Influence of acute coronary syndromes in clinical outcome of patients with acute heart failure

Abstract

Background: A substantial proportion of hospitalized patients with coronary heart disease develop acute heart failure (HF) during the hospital stay, and have a worse prognosis in those patients who develop acute HF at the initial presentation.

Methods: In a retrospective cohort study, we stratified the enrolled acute HF patient population into two groups: Group I: adult patients admitted with a diagnosis of acute HF secondary to acute coronary syndrome (ACS). Group II: adult patients admitted with a diagnosis of acute HF secondary to other causes not related to ACS, during January 2015 to December 2016. We aimed to determine the frequency of cardiovascular complications, triggering causes, clinical profile and cardiovascular risk factors in patients with acute HF secondary to ACS in comparison to those patients without ACS.

Results: We found significant differences in the forms of presentation of acute HF. In the acute HF with ACS cohort the de novo HF was more frequent (19%), while the acute on chronic HF was more frequent in the acute HF without ACS cohort (94%) (p=0.001). 90% of patients with ACS and 99% of patients without ACS were in hospital admission in NYHA functional class III-IV (p=0.001). Patients with ACS were more susceptible to develop shock during hospitalization (cardiogenic, non-cardiogenic and mixed), with a RR of 1.2 (0.7-2.1) for the overall shock (p=0.4). Complications such as acute pulmonary edema, malignant ventricular arrhythmias, stroke, and cardio-renal syndrome were more frequent in patients with acute HF with ACS. Cardiovascular death, multifactorial death, and sudden death were more frequent in patients in the acute HF with ACS group. Septic deaths were more frequent in HF patients without ACS.

Conclusion: ACS as a precipitating factor of HF is a distinct clinical entity, with a particular Pathophysiology and a different clinical outcome compared with other causes of heart failure. In addition, it is associated with more adverse cardiovascular events in the evolution, therefore, it should be considered for the early initiation of therapeutic strategies in order to improve the prognosis of this group of patients.

Keywords: acute heart failure, acute coronary syndrome, clinical outcome, cardiovascular death

Abbreviations: ACS, acute coronary syndrome; DM, Diabetes mellitus; TIA, transient ischemic attack.

Introduction

Heart failure (HF) is a clinical syndrome characterized by typical symptoms such as dyspnea, edema and fatigue, which may be accompanied by signs such as elevated jugular venous pressure, pulmonary crackles and peripheral edema. HF may be caused by a structural and/or functional cardiac abnormality, resulting in a reduced cardiac output and/or elevated intracardiac pressures at rest or during stress.1-5 Acute heart failure is defined as a rapid worsening of the signs and symptoms of HF and it needs urgent treatment.6-10 It is a clinical syndrome with a complex Pathophysiology which is not completely understood. Given the diversity of clinical presentations, several different physio-pathological mechanisms along with triggering factors of circulatory decomposition are involved. Events that precipitate acute decomposition mainly consist of ischemia, hypertension, arrhythmias, non-cardiac comorbidities, and administered drug agents.4 Up to 15-20% of patients admitted for acute coronary syndrome (ACS) have signs and symptoms of HF at presentation, and another 10% develop HF during the hospital stay. The incidence is even higher in studies that focus on patients diagnosed with acute HF, in these patients the triggering factor is an acute coronary syndrome in up to 40%. It is interesting to note that currently ACS complicated by acute HF is often considered a distinct clinical entity, characterized by structural, hemodynamic and neurohormonal complex interactions, which require urgent coronary intervention. It is usually associated with poor outcome and worse prognosis compared with other patients with acute HF triggered by other causes.4

Heart failure continues to be a significant cause of morbidity and mortality worldwide,11-12 and over a half of patients with acute HF admitted to the hospital have a history of coronary heart disease.16-20 ACS complicated by acute HF leads to a several-fold increase in hospital mortality compared to those without acute HF. In addition, a substantial proportion of hospitalized patients with coronary heart disease develop acute HF during the hospital stay, and have a worse prognosis in those patients who develop acute HF at the initial presentation.21
Acute HF represents a syndrome that has a heterogeneous Pathophysiology with variable results. Therefore, it is interesting to compare a group with unique Pathophysiology and therapeutic objectives as represented by acute HF due to ACS, with those patients who present with acute HF without ACS. Consequently, the prevalence, clinical evolution and the therapies established will be determined. In addition, the incidence of cardiovascular complications and mortality will be determined by comparing both groups of patients with acute HF.

