

# Impact of Different Electronic Cohort Definitions to Identify Patients With Atrial Fibrillation From the Electronic Medical Record

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**Background**—Electronic medical records (EMRs) allow identification of disease-specific patient populations, but varying electronic cohort definitions could result in different populations. We compared the characteristics of an electronic medical record–derived atrial fibrillation (AF) patient population using 5 different electronic cohort definitions.

*Methods and Results*—Adult patients with at least 1 AF billing code from January 1, 2010, to December 31, 2017, were included. Based on different electronic cohort definitions, we trained 5 different logistic regression models using a labeled training data set (n=786). Each model yielded a predicted probability; patients were classified as having AF if the probability was higher than a specified cut point. Test characteristics were calculated for each model. These models were then applied to the full cohort and resulting characteristics were compared. In the training set, the comprehensive model (including demographics, billing codes, and natural language processing results) performed best, with an area under the curve of 0.89, sensitivity of 0.90, and specificity of 0.87. Among a candidate population (n=22 000), the proportion of patients identified as having AF varied from 61% in the model using diagnosis or procedure *International Classification of Diseases (ICD)* billing codes to 83% in the model using natural language processing of clinical notes. Among identified AF patients, the proportion of patients with a CHA<sub>2</sub>DS<sub>2</sub>-VASc score  $\geq$ 2 varied from 69% to 85%; oral anticoagulant treatment rates varied from 50% to 66% depending on the model.

*Conclusions*—Different electronic cohort definitions result in substantially different AF study samples. This difference threatens the quality and reproducibility of electronic medical record—based research and quality initiatives. (*J Am Heart Assoc.* 2020;9: e014527. DOI: 10.1161/JAHA.119.014527.)

Key Words: atrial fibrillation • electronic health records • health services research • informatics • quality of care

E lectronic medical records (EMR) are increasingly prevalent, resulting in an explosion of electronic health data available for research and quality initiatives. These data allow healthcare systems to capture large patient populations in order to study diagnoses, treatments, and outcomes.

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Accompanying Tables S1 through S3 are available at https://www.ahajourna ls.org/doi/suppl/10.1161/JAHA.119.014527

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© 2020 The Authors. Published on behalf of the American Heart Association, Inc., by Wiley. This is an open access article under the terms of the Creative Commons Attribution-NonCommercial License, which permits use, distribution and reproduction in any medium, provided the original work is properly cited and is not used for commercial purposes. Specifically, atrial fibrillation (AF) is a common condition, and large patient cohort studies may allow health systems and researchers to monitor quality and outcomes. Using the EMR, a health system could monitor the number of AF patients treated with an oral anticoagulant (OAC) for quality improvement efforts, such as increasing appropriate treatment rates among eligible patients. However, no common method exists to identify patients for inclusion in EMR-based initiatives. Different approaches could result in different patient cohorts with respect to characteristics and apparent outcomes, and this would limit the potential of EMR-based initiatives.

Prior studies used varying methods to identify AF patients.<sup>1–</sup> <sup>4</sup> Medicare studies, for example, include patients who have at least 1 inpatient or 2 outpatient *International Classification of Diseases (ICD)* codes for AF.<sup>2</sup> Reports from Kaiser Permanente also include ECG results in the patient-selection process.<sup>4</sup> Other studies have included EMR data such as ablation codes or antiarrhythmic medication treatment, but these studies report only positive predictive value and include outcomes (eg, anticoagulation use) in the prediction model.<sup>3</sup> Although data collected for billing purposes, including *ICD* codes, follow a

# **Clinical Perspective**

#### What Is New?

- We evaluated 5 different electronic definitions for identifying atrial fibrillation patients from the medical record.
- The characteristics and outcomes of the population differ substantially between different definitions.
- Of 22 000 possible patients, the number of included atrial fibrillation patients varied by up 6690 patients, and the apparent oral anticoagulant treatment rate in patients with CHA<sub>2</sub>DS<sub>2</sub>-VASc  $\geq$ 2 varied from half (49.6%) to two-thirds (66.3%) of the population, depending on the electronic cohort definition.

#### What Are the Clinical Implications?

- Quality improvement, learning healthcare systems, and realworld evidence require electronic cohort definitions to identify disease-specific patient populations.
- Different definitions will result in different populations and different apparent outcome rates.
- Validated, consistent electronic definitions are needed to ensure reproducibility and accuracy of studies that rely on electronic medical records.

controlled vocabulary, this approach may be inaccurate<sup>5,6</sup> and does not exploit other types of data, such as demographic information and non-AF diagnoses.

As an alternative to structured data, clinical notes are an untapped resource for detailed clinical information. Notes often include narrative references to patient conditions, such as "Patient was diagnosed with afib last year." Text mining with natural language processing (NLP) leverages the unstructured narrative from routine care and is another option for identifying patient cohorts. An advantage of NLP is that the clinical narrative may be less prone to some types of variation seen with billing codes,<sup>6,7</sup> which could support more precise patient selection and portability between institutions. The overall goal of this study was (1) to develop and train various models using different electronic-cohort definitions to identify AF patients from the EMR, incorporating structured and unstructured data; (2) to compare the resulting patient samples and characteristics from each model; and (3) to compare apparent OAC treatment rates in each sample.

# **Methods**

We developed and compared the performance of 5 cohort definitions to identify AF patients from the EMR:

- 1. Outpatient and inpatient *ICD* AF diagnosis billing codes (Medicare methodology)<sup>2</sup>:  $\geq$ 1 inpatient billing code or  $\geq$ 2 outpatient billing codes within 365 days
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- Outpatient AF diagnosis billing codes and ECG (Kaiser methodology)<sup>4</sup>: >1 outpatient diagnosis billing code, 1 outpatient diagnosis billing code and ECG consistent with AF
- 3. Demographics and *ICD* AF diagnosis billing codes: logistic regression model using patient demographics, presence of an inpatient AF diagnosis billing code, presence of an AF diagnosis billing code in the first position (primary), number of outpatient AF diagnoses billing codes, comorbid conditions and procedures from *ICD* codes, year-of-index-AF diagnosis billing code
- NLP: at least 1 nonnegated mention of AF in the clinical text (negated AF mentions use phrases such as "patient denies AF," whereas nonnegated references use phrases such as "Holter monitor showed AF")
- 5. Comprehensive: comprehensive logistic regression model combining patient demographics, presence of an inpatient AF diagnosis billing code, presence of a primary AF diagnosis billing code, number of outpatient AF diagnoses billing codes, comorbid conditions and procedures from *ICD* codes, year of index AF diagnosis billing code, at least 1 nonnegated mention of AF in clinical text, ECG with reference to AF, Current Procedural Terminology (CPT) codes for ablation or cardioversion

*ICD-9* codes were used through September, 2015 and *ICD-10* codes were used from October, 2015 onward.

