Specific prodromal symptoms in patients with acute coronary syndrome

Mehdi Heidarzadeh1 | Shahla Elyaszadeh2 | Behrouz Dadkhah1 | Hossein Doustkami3

Aims: Assessing the prodromal cardiac symptoms in patients with acute coronary syndrome (ACS) and compare them with healthy population.

Background: Identifying specific prodromal cardiac symptoms can play an important role in screening people at risk.

Design: A comparative study of prodromal symptoms.

Methods: In this comparison study, an 80-item checklist of prodromal symptoms was designed and completed by 337 participants in three groups (Patient group I, Patient group II and Healthy group). All participants were studied over a period of six months (from May to October 2017). SPSS-15 software was used to analyse the data.

Results: The symptoms of pain/discomfort in chest, pain/discomfort centred in the superior part of chest, pain/discomfort in the left breast and numbness or burning of both arms were the most important symptoms to predict ACS incidence in the two patient groups (odds ratio > 4 and p ≤ .05).

KEYWORDS
acute coronary syndrome, ischemic heart diseases, nurses, nursing, prodromal symptoms

1 | INTRODUCTION

Cardiovascular diseases are the world’s leading cause of death (World Health Organization, [WHO], 2017), with estimated annual mortality rate of 17.3 million people, which is expected to exceed 23.6 million people by 2030 (Mozaffarian et al., 2015). Acute coronary syndrome (ACS) as a subset of cardiovascular diseases often is accompanied by clinical symptoms such as chest pain or discomfort, shortness of breath and fatigue (Kasper et al., 2015). Nurses can play an important role in reducing mortality and irreparable complications by early diagnosis of ACS and timely therapeutic and diagnostic measures (Canto et al., 2014; McSweeney et al., 2014, 2017). Getting know about “prodromal symptoms” can help nurses timely identify people at risk with minimum damage. In fact, prodromal symptoms are warning signs of an impending ACS (McSweeney et al., 2017) and signs of an impending cardiac event that have a new onset or change in severity or frequency before the cardiac event and then disappear or return to its previous levels (McSweeney et al., 2003). Forty-nine to ninety-five per cent of the people at risk experience some undiagnosed symptoms as prodromal symptoms days, weeks, months and even 2 years before the onset of cardiac events (Lee et al., 2020; O’Keefe-McCarthty et al., 2019; Soltani et al., 2016).

2 | BACKGROUND

O’Keefe-McCarthy and Ready (2016) investigated the prevalence of heart disease prodromal symptoms in a review study and reported...
anxiety, unusual fatigue, pain or discomfort in arms etc. However, McSweeney et al. (2017) showed that the most common prodromal symptoms are not necessarily the most predictive symptoms of cardiac events. Despite the importance of the study by McSweeney et al. (2017), they examined the predictive properties of prodromal symptoms of cardiac events only among females and there is no information available on the specific prodromal symptoms among males. In general, not many studies have examined the specific prodromal symptoms in both genders.

Also, many factors can affect the predictive power of each of the prodromal symptoms. For example, previous studies have shown prodromal symptom in patients with ACS, but these symptoms may be related to other factors such as demographic characteristics and underlying diseases (Elyaszadeh et al., 2018). Therefore, it is important to evaluate these symptoms in other population especially in people without ACS experience.

The present study has two research questions as follows: (a) What are the prevalence of prodromal symptoms in patients (men and women) with ACS? (b) In compared with people without ACS (healthy population), which of these prodromal symptoms are specific for patients with ACS?

3 | METHODS

3.1 | Design

This comparison study was conducted to assess the prodromal cardiac symptoms in ACS patients in Imam Khomeini Hospital, Ardabil, Iran.

3.2 | Participants

The 337 participants were divided into three groups: two patient groups diagnosed with ACS by a cardiologist at CCU and cardiac wards including 111 patients without a history of IHD as Patient group I and 107 with a history of myocardial infarction or angina pectoris as Patient group II. The third group included 119 patients with no ACS presenting to the heart clinics for heart checkup and were diagnosed as healthy after cardiac assessment by a cardiologist (healthy group). The inclusion criteria for Patient groups I and II were as follows: (a) definitive diagnosis of ACS by a cardiologist; (b) ability to speak Persian language; (c) normal cognitive status (according to medical history and clinical records); (d) favourable and stable physical conditions for conducting interviews; and (e) no history of IHD in Patient group I and a history of IHD in patient Group II. The inclusion criteria for the Healthy group were as follows: (a) definitive diagnosis of ACS by a cardiologist; (b) ability to speak Persian language; (c) normal cognitive status (according to medical history and clinical records); (d) favourable and stable physical conditions for conducting interviews; and (e) no history of underlying diseases (Elyaszadeh et al., 2018). Therefore, it is important to evaluate these symptoms in other population especially in people without ACS experience.

