



ORIGINAL ARTICLE

Pharmacological treatment after acute coronary syndrome: Baseline clinical characteristics and gender differences in a population-based cohort study



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KEYWORDS

Electronic health records;
Acute coronary syndrome;
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Abstract

Objective: To describe baseline socio-demographic and clinical characteristics and drugs prescribed for secondary prevention after a first episode of ACS and to assess differences between men and women.

Setting: PHC in Catalonia. Data source: SIDIAP (Information System for Research in Primary Care).

Participants: Patients who suffered an ACS during 2009–2016 and followed-up in PHC centres of the Catalan Health Institute in Catalonia.

Interventions: Not applicable.

Main measures: Socio-demographic and clinical characteristics at baseline: sex, age, socioeconomic index, toxic habits, comorbidities, study drugs (prescribed for cardiovascular secondary prevention: antiplatelets, betablockers, statins, drugs acting on the renin–angiotensin system) and comedications.

Abbreviations: ACEI, angiotensin converting enzyme inhibitors; ACS, acute coronary syndrome; AMI, acute myocardial infarction; ARB, angiotensin receptor blockers; ATC, chemical classification system; BMI, body mass index; CMBD-HA, minimum basic dataset at hospital discharge; ECAP, electronic health records in Primary Health Care of the Catalan Health Institute; ICD, international classification of disease; MEDEA, socioeconomic index; R, rural; SIDIAP, information system for research in primary care; U, urban.

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Results: 8071 patients included, 71.3% of them were men and 80.2% had an acute myocardial infarction. Their mean age was 65.3 and women were older than men. The most frequent comorbidities were hypertension, dyslipidaemia and diabetes and they were more common in women. Antiplatelets (91.3%) and statins (85.7%) were the study drugs most prescribed. The uses of all comedications were significantly higher in women, except for nitrates. The combination of four study groups was initially prescribed in 47.7% of patients and combination of betablockers, statins and antiplatelets was prescribed in 18.4%. More men than women received all recommended pharmacological groups.

Conclusion: Women were older, had more comorbidities and received more comedications. Most patients were treated with a combination of four or three study drugs for secondary prevention. Men initiated more drug treatments for secondary prevention and dual antiplatelet therapy than women.

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PALABRAS CLAVE

Registros electrónicos de salud;
Síndrome coronario agudo;
Adherencia a la medicación;
Prevención secundaria

Tratamiento farmacológico después de un síndrome coronario agudo: características clínicas y diferencias de género en un estudio poblacional de cohortes

Resumen

Objetivos: Describir las características sociodemográficas y clínicas basales, y los fármacos prescritos para la prevención cardiovascular secundaria tras un síndrome coronario agudo (SCA). Analizar si existen diferencias entre varones y mujeres.

Emplazamiento: Atención primaria (AP) en Cataluña. Fuente de datos: Sistema de Información para el Desarrollo de la Investigación en AP (SIDIAP).

Participantes: Pacientes que hayan sufrido un primer SCA durante 2009-2016, seguidos en AP del Instituto Catalán de la Salud en Cataluña.

Intervenciones: No aplica.

Mediciones principales: Características sociodemográficas y clínicas al inicio: sexo, edad, índice socioeconómico, hábitos tóxicos, comorbilidades, fármacos de estudio (prescritos para prevención secundaria: antiagregantes, betabloqueantes, estatinas, fármacos del sistema renina-angiotensina) y fármacos concomitantes.

Resultados: Se incluyeron 8.071 pacientes; 71,3% varones y 80,2% habían sufrido infarto. La edad media era de 65,3 años y las mujeres eran mayores que los varones. Las comorbilidades más frecuentes fueron hipertensión, dislipemia y diabetes; más comunes en mujeres. Antiagregantes (91,3%) y estatinas (85,7%) fueron los fármacos más prescritos. El uso de todas las comedicaciones era más frecuente en mujeres, excepto nitratos. La combinación de los 4 grupos farmacológicos de estudio se prescribió al 47,7% de los pacientes incluidos y la combinación de antiagregante, betabloqueante y estatina al 18,4%. Más varones que mujeres recibieron los fármacos recomendados.

Conclusiones: Las mujeres incluidas eran mayores, con más comorbilidad y mayor uso de comedicaciones. La mayoría de pacientes eran tratados con la combinación de 3 o 4 fármacos para prevención secundaria. Los varones iniciaban más fármacos para prevención secundaria y más terapia antiagregante doble que las mujeres.

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Introduction

Cardiovascular disease remains the most common cause of death worldwide, 31.5% of all deaths and 45% for non-communicable disease deaths in Europe.^{1,2} Despite these numbers, the incidence of cardiovascular disease has decreased over the last four decades, due to population-level lifestyle changes and the development of effective

interventions to treat individuals and invasive procedures and effective drugs to tackle modifiable risk factors.³

Several randomised clinical trials, meta-analyses and cohort studies have shown that long-term administration of aspirin, statins, beta-blockers, and angiotensin-converting enzyme inhibitors (ACEI) or angiotensin-receptor blockers (ARB) improve survival in high risk patients, particularly those with established cardiovascular disease.⁴⁻⁷

Therefore, the European and American Cardiology guidelines recommend in both genders this long-term pharmacological therapy for an acute coronary syndrome (ACS) secondary prevention.^{8–12}

Several population-based studies have analysed the pharmacological secondary prevention in the real-world practice. In Lafeber et al. study, 67% of patients with cardiovascular disease were treated with a combination of aspirin, statin and at least one blood pressure-lowering agent for secondary prevention.¹³ Sanfélix-Gimeno et al. showed that after an ACS 92.8% of patients were treated with an antiplatelet, 74.7% with beta-blocker, 87.1% with statins and 77.2% with an ACEI or ARB.⁵

Some population-based studies have described differences between men and women in clinical characteristics and pharmacological treatment received after ACS. Women have been reported to be older than men and have greater comorbidities, such as hypertension, diabetes and dyslipidaemia.^{14–17} Some differences between genders in secondary prevention have also been described and found that women were less likely to be treated.^{14–16}

This work is part of IMPACT study and the protocol has been previously published.¹⁸ The objective of IMPACT study is to assess the impact of the four recommended drugs adherence on mortality and cardiovascular morbidity. This study aims to describe the baseline socio-demographic and clinical characteristics and the medication prescribed for secondary prevention after a first episode of ACS in a Primary Health Care (PHC) cohort in Catalonia (Spain) and to assess differences in these characteristics between women and men.

