Predicting acute coronary syndrome in males and females with chest pain who call an emergency medical communication centre

Paul-Georges Reuter 1,2,3*, Catherine Pradeau 4, Samantha Huo Yung Kai 2,5, Thibault Lhermusier 6, Arnaud Bourdé 7, Eric Tentillier 4, Xavier Combes 7, Vanina Bongard 2,5, Jean-Louis Ducassé 1 and Sandrine Charpentier 1,2

Abstract

Background: Chest pain is a frequent reason for calls in emergency medical communication centre (EMCC). Detecting a coronary origin by phone is a challenge. This is especially so as the presentations differ according to gender. We aimed to establish and validate a sex-based model to predict a coronary origin of chest pain in patients calling an EMCC.

Methods: This prospective cohort study enrolled patients at 18 years of age or older who called the EMCC because of non-traumatic chest pain. The main outcome was the diagnosis of acute coronary syndrome (ACS) determined by expert evaluation of patient files.

Results: During 18 months, 3727 patients were enrolled: 2097 (56%) men and 1630 (44%) women. ACS was diagnosed in 508 (24%) men and 139 (9%) women. For men, independent factors associated with an ACS diagnosis were age, tobacco use, severe and permanent pain; retrosternal, breathing non-related and radiating pain; and additional symptoms. The area under the receiver operating characteristic curve (AUC) was 0.76 (95% confidence interval [CI] 0.73–0.79) for predicting ACS. The accuracy of the male model to predict ACS was validated in a validation dataset (Hosmer-Lemeshow test: p = 0.554); the AUC was 0.77 (95%CI 0.73–0.80). For women, independent factors associated with an ACS diagnosis were age ≥ 60 years, personal history of coronary artery disease, and breathing non-related and radiating pain. The AUC was 0.79 (95%CI 0.75–0.83). The accuracy of the female model to predict ACS was not validated in the validation dataset (Hosmer-Lemeshow test: p = 0.035); the AUC was 0.67 (95%CI 0.60–0.74).

Conclusions: Predictors of an ACS diagnosis in patients calling an EMCC for chest pain differ between men and women. We developed an accurate predictive model for men, but for women, the accuracy was poor.

Trial registration: This study is registered with ClinicalTrials.gov (NCT02042209).

Keywords: Chest pain, Acute coronary syndrome, Emergency medical communication Centre, Accuracy, Sex disparity

Background

Acute coronary syndrome (ACS) is a frequent pathology worldwide. Management consists in coronary reperfusion, by percutaneous intervention or fibrinolysis. The worst evolution consists in ventricular fibrillation and cardiac arrest. Survival depends on the total ischemic time: delay between chest pain onset and reperfusion [1]. The main symptom is chest pain or discomfort [2]. The first link for out-of-hospital chain of survival is patient education to call the emergency medical communication centre (EMCC) in case of chest pain. The second is the ability of the dispatcher, medical or not, to identify patients with ACS and to send an ambulance (with an emergency physician or not, depending on the country).

In a Copenhagen EMCC, chest pain was the second identified reason after minor trauma for calls, with 11% of the requests [3]. The sex ratio was 1:1 and over half of the...
patients were over 65 years of age [4]. When a contact occurred with an EMCC, the prevalence at 30 days of ACS ranged from 12 to 16% [5, 6]. For patients with ACS, the use of the EMCC was in the range of 23 to 43% [7, 8]. Patients with ST elevation myocardial infarction (STEMI) were given the highest priority in 82% of cases [9].

ACS diagnosis depends on electrocardiography (ECG) findings and biomarkers. At an EMCC, these data are not available. Only patient history and characteristics of chest pain can be investigated. A persistent pain or a typical location with radiation without associated symptoms influences the dispatcher to send a Mobile Intensive Care Unit (MICU) in France [10]. In Sweden, the intensity, the localisation of the pain and a history of ischemic heart disease were associated with the final diagnosis of ACS [11]. The accuracy of a computer-based decision support was compared with dispatchers' decisions to predict ACS [5]. Sensitivity was greater with the computer- than dispatcher-based decision (90% vs. 83%), and that under triage (false negative) was 10 and 17%, respectively. The factors correlated to under triage were the time of the call (lunch time) and the level of medical knowledge of the dispatcher (assistant nurse versus dispatcher with no medical training) [12]. These findings confirm the need for a decision support tool to help dispatchers identify patients at risk of ACS.

