Impact of gender difference in hospital outcomes following percutaneous coronary intervention. Results of the Belgian Working Group on Interventional Cardiology (BWGIC) registry

Mathieu Lempereur[1], MD; Julien Magne[2], PhD; Kristoff Cornelis[3], MD; Claude Hanet[4], MD, PhD; Yves Taeymans[5], MD, PhD; Mathias Vrolix[6], MD, FSAIC; Victor Legrand[1]*, MD, PhD; on behalf of the Belgian Working Group on Interventional Cardiology

1. University Hospital of Liege, Liege, Belgium; 2. CHU Limoges, Hôpital Dupuytren and INSERM 1094, Faculté de Médecine de Limoges, Limoges, France; 3. AZ Maria Middelares Ghent, Ghent, Belgium; 4. Cliniques Universitaires Mont-Godinne, Yvoir, Belgium; 5. University Hospital Ghent, Ghent, Belgium; 6. Ziekenhuis Oost Limburg, Genk, Belgium

KEYWORDS

- coronary artery disease
- gender
- outcomes
- percutaneous coronary intervention

Abstract

Aims: To determine whether there are gender-based differences in in-hospital outcomes among patients undergoing percutaneous coronary intervention (PCI).

Methods and results: We studied a large cohort using clinical data from a registry of 130,985 PCI procedures in Belgium, from January 2006 to February 2011. Compared to males, females were significantly older (70.3 vs. 64.8 years), and were more frequently diabetic or hypertensive. Men smoked more and more frequently had previous myocardial infarction (MI), previous PCI or previous coronary artery bypass graft (CABG) surgery. Coronary artery disease (CAD) was less severe in women, and PCI to the left anterior descending artery was more common in female patients. Unadjusted in-hospital mortality rates were higher in females versus males (2.5% for women and 1.6% for men, p<0.0001). After multivariable analysis, female gender remained an independent predictor of mortality (odds ratio 1.35, 95% CI: 1.22-1.49, p<0.0001).

Conclusions: Gender-based differences in hospital mortality rates after PCI were observed in this large registry. Female sex remained an independent predictor of mortality after multivariable adjustment.

*Corresponding author: Department of Cardiology, University Hospital Sart-Tilman, B-4000 Liege, Belgium.
E-mail: vlegrand@chu.ulg.ac.be*

Introduction

Despite significant advances in the management of coronary artery disease (CAD), this pathology is still one of the major causes of death worldwide today[1]. Cardiovascular disease is the first cause of death in women in Europe. According to the European Heart Network, cardiovascular disease accounts for 42% of deaths in women under 75 years of age[2]. Even though the prevalence of CAD is greater in men than in women at any given age, several reports have shown that female gender is associated with worse outcomes in case of ischaemic heart disease[3,4]. Gender-related differences in the pathophysiology of atherosclerosis[5], in cardiovascular risk factors[6], in clinical presentation[7] and in the diagnosis of coronary artery disease[8] are well known. Recently, analyses from the PROSPECT study showed that there were gender-specific differences in the extent and composition of coronary plaques in patients <65 years of age, suggesting that atherosclerosis development and progression were influenced by gender[9,10]. Differences according to sex have also been outlined in the management and outcomes after acute coronary syndromes[11]. Concerning outcomes after percutaneous coronary intervention (PCI), the question is still debated and controversial. In earlier studies, female gender was identified as an independent predictor of adverse outcome after PCI[12]. In contrast, more recent studies have demonstrated that female gender had no or minimal impact on outcomes after adjustment for clinical and anatomical variables[13,14].

The aim of this study was to analyse gender differences in in-hospital outcomes in patients who underwent PCI in Belgium from January 2006 to February 2011.

Methods

PATIENT POPULATION

The Belgian Working Group on Interventional Cardiology registry is a prospective procedure-based database. A web-based standardised case record form (CARDS database, European Heart House, Nice, France) is completed for each procedure, allowing collection of detailed data in all patients undergoing PCI in the 36 Belgian hospitals performing PCI. Data collection includes clinical, demographic, procedural and angiographic characteristics as well as in-hospital outcomes.

We analysed data from January 2006 to February 2011. During this period, 131,119 procedures were performed. Of these, 134 procedures were excluded from the analysis due to missing gender data.

ENDPOINTS AND STUDY DEFINITIONS

The primary outcome for this study was all-cause in-hospital mortality. Secondary endpoints included the need for coronary artery bypass grafting, cerebrovascular accident (CVA), major bleeding, major vascular complication, renal failure requiring dialysis, post-procedural myocardial infarction and acute stent thrombosis. A composite secondary endpoint only including major adverse cardiac and cerebrovascular events (MACCE) was also analysed. Standard definitions were used to define the most common clinical scenarios.

