

Novel Method of Atrial Fibrillation Case Identification and Burden Estimation Using the MIMIC-III Electronic Health Data Set

Journal of Intensive Care Medicine
1-7
© The Author(s) 2019
Article reuse guidelines:
sagepub.com/journals-permissions
DOI: 10.1177/0885066619866172
journals.sagepub.com/home/jic


Eric Y. Ding, MS¹ , Daniella Albuquerque, BS², Michael Winter, MS³,
Sophia Binici, BS², Jaclyn Piche, RN², Syed Khairul Bashar, BS⁴, Ki Chon, PhD⁴,
Allan J. Walkey, MD, ScM⁵, and David D. McManus, MD, ScM^{1,2}

Abstract

Background: Atrial fibrillation (AF) portends poor prognoses in intensive care unit patients with sepsis. However, AF research is challenging: Previous studies demonstrate that *International Classification of Disease (ICD)* codes may underestimate the incidence of AF, but chart review is expensive and often not feasible. We aim to examine the accuracy of nurse-charted AF and its temporal precision in critical care patients with sepsis. **Methods:** Patients with sepsis with continuous electrocardiogram (ECG) waveforms were identified from the Medical Information Mart for Intensive Care (MIMIC-III) database, a de-identified, single-center intensive care unit electronic health record (EHR) source. We selected a random sample of ECGs of 6 to 50 hours' duration for manual review. Nurse-charted AF occurrence and onset time and ICD-9-coded AF were compared to gold-standard ECG adjudication by a board-certified cardiac electrophysiologist blinded to AF status. Descriptive statistics were calculated for all variables in patients diagnosed with AF by nurse charting, ICD-9 code, or both. **Results:** From 142 ECG waveforms (58 AF and 84 sinus rhythm), nurse charting identified AF events with 93% sensitivity (95% confidence interval [CI]: 87%-100%) and 87% specificity (95% CI: 80%-94%) compared to the gold standard manual ECG review. Furthermore, nurse-charted AF onset time was within 1 hour of expert reader onset time for 85% of the reviewed tracings. The ICD-9 codes were 97% sensitive (95% CI: 88-100%) and 82% specific (95% CI: 74-90%) for incident AF during admission but unable to identify AF time of onset. **Conclusion:** Nurse documentation of AF in EHR is accurate and has high precision for determining AF onset to within 1 hour. Our study suggests that nurse-charted AF in the EHR represents a potentially novel method for AF case identification, timing, and burden estimation.

Keywords

atrial fibrillation, nurse documentation, sepsis, accuracy, case identification

Introduction

Atrial fibrillation (AF) is the most common cardiac arrhythmia among critically ill patients,¹ with an especially high incidence among patients with sepsis.² New-onset AF during sepsis is associated with higher in-hospital mortality and important postdischarge outcomes,³⁻⁶ including incident heart failure, stroke, and death.⁷ Furthermore, AF that occurs in the context of suspected infection may be an indicator of acute cardiac dysfunction consistent with sepsis.⁸ It is therefore important to accurately identify cases of AF in health records to facilitate research regarding triggers, treatments, and associated outcomes.

Case identification of AF often relies on administrative databases^{9,10} that use *International Classification of Disease (ICD-9 or ICD-10)* systems primarily used for billing purposes. Although administrative data are widely available, previous studies have shown that the accuracy of using the ICD-9 and ICD-10 codes of AF varies widely across populations and

suffers from poor sensitivity.^{9,11-13} Another key limitation of ICD codes is the inability to determine AF onset time, an important factor in studies of time-varying AF triggers and

¹ Department of Population and Quantitative Health Sciences, University of Massachusetts Medical School, Worcester, MA, USA

² Division of Cardiology, Department of Medicine, University of Massachusetts Medical School, MA, USA

³ Biostatistics and Epidemiology Data Analytics Center, Boston University School of Public Health, MA, USA

⁴ Department of Biomedical Engineering, University of Connecticut, CT, USA

⁵ Pulmonary Center, Boston University School of Medicine, MA, USA

Received March 18, 2019. Received revised July 03, 2019. Accepted July 08, 2019.