Atypical clinical presentations are difficult to diagnose [13]. In older and diabetic patients, chest discomfort can be absent [14, 15]. Males and females also differ in clinical presentation of ACS [2]. Women, particularly those < 55 years old, most often describe atypical chest pain, such as discomfort, pinching, or burning, [16–20]. These differences in initial presentation lead to an increase in mortality among women (10% versus 5%) [21]. Total ischemic time is longer in women than men [8, 9, 22]. One explanation is a lower prioritization for women when calling call centres (79% for women and 89% for men) [9]. These results suggest that attention should be paid to recognize these patients as soon as possible. In creating a “by-phone” predictive score of ACS, items should differ according to sex.

We aimed to establish and validate a model to predict ACS for men and women calling an EMCC from information that can be recorded by phone.

**Methods**

**Study design and setting**

The DOREMI 2 prospective cohort study was conducted in three French university hospitals. This study is a follow-up to DOREMI 1, which was a pilot and feasibility study. The three participating EMCCs were located in Toulouse, Bordeaux and Saint-Denis de la Réunion. In 2017, the EMCCs served 1.318, 1.506 and 0.843 million inhabitants, respectively. After an evaluation by a dispatcher assistant, a physician manages every call for medical reasons. Depending on the dispatcher prioritization, calls are handled by a general practitioner or an emergency physician. In France, calls from patients with chest pain or discomfort are generally transferred to the emergency physician dispatcher. In response to a call, a medical dispatcher can give medical advice, recommend going into a medical care structure or send an ambulance, fire brigade, physician or MICU (ambulance with an emergency physician on board).

**Selection of participants**

From May 2010 to November 2011, we consecutively included adults at 18 years of age or older who called the EMCC for non-traumatic chest pain. The “non-traumatic” characteristic was verified directly by asking the patient. The exclusion criterion was any difficulty in communicating: uncommunicative patient, language barrier, or inability to speak with the patient.

**Measurements**

At the first call to the EMCC, the emergency physician recorded patient characteristics, cardiovascular risk factors, medical history and clinical presentation on a standardized form (Additional file 1). Follow-up data were collected 30 days after the call (D30) by telephone interview. A research assistant contacted the patient’s general practitioner and/or the patient directly. The patient was contacted in case of non-response from the general practitioner or in case of incomplete information.

At the D30 follow-up, the research assistant retrieved reports from the emergency department, hospitalization and additional examinations. They collected data on major adverse cardiovascular events (rehospitalisation, myocardial infarction, urgent revascularization or death), admission or a consultation in a cardiology unit, non-invasive imaging (transthoracic echocardiography, stress imaging with exercise or drug, cardiac MRI), coronary angiography and the final diagnosis during hospitalization. The final diagnosis of ACS was based on these data.

**Outcome**

The outcome was a diagnosis of ACS by experts according to current guidelines [23].

STEMI was defined by the onset of a persistent ST-elevation on ECG, considered suggestive in the following cases: 1) at least two continuous leads with ST-segment elevation > 0.2 mV in leads V1-V3 or > 0.1 mV in leads V4-V9, V3R and V4R or 2) left bundle branch block with the presence of concordant ST-segment elevation.

Non–STEMI was diagnosed with compatible clinical presentation and ECG abnormalities in two continuous leads such as ST-segment depression or T-wave changes and elevated cardiac troponin level higher than the 99th percentile.

Unstable angina was considered when the patient had a compatible clinical presentation and ECG abnormalities.
without elevated cardiac troponin level and at least one of the following abnormalities: 1) dynamic changes of the ST-segment within 30 days or during the stress test; 2) a positive test result from stress echocardiography, cardiac MRI, coronary CT angiography; 3) coronary angiography with >70% occlusion, 4) death within 30 days and, 5) rehospitalisation within 30 days with a diagnosis of ACS.

All patient files were retrospectively analysed by two experts to determine the final diagnosis of ACS or not. Three pairs of experts were recruited from the three centres. They did not belong to the team that included or cared for the patient. Files were randomly assigned. In case of discordance between the two experts, a third one was consulted. The diagnosis was based on pre-hospital data, reports of emergency departments, hospitalization and/or additional examinations, and follow-up on D30.