STATISTICAL ANALYSIS

Categorical variables are presented as counts and percentages. Continuous variables are presented as mean±SD. Differences were deemed significant if p<0.05. We used the chi-square test and Student's t-test to compare groups, as appropriate. Regression analysis was performed using univariable then multivariable logistic regression modelling in order to identify the independent predictors of death and secondary endpoints. Multivariable models were formed by including significant univariable predictors for the outcome variable of interest. Variables with a value of p<0.10 in univariate analysis were incorporated into the model in a forward stepwise manner. All statistical analyses were performed with SPSS version 17 (SPSS Inc., Chicago, IL, USA).

Results

BASELINE, ANGIOGRAPHIC AND INTERVENTION CHARACTERISTICS

Among the 130,985 PCI performed, 35,955 were performed in women. Compared with men, women were significantly older and were more likely to have diabetes, hypertension, chronic renal failure (CRF), prior stroke, previous valvular heart disease (VHD) or previous congestive heart failure. Conversely, history of prior MI, prior PCI, prior CABG surgery or prior peripheral vascular disease (PVD) were less frequent in women than in men. Women were also less likely to be smokers. Of note, the proportion of patients with hypercholesterolaemia was similar in both groups **(Table 1)**.

Regarding PCI indication, most cases were elective procedures in both groups but men were more likely to present with ST-segment elevation myocardial infarction (STEMI). Cardiogenic shock and inotropic support at presentation were more frequent in women. Reduced left ventricle function was more common in male patients. A significant gender-based difference was observed in the severity of CAD, with more frequent single-vessel disease in women and more high-risk coronary artery disease (defined as left main artery stenosis, 3-vessel disease, 2-vessel disease with proximal left anterior descending artery disease) in men. The lesions more often involved the left anterior descending coronary artery in women and more frequently the left circumflex artery and grafts in men. The procedural data showed a slightly greater number of treated vessels and treated lesions per patient in men **(Table 2)**.

CLINICAL OUTCOMES

In-hospital death rate was 1.8% (n=2,297), and women had a significantly higher risk of death: 2.5% vs. 1.6% in men (p<0.0001), odds ratio (OR) 1.57, 95% confidence interval (CI): 1.44-1.70, p<0.0001 **(Table 3**, **Figure 1)**. Women had a higher risk of in-hospital death after PCI in all clinical settings (i.e., acute coronary syndrome [ACS] and stable angina), but the difference in death rate was particularly high in case of acute coronary syndrome, especially in case of STEMI. There was a graded relationship between mortality rates and clinical indication for PCI: elective PCI (0.7% vs. 0.5%, p=0.003), to NSTE-ACS (2.4% vs.

Table 1. Baseline clinical characteristics, clinical indication for PCI and lesion characteristics for the entire cohort.

Characteristic		Total (n=130,985)	Males (n=95,030) 72.6%	Females (n=35,955) 27.4%	*p*-value for females vs. males
Baseline clinical characteristics					
Age, years		66.3±11.8	64.8±11.6	70.3±11.3	<0.0001
Height, cm		169.8±8.7	173.2±6.9	160.9±6.5	<0.0001
Weight, kg		79.3±15.1	82.6±14.2	70.8±13.9	<0.0001
BMI, kg/m²		27.4±4.5	27.5±4.3	27.4±5.2	0.008
Obesity, %		24.2	23.4	26.3	<0.0001
BSA, m²		1.93±0.21	1.99±0.19	1.77±0.19	<0.0001
Previous angina, %		4.9	4.8	5.2	0.002
Previous MI, %		16.3	17.9	12.3	<0.0001
CRF, %		3.3	3.1	3.6	<0.0001
Previous CHF, %		2.1	2.1	2.4	0.001
Previous PCI, %		28.6	29.9	25.4	<0.0001
Previous CVA, %		3.1	3.0	3.4	<0.0001
Previous CABG surgery, %		9.9	10.8	7.4	<0.0001
Previous PVD, %		7.6	7.9	6.8	<0.0001
Previous VHD, %		2.1	1.8	2.9	<0.0001
Current smoking, %		27.8	30.8	19.8	<0.0001
Former smoking, %		31.0	36.8	15.4	<0.0001
Diabetes, %		22.1	20.3	26.8	<0.0001
Insulin-dependent diabetes, %		5.0	4.2	7.4	<0.0001
Non-insulin-dependent diabetes, %		17.1	16.1	19.4	
Hypertension, %		55.8	52.9	63.3	<0.0001
Hypercholesterolaemia, %		58.6	58.4	58.9	0.112
Initial evaluation					
Pre-procedural elevated biomarkers, %		24.4	24.2	24.8	0.226
Pre-procedural thrombolysis (STEMI), %		3.7	3.9	3.1	0.002
Cardiogenic shock, %		1.9	1.9	2.1	0.001
Pre-procedural cardiopulmonary reanimation, %		1.5	1.6	1.5	0.748
Haemodynamic support, %		1.1	1.1	1.3	0.003
LV ejection fraction, %	>50 %	78.1	77.1	80.5	<0.0001
	≤50%	21.9	22.9	19.5	<0.0001
	41-50%	12.1	12.7	10.4	
	31-40%	6.0	6.2	5.6	
	<= 30%	3.8	3.9	3.4	
Clinical indications for PCI, %					
STEMI		16.6	17.3	14.9	<0.0001
NSTE-ACS		13.6	13.5	14.0	
Stabilised ACS		19.0	18.7	19.6	
Elective indications, others		50.8	50.5	51.5	
Lesion characteristics					
Number of diseased vessels, %	1	47.1	45.4	51.5	<0.0001
	2	31.0	31.4	29.8	
	3	21.9	23.1	18.7	
Coronary anatomic risk *, %	High	24.3	25.4	21.2	<0.0001
	Low	75.7	74.6	78.8	

