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# **ORIGINAL RESEARCH**

# RHYTHM DISORDERS AND ELECTROPHYSIOLOGY

# Sex Differences in Ventricular Arrhythmias and Adverse Outcomes Following Acute Myocardial Infarction

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## ABSTRACT

**BACKGROUND** Ventricular arrhythmias (VAs) are a common cause of death in patients with acute myocardial infarction (AMI). Studies have shown sex differences in the incidence, presentation, and outcomes of AMI. However, less is known about sex differences in patients with AMI who develop VAs.

**OBJECTIVES** The authors assessed sex differences in incidence and in-hospital outcomes of patients with AMI and VAs.

**METHODS** Using the National Inpatient Sample 2016 to 2020, we conducted a retrospective analysis of patients admitted for AMI with a secondary diagnosis of VAs. Multivariable logistic regression was performed to estimate the sex-specific differences in the rates and in-hospital outcomes of VAs post-AMI.

**RESULTS** We identified 1,543,140 patients admitted with AMI. Of these, (11.3%) 174,565 patients had VAs after AMI. The odds of VAs after AMI were higher among men (12.6% vs 8.8% adjusted odds ratio [AOR]: 1.72; CI: 1.67-1.78; P < 0.001). Women had significantly higher odds of in-hospital mortality (AOR: 1.32; CI: 1.21-1.42; P < 0.001), cardiogenic shock (AOR: 1.08; CI: 1.01-1.15; P < 0.022), and cardiac arrest (AOR: 1.11; CI: 1.03-1.18; P < 0.002). Women were less likely to receive an implantable cardioverter-defibrillator (ICD) (AOR: 0.57; CI: 0.47-0.68; P < 0.001) or undergo catheter ablation (AOR: 0.51; CI: 0.27-0.98; P < 0.001) during the index admission.

**CONCLUSIONS** We found important sex differences in the incidence and outcomes of VAs among patients with AMI. Women had lower odds of VAs but worse hospital outcomes overall. In addition, women were less likely to receive ICD. Further studies to address these sex disparities are needed. (JACC Adv 2024;3:101042) © 2024 The Authors. Published by Elsevier on behalf of the American College of Cardiology Foundation. This is an open access article under the CC BY-NC-ND license (http://creativecommons.org/licenses/by-nc-nd/4.0/).

entricular arrhythmias (VAS), including ventricular tachycardia (VT) and ventricular fibrillation (VF) occur following 10 to 20% of acute myocardial infarction (AMI) cases and are associated with an increased risk of poor hospital outcomes.<sup>1-4</sup> Data from the SPACE (Saudi Project for Assessment of Coronary Events) registry have shown that patients with AMI who develop VAs are more

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The authors attest they are in compliance with human studies committees and animal welfare regulations of the authors' institutions and Food and Drug Administration guidelines, including patient consent where appropriate. For more information, visit the Author Center.

## ABBREVIATIONS AND ACRONYMS

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AMI = acute myocardial infarction

AOR = adjusted odds ratio

HCUP = Healthcare Cost and Utilization Project

ICD = Implantable cardiac defibrillator

IHD = ischemic heart disease
NIS = National Inpatient

Sample

VA = ventricular arrhythmia VF = ventricular fibrillation

VT = ventricular tachycardia

likely to experience reinfarction, cardiogenic shock, congestive heart failure, stroke, and in-hospital mortality.<sup>5</sup> Congruently, a study using data from the GRACE (Global Registry of Acute Coronary Events) registry showed similar outcomes in patients with AMI who develop VAs, with in-hospital mortality rates as high as 52% as compared to 1.6% in those who did not develop VAs.<sup>6</sup>

Previous studies have explored the sexspecific differences in AMI outcomes, including a higher incidence of VAs in men.<sup>5</sup> However, data addressing sex-specific differences in the in-hospital outcomes of the subset of patients with VAs have been lacking. The FAST-MI (French Registry of Acute

ST-Elevation or Non-ST elevation MI found that women with VAs had higher 1-year mortality but were less likely to receive cardiac interventions.<sup>7</sup> Given these findings, using a national representative sample of the United States, we sought to determine whether sex differences exist in VA incidence and in-hospital outcomes in patients admitted for AMI.