Methods

Study design

Population-based observational cohort study of patients with a first episode of ACS admitted in hospitals of the Catalan Health Institute during 2009–2016, followed-up in PHC. The data source is Information System for Research in Primary Care (SIDIAP) database, which includes PHC data of more than 5.8 million people from Catalonia (approximately 80% of the Catalan population).¹⁸

Data source

SIDIAP database,¹⁹ which contains pseudonymized information coming from different data sources: ECAP (electronic health records in PHC of the Catalan Health Institute, including) socio-demographic characteristics, comorbidities registered as International Classification of Disease (ICD) 10 codes (Table S1, *Appendix*),²⁰ specialist referrals, clinical parameters, toxic habits (smoking and alcohol intake), sickness leave, date of death, laboratory test data; general practitioners' prescriptions and their corresponding pharmacy invoice data registered as chemical classification system (ATC) codes²¹; and the CMBD-HA (minimum basic dataset at hospital discharge),²² which includes diagnoses at hospital discharge registered as ICD9 codes (Table S1, *Appendix*).²³

Study population

All adults with a first episode of ACS (acute myocardial infarction (AMI) or unstable angina) registered in CMBD-HA from 2009 to 2016 with at least two months of follow-up in SIDIAP after the index date were included. The individuals lost in follow-up during the first two months have no information available in the database to be captured. *Exclusion criteria:* patients with a recorded diagnosis of a previous ischaemic stroke.

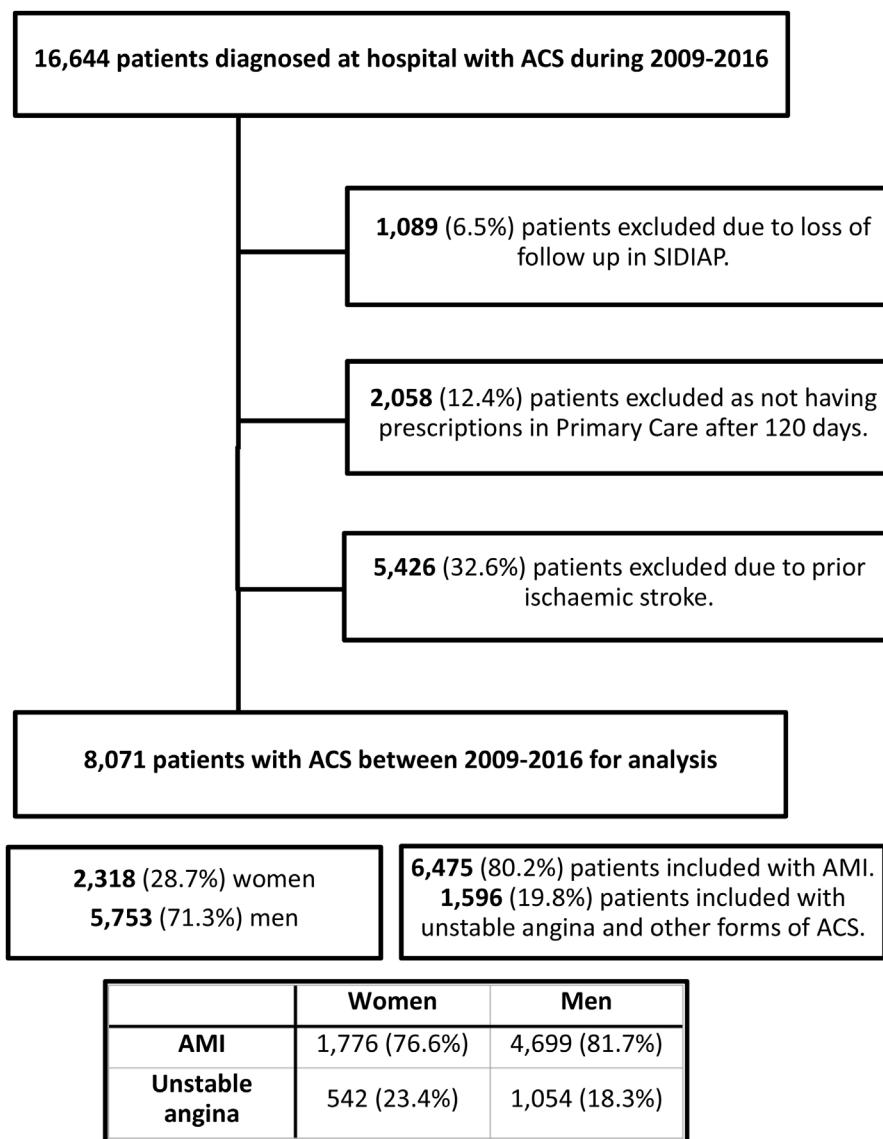
Study variables

At index date: age, gender, socioeconomic MEDEA Index,^{24,25} toxic habits (smoking and alcohol), body mass index (BMI), type of ACS event (AMI, unstable angina or other forms of ACS), laboratory data (cholesterol, other lipid parameters and glomerular filtration rate), and comorbidities of interest. MEDEA socioeconomic index is a deprivation index built with the information of five cities in Spain (Barcelona, Bilbao, Madrid, Sevilla, Valencia), using the census section as the unit of analysis and 2001 census data, based on five indicators of socioeconomic position: manual workers, unemployment, temporary workers, overall insufficient education and insufficient education in young people. MEDEA is able to detect small areas with socioeconomic inequalities in large cities, allowing the study of associations between socioeconomic indicators and mortality. MEDEA is categorised in five urban quintiles, with quintile 1 (U1) corresponding to the least deprived population and quintile 5 (U5), the most deprived.²⁴ In order to facilitate the presentation of our results, we grouped categories U1 to U3, and U4 to U5. The rural category (R) includes municipalities with less than 10,000 inhabitants and a population density lower than 150/km². The use of MEDEA index has not been analysed for rural areas. Socioeconomic deprivation measured with MEDEA was associated with an increase in total mortality in urban areas of Catalonia.²⁵

The study drugs were those recommended for secondary prevention: antiplatelets, beta-blockers, statins and ACEI/ARB. Study drugs prescribed after the ACS event and other concomitant drugs were collected after the index date. The initiation of exposure to the study drugs was defined according to the drugs firstly prescribed during the period spanning from index day to 120 days after the event in order to capture all prescriptions in PHC, due to the length of hospital's prescriptions and the delay in the register of the dispensing in our records.