### Analysis

#### Sample size calculation

At least 10 events per independent variable are recommended to ensure satisfactory statistical power in multivariate regression models [24, 25]. Because we planned to include a maximum of 15 independent variables in the final predictive model for each sex, we needed 150 calls with ACS for men and 150 for women. Given that approximately 16% of calls for non-traumatic chest pain have a definite diagnosis of ACS in the French EMCC, we needed 938 calls for each sex [6]. Given that the percentage of lost to follow-

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**Fig. 1** Flowchart based on final diagnosis for men (**A**) and women (**B**). ACS: Acute Coronary Syndrome; NSTEMI: Non ST Elevation Myocardial Infarction; STEMI: ST Elevation Myocardial Infarction
| Characteristics and type of pain for male patients in the derivation set with and without a diagnosis of acute coronary syndrome (ACS) |
|---------------------------------------------------------------|
| Total (\(n=1398\)) | ACS (\(n=329\)) | No ACS (\(n=1069\)) | \(p\) value |
|---|---|---|---|
| Age, mean (SD), y, No. (%) | 59.2 (16.1) | 59.4 (13.2) | 50.9 (16.4) | < 0.001 |
| < 40 | 291 (20.8) | 18 (5.5) | 273 (25.5) | < 0.001 |
| 40–50 | 317 (22.7) | 59 (17.9) | 258 (24.1) | |
| 50–60 | 320 (22.9) | 95 (28.9) | 225 (21.0) | |
| \(\geq\) 60 | 470 (33.6) | 157 (47.7) | 313 (29.3) | |
| Coexisting conditions, No. (%) | | | | |
| Hypertension | 405 (29.0) | 123 (37.4) | 282 (26.4) | < 0.001 |
| Personal coronary artery disease | 387 (27.7) | 126 (38.3) | 261 (24.4) | < 0.001 |
| Family coronary artery disease | 231 (16.5) | 51 (15.5) | 180 (16.8) | 0.568 |
| Tobacco use | 545 (39.0) | 133 (40.4) | 412 (38.5) | 0.540 |
| Diabetes | 183 (13.1) | 59 (17.9) | 124 (11.6) | 0.003 |
| Dyslipidaemia | 337 (24.1) | 97 (29.5) | 240 (22.5) | 0.009 |
| Medication therapy, No. (%) | | | | |
| Aspirin | 296 (21.2) | 100 (30.4) | 196 (18.3) | < 0.001 |
| Clopidogrel | 209 (14.9) | 77 (23.4) | 132 (12.3) | < 0.001 |
| Thyroid hormone | 12 (0.9) | 2 (0.6) | 10 (0.9) | 0.743 |
| Statin | 263 (18.8) | 87 (26.4) | 176 (16.5) | < 0.001 |
| Severity of pain, median (IQR) | 5 [3–7] | 5 [4–7] | 5 [3–7] | < 0.001 |
| Type of pain, No. (%) | | | | |
| Severe pain (NRS \(\geq\) 6) | 592 (42.4) | 162 (49.2) | 430 (40.3) | 0.004 |
| Permanent | 350 (25.0) | 102 (31.0) | 248 (23.2) | 0.004 |
| Pain onset: | | | | 0.066 |
| Crescendo | 307 (22.0) | 67 (20.4) | 240 (22.5) | |
| Abruptly | 590 (42.2) | 157 (47.7) | 433 (40.5) | |
| Unclassified | 501 (35.8) | 105 (31.9) | 396 (37.0) | |
| Circumstance of pain, No. (%) | | | | 0.012 |
| At rest | 1106 (79.1) | 244 (74.2) | 862 (80.6) | |
| Sport or stress-related | 180 (12.9) | 58 (17.6) | 122 (11.4) | |
| Unclassified | 112 (8.0) | 27 (8.2) | 85 (8.0) | |
| Pain typography, No. (%) | | | | |
| Typical chest pain\(^a\) | 799 (57.2) | 224 (68.1) | 575 (53.8) | < 0.001 |
| Retrosternal | 671 (48.0) | 201 (61.1) | 470 (44.0) | < 0.001 |
| Post-myocardial infarction angina | 119/387 (30.8) | 50/126 (39.7) | 69/261 (24.4) | 0.008 |
| Peak type | 260 (18.6) | 28 (8.5) | 232 (21.7) | < 0.001 |
| Burning | 182 (13.0) | 51 (15.5) | 131 (12.3) | 0.126 |
| Pinching | 81 (5.8) | 7 (2.1) | 74 (6.9) | 0.001 |
| Increasing at position change | 252 (18.0) | 29 (8.8) | 223 (20.9) | < 0.001 |
| Breathing non-related | 1037 (74.2) | 299 (90.9) | 738 (69.0) | < 0.001 |
| Radiating | 592 (42.3) | 179 (54.4) | 413 (38.6) | < 0.001 |
| Additional symptoms | 937 (67.1) | 249 (73.7) | 688 (64.4) | < 0.001 |