# METHODOLOGY

**STUDY DESIGN AND DATA SOURCE**. This study is reported following the STROBE (Strengthening the Reporting of Observational Studies in Epidemiology) reporting guidelines.<sup>8</sup>

We analyzed hospitalizations between January 1, 2016, and December 31, 2020, in the U.S. National Inpatient Sample (NIS). The NIS is the largest publicly available all-payer inpatient database in the United States and is maintained by the Agency for Healthcare Research and Quality; it is designed as a stratified probability sample of discharges representing nonfederal acute care hospitals nationwide. Samples from these hospitals are recorded and then weighted to ensure that they are nationally representative. Further details on the content and design of the NIS database can be found on the Healthcare Cost and Utilization Project (HCUP) website.<sup>9</sup> Institutional Review Board approval was waived since all patient data in NIS were deidentified.

**STUDY SAMPLE**. The study population consisted of all inpatient hospitalizations with a primary diagnosis



TABLE 1 Gender Differences in the Sociodemographic and Comorbid Characteristics of

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of type 1 AMI who underwent revascularization (percutaneous coronary intervention or coronary artery bypass grafting) and had a secondary diagnosis of VAs (VT or VF). Diagnoses were identified using ICD-10-CM codes (type 1 AMI "I21.0," "I21.1," "I21.2," "I21.3," "I21.4," "I21.9," ventricular tachycardia-"I47.2" and ventricular fibrillation-"I49.0") as recommended by the American College of Cardiology.<sup>10</sup> We excluded patients <18 years and those who had infiltrative cardiac diseases (amyloidosis, sarcoidosis), Lyme disease, or long QT syndrome (Figure 1). Additional codes on revascularization documented are given in Supplemental Table 1.

**VARIABLES.** The study population was divided into 2 subgroups based on the gender/sex. In the NIS, the sex of the patient was recorded as binary, either "female or male," with all nonfemale and nonmale (eg, "other") values set to missing. Hence to maintain uniformity, we consistently used "sex" instead of "gender" throughout this article. Baseline demographic variables including age, sex, race, hospital characteristics, and medical comorbidities (computed from Charlson index for comorbidities) were identified as variables present in the data set. Other comorbidities were obtained using ICD-10-CM and ICD-10-PCS codes (Supplemental Table 1). These codes have been used in prior nationally representative studies and recommended for use by the Centers for Medicare and Medicaid Services.<sup>11,12</sup>

**STUDY OUTCOMES.** The primary endpoint of this study was to determine the sex differences in the rates of VAs among patients with AMI, while the secondary outcomes were sex differences in rates of in-hospital mortality, cardiogenic shock, cardiac arrest, implantable cardiac defibrillator insertion, palliative care consultation, catheter ablation, and length of hospitalization. We also performed a subanalysis to assess the sex differences in the outcomes of patients with AMI and either VT or VF, and the VA outcomes in patients with NSTEMI or STEMI.

**STATISTICAL ANALYSIS.** Data analysis was performed using the HCUP STATA package, which incorporates the NIS-specific variables, including hospital identifiers, stratum, and discharge weights to account for clustering and large survey-weighted data analysis according to HCUP recommendations.<sup>13</sup> Stratified weighted data were used in our analysis using the Svy command to obtain nationwide estimates. Baseline sociodemographic, hospital, and clinical characteristics of patients with AMI and VAs by sex were reported. In our descriptive statistics, continuous variables were presented as means  $\pm$  SD and compared using a Student's *t*-test while