**Objectives**

**Primary**

To determine the frequency of cardiovascular complications in patients with acute HF secondary to ACS in comparison with those patients without ACS.

**Secondary**

i. To describe cardiovascular complications in patients with acute HF.

ii. To determine the frequency of patients admitted to the hospital with the diagnosis of acute HF with and without ACS.

iii. Describe the most frequent triggering causes of acute HF.

iv. Describe the demographic characteristics, clinical profile and cardiovascular risk factors, therapies and outcomes in patients presenting acute HF with and without ACS.

**Material and methods**

**Design:** A retrospective cohort study

**Population:** For this analysis, we stratified the enrolled acute HF patient population into two groups:

- **Cohort 1:** adult patients admitted to the “Hospital de Clínicas” with a diagnosis of acute HF secondary to ACS, from January 2015 to December 2016.

- **Cohort 2:** adult patients admitted to the “Hospital de Clínicas” with a diagnosis of acute HF secondary to other causes not related to ACS, from January 2015 to December 2016.

**Inclusion criteria**

i. Patients diagnosed with acute HF according to the Guidelines of the European Society of Cardiology and the American College of Cardiology for the diagnosis and management of acute HF.

ii. Patients diagnosed with ACS based on the symptoms and clinical context, on electrocardiography and the levels of cardiac biomarkers.

iii. The definitions for STEMI and NSTEMI were based on the European Society of Cardiology and the American College of Cardiology clinical data standards.

**Exclusion criteria:** Patients who do not meet the inclusion criteria.

**Sample:** Non-Probability Sampling. Consecutive sampling.

**Variables**

i. Independent variable: acute coronary syndrome (dichotomous)

ii. Dependent: days of hospitalization, refractory or recurrent heart failure, acute pulmonary edema, cardiogenic or non-cardiogenic or mixed shock, stroke, sepsis, requirement of mechanical ventilation in the ICU, malignant arrhythmias (VT/VF), hospital death.

**Other variables**

a. **Demographic:** age, sex, origin.

b. **Medical history:** heart failure, ischemic heart disease, valvular heart disease, chronic atrial fibrillation, stroke/TIA, COPD, chronic kidney disease, chronic liver disease, thyroid disorders, rheumatologic disease, chronic anemia. Risk factors for atherosclerosis: diabetes mellitus, hypertension, smoking, dyslipidemia.

c. **Triggering factors:** acute coronary syndrome, arrhythmias, pulmonary embolism, anemia, drugs, non-adherence to medication, non-adherence to diet, abandonment of medication, infection, volume overload, valvular, pericardial, cardiomyopathy.

d. **Clinical parameters on presentation:** systolic blood pressure, diastolic blood pressure, heart rate, body mass index.

e. **Acute Heart Failure type in presentation:** Acute de novo and acute on chronic HF.

f. **Functional Classification:** NYHA class. Findings in Electrocardiography: Atrial fibrillation, atrial Flutter, Left Bundle Branch Block, prolonged QRS (>120 ms).

g. **Biochemical parameters:** plasma sodium, plasma potassium, urea, creatinine, estimated glomerular filtration rate, hemoglobin, glycermia, albumin, NT-PRO-BNP, positive troponins, echocardiography findings (Left ventricular ejection fraction), coronary angiography findings.

h. **Treatments received in hospital:** beta-blockers, ACEI, ARB II, aldosterone antagonist, loop diuretics, inotropic and vasopressors.