# **Population and Reference Standard**

We used data from the Enterprise Data Warehouse (EDW) from University of Utah Health for this study. Enterprise data warehouses are storage systems that integrate numerous data sources within an organization (eg, inpatient and outpatient facilities, radiology reporting, or laboratory result systems) into a central repository.<sup>8</sup> Our health system uses an internally developed EDW (as opposed to a third-party data warehousing solution). The candidate population included patients with at least 1 ICD-9 or ICD-10 code for AF between 2010 and 2017 (427.31, 148.0, 148.1, 148.2, 148.9, 148.91), and without an AF diagnosis from January 1, 2007, to December 31, 2009. For model development and training, 786 patients were randomly selected from the candidate population. Chart review by a team of 5 clinicians was used to classify each patient as AF present or AF absent, which served as the reference standard. This reference standard served as the outcome for all 5 models. Each patient was classified as having AF (1) if AF was referenced in a problem list or past medical history, (2) if AF was documented but appeared only as a transient event, as part of other acute conditions (eg, cardiac surgery or sepsis), or (3) if clinic notes described active AF management. Examples of active management include procedures or medications (eg, cardioversion, anticoagulation), outside records or procedures, or listing in the assessment and plan. Otherwise, the patient was classified as not having AF. At least 2 clinicians reviewed each patient. In case of disagreement, a third reviewer adjudicated the classification. If uncertainty was still present, the team discussed the case to arrive at consensus.

#### **Feature Specification**

The feature specifications for the different models are provided in Table S1. Model features included demographics, comorbid conditions, procedures, ECG findings, and textderived features. Each of the 5 models included some combination of these features, and all features were extracted from structured data fields in the EDW (except for the textderived features). Table S1 specifies which features were included in which models. Briefly, demographic features included age, sex, race, Hispanic ethnicity, and primary insurance at the time of the index AF diagnosis. Comorbid conditions and procedures were identified based on the presence of an ICD diagnostic or procedure billing code any time during the study period. Codes were grouped into clinically meaningful groups according to the Clinical Classification Software (CCS) for the US Agency for Healthcare Research and Quality.<sup>9</sup> We used CPT codes to identify patients who had cardioversions (92960) or ablations (93651, 93655, 93656, 93657) at any time during the study period. For ECGs, we used the text interpretations and a simple regular-expression matching approach. If "atrial fibrillation," "afib," or "a fib" were present, the ECG was classified as positive.

# **Model Training**

In the 786-patient training set, we trained 5 different logistic regression models using the definitions predict the presence of AF. The models yielded predicted probabilities of AF for each training case, and the optimal cut point for each model was identified using Liu's<sup>10</sup> method, which maximizes the product of the sensitivity and specificity. In other words, each model had its own cut point. If the predicted probability was higher than the cut point, the case was classified as AF present. Accuracy, sensitivity, specificity, positive predictive value, and negative predictive value were calculated for each model, compared with the reference standard. Accuracy was defined as the number of correctly classified patients over the total number of patients. We generated and compared the area under the receiver operating characteristic curve for each model to the reference standard using Stata's "roccomp" command.

We used a rules-based NLP approach based on the pyConText algorithm, a freely available Python software

package.<sup>11–13</sup> Using the training data and clinical expertise, we identified AF-specific target terms and relevant modifiers that allow classification of each AF mention as present or absent (Table S2).<sup>14</sup> For each patient, each note was analyzed for AF-specific mentions, and each mention was classified as AF present or absent based on the modifiers surrounding the AF target term. If 1 nonnegated mention was present in any note (eg, "Patient has had long standing AF for the past 10 years"), that patient was classified as AF present. In addition, we created a summary variable for each patient, counting the total number of times an AF-specific target term appeared in the notes, regardless of negation.

# Application to Full Candidate Population

The candidate population included patients with at least 1 ICD billing diagnosis code for AF, seen between 2010 and 2017, excluding those with an AF diagnosis code going back to 2007. In other words, patients with an AF billing code in 2008 and again in 2011 would be excluded. Model training resulted in coefficients for each term in the models, which were then applied to the full candidate population. Patients with a predicted probability of AF higher than the cut point specified during training were classified as having AF. We evaluated the number of patients identified as AF per model and the apparent OAC treatment rates according to model. Patients were classified as treated with an OAC if they had an order for an OAC, including warfarin, dabigatran, apixaban, rivaroxaban, and edoxaban, in the EMR. The OAC treatment rate was calculated for patients with a CHA<sub>2</sub>DS<sub>2</sub>-VASc score  $\geq 2.^{15,16}$ The CHA2DS2-VASc score is automatically calculated in our EDW using all available prior diagnosis codes as well as ejection fraction from echocardiogram for classifying heart failure (R.U.S., unpublished data, 2019).

To compare patient characteristics between the different models, we created regression models with each characteristic as the outcome and each model as the predictor variables. In addition, we added a predictor variable that indicates whether all models agreed on whether a patient was included in the final cohort. In other words, the indicator variable equals 1 if all 5 models resulted in patients being included or excluded from the AF cohort. When the indicator variable was 1, all other predictors were reassigned to 0. Therefore, the indicator variable serves as a reference variable, or dummy variable. The likelihood ratio test was used to compare this model with a nested reduced model limited to the y-intercept. Thus, a significant P value with the likelihood ratio test indicates that at least 1 model differed in terms of inclusion of patients with a given outcome (eg. characteristic, in this case).