* High anatomic risk: left main artery stenosis, 3-vessel disease, 2-vessel disease with proximal left anterior descending artery disease; low anatomic risk: other 2-vessel disease, 1-vessel disease. ACS: acute coronary syndrome; BMI: body mass index; BSA: body surface area; CABG: coronary artery bypass graft; CHF: congestive heart failure; CRF: chronic renal failure; CVA: cerebrovascular accident; LV: left ventricle; MI: myocardial infarction; NSTE-ACS: non-ST-segment elevation acute coronary syndrome; PCI: percutaneous coronary intervention; PVD: peripheral vascular disease; STEMI: ST-segment elevation myocardial infarction; VHD: valvular heart disease

Table 2. Lesion and procedural characteristics for the entire cohort.

	Characteristic	Total (n=130,985)	Males (n=95,030) 72.6%	Females (n=35,955) 27.4%	*p*-value for females vs. males
Lesion characteristics					
Number of diseased vessels, %	1	47.1	45.4	51.5	
	2	31.0	31.4	29.8	<0.0001
	3	21.9	23.1	18.7	
Coronary anatomic risk*, %	High	24.3	25.4	21.2	
	Low	75.7	74.6	78.8	<0.0001
Procedural characteristics					
Number of stented lesions (number of stented coronary segments), %	1	73.5	73.0	74.8	
	2	21.5	21.9	20.4	
	3	4.0	4.0	3.8	<0.0001
	>3	0.6	0.6	0.7	
Treated vessel, %	Right coronary	34.8	34.6	35.4	0.005
	Left anterior descending	41.4	40.0	45.2	<0.0001
	Left circumflex	26.6	28.1	22.9	<0.0001
	Left main	2.2	2.1	2.2	0.246
	Arterial graft	0.4	0.5	0.3	<0.0001
	Venous graft	2.9	3.3	1.9	<0.0001
Number of treated vessels, %	1	91.3	91.1	91.8	
	2	7.8	8.0	7.3	
	3	0.5	0.5	0.5	<0.0001
	4	0.0	0.0	0.0	
Bifurcation lesion, %		15.3	15.5	14.9	0.085
Intra-stent restenosis, %		7.2	7.5	6.5	<0.0001
Vascular access, %	Femoral	81.2	80.6	82.7	
	Radial	15.3	15.9	13.8	<0.0001
	Brachial, others	3.5	3.5	3.5	
	Closure device, %	26.1	26.4	25.5	<0.0001

* High anatomic risk: left main artery stenosis, 3-vessel disease, 2-vessel disease with proximal left anterior descending artery disease; low anatomic risk: other 2-vessel disease, 1-vessel disease.
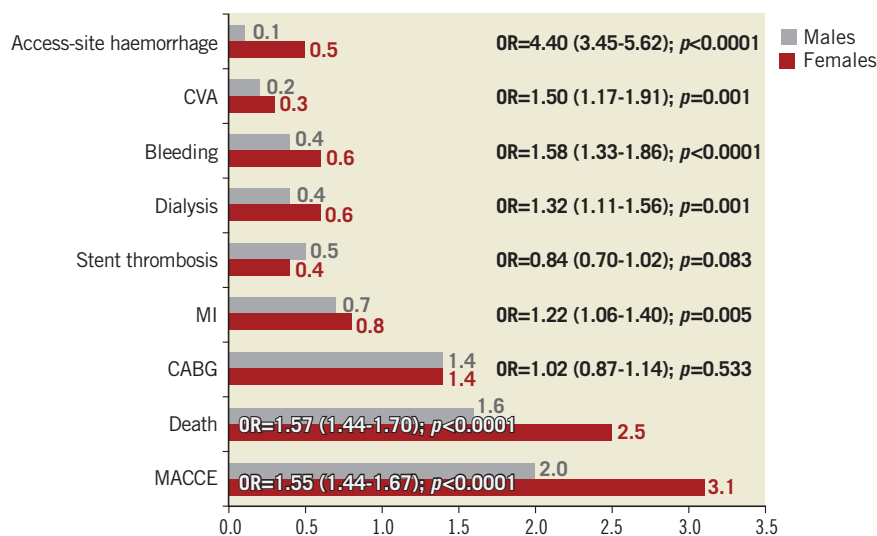