**Sample size estimation:** the sample size was estimated using EPI Info statistical software version 7.2.0.1 (EPI Info, Center for Disease Control). We estimated the strength of the associations of these groups using 95% confidence intervals, power of 80%, Reason of 1:1. Mortality rate of 5% in cohort 1, and 12% in cohort 2. We calculated a minimum sample size (n) of 200 patients.13,14

**Statistical analysis:** The variables were recorded in the Excel 2007 spreadsheets. The analysis was performed using EPI Info statistical version 7.2.0.1 and Epidat 3.1 software’s. The qualitative variables were expressed in frequencies and percentages, and the quantitative variables in means and standard deviations (SD); or as medians and interquartile ranges. The comparisons between the two cohorts were performed using the chi-square statistical test for the qualitative variables, expressed as Relative Risk. We estimated the strength of the associations of these groups using 95% confidence intervals and a p-value <0.05 was considered statistically significant. Comparisons of the quantitative variables between both cohorts were performed using the Bartlett’s test to verify the homogeneity of variances. In the cases of variance homogeneity, statistical significance was determined with the ANOVA test. In cases without homogeneity of variances, statistical significance was determined using the nonparametric tests of Mann Whitney Wilcoxon and Kruskal Wallis.

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Results

We included 233 patients with acute HF in this study, of which 72 (31%) were secondary to ACS and 161 (69%) were triggered by other causes. The precipitating causes of acute HF are depicted in Table 1.

Of the cases diagnosed at admission with ACS 65% were NSTEMI, 19% STEMI and 15% Unstable Angina. Comparing both groups, cohort 1 (with ACS) had a mean age 2 years older (66±12, p=0.2). The female sex was higher in cohort 1 (57%), while the male sex predominated in cohort 2 (without ACS) (57%) (p=0.04). The baseline characteristics can be seen in Table 2.

Previous history of HF was significantly higher in the acute HF without ACS cohort (p <0.001). Comorbidities such as anemia, rheumatological disease, atrial fibrillation, valvular heart disease, chronic lung disease, chronic kidney disease and liver disease were predominant in these patients without statistically significant difference, except for valvular disease (p=0.004).

We found significant differences in the forms of presentation of acute HF. In the acute HF with ACS cohort the de novo HF was more frequent (19%), while the acute on chronic HF was more frequent in the acute HF without ACS cohort (94%) (p=0.001). 90% of patients with ACS and 99% of patients without ACS were in hospital admission in NYHA functional class III-IV (p=0.001). Patients without ACS tended to be more obese, median BMI of 31 (IQR 29-33) (p=0.6), had a higher heart rate (mean 91±20, p=0.2), and had a higher proportion of atrial fibrillation / flutter (35%) (p=0.2). In contrast, patients with ACS had higher blood pressure (mean SBP 135±22, p=0.3, mean DBP 80±15, p=0.5) and had a higher proportion of complete left bundle branch block (15%) (p=0.09) and a longer QRS interval ≥120 ms (15%) (p=0.007).

Regarding the biochemical parameters, we did not find statistically significant differences between both groups, except in the presence of positive troponin, which predominated in the acute HF with ACS cohort (83%) (p <0.001). Also, hyponatremia (Na <135) that predominated in the AHF without ACS cohort (39%) (p=0.03). Patients with ACS had a tendency to a higher glycemia with a mean of 142±72 (p=0.05) and to have a lower eGFR, mean of 58±34 (p=0.2). The mean of EF was lower in the patients with ACS (mean 48±14, p=0.1). In addition, ventricular dysfunction (FE <50%) was found more frequently in patients in this group (54%, p=0.2). There was no statistically significant difference between both groups regarding the treatment with ACEI, ARB and beta-blockers, as can be seen in Table 3.

Abbreviations: ACS, acute coronary syndrome; ACEI, angiotensin-converting enzyme inhibitor; AHF, acute heart failure; ARB, Angiotensin II receptor blockers; CAD, coronary artery disease; PCI, Percutaneous Coronary Intervention.

In contrast, patients without ACS used loop diuretics (p=0.005) and antialdosterone diuretics more frequently (p <0.001). Patients with ACS required more vasopressors (14% vs. 10%, p=0.3), and similar inotropic requirement with the other group (12%) (p=0.8).