Corresponding Author:

Eric Y. Ding, MS, Department of Population and Quantitative Health Sciences, University of Massachusetts Medical School, 368 Plantation St, Worcester, MA 01655, USA.

Email: eric.ding@umassmed.edu

treatment outcomes. Current AF guidelines recommend clinician interpretation¹⁴ of a 12-lead electrocardiogram (ECG)¹⁵ for the determination of AF onset, but large-scale ECG clinical databases are rare and large-scale manual review of ECG data is costly, time-consuming, and generally infeasible.¹⁶ The widespread integration of electronic health records (EHRs) into hospital systems provides the opportunity to develop more efficient and accurate tools that facilitate the detection of AF.

Clinical information technologies in intensive care units (ICUs) have increased the speed of nurse charting¹⁷ and enabled the timely documentation of a number of clinical parameters by nursing staff, including cardiac rhythm status.¹⁸ However, AF events adjudicated by routine nurse charting have not previously been validated against gold standard manual ECG interpretation. We sought to determine the accuracy of nurse-charted AF compared with gold standard manual ECG interpretation by an electrophysiologist and *ICD-9* and evaluate codes in an electronic health database of critically ill patients with sepsis.

Methods

Study Population

The study cohort was derived from the Medical Information Mart for Intensive Care (MIMIC-III) database.¹⁸ The MIMIC-III is a large, single-center database of critical care admissions to a tertiary care hospital in Boston, Massachusetts. The database consists of patients admitted to surgical and medical ICUs from 2001 to 2012 and contains hourly data on vital signs, charted events, laboratory values, clinical notes, and waveform data from continuous ECG telemetry recordings. The use of open-source, deidentified MIMIC-III data for the current study has been deemed not human subjects research by the institutional review boards at both University of Massachusetts Medical School and Boston University, with data use approved by MIMIC-III database administration staff.

We included adult patients over 18 years of age with available continuous ECG waveforms who were diagnosed with sepsis according to the Angus definition combining acute infection and organ dysfunction *ICD-9* codes or *ICD-9* codes specifically used for severe sepsis (995.92) or septic shock (785.52),^{19,20} Patients were excluded from study analysis if they had less than 6 hours of telemetry data (to ensure sufficient data for analysis) or more than 50 hours of telemetry data (to enhance the feasibility of manual ECG analysis) or if the rhythm status was not clearly discernable (poor data quality). Patients with ECG waveforms showing atrial flutter, electronic ventricular pacemaker, or significant artifact obscuring ECG interpretation were also excluded. Finally, ECG waveforms were excluded from accuracy analysis if the nurse-charted AF onset time did not occur within 1 hour of available telemetry data. The duration of available telemetry data does not necessarily reflect the length of the patient's ICU stay or the duration of their sepsis. All exclusionary criteria were implemented to facilitate and ensure the accuracy of the review. If a patient had multiple admissions with sepsis, only the first admission was analyzed.

Clinical Variables

We abstracted vital signs, laboratory data, inpatient medication, diagnosis, and procedure information from MIMIC-III. We used *ICD-9* codes to identify comorbid medical conditions, including hypertension (401.X), diabetes (250.X), congestive heart failure (CHF) (428.X), and ischemic stroke (433-434, 436), and procedures including electrical cardioversion (99.61), cardiac catheterization (37.2X), intubation (96.01-96.06), pulmonary artery catheterization (89.64), and coronary artery bypass graft procedures (36.1X) were also determined through their *ICD-9* procedure codes. Vital signs were collected from the first 24 hours of ICU stay, including mean arterial pressure, systolic and diastolic blood pressure, and heart rate. Laboratory values included creatinine, troponin, brain natriuretic peptide, and hemoglobin on admission. We collected medication dosing information for oral anticoagulants (including vitamin K antagonists and direct oral anticoagulants), digoxin, β -blockers, calcium channel blockers, antiarrhythmics, and vasopressors.