The present study has two research questions as follows: (a) What are the prevalence of prodromal symptoms in patients (men and women) with ACS? (b) In compared with people without ACS (healthy population), which of these prodromal symptoms are specific for patients with ACS?

3.3 | Instrument and data collection

Data were collected with a demographic questionnaire and prodromal symptoms questionnaire. The prodromal symptoms questionnaire was designed in two phases. In the first step, a review of printed and electronic literature yielded 73 prodromal symptoms among ACS patients (Bahr et al., 2000; Bonow et al., 2011; Cole et al., 2012; Graham et al., 2008; Hofgren et al., 1995; Levlen et al., 2009; McSweeney et al., 2003; McSweeney & Crane, 2000; Milner et al., 2001; Norris et al., 2008; O’Keefe-McCarthy & Guo, 2016; Ottolini et al., 2005).

In the second step, for developing the questionnaire, a qualitative assessment was performed.

In the qualitative phase, 30 patients with ACS (15 men & 15 women) after receiving enough information about the study goal filled the 73-item questionnaire prepared in the first step. Then, they were asked to express their experience of other symptoms before the incidence of ACS which they believed were related to their heart attack but were not listed in the 73-item list. During analyses of their statements, seven new prodromal symptoms were extracted including tingling toes, hot flashes, a feeling of heat on the foot, dry mouth, foot pain, numbness toes and knee pain. At the end of this part, an 80-item questionnaire was developed.

In addition to examining the incidence of 80 symptoms, this questionnaire assessed their severity and frequency. For this purpose, the mean score of each symptom was scored based on the incidence, severity and frequency and ranged from 0–10 for each item based on incidence (0 and 1 for No and Yes, respectively), intensity (1, 2 and 3 for mild, moderate and severe, respectively) and frequency (6, 5, 4, 3, 2 and 1 for daily, several times a week, once a week, 2–3 times a month, once a month and less than once a month, respectively). High score of each item shows more severity and frequency of the symptom. This way of scoring has been introduced by O’Keefe-McCarthy and Guo (2016). For both patient Groups I and II, items were asked retrospectively with a focus on the past 3 months. Meanwhile, the subjects in the healthy group were monitored prospectively for 3 months after sampling in terms of ACS incidence and after the three-month follow-up, none of the participants in the healthy group had ACS.

The content validity index (CVI) and content validity ratio (CVR) were used to determine the content validity of the questionnaire. The questionnaire was distributed to five nursing faculty members and five expert clinicians in the cardiovascular field. The CVI and CVR scores for the total scale were determined as 0.8 and 0.79, respectively. To determine the internal consistency reliability, the Cronbach’s alpha coefficient was calculated 0.8 among 30 patients with ACS presenting.

All the participants were interviewed to complete the questionnaires in bedside (at CCU and cardiac wards) or inside the nursing room (at heart clinic). Every interview and completing the questionnaires lasted approximately 20–30 min in all groups.
3.4 | Statistical analyses

Data were analysed using descriptive and inferential statistics. Analysis of variance (ANOVA) with post hoc Tukey test was used for determining any significant differences between the scores of every symptom in the groups. Also, logistic regression was used to calculate odds ratios at a 95% confidence interval for the association between the presence/absence of each of the prodromal symptoms with ACS incidence (and lack of it in healthy population) as a dependent variable and each symptom tested separately. All the symptoms were analysed twice; first, we considered group I against the healthy group as a dependent variable; then, we did it in group II and healthy group in the same way. Data were analysed in SPSS-15 software.