Patients in the acute HF with ACS arm tended to present some type of significant coronary lesion, with the predominant finding being three-vessel CAD (35%). Non-significant coronary stenosis was more frequent in the other arm acute HF without ACS (44%).

No statistically significant differences were found when comparing both groups in terms of the complications presented during hospitalization, as can be seen in Table 4.

Table 1 Precipitating causes of acute heart failure

| Precipitating factors       | % patients (n=233) | % patients with ACS (n=72) | % patients without ACS (n=161) |
|-----------------------------|-------------------|---------------------------|------------------------------|
| Non-adherence to medication | 52                | 35                        | 60                           |
| Infection                   | 48                | 36                        | 53                           |
| ACS                         | 31                | 100                       | 0                            |
| Arrhythmia                  | 16                | 8                         | 19                           |
| Valvular                    | 10                | 4                         | 13                           |
| Non-adherence to diet       | 8                 | 0                         | 11                           |
| Abandonment of treatment    | 6                 | 1                         | 7                            |
| Uncontrolled hypertension   | 5                 | 4                         | 5                            |
| No treatment                | 4                 | 3                         | 4                            |
| Anemia                      | 3                 | 1                         | 4                            |
| Pericarditis/Pericardial effusion | 2               | 0                         | 3                            |
| Stroke/TIA                  | 2                 | 0                         | 2                            |
| Decompensated DM            | 2                 | 1                         | 2                            |
| Pulmonary hypertension      | 2                 | 0                         | 2                            |
Table 2 Demographic and clinical characteristics of patients with acute heart failure with and without acute coronary syndrome

|                | Overall (n: 233) | AHF with ACS n:72 (31%) | AHF without ACS n:161 (69%) | P value |
|----------------|------------------|-------------------------|-----------------------------|---------|
| **Demographics** |                  |                         |                             |         |
| Age, years; mean ± SD | 65 ± 14          | 66 ± 12                  | 64 ± 14                     | 0.2     |
| Male, n (%)        | 123 (53)         | 31 (43)                  | 92 (57)                     | 0.04    |
| Female, n (%)      | 110 (47)         | 41 (57)                  | 69 (43)                     |         |
| **Medical history** |                  |                         |                             |         |
| Heart Failure, n (%) | 212 (91)        | 58 (81)                  | 154 (96)                   | <0.001  |
| Ischemic heart disease, n (%) | 111 (48) | 40 (56)                  | 71 (44)                    | 0.1     |
| Valvular disease, n (%) | 113 (49)       | 25 (35)                  | 88 (55)                    | 0.004   |
| Rheumatologic disease, n (%) | 7 (3)         | 1 (1)                    | 6 (4)                      | 0.5     |
| Chronic AF, n (%)  | 73 (31)          | 19 (26)                  | 54 (34)                    | 0.2     |
| Stroke/TIA, n (%)  | 10 (4)           | 4 (6)                    | 6 (4)                      | 0.7     |
| PAD, n (%)         | 19 (8)           | 7 (10)                   | 12 (7)                     | 0.5     |
| Anemia, n (%)      | 68 (29)          | 16 (22)                  | 52 (32)                    | 0.1     |
| COPD, n (%)        | 42 (18)          | 8 (11)                   | 34 (21)                    | 0.06    |
| CKD, n (%)         | 58 (25)          | 18 (25)                  | 40 (25)                    | 0.9     |
| Liver disease, n (%) | 4 (2)            | 0 (0)                    | 4 (2)                      | -       |
| Thyroid disorders, n (%) | 7 (3)           | 4 (6)                    | 8 (5)                      | 0.8     |
| **Risk factors for atherosclerosis** |                  |                         |                             |         |
| Smoking, n (%)     | 75 (32)          | 21 (29)                  | 54 (34)                    | 0.5     |
| Hypertension, n (%) | 220 (94)         | 70 (97)                  | 150 (93)                   | 0.3     |
| Dyslipidemia, n(%); n=228 | 152 (67) | 54 (79)                  | 98 (61)                    | 0.007   |
| Diabetes mellitus, n (%) | 86 (37)        | 29 (40)                  | 57 (35)                    | 0.4     |
| **Clinical parameters on presentation** |                  |                         |                             |         |
| Acute de novo HF, n (%) | 23 (10)         | 14 (19)                  | 9 (6)                      | 0.001   |
| Acute on chronic HF, n (%) | 210 (90)       | 58 (81)                  | 152 (94)                   |         |