Atrial Fibrillation Status Based on Electronic Hospital and Administrative Records

Atrial fibrillation status was determined through 2 methods, based on data from the nursing records in the EHR. Nurse-charted rhythm status is encoded in the MIMIC-III data set and is recorded hourly, consistent with standard nursing practice and EHR documentation. We defined "nurse-charted AF" as any documentation by the nurse of "AF," "atrial fibrillation," "AFib," or "atrial fib." Only the first instance of AF was identified for each patient. The *ICD* codes in MIMIC, as with many EHR data sets, were coded at the time of hospital discharge and thus included all cases present on admission or occurring during hospitalization. As has previously been conducted, the *ICD-9* code of 427.31 was used to define AF based on claims data.

Telemetry Data Analysis and Gold Standard AF Ascertainment

Approximately 30% of all telemetry ECG waveforms were randomly selected for manual screening by 3 trained study personnel (D.A., S.B., and E.Y.D.). Rhythm status was ascertained by trained study staff at baseline and every 60 minutes thereafter for the entirety of the telemetric ECG recording. If over 30 seconds of uninterrupted AF was detected at any time in this interval scanning, the exact onset and offset times of the AF episode were then ascertained. Similar to the identification of nurse-charted AF, only the first instance of AF was reviewed. A board-certified cardiac electrophysiologist (D.D.M.) who was blinded to AF status was then presented with approximately equal numbers of AF and non-AF ECGs as determined by research staff. An unequal distribution was chosen to maintain the integrity of the expert reviewer's blinding to rhythm status throughout all

Table 1. Patient Characteristics of ICU Patients With Sepsis.

Demographics	No AF (n = 1704)	Any AF (n = 1270)	P Value
Age, mean (SD)	60.7 (16.0)	71.4 (12.0)	<.001
Male sex (%)	53.1%	56.7%	.052
Race			<.001
White	65.6%	73.9%	
Black	11.3%	7.9%	
Latino	3.8%	2.0%	
Asian	2.9%	2.8%	
Other	16.4%	13.5%	
Discharge location			<.001
Home	32.9%	22.9%	
SNF	17.7%	18.3%	
Transfer	3.5%	3.1%	
Long term	12.3%	12.0%	
Hospice	1.3%	1.3%	
Died	16.6%	24.3%	
Other	15.8%	18.3%	
In-hospital factors, diagnoses, and procedures			
Length of stay, days, mean (SD)	15.1 (15.1)	16.2 (14.1)	.053
Hypertension	43.1%	47.9%	.010
CHF	29.3%	60.7%	<.001
Diabetes	31.6%	38.6%	<.001
Ischemic stroke	4.9%	6.8%	.032
Electrical cardioversion	0.1%	3.5%	<.001
CABG	3.8%	10.6%	<.001
Intubation	39.5%	45.2%	.002
Echocardiogram	8.7%	16.5%	<.001
Pulmonary artery catheterization	4.0%	7.2%	<.001
Cardiac catheterization	11.7%	20.7%	<.001
Medications			
Warfarin	12.9%	43.7%	<.001
Other anticoagulants	89.3%	90.3%	.38
Digoxin	2.2%	24.4%	<.001
β-blocker	63.6%	85.7%	<.001
Calcium channel blocker	15.6%	40.3%	<.001
Antiarrhythmic	4.0%	43.7%	<.001
Vasopressor	49.9%	69.2%	<.001
Physiologic and laboratory values on admission			
Mean arterial pressure, mm Hg, mean (SD)	81.4 (43.6)	79.1 (43.0)	.33
Systolic blood pressure, mm Hg, mean (SD)	124.6 (29.8)	120.1 (27.5)	.002
Diastolic blood pressure, mm Hg, mean (SD)	60.9 (18.5)	57.2 (16.8)	<.001
Heart rate, bpm, mean (SD)	92.2 (20.4)	89.3 (20.7)	<.001
Creatinine, ng/dL, mean (SD)	1.8 (1.9)	1.9 (1.7)	.044
Troponin, ng/mL, mean (SD)	0.4 (1.7)	0.4 (1.6)	.84
NT-proBNP, pg/mL, mean (SD)	6588 (9796)	10 715.0 (11 302.0)	<.001
Hemoglobin, g/dL, mean (SD)	11.5 (2.3)	11.3 (2.2)	.019