**ACS** Acute coronary syndrome, **NRS** Numeric rating scale, **IQR** Interquartile range

\(^a\)Typical chest pain is characterized by a retrosternal sensation of pressure or heaviness ("angina"), which may be intermittent (usually lasting several minutes) or persistent
up is estimated to reach 20% (undetermined diagnosis), we needed to include 1123 calls for non-traumatic chest pain for each sex. This sample size estimation concerns the training (derivation) dataset, which corresponds to two-thirds of the overall included patients. Thus, we needed to include a total of 1705 men and 1705 women for the derivation and validation datasets. Owing to an expected sex ratio of women/men of 40%/60%, we needed to include 4263 consecutive calls for non-traumatic chest pain.

Analysis
Data are expressed as numbers with percentages for categorical variables or means with standard deviation or medians with interquartile range [IQR] for continuous variables. Categorial data were compared by chi-square or Fisher exact test when appropriate and continuous data by Student t or Mann and Whitney test as appropriate. Inter-expert agreement for the final diagnosis was calculated with the Kappa coefficient and its 95% confidence interval.

Model development
For each sex, we randomly selected a training (derivation) dataset from two-thirds of the data. Potential predictors were identified as variables associated with an ACS diagnosis significant at \( p < 0.2 \) on bivariate analysis or already known to be associated in the literature. We used a backward stepwise logistic regression to retain the final independent predictive variables, based on both \( p < 0.05 \) and the log-likelihood test. Then, we built a receiver operating characteristic (ROC) curve for each sex and defined the area under the ROC curve (AUC).

Model validation
The main characteristics of the derivation and validation models were compared for each sex by using appropriate bivariate statistical tests. The internal validity of the predictive model for each sex was tested in the validation dataset. First the discriminative performance of the score was evaluated with the validation dataset. Then, mean predictive probabilities were plotted against observed proportions of ACS in each quintile of predictive probabilities. Differences between observed and predicted probabilities were tested with the Hosmer-Lemeshow test.

All tests were two-sided, with statistical significance set at \( p < 0.05 \). All analyses were performed with STATA v11.2 (StataCorpLP, College Station, TX, USA) and CART software (Salford System, CA 92126 USA).

Results
Characteristics of study subjects
Over the 18 months of the study, 4205 patients were enrolled. A final diagnosis was established for 3727 (89%) (1630 [44%] women and 2097 [56%] men). Flowcharts of the final diagnosis by sex are presented in Fig. 1. Sensitivity analyses are presented in Appendix. Overall, 647 (17%) participants had an ACS diagnosis (508 [24%] men and 139 [9%] women), including 260 (7%) with STEMI. The inter-expert agreement was excellent (Kappa = 0.91 [0.89–0.93]) for diagnosing ACS.

Main results
Males with ACS
The derivation dataset consisted of 1398 men. In total, 324 (23%) men had an ACS diagnosis, 135/324 (42%) with

**Table 2** Final model for predicting ACS in males after multivariate analysis

| Variables                          | Regression coefficient | OR     | 95% CI          | P-value |
|-----------------------------------|------------------------|--------|-----------------|---------|
| Age, y                            |                        |        |                 |         |
| < 40                              | 0                      | 1      |                 |         |
| 40–50                             | 1.085                  | 2.958  | [1.668–5.246]   | < 0.001 |
| 50–60                             | 1.692                  | 5.431  | [3.125–9.439]   | < 0.001 |
| ≥ 60                              | 1.969                  | 7.166  | [4.162–12.336]  | < 0.001 |
| Tobacco use                       | 0.359                  | 1.432  | [1.064–1.927]   | 0.018   |
| Severe pain (NRS ≥ 6)             | −1.016                 | 0.362  | [0.143–0.917]   | 0.032   |
| Permanent pain                    | 0.385                  | 1.469  | [1.09–1.981]    | 0.012   |
| Breathing non-related pain        | 0.813                  | 2.254  | [1.281–3.967]   | 0.005   |
| Retrosternal pain                 | 0.457                  | 1.580  | [1.203–2.075]   | 0.001   |
| Radiating pain                    | 0.465                  | 1.592  | [1.209–2.097]   | 0.001   |
| Additional symptoms               | 0.066                  | 1.068  | [0.735–1.551]   | 0.729   |
| Severe pain* Breathing non-related| 0.838                  | 2.313  | [1.015–5.271]   | 0.046   |
| Severe pain* Additional symptoms  | 0.850                  | 2.339  | [1.202–4.553]   | 0.012   |

Because of missing values for one man, the analyses were performed on 1397 males

OR Odds ratio, CI Confidence interval, NRS Numeric rating scale
The symbol ‘*’ indicate the interaction
STEMI. The flowchart of the derivation dataset is in the Additional file 2. The general characteristics and type of pain for males are in Table 1. Men with an ACS diagnosis were older, more frequently had cardiovascular risk factors (except family history of coronary diseases and tobacco use) and had more typical pain typography characteristics than those without a diagnosis (Table 1).