Statistical analysis

Demographic and baseline characteristics of the participants were described using counts and proportions for categorical variables and for continuous variables mean with standard deviation (SD) for normally distributed variables and median and interquartile range (IQR) for skewed distributions. Univariate analysis between genders was performed by means of Pearson's Chi-square test and we compared mean or median between groups using Student's *T* test and Mann-Whitney *U* test, respectively. The analysis between groups according to the number of study drugs



Scheme 1. General outline of the study: Study flowchart. Includes the flowchart of patient's inclusion and exclusion for the study. ACS, acute coronary syndrome; AMI, acute myocardial infarction.

was performed using the ANOVA test (under equal variance assumption) for continuous variables and Pearson's Chi-square test (with continuity correction) for categorical variables.

Regarding to the missing data, we assumed that if data was missing, it meant that the patient did not had that condition.

All analyses were performed using R 3.5.1 (R Core Team, 2020. A language and environment for statistical computing. R Foundation for Statistical Computing, Vienna, Austria. URL <https://www.R-project.org/>), under a significance level of 0.05.

Results

There were 16,644 patients admitted to hospital with a first episode of ACS from 2009 to 2016 and 8573 of them were excluded (Scheme 1). 8071 patients were included,

71.3% of them were men and 80.2% had an AMI (men: 81.7%; women: 76.6%). Their mean age was 65.3, women were older than men (71.1 vs 63.0, $p < 0.001$) and 45.1% older than 75. The most frequent comorbidities were hypertension, dyslipidaemia and diabetes and they were all significantly more common in women. Heart failure and renal impairment were also common in women (**Table 1**).

Antiplatelet agents (91.3%) were the most prescribed drugs, followed by statins (85.7%), beta-blockers (76.7%), and lastly, ACEI/ARBs (66.3%). More men than women received all study drugs. Nitrates were the comedication most prescribed overall after the event. The use of all comediations was significantly higher in women, except for nitrates (**Table 2**). The combination of four study drugs was initially prescribed in 47.7% of patients and 31.8% of total prescriptions were with three study drugs. Beta-blockers, statins and antiplatelets was the more frequent combination of three components (18.4%) (**Fig. 1**). More men were

Table 1 Gender differences in socio-demographic characteristics, laboratory data and comorbidities.

N (%)	Overall	Women	Men	p-Value
	8071	2318 (28.7)	5753 (71.3)	
<i>Acute myocardial infarction</i>	6475 (80.2)	1776 (76.6)	4699 (81.7)	<0.001
<i>Unstable angina</i>	1596 (19.8)	542 (23.4)	1054 (18.3)	<0.001
<i>Age in years, mean (SD)</i>	65.3 (13.6)	71.1 (13.1)	63.0 (13.0)	<0.001
Median (IQR, Range)	71.0 (22, 82)	80.0 (21, 82)	68.0 (19, 73)	<0.001
>75 years	2198 (27.2)	1046 (45.1)	1152 (20.0)	<0.001
<i>MEDEA</i> ^{24,25}				0.009
R	1427 (17.7)	386 (16.7)	1041 (18.1)	
U1-3	3366 (41.7)	924 (39.9)	2442 (42.5)	
U4-5	2785 (34.5)	851 (36.7)	1934 (33.6)	
<i>Smokers*</i>	2320 (32.1)	335 (15.5)	1985 (39.1)	<0.001
Missing (10.3%)				
<i>High alcohol intake**</i>	5 (0.1)	0 (0.0)	5 (0.1)	<0.001
Missing (21.8%)				
<i>BMI (kg/m²; mean, SD)</i>	29.0 (4.7)	29.9 (5.5)	28.7 (4.3)	<0.001
Missing (20.8%)				
<i>BMI ≥ 30: obesity</i>	2387 (37.4)	903 (45.1)	1484 (33.8)	<0.001
<i>Cholesterol Total mg/dL, mean (SD)</i>	208.00 [180.00, 235.00]	211.00 [183.00, 238.00]	206.00 [179.00, 235.00]	<0.001
Missing (14.8%)				
<i>Cholesterol LDL mg/dL, median (IQR, Range)</i>	128.00 [104.00, 153.00]	128.00 [103.00, 152.00]	129.00 [104.00, 153.00]	0.510
Missing (21.5%)				
<i>Cholesterol HDL mg/dL, median (IQR, Range)</i>	47.00 [40.00, 56.00]	53.00 [44.00, 62.00]	45.00 [38.00, 53.00]	<0.001
Missing (19.0%)				
<i>Triglycerides mg/dL, median (IQR, Range)</i>	127.00 [94.00, 183.00]	124.00 [93.00, 178.00]	128.00 [95.00, 185.00]	<0.001
Missing (17.7%)				
<i>Diabetes mellitus</i>	2169 (26.9)	743 (32.1)	1426 (24.8)	<0.001
<i>Dyslipidaemia</i>	3450 (42.7)	1134 (48.9)	2316 (40.3)	<0.001
<i>Heart failure</i>	296 (3.7)	159 (6.9)	137 (2.4)	<0.001
<i>Hypertension</i>	4294 (53.2)	1540 (66.4)	2754 (47.9)	<0.001
<i>Peripheral artery disease</i>	385 (4.8)	90 (3.9)	295 (5.1)	0.021
<i>Renal impairment; eGFR <45 ml/min/1.73 m²</i>	528 (7.6)	274 (12.8)	254 (5.4)	<0.001
Missing (14.9%)				

p-Value from Pearson's Chi-square test (categoric variables) and t-test or Mann-Whitney U test (numeric variables) comparing women versus men. BMI, body mass index; LDL-C, low density lipoprotein-cholesterol; HLD-C, high density lipoprotein- cholesterol; eGFR, estimated glomerular filtration rate; R (Rural); U (Urban).

treated with the combination of four (2879 [50.0%] vs 968 [41.8%, $p < 0.001$] and with the most frequent combination of three drugs: antiplatelets, statins and beta-blockers (1115 [19.4%] vs 368 [15.9%]; $p < 0.001$); and antiplatelets, statins and ACEI/ARB (492 [8.6%] vs 210 [9.1], $p = 0.491$).