Abbreviations: AF, atrial fibrillation; CABG, coronary artery bypass surgery; CHF, congestive heart failure; ICU, intensive care unit; NT-proBNP, N-terminal prohormone of brain natriuretic peptide; SD, standard deviation; SNF, skilled nursing facility.

adjudications. The physician then manually adjudicated all potential ECG waveforms to confirm the rhythm status and AF onset times.

Statistical Analysis

Descriptive statistics were calculated for all abstracted clinical, physiological, and laboratory parameters. These data were categorized according to AF status. We evaluated differences in characteristics using analysis of variance and chi-square

across 4 categories of AF status: (1) patients without AF, (2) those with AF identified by both *ICD-9* codes and nurse-charted AF, (3) those with only *ICD-9* identified AF, and (4) those with only nurse-identified AF. To directly compare 2 samples, the Tukey test and chi-squared were used.

Sensitivity and specificity were calculated for “nurse-charted AF” and *ICD*-coded AF was calculated via comparison with gold standard rhythm assessment by a blinded board-certified cardiac electrophysiologist (gold standard). Exact Clopper–Pearson confidence intervals were calculated based

Table 2. Characteristics of MIMIC-III Patients by Atrial Fibrillation Status.

	Nurse-Charted AF Plus ICD-coded AF (n = 835)	Only Nurse-Charted AF (n = 227)
Age, mean (SD)	72.6 (11.3)	69.1 (13.0)
Male sex, n (%)	55.8%	58.1%
Race		
White	73.8%	73.1%
Black	7.9%	7.9%
Latino	1.8%	3.1%
Asian	3.0%	3.1%
Other	13.5%	12.8%
Discharge location		
Home	21.6%	34.4%
SNF	20.6%	17.2%
Transfer	2.8%	2.2%
Long-term facility	12.7%	7.0%
Hospice	1.3%	1.3%
Died	22.5%	20.7%
Other	18.6%	17.2%
Length of stay in days, mean (SD)	16.1 (13.4)	13.0 (12.0)
Hypertension	51.0%	46.3%
CHF	63.4%	59.0%
Diabetes	38.3%	45.8%
Ischemic stroke	6.3%	7.5%
Electrical cardioversion	4.9%	0.4%
CABG	11.5%	7.5%
Intubation	42.3%	43.3%
Echocardiogram	16.6%	15.9%
Pulmonary artery catheterization	7.3%	6.2%
Cardiac catheterization	20.1%	22.0%
Warfarin	49.9%	42.7%
Digoxin	30.4%	14.1%
β-blocker	88.6%	84.6%
Calcium channel blocker	47.8%	22.5%
Antiarrhythmic	49.5%	32.2%
Other anticoagulants	92.6%	86.3%
Vasopressors	71.6%	56.4%
Heart rate, bpm, mean (SD)	91.0 (20.8)	82.0 (17.7)
Mean arterial pressure, mm Hg, mean (SD)	80.2 (47.5)	80.9 (30.7)
Systolic blood pressure, mm Hg, mean (SD)	118.7 (25.6)	127.1 (29.3)
Diastolic blood pressure, mm Hg, mean (SD)	57.4 (16.2)	57.6 (16.3)
Creatinine, ng/dL, mean (SD)	1.9 (1.7)	2.0 (1.7)
Troponin, ng/mL, mean (SD)	0.4 (1.8)	0.4 (1.1)
Hemoglobin, g/dL, mean (SD)	11.3 (2.3)	11.4 (2.1)
NT-proBNP, pg/mL, mean (SD)	11153.2 (11270.1)	9351.7 (11592.8)

