

1 **Sex differences in 1-year rehospitalization for heart failure and myocardial infarction**
2 **after primary percutaneous coronary intervention**

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38 Abstract

39 It is unclear whether universal access to primary percutaneous coronary intervention (pPCI)
40 may reduce sex differences in 1-year rehospitalization for heart failure (HF) and myocardial
41 infarction (MI) after ST-elevation myocardial infarction (STEMI). We studied 7,597
42 consecutive STEMI patients (13.8% women, N=1,045) who underwent pPCI from January
43 2007 to December 2013. Cox regression models adjusted for competing risk from death
44 were used to assess sex differences in rehospitalization for HF and MI within 1 year from
45 discharge. Compared with men, women were older (median age 67.6 vs 56.0 years,
46 $P<0.001$) with higher prevalence of co-morbidities and multivessel disease. Women had
47 longer median door-to-balloon time (median 76 vs 66 minutes, $P<0.001$) and were less likely
48 to receive drug-eluting stents (19.5% vs 24.1%, $P=0.001$). Of the medications prescribed at
49 discharge, fewer women received aspirin (95.8% vs 97.6%, $P=0.002$) and P2Y₁₂ antagonists
50 (97.6% vs 98.5%, $P=0.039$), but there were no significant sex differences in other discharge
51 medications. After adjusting for differences in baseline characteristics and treatment, sex
52 differences in risk of rehospitalization for HF attenuated (HR 1.05, 95% CI 0.79-1.40), but
53 persisted for MI (HR 1.68, 95% CI 1.22-2.33), with greater disparity among patients aged
54 ≥ 60 years (HR 1.83, 95% CI 1.18-2.85) than those aged <60 years (HR 1.45, 95% CI 0.84-
55 2.50). In conclusion, in a setting of universal access to pPCI, the adjusted risk of 1-year
56 rehospitalization for HF was similar among the sexes, but women had higher adjusted risk of
57 1-year rehospitalization for MI, especially older women.

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59 Key words: sex differences; rehospitalization for heart failure; rehospitalization for
60 myocardial infarction; primary percutaneous coronary intervention

61 Introduction

62 1 in 4 patients with acute myocardial infarction (MI) are rehospitalized within 12
63 months of discharge¹. Women are known to have higher rate of rehospitalization after acute
64 MI than men^{2,3}. Sex disparities in post-MI rehospitalization were often attributed by women
65 being less likely to receive primary percutaneous coronary intervention (pPCI)⁴. It is unclear
66 if sex differences in outcomes persist in the contemporary era where pPCI is the universal
67 treatment among patients with ST-segment elevation MI (STEMI).

68 Singapore is a country in South East Asia with a population of 5.5 million and a
69 balanced sex distribution (49% men and 51% women in 2017)⁵. The combination of small
70 land mass and economic resources has enabled Singapore's public healthcare system to
71 provide round-the-clock and universal access to pPCI for all patients with STEMI through its
72 nationwide network of pPCI-capable public hospitals since 2007.

73 We sought to determine whether sex differences in 1-year rehospitalization exist
74 among STEMI patients with pPCI in Singapore. Specifically, we assessed the relationship of
75 baseline characteristics and variables associated with STEMI care with 2 cardiac-specific
76 causes of rehospitalization: heart failure (HF) and MI. Furthermore, since prior studies have
77 shown that younger women have poorer outcomes after MI^{6,7,8}, we sought to ascertain
78 whether sex disparities in rehospitalization for HF and MI differed among young (aged <60
79 years) and older (aged ≥ 60 years) patients.

80 Methods

81 This is a retrospective study of patients enrolled in the Singapore MI Registry (SMIR).
82 The SMIR is an ongoing population-based registry established in 2007⁹. It captures acute MI
83 treated by the public and private hospitals, as well as out-of-hospital acute MI deaths. More
84 than 95% of acute MI in Singapore are managed at the public hospitals each year. State
85 legislature mandates data collection on acute MI without the need for prior written informed
86 consent from patients and the quality of acute MI care is closely monitored across all public

87 hospitals by the Ministry of Health. The SMIR identifies MI cases from (i) Hospital Inpatient
88 Discharge Summaries and (ii) cardiac biomarker lists from all hospitals, (iii) claims data and
89 (iv) Casemix and Subvention data from the Ministry of Health, and (v) death data from the
90 Ministry of Home Affairs, based on International Classification of Diseases (ICD) 9th (Clinical
91 Modification) code of 410 and ICD 10th (Australian Modification) code of I21 and I22. To
92 ensure data accuracy and consistency, yearly internal audit is performed by the Registry to
93 ensure interrater reliability of $\geq 95\%$. Detailed data collection method has been described in
94 previous publications^{10,11}.