Eight factors mostly contributed to the final model for predicting ACS in males: age, tobacco use, severe and permanent pain; retrosternal, breathing non-related and radiating pain; and additional symptoms (Table 2). The AUC value for the final male model was 0.76 (95% CI 0.73–0.79), with no differences between observed and predicted probabilities (Hosmer-Lemeshow test: \( p = 0.78 \)).

General characteristics were well balanced between the derivation dataset (\( n = 1398 \)) and the validation dataset (\( n = 699 \)). The accuracy of the male model to predict ACS was validated, with no differences between observed and predicted probabilities (Hosmer-Lemeshow test: \( p = 0.554 \), Fig. 2a). The AUC value for the male prediction score was 0.76 (0.73–0.80).

**Females with ACS**
The derivation dataset consisted of 1087 women. Overall, 92 (8%) women had an ACS diagnosis, 32/92 (35%) with STEMI. The flowchart of the derivation dataset is in the Additional file 2. The general characteristics and type of pain for females are in Table 3. Women with an ACS diagnosis were older, with more cardiovascular risk factors (except family history of coronary diseases and tobacco use) and had more typical pain typography than those without a diagnosis (Table 3).

Four factors mostly contributed to the final model for predicting ACS in females: age \( \geq 60 \) years,
personal history of coronary artery disease, and breathing non-related and radiating pain (Table 4). The AUC value for the final female model was 0.79 (95% CI: 0.75–0.83), with no differences between observed and predicted probabilities (Hosmer-Lemeshow test: $p = 0.70$).

General characteristics were well balanced between the derivation dataset ($n = 1087$) and the validation dataset ($n = 995$).

### Table 3 Characteristics and type of pain for female patients in the derivation set with and without a diagnosis of ACS