Abbreviations: AF, atrial fibrillation; CABG, coronary artery bypass surgery; CHF, congestive heart failure; ICD, *International Classification of Disease*; MIMIC-III, Medical Information Mart for Intensive Care; NT-proBNP, N-terminal prohormone of brain natriuretic peptide; SD, standard deviation; SNF, skilled nursing facility.

on the binomial distribution. The proportion of “charted AF” onset times that were within 60 minutes of manual adjudication of AF onset was also calculated in order to evaluate the accuracy of timing in nurse-charted AF.

Results

The characteristics of the 2974 patients with sepsis and available ECG telemetry data included in our analyses are shown in Table 1. The average age of the study population was 65.2 ± 15.4 years, 54.6% were male, and a majority were Caucasian. The cohort had a median length of hospital stay

of 12 (interquartile range: 7-20) and more than 40% of patients required intubation and mechanical ventilation. Patients had a high burden of cardiovascular comorbidity, 45% had hypertension, 34% diabetes, and 43% CHF. The clinical profiles of patients with AF compared to those without AF are presented in Table 1.

Patient characteristics differed according to the presence of AF and the manner in which AF was identified. Patients with ICD-9-coded AF as well as nurse charting had higher rates of electrical cardioversion, longer length of stay, lower rates of discharge home, and higher rates of medication use than patients with only nurse-charted AF (Table 2).

Table 3. Accuracy of Nurse-Documented AF Compared to Manual Adjudication.^a

		Manual adjudication		
		AF	No AF	Total
Nurse documentation	AF	54	11	65
	No AF	4	73	77
	Total	58	84	142

Abbreviations: AF, atrial fibrillation; CI, confidence interval; PPV, Positive predictive value.

^aSensitivity = 93% (95% CI: 87%-100%), specificity = 87% (95% CI: 80%-94%), PPV = 83%.

Table 4. Accuracy of ICD-9 AF Compared to Manual Adjudication.^a

		Manual adjudication		
		AF	No AF	Total
ICD-9 defined	AF	56	15	71
	No AF	2	69	71
	Total	58	84	142

Abbreviations: AF, atrial fibrillation; CI, confidence interval; ICD, *International Classification of Disease*; PPV, Positive predictive value.

^aSensitivity = 97% (95% CI: 88%-100%), specificity = 82% (95% CI: 74%-90%), PPV = 79%.

Among all ECG waveforms reviewed, 142 (58 AF and 84 sinus rhythm) met inclusion and exclusion criteria (see Supplemental Figure for flow diagram detailing waveform selection). Nurse-charted AF had 93% sensitivity (95% confidence interval [CI]: 87%-100%) and 87% specificity (95% CI: 80%-94%) for the detection of AF (Table 3). Forty-six (85%) of nurse-documented AF episodes had onset times within 60 minutes of manually adjudicated AF based on primary analysis of continuous ECG data. The ICD-9 showed 97% sensitivity (95% CI: 88%-100%) and 82% specificity (95% CI: 74%-90%) (Table 4).