This study was approved by the institutional review board at the University of Utah, with a waiver of consent for patient

Characteristic	AF Present (n=632)	AF Absent (n=154)	P Value
Age, y, mean (SD)	69.0 (14.2)	61.3 (17.9)	<0.01
Female sex	249 (39.4)	81 (52.6)	<0.01
White race	563 (89.1)	132 (85.7)	0.24
Medicare insured	411 (65.0)	81 (52.6)	<0.01
No. of outpatient AF diagnoses, mean (SD)	10.1 (21.6)	1.2 (1.7)	<0.01
Primary AF diagnosis <sup>†</sup>	404 (63.9)	99 (64.3)	0.93
Comorbid conditions <sup>‡</sup>	1	1	
Acute myocardial infarction	80 (12.7)	13 (8.4)	0.15
Coronary artery disease	302 (47.8)	53 (34.4)	<0.01
Valvular heart disease	216 (34.3)	47 (30.5)	0.39
Congestive heart failure	222 (35.1)	32 (20.8)	<0.01
Cerebrovascular disease	156 (24.7)	60 (39.0)	<0.01
Dementia	183 (29.0)	39 (25.3)	0.37
Liver disease	132 (20.9)	36 (23.4)	0.50
Diabetes mellitus	289 (45.7)	62 (40.3)	0.22
Acute renal failure	161 (25.5)	25 (16.2)	0.02
Chronic kidney disease	165 (26.1)	19 (12.3)	<0.01
Pulmonary heart disease	158 (25.0)	25 (16.2)	0.02
Hypertension	465 (73.6)	103 (66.9)	0.10
Thyroid disease	190 (30.1)	29 (18.8)	<0.01
Anemia	219 (34.7)	36 (23.4)	<0.01
Cancer	235 (37.2)	33 (21.4)	<0.01
Procedures, ICD codes		-	
Heart valve surgery	27 (4.3)	5 (3.3)	0.56
Coronary artery bypass grafting	21 (3.3)	4 (2.6)	0.65
Percutaneous coronary intervention	17 (2.7)	3 (1.8)	0.60
Angioplasty	56 (8.9)	8 (5.2)	0.14
Pacemaker/defibrillator	23 (3.6)	5 (3.3)	0.81
Cardioversion	65 (10.3)	2 (1.3)	<0.01
Procedures, CPT codes			
Ablation	19 (3.0)	2 (1.3)	0.24
Cardioversion	234 (37.0)	7 (4.6)	<0.01
Natural language processing			
At least 1 nonnegated mention	614 (97.2)	57 (37.0)	<0.01
No. of AF mentions $^{\$}$			
None	16 (2.5)	76 (49.3)	<0.01
First quartile	123 (19.5)	57 (37.0)	
Second quartile	159 (25.2)	15 (9.7)	
Third quartile	162 (25.6)	6 (3.9)	
Fourth quartile	172 (27.2)	0 (0)	
ECG with reference to AF	234 (37.0)	7 (4.6)	<0.01

Values are shown as n (%), unless otherwise specified. AF indicates atrial fibrillation; CPT, Current Procedural Terminology; *ICD, International Classification of Diseases.* <sup>†</sup>Primary diagnosis refers to position 1 in the order of the billed codes.

<sup>‡</sup>Comorbid conditions were identified from *ICD* billing codes present in the patient medical record.

<sup>§</sup>Refers to the number of times a target term for AF was present in the clinical notes. The ranges are as follows: none, no mentions; first, 1–6; second, 7–19; third, 20–46; fourth, 48–670.

participation. Data set cleaning and analyses were completed using Stata v14.2, and the NLP was executed using Python. The Stata output for the model training is included in Table S3, along with the cut points and regression coefficients. The data that support the findings of this study are available from the corresponding author upon reasonable request. To protect patient information, sharing will be limited to Python scripts, in most cases.

# Results

A total of 786 patients were included in the training set, with an AF prevalence of 80.4% per our reference standard. The mean age of the training population was 67.5 years (SD: 15.3), and 42.0% of participants were female. Comorbid conditions varied between patients with and without AF, including higher rates of coronary artery disease, congestive heart failure, thyroid disease, and cancer among AF patients. Aside from cardioversion, cardiac procedures did not differ significantly between patients with and without AF (Table 1).

The test characteristics for the training models are seen in Figures 1 and 2. Compared with the reference standard, accuracy, sensitivity, and negative predictive value were highest using the NLP model, whereas specificity and positive predictive value were highest using the comprehensive model. Figure 1 shows the receiver operating characteristic curves for each model compared with the reference standard. The areas under the receiver operating characteristic curve were highest for the comprehensive and NLP models, at 0.887 and 0.801, respectively (P<0.01). The *ICD* and NLP models did not differ significantly regarding discrimination (area under the receiver operating characteristic curve: 0.801 versus 0.798, respectively; P=0.91); the *ICD* model had higher specificity at the cost of lower sensitivity. The previously published models using AF-specific diagnosis codes and ECGs resulted in high false-negative rates (Figure 2).

The full candidate population included 22 000 patients, with a mean age of 67.1 years (SD: 15.1); 42.3% were female. The number of patients, patient characteristics, and OAC treatment rates varied substantially when the models were applied to the candidate population (Table 2, Figure 3). The number of patients who could be included in an AF sample varied by up to 6690 patients. The model using outpatient AF codes and ECG resulted in the smallest AF sample, including 11 512 patients, or 52.3% of the candidate population. Comparatively, the NLP model resulted in the largest AF sample, including 18 202 patients, or 82.7% of the candidate population. The mean age of patients identified as AF was



**Figure 1.** Receiver operating characteristic curves for different models to identify atrial fibrillation patients using the electronic medical record. In the training set (n=786), the AUC was highest for the comprehensive model and lowest for the Medicare model. AUC indicates area under the receiver operating characteristic curve; *ICD, International Classification of Diseases*; NLP, natural language processing; Sens, sensitivity; Spec, specificity.



