (±5:16)	7:47 (±3:48)	0.021	1.00 (0.19-1.80)	6:31 (±3:13)	7:33 (±3:42)) 2000 1000 1000 1000 1000 1000 1000 100	1.02 (0.37-1.68)	0.803
18							<u> </u>		
19 Mean patient's introduction in min (SD)	0:14 (± 0:08)	0:19 (±0:13)	<0.001	0.05 (0.02-0.09)	0:16 (±0:11)	0:19 (±0:14)	0.0050	0.03 (0.00-0.06)	0.042
20							-		
2 Triage nurse consulted the GP	35 (41.2)	483 (53.2)	0.034	12.0 (0.3-22.9)	64 (53.3)	381 (55.9)	0.	2.5 (-7.3-12.6)	0.208
22							† 5		
23 Someone else called on behalf of patient	59 (69.4)	433 (47.7)	<0.001	21.7 (10.1-31.6)	79 (65.8)	360 (52.8)	0.008	13.0 (3.0-22.2)	0.251
24							₽ <mark>8</mark>		
25 The patient or person who called	36 (97.3)	409 (88.9)	0.109	8.4 (-5.0-12.6)	52 (96.3)	321 (86.5)	0.041	9.8 (-0.9-14.8)	0.935
26 27 expressed concerned (n=922)							į į		
27 expressed concerned (n=322)						0	<u> </u>		
28 29 Chest pain (n=1739)	81 (98.8)	817 (93.1)	0.055	5.7 (-0.8-8.0)	110 (92.4)	624 (94.5)	0 34	2.1 (-2.3-9.0)	0.048
29 Chest Pain (11–1739)	01 (90.0)	017 (93.1)	0.033	3.7 (-0.6-6.0)	110 (92.4)	024 (94.3)	0. 36 4	2.1 (-2.3-9.0)	0.048
30 31 Severe Pain (>7 on a scale 0-10) (n=753)	17 (65.4)	147 (38.1)	0.006	27.3 (5.7-44.7)	13 (23.6)	87 (30.4)	0.342	6.8 (-7.9-18.3)	0.007
3 Severe Pain (>7 on a scale 0-10) (11-755)	17 (05.4)	147 (50.1)	0.000	27.3 (3.7-44.7)	15 (25.0)	67 (50.4)	0.3 <u>1</u>	0.6 (-7.9-16.5)	0.007
32 33 Duration						87 (30.4)	13		
33 Duration							8, 20		
34 > 15 min (n=1501)	66 (100)	720 (05 8)	0.102	42/2860	00 (07 1)		1 0	25/2056)	0.008
35 > 15 min (n=1501)	66 (100)	729 (95.8)	0.102	4.2 (-2.8-6.0)	99 (97.1)	541 (94.6)	0. 29 2	2.5 (-3.8-5.6)	0.998
36	60 (05.7)	 F7F (70 7)	0.040	42.0 (4.7.20.6)	00 (04 5)	424 (72.6)	ر کی ا	0.0 (0.6 46 5)	0.400
37 < 12 hrs (n=1563)	60 (85.7)	575 (72.7)	0.018	13.0 (1.7-20.6)	88 (81.5)	431 (72.6)	0. @partm	8.9 (-0.6-16.5)	0.490
38							17		
39 Location (n=1267)*							nent		
40							<u>nt</u> G		
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Retrosternal	33 (62.3)	263 (40.3)	0.002	22.0 (7.1-35.2)	42 (52.5)	191 (39.7)	<u>ig</u> n- - 1 0.662	12.8 (0.5-24.8)	0.323
Right or left side thorax	14 (26.4)	260 (39.8)	0.054	13.4 (-1.2-24.8)	26 (32.5)	211 (43.9)	2 2020-0#2406 0.0.0.0.0.0.0.0.0.0.0.0.0.0.0.0.0.0.0	11.4 (-1.0-22.2)	0.757
Type of pain (n=1334)**							406 on ng for		
Pressing/heavy/tightening	44 (78.6)	423 (61.5)	0.011	17.1 (3.2-27.2)	78 (82.1)	284 (57.4)	n 250June 2021. © Erasmu ruses related to	24.7 (14.3-32.9)	0.368
Stabbing	8 (14.3)	155 (22.5)	0.152	8.2 (-4.6-16.4)	8 (8.4)	131 (26.5)	Ine 201	18.1 (9.2-24.1)	0.141
Radiation of chest pain (n=1391) ***		04)21. D smush ad to t		
Any location	61 (85.9)	485 (68.6)	0.002	17.3 (6.0-25.0)	65 (63.7)	292 (57.1)	. Dowizioaded f ushogeschool o text and data	6.6 (-4.5-16.7)	0.071
Arm	31 (75.6)	188 (45.9)	<0.001	29.7 (12.8-42.2)	47 (56.0)	117 (34.8)	adea f chool of data	21.1 (8.7-32.9)	0.339