Figure 1. *In-hospital complication rates after PCI for females vs. males and odds ratios for in-hospital complications for females vs. males. CABG: coronary artery bypass graft; CVA: cerebrovascular accident; MACCE: major adverse cardiac and cerebrovascular events; MI: myocardial infarction*

Table 3. Major in-hospital complications and outcomes: incidence and odds ratios for females vs. males.

Variable	Total	Males	Females	*p*-value for females vs. males
Post-procedural MI, %	0.7	0.7	0.8	0.005
Bleeding during hospital stay, %	0.4	0.4	0.6	<0.0001
CVA, %	0.2	0.2	0.3	0.001
Renal failure requiring dialysis, %	0.5	0.4	0.6	0.001
Stent thrombosis, %	0.5	0.5	0.4	0.083
CABG, %	1.4	1.4	1.4	0.533
In-hospital death, %	1.8	1.6	2.5	<0.0001
STEMI, %	6.6	5.4	10.1	<0.0001
NSTE-ACS, %	2.0	1.8	2.4	0.018
Stabilised ACS, %	1.2	1.0	1.6	<0.0001
Elective PCI, others, %	0.5	0.5	0.7	0.003
MACCE (death, stroke, bleeding), %	2.3	2.0	3.1	<0.0001
STEMI, %	7.4	6.1	11.3	<0.0001
NSTE-ACS, %	2.5	2.3	3.1	0.004
Stabilised ACS, %	1.7	1.4	2.3	<0.0001
Elective PCI, others, %	0.9	0.8	1.1	0.001

Odds ratios for females vs. males	OR	95% CI		*p*-value
Post-procedural MI	1.22	1.06	1.40	0.005
Bleeding during hospital stay	1.58	1.33	1.86	<0.0001
CVA	1.50	1.17	1.91	0.001
Renal failure requiring dialysis	1.32	1.11	1.56	0.001
Stent thrombosis	0.84	0.70	1.02	0.083
CABG	0.97	0.87	1.07	0.533
In-hospital death	1.57	1.44	1.70	<0.0001
MACCE (death, stroke, bleeding)	1.55	1.44	1.67	<0.0001

ACS: acute coronary syndrome; CABG: coronary artery bypass graft; CI: confidence interval; CVA: cerebrovascular accident; MACCE: major adverse cardiac and cerebrovascular events; MI: myocardial infarction; NSTE-ACS: non-ST-segment elevation acute coronary syndrome; STEMI: ST-segment elevation myocardial infarction

Table 4. Odds ratios for multivariable predictors of in-hospital events.

Variables	OR	95% CI		*p*-value
Death* (n=2,297)				
Hypercholesterolaemia	0.590	0.534	0.653	<0.0001
Previous PCI	0.634	0.554	0.726	<0.0001
Age	1.047	1.043	1.052	<0.0001
Female gender	1.346	1.216	1.490	<0.0001
Previous CVA	1.346	1.088	1.664	0.006
Previous MI	1.348	1.172	1.551	<0.0001
Previous CHF	1.351	1.074	1.699	0.010
Previous PVD	1.465	1.253	1.712	<0.0001
High coronary anatomic risk	1.642	1.487	1.813	<0.0001
Valvulopathy	1.737	1.392	2.167	<0.0001
CRF	2.471	2.064	2.959	<0.0001
STEMI	4.322	3.895	4.796	<0.0001
Cardiogenic shock	29.492	26.362	32.994	<0.0001
CVA¶ (n=260)				
Previous PCI	0.683	0.500	0.933	0.017
Age	1.020	1.009	1.032	<0.0001
Female gender	1.366	1.051	1.774	0.019
Previous CABG	1.600	1.106	2.314	0.013
CRF	1.771	1.100	2.851	0.019
STEMI	3.033	2.324	3.958	<0.0001
Previous CVA	4.193	2.893	6.076	<0.0001
Bleeding‡ (n=552)				
Age	1.024	1.016	1.032	<0.0001
Female gender	1.365	1.141	1.632	0.001
Previous PVD	1.490	1.144	1.941	0.003
Previous CVA	1.524	1.069	2.170	0.020
CRF	2.244	1.650	3.052	<0.0001
STEMI	2.667	2.202	3.230	<0.0001
Cardiogenic shock	4.186	3.187	5.499	<0.0001
MACCE (n=2,909)**				
Hypercholesterolaemia	0.648	0.595	0.706	<0.0001
Previous PCI	0.686	0.612	0.768	<0.0001
Age	1.040	1.036	1.044	<0.0001
Previous MI	1.253	1.110	1.413	<0.0001
Female gender	1.352	1.239	1.477	<0.0001
High-risk coronary anatomy	1.433	1.313	1.564	<0.0001
Previous CVA	1.529	1.279	1.828	<0.0001
Valvulopathy	1.568	1.287	1.910	<0.0001
Previous PVD	1.579	1.382	1.804	<0.0001
CRF	2.431	2.081	2.840	<0.0001
STEMI	3.578	3.267	3.918	<0.0001
Cardiogenic shock	23.829	21.452	26.468	<0.0001