**Figure 2.** Proportion of correct, false-positive, and false-negative classifications for each model in the training set. In the training set (n=786), the NLP model resulted in the highest number of correctly classified patients, at the expense of a high false-positive rate. The outpatient billing codes and ECG method had the lowest number of correctly classified patients and the highest number of false negatives. AF indicates atrial fibrillation; *ICD, International Classification of Diseases*; NLP, natural language processing.

lowest using the outpatient AF/ECG model (68.1 years) and highest using the demographics/*ICD* codes model (70.1 years), with additional variation in comorbid conditions (Table 2). Finally, the proportion of AF patients with a CHA<sub>2</sub>DS<sub>2</sub>-VASc score  $\geq$ 2 ranged from 78.5% to 85.3%. The OAC treatment rates for patients with a score  $\geq$ 2 also varied, from 49.6% in the NLP model to 66.3% in the outpatient AF/ECG model.

# Discussion

Accurate identification of patient populations is critical for effective quality-improvement efforts. The EMR provides important opportunities to identify patient populations, but standard, electronic cohort definitions do not exist. We found that the number of AF patients included in a cohort varies by an absolute range of up to 30%, depending on which electronic cohort definition is used. In large health systems, this translates into cohorts that differ by thousands of patients. In addition, quality measures such as OAC treatment rates varied by 16.7%, between 49.6% and 66.3%, depending on the cohort definition. These findings have important implications for quality-improvement initiatives, research endeavors, and case-mix analyses.

From a quality perspective, health systems use EMR-based tools to characterize patient populations and find

opportunities for improvement.<sup>17</sup> In addition, OAC treatment rate in AF is a quality measure in the federal government's Merit-Based Incentive Payment System (MIPS)<sup>18</sup>; accurate estimation of the denominator-the number of patients with AF, in this case-is critical to its success. The AF quality measure for MIPS relies on AF billing codes and outpatient CPT codes for evaluation and management.<sup>18</sup> Based on our findings, this type of cohort selection could result in underestimating the true AF population; some electronic cohort definitions are biased and can omit a substantial number of patients, affecting the impact of the quality measure. Less sensitive AF cohort definitions would omit a large proportion of patients from any assessment of the quality of their care or related interventions. Conversely, false positives are also problematic for quality reporting because patients who do not truly have AF are unlikely to receive (or benefit from) guideline-recommended treatment.

Just as clinical trials and disease registries have specific inclusion and exclusion criteria, the same is needed for EMR-based research and initiatives. Trials and registries use "human-readable" definitions; for example, persistent AF is defined as "sustained for  $\geq$ 7 days."<sup>19</sup> These definitions should have corresponding "machine-readable" definitions to increase uniformity and reproducibility in EMR-based initiatives. The challenge is creating machine-readable definitions that are portable across institutions. In this study, for example, the outpatient AF/ECG model had suboptimal

Table 2.	Population	Characteristics	Based	on	the	Patient-Selection	Model
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Selected Characteristics	Medicare	Outpatient AF Codes, ECG	Demographics, ICD Codes	NLP	Comprehensive	P Value
Proportion identified as AF, %	18 030 (82.0)	11 512 (52.3)	13 427 (61.0)	18 202 (82.7)	15 962 (72.6)	<0.01
Age, y, mean (SD)	67.8 (14.3)	68.1 (14.1)	70.8 (12.4)	68.7 (13.8)	69.8 (13.1)	<0.01
Female sex	7434 (41.2)	4846 (42.1)	5113 (38.1)	7538 (41.4)	6528 (40.9)	<0.01
White race	15 707 (87.2)	10 143 (88.1)	11 980 (89.2)	15 957 (87.7)	14 110 (88.4)	<0.01
Medicare	11 092 (61.5)	7116 (61.8)	8874 (66.1)	11 481 (63.1)	10 389 (65.1)	<0.01
$CHA_2DS_2$ -VASc $\geq$ 2	14 920 (82.8)	9156 (79.5)	11 450 (85.3)	15 110 (83.0)	13 286 (83.2)	<0.01
OAC prescribed <sup>†</sup>	7838 (52.5)	6074 (66.3)	6572 (57.4)	7502 (49.6)	8127 (61.2)	<0.01
Comorbid conditions						
Acute myocardial infarction	2690 (14.9)	1493 (13.0)	2198 (16.4)	2567 (14.2)	2356 (14.8)	<0.01
Coronary artery disease	8463 (46.9)	5365 (46.6)	6809 (50.7)	8431 (46.3)	7496 (47.0)	<0.01
Valvular heart disease	6801 (37.7)	4001 (34.7)	5024 (37.4)	6604 (36.3)	5665 (35.5)	<0.01
Congestive heart failure	6859 (38.0)	4352 (37.8)	5766 (42.9)	3828 (37.5)	6173 (38.7)	<0.01
Cerebrovascular disease	5914 (32.8)	3077 (26.7)	3132 (23.3)	5506 (30.3)	4265 (27.7)	<0.01
Dementia	2488 (13.8)	1340 (11.6)	1776 (13.2)	2386 (13.1)	2092 (13.1)	<0.01
Diabetes mellitus	8283 (45.9)	4779 (41.5)	6219 (46.3)	8106 (44.5)	7080 (44.4)	<0.01
Chronic kidney disease	4487 (24.9)	2610 (26.7)	4082 (30.4)	4504 (24.7)	4306 (27.0)	<0.01
Hypertension	14 109 (78.3)	8729 (75.8)	10 797 (80.4)	14 068 (77.3)	12 261 (76.8)	<0.01
Cancer	6116 (33.9)	3886 (33.8)	5387 (40.1)	6257 (34.4)	5631 (35.3)	<0.01
Procedures						
Heart valve surgery	867 (4.8)	502 (4.4)	627 (4.7)	844 (4.6)	672 (4.2)	<0.01
Coronary artery bypass grafting	644 (3.6)	325 (2.8)	457 (3.4)	615 (3.4)	583 (3.7)	<0.01
Percutaneous coronary intervention	608 (3.4)	333 (2.9)	553 (4.1)	558 (3.1)	481 (3.0)	<0.01
Pacemaker/defibrillator	812 (4.5)	563 (4.9)	582 (4.3)	783 (4.3)	706 (4.4)	<0.01

Values shown as n (%), unless otherwise specified. AF indicates atrial fibrillation; *ICD, International Classification of Diseases*; NLP, natural language processing; OAC, oral anticoagulant. <sup>†</sup>Including only patients with CHA<sub>2</sub>DS<sub>2</sub>-VASc ≥2.

performance, whereas it may perform well in the system for which it was designed. We showed it cannot easily be applied to a system in which patients receive fragmented care from different institutions using different EMRs.