Discussion

Methods of identifying the presence of AF from administrative claims data may be limited by low sensitivity and lack of information regarding the timing of AF. We explored the performance of novel methods of AF identification using hourly nurse charting of events in EHR data. We evaluated the accuracy of nurse-charted AF, which were 93% sensitive and 87% specific, as well as ICD-9-identified AF, which were 97% sensitive and 82% specific. In addition, the timing associated with nurse documentation allows for the estimation of AF onset, which matched manual review in 85% of cases. Our findings suggest that nurse-charted AF was accurate for identifying AF cases, performing as well as administrative claims data, but with the additional benefit of reasonable temporal precision unavailable with administrative data. Our study also suggests that nurse chart reporting of AF may be a useful tool in research seeking to evaluate triggers for AF and near-term

outcomes associated with new-onset AF during critical illness.^{3,21}

Although the accuracy of AF detected by nurse events and ICD-9 codes was similar, characteristics of patients differed based upon the methods used to identify AF. In addition, patients with AF identified via both nurse charting and ICD-9 codes had generally worse health outcomes as compared to patients identified through only nurse charting (Table 2). Because the method of AF detection yielded cohorts of patients with different characteristics and outcomes, understanding the methods of AF identification in epidemiological research is critical. Mechanisms driving differences in outcomes associated with these methods of detection would require further study.

Previous work detailing the duration and frequency of AF episodes required manual ECG waveform interpretation,^{22,23} a labor-intensive process that is not conducive to research involving large electronic data sets. Nurse-charted AF performs adequately for the accurate determination of AF onset timing during a patient's stay in the ICU, thus circumventing this limitation. Future research is needed to establish if nurse-abstracted onset times can accurately enable estimation of the length of AF episodes. In addition, developments in machine learning algorithms and deep neural networks in recent years also provide an additional avenue for more granular estimation of AF burden. Further investigations comparing these technologies with nurse documentation may provide valuable insight into novel and effective strategies for AF identification in the EHR.

Our study has numerous strengths. To the best of our knowledge, only one large study (n = 1782) has used hourly ICU nurse charting for AF case identification,⁵ and no study to date has manually validated its accuracy in an EHR database. Additionally, unlike many other validation studies that use chart review as a comparison group, our study's gold standard is manual ECG adjudication by a board-certified cardiac electrophysiologist, which strengthens the validity of our results. We also examined AF onset times, which offers a more complete picture of the accuracy of nurse charting than just AF status alone. The temporal precision of AF episodes is critical for studies aiming to assess acute AF triggers and determine the length of the latency period between trigger and AF onset. This timing is also crucial in comparative effectiveness studies for the treatment of acute AF, as the precise timing of AF episodes is necessary to accurately determine the efficacy and time course of its treatments.

The study also has several limitations. The available ECG waveform data only reflect the patients' ICU stay, and thus, any AF that may have occurred outside of the ICU, and subsequently prompting ICD-9 designation, would be unable to be captured. Our study has potentially limited generalizability to other EHR databases. The data set we used originates from a large tertiary care center with significant resources invested into its ICU clinical information systems. However, with rapid advancements in EHRs and better integration into hospital systems, we expect similar quality data to be more widely

available across different health systems in the future. In addition, the collected data have undergone extensive cleaning and processing, which enables its usability. Curation of the ECG waveforms to focus on interpretable tracings limits the scope of our conclusion to patients whose ECG waveform is of high quality and thus may not be fully representative of real-world settings. By restricting our data set to patients with AF and excluding those with other atrial arrhythmias, we may have overestimated the accuracy of nurse-charted AF. Lastly, some patients in the database with nurse-charted AF without *ICD* diagnoses could also potentially be misclassification of similar electrocardiographic waveforms such as atrial flutter or atrial tachycardia because not all waveforms in the database were formally adjudicated.

We have identified and validated a novel and accurate method for AF case identification, timing, and burden estimation from administrative databases. When available, nurse-charted AF may be a useful tool to augment *ICD-9* codes for identifying, classifying, and timing AF episodes.

Declaration of Conflicting Interests

The author(s) declared no potential conflicts of interest with respect to the research, authorship, and/or publication of this article.