3.5 | Ethical considerations

This study was approved by the Ethics Committee of the Ardabil University of Medical Sciences. One of the researchers (first author) was referred to the clinics and invited eligible participants to the study (May–October in 2017). Then, willing participants were also asked to answer questionnaire by interview. Before completing the questionnaires, participants were briefed on the goals and method of the study before signing a written consent. Patients were assured that the study would not disturb their treatment process, the collected information would be completely confidential and the results would be published collectively.

4 | RESULTS

4.1 | Sample description

A total of 337 subjects were recruited in three groups of Healthy (N = 119), Patient I (N = 111) and Patient II (N = 107) for data analysis. Healthy group did not differ significantly with any of the Patient groups I and II in terms of age, sex and underlying diseases (p > .05). Demographic characteristics and medical history of the groups are presented in Table 1.

In this study, 80 prodromal symptoms were analysed in terms of frequency and mean score in three groups. By comparing the results (frequency and mean score of prodromal symptoms) in patient groups with healthy group, the true value of each symptom became clear.

| Parameter | Healthy group | Patient group I | Patient group II | p-value |
|-----------|---------------|-----------------|-----------------|---------|
| Mean Age, year (SD) | 63.15 (11.34) | 62.14 (10.72) | 64.75 (10.45) | .206a |
| Gender | | | | |
| Female | 58 (48.7) | 50 (45) | 51 (47.7) | .848b |
| Male | 61 (51.3) | 61 (55) | 56 (52.3) | |
| ACS | | | | |
| Myocardial infarction | — | 59 (53.2) | 45 (42.1) | — |
| Unstable angina | — | 52 (46.8) | 62 (57.9) | |
| Medical history | | | | |
| Hypertension | 83 (69.7) | 73 (65.8) | 84 (78.5) | .105b |
| Diabetes | 31 (26.1) | 29 (26.1) | 33 (30.8) | .662b |
| Hyperlipidemia | 40 (33.6) | 37 (33.3) | 41 (38.3) | .688b |
| Low back pain | 18 (15.1) | 11 (9.9) | 14 (13.1) | .492b |
| Rheumatoid arthritis | 27 (22.7) | 20 (18) | 18 (16.8) | .492b |
| Thyroid condition | 9 (7.6) | 7 (6.3) | 6 (5.6) | .833b |
| IBS, Peptic Ulcer, Oesophageal Reflux | 4 (3.4) | 3 (2.7) | 3 (2.8) | .951b |
| COPD, Asthma | 3 (2.5) | 3 (2.7) | 6 (5.6) | .408b |
| Sinusitis, migraine | 6 (5.6) | 5 (4.5) | 3 (2.5) | .480b |
| BPH | 4 (6.6) | 3 (4.9) | 6 (10.7) | .478b |
| Other | 12 (10.1) | 11 (9.9) | 13 (12.1) | .837b |

aUsing ANOVA. 
bUsing chi-square tests; other: depression, anaemia, cerebrovascular accident, epilepsy, cholecystitis, fatty liver disease and a cold. Patient group I: participants with ACS incidence and without a history of ischaemic heart disease, patient group II: participants with ACS incidence and with a history of ischaemic heart disease; healthy group: participants without ACS incidence; ACS, acute coronary syndrome; IBS, irritable bowel syndrome; COPD, chronic obstructive pulmonary disease; BPH, benign prostatic hyperplasia; SD, standard deviation.

TABLE 1 Demographic characteristics and medical history of the participants by groups
## Table 2: Prevalence and mean score of each prodromal symptom by groups and significant association of the prodromal symptom with incidence of ACS