95 The SMIR data was matched with procedural data from the Singapore Cardiac
96 Databank. Further matching was done with claims data from the Ministry of Health to
97 ascertain rehospitalization outcomes.

98 We included patients who were admitted for STEMI and underwent pPCI from
99 January 2007 to December 2013 in all public hospitals with onsite pPCI capabilities in
100 Singapore. We excluded patients transferred from non-pPCI capable hospitals (N=417,
101 Supplemental Figure 1).

102 The 2 primary endpoints of interest were unplanned fatal and non-fatal
103 rehospitalization for HF and MI within 1 year after discharge for STEMI. The rehospitalization
104 diagnoses were based on claims data submitted to the Ministry of Health by the hospitals.
105 Each patient has a primary diagnosis, which the clinical team deems to be the primary cause
106 of hospitalization, and ≥ 1 secondary diagnoses, which are deemed to be complications that
107 may have arisen during hospitalization. Rehospitalizations with a primary diagnosis of HF or
108 MI were considered in our study. The full list of primary diagnoses is shown in the
109 Supplemental Materials.

110 Baseline characteristics and variables related to STEMI care were compared
111 between the sexes using Wilcoxon Rank Sum test for numeric variables and Chi-Square test
112 for categorical variables. To account for attrition from mortality and circumvent

113 overestimation of event rate, death was treated as a competing event when examining the
114 relationships between sex and time to rehospitalization for HF or MI using cox regression¹².
115 The cox regression models were built hierarchically, starting with sex only (model 1).
116 Subsequently, other demographic variables (age, ethnicity), past medical history
117 (hypertension, diabetes, hyperlipidemia, cardiovascular disease i.e. MI/ PCI/ coronary artery
118 bypass grafting/ stroke/ peripheral arterial disease) and presenting features on admission
119 (Killip class, creatinine, number of narrowed coronary arteries, pre-PCI thrombolysis-in-
120 myocardial-infarct (TIMI) flow grade, coronary artery intervened) were added (model 2).
121 Finally, variables related to STEMI care (door-to-balloon time, use of stent, use of
122 thrombectomy, use of intra-aortic balloon pump, procedure success, use of glycoprotein IIB-
123 IIIA inhibitor, aspirin given at discharge, beta blockers given at discharge, lipid lowering
124 drugs given at discharge, renin-angiotensin system inhibitors given at discharge, P2Y₁₂
125 antagonists given at discharge, highest Killip class during hospitalization, lowest left
126 ventricular ejection fraction (LVEF) during hospitalization) were added (model 3). Among the
127 independent variables included in the cox regression models, pre-procedure TIMI flow grade
128 had the highest proportion of missing data (10%). Missing data were addressed using
129 multiple imputation with 20 imputed datasets and no auxiliary variable based on the Markov
130 Chain Monte Carlo procedure, which assumes that all variables in the imputation models
131 have a joint multivariate normal distribution^{13,14}. Kaplan-Meier survival curves were used to
132 visually assess sex differences in time to rehospitalization for HF and MI. To ascertain
133 whether sex disparities in rehospitalization for HF and MI were equally prevalent among
134 patients aged <60 years and ≥60 years, we tested for interaction between sex and age
135 based on model 3.

136 This study was conducted according to the Helsinki declaration, and the National
137 Healthcare Group Domain Specific Review Board allowed for waiver of patients' consent as
138 the data used were anonymized and analyses were done at a central data repository
139 (National Registry of Diseases Office) with data protection measures in place. All statistical

140 analyses were done using STATA SE (version 13) software. All reported P-values were 2-
141 sided and P-values <0.05 were considered to be statistically significant.

142 Results

143 Between January 2007 and December 2013, there were 7,597 consecutive STEMI
144 patients who underwent pPCI from all public hospitals in Singapore. 1,045 (13.8%) of them
145 were women (Table 1). Compared to men, the median age of women at STEMI onset was a
146 decade older (67.6 vs 56.0 years, $P<0.001$). Women were less likely to have history of MI
147 (7.8% vs 12.1%, $P<0.001$), prior PCI (6.7% vs 10.7%, $P<0.001$) or being current or former
148 smokers (14.0% vs 72.4%; $P<0.001$), but were more likely to have history of hypertension
149 (69.9% vs 48.6%, $P<0.001$), diabetes (43.2% vs 24.5%, $P<0.001$), hyperlipidemia (54.1% vs
150 42.8%, $P<0.001$) and stroke (6.7% vs 3.1%, $P<0.001$). The median creatinine was lower
151 among women (75 vs 92 $\mu\text{mol/L}$, $P<0.001$). Women were less likely to have pre-procedure
152 complete occlusion of the infarct-related artery (TIMI flow grade 0: 66.8% vs 73.0%,
153 $P<0.001$), but more likely to have HF on admission (Killip class \geq II: 20.6% vs 14.9%,
154 $P<0.001$). Although multivessel disease, defined as \geq 2 major epicardial arteries with >50%
155 stenosis, was more common among women at the time of emergent coronary angiography
156 (33.4% vs 32.5% with double vessel disease and 32.1% vs 28.9% with triple vessel disease,
157 $P=0.030$), there was no significant difference in the rate of multivessel PCI performed during
158 hospitalization (4.9% vs 4.6%, $P=0.666$).