Table 3 compares the baseline characteristics difference of patients by study drug number prescribed. Patients with AMI significantly received four study drugs more frequently (86%) than other combination of three (79.2%) or \leq two study drugs (68.3%, $p < 0.001$). More women initiated \leq two study drugs (38.9%) than three (27.5%) or four (25.2%). Patients receiving \leq two study drugs were older (68.9 years). There were more patients treated with other comediations after the event in the group of \leq two study drugs than the other combinations (**Table 3**).

Fig. 2 represents the different drugs prescribed overall, in men and women. Men received dual antiplatelet therapy more frequently than women; the most used antiplatelets were aspirin and clopidogrel. The most prescribed beta-blocker was bisoprolol both in men and women. Atorvastatin was the most prescribed statin for all patients. Enalapril and ramipril were the most used ACEI, being ramipril more frequent in men. Losartan is the most prescribed ARB, followed by valsartan and olmesartan (**Fig. 2**).

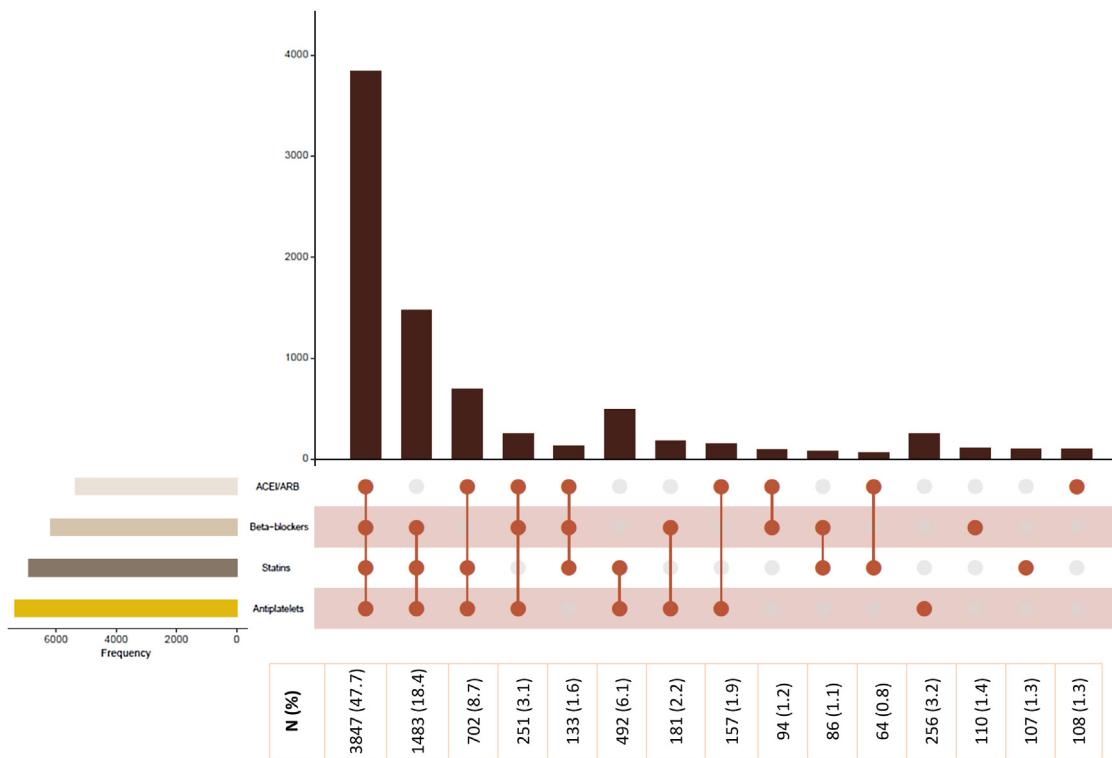
Discussion

We report baseline socio-demographic and clinical characteristics of 8071 patients from a Primary Health Care cohort who had a first ACS. Patients' characteristics have been

Table 2 Gender differences in population that initiate treatment for secondary prevention: study drugs and comediations after the event.

N (%)	Overall	Women	Men	p-Value
Study drugs				
Antiplatelets	7369 (91.3)	1998 (86.4)	5371 (93.3)	<0.001
Statins	6914 (85.7)	1864 (80.5)	5050 (87.8)	<0.001
Beta-blockers	6185 (76.7)	1675 (72.4)	4510 (78.4)	<0.001
ACEI/ARB	5356 (66.3)	1505 (65.1)	3851 (66.9)	0.2223
Comediations				
Anticoagulants	602 (7.5)	260 (11.2)	342 (5.9)	<0.001
Calcium channel-blockers	1309 (16.2)	471 (20.3)	838 (14.6)	<0.001
Diuretics	1754 (21.7)	792 (34.2)	962 (16.7)	<0.001
Drug used in diabetes mellitus	1997 (24.7)	679 (29.3)	1318 (22.9)	<0.001
NSAID	1627 (20.2)	655 (28.3)	972 (16.9)	<0.001
Nitrates	3005 (37.2)	811 (35.0)	2194 (38.1)	0.009

p-Value from Pearson's Chi-square test comparing women versus men. ACEI, angiotensin-converting enzyme inhibitors; ARB, angiotensin-receptor blockers; NSAID, non-steroidal anti-inflammatory drugs.

**Figure 1** Study drugs combinations. This figure depicts N and % of patients initiating any possible combination of the drugs used for secondary prevention. ACEI, angiotensin-converting enzyme inhibitors; ARB, angiotensin-receptor blockers; AntiPL, antiplatelets.

analysed overall, divided into genders and number of study drugs prescribed. We found that women were older, had greater comorbidity at baseline and received more comediations after the study event than men, probably because they were older when had the first ACS, as described in a similar cohort by Ribas et al.²⁶ In agreement with similar studies, we found a higher prevalence of comorbidities in women,^{15,27,28} while men had a higher prevalence of peripheral artery disease,²⁹ possibly related with the higher frequency of smoking habit.