Furthermore, many administrative data sets and definitions are used to calculate observed-to-expected event ratios in efforts to understand quality of care (and, on occasion, payment, scoring, etc). Underlying these calculations is a case mix, to account for severity of illness, on which to base expected outcomes. Once again, such case-mix analyses could vary dramatically with the definition of the underlying disease-based cohort, leading to wide variability in expected outcomes, observed outcomes, and downstream effects. With more precise, portable, cohort-definition methods, precision and utility of such analyses could improve dramatically.

NLP may have some advantages over billing data models because the clinical narrative may be less prone to certain types of variation, given that there are relatively few ways that clinicians state that a patient has AF. However, we were limited by low specificity with our rules-based approach. Machine learning, as opposed to rules-based approaches, can also be used for NLP but often requires large sets of labeled training data. Our future efforts will focus on improving NLP specificity by using the comprehensive model to automatically label patients and create a large training set for a machine learning approach. Still, whatever methods result in the ideal performing model (text or structured data, eg, ICD codes), we will need to ensure that the model is calibrated and portable. Efforts are underway using common data models such as the Observational Medical Outcomes Partnership (OMOP) common data model,<sup>20</sup> but standardization of the data that go into the common data model must also be a part of the process. For example, 2 different NLP systems can extract AF patients and map the concept to OMOP, but the systems are different; the common data model alone does not solve the portability issue. In the



**Figure 3.** Proportion of patients included with  $CHA_2DS_2$ -VASc score  $\geq 2$  and treated with an OAC for each model. When applied to the candidate population, different patient-selection models resulted in populations with different sizes, stroke risks, and OAC treatment rates. The corresponding values are found in Table 2. "Outpatient AF codes, ECG" refers to the method used in prior publications from Kaiser Permanente. AF indicates atrial fibrillation; *ICD, International Classification of Diseases*; NLP, natural language processing; OAC, oral anticoagulant.

future, if EMRs become more similar, an option is for guidelines and regulations to include validated algorithms along with recommendations.

# Limitations

We used diverse approaches to patient electronic cohort definitions in this study, and this is only 1 factor that can skew outcome results. CHA2DS2-VASc score calculations and OAC treatment classification methods can also vary and yield different apparent treatment rates. In addition, we used only a small fraction of the variables available in the EMR. We chose features based on widespread availability (eg. demographics) and controlled vocabularies (eg, ICD and CPT) for this demonstration project. Additional features, such as ejection fraction, have varying capture and format across institutions; adding features and increasing model complexity could decrease bias but would probably increase model overfitting and result in site-specific, nonportable models. Our candidate population, including patients with at least 1 AF billing code, was enriched with a high prevalence of "AF present" patients. Generation of models that accurately identify low-prevalence conditions, such as AF patients in an entire health system, is limited by challenges in creating a reference standard; manual chart review to identify 1% of the population is cumbersome, if not impossible. Our reference standard definition of AF was broad, and the results would differ with narrower definitions. From this larger group, health systems could apply criteria to select patient subsets, such as patients who have at least 2 outpatient encounters, a designated primary care physician within the health system, or a first AF encounter during admission for cardiac surgery. Finally, both billing codes and text-based terms vary between institutions. We did not include internal and external validation populations for each model because the purpose of this study was not to identify the optimal model to select AF patients but rather to compare population characteristics and outcomes from different approaches.

# Conclusions

EMRs provide an opportunity to identify large patient cohorts for research and quality initiatives. Cohort selection is a critical step to realizing the potential of EMRs for quality improvement and research and a prerequisite to developing learning healthcare systems. Cohort definitions should be based on validated portable definitions to maximize comparability. In the case of AF, number of patients, characteristics, and outcomes vary depending on the patient-selection method. To optimize the impact of EMR-driven research and quality improvement, we need an unbiased, portable approach to identify patient populations. Combining multiple types of data from EMRs may serve this goal. Nevertheless, regardless of the data sources structured data like *ICD* codes or unstructured data like text we will ultimately require a common AF definition for use in research and quality improvement.

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# SUPPLEMENTAL MATERIAL

# Table S1. Model specifications.

Feature	Definition	Models
Kaiser	>1 outpatient ICD AF diagnosis billing code OR	Model 2
	1 outpatient ICD AF diagnosis billing code and ECG consistent with AF	
Medicare	≥1 inpatient ICD AF diagnosis billing code OR ≥2 outpatient ICD AF diagnoses billing codes within 365 days	Model 1
Demographics: Based on status at the	time of the index AF encounter	
Age	Ordinal; roughly according to decade (18 to 29, 30 to 39, 40 to 49, 50 to 59, 60 to 69, 70 to 79, 80 and older)	Model 3 Model 5
Sex	Binary; female (reference) or male	Model 3
		Model 5
Race	Categorical; white (reference), black, Asian, other/missing	Model 3
		Model 5
Hispanic	Categorical; Hispanic, not Hispanic, or missing	Model 3
		Model 5
Primary payer	Categorical; Medicare (reference), Medicaid, private, self, other/missing	Model 3
		Model 5
Inpatient AF diagnosis	Binary; presence of an inpatient billing code at any time during the study period	Model 1
		Model 3
		Model 5
Number of outpatient AF diagnoses	Numeric; the total number of outpatient AF billing codes during the study period	Model 3
		Model 5
Primary AF diagnosis	Binary; presence of an AF billing code in the first billing position any time during the study	Model 3
	penda	Model 5
Acute myocardial infarction	Binary; presence of an ICD diagnosis billing code for condition any time during the study	Model 3
	period	Model 5
Comorbid Conditions: Based on ICD b	lling codes	
Coronary artery disease	Binary; presence of an ICD diagnosis billing code for condition any time during the study	Model 3
Valvular heart disease		Model 5
Congestive heart failure		