*All variables included in the multivariate model were significant and reported in the table.
¶History of peripheral vascular disease was also included in the multivariate model (*p*=NS).
‡History of valvular heart disease and prior PCI were also included in the multivariate model (*p*=NS for both). **MACCE: death, CVA and bleeding. CABG: coronary artery bypass graft; CHF: congestive heart failure; CRF: chronic renal failure; CVA: cerebrovascular accident; MACCE: major adverse cardiac and cerebrovascular events; PCI: percutaneous coronary intervention; PVD: peripheral vascular disease; STEMI: ST-segment elevation myocardial infarction
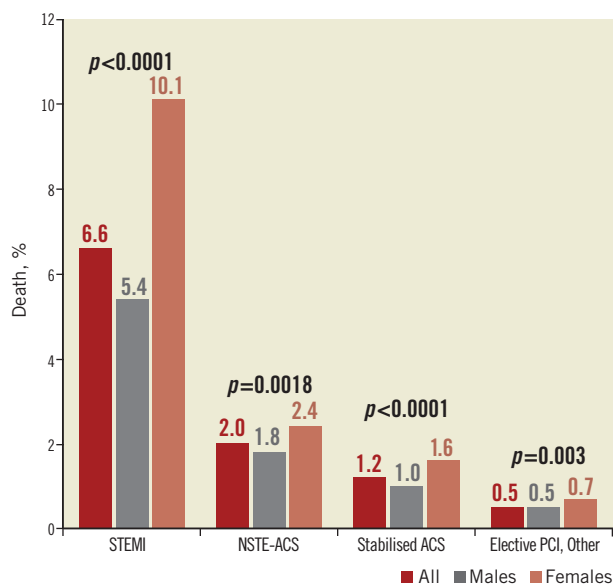
1.8%, p=0.018), to STEMI (10.1% vs. 5.4%, p<0.0001). In each clinical subgroup, a significant gender-related difference was seen, with the worst prognosis in women (p<0.0001) **(Figure 2)**. Women were more likely to have a MACCE (death, stroke or severe bleeding) during hospital stay, with a rate of 3.1% in women and 2.0% in men (OR 1.55, 95% CI: 1.44-1.67, p<0.0001). The rate of severe bleeding was higher in female patients **(Figure 1)**.

After univariate analysis of clinical, angiographic and procedural analyses, we identified independent risk factors for death in multivariate analysis **(Table 4)**. In multivariate analysis, female gender was a significant independent risk factor for in-hospital death (OR 1.35, 95% CI: 1.22-1.49, p<0.0001). The other independent risk factors were age, prior MI (OR 1.35, 95% CI: 1.12-1.56, p<0.0001), hypercholesterolaemia, prior PCI, prior PVD, prior VHD, CRF, high coronary anatomic risk, STEMI and cardiogenic shock. In multivariate analysis, female gender was an independent

Figure 2. *In-hospital mortality rate according to gender and indication for PCI. ACS: acute coronary syndrome; NSTE-ACS: non-ST-segment elevation acute coronary syndrome; PCI: percutaneous coronary intervention; STEMI: ST-segment elevation myocardial infarction.*

risk factor for bleeding (OR 1.37, 95% CI: 1.14-1.63, p=0.001), stroke (OR 1.37, 95% CI: 1.05-1.77, p=0.019) and MACCE (OR 1.35, 95% CI: 1.24-1.48, p<0.0001) **(Table 4)**.

Discussion

From the Belgian PCI registry, the main results of this study are marked gender-based differences in in-hospital mortality rates following PCI. Even after robust multivariable adjustment, female gender remained an independent predictor of mortality, bleeding, stroke and MACCE.