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|                           | Overall (n: 233) | AHF with ACS n=72 (31%) | AHF without ACS n=161 (69%) | P value |
|---------------------------|------------------|-------------------------|-----------------------------|---------|
| NYHA class I-II, n (%)    | 8 (3)            | 7 (10)                  | 1 (1)                       | 0.001   |
| NYHA class III-IV, n (%)  | 225 (97)         | 65 (90)                 | 160 (99)                    | 0.001   |
| BMI, kg/m²; median (IQR)  | 31 (28-33)       | 30.5 (28-33)            | 31 (29-33)                  | 0.6**   |
| SBP; mean ± SD            | 132 ± 24         | 135 ± 22                | 131 ± 24                    | 0.3*    |
| DBP; mean ± SD            | 79 ± 14          | 80 ± 15                 | 79 ± 13                     | 0.5*    |
| HR; mean ± SD             | 90 ± 20          | 88 ± 20                 | 91 ± 20                     | 0.2*    |

Electrocardiography

|                           |                  |                       |                             |         |
|---------------------------|------------------|-----------------------|------------------------------|---------|
| AF/Flutter, n (%)         | 77 (33)          | 20 (28)               | 57 (35)                     | 0.2     |
| QRS ≥120 msec, n (%)      | 24 (10)          | 11 (15)               | 13 (8)                      | 0.09    |
| LBBB, n (%)               | 19 (8)           | 11 (15)               | 8 (5)                       | 0.007   |

Biochemical parameters

|                           |                  |                       |                             |         |
|---------------------------|------------------|-----------------------|------------------------------|---------|
| Sodium, mEq/L; media ±SD  | 136 ± 4.6        | 136.9 ± 4.8           | 135.6 ± 4.5                  | 0.05*   |
| Sodium <135 mEq/L, n (%)  | 81 (35)          | 18 (25)               | 63 (39)                     | 0.03    |
| Potassium; mEq/L ±SD      | 4.2 ± 0.7        | 4.2 ± 0.7             | 4.3 ± 0.6                    | 0.3*    |
| Urea, mg/dl               | 81.1 ± 59.9      | 81.1 ± 67.1           | 81.2 ± 56.7                  | 0.9*    |
| Creatinine, g/dl          | 1.8 ± 1.8        | 2 ± 2                 | 1.7 ± 1.7                    | 0.3*    |
| eGFR, mg/ml/1.73 m2; mean ± SD | 62.2 ± 36.9   | 58.3 ± 34.3           | 63.9 ± 38                    | 0.2*    |
| eGFR <60 mg/ml/1.73 m2, n (%) | 113 (49)      | 32 (44)               | 81 (50)                      | 0.4     |
| Hemoglobin, g/dl; mean ± SD | 12.1 ± 2.8      | 12.6 ± 2.7            | 12 ± 2.9                     | 0.1*    |
| Glycemia, mg/dl; mean ± SD | 130.2 ± 64.9    | 142.4 ± 71.8          | 124.8 ± 61                   | 0.05*   |
| Albumin, mg/dl; mean ± SD | 3 ± 0.5          | 3.1 ± 0.6             | 3 ± 0.5                      | 0.8*    |
| NT-PRO-BNP elevated, n (%), n=4 | 4 (2)            | 1 (1)                 | 3 (2)                        | 0.7     |
| Positive troponin, n (%)  | 59 (49)          | 58 (83)               | 7 (14)                       | <0.001  |

Echocardiography n=228

|                           |                  |                       |                             |         |
|---------------------------|------------------|-----------------------|------------------------------|---------|
| EF%, mean ± SD            | 49 ± 14          | 48 ± 14                | 50 ± 15                      | 0.1*    |
| EF ≥50%, n (%)            | 116 (51)         | 32 (46)                | 84 (53)                      | 0.2     |
| EF <50%, n (%)            | 112 (49)         | 38 (54)                | 74 (47)                      |         |