With regard to socio-demographic characteristics, the proportion of men and women in our study is not balanced (28.7% of women) and it is similar to previous studies.^{15,16,26,30}

Most patients in our study (91.3%) initiated treatment for secondary prevention with antiplatelets after the first ACS, mainly with dual antiplatelet therapy, as recommended by guidelines.^{8–10} Statins were the second drug more prescribed (85.7% of patients) and beta-blockers and ACEI/ARB were less prescribed. All patients with established

Table 3 Socio-demographic characteristics, laboratory data, comorbidities and comedications stratified by study drugs number.

N (%)	4	3	≤ 2	p-Value
	3847 (47.7)	2569 (31.8)	1655 (20.5)	
<i>Acute myocardial infarction</i>	3310 (86.0)	2035 (79.2)	1130 (68.3)	<0.001
<i>Unstable angina</i>	537 (14.0)	534 (20.8)	525 (31.7)	<0.001
<i>Gender; women</i>	968 (25.2)	706 (27.5)	644 (38.9)	<0.001
<i>Age in years, mean (SD)</i>	63.9 (13.0)	65.2 (13.6)	68.9 (14.4)	<0.001
>75 years	869 (22.6)	695 (27.1)	634 (38.3)	<0.001
<i>MEDEA^{24,25}</i>				<0.001
R	683 (17.8)	412 (16.1)	332 (20.1)	
U1-3	1638 (42.6)	1056 (41.2)	672 (40.6)	
U4-5	1335 (34.7)	929 (36.2)	521 (31.5)	
<i>Smokers*</i>	1234 (35.5)	745 (32.9)	341 (22.8)	<0.001
Missing (10.3%)				
<i>High alcohol intake**</i>	3 (0.1)	1 (0.1)	1 (0.1)	<0.001
Missing (21.8%)				
<i>BMI (kg/m²; mean, SD)</i>	29.3 (4.7)	28.8 (4.7)	28.7 (4.9)	<0.001
Missing (20.8%)				
<i>BMI ≥ 30: obesity</i>	1194 (39.3)	712 (35.8)	481 (35.3)	<0.001
<i>Cholesterol total mg/dL, mean, (SD)</i>	211.7 (42.7)	210.0 (42.8)	201.10 (44.4)	<0.001
Missing (14.8%)				
<i>Cholesterol LDL mg/dL, mean, (SD)</i>	131.5 (35.3)	131.3 (37.6)	122.1 (36.9)	<0.001
Missing (21.5%)				
<i>Cholesterol HDL mg/dL, mean, (SD)</i>	48.5 (12.8)	49.1 (13.1)	50.2 (15.0)	0.001
Missing (19.0%)				
<i>Triglycerides mg/dL, mean, (SD)</i>	159.5 (108.5)	154.00 (102.9)	145.1 (95.4)	<0.001
Missing (17.7%)				
<i>Diabetes mellitus</i>	1077 (28.0)	640 (24.9)	452 (27.3)	0.022
<i>Dyslipidaemia</i>	1686 (43.8)	1108 (43.1)	656 (39.6)	0.014
<i>Heart failure</i>	75 (1.9)	92 (3.6)	129 (7.8)	<0.001
<i>Hypertension</i>	2189 (56.9)	1230 (47.9)	875 (52.9)	<0.001
<i>Peripheral artery disease</i>	164 (4.3)	120 (4.7)	101 (6.1)	0.013
<i>Renal impairment; eGFR <45 ml/min/1.73 m²</i>	156 (4.8)	179 (8.3)	193 (13.3)	<0.001
Missing (14.9%)				
<i>Comedications after the event</i>				
Anticoagulants	188 (4.9)	170 (6.6)	244 (14.7)	<0.001
Calcium channel-blockers	541 (14.1)	405 (15.8)	363 (21.9)	<0.001
Diuretics	748 (19.4)	510 (19.9)	496 (30.0)	<0.001
Drug used in diabetes mellitus	1008 (26.2)	577 (22.5)	412 (24.9)	0.003
NSAID	734 (19.1)	538 (20.9)	355 (21.5)	0.065
Nitrates	1544 (40.1)	940 (36.6)	521 (31.5)	<0.001

p-Value from ANOVA test comparing samples with 4, 3 or 2-1 drugs of interest. ACH, acute coronary heart disease; BMI, body mass index; LDL-C, low density lipoprotein-cholesterol; HDL-C, high density lipoprotein-cholesterol; eGFR, estimated glomerular filtration rate; NSAID, non-steroidal anti-inflammatory drugs; R, Rural; U, Urban.

cardiovascular disease should be treated during hospital admission and after discharge with statins, regardless of their cholesterol values.³¹ ACEI/ARB might be less prescribed as they are not always recommended for all patients, they should be considered in all ST-Elevation Myocardial Infarction patients.⁸⁻¹⁰ All study drugs were more commonly

prescribed in men than women, except for ACEI/ARB, that difference between women and men was slight and not significant, probably related to higher frequency of hypertension in women in our study population, because women were older than men. These results were similar to Lafeber et al.,³² and Sanfélix-Gimeno et al. studies.⁵ Regarding