Peripheral vascular disease		
Cerebrovascular disease		
Dementia		
Pulmonary heart disease		
Rheumatologic disease		
Gastrointestinal ulcer		
Liver disease		
Diabetes mellitus		
Acute renal disease		
Chronic renal disease		
Lymphoma		
Hypertension		
Coagulopathy		
Electrolyte disorder		
Anemia		
Cancer		
Dialysis		
Procedures: Based on ICD billing codes		
Heart valve surgery	Binary; presence of an ICD procedure billing code for procedure any time during the study	Model 3
Coronary artery bypass grafting	period	Model 5
Percutaneous coronary intervention		
Angiogram		
Pacemaker/defibrillator		
Cardioversion (ICD based)		
Procedures: Based on CPT billing code:	5	
Cardioversion (CPT based)	Binary; presence of a CPT billing code for procedure any time during the study period	Model 5
Ablation		
Electrocardiograms and text		
Electrocardiogram	Binary; presence of an ECG interpretation that includes an AF-specific term any time during	Model 2
	the study period	Model 5
Number of AF mentions in the text	Categorical, split into zero and quartiles for values >0; the total number of AF mentions, as extracted by NLP, in the available text any time during the study period	Model 5
Non-negated AF mention in the text	Binary; the presence of a non-negated reference to AF in the available text any time during the study period	Model 4

Model Definitions (also see Methods section of manuscript):

Model #1: Outpatient and inpatient AF billing codes (Medicare methodology)<sup>2</sup>:  $\geq$ 1 inpatient diagnosis or  $\geq$ 2 outpatient diagnoses within 365 days

Model #2: Outpatient AF codes and electrocardiogram (Kaiser methodology)<sup>4</sup>: >1 outpatient ICD code, 1 outpatient ICD code and ECG consistent with AF

Model #3: Demographics and International Classification of Diseases (ICD) billing AF codes: A logistic regression model using patient demographics, presence of an inpatient AF diagnosis, presence of a primary AF diagnosis, number of outpatient AF diagnoses, comorbid conditions and procedures from ICD codes, year of index AF diagnosis

Model #4: Natural language processing: At least one non-negated mention of AF in the clinical text

Model #5: Comprehensive: A comprehensive logistic regression model combining patient demographics, presence of an inpatient AF diagnosis, presence of a primary AF diagnosis, number of outpatient AF diagnoses, comorbid conditions and procedures from ICD codes, year of index AF diagnosis, at least one non-negated mention of AF in clinical text, ECG with reference to AF, CPT codes for ablation or cardioversion

AF=atrial fibrillation; CPT=current procedural terminology; ICD=International Classification of Diseases

Table S2. Target terms used in natural language processing task to identify atrial fibrillation patients from clinical notes.

Target term	Regular Expression
afib	\bafib\b \batrial\sfib a-fib a\.\sfib a\.fib \ba\sfib\b
Modifier terms	
no	\bno(?!\sfurther)\b
not	\bnot\b
none	\bnone\b
negative	\bnegative\b
denies	denies denied denying
family	\bmother\b \bfather\b \bsister\b \bbrother\b \bdaughter\b \bson\b \baunt\b \ buncle\b \bgranddaughter\b \bgrandson\b
rule out	r/o r\\o \brule\s+out\b \brules\s+out\b \bruled\s+out\b
unlikely	\bunlikely\b
investigate	\binvestigate\b \binvestigating\b
look for	\blook\s+for\b\b
differential	\bdifferential\b\b ddx
possible	\bpossible\b
holter	\b(holter event)\s+(monitor(ing)?\s+)?ordered\s+for\b\b ddx
etc	\betc\b
screen for	\bscreen\s+for\b
risk of	\brisk\s+(of for)\b
suspicious	\bsuspicious\b
question of	\bquestion\s+of\b

Table S3. Training model regression results.

ogistic regression		Numbe	er of obs	=	786
		LR ch	ni2(1)	=	78.77
		Prob	> chi2	= 0	.0000
og likelihood = -349.45524		Pseud	lo R2	= 0	.1013
inary_adj_goldstd   Coef. St	d. Err.	Z	P> z	[95% Conf	. Interval]
++					
kaiser   1.723243 .2	125683	8.11	0.000	1.306616	2.139869
_cons   .7256704 .	110745	6.55	0.000	.5086141	.9427267
mpirical cutpoint estimation					
ethod:	Liu				
eference variable:	binary_	_adj_gol	.dstd (0=ne	eg, 1=pos)	
lassification variable:	kaiser_	_lr			
pirical optimal cutpoint:	. /9/168	32			
pecificity at cutpoint:	0.80				
rea under ROC curve at cutpoint:	0.70				
415 real changes made)					
etailed report of sensitivity and s	pecificity	7			
	Cor	rectly			

60.44% 78.57% 63.99% 2.8207 0.5035 ( >= 1 ) \_\_\_\_\_ ROC -Asymptotic Normal--Obs Area Std. Err. [95% Conf. Interval] \_\_\_\_\_ 786 0.6951 0.0192 0.65738 0.73276 binary\_adj | kaiser\_class \_goldstd | Pos. Neg. | Total Abnormal | 382 250 | 632 Normal | 33 121 | 154 Total | 415 371 | 786 True abnormal diagnosis defined as binary adj goldstd = 1 [95% Confidence Interval] \_\_\_\_\_ 80% 77% Prevalence Pr(A) 83.1% \_\_\_\_\_ Sensitivity Pr(+|A) 60.4% 56.5% 64.3% Pr(-|N) 78.6% 71.2% 84.8% Specificity (Sens. + Spec.)/2 .695 .657 .733 ROC area \_\_\_\_\_ Likelihood ratio (+) Pr(+|A)/Pr(+|N) 2.82 2.07 Likelihood ratio (-) Pr(-|A)/Pr(-|N) .503 .443 3.84 .572 5.6 3.7 8.48 LR(+)/LR(-) Odds ratio Pr(A|+) 92% 89% 94.5% Positive predictive value Pr(N|-) 32.6% 27.9% 37.6% Negative predictive value \_\_\_\_\_ MEDICARE MODEL Number of obs = 786 Logistic regression LR chi2(1) = 65.41 Prob > chi2 = 0.0000