We observed significant differences between men and women in terms of clinical characteristics, angiographic characteristics, interventions and in-hospital outcomes. Baseline characteristics in our study showed that women were older at the time of PCI. It is known that women develop cardiovascular disease seven to ten years later than men, a difference thought to result, at least in part, from the protective role of endogenous oestrogen[15]. As hypertension and diabetes prevalence increases with age, the higher prevalence of hypertension and diabetes in our female cohort is partly in relation to the older age of female patients undergoing PCI. On the other hand, the higher prevalence of CAD in men[16] is consistent with the higher rate of prior MI and PCI observed in this subset of patients. In the literature, the rate of severe CAD and multivessel disease is lower in the female population compared to males[17,18]. This was confirmed in our study: the OR for a high-risk coronary anatomy was 0.79 for women (95% CI: 0.77-0.81, p<0.0001) and coronary multivessel disease was less frequent in female patients. This apparent dissociation between a higher prevalence of cardiovascular risk factors and lower angiographic

severity of ischaemic heart disease in women has been previously described[19]. However, the mechanism involved and the reasons explaining such discrepancy remain unclear. In cohort studies, gender bias could be responsible for decreased access to PCI for women with advanced CAD[11]. The mode of presentation of CAD, which varies according to gender, could also explain this observation: women more frequently have angina pectoris as first symptom whereas men present more frequently with MI, influencing the access to coronary angiography[20]. This difference in the proportions of acute and elective indications for PCI between male and female patients is confirmed in the present cohort.

There is controversy about mortality and adverse event rates between men and women undergoing PCI. Early studies analysing gender differences in clinical outcomes showed that interventions in women were associated with higher rates of complications and higher rates of intervention failure[21]. It has been argued that these results were secondary to unadjusted analyses, clinical characteristics and comorbidities being responsible for the observed differences. Currently, it remains clear that in unadjusted analysis women have poorer outcomes than men following PCI. To identify the role of gender on outcomes, several studies with adjusted analyses for clinical characteristics (i.e., robust multivariate analysis or propensity score analysis) have been reported recently with varying results. Some studies show a persistent gender-related difference in outcomes after adjustment[12,22], while others show that sex is not an independent factor of outcomes[13,23]. The most recent large studies performed in the DES era do not show gender-specific differences in mortality following PCI after adjustments[14,24]. Conversely, female gender was an independent correlate of long-term mortality in the PCI arm of the SYNTAX trial, despite adjustments for risk factors[25,26]. In our study, there is a significant difference in in-hospital mortality between men and women after PCI. Even after multivariate analysis and multiple adjustment for confounding factors, female gender remains an independent predictor of in-hospital death and MACCE.

Mortality rate after PCI varies according to PCI indication, as observed in the European registry Euro Heart Survey of PCI[27]. Significant sex-related difference in mortality is observed in each clinical subgroup, particularly in patients presenting with STEMI. A worse prognosis in women admitted for acute coronary syndrome (ACS) contributes to the difference in mortality rates between women and men.

The difference in mortality rate between genders can be explained partly by older age and more frequent comorbidities in the female cohort. Nevertheless, in our study, female gender is an independent predictor of death after multivariate analysis with an increase in risk of death of 35%. Complications, such as severe bleeding, acute renal failure or stroke are more frequent in female patients, particularly in patients presenting with ACS. These complications may be considered as a factor contributing to the increased mortality rate in females undergoing PCI.

The influence of female gender on outcomes might be explained by differences in coronary physiology, anatomy and in

cardiovascular risk factors. As some of these factors may be under-evaluated or unknown, female gender may appear as an independent predictor of adverse outcome even though it may be associated with other causative factors (hormonal status, endothelial dysfunction, micro-circulation disorders, etc.)[5].

There is a lack of consistency across studies concerning the influence of gender on clinical outcome after PCI. Many earlier studies relied on relatively small samples, were single-centre studies, focused on different populations or did not analyse the same factors. Additionally, thanks to better PCI techniques and medical management, adverse event rates have decreased in high-risk populations, including women. Nowadays, it is statistically more difficult to show significant event rate difference among groups, and large population analyses are needed[28].

Limitations

Our study has some limitations. First, as a multicentre registry, it offers observational data on selected patients undergoing PCI. Thus, it is limited to patients who have had PCI and does not account for the outcomes of women who undergo cardiac catheterisation without intervention and women who are referred for CABG. Second, even though multivariate analysis was performed, we cannot exclude the influence of confounding factors in our results. Finally, SYNTAX scores were not registered and are lacking in the analysis. SYNTAX scores would probably be more sensitive to determine the influence of lesion severity on the outcomes.

Conclusions

In conclusion, the results of this study demonstrate a significant difference in in-hospital mortality rates between men and women undergoing PCI. This difference is particularly marked in patients undergoing PCI for ACS. After robust adjustment for known predictors of mortality, female gender remained an independent predictor of death, being associated with a 35% increase in risk of death as compared to male gender. However, there is a lack of consistency in findings across studies evaluating gender difference in outcomes after PCI. Larger studies in populations treated with current PCI techniques and cardiovascular medical treatment are needed to identify factors explaining these differences in outcomes.