Between the shoulder blades	11 (52.4)	159 (41.7)	0.336	10.7 (-12.0-32.5)	14 (27.5)	91 (29.4)	frogs http	1.9 (-13.4-14.3)	0.352
Jaws	10 (50.0)	66 (22.9)	0.007	27.1 (4.3-49.7)	1 (2.6)	28 (11.3)	ttp%/bn 0.7	8.7 (-4.6-14.1)	0.015
Shortness of breath (n=1365)	47 (72.3)	455 (66.3)	0.328	6.0 (-7.2-16.7)	53 (60.9)	329 (62.4)	min 0. 78 8	1.5 (-9.5-13.3)	0.355
Symptoms similar to previous cardiac	13 (32.5)	79 (21.4)	0.108	11.1 (-3.0-28.4)	27 (52.9)	92 (32.1)	9 5 2 0. 9 4	20.8 (5.4-35.8)	0.532
event (n=748)							.@mj.com/ o. and simi		
ANS-related symptoms							nii on		
Sweating (n=1130)	28 (49.1)	234 (42.5)	0.340	6.6 (-7.4-20.7)	43 (52.4)	170 (38.5)	on Juge 13,20, on Juge 13,20, o. ar technologies	13.9 (16.7-25.8)	0.418
Nausea or vomiting (n=808)	16 (44.4)	240 (56.2)	0.173	11.8 (-6.2-28.6)	27 (45.8)	122 (42.7)	13,\(\varphi\)025 ologies.	3.1 (-11.1-17.7)	0.186
Pallor or ashen skin (n=652)	20 (55.6)	110 (35.5)	0.019	20.1 (1.9-37.0)	28 (53.8)	103 (40.6)	0. 047 8	13.2 (-2.3-28.3)	0.545
Dizziness or near fainting n=1599)	13 (18.6)	197 (24.4)	0.277	5.8 (-6.0-14.3)	17 (15.9)	127 (20.7)	Degartment (4.8 (-4.3-11.9)	0.963
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3 CV disease or risk factors (n=1461)	54 (78.3)	407 (57.5)	0.001	20.8 (8.4-30.2)	75 (72.1)	359 (61.9)	, 0. 22 6	10.2 (-0.3-19.3)	0.179
5 Coronary artery disease (n=966)	17 (42.5)	114 (25.1)	0.017	17.4 (1.7-34.3)	45 (57.0)	151 (38.4)	-2820-08240639n	18.6 (5.9-30.5)	0.927
7 Hypertension (n=756)	22 (71.0)	142 (35.1)	<0.001	35.9 (16.0-50.7)	17 (47.2)	94 (33.0)	0.891	14.2 (-3.0-32.1)	0.094
9 Diabetes mellitus (n=745) 10	12 (41.4)	56 (14.6)	<0.001	26.8 (9.0-46.5)	12 (28.6)	59 (20.3)	252Jun o. J uses re	8.3 (-5.1-25.1)	0.079
11 Hypercholesterolemia (n=679) 12	8 (36.4)	81 (23.2)	0.161	13.1 (-5.8-36.4)	19 (45.2)	62 (23.3)	ne202 Ecasm	21.9 (5.8-38.6)	0.527
13 *P-value comparing retrosternal or left, 14	/right side thorax v	s. others locations	of pain togeth	her (restrosternal, left,	/right side thorax,	back/shoulder, epi)	
15 ** Prossing/hogyy/tightoning nain vs. o	other types of pain	(stabbing, burning	, cramping, te	aring). Stabbing pain:	stabbing vs. other	types of pain (pres	x+ 00 M sing e h@a ⊋ v/tig	ghtening, burning, cram	nping, tearing)
17				G. G.	Ü		oaded school nd data		
18 *** P-value comparing radiation arm or 18	back/shoulder or	jaws vs. 110 radiatio	on .				d fro ol . ita m		
19 20 NRS: Numeric Rating Scale							m htt ining,		
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26 Coronary artery disease: History of prio	r MI, PCI, CABG, st	able or unstable an	igina pectoris	(patient reported)			en.bmj. 1g, and		
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Table 2. Diagnosis of 1,795 patients who contacted the OHS-PC for chest discomfort (pain, pressure, tightness, or discomfort), divided in women and men