Pseudo R2 = 0.0841 Log likelihood = -356.13488\_\_\_\_\_ Coef. Std. Err. z P>|z| [95% Conf. Interval] binary\_adj\_goldstd | simpleicd | 1.675786 .2039045 8.22 0.000 1.27614 2.075431 cons | .1563462 .1689453 0.93 0.355 -.1747804 .4874729 \_\_\_\_\_ Logistic model for binary adj goldstd number of observations = 786 area under ROC curve = 0.6509 Empirical cutpoint estimation Method: Liu Reference variable: binary adj goldstd (0=neg, 1=pos) Classification variable: medicare lr Empirical optimal cutpoint: .70051131 Sensitivity at cutpoint: 0.88 Specificity at cutpoint: 0.42 Area under ROC curve at cutpoint: 0.65 (645 real changes made) Detailed report of sensitivity and specificity \_\_\_\_\_ Correctly Cutpoint Sensitivity Specificity Classified LR+ LR-\_\_\_\_\_ ( >= 1 ) 87.97% 42.21% 79.01% 1.5223 0.2849 \_\_\_\_\_ ROC -Asymptotic Normal--Obs Area Std. Err. [95% Conf. Interval] \_\_\_\_\_ 786 0.6509 0.0210 0.60978 0.69205

binary_adj	medica	are_class						
_goldstd	Pos.	. Neg.	Total					
+			+					
Abnormal	556	5 76	632					
Normal	89	65	154					
+			+					
Total	645	5 141	786	i				
True abnorma	al diagnosi	is defined as	binary_adj	_goldst	:d = 1			
				[ 9	95% Confi	dence Inter	val]	
Prevalence			Pr(A)	80%	778	83.1%	5	
Sensitivity		P	r(+ A)	88%	85.28	90.4%		
Specificity		P	r(- N)	42.2%	34.38	50.4%		
ROC area		(Sens. + Sp	ec.)/2	.651	.61	.692		
Likelihood r	ratio (+)	Pr(+ A)/P	r(+ N)	1.52	1.33	1.75		
Likelihood r	ratio (-)	Pr(- A)/P	r(- N)	.285	.215	.377		
Odds ratio		LR(+)	/LR(-)	5.34	3.59	7.96		
Positive pre	edictive va	alue P.	r (A +)	86.2%	83.38	88.8%		
Negative pre	edictive va	alue P.	r(N -)	46.1%	37.78	54.7%		
ICD MODEI								
Logistic rec	ression			Numł	er of ob	)s =	78	6
20920020 209	,10001011			LR (	:hi2(43)	=	2.60.7	3
				Prob	$\rightarrow$ chi2	=	0.000	- 0
Log likeliho	pod = -258	47662		Psei	ido R2	=	0.335	3
								-
binary ad	hteblon i	l Coef	Std Err		7 P>1	71 [95%	Conf	Intervall
		+				_, [,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,		
	agegrp1	.0446484	.0099897	4	47 0.0	دە. 00	25069	.0642279
		358715	.2435280	1	47 0 1	41 - 119	35929	.8360229
	JEA		.2133203	1.			, , , , , , , , , , , , , , , , , , , ,	.0300229

I							
race_categ							
Black	-1.942464	1.308129	-1.48	0.138	-4.506349	.6214212	
Asian	-1.375892	1.244579	-1.11	0.269	-3.815221	1.063437	
Other/missing	4826733	.4840383	-1.00	0.319	-1.431371	.4660243	
1							
hispanic	.0927108	.1684888	0.55	0.582	2375211	.4229427	
I							
index_pay1_categ							
2	.8464419	.6084834	1.39	0.164	3461637	2.039047	
3	.0162844	.3308754	0.05	0.961	6322194	.6647882	
4	.4823887	.6985767	0.69	0.490	8867964	1.851574	
5	2.056091	1.095191	1.88	0.060	0904435	4.202626	
I							
inpatientdx	1.640814	.3087453	5.31	0.000	1.035684	2.245943	
countoutpatient_afib	.4895864	.0782739	6.25	0.000	.3361723	.6430004	
afibicd_primary	9130516	.2387842	-3.82	0.000	-1.38106	4450431	
index_year	.0646928	.0550775	1.17	0.240	0432571	.1726428	
amidiag_all	.3809207	.4520047	0.84	0.399	5049923	1.266834	
caddiag_all	.0267004	.2832404	0.09	0.925	5284405	.5818414	
valvediag_all	121885	.2692074	-0.45	0.651	6495218	.4057518	
chfdiag_all	.2366036	.3143618	0.75	0.452	3795342	.8527414	
pvddiag_all	6911086	.3083174	-2.24	0.025	-1.2954	0868175	
cvddiag_all	-1.278336	.2852031	-4.48	0.000	-1.837324	7193479	
dementiadiag_all	0838705	.3846242	-0.22	0.827	8377201	.669979	
pulmdzdiag_all	2523725	.2855902	-0.88	0.377	812119	.3073739	
rheumdiag_all	.7370864	.5824089	1.27	0.206	404414	1.878587	
ulcerdiag_all	.0170389	.5962036	0.03	0.977	-1.151499	1.185576	
liverdiag_all	4056906	.3076358	-1.32	0.187	-1.008646	.1972644	
dmdiag_all	0513154	.2568892	-0.20	0.842	554809	.4521781	
renaldiag_all	.0839594	.4110108	0.20	0.838	7216069	.8895257	
ckddiag_all	.7758547	.4285092	1.81	0.070	0640079	1.615717	
lymphdiag_all	.7620735	.8795399	0.87	0.386	961793	2.48594	
pulmhtndiag_all	.1058361	.3386697	0.31	0.755	5579443	.7696164	
htndiag_all	.0271604	.2786255	0.10	0.922	5189355	.5732563	