Impact on daily practice

Female gender is associated with worse outcomes in case of ischaemic heart disease and after PCI, particularly in patients undergoing PCI for ACS. After adjustment for clinical and anatomical variables, results vary between studies and the influence of gender as an independent risk factor for outcomes after PCI is still under debate. In daily practice, gender differences in terms of presentation of CAD and outcomes after PCI have to be considered.

Conflict of interest statement

The authors have no conflicts of interest to declare.

References

1. Shaw LJ, Bairey Merz CN, Pepine CJ, Reis SE, Bittner V, Kelsey SF, Olson M, Johnson BD, Mankad S, Sharaf BL, Rogers WJ, Wessel TR, Arant CB, Pohost GM, Lerman A, Quyyumi AA, Sopko G; WISE Investigators. Insights from the NHLBI-Sponsored Women's Ischemia Syndrome Evaluation (WISE) Study: Part I: gender differences in traditional and novel risk factors, symptom evaluation, and gender-optimized diagnostic strategies. *J Am Coll Cardiol.* 2006;47:S4-S20.

2. European Heart Network, Brussels, www.ehnheart.org/media/press-releases/publication/148.html and European Society of Cardiology, Sophia Antipolis. European Cardiovascular Disease Statistics 2008.

3. Hochman JS, Tamis JE, Thompson TD, Weaver WD, White HD, Van de Werf F, Aylward P, Topol EJ, Califf RM. Sex, clinical presentation, and outcome in patients with acute coronary syndromes. Global Use of Strategies to Open Occluded Coronary Arteries in Acute Coronary Syndromes IIb Investigators. *N Engl J Med.* 1999;341:226-32.

4. Roger VL, Farkouh ME, Weston SA, Reeder GS, Jacobsen SJ, Zinsmeister AR, Yawn BP, Kopecky SL, Gabriel SE. Sex differences in evaluation and outcome of unstable angina. *JAMA.* 2000;283:646-52.

5. Shaw LJ, Bugiardini R, Merz CN. Women and ischemic heart disease: evolving knowledge. *J Am Coll Cardiol.* 2009;54:1561-75.

6. Anand SS, Islam S, Rosengren A, Franzosi MG, Steyn K, Yusufali AH, Keltai M, Diaz R, Rangarajan S, Yusuf S; INTERHEART Investigators. Risk factors for myocardial infarction in women and men: insights from the INTERHEART study. *Eur Heart J.* 2008;29:932-40.

7. Regitz-Zagrosek V. Sex and gender differences in symptoms of myocardial ischaemia. *Eur Heart J.* 2011;32:3064-6.

8. Stangl V, Witzel V, Baumann G, Stangl K. Current diagnostic concepts to detect coronary artery disease in women. *Eur Heart J.* 2008;29:707-17.

9. Ruiz-Garcia J, Lerman A, Weisz G, Maehara A, Mintz GS, Fahy M, Xu K, Lansky AJ, Cristea E, Farah TG, Teles R, Botker HE, Templin B, Zhang Z, de Bruyne B, Serruys PW, Stone GW. Age- and gender-related changes in plaque composition in patients with acute coronary syndrome: the PROSPECT study. *EuroIntervention.* 2012;8:929-38.

10. Lansky AJ, Ng VG, Maehara A, Weisz G, Lerman A, Mintz GS, De Bruyne B, Farhat N, Niess G, Jankovic I, Lazar D, Xu K, Fahy M, Serruys PW, Stone GW. Gender and the extent of coronary atherosclerosis, plaque composition, and clinical outcomes in acute coronary syndromes. *JACC Cardiovasc Imaging.* 2012;5:S62-72.

11. Daly C, Clemens F, Lopez Sendon JL, Tavazzi L, Boersma E, Danchin N, Delahaye F, Gitt A, Julian D, Mulcahy D, Ruzyllo W, Thygesen K, Verheugt F, Fox KM; Euro Heart Survey Investigators. Gender differences in the management and clinical outcome of stable angina. *Circulation.* 2006;113:490-8.

12. Peterson ED, Lansky AJ, Kramer J, Anstrom K, Lanzilotta MJ. National Cardiovascular Network Clinical Investigators. Effect of

gender on the outcomes of contemporary percutaneous coronary intervention. *Am J Cardiol.* 2001;88:359-64.

13. Duvernoy CS, Smith DE, Manohar P, Schaefer A, Kline-Rogers E, Share D, McNamara R, Gurm HS, Moscucci M. Gender differences in adverse outcomes after contemporary percutaneous coronary intervention: an analysis from the Blue Cross Blue Shield of Michigan Cardiovascular Consortium (BMC2) percutaneous coronary intervention registry. *Am Heart J.* 2010;159:677-683.e1.