	Women n= 993 (55.3%)	Men n=802 (44.7%)	p-value
	, , ,	, ,	•
Acute coronary syndrome	85 (8.6)	120 (15.0)	<0.001
STEMI	16 (18.8)	39 (32.5)	0.037
NSTEMI	41 (48.2)	44 (36.7)	0.114
UAP	17 (20.0)	33 (27.5)	0.250
Non-classified ACS	11 (13.0)	4 (3.3)	0.013
Life threatening events (LTE's)	22 (2.2)	23 (2.9)	0.448
Pulmonary embolism	6 (27.3)	7 (30.4)	0.815
Thoracic aortic dissection	4 (18.2)	2 (8.7)	0.349
Acute abdominal aneurysm	3 (13.6)	2 (8.7)	0.598
Other*	9 (40.9)	12 (52.2)	0.449
Non-urgent cardiovascular	194 (19.5)	170 (21.2)	0.384
diseases**		4	
Non-cardiac chest pain, not further	163 (16.4)	159 (19.8)	0.061
specified ***		0,	
Musculoskeletal pain	199 (20.0)	113 (14.1)	0.001
Psychogenic disorders	139 (14.0)	67 (8.4)	<0.001
Gastrointestinal tract disorders	76 (7.7)	62 (7.7)	0.951
Respiratory tract disorders	52 (5.2)	45 (5.6)	0.727
Other non-urgent diagnoses***	63 (6.3)	43 (5.4)	0.380