| Symptom                              | Frequency | Score (Mean±SD) | Significance |
|--------------------------------------|-----------|----------------|--------------|
|                                      | n (%      | Patient group I (N = 111) | Patient group II (N = 107) | Healthy group (N = 119) | Patient group I (N = 111) | Patient group II (N = 107) | Healthy group (N = 119) |
| Pain/discomfort in chest             | 66 (59.5)*** | 4.55 (4.02)a | 6.78 (3.58)a | 0.82 (2.19) |
| Pain/discomfort centred in the superior part of chest | 51 (45.9)*** | 3.52 (4.01)a | 5.06 (4.23)a | 0.80 (2.18) |
| Pain/discomfort in left breast       | 32 (28.8)*** | 2.29 (3.75)a | 3.01 (4.05)a | 0.44 (1.60) |
| Pain/discomfort centred in left part of chest | 19 (17.1)*** | 1.28 (2.93)a | 2.60 (3.89)a | 0.14 (1.02) |
| Vomiting                             | 22 (19.8)*** | 1.43 (3.01)a | 0.77 (2.00) | 0.33 (1.37) |
| Numbness or burning of both arms     | 16 (14.4)*** | 1.08 (2.77)a | 1.65 (3.44)a | 0.25 (1.42) |
| Pain/discomfort in back, between/under shoulder blades | 68 (61.3)** | 4.98 (4.18)a | 5.46 (4.20)a | 1.97 (3.24) |
| Pain/discomfort in both arms         | 43 (38.7)** | 3.29 (4.30)a | 3.53 (4.39)a | 1.36 (3.05) |
| Pain/discomfort in neck/throat       | 54 (48.6)** | 3.85 (4.16)a | 4.23 (4.17)a | 2.31 (3.64) |
| Diaphoresis                          | 69 (62.2)** | 5.18 (4.28)a | 6.05 (4.22)a | 3.76 (4.40) |
| Shortness of breath                  | 52 (46.8)** | 3.78 (4.23)a | 5.18 (4.38)a | 2.05 (3.37) |
| Arms ache                             | 33 (29.7)** | 2.60 (4.07)a | 3.42 (4.51)a | 1.02 (2.76) |
| Difficulty breathing at night         | 30 (27)** | 2.19 (3.76)a | 3.04 (4.27)a | 0.70 (2.06) |
| Nausea                               | 28 (25.2)** | 1.91 (3.43)a | 1.63 (3.10) | 0.85 (2.44) |
| Orthopnea                            | 28 (25.2)** | 2.05 (3.63)a | 3.24 (4.31)a | 0.76 (2.04) |
| Pain/discomfort in jaw/teeth         | 21 (18.9)** | 1.49 (3.20) | 2.28 (3.70)a | 0.63 (2.08) |
| Panic                                 | 10 (9)** | 0.55 (1.87) | 0.77 (2.35)a | 0.16 (1.10) |