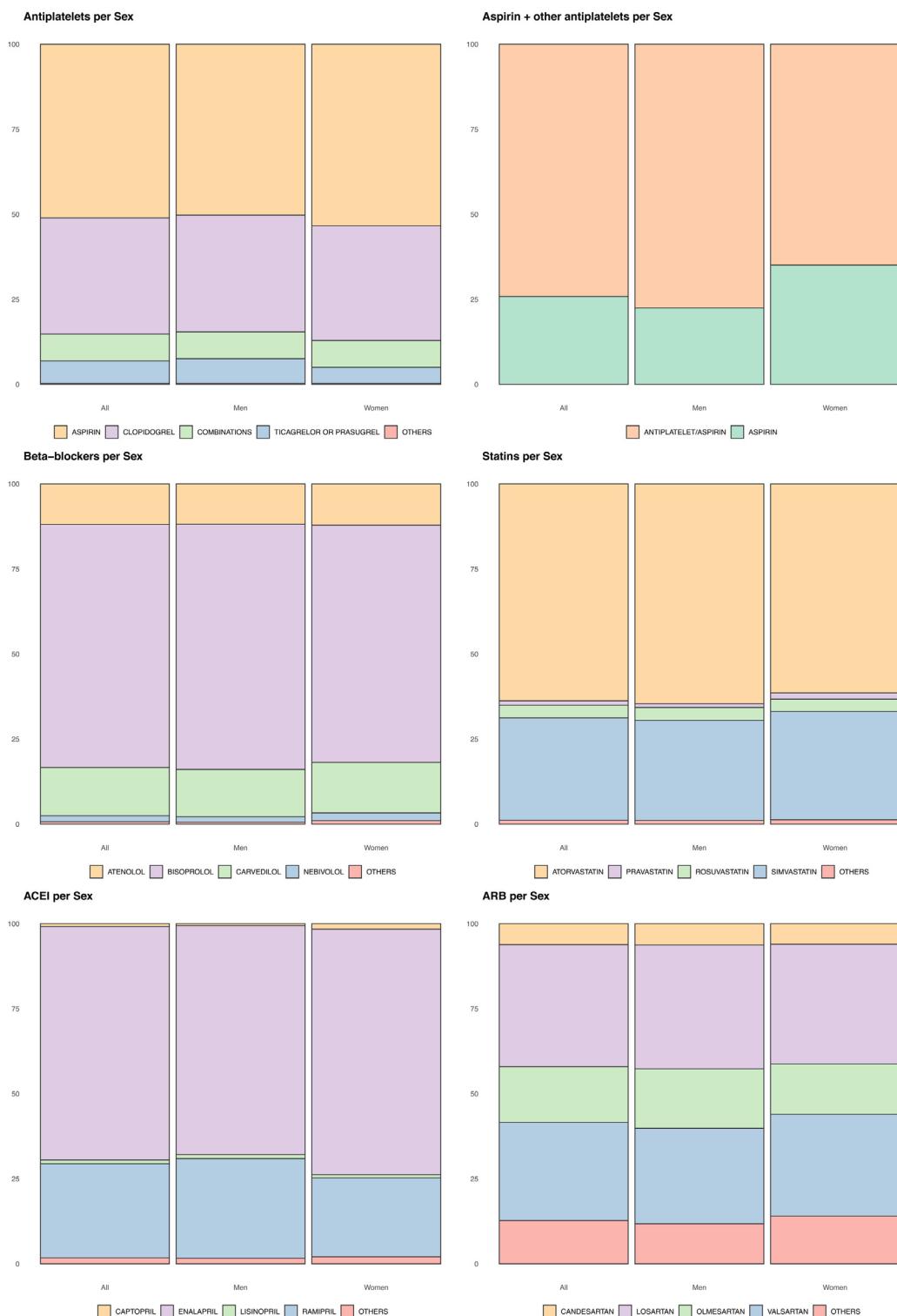


Figure 2 Drugs prescribed per gender. Fig. 2 depicts the different drugs prescribed overall, in men and women. Distribution between genders was compared using the Chi-Square test with all p -values < 0.001 . ACEI, angiotensin-converting enzyme inhibitors; ARB, angiotensin-receptor blockers.

comedications, anticoagulants and diuretics were the most prescribed in women, possibly related with their higher frequency of atrial fibrillation and heart failure than in men. Women initiated secondary prevention less frequently than men.^{14–16,33–35} Nevertheless, the majority of our

population (79.5%) initiated treatment with three or four drugs combined, and almost half (47.7%) with four study drugs, although we found more women treated with \leq two study drugs than with three or four. This may perhaps occur because physicians prescribed fewer drugs to older patients

who were multimorbid and polymedicated.³⁶ Probably, the same assumption could be extended to our finding found for women and the number of drugs prescribed, because men usually suffer ACS at an earlier age.^{37–39}

Zeymer et al.⁴⁰ conducted an observational prospective study including 9998 patients with ACS from June 2000 until December 2002. They reported that patients receiving four drugs were younger and patient's characteristics according to the number of drugs prescribed were similar to our population. They found higher percentage (92.5%) with combination of four or three components and 62.6% with combination of four. The combination of beta-blockers, statins and antiplatelets was also high (39.5%). Also, they suggested that age > 75 years old is a potent predictor for not receiving therapy with four components.^{35,40,41}

Other author already mentioned, Lafeber et al.³² conducted an observational prospective cohort study of 2706 recently diagnosed patients clinically manifest coronary artery disease between January 1996 and February 2010. They found fewer patients (67.0%) treated with the combination of aspirin, a statin and ≥ one blood-pressure lowering agent(s).³²

Aspirin and clopidogrel were the most frequently antiplatelets prescribed. Dual antiplatelet therapy was less frequently prescribed to women as described by previous studies,^{41–43} probably because women were older.⁴⁴ Bisoprolol, enalapril, and losartan were the most prescribed beta-blockers with slight differences between genders. The statins most commonly prescribed overall were atorvastatin and simvastatin, probably because they are the statins with more experience of use.

We found a strong relation in the medication prescribed between being women and older in our population, probably because women had the first ACS in older age than men. Consequently, women had lower probability to be treated with study drugs and higher probability to be treated with other comedications.

This study has some limitations inherent to electronic database studies, such as data incompleteness, loss of follow-up of patients suffering an ACS, potential confounders, non-randomised data and possible selection biases. Other limitation is that prescriptions are not linked with diagnoses in SIDIAP database. Our database has PHC's data, therefore some hospital's data is not available. On the other hand, the strengths of our study are the large number of patients included, representativeness for the general population, complete socio-demographic and health records, long follow-up periods and real-world data. Our data is supported by previous studies and the presence of cardiovascular risk factors and outcomes has been previously validated in SIDIAP.^{45–47}

This is the first work conducted with SIDIAP database which analyses the drugs prescribed for secondary prevention of cardiovascular disease providing high value knowledge about the cardiovascular disease in Catalonia (North-East Spain), which represents more than 5.8 million inhabitants in south Europe. The results can be extrapolated to all population in Catalonia and the rest of Spain, as the health systems and population characteristics are similar.