14. Stefanini GG, Kalesan B, Pilgrim T, Räber L, Onuma Y, Silber S, Serruys PW, Meier B, Jüni P, Windecker S. Impact of sex on clinical and angiographic outcomes among patients undergoing revascularization with drug-eluting stents. *JACC Cardiovasc Interv.* 2012;5:301-10.

15. Mendelsohn ME, Karas RH. The protective effects of estrogen on the cardiovascular system. *N Engl J Med.* 1999;340:1801-11.

16. Wenger NK, Speroff L, Packard B. Cardiovascular health and disease in women. *N Engl J Med.* 1993;329:247-56.

17. Hochman JS, McCabe CH, Stone PH, Becker RC, Cannon CP, DeFeo-Fraulini T, Thompson B, Steingart R, Knatterud G, Braunwald. Outcome and profile of women and men presenting with acute coronary syndromes: a report from TIMI IIIB. TIMI Investigators. Thrombolysis in Myocardial Infarction. *J Am Coll Cardiol.* 1997;30:141-8.

18. Zhao XQ, Theroux P, Snapinn SM, Sax FL. Intracoronary thrombus and platelet glycoprotein IIb/IIIa receptor blockade with tirofiban in unstable angina or non-Q-wave myocardial infarction. Angiographic results from the PRISM-PLUS trial (Platelet receptor inhibition for ischemic syndrome management in patients limited by unstable signs and symptoms). PRISM-PLUS Investigators. *Circulation.* 1999;100:1609-15.

19. Mehilli J, Kastrati A, Dirschinger J, Bollwein H, Neumann FJ, Schömig A. Differences in prognostic factors and outcomes between women and men undergoing coronary artery stenting. *JAMA.* 2000;284:1799-805.

20. Lerner DJ, Kannel WB. Patterns of coronary heart disease morbidity and mortality in the sexes: a 26-year follow-up of the Framingham population. *Am Heart J.* 1986;111:383-90.

21. Cowley MJ, Mullin SM, Kelsey SF, Kent KM, Gruentzig AR, Detre KM, Passamani ER. Sex differences in early and long-term results of coronary angioplasty in the NHLBI PTCA Registry. *Circulation.* 1985;71:90-7.

22. Watanabe CT, Maynard C, Ritchie JL. Comparison of short-term outcomes following coronary artery stenting in men versus women. *Am J Cardiol.* 2001;88:848-52.

23. Singh M, Rihal CS, Gersh BJ, Roger VL, Bell MR, Lennon RJ, Lerman A, Holmes DR Jr. Mortality differences between men and women after percutaneous coronary interventions. A 25-year, single-center experience. *J Am Coll Cardiol.* 2008;51:2313-20.

24. Kovacic JC, Mehran R, Karajgikar R, Baber U, Suleman J, Kim MC, Krishnan P, Dangas G, Sharma SK, Kini A. Female gender and mortality after percutaneous coronary intervention: results from a large registry. *Catheter Cardiovasc Interv.* 2012;80:514-21.

25. Farooq V, Serruys PW, Bourantas C, Vranckx P, Diletti R, Garcia Garcia HM, Holmes DR, Kappetein AP, Mack M, Feldman T, Morice MC, Colombo A, Morel MA, de Vries T, van Es GA, Steyerberg EW, Dawkins KD, Mohr FW, James S, Ståhle E. Incidence and multivariable correlates of long-term mortality in patients treated with surgical or percutaneous revascularization in the synergy between percutaneous coronary intervention with taxus and cardiac surgery (SYNTAX) trial. *Eur Heart J.* 2012;33:3105-13.

26. Farooq V, van Klaveren D, Steyerberg EW, Meliga E, Vergouwe Y, Chieffo A, Kappetein AP, Colombo A, Holmes DR Jr, Mack M, Feldman T, Morice MC, Ståhle E, Onuma Y, Morel MA, Garcia-Garcia HM, van Es GA, Dawkins KD, Mohr FW, Serruys PW. Anatomical and clinical characteristics to guide decision making between coronary artery bypass surgery and percutaneous coronary intervention for individual patients: development and validation of SYNTAX score II. *Lancet.* 2013;381: 639-50.

27. de Mulder M, Gitt A, van Domburg R, Hochadel M, Seabra-Gomes R, Serruys PW, Silber S, Weidinger F, Wijns W, Zeymer U, Hamm C, Boersma E. EuroHeart score for the evaluation of in-hospital mortality in patients undergoing percutaneous coronary intervention. *Eur Heart J.* 2011;32:1398-408.

28. Vogt A, Engel HJ, Glunz HG, Sattelberger U, Reil GH, Sechtem U, Sabin G, Senges J, Hanrath P, Neuhaus KL; ArbeitSgemeinschaft Leitender Kardiologischer Krakenhausärzte (ALKK). Early results of coronary angioplasty despite more complex interventions (Registry of The German Community Hospitals 1993-2000). *Am J Cardiol.* 2002;90:1005-9.