(Continues)
| Symptom                                      | Frequency n (%) | Score Mean(SD) |
|----------------------------------------------|-----------------|----------------|
|                                              | Patient group I (N = 111) | Patient group II (N = 107) | Healthy group (N = 119) |
| Headache frequency change (Increased frequency of headaches) | 28 (25.2)**     | 33 (30.8)**     | 17 (14.3)    | 2.14 (3.80) | 2.71 (4.13)*  | 1.04 (2.65) |
| Headache intensity change (Increased intensity of headaches) | 28 (25.2)**     | 22 (20.6)**     | 13 (10.9)    | 2.08 (3.69)*  | 1.92 (3.80)*  | 0.86 (2.51) |
| Sleep disturbance                            | 62 (55.9)*      | 73 (68.2)**     | 50 (42)      | 4.75 (4.42)*  | 5.95 (4.28)*  | 3.19 (3.94) |
| Arms weak/heavy                              | 49 (44.1)**     | 53 (49.5)       | 28 (23.5)    | 3.82 (4.38)*  | 4.36 (4.53)*  | 1.86 (3.49) |
| Pain/discomfort at top of shoulders          | 46 (41.4)       | 58 (54.2)**     | 32 (26.9)    | 3.46 (4.28)*  | 4.49 (4.35)*  | 2.03 (3.48) |
| Frequent indigestion                         | 42 (37.8)*      | 47 (43.9)**     | 29 (24.4)    | 3.04 (4.10)   | 3.77 (4.43)*  | 1.87 (3.41) |
| A feeling of fatigue at waking up            | 43 (38.7)*      | 40 (37.4)       | 30 (25.2)    | 3.30 (4.27)*  | 3.28 (4.32)*  | 1.98 (3.50) |
| Unusual fatigue                              | 60 (54.1)       | 69 (64.5)**     | 55 (46.2)    | 4.56 (4.41)   | 5.69 (4.38)*  | 3.71 (4.17) |
| Heart racing                                 | 51 (45.9)       | 68 (63.9)**     | 43 (36.1)    | 3.65 (4.10)*  | 5.20 (4.20)*  | 2.39 (3.42) |
| Waking up repeatedly during the night        | 46 (41.4)       | 63 (58.9)**     | 46 (38.7)    | 3.51 (4.32)   | 5.27 (4.55)*  | 3.09 (4.03) |
| Dizziness                                    | 45 (40.5)       | 58 (54.2)**     | 43 (36.1)    | 2.85 (3.71)   | 4.30 (4.15)*  | 2.49 (3.47) |
| Insomnia                                     | 47 (42.3)       | 55 (51.4)       | 39 (32.8)    | 3.67 (4.43)   | 4.66 (4.67)*  | 2.50 (3.78) |
| Trouble falling asleep                       | 45 (40.5)       | 56 (52.3)**     | 37 (31.1)    | 3.57 (4.43)   | 4.74 (4.65)*  | 2.50 (3.87) |
| Loss of appetite                              | 32 (28.8)       | 45 (42.1)**     | 23 (19.3)    | 2.49 (4.03)   | 3.82 (4.60)*  | 1.59 (3.55) |
| Numbness or burning of fingers on both hands | 31 (27.9)       | 41 (38.3)**     | 21 (17.6)    | 2.22 (3.70)   | 3.07 (4.09)*  | 1.34 (2.97) |
| Hands/arms tingling                          | 28 (25.2)       | 42 (39.3)**     | 27 (22.7)    | 2.09 (3.61)   | 3.13 (7.05)*  | 1.70 (3.29) |
| Confusion                                    | 8 (7.2)         | 15 (14)*        | 6 (5)        | 0.42 (1.62)   | 1.05 (2.70)*  | 0.26 (1.17) |
| Agoraphobia                                  | 8 (7.2)         | 10 (9.3)**      | 2 (1.7)      | 0.48 (1.81)   | 0.78 (2.51)*  | 0.15 (1.17) |
| Headaches                                    | 48 (43.2)       | 61 (57)         | 50 (42)      | 3.24 (3.97)   | 4.35 (4.05)*  | 2.79 (3.54) |
| Heartburn                                    | 48 (43.2)       | 48 (44.9)       | 38 (31.9)    | 3.40 (4.16)   | 3.60 (4.19)   | 2.26 (3.45) |
| Sputum                                       | 43 (38.7)       | 48 (44.9)       | 36 (30.3)    | 3.08 (4.07)   | 3.83 (4.37)*  | 2.13 (3.47) |
| Symptom                                  | Frequency n (%) | Score Mean (SD) | Patient group I (N = 111) | Patient group II (N = 107) | Healthy group (N = 119) |
|------------------------------------------|----------------|----------------|--------------------------|---------------------------|------------------------|
| **Cough**                                | 35 (31.5)      | 2.47 (3.77)    | 3.07 (4.13)              | 1.75 (3.18)               |
| **Unusually located aches and/or pains** | 69 (62.2)      | 5 (4.12)       | 5.13 (4.45)              | 4.4 (4.35)                |
| **Nervousness**                          | 61 (55)        | 4.49 (4.27)    | 4.34 (4.27)              | 4.31 (3.89)               |
| **Anxiety**                              | 58 (52.3)      | 4.26 (4.29)    | 4.72 (4.26)              | 3.96 (3.87)               |
| **Stress**                               | 49 (44.1)      | 3.78 (4.40)    | 3.34 (4.37)              | 3.44 (4.07)               |
| **Changes in thinking or remembering**  | 48 (43.2)      | 3.33 (4.07)    | 3.55 (4.30)              | 3.38 (3.81)               |
| **Irritation**                           | 47 (42.3)      | 3.48 (4.24)    | 3.24 (4.33)              | 2.38 (3.78)               |
| **Pain in back (backache)**              | 47 (42.3)      | 3.28 (4.10)    | 3.50 (4.41)              | 3.86 (4.20)               |
| **Gastric reflux**                       | 45 (40.5)      | 3.08 (3.96)    | 2.99 (3.82)              | 2.59 (3.53)               |
| **Apprehension**                         | 44 (39.6)      | 3.09 (4.01)    | 2.88 (4.16)              | 2.93 (3.91)               |
| **Weakness**                             | 43 (38.7)      | 3.09 (4.05)    | 3.91 (4.38)              | 2.55 (3.63)               |
| **General malaise**                      | 40 (36)        | 2.80 (3.89)    | 2.81 (4.21)              | 2.56 (3.77)               |
| **Pain/ discomfort in legs**             | 40 (36)        | 3.13 (4.32)    | 2.92 (4.29)              | 4.29 (4.44)               |
| **Hot flashes**                          | 40 (36)        | 2.97 (4.14)    | 3.25 (4.29)              | 2.78 (4.07)               |
| **Flatulence**                           | 35 (31.5)      | 2.70 (4.08)    | 3.40 (4.35)              | 2.85 (3.79)               |