159 Women had longer median door-to-balloon (76 vs 66 minutes, $P<0.001$) and
160 symptom-to-balloon (233 vs 192 minutes, $P<0.001$) time (Table 2). HF during hospitalization
161 was more common among women (Killip class \geq II: 15.4% vs 10.2%, $P<0.001$), but not left
162 ventricular systolic dysfunction, defined as left ventricular ejection fraction <50% (64.1% vs
163 61.6%, $P=0.137$). Use of drug-eluting stent (19.5% vs 24.0%, $P=0.001$) and thrombectomy
164 (50.3% vs 56.9%, $P<0.001$) were less common among women. Although the prescription
165 rates of aspirin (95.8% vs 97.6%, $P=0.002$), P2Y₁₂ antagonists (97.6% vs 98.5%, $P=0.039$)

166 and glycoprotein IIb-IIIa inhibitors (24.9% vs 31.9%, $P<0.001$) were lower among women,
167 there were no significant sex differences in beta blockers, lipid lowering drugs and renin-
168 angiotensin system inhibitors prescription at discharge.

169 Women had higher unadjusted risk of rehospitalization for both HF (hazard ratio 1.83,
170 95% confidence interval 1.42-2.35) and MI (HR 1.78, 95% CI 1.30-2.45) (Table 3). After
171 adjusting for baseline characteristics, sex differences were no longer observed for HF
172 rehospitalization (HR 1.05, 95% CI 0.79-1.40), but persisted for MI rehospitalization (HR
173 1.68, 95% CI 1.22-2.33). Further accounting for variables related to STEMI care yielded
174 similar results (HF rehospitalization: HR 1.04, 95% CI 0.78-1.40; MI rehospitalization: HR
175 1.71, 95% CI 1.23-2.38) (Figures 1 and 2).

176 Stratifying by age, women had higher unadjusted risk of rehospitalization for HF than
177 men in the <60 years age group (HR 1.89, 95% CI 1.14-3.13), but not in the ≥ 60 years age
178 group ($P=0.120$) (Table 3). The interaction between sex and age for HF rehospitalization
179 was not statistically significant (P for interaction=0.671). In contrast, women had higher
180 unadjusted risk of rehospitalization for MI than men in the ≥ 60 years age group (HR 1.91,
181 95% CI 1.26-2.91), but not in the <60 years age group ($P=0.090$) (Table 3 and Supplemental
182 Figure 3). The interaction between sex and age for MI rehospitalization was statistically
183 significant (P for interaction=0.029). After adjusting for baseline characteristics, the higher
184 risk of rehospitalization for HF among women in the <60 years age group attenuated
185 ($P=0.272$) (Supplemental Figure 2). However, the higher risk of rehospitalization for MI
186 among women in the ≥ 60 years age group persisted after adjusting for baseline
187 characteristics (HR 1.83, 95% CI 1.18-2.85). The sex difference in risk of rehospitalization
188 for MI among patients aged ≥ 60 years persisted after further accounting for variables related
189 to STEMI care (HR 1.82, 95% CI 1.16-2.86) (Supplemental Figure 3).

190 Discussion

191 Our study found that among patients with STEMI who underwent pPCI, women were
192 older with a higher prevalence of co-morbidities, multivessel disease and heart failure on
193 admission, compared to men. Women had longer ischemic time and were less likely to
194 receive drug-eluting stents (DES), thrombectomy and antiplatelet agents. Women were
195 approximately 1.8 times more likely to be rehospitalized for HF or MI within 1 year of
196 discharge in unadjusted analyses. After adjusting for baseline characteristics, sex
197 differences in risk of rehospitalization for HF attenuated, but persisted in rehospitalization for
198 MI. Further accounting for variables related to STEMI care yielded similar results. The higher
199 risk of rehospitalization for MI among women than men was more pronounced in the ≥ 60
200 years age group than in the < 60 years age group.