|                         | Total ($n = 1087$) | ACS ($n = 92$) | No ACS ($n = 995$) | $p$ value |
|-------------------------|--------------------|---------------|-------------------|-----------|
| Age, mean (SD), y, No. (%) | 55.2 (18.1)        | 69.7 (13.7)   | 53.9 (17.9)       | < 0.001   |
| < 60                    | 215 (19.8)         | 9 (9.8)       | 206 (20.7)        | < 0.001   |
| ≥ 60                    | 445 (40.9)         | 75 (81.5)     | 370 (37.2)        |           |
| Coexisting conditions, No. (%) |                   |               |                   |           |
| Hypertension            | 382 (35.1)         | 49 (53.3)     | 333 (33.5)        | < 0.001   |
| Personal coronary artery disease | 179 (16.5)        | 35 (38.0)     | 144 (14.5)        | < 0.001   |
| Family coronary artery disease | 178 (16.4)        | 10 (10.9)     | 168 (16.9)        | 0.136     |
| Tobacco use             | 249 (22.9)         | 12 (13.0)     | 237 (23.8)        | 0.019     |
| Diabetes                | 130 (12.0)         | 17 (18.5)     | 113 (11.4)        | 0.044     |
| Dyslipidaemia           | 207 (19.0)         | 28 (30.4)     | 179 (18.0)        | 0.004     |
| Medication therapy, No. (%) |                   |               |                   |           |
| Aspirin                 | 129 (11.9)         | 24 (26.1)     | 105 (10.6)        | < 0.001   |
| Clopidogrel             | 64 (5.9)           | 16 (17.4)     | 48 (4.8)          | < 0.001   |
| Thyroid hormone         | 90 (8.3)           | 7 (7.6)       | 83 (8.3)          | 0.807     |
| Statin                  | 132 (12.1)         | 20 (21.7)     | 112 (11.3)        | 0.003     |
| Severity of pain, median (IQR) | 5 [4–7]          | 5 [4–7]       | 5 [4–7]           | 0.496     |
| Type of pain, No. (%)   |                    |               |                   |           |
| Severe pain (NRS ≥6)    | 489 (45.0)         | 42 (45.7)     | 447 (45.0)        | 0.900     |
| Permanent               | 272 (25.0)         | 24 (26.1)     | 248 (24.9)        | 0.805     |
| Pain onset:             |                    |               |                   | 0.620     |
| Crescendo               | 207 (19.1)         | 14 (15.2)     | 193 (19.4)        |           |
| Abruptly                | 483 (44.4)         | 43 (46.7)     | 440 (44.2)        |           |
| Unclassified            | 397 (36.5)         | 35 (38.1)     | 362 (36.4)        |           |
| Circumstance of pain, No. (%) |                 |               |                   | 0.210     |
| At rest                 | 914 (84.1)         | 75 (81.5)     | 839 (84.3)        |           |
| Sport or stress-related | 81 (7.5)           | 5 (5.4)       | 76 (7.6)          |           |
| Unclassified            | 92 (8.5)           | 12 (13.1)     | 80 (8.0)          |           |
| Pain typography, No. (%) |                    |               |                   |           |
| Typical chest pain*     | 616 (56.7)         | 64 (69.6)     | 552 (55.5)        | 0.009     |
| Retrosternal            | 461 (42.4)         | 52 (56.5)     | 409 (41.1)        | 0.004     |
| Post-myocardial infarction angina | 41/179 (22.9) | 9/35 (25.7) | 32/144 (22.2) | 0.659 |
| Peak type               | 216 (19.9)         | 6 (6.5)       | 210 (21.1)        | 0.001     |
| Burning                 | 132 (12.1)         | 14 (15.2)     | 118 (11.9)        | 0.345     |
| Pinching                | 53 (4.9)           | 1 (1.1)       | 52 (5.2)          | 0.122     |
| Increasing at position change | 229 (21.1) | 5 (5.4)       | 224 (22.5)        | < 0.001   |
| Breathing non-related   | 769 (70.7)         | 83 (90.2)     | 686 (68.9)        | < 0.001   |
| Radiating              | 475 (43.7)         | 52 (56.5)     | 423 (42.5)        | 0.010     |
| Additional symptoms    | 737 (67.9)         | 70 (76.1)     | 667 (67.1)        | 0.077     |

ACS Acute coronary syndrome, NRS Numeric rating scale, IQR Interquartile range

*Typical chest pain is characterized by a retrosternal sensation of pressure or heaviness ("angina"), which may be intermittent (usually lasting several minutes) or persistent.
dataset \( (n = 543) \). The female model’s accuracy to predict ACS was not validated: predicted probabilities significantly differed from observed values (Hosmer-Lemeshow test: \( p = 0.035 \), Fig. 2b). The AUC value for the female prediction score was 0.67 (0.60–0.74).

**Discussion**

**Main results**

For adults calling an EMCC with chest pain or discomfort, predictors of a final ACS diagnosis differed by sex. The discriminative performance of the model was poor for women and good for men.

**Explanation of the findings**

In our study, a predictive variable for ACS in males agreed with traditional typical angina. The pain characteristics were so typical that even the coexisting conditions, such previous coronary artery disease, did not significantly add to the prediction in the multivariate model. Therefore, decision-making in men is based on the characteristics of pain. For females, except for age and personal history of ACS, factors were not related to typical angina. Thus, decision-making in women is mainly based on criteria other than the pain characteristics. The initial presentation of ACS is well known to differ by sex. In contrast to men, women complain of discomfort or pain due to pinching or burning [16–20]. These discrepancies could be due to disparities in pathophysiology and aetiologies [26, 27]. Women with an ACS diagnosis were more likely to have a normal or mild angiographic coronary heart disease [19, 21]. One-tenth of ACS cases involve spontaneous coronary artery dissection, mainly in young women [28]. Myocardial infarction with a non-obstructive coronary artery (MINOCA) occurs mostly in women and includes coronary endothelial dysfunction, myocarditis or Takostubo syndrome [1]. In these pathologies, traditional cardiovascular risk factors have a low implication.

The identification of a woman presenting an ACS remains a challenge. The mortality rate is higher in women than men because of their atypical initial presentation, older age, and less recourse to coronary angiography [21, 29, 30]. When adjusted on the same level of care, mortality is similar between the sexes, which led the authors of one study to advocate for a diagnosis of ACS in women [31, 32]. In our study, the discriminative performance of our model to predict ACS in women was not reproducible. Determinants were limited in number, not typical and above all not reproducible. An explanation is a potential lack of knowledge regarding variables to investigate to detect ACS in women. Currently, women are assessed with the variables used for men. Specific factors in women need to be investigated.