| **Vision change**                        | 34 (30.6)      | 2.39 (3.72)    | 3.03 (3.86)              | 2.26 (3.44)               |
| **Demoralization**                       | 28 (25.2)      | 2.08 (3.71)    | 2.83 (4.13)              | 2.24 (3.58)               |
| **A feeling of hopelessness**            | 27 (24.3)      | 1.99 (3.63)    | 1.56 (3.41)              | 1.52 (3.25)               |
| **Depression**                           | 26 (23.4)      | 1.94 (3.57)    | 2.29 (3.81)              | 1.91 (3.37)               |
| **Pain/ discomfort in left arm or shoulder** | 22 (19.8) | 1.52 (3.14)    | 1.44 (3.28)              | 0.84 (2.43)               |
| **Knee (s) pain**                        | 19 (17.1)      | 2.97 (4.14)    | 0.88 (2.76)              | 1.23 (3.10)               |
| **Wanting to be dead at times**          | 18 (16.2)      | 1.20 (2.93)    | 1.30 (3.12)              | 0.77 (2.44)               |
| **Hand (s)/arm (s) oedema**              | 17 (15.3)      | 1.05 (2.72)    | 1.34 (3.13)              | 0.79 (2.47)               |
| **Dry mouth**                            | 17 (15.3)      | 1.29 (3.09)    | 0.81 (2.50)              | 1.08 (2.80)               |
| Symptom                                      | Frequency n (%) | Patient group I (N = 111) | Patient group II (N = 107) | Healthy group (N = 119) | Score Mean (SD) | Patient group I (N = 111) | Patient group II (N = 107) | Healthy group (N = 119) |
|----------------------------------------------|-----------------|---------------------------|---------------------------|-------------------------|-----------------|---------------------------|---------------------------|-------------------------|
| Not being able to cope with everyday problems as well as before | 17 (15.3)       | 22 (20.6)                 | 16 (13.4)                 | 1.23 (3.01)             | 1.73 (3.49)     | 1.01 (2.66)               |                           |                         |
| Numbness of toes                             | 16 (14.4)       | 18 (16.8)                 | 21 (17.6)                 | 1.18 (2.97)             | 1.52 (3.45)     | 1.68 (3.24)               |                           |                         |
| Tingling toes                                | 16 (14.4)       | 18 (16.8)                 | 27 (22.7)                 | 1.48 (3.30)             | 1.28 (3.15)     | 1.68 (3.24)               |                           |                         |
| A feeling of heat on the foot                | 14 (12.6)       | 15 (14)                   | 32 (26.9)                 | 1.06 (2.89)             | 1.23 (3.15)     | 2.23 (3.80)               |                           |                         |
| Numbness or burning of fingers left hand     | 13 (11.7)       | 13 (12.1)                 | 7 (5.9)                   | 0.87 (2.57)             | 0.91 (2.63)     | 0.15 (1.05)               |                           |                         |
| Pain/ discomfort in the general chest        | 12 (10.8)       | 17 (15.9)                 | 0 (0)                     | 0.86 (2.58)             | 1.36 (3.22)     | 0 (0)                     |                           |                         |
| Increasing difficulty in concentrating on a single subject for long | 9 (8.1)         | 11 (10.3)                 | 11 (9.2)                  | 0.65 (2.23)             | 0.81 (2.49)     | 0.63 (2.10)               |                           |                         |
| Leg/foot oedema                              | 8 (7.2)         | 10 (9.3)                  | 17 (14.3)                 | 0.50 (1.95)             | 0.74 (2.42)     | 1.02 (2.64)               |                           |                         |
| Numbness or burning of fingers right hand    | 7 (6.3)         | 10 (9.3)                  | 8 (6.7)                   | 0.45 (1.87)             | 0.66 (2.27)     | 0.30 (1.50)               |                           |                         |
| Pain/ discomfort in right arm or shoulder    | 7 (6.3)         | 6 (5.6)                   | 19 (16)                   | 0.56 (2.19)             | 0.47 (1.94)     | 1.31 (3.08)               |                           |                         |
| Foot pain                                    | 7 (6.3)         | 4 (3.7)                   | 5 (4.2)                   | 0.58 (2.28)             | 0.30 (1.55)     | 0.38 (1.83)               |                           |                         |
| Abdominal discomfort                         | 6 (5.4)         | 12 (11.2)                 | 15 (12.6)                 | 0.44 (1.90)             | 0.87 (2.55)     | 0.87 (2.35)               |                           |                         |
| Numbness or burning of left arm             | 6 (5.4)         | 7 (6.5)                   | 2 (1.7)                   | 0.39 (1.67)             | 0.50 (2.00)     | 0.05 (0.55)               |                           |                         |
| Numbness or burning of right arm            | 4 (3.6)         | 2 (1.9)                   | 2 (1.7)                   | 0.31 (1.60)             | 0.18 (1.29)     | 0.13 (0.97)               |                           |                         |
| Pain/ discomfort centred right part of the chest | 4 (3.6)     | 2 (1.9)                   | 1 (0.8)                   | 0.25 (1.35)             | 0.16 (1.17)     | 0.05 (0.55)               |                           |                         |
| Syncope                                      | 2 (1.8)         | 6 (5.6)                   | 1 (0.8)                   | 0.08 (0.63)             | 0.23 (1.00)     | 0.07 (0.73)               |                           |                         |
4.2 | Comparing the frequency