Ethical requirements

In accordance with European and Spanish legislation on confidentiality and data protection ([EU] 2016/679), the data contained in SIDIAP are always pseudonymised. For the cross-over with the CMBD database, SIDIAP uses a third party to ensure confidentiality.

This study follows all national and international regulations: Declaration of Helsinki and Principles of Good Research Practice.

The study was approved by the IDIAPJGol Research Ethics Committee on May 3, 2017.

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Conflict of interests

The authors declare that they have no conflict of interest in conducting this study.

Appendix A. Supplementary data

Supplementary data associated with this article can be found, in the online version, at doi:10.1016/j.aprim.2021.102157.

References

1. World Health Organization. Global status report on noncommunicable diseases; 2014. Available from: <https://www.who.int/nmh/publications/ncd-status-report-2014/en/>
2. Khan MA, Hashim MJ, Mustafa H, Baniyas MY, Al Suwaidi SKBM, AlKatheeri R, et al. Global epidemiology of ischemic heart disease: results from the global burden of disease study. Cureus. 2020;12:e9349.
3. Mensah GA, Wei GS, Sorlie PD, Fine LJ, Rosenberg Y, Kaufmann PG, et al. Decline in cardiovascular mortality. Circ Res. 2017;120:366–80.
4. Crowley MJ, Zullig LL, Shah BR, Shaw RJ, Lindquist JH, Peterson ED, et al. Medication non-adherence after myocardial infarction: an exploration of modifying factors. J Gen Intern Med. 2015;30:83–90.
5. Sanfélix-Gimeno G, Peiró S, Ferreros I, Pérez-Vicente R, Librero J, Catalá-López F, et al. Adherence to evidence-based therapies after acute coronary syndrome: a retrospective population-based cohort study linking hospital, outpatient, and pharmacy health information systems in Valencia, Spain. J Manag Care Pharm. 2013;19:247–57.
6. Bansilal S, Castellano JM, Garrido E, Wei HG, Freeman A, Spettell C, et al. Assessing the impact of medication adherence on long-term cardiovascular outcomes. J Am Coll Cardiol. 2016;68:789–801.
7. Banerjee A, Khandelwal S, Nambiar L, Saxena M, Peck V, Moniruzzaman M, et al. Health system barriers and facilitators to medication adherence for the secondary prevention of cardiovascular disease: a systematic review. Open Hear. 2016;3:e000438.
8. Ibanez B, James S, Agewall S, Antunes MJ, Bucciarelli-Ducci C, Bueno H, et al. 2017 ESC Guidelines for the management