* T-test
** Kruskal Wallis

**Abbreviations:** ACS, acute coronary syndrome; AF, atrial fibrillation; AHF, acute heart failure; BMI, body mass index; COPD, chronic obstructive pulmonary disease; CKD, chronic kidney disease; DBP, diastolic blood pressure; EF, ejection fraction; eGFR, estimated glomerular filtration rate; HR, heart rate; LBBB, left bundle branch block; NYHA, New York Heart Association functional class; PAD, peripheral artery disease; SBP, systolic blood pressure; TIA, transient ischemic attack. Ischemic heart disease was more common in the acute HF with ACS cohort (p=0.1), as well as risk factors for atherosclerotic disease such as Hypertension (p=0.3), dyslipidemia (p=0.007) and diabetes mellitus (p=0.4). Smoking or smoking history (p=0.5) was more frequent in the acute HF without ACS cohort, as well as the history of cerebrovascular disease, peripheral arterial disease and thyroid disease were more frequent in this cohort.

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Table 3 Treatments received and procedures performed during hospitalization in patients with AHF with and without ACS

| Treatments                      | Overall n=233 | AHF with ACS n=72 (31%) | AHF without ACS n=161 (69%) | P value |
|---------------------------------|--------------|-------------------------|----------------------------|---------|
| ACEI, n (%)                     | 100 (43)     | 30 (42)                 | 70 (43)                    | 0.7     |
| ARB, n (%)                      | 40 (17)      | 10 (14)                 | 30 (19)                    | 0.3     |
| Aldosterone antagonist, n (%)   | 88 (38)      | 12 (17)                 | 76 (47)                    | <0.001  |
| Beta-blockers, n (%)            | 154 (66)     | 53 (74)                 | 101 (63)                   | 0.1     |
| Loop diuretics, n (%)           | 226 (97)     | 66 (92)                 | 160 (99)                   | 0.005   |
| Inotropic, n (%)                | 28 (12)      | 9 (12)                  | 19 (12)                    | 0.8     |
| Vasopressors, n (%)             | 26 (11)      | 10 (14)                 | 16 (10)                    | 0.3     |

Procedures

| Coronary angiography, n (%)     | 35 (15)      | 26 (36)                 | 9 (6)                      | <0.001  |
| Non-significant CAD, n (%)     | 12 (34)      | 8 (31)                  | 4 (44)                     | 0.7     |
| Single-vessel CAD, n (%)       | 4 (11)       | 4 (15)                  | 0                          | 0.5     |
| Double-vessel CAD, n (%)       | 6 (17)       | 5 (19)                  | 1 (11)                     | 0.9     |
| Three-vessel CAD, n (%)        | 13 (37)      | 9 (35)                  | 4 (44)                     | 0.8     |
| PCI, n (%)                     | 2 (1)        | 2 (11)                  | 0                          | 0.8     |