Funding

The author(s) disclosed receipt of the following financial support for the research, authorship, and/or publication of this article: Eric Y. Ding is funded through NIH Grants 1T32GM107000-01 and 5T32HL120823-05. David D. McManus's time was supported by 1R15HL121761, R01HL126911, R01HL137734, R01HL137794, and R01HL136660 from the National Heart, Lung and Blood Institute. Allan J Walkey was supported by R01HL136660, R01HL139751, and K01HL116768 from the National Heart, Lung and Blood Institute. This research was supported through the NIH Grant R01HL136660. Dr David D. McManus receives research support from Grants U54HL143541, R01HL126911, R01HL137734, R01HL137794, R01HL135219 from the National Heart, Lung and Blood Institute and National Center for Complementary and Integrative Health (National Institute of Health), and Grant NSF-12-512 from the National Science Foundation. Dr David D. McManus has received research support from Bristol Myers Squibb, Care Evolution, Samsung, Apple Computer, Pfizer, Biotronik, Boehringer Ingelheim, Philips Research Institute; has consulted for Bristol Myers Squibb, Pfizer, Philips, Samsung Electronics, and FlexCon.

ORCID iD

Eric Y. Ding, MS  <https://orcid.org/0000-0002-6904-8534>

Supplemental Material

Supplemental material for this article is available online.

References

1. Kanji S, Williamson DR, Yaghchi BM, Albert M, McIntyre L; Canadian Critical Care Trials Group. Epidemiology and management of atrial fibrillation in medical and noncardiac surgical adult intensive care unit patients. *J Crit Care*. 2012;27(3):326.e1-e8. doi:10.1016/j.jcrc.2011.10.011.
2. Walkey AJ, Greiner MA, Heckbert SR, et al. Atrial fibrillation among Medicare beneficiaries hospitalized with sepsis: incidence and risk factors. *Am Heart J*. 2013;165(6):949-955.e3. doi:10.1016/j.ahj.2013.03.020.
3. Walkey AJ, Wiener RS, Ghobrial JM, Curtis LH, Benjamin EJ. Incident stroke and mortality associated with new-onset atrial fibrillation in patients hospitalized with severe sepsis. *JAMA J Am Med Assoc*. 2011;306(20):2248-2254. doi:10.1001/jama.2011.1615.
4. Gandhi S, Litt D, Narula N. New-onset atrial fibrillation in sepsis is associated with increased morbidity and mortality. *Neth Heart J*. 2015;23(2):82-88. doi:10.1007/s12471-014-0641-x.
5. Klein Klouwenberg PMC, Frencken JF, Kuipers S, et al. Incidence, predictors, and outcomes of new-onset atrial fibrillation in critically ill patients with sepsis: a cohort study. *Am J Respir Crit Care Med*. 2017;195(2):205-211. doi:10.1164/rccm.201603-0618OC.
6. Kuipers S, Klouwenberg PMK, Cremer OL. Incidence, risk factors and outcomes of new-onset atrial fibrillation in patients with sepsis: a systematic review. *Crit Care*. 2014;18(6):688. doi:10.1186/s13054-014-0688-5.
7. Walkey AJ, Hammill BG, Curtis LH, Benjamin EJ. Long-term outcomes following development of new-onset atrial fibrillation during sepsis. *Chest*. 2014;146(5):1187-1195. doi:10.1378/chest.14-0003
8. Bosch NA, Cimini J, Walkey AJ. Atrial Fibrillation in the ICU. *Chest*. 2018;154(6):1424-1434. doi:10.1016/j.chest.2018.03.040.
9. Jensen PN, Johnson K, Floyd J, Heckbert SR, Carnahan R, Dublin S. A systematic review of validated methods for identifying atrial fibrillation using administrative data. *Pharmacoepidemiol Drug Saf*. 2012;21(suppl 1):141-147. doi:10.1002/pds.2317.
10. Tu K, Nieuwlaat R, Cheng SY, et al. Identifying patients with atrial fibrillation in administrative data. *Can J Cardiol*. 2016;32(12):1561-1565. doi:10.1016/j.cjca.2016.06.006.
11. Alonso A, Agarwal SK, Soliman EZ, et al. Incidence of atrial fibrillation in whites and African-Americans: the Atherosclerosis Risk in Communities (ARIC) study. *Am Heart J*. 2009;158(1):111-117. doi:10.1016/j.ahj.2009.05.010.
12. Thigpen JL, Dillon C, Forster KB, et al. Validity of *International Classification of Disease* codes to identify ischemic stroke and intracranial hemorrhage among individuals with associated diagnosis of atrial fibrillation. *Circ Cardiovasc Qual Outcomes*. 2015;8(1):8-14. doi:10.1161/CIRCOUTCOMES.113.000371.
13. Horsky J, Drucker EA, Ramelson HZ. Accuracy and completeness of clinical coding using *ICD-10* for ambulatory visits. *AMIA Annu Symp Proc*. 2018;2017:912-920.
14. Kligfield P, Gettes LS, Bailey JJ, et al. Recommendations for the standardization and interpretation of the electrocardiogram: part I: the electrocardiogram and its technology: a scientific statement from the American Heart Association Electrocardiography and Arrhythmias Committee, Council on Clinical Cardiology; the American College of Cardiology Foundation; and the Heart Rhythm Society endorsed by the International Society for Computerized Electrocardiology. *Circulation*. 2007;115(10):1306-1324. doi:10.1161/CIRCULATIONAHA.106.180200.