- of acute myocardial infarction in patients presenting with ST-segment elevation. *Eur Heart J.* 2018;39:119–77.
9. Roffi M, Patrono C, Collet J-P, Mueller C, Valgimigli M, Andreotti F, et al. Guía ESC 2015 sobre el tratamiento de los síndromes coronarios agudos en pacientes sin elevación persistente del segmento ST. *Rev Española Cardiol.* 2015;68:1125e1–64.
 10. National Institute for Health and Care Excellence (NICE). Myocardial ocardial infarction: cardiac rehabilitation and prevention of further cardiovascular disease; 2013.
 11. Amsterdam EA, Wenger NK, Brindis RG, Casey DE, Ganiats TG, Holmes DR, et al. 2014 AHA/ACC guideline for the management of patients with non-ST-elevation acute coronary syndromes: executive summary. *J Am Coll Cardiol.* 2014;64:2645–87.
 12. Ghimire G, Gupta A, Hage FG. Guidelines in review: 2013 ACCF/AHA guideline for the management of ST-elevation myocardial infarction. *J Nucl Cardiol.* 2014;21:190–1.
 13. Lafeber M, Spiering W, Graaf Y, Van Der. The combined use of aspirin, a statin, and blood pressure-lowering agents (polypill components) and the risk of vascular morbidity and mortality in patients with coronary artery disease. *Am Heart J.* 2013;166:282–9, e1.
 14. García-García C, Molina L, Subirana I, Sala J, Bruguera J, Arós F, et al. Sex-based differences in clinical features management, and 28-day and 7-year prognosis of first acute myocardial infarction. RESCATE II study. *Rev Esp Cardiol.* 2014;67:28–35.
 15. Lahoz C, Mantilla T, Taboada M, Soler B, Tranche S, López-Rodríguez I, et al. Gender differences in evidence-based pharmacological therapy for patients with stable coronary heart disease. *Int J Cardiol.* 2009;133:336–40.
 16. Plaza-Martín M, Sanmartín-Fernandez M, Álvarez-Álvarez B, Andrea R, Seoane-García T, González-D'Gregorio J, et al. Contemporary differences between men and women with acute coronary syndromes: CIAM multicenter registry. *J Cardiovasc Med (Hagerstown).* 2019;20:525–30.
 17. ten Haaf ME, Bax M, ten Berg JM, Brouwer J, van't Hof AW, van der Schaaf RJ, et al. Sex differences in characteristics and outcome in acute coronary syndrome patients in the Netherlands. *Netherlands Hear J.* 2019;27:263–71.
 18. Giner-Soriano M, Sotorra Figuerola G, Cortés J, Pera Pujadas H, García-Sangenís A, Morros R. Impact of medication adherence on mortality and cardiovascular morbidity: protocol for a population-based cohort study. *JMIR Res Protoc.* 2018;7:e73.
 19. SIDAPI. SIDAPI. Information system for research in Primary Care. SIDAPI. 2020.
 20. WHO. ICD-10 Version: 2016 [Internet]. International Statistical Classification of diseases and Related Health Problems 10th Revision. s.f.
 21. WHO Collaborating Centre for Drug Statistics Methodology. ATC/DDD Index. 2019;2019.
 22. CatSalut. Servei Català de la Salut. Conjunt mínim bàsic de dades (CMBD). 2019.
 23. MSSSI M de SSS e I. Clasificación internacional de enfermedades 9a revisión, modificación clínica (eCIE9MC). eCIE9MC. 2014.
 24. Domínguez-Berjón M, Borrell C, Cano-Serral G, Esnaola S, Nolasco A, Pasarin M, et al. Construcción de un índice de privación a partir de datos censales en grandes ciudades españolas (Proyecto MEDEA). *Gac Sanit.* 2008;22:179–87.
 25. Caro-Mendivelso J, Elorza-Ricart JM, Hermosilla E, Méndez-Boo L, García-Gil MP-AD. Associations between socioeconomic index and mortality in rural and urban small geographic areas of Catalonia Spain: ecological study. *J Epidemiol Res.* 2015;2:80.
 26. Ribas N, García-García C, Meroño O, Recasens L, Pérez-Fernández S, Bazán V, et al. Secondary prevention strategies after an acute ST-segment elevation myocardial infarction in the AMI code era: beyond myocardial mechanical reperfusion. *BMC Cardiovasc Disord.* 2017;17:54.
 27. Valero-Masa MJ, Velásquez-Rodríguez J, Diez-Delhoyo F, Devesa C, Juárez M, Sousa-Casasnovas I, et al. Sex differences in acute myocardial infarction: is it only the age? *Int J Cardiol.* 2017;231:36–41.
 28. Plaza-Martín M, Sanmartín-Fernandez M, Álvarez-Álvarez B, Andrea R, Seoane-García T, González-D'Gregorio J, et al. Contemporary differences between men and women with acute coronary syndromes: CIAM multicenter registry. *J Cardiovasc Med.* 2019;20:525–30.
 29. Wilkinson C, Bebb O, Dondo TB, Munyombwe T, Casadei B, Clarke S, et al. Sex differences in quality indicator attainment for myocardial infarction: a nationwide cohort study. *Heart.* 2019;105:516–23.
 30. Zagnoni S, Casella G, Pallotti MG, Gonçzini L, Abrignani MG, Caldarola P, et al. Sex differences in the management of acute coronary syndromes in Italy: data from the MANTRA registry. *J Cardiovasc Med.* 2017;18:178–84.
 31. Catapano AL, Graham I, De Backer G, Wiklund O, Chapman MJ, Drexel H, et al. 2016 ESC/EAS guidelines for the management of dyslipidemias. *Rev Esp Cardiol.* 2017;70, 115.e.
 32. Lafeber M, Spiering W, Van Der Graaf Y, Nathoe H, Bots ML, Grobbee DE, et al. The combined use of aspirin, a statin, and blood pressure-lowering agents (polypill components) and the risk of vascular morbidity and mortality in patients with coronary artery disease. *Am Heart J.* 2013;166:282–9, e1.
 33. Timmis A, Townsend N, Gale C, Grobbee R, Maniadakis N, Flather M, et al. European Society of Cardiology: cardiovascular disease statistics 2017. *Eur Heart J.* 2018;39:508–77.
 34. Hao Y, Liu J, Liu J, Yang N, Smith SC, Huo Y, et al. Sex differences in in-hospital management and outcomes of patients with acute coronary syndrome. *Circulation.* 2019;139:1776–85.
 35. Lee HY, Cooke CE, Robertson TA. Use of secondary prevention drug therapy in patients with acute coronary syndrome after hospital discharge. *J Manag Care Pharm.* 2008;14: 271–80.
 36. Cherubini A, Corsonello A, Lattanzio F. Underprescription of beneficial medicines in older people: causes, consequences and prevention. *Drugs Aging.* 2012;29:463–75.
 37. Vasiljevic Pokrajcic Z, Mickovski N, Davidovic G, Asanin M, Stefanovic B, Krljanac G, et al. Sex and age differences and outcomes in acute coronary syndromes. *Int J Cardiol.* 2016;217:S27–31.
 38. Vogel B, Farhan S, Hahne S, Kozanli I, Kalla K, Freynhofer MK, et al. Sex-related differences in baseline characteristics, management and outcome in patients with acute coronary syndrome without ST-segment elevation. *Eur Hear J Acute Cardiovasc care.* 2016;5:347–53.
 39. Cenko E, Yoon J, Kedev S, Stankovic G, Vasiljevic Z, Krljanac G, et al. Sex differences in outcomes after STEMI: effect modification by treatment strategy and age. *JAMA Intern Med.* 2018;178:632–9.
 40. Zeymer U, Jünger C, Zahn R, Bauer T, Bestehorn K, Senges J, et al. Effects of a secondary prevention combination therapy with an aspirin, an ACE inhibitor and a statin on 1-year mortality of patients with acute myocardial infarction treated with a beta-blocker Support for a polypill approach. *Curr Med Res Opin.* 2011;27:1563–70.
 41. Pereira M, Araújo C, Dias P, Lunet N, Subirana I, Marrugat J, et al. Age and sex inequalities in the prescription of evidence-based pharmacological therapy following an acute coronary syndrome in Portugal: the EURHOBOP study. *Eur J Prev Cardiol.* 2014;21:1401–8.
 42. Wilkinson C, Bebb O, Dondo TB, Munyombwe T, Casadei B, Clarke S, et al. Sex differences in quality indicator attainment for myocardial infarction: a nationwide cohort study Coronary artery disease. *Heart.* 2019;105:516–23.
 43. Chow CK, Brieger D, Ryan M, Kangaharan N, Hyun KK, Briffa T. Secondary prevention therapies in acute coronary syndrome and relation to outcomes: observational study. *Heart Asia.* 2019;11.

44. Saunderson CED, Brogan RA, Simms AD, Sutton G, Batin PD, Gale CP. Acute coronary syndrome management in older adults: guidelines, temporal changes and challenges. *Age Ageing.* 2014.
45. Ramos R, Balló E, Garcí M, Martí R, Fina F, Hermosilla E, et al. Validity for use in research on vascular diseases of the SIDIAP (information system for the development of research in primary care): the EMMA study. *Rev Esp Cardiol (Engl Ed).* 2017;65:29–37.
46. Vinagre I, Mata-Cases M, Hermosilla E, Morros R, Fina F, Rosell M, et al. Control of glycemia and cardiovascular risk factors in patients with type 2 diabetes in primary care in Catalonia (Spain). *Diabetes Care.* 2012;35:774–9.
47. Bolíbar B, Fina Avilés F, Morros R, del Mar Garcia-Gil M, Hermosilla E, Ramos R, et al. Base de datos SIDIAP: la historia clínica informatizada de Atención Primaria como fuente de información para la investigación epidemiológica. *Med Clin (Barc).* 2012;138:617–21.