Table 4 Clinical outcomes during hospitalization in acute HF

| Overall n=233 | AHF with ACS n=72 (31%) | AHF without ACS n=161 (69%) | RR (CI 95%) | P value |
|---------------|-------------------------|----------------------------|--------------|---------|
| Overall shock, n (%)      | 42 (18)     | 15 (21)                 | 27 (17)      | 1.2 (0.7-2.1) | 0.4     |
| Cardiogenic shock, n (%)  | 21 (9)      | 7 (10)                  | 14 (9)       | 1.1 (0.4-2.6) | 0.8     |
| Non-cardiogenic shock, n (%) | 5 (2)     | 2 (3)                   | 3 (2)        | 1.4 (0.2-8.7) | 0.9     |
| Mixed shock, n (%)        | 17 (7)      | 7 (10)                  | 10 (6)       | 1.5 (0.6-3.94) | 0.3     |
| VT/VF, n (%)              | 6 (3)       | 2 (3)                   | 4 (2)        | 1.1 (0.2-5.9) | 0.7     |
| Stroke, n (%)             | 3 (1)       | 2 (3)                   | 1 (1)        | 4.4 (0.4-48.5) | 0.4     |
| Major bleeding, n (%)     | 2 (1)       | 1 (1)                   | 1 (1)        | 2.2 (0.1-35.2) | 0.8     |
| Cardiorenal syndrome, n (%) | 15 (6)     | 5 (7)                   | 10 (6)       | 1.1 (0.3-3.1) | 0.8     |
| Hemodialysis requirement, n (%) | 14 (6)   | 4 (6)                   | 10 (6)       | 0.8 (0.2-2.7) | 0.9     |
| Acute Pulmonary Edema, n (%) | 24 (10)   | 8 (11)                  | 16 (10)      | 1.1 (0.5-2.4) | 0.7     |
| Sepsis, n (%)             | 16 (7)      | 2 (3)                   | 14 (9)       | 0.3 (0.07-1.3) | 0.1     |
| ICU requirement, n (%)     | 7 (3)       | 2 (3)                   | 5 (3)        | 0.8 (0.1-4.5) | 0.7     |
| Days of hospitalization, mean ± SD | 12 ± 8   | 12 ± 9                  | 12 ± 8       | 0.9*    |
| MI, n (%)                 | 3 (1)       | 0                       | 3 (2)        | NC      |
| Emergency cardiac surgery, n (%) | 3 (1)     | 0                       | 3 (2)        | NC      |
| Complete heart block, n (%) | 2 (1)       | 1 (1)                   | 1 (1)        | 2.2 (0.1-35.2) | 0.8     |
| Global death, n (%)        | 30 (13)     | 11 (15)                 | 19 (12)      | 1.2 (0.6-2.5) | 0.4     |
| Cardiovascular death, n (%) | 15 (50)    | 6 (55)                  | 9 (47)       | 1.1 (0.5-2.3) | 0.7     |
| Sudden death, n (%)        | 5 (17)      | 3 (27)                  | 2 (11)       | 2.5 (0.5-13.1) | 0.4     |
| Death of multifactorial cause, n (%) | 13 (43) | 5 (45)                  | 8 (42)       | 1 (0.4-2.4) | 0.8     |
| Death by sepsis, n (%)     | 2 (7)       | 0                       | 2 (11)       | NC      |

* T-test

Abbreviations: ACS, acute coronary syndrome; AHF, acute heart failure; ICU, intensive care unit; MI, myocardial infarction; NC, not calculable; VF, ventricular fibrillation; TV, ventricular tachycardia

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However, patients with ACS were more susceptible to develop shock during hospitalization (cardiogenic, non-cardiogenic and mixed), with a RR of 1.2 (0.7-2.1) for the overall shock (p=0.4). Complications such as acute pulmonary edema, malignant ventricular arrhythmias, stroke, and cardiorenal syndrome were more frequent in patients with acute HF with ACS. Other complications such as: major bleeding, hemodialysis requirement, complete heart block, mechanical ventilation requirement and admission to intensive care unit were similar in both groups. Hospital-acquired infection, myocardial infarction, emergency cardiac surgery were more frequent in acute HF without ACS patients. Days of hospitalization were similar in both groups, mean 12±8 (p=0.9).

We did not find significant differences in terms of overall mortality, RR of 1.2 (0.6-2.5) (p=0.4). However, cardiovascular death, multifactorial death, and sudden death were more frequent in patients in the acute HF with ACS group. Septic deaths were more frequent in patients without ACS.

Discussion

In this study we compared the results of patients admitted to the hospital with acute heart failure triggered by acute coronary syndrome opposed to cases of acute heart failure due to a different trigger. Approximately 30% of the total enrolled patients belonged to the acute HF with ACS cohort, which agrees with the findings of previous studies.21-23 The patients belonging to this cohort were on average older, had more associated cardiovascular risk factors, which was consistent with the greater history of ischemic heart disease.20-33 In addition, they had fewer associated comorbidities, and at the time of hospital admission they were more likely to have de novo acute HF compared with patients without ACS.