AMI Patients With Ventricular Arrhythmias					
	All Ventricula (N = 174.5				
	Male (n = 132,385, 75.8%)	Female (n = 42,180, 24.2%)	P Value		
Age (y)	$62 \pm 0.2$	$66 \pm 0.3$	< 0.001		
Race			< 0.0001		
White	78.4	77.3			
Blacks	7.76	11.7			
Hispanics	6.99	6.2			
Asians	2.63	1.6			
Native Americans	0.6	0.5			
Others	3.7	2.7			
Median household income			<0.0001		
1	25.8	29.1			
2	26.4	29.0			
3	25.4	23.8			
4	22.3	18.1			
Primary insurance payer	10.04	56.5	<0.0001		
Medicaid	40.84	56.5			
Medicare	10.03	10.86			
Private pay	37.78	25.98			
Self-pay	0.80	4.83			
No-charge	0.54	0.3			
Hospital had size	3.95	1.55	0 6 2 2		
Small	15 / 0	15 85	0.025		
Medium	13.43	13.83			
Large	54.73	54.81			
Hospital location/teaching status	5.175	5.101	0.093		
Rural	5.08	5.69			
Urban nonteaching	20.57	20.41			
Urban teaching	74.36	73.9			
Hospital region			< 0.0001		
Northeast	16.62	16.39			
Midwest	24.53	26.67			
South	37.93	38.8			
West	20.91	18.14			
Old MI	14.3	11.2	< 0.0001		
Old PCI	1.1	0.9	0.056		
Old CABG	5.6	3.8	<0.0001		
Old pacemaker	1.4	1.6	0.145		
Old catheter ablation	2.7	1.6	<0.0001		
Hypertension	39.9	39.4	0.379		
Diabetes mellitus	11.2	12.2	0.013		
Peripheral vascular disease	3.6	5.1	<0.0001		
Smoking	25.9	21.2	< 0.0001		
Obesity	16.6	20.5	<0.0001		
Cocaine use	1.0	0.6	0.0005		
CHF	34.0	39.5	<0.0001		
	F 67	2 57	<0.0001		
י כ	13.07	3.37			
∠ >3	08.61	0.00			
 Dvslinidemia	63.3	61 9	0.025		
Prior catheter ablation	73	27	<0.0001		
Atrial fibrillation	19.5	21.5	0.0001		
Palliative care	4.5	5.9	< 0.0001		

Continued on the next page

TABLE 1 Continued			
	All Ventricula (N = 174,5		
	Male (n = 132,385, 75.8%)	Female (n = 42,180, 24.2%)	P Value
DNR	6.8	10.0	< 0.0001
ADPRG severity scale			
1	1.5	2.1	
2	34.4	26.9	
3	24.7	25.6	
4	39.4	45.4	

Values are percentage unless otherwise indicated. Estimated median household incomes are zip code-specific, updated annually and are classified into 4 quartiles indicating the poorest to wealthiest populations.

ADPRG = All-Payer Refined Diagnostic Related Group; AMI = acute myocardial infarction; CABG = coronary artery bypass grafting; CHF = congestive heart failure; DNR = do-not-resuscitate; MI = myocardial infarction; PCI = percutaneous coronary intervention.

categorical variables were presented in percentages (%) and compared using the chi-square test. Multivariable logistic and linear regression was used to determine the adjusted odds ratio (AOR)/adjusted risk ratio of the categorical and continuous outcomes of interest, respectively. Confounders were obtained from a literature review of prior similar studies and included age, race, income status, hospital region, hyperlipidemia, heart failure, history of MI, chronic kidney disease, diabetes mellitus, hypertension, anemia, and obesity.<sup>14-16</sup> A *P* value of <0.05 was considered statistically significant, and all tests were 2-sided with a 95% CI.

#### RESULTS

**BASELINE CHARACTERISTICS.** From 2016 to 2020, there were 3,191,049 admissions for AMI. After applying exclusions, our final cohort comprised of 1,542,910 patients with AMI. Of these, 174,565 (11.3%) patients had VAs. Among patients with AMI and VAs 132,385 (75.8%) were men. Baseline sociodemographic, hospital, and clinical characteristics of patients with AMI who had VAs are summarized by sex in (**Table 1**). Overall, women were older, more likely to have congestive heart failure, diabetes mellitus, obesity, atrial arrhythmia, and had higher index of illness severity by the All-Payer Refined Diagnostic Related Group severity of illness scale, and more likely to have do-not-resuscitate status.