**Strengths**

Our study is the first prospective multicentric study to focus on predicting an ACS diagnosis in patients calling an EMCC for chest pain or discomfort. Other studies analysed patients with an established ACS diagnosis and cared for in an emergency department or cardiology department. Furthermore, this large study improves on the small number of studies evaluating the effectiveness of EMCC [33].

**Limitations**

We excluded uncommunicative patients because they could experience impending death, an at-risk sign. This state has already been highlighted in the evaluation of imminent delivery at an EMCC [34]. Thus, we investigated ACS as the only outcome without considering other life threatening causes of chest pain. The outcome was established retrospectively by experts, which could be considered as subjective and potentially biased. Nevertheless, decision was based on medical records that were collected prospectively, limiting this bias. Lastly, in women we failed to propose a model that accurately predicted an ACS diagnosis in the validation sample.

**Conclusions**

A sex disparity exists in screening for ACS in people calling an EMCC because of chest pain. A score could be proposed for men. For women, a better understanding of pathophysiology and symptomatology are needed to increase the detection of ACS.

**Table 4** Final model for predicting ACS in males after multivariate analysis

| Variables                                      | Regression coefficient | OR   | 95% CI          | \( p \) value |
|------------------------------------------------|------------------------|------|-----------------|--------------|
| Age \( \geq 60 \) y                            | 1.716                  | 5.564| [3.160–9.800]   | < 0.001      |
| Personal history of coronary artery disease    | 0.603                  | 1.828| [1.120–2.982]   | 0.016        |
| Breathing non-related pain                     | 1.017                  | 2.765| [1.346–5.678]   | 0.006        |
| Radiating pain                                 | 0.469                  | 1.598| [1.017–2.513]   | 0.042        |

Because of missing values for one woman, the analyses were performed on 1086 females

OR Odds ratio, CI Confidence interval
### Table 5: Characteristics and type of pain by availability of diagnosis

| Characteristic                          | Diagnosis unknown (n = 478) | Diagnosis established (n = 3727) | P value |
|----------------------------------------|-----------------------------|---------------------------------|---------|
| Female, No. (%)                        | 212 (44.4)                  | 1630 (43.7)                     | 0.789   |
| Age, mean (SD), y, No. (%)             |                             |                                 |         |
| <40                                    | 147 (30.8)                  | 764 (20.5)                      | <0.001  |
| 40-50                                   | 133 (27.8)                  | 804 (21.6)                      |         |
| 50-60                                   | 95 (19.9)                   | 790 (21.2)                      |         |
| ≥60                                     | 103 (21.5)                  | 1369 (36.7)                     |         |
| Coexisting conditions, No. (%)         |                             |                                 |         |
| Hypertension                           | 84 (17.6)                   | 1181 (31.7)                     | <0.001  |
| Personal coronary artery disease        | 71 (14.9)                   | 857 (23.0)                      | <0.001  |
| Family coronary artery disease          | 82 (17.2)                   | 603 (16.2)                      | 0.587   |
| Tobacco use                            | 220 (46.0)                  | 1244 (33.4)                     | <0.001  |
| Diabetes                               | 20 (4.2)                    | 462 (12.4)                      | <0.001  |
| Dyslipidaemia                          | 77 (16.1)                   | 838 (22.5)                      | 0.001   |
| Medication therapy, No. (%)            |                             |                                 |         |
| Aspirin                                | 55 (11.5)                   | 624 (16.7)                      | 0.003   |
| Clopidogrel                            | 31 (6.5)                    | 410 (11.0)                      | 0.002   |
| Thyroid hormone                        | 20 (4.2)                    | 152 (4.1)                       | 0.912   |
| Statin                                 | 40 (8.4)                    | 589 (15.8)                      | <0.001  |
| Severity of pain, median (IQR)         | 5 [3–6]                     | 5 [3–7]                         | 0.032   |
| Type of pain, No. (%)                  |                             |                                 |         |
| Severe pain (NRS ≥6)                   | 177 (37.1)                  | 1616 (43.4)                     | 0.009   |
| Permanent                              | 112 (23.4)                  | 947 (25.4)                      | 0.348   |
| Pain onset:                             |                             |                                 | 0.057   |
| Crescendo                              | 113 (23.6)                  | 782 (21.0)                      |         |
| Abruptly                               | 179 (37.4)                  | 1608 (43.1)                     |         |
| Unclassified                           | 186 (38.9)                  | 1337 (35.9)                     |         |
| Circumstance of call, No. (%)          |                             |                                 | 0.733   |
| At rest                                | 382 (79.9)                  | 3026 (81.2)                     |         |
| Sport or stress related                | 52 (10.9)                   | 395 (10.6)                      |         |
| Unclassified                           | 44 (9.2)                    | 306 (8.2)                       |         |
| Pain typography, No. (%)               |                             |                                 |         |
| Typical chest pain*                    | 231 (48.3)                  | 2121 (56.9)                     | <0.001  |
| Retrosternal                           | 170 (35.6)                  | 1721 (46.2)                     |         |
| Post-myocardial infarction angina      | 147 (19.7)                  | 241/857 (28.1)                  | 0.127   |
| Peak type                              | 143 (29.9)                  | 701 (18.8)                      | <0.001  |
| Burning                                | 59 (12.3)                   | 478 (12.8)                      | 0.766   |
| Pinching                               | 19 (4.0)                    | 204 (5.5)                       | 0.169   |
| Increasing at position change          | 119 (24.9)                  | 726 (19.5)                      | 0.005   |
| Breathing non-related                  | 311 (65.1)                  | 2738 (73.5)                     | <0.001  |
| Radiating                              | 198 (41.4)                  | 1637 (43.9)                     | 0.299   |
| Additional symptoms                    | 302 (63.3)                  | 2521 (67.7)                     | 0.056   |