^{*}Acute heart failure, stroke, severe COPD exacerbation, sepsis, coronary spasm probably caused by hypokalaemia, diabetic ketoacidosis, epileptic insult, bleeding from oesophageal varices, ovarian torsion, ventricular fibrillation.

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**** Amongst others: anaemia, malignancy, vasovagal collapse, side effects medication, dermatologic diseases



Table 3. Association between urgency allocation, diagnose ACS and other life threatening events (LTE)

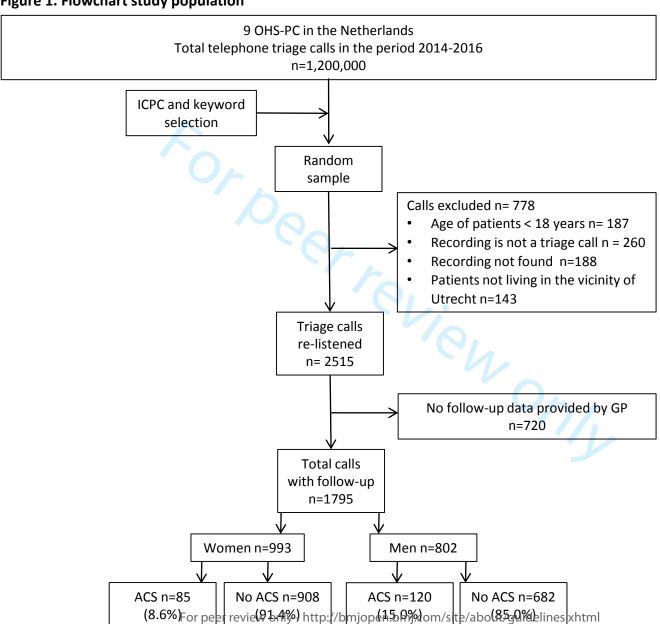
Women	ACS n=85 (8.6%)	No ACS n=908 (91.4%)	p-value 1*	p-value 2**
U1	62 (72.9)	371 (40.9)	<0.001	<0.001
U2	12 (14.1)	231 (25.4)	_	
U3-U5	11 (13.0)	306 (33.6)		
Men	ACS n=120 (15.0%)	No ACS n=682 (85.0%)		
U1	84 (70.0)	290 (42.5)	<0.001	<0.001
U2	19 (15.8)	142 (20.8)	_	
U3-U5	17 (14.2)	250 (36.7)	_	
Women	ACS or LTE n=107 (10.8%)	No ACS or LTE n=886 (89.2%)		
U1	71 (66.4)	362 (40.9)	<0.001	<0.001
U2	21 (19.6)	222 (25.1)	_	
U3-U5	15 (14.0)	302 (34.0)		
Men	ACS or LTE n=143 (17.8%)	No ACS or LTE n=659 (82.2%)		
U1	96 (67.1)	278 (42.2)	<0.001	<0.001
U2	24 (16.8)	137 (20.8)		
U3-U5	23 (16.1)	244 (37.0)	5,	

^{*} P-value 1: U1 vs. U2, U3, U4 and U5

^{**} P-value 2: U1,U2 vs. U3,U4,U5

^{***}LTE = life threatening event. Life threatening events consist of: ACS; pulmonary embolism; thoracic aortic dissection; acute heart failure; stroke; abdominal aortic aneurysm; severe COPD exacerbation; diabetic ketoacidosis; coronary spasm probably caused by hypokalaemia; epileptic insult; bleeding from oesophageal varices; ovarian torsion; ventricular fibrillation.

Figure 1. Flowchart study population



2020-042406 on 25 June 2021. Downloaded from http://bmjopen.bmj.com/ on June 13, 2025 at Department GEZ-LTA Erasmushogeschool . t, including for uses related to text and data mining, Al training, and similar technologies.

Supplementary file

Appendix-Table 1: Overview of ICPC-codes used to select calls

ICPC code	Calls of patients with ACS	Calls of patients without ACS
	N=205 (%)	N=1,590 (%)
K01 Pain attributed to the heart	112 (54.6)	485 (30.5)
K02 Pressure/tightness attributed to the heart	39 (19.0)	184 (11.6)
K03 Other cardiovascular pain	1 (0.5)	6 (0.4)
K24 Fear of heart attack	1 (0.5)	8 (0.5)
K74 Angina pectoris	19 (9.3)	101 (6.4)
K75 Acute myocardial infarction	7 (3.4)	22 (1.4)
K76 Other/chronic ischaemic heart disease	0 (0.0)	2 (0.1)
K77 Heart failure	0 (0.0)	11 (0.7)
K93 Pulmonary embolism	0 (0.0)	15 (0.9)
L04 Chest discomfort	25 (12.2)	689 (43.9)
P74 Anxiety disorder	0 (0.0)	12 (0.8)
R02 Shortness of breath	1 (0.5)	36 (2.3)
R98 Hyperventilation	0 (0.0)	10 (0.6)
	7007	

Implication
Reanimation
Life-threatening, GP/ ambulance should arrive within 15 minutes
Emergency, GP should arrive within 60 minutes
Urgent, consultation by GP within three hours
Routine, consultation by GP the same day
Advice given by triage nurse

STROBE Statement—Checklist of items that should be included in reports of cross-sectional studies