Logistic regression was used for investigating the predictive power of the prodromal symptoms. To this end, the frequency of each symptom in the Patient group I and Patient group II was compared with its frequency in the healthy group. Based on the results, the frequency of 24 symptoms in the Patient group I was significantly higher than those of the Healthy group. Furthermore, the Patient group II had higher scores in the same 24 symptoms plus 15 other symptoms (a total of 39 symptoms) compared with the Healthy group (Table 2).

4.3 | Comparing the mean score

For more analyses, significant differences between the mean score of every prodromal symptom in three groups were calculated. ANOVA with post hoc Tukey test showed that the scores of 23 prodromal symptoms (20 of 24 predictive symptoms) in patients group I and 42 (36 of 39 predictive symptoms) in patients group II were significantly higher than healthy group (Table 2).

5 | DISCUSSION

The results of the study showed that 24 specific symptoms predicting ACS both in patients with IHD and patients without IHD, some of which are pain/discomfort in chest, heart racing, diaphoresis, vomiting and shortness of breath. Some other researchers found contradictory results in their study. For example, they have reported vomiting, heart racing and diaphoresis as the non-specific prodromal symptoms (McSweeney et al., 2014; O’Keefe-McCarthy & Guo, 2016). The difference in the characteristics of the participants and data analysis might attribute to this inconsistency. In the most of these studies, only female enrolled and higher probability of occurrence and high frequency of symptoms were the most criteria for determining specific symptoms in these studies. In the present study, both men and women enrolled. In addition, indicators such their higher probability of occurrence compared with the healthy subjects their abundance and mean score of symptoms (severity and frequency) were also calculated.

The results of this study showed that a history of ischaemic heart disease can be effective in the type of symptom experienced and its predictive power. For example, among common predictive symptoms in patient groups, vomiting in people without a history of IHD (patient group I) was higher than in those with a history of IHD (patient group II) and, on the other hand, shortness of breath, discomfort in the arms, difficulty breathing at night, orthopnea, pain/discomfort in jaw/teeth and panic in people with a history of IHD (patient group II) had a higher predictive power than in those without a history of IHD (patient group I). Also, individuals with a history of IHD (patient group II) experienced 15 other symptoms in addition to the 24 common symptoms (Table 2).