NRS: Numeric rating scale

*Typical chest pain is characterized by a retrosternal sensation of pressure or heaviness ('angina'), which may be intermittent (usually lasting several minutes) or persistent.
Supplementary information

Supplementary information accompanies this paper at https://doi.org/10.1186/s13049-019-0670-y.

Additional file 1. Standardized form for gathering data on patient characteristics, cardiovascular risk factors, medical history and clinical presentation.

Additional file 2. Flowchart in the derivation set by sex.

Abbreviations

ACSM: American College of Sports Medicine; ACS: Acute Coronary Syndrome; AUC: Area Under the ROC Curve; ECG: Electrocardiography; EMCC: Emergency Medical Communication Centre; IQR: Interquartile Range; MICU: Mobile Intensive Care Unit; MINOCA: Myocardial infarction with a non-obstructive coronary artery; ROC: Receiver Operating Characteristic; STEMI: ST Elevation Myocardial Infarction

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Transparency declaration

The lead authors (the manuscript’s guarantors) affirm 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.

Authors’ contributions

Conception and design: SHYK, VB, JLD and SC; acquisition of data: CP, AB, ET, XC and SC; analysis: SHYK and VB; interpretation of data: PGR, SHYK, VB and SC; drafting the article: PGR and SC; critical revision for important intellectual content: all authors; final approval of the version to be published: all authors.

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Availability of data and materials

The statistical code and technical processes are available from the time of publication. Appropriate institutional agreements will be required for anonymised participant data transfer. Requests should be made via email to the corresponding author along with an analysis proposal.

Ethics approval

In accordance with French law, our local ethics committee considered that patient oral consent could be sufficient for participation in this observational study. Data collection and storage were approved by the “Comité consultatif sur le traitement de l’information en matière de recherche” (CCTRIS) and the “Commission nationale informatique et liberté” (CNIL). This study is registered with ClinicalTrials.gov, number NCT02042209.

Competing interests

All authors have completed the Unified Competing Interest form (available on request from the corresponding author) and declare: no support from any organisation for the submitted work; no financial relationships with any organisations that might have an interest in the submitted work in the previous 3 years; no other relationships or activities that could appear to have influenced the submitted work.

Author details

1Emergency Department, Toulouse University Hospital, 31000 Toulouse, France. 2UMR 1027, Paul Sabatier University Toulouse III, Inserm, Toulouse, France. 3SAMU 92, Assistance Publique-Hôpitaux de Paris, Hôpital Raymond Poincaré, 92380 Garches, France. 4SAMU 33, CHU de Bordeaux, 33076 Bordeaux cedex, France. 5Unité de Soutien Méthodologique à la Recherche (USMR), Centre Hospitalier Universitaire de Toulouse (CHU de Toulouse), Toulouse, France. 6Department of Cardiology, Rangueil University Hospital, Toulouse, France. 7Department of Emergency, CHU de la Réunion, allée des Topazes, Université de la Réunion, 97400 Saint Denis, France.

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Supplementary information

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