	Item No	Recommendation	Page No
Title and abstract	1	(a) Indicate the study's design with a commonly used term in the	1,3
		title or the abstract	
		(b) Provide in the abstract an informative and balanced summary of	3
		what was done and what was found	
Introduction			
Background/rationale	2	Explain the scientific background and rationale for the investigation being reported	5,6
Objectives	3	State specific objectives, including any prespecified hypotheses	6
Methods			
Study design	4	Present key elements of study design early in the paper	6
Setting	5	Describe the setting, locations, and relevant dates, including periods	6,7
C		of recruitment, exposure, follow-up, and data collection	
Participants	6	(a) Give the eligibility criteria, and the sources and methods of	6,7
•		selection of participants	
Variables	7	Clearly define all outcomes, exposures, predictors, potential	6,7
		confounders, and effect modifiers. Give diagnostic criteria, if	
		applicable	
Data sources/	8*	For each variable of interest, give sources of data and details of	6,7
measurement		methods of assessment (measurement). Describe comparability of	
		assessment methods if there is more than one group	
Bias	9	Describe any efforts to address potential sources of bias	8
Study size	10	Explain how the study size was arrived at	7
/Quantitative	11	Explain how quantitative variables were handled in the analyses. If	8
variables		applicable, describe which groupings were chosen and why	
Statistical methods	12	(a) Describe all statistical methods, including those used to control	8
		for confounding	
		(b) Describe any methods used to examine subgroups and	8
		interactions	
		(c) Explain how missing data were addressed	14
		(d) If applicable, describe analytical methods taking account of	
		sampling strategy	
		(\underline{e}) Describe any sensitivity analyses	8
Results			
Participants	13*	(a) Report numbers of individuals at each stage of study—eg	19 (figure 1)
•		numbers potentially eligible, examined for eligibility, confirmed	
		eligible, included in the study, completing follow-up, and analysed	
		(b) Give reasons for non-participation at each stage	19 (figure 1)
		(c) Consider use of a flow diagram	19
Descriptive data	14*	(a) Give characteristics of study participants (eg demographic,	7
•		clinical, social) and information on exposures and potential	
		confounders	
		(b) Indicate number of participants with missing data for each	20,21 (table 1)
		variable of interest	
Outcome data	15*	Report numbers of outcome events or summary measures	9
	-	1	

Main results	16	(a) Give unadjusted estimates and, if applicable, confounder-	n.a.
		adjusted estimates and their precision (eg, 95% confidence interval).	
		Make clear which confounders were adjusted for and why they were	
		included	
		(b) Report category boundaries when continuous variables were	20,21 (table 1)
		categorized	
		(c) If relevant, consider translating estimates of relative risk into	NA
		absolute risk for a meaningful time period	
Other analyses	17	Report other analyses done—eg analyses of subgroups and	9,10 (subgroup
		interactions, and sensitivity analyses	diabetes)
Discussion			
Key results	18	Summarise key results with reference to study objectives	11
Limitations	19	Discuss limitations of the study, taking into account sources of	14
		potential bias or imprecision. Discuss both direction and magnitude	
		of any potential bias	
Interpretation	20	Give a cautious overall interpretation of results considering	13,14
		objectives, limitations, multiplicity of analyses, results from similar	
		studies, and other relevant evidence	
Generalisability	21	Discuss the generalisability (external validity) of the study results	14
Other information			
Funding	22	Give the source of funding and the role of the funders for the present	2
		study and, if applicable, for the original study on which the present	
		article is based	

^{*}Give information separately for exposed and unexposed groups.

Note: An Explanation and Elaboration article discusses each checklist item and gives methodological background and published examples of transparent reporting. The STROBE checklist is best used in conjunction with this article (freely available on the Web sites of PLoS Medicine at http://www.plosmedicine.org/, Annals of Internal Medicine at http://www.annals.org/, and Epidemiology at http://www.epidem.com/). Information on the STROBE Initiative is available at www.strobe-statement.org.