**SEX DIFFERENCES IN IN-HOSPITAL OUTCOMES.** Compared with men, women were found to have a lower incidence of VAs following AMI (8.6% for women vs 12.6% for men; AOR: 0.67 [0.65-0.69]; P < 0.001). However, women had worse secondary outcomes with higher rates in hospital mortality (AOR: -1.31 [1.21-1.42] P < 0.0001), cardiogenic shock (AOR: 1.08 [1.01-1.15]; P = 0.022), and cardiac arrest (AOR: 1.11 [1.03-1.18]; P < 0.002) (Figure 2).

**PROCEDURAL OUTCOMES.** Women were also found to have lower rates of ICD insertion (AOR: 0.57 [0.47-0.68] P < 0.0001) and less likely to receive ablative therapies (AOR: 0.51 [0.27-0.98] P < 0.0001) following VAs, as compared to men (**Figure 2**). A subanalysis considering VT and VF patients separately found the disparity to apply to both types of VAs: (2.0% vs 2.9%), AOR 0.60 [0.49-0.74] P < 0.001 for VT and (1.7% vs 2.7%), AOR 0.56 [0.47-0.68] P < 0.001 for VF. Similarly, women were less likely to undergo catheter ablation during the index hospitalization as compared to men (AOR: 0.51; CI: 0.27-0.98; P < 0.001).

SUBANALYSIS OF OUTCOMES IN NSTEMI VS STEMI PATIENTS. Subgroup analysis on the sex differences in outcomes of STEMI patients revealed that women were less likely to experience VAs post-STEMI (AOR: 0.63 [0.60-0.66] P < 0.001), but they had worse inhospital mortality and were less likely to receive an ICD (Table 2). Similar findings were also observed in the NSTEMI subgroup of patients as shown in (Table 3).

## DISCUSSION

Our study set out to explore the sex differences in the incidence and in-hospital outcomes of VAs following an AMI using a nationally representative sample of patients admitted for AMI in the United States. The salient findings of this study are that while women were less likely to have VAs following AMI, they suffered worse in-hospital outcomes in terms of in-hospital cardiogenic shock, cardiac arrest, and in-hospital mortality, and were less likely to receive procedural care such as ICD implantation or ablation therapy (Figure 2, Central Illustration).

VAs are known complications of AMI. However, to the best of our knowledge, this study represents the largest study to date to examine sex differences in the incidence and in-hospital outcomes of VAs in patients admitted with AMI. Consistent with the findings of other studies, our work confirms observations that women are less likely to experience VAs after AMI.<sup>17,18</sup> While the reasons for sex differences in the risk of VAs post-AMI is not entirely understood, our study showed a higher rate of prior MI and ischemic heart disease (IHD) among men, which might result in a greater myocardial scar burden as a substrate for potential re-entrant circuits. Our findings are



supported by reports from the landmark MADIT-CRT (Multicenter Automatic Defibrillator Implantation with Cardiac Resynchronization Therapy Trial), which also reported higher rates of IHD in men. They also noted that the probability of ICD shocks due to VT/VF among women was half that observed in men (HR: 0.51; P = 0.003).<sup>19</sup> In addition to a greater burden of proarrhythmic substrate, other sex-dependent characteristics due to hormonal influences on ion channel expression and action potential may also help explain our findings.<sup>20-23</sup>

Although women had a lower likelihood of VAs following AMI, women had higher odds of in-hospital mortality, cardiogenic shock, and cardiac arrest among those with VAs. These findings can be partly explained by the general comorbid profile of women with AMI, as they were older, had higher rates of congestive heart failure, diabetes mellitus and worse Charlson comorbidity index scores. However, this excess risk was observed even after accounting for these differences. In a similar study by Shih et al,<sup>24</sup> as compared to men, women had higher mortality rates despite equivalent rates of VAs with primary drivers of these higher rates being older age, post-MI Heart Failure, and less frequent utilization of guideline-directed therapies. We noted persistence of these worse outcomes in women with VAs even after subdivision into AMI groups (STEMI and non-STEMI) (Tables 2 and 3).