In contrast, patients without ACS had a history of chronic heart failure, and therefore presented in almost all cases as an exacerbation of heart failure. Among the biochemical parameters of both groups, significant differences were not found, therefore, comparable for the purpose of the study. Regarding the treatments received, no significant differences were found for most of them, which would be explained by the similar management of coronary syndrome and heart failure.1,2,20-21 On the other hand, there were differences regarding the use of loop diuretics and antialdosterone diuretics, which predominated in patients without ACS, this could be explained by the higher proportion of patients with chronic heart failure in this group, because the congestive symptoms are more frequent in this entity.

There were no differences regarding the days of hospitalization. It is expected that the group with ACS have a longer hospital stay according to previous studies.14 The reason why patients without ACS presented similar days of hospitalization could be explained by the high proportion of the infectious cause as precipitants of acute HF in this group, in addition they presented more frequently hospital-acquired infection, which supposes the requirement of several days of hospitalized antibiotic treatment. We did not find significant differences in hospital adverse cardiovascular outcomes comparing both groups, however, patients with ACS required more vasopressors, so we can assume a more aggressive course of this entity15 which relates with an increased frequency of overall shock, excepting septic shock, for reasons previously discussed. Other complications were also recorded more frequently in these patients, such as the appearance of malignant arrhythmias, stroke, renal failure, although the requirement for rescue hemodialysis was similar in both groups, possibly due to the greater history of chronic renal failure in the group without ACS.

The requirement for mechanical ventilation and admission to the ICU was similar in both cohorts. Three events of myocardial infarction were recorded in the in-hospital setting in 3 patients belonging to the group without ACS. This was associated with worse evolution during hospitalization, all of them were complicated by cardiogenic shock, 2 died during hospitalization, and the other patient required mechanical ventilation and admission to the ICU, leading us to suppose that infarction represented a triggering event for the appearance of complications in this group of patients.

Emergency cardiac surgery was only performed in 3 patients without ACS, due to the etiology of heart failure, 2 of them of valvular cause secondary to endocarditis requiring valve replacement intervention, and another patient due to cardiac tamponade requiring surgical pericardiectomy. There were no significant differences when comparing the mortality between both groups; however, greater tendency was found in patients with ACS. Overall in-hospital mortality was 13%, comparable with other reports.31,32 The acute HF cohort with ACS presented a greater proportion of cardiovascular death, death of multifactorial cause and sudden death, which is related to the worse prognosis in this entity.

With these findings we could assume that an acute coronary event represents a strong risk factor for the appearance of major adverse cardiovascular events in patients with acute HF during the course of the disease.30 Therefore, a strategy for early diagnosis, risk stratification, and early initiation of aggressive therapy could improve the prognosis of this group of patients.

The limitations of this study are that they represent only the tendency of a single hospital center, while the reports referring to this topic were carried out in a multicentric and multinational manner with the recruitment of a much higher sample. However, the proportion of patients with acute HF with and without ACS in this study, as well as the results found are to a large extent comparable with previous studies.13,14,30-32 We did not find a national or a regional record related to the subject of this study, so this report would be useful as an initiative for future research, in order to have a registration system on a very prevalent disease with high morbidity and mortality.

Conclusion

Adverse cardiovascular outcomes were more frequent in patients with ACS. These patients were more likely to suffer shock, especially cardiogenic and mixed shock type. In addition, these patients developed more frequently acute pulmonary edema, cardiorenal syndrome, stroke and malignant arrhythmias. All causes of death were more prevalent in patients with ACS, especially cardiovascular death. The most frequent precipitating causes of acute HF were non-adherence to medication, infection, ACS, arrhythmias, especially tachyarrhythmias, and valvular etiology, among others.

We can conclude that ACS as a precipitating factor of HF is a distinct clinical entity, with a particular Pathophysiology and a different clinical outcome compared with other causes of heart failure. In addition, it is associated with more adverse cardiovascular events in the evolution, therefore, it should be considered for the early initiation of therapeutic strategies in order to improve the prognosis of this group of patients.
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Conflict of interest

The author declares that there is no conflict of interest.

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