201 HF is a common complication of STEMI¹⁵. Besides being generally older than men at
202 the onset of STEMI, women also tend to have HF on admission, longer ischemic time, and
203 higher burden of hypertension and diabetes- all of which are risk factors that have an
204 established association with HF¹⁶. The attenuation of statistical significance after accounting
205 for differences in baseline characteristics suggests that the higher risk of rehospitalization for
206 HF among women was largely explained by their higher baseline risk. We further observed
207 high prescription rates of beta blockers and renin-angiotensin system inhibitors at discharge
208 for both sexes. Notably, $>75\%$ of women and men received beta blockers and renin-
209 angiotensin system inhibitors at discharge, which was in excess to the prevalence of left
210 ventricular systolic dysfunction among women (64.1%) and men (61.6%). As both beta
211 blockers and renin-angiotensin system inhibitors are known to reduce the risk of adverse
212 ventricular remodeling after a large infarct¹⁷, it is plausible that the similarly high prescription
213 rates of beta blockers and renin-angiotensin system inhibitors among both sexes may also
214 be responsible in part for the similar adjusted risk of rehospitalization for HF.

215 In contrast to rehospitalization for HF, the risk of rehospitalization for MI remained
216 higher among women after adjusting for sex differences in baseline characteristics. Despite
217 studies showing that women derive greater benefit from DES than men¹⁸, there were

218 significantly lower use of DES among women in our study. The lower use of DES among
219 women was limited to patients ≥ 60 years of age (women 16.9%, men 23.9%, $P < 0.001$).
220 Procedure success was similarly high in both sexes despite higher rate of pre-procedure
221 complete occlusion of the infarct-related artery among men. In the same vein, the rate of
222 multivessel PCI was similarly low in both sexes despite multivessel disease being more
223 common among women. However, the interaction between sex and complete
224 revascularization for MI rehospitalization was not statistically significant (P for
225 *interaction* = 0.441). Unmeasured variables may be postulated to explain the higher risk of
226 rehospitalization for MI among women in our study. Having typically smaller and less
227 compliant conduit arteries, coupled with concomitant risk factors such as diabetes and
228 complex lesions, put women at a higher risk of restenosis^{19,20}. Hormonal fluctuation,
229 especially during menopause in women, may lead to macrovascular and microvascular
230 alterations, leaving older women vulnerable to a decreased ability to sustain adequate
231 vascular repair²¹. Smooth muscle cell dysfunction is more commonly seen in women, which
232 may lead to impairment of coronary flow reserve²². Spontaneous coronary artery dissection
233 also occurs more frequently in women²³. Although it is a rare cause of MI, a study by Tweet
234 et al. found that revascularization did not protect against recurrent spontaneous coronary
235 artery dissection even in patients presenting with preserved vessel flow²⁴. Women are
236 known to have a higher risk of bleeding than men, which could lead to early discontinuation
237 of dual-antiplatelet therapy (DAPT) with a subsequent increased risk of MI²⁵. Moreover,
238 women tend to have poorer medical adherence than men^{26,27}. Dreyer et al. further observed
239 that women had poorer health and psychosocial status after MI and adjusting for health
240 status and psychosocial status attenuated sex differences in post-MI rehospitalization².

241 Our study covered an unselected population using national data, which were
242 captured in a standardized manner across all hospitals and is expected to yield results with
243 high internal validity. While the long study period of seven years (2007 to 2013) is a strength,
244 progress in management of STEMI has been rapid and hence there are likely to be time-

245 varying trends that may not be fully accounted in our study. Studies have questioned the
246 accuracy of primary diagnoses coded by hospitals²⁸, yet others have shown that major
247 contributory disease conditions can be reliability identified with a positive predictive
248 value >80%²⁹. As higher risk of rehospitalization for MI among women persisted after
249 adjusting for all potential confounders available in our study, we can only conclude that these
250 variables are able to only partially explain the association between sex and rehospitalization
251 for MI and attribute the unexplained sex disparity to other unmeasured variables, such as
252 duration of DAPT, medication adherence and psychosocial status.

253 In conclusion, in a setting of universal access to pPCI, sex disparities persist in
254 STEMI treatment and outcomes. Compared with men, women with STEMI are more likely to
255 experience treatment delay, less likely to receive DES and DAPT, and more likely to be
256 rehospitalized for HF and MI. Sex differences in rehospitalization for HF but not MI appear to
257 be largely explained by their differing baseline risk. To better understand how to mitigate the
258 sex disparity in rehospitalization for MI after pPCI, future studies should focus on sex
259 differences in the complexity and severity of coronary artery disease and evaluate their
260 interaction with the impact of intensification of secondary preventive medical therapy and
261 more complete revascularization in women.

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269 Disclosures

270 There are no potential conflicts of interest, including related consultancies,
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272 Figure legends

273 **Figure 1: Time to rehospitalization for heart failure**

274 Event curves show the adjusted rehospitalization events

275 **Figure 2: Time to rehospitalization for myocardial infarction**

276 Event curves show the adjusted rehospitalization events

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