In addition, women were less likely to receive ICDs or catheter ablation therapies during hospital admission. While ICD is recommended as a secondary prevention therapy for women as it is for men with ventricular tachyarrhythmias in the setting of AMI, there continue to be sex differences in the recruitment to ICD trials and implantation in women as previously demonstrated.<sup>18,25</sup> Whether the difference in implantation is related to suggestions of higher

TABLE 2Sex Differences in the Outcomes of Patients WithSTEMI and Ventricular Arrhythmias					
	Males <sup>a</sup>	Females	OR (95% CI)	P Value	
In-hospital mortality	12.5	17.9	1.24 (1.13-1.35)	< 0.001	
Cardiogenic shock	26.4	30.8	1.03 (0.95-1.11)	0.435	
Cardiac arrest	21.8	23.4	1.05 (0.97-1.13)	0.209	
ICD insertion	2.0	1.30	0.57 (0.43-0.72)	< 0.001	
Ablation	0.1	0.01	0.49 (0.18-1.33)	0.165	
<sup>a</sup> In males represent "reference variable".					

TABLE 3 Sex Differences in the Outcomes of Patients With NSTEMI and Ventricular Arrhythmias

	Males <sup>a</sup>	Females	OR (95% CI)	P Value	
In-hospital mortality	8.5	13.4	1.48 (1.27-1.74)	< 0.001	
Cardiogenic shock	14.7	17.7	1.15 (1.01-1.32)	0.04	
Cardiac arrest	13.3	17.3	1.28 (1.12-1.46)	< 0.001	
ICD insertion	4.9	3.1	0.61 (0.47-0.78)	< 0.001	
Ablation	0.5	0.3	0.57 (0.26-1.25)	0.163	
<sup>a</sup> ln males represent "reference variable"					



complications of ICD implantation in women or a greater benefit in men, as suggested in some trials. The available data indicate an underrepresentation of women in these studies, with no definite sex differences in mortality and an equivocal benefit of ICD therapy postimplantation.<sup>26-28</sup> Therefore, future efforts should be directed toward minimizing the sex disparities in ICD implantation in AMI patients with VAs.

This study should be interpreted in the context of some limitations. First, this is an observational and nonrandomized study, utilizing an administrative data set. Although we used a robust statistical model in the analysis, the effect of residual confounders cannot be fully excluded. Secondly, our clinical and procedural variables were defined by ICD-10-CM and PCS codes which are vulnerable to coding errors. Third, the data available to us lack the granularity necessary to determine the clinical severity on presentation, laboratory, and echocardiographic parameters. Similarly, lack of data on medications limited our assessment of sex disparities in treatment with guideline medical therapy. Finally, due to the lack of longitudinal data on individual patients, we cannot extrapolate our findings to the posthospitalization period.

# CONCLUSIONS

In this large nationwide analysis of patients with AMI, we found a lower incidence of complicating VAs in women as compared to men. While the lower prevalence of previous IHD might partially explain this, further studies are needed to understand the potential contributions of other factors, such as endocrine and genetic factors to explain this sex difference in VAs following AMI. Despite having less VAs, women were found to have worse in-hospital outcomes (mortality, cardiac arrest, and cardiogenic shock) in the setting of AMI and VAs but were less likely to receive ICD implantation or ablation therapy. These findings highlight the importance of assessing for sex differences in cardiovascular disease to improve the care of both women and men. Further investigation is needed to determine what treatments can improve the outcomes in women.

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#### PERSPECTIVES

**COMPETENCY IN MEDICAL KNOWLEDGE:** Women are less likely to develop VAs following an acute ischemic event. However, among persons who sustained VAs post-MI, women are more likely to have worse in-hospital outcomes and less likely to receive implantable cardiac defibrillator therapy.

**TRANSLATION OUTLOOK:** Additional real-world studies are needed to replicate the findings from this study. In addition, further research geared toward targeted interventions to address sex-specific risk factors associated with outcomes following MI and improve postarrhythmia outcomes among women are important next steps.