15. January CT, Wann LS, Alpert JS, et al. 2014 AHA/ACC/HRS Guideline for the management of patients with atrial fibrillation. *J Am Coll Cardiol*. 2014;64(21):e1. doi:10.1016/j.jacc.2014.03.022.
16. Aronsson M, Svennberg E, Rosenqvist M, et al. Cost-effectiveness of mass screening for untreated atrial fibrillation using intermittent ECG recording. *EP Eur*. 2015;17(7):1023-1029. doi:10.1093/europace/euv083.
17. Saarinen K, Aho M. Does the implementation of a clinical information system decrease the time intensive care nurses spend on documentation of care? *Acta Anaesthesiol Scand*. 2005;49(1):62-65. doi:10.1111/j.1399-6576.2005.00546.x.
18. Johnson AEW, Pollard TJ, Shen L, et al. MIMIC-III, a freely accessible critical care database. *Sci Data*. 2016;3:160035. doi:10.1038/sdata.2016.35.
19. Rhee C, Dantes R, Epstein L, et al. Incidence and trends of sepsis in US Hospitals using clinical vs claims data, 2009-2014. *JAMA*. 2017;318(13):1241-1249. doi:10.1001/jama.2017.13836.
20. Iwashyna TJ, Odden A, Rohde J, et al. Identifying patients with severe sepsis using administrative claims: patient-level validation of the angus implementation of the international consensus conference definition of severe sepsis. *Med Care*. 2014;52(6):e39-43. doi:10.1097/MLR.0b013e318268ac86.
21. Moskowitz A, Chen KP, Cooper AZ, Chahin A, Ghassemi MM, Celi LA. Management of atrial fibrillation with rapid ventricular response in the intensive care unit: a secondary analysis of electronic health record data. *Shock*. 2017;48(4):436-440. doi:10.1097/SHK.0000000000000869.
22. Patti G, Chello M, Candura D, et al. Randomized trial of atorvastatin for reduction of postoperative atrial fibrillation in patients undergoing cardiac surgery: results of the ARMYDA-3 (Atorvastatin for Reduction of MYocardial dysrhythmia After cardiac surgery) study. *Circulation*. 2006;114(14):1455-1461. doi:10.1161/CIRCULATIONAHA.106.621763.
23. Kottkamp H, Tanner H, Kobza R, et al. Time courses and quantitative analysis of atrial fibrillation episode number and duration after circular plus linear left atrial lesions: trigger elimination or substrate modification: early or delayed cure? *J Am Coll Cardiol*. 2004;44(4):869-877. doi:10.1016/j.jacc.2004.04.049.