Few studies have been conducted on the specific prodromal symptoms with a focus on the history of heart disease. In one of
the few studies in this area, McSweeney et al. (2017) found similar results in a study to identify predictive prodromal symptoms of cardiac events in women without a history of heart diseases. In their study, similar to the present study, the symptoms of discomfort in arms and discomfort in teeth/jaw were significantly related to the incidence of cardiac events and symptoms such as pain in feet, back pain, pain/discomfort in the general chest, vision change, changes in thinking or remembering, anxiety, abdominal discomfort and pain/discomfort centred in the right part of the chest were not related to ACS incidence (McSweeney et al., 2017). Despite these similarities, some results in the present study were contradictory to those of McSweeney's study in that symptoms such as discomfort in arms, difficult breathing at night, shortness of breath, numbness in both arms, weak/heavy feeling in arms, sleep disturbances, frequent dyspepsia, pain/discomfort at top of shoulders, pain/discomfort in neck/throat, headache intensity change, headache frequency change, pain/discomfort centred in the superior part of chest, loss of appetite, cough, dizziness and unusual fatigue were correlated with ACS incidence in the present study, while these symptoms were not significantly correlated with ACS in the study by McSweeney et al. (2017).

We found that the prevalence of a symptom does not indicate the specificity of the symptom, since symptoms such as unusually located aches and/or pains, nervousness, anxiety, changes in thinking or remembering and back pain, although highly prevalent in patient groups, were prevalent in the healthy group, too and their scores were close to those of the patient groups. Due to their nature, these symptoms appear to lack the credibility of ACS prediction. For instance, symptoms of anxiety and unusually located aches and/or pains have several causes, including physiological, situational and psychological factors, in addition to heart status (Elyaszadeh et al., 2018; Faravelli et al., 2013; Kasper et al., 2015). Impending ACS is just one of the factors affecting their experience. Therefore, people who are not at risk of ACS can also experience them due to a variety of other causes and experiencing these symptoms is not specifically related to an impending ACS.

### 5.1 | Limitations

The limitations of the study included the use of convenience sampling, which makes the generalization of the results subject to caution. The gold standard for ruling out coronary artery disease is angiography, but it was not possible to perform it in the healthy group to rule out the coronary artery disease. As a result, according to a cardiologist, we only used diagnostic tests (echocardiogram, exercise treadmill test and electrocardiography) and physical examination; also, subjects were monitored for ACS up to 3 months after sampling. Retrospective design is another limitation which led participants recall recurrent symptoms in previous months and participants experiencing ACS remember symptoms better than who did not. Although we tried to set a time limit for remembering experienced symptoms up to three months ago which can decrease the chance of forgetfulness. On the other hand, researches support the accurate recall of significant medical life events up to 6 years (Githens et al., 1993), it is possible that some details of the symptoms were forgotten.

Another limitation of this study was conducting the research in the community of ACS patients in Ardabil, Iran, which limits the generalizability of the results to other populations, and it is suggested that the specificity of these symptoms be investigated in other populations.

### 6 | CONCLUSION

The present study showed that both patients group I and II had experienced same prodromal symptom, but the mean score of them (severity and frequency) was different. Also, individuals with a history of IHD (patient group II) reported experiencing of 15 more specific symptoms than patient group I.

According to this study, although there are some symptoms experienced in both patients group, the healthy population experienced them, too. It indicates, these symptoms maybe emerge by other causes than ischaemic heart diseases.

### 7 | IMPLICATIONS FOR PRACTICE

The findings of this study have certain applications in nursing practice and research. We tried to assess all possible prodromal symptoms collected by literature review and qualitative interviews and to give evidence for frequency and mean score of all 80 prodromal symptoms. Comparing both frequency and mean score of prodromal symptoms in patient groups with healthy group, provides valuable information about predicting power of prodromal symptoms and determined the specific prodromal symptoms in each group.

This study also showed that the frequency and mean score of prodromal symptoms cannot be appropriate criteria for predicting ACS, so it is necessary to consider the predicting value of the symptoms. Knowing about these specific prodromal symptoms by nurses, other healthcare providers and patients at risk of ACS can be helpful in the early diagnosis of IHD and prevent ACS, so this can lead to timely therapeutic measures and a reduction in irreparable complications and mortality from IHD.

Another application of the findings is that frequency and severity of each specific prodromal symptom in this study can provide well information for researchers to design or develop valid tools for predicting ACS in the people at risk of ACS.

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CONFLICTS OF INTEREST
The authors have no funding or conflicts of interest to disclose.

DATA AVAILABILITY STATEMENT
The data used to support the findings of this study are available from the corresponding author on reasonable request.

ORCID
Shahla Elaysazdeh https://orcid.org/0000-0003-2320-5503

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