#### REFERENCES

**1.** Bhar-Amato J. Ventricular arrhythmia after acute myocardial infarction: 'the Perfect Storm'. 2017. Accessed August 26, 2023. https://www.aerjournal.com/articles/ventricular-arrhythmia-after-acute-myocardial-infarction-perfect-storm

 Piccini JP, Berger JS, Brown DL. Early sustained ventricular arrhythmias complicating acute myocardial infarction. *Am J Med.* 2008;121(9): 797-804. https://doi.org/10.1016/j.amjmed. 2008.04.024

**3.** Masuda M, Nakatani D, Hikoso S, et al. Clinical impact of ventricular tachycardia and/or fibrillation during the acute phase of acute myocardial infarction on in-hospital and 5-year mortality rates in the percutaneous coronary intervention era. *Circ J.* 2016;80(7):1539-1547. https://doi.org/10.1253/ circj.CJ-16-0183

**4.** Thomas DE, Jex N, Thornley AR. Ventricular arrhythmias in acute coronary syndromes—mechanisms and management. *Contin Cardiol Educ.* 2017;3(1):22-29. https://doi.org/10.1002/cce2.51

**5.** Hersi AS, Alhabib KF, AlFaleh HF, et al. Incidence of ventricular arrhythmia and associated patient outcomes in hospitalized acute coronary syndrome patients in Saudi Arabia: findings from the Registry of the Saudi Project for Assessment of Acute Coronary Syndrome (SPACE). *Ann Saudi Med.* 2012;32(4):372-377. https://doi.org/10. 5144/0256-4947.2012.372

**6.** Avezum Á, Piegas LS, Goldberg RJ, et al. Magnitude and prognosis associated with ventricular arrhythmias in patients hospitalized with acute coronary syndromes (from the GRACE Registry). Am J Cardiol. 2008;102(12):1577-1582. https://doi.org/10.1016/j.amjcard.2008.08.009

**7.** Weizman O, Marijon E, Narayanan K, et al. Incidence, characteristics, and outcomes of ventricular fibrillation complicating acute myocardial infarction in women admitted alive in the hospital. *J Am Heart Assoc.* 2022;11(17):e025959. https:// doi.org/10.1161/JAHA.122.025959

**8.** von Elm E, Altman DG, Egger M, Pocock SJ, Gøtzsche PC, Vandenbroucke JP. The Strengthening the Reporting of Observational Studies in Epidemiology (STROBE) statement: guidelines for reporting observational studies. *J Clin Epidemiol.* 2008;61(4):344-349. https://doi.org/10.1016/ j.jclinepi.2007.11.008

9. NIS database documentation. Accessed August 20, 2023. https://hcup-us.ahrq.gov/db/nation/ nis/nisdbdocumentation.jsp

**10.** ICD-10. American College of Cardiology. Accessed August 20, 2023. https://www.acc.org/ http%3a%2f%2fwww.acc.org%2ficd10

**11.** 2023 ICD-10-CM | CMS. Accessed August 20, 2023. https://www.cms.gov/medicare/icd-10/2 023-icd-10-cm

12. Markson FE, Akuna E, Lim CY, Khemani L, Amanullah A. The impact of COVID-19 on hospitalization outcomes of patients with acute myocardial infarction in the USA. *Am Heart J Plus.* 2023;32:100305. https://doi.org/10.1016/j.ahjo. 2023.100305

**13.** HCUP-US NIS overview. Accessed August 20, 2023. https://hcup-us.ahrq.gov/nisoverview.jsp

**14.** Saxena S, Goldenberg I, McNitt S, et al. Sex differences in the risk of first and recurrent ventricular tachyarrhythmias among patients receiving an implantable cardioverter-defibrillator for primary prevention. *JAMA Netw Open.* 2022;5(6):e2217153. https://doi.org/10.1001/jamanetworkopen.2022.17153

**15.** Tarantino N, Rocca DGD, Zou F, Lin A, Natale A, Biase LD. Prevalence, outcomes, and management of ventricular arrhythmias in COVID-19 patients. *Card Electrophysiol Clin*. 2022;14(1):11-20. https:// doi.org/10.1016/j.ccep.2021.10.002

**16.** Matetic A, Shamkhani W, Rashid M, et al. Trends of sex differences in clinical outcomes after myocardial infarction in the United States. *CJC Open*. 2021;3(12, Supplement):S19-S27. https:// doi.org/10.1016/j.cjco.2021.06.012

**17.** Buxton AE, Hafley GE, Lehmann MH, et al. Prediction of sustained ventricular tachycardia inducible by programmed stimulation in patients with coronary artery disease. *Circulation*. 1999;99(14):1843-1850. https://doi.org/10.1161/ 01.CIR.99.14.1843

18. Zaman S, Deshmukh T, Aslam A, Martin C, Kovoor P. Sex differences in electrophysiology, ventricular tachyarrhythmia, cardiac arrest and sudden cardiac death following acute myocardial infarction. *Heart Lung Circ.* 2020;29(7):1025-1031. https://doi.org/10.1016/j.hlc.2019.07.017

19. Tompkins CM, Kutyifa V, Arshad A, et al. Sex differences in device therapies for ventricular arrhythmias or death in the multicenter automatic defibrillator implantation trial with cardiac resynchronization therapy (MADIT-CRT) trial. J Cardiovasc Electrophysiol. 2015;26(8):862-871. https://doi.org/10.1111/jce.12701

**20.** Dogan M, Yiginer O, Uz O, et al. The effects of female sex hormones on ventricular premature beats and repolarization parameters in physiological menstrual cycle. *Pacing Clin Electrophysiol*. 2016;39(5):418-426. https://doi.org/10.1111/pace. 12821

**21.** Gillis AM. Atrial fibrillation and ventricular arrhythmias. *Circulation*. 2017;135(6):593-608. https:// doi.org/10.1161/CIRCULATIONAHA.116.025312

22. Linde C, Bongiorni MG, Birgersdotter-Green U, et al. Sex differences in cardiac arrhythmia: a consensus document of the European Heart Rhythm Association, endorsed by the Heart Rhythm Society and Asia Pacific Heart Rhythm Society. *EP Europace*. 2018;20(10):1565-1565ao. https://doi.org/10.1093/europace/euy067 **23.** Tadros R, Ton AT, Fiset C, Nattel S. Sex differences in cardiac electrophysiology and clinical arrhythmias: epidemiology, therapeutics, and mechanisms. *Can J Cardiol.* 2014;30(7):783-792. https://doi.org/10.1016/j.cjca.2014.03.032

24. Shih JY, Chen ZC, Chang HY, Liu YW, Ho CH, Chang WT. Risks of age and sex on clinical outcomes post myocardial infarction. *Int J Cardiol Heart Vasc*. 2019;23:100350. https://doi.org/10. 1016/j.ijcha.2019.100350

**25.** Barra S, Providência R, Boveda S, et al. Device complications with addition of defibrillation to cardiac resynchronisation therapy for primary prevention. *Heart*. 2018;104(18):1529-1535. https://doi.org/10.1136/heartjnl-2017-312546

**26.** Yoder M, Dils A, Chakrabarti A, et al. Gender and race-related disparities in the management of ventricular arrhythmias. *Trends Cardiovasc Med.* 2023. https://doi.org/10.1016/j.tcm.2023.10.001 27. Dêbski M, Ulman M, Z¥Bek A, Haberka K, Lelakowski J, Ma£ecka B. Gender differences in dual-chamber pacemaker implantation indications and long-term outcomes. Acta Cardiol. 2016;71(1):41-45. https://doi.org/10.1080/AC.71. 1.3132096

28. Elango K, Curtis AB. Cardiac implantable electrical devices in women. *Clin Cardiol*. 2018;41(2):232-238. https://doi.org/10.1002/clc. 22903

**KEY WORDS** acute myocardial infarction, sex or gender disparities, ventricular arrhythmias

**APPENDIX** For a supplemental table, please see the online version of this paper.