thyroiddiag_all	.2414638	.2949052	0.82	0.413	3365399	.8194674	
coagdiag_all	.3509488	.3714666	0.94	0.345	3771123	1.07901	
elecdiag_all   -	.1480849	.3050795	-0.49	0.627	7460297	.44986	
anemiadiag_all	.1875522	.3232971	0.58	0.562	4460984	.8212028	
cancerdiag_all	.3973601	.2820185	1.41	0.159	155386	.9501062	
dialysis_icdproc_all   -	.3362597	.7277529	-0.46	0.644	-1.762629	1.09011	
valve_icdproc_all   -	.1382759	.6785085	-0.20	0.839	-1.468128	1.191576	
cabg_icdproc_all   -	.0303663	.8015717	-0.04	0.970	-1.601418	1.540685	
pci_icdproc_all	.2712517	.9442262	0.29	0.774	-1.579398	2.121901	
angio_icdproc_all	.0102707	.6437773	0.02	0.987	-1.25151	1.272051	
ppm_defib_icdproc_all   -	.8168417	.6898823	-1.18	0.236	-2.168986	.5353027	
dccv_icdproc_all	1.196939	.8358149	1.43	0.152	4412278	2.835106	
_cons	-132.932	110.9735	-1.20	0.231	-350.436	84.57198	
number of observations =	786 0.8738						
Empirical cutpoint estimat	ion						
Method:		Liu					
Reference variable:		binary_a	dj_goldst	d (0=neg	, 1=pos)		
Classification variable:		icd_lr					
Empirical optimal cutpoint	:	.7704673	4				
Sensitivity at cutpoint:		0.78					
Specificity at cutaciat.							
specificity at cutpoint:		0.81					
Area under ROC curve at cu	itpoint:	0.81					
Area under ROC curve at cu (525 real changes made)	tpoint:	0.81					
Area under ROC curve at cu (525 real changes made) Detailed report of sensiti	vity and s	0.81 0.80 pecificity					
Area under ROC curve at cu (525 real changes made) Detailed report of sensiti	vity and s	0.81 0.80 pecificity Corr					
Area under ROC curve at cu (525 real changes made) Detailed report of sensiti  Cutpoint Sensitivity	vity and s Specific	0.81 0.80 pecificity  Corr ity Class	 ectly ified	LR	+ 1	 LR-	

( >= 1 )		78.48%	81.17%		79.01%	4.167	6 0.26
		ROC			-Asymptot	ic Normal-	_
	Obs	Area	Std. Err.		[95% Conf	. Interval	]
							_
	786	0.7982	0.0178		0.76337	0.8331	3
binary adj	i I	icd class					
_goldstd	1   :	Pos.	Neg.	Total			
	+		+				
Abnormal	-	496	136	632			
Normal	-	29	125	154			
Total	_	525	261	786			
					[95%	Confidence	e Interval]
Prevalence	2		Pr (A)		80%	77%	83.1%
Sensitivit	су		Pr(+ A)		78.5%	75.1%	81.6%
Specificit	у		Pr(- N)	:	31.2%	74.1%	87%
ROC area		(Sens	. + Spec.)/2	2	.798	.763	.833
Likelihood	l ratio (	+) Pr <i>l</i>	+ A)/Pr(+ N)		4.17	2.99	5,8
Likelihood Likelihood	l ratio (· l ratio (·	+) Pr( -) Pr(	+   A) / Pr (+   N) -   A) / Pr (-   N)		4.17 .265	2.99	5.8 .313
Likelihood Likelihood Odds ratic	d ratio (* d ratio (*	+) Pr( -) Pr(	+ A)/Pr(+ N) - A)/Pr(- N) LR(+)/LR(-)		4.17 .265 15.7	2.99 .224 10.1	5.8 .313 24.5
Likelihood Likelihood Odds ratic Positive p	d ratio (* d ratio (* o predictive	+) Pr( -) Pr( e value	+ A)/Pr(+ N) - A)/Pr(- N) LR(+)/LR(-) Pr(A +)		4.17 .265 15.7 94.5%	2.99 .224 10.1 92.2%	5.8 .313 24.5 96.3%
Likelihood Likelihood Odds ratic Positive p Negative p	d ratio (* d ratio (* o predictive predictive	+) Pr( -) Pr( e value e value	+   A) /Pr (+   N) -   A) /Pr (-   N) LR (+) /LR (-) Pr (A   +) Pr (N   -)		4.17 .265 15.7 94.5% 47.9%	2.99 .224 10.1 92.2% 41.7%	5.8 .313 24.5 96.3% 54.1%
Likelihood Likelihood Odds ratic Positive p Negative p	d ratio ( d ratio ( ) predictive predictive	+) Pr( -) Pr( e value e value	+   A) / Pr (+   N) -   A) / Pr (-   N) LR (+) / LR (-) Pr (A   +) Pr (N   -)		4.17 .265 15.7 94.5% 47.9%	2.99 .224 10.1 92.2% 41.7%	5.8 .313 24.5 96.3% 54.1%
Likelihood Likelihood Odds ratic Positive p Negative p	d ratio (* d ratio (* opredictive predictive EL	+) Pr( -) Pr( e value e value	+ A)/Pr(+ N) - A)/Pr(- N) LR(+)/LR(-) Pr(A +) Pr(N -)		4.17 .265 15.7 94.5% 47.9%	2.99 .224 10.1 92.2% 41.7%	5.8 .313 24.5 96.3% 54.1%

	LR ch	i2(1)	=	287.78
	Prob	> chi2	=	0.0000
Log likelihood = -244.94682	Pseud	o R2	=	0.3701
binary_adj_goldstd   Coef. Std. Err	r. z	₽> z	[95% Cc	onf. Interv
afnlp_mrn_predict   4.061283 .2916117	7 13.93	0.000	3.48973	4.632
_cons   -1.684339 .2566414	4 -6.56	0.000	-2.18734	-1.181
Logistic model for binary adj goldstd				
number of observations = 786				
area under ROC curve = 0.8007				
Empirical cutpoint estimation				
Method: Liu	L			
Reference variable: bir	nary_adj_gol	dstd (O=ne	g, 1=pos	5)
Classification variable: afr	nlp_lr			
Empirical optimal cutpoint: .53	3578696			
Sensitivity at cutpoint: 0.9	97			
Specificity at cutpoint: 0.6	63			
Area under ROC curve at cutpoint: 0.8	30			
(671 real changes made)				
Detailed report of sensitivity and specifi	icity			
Cutnoint Consistivity Consistivity	Classified	Ŧ	DT	מ ד
calpoint sensitivity specificity		ىل = = = = = = = = = = = = =	лт 	шк-
( >= 1 ) 97.15% 62.99%	90.46%	2.62	48	0.0452
ROC	-Asympto	tic Normal		

	786	0.80	)7 (	0.0198		0.7618	9 0.8	3949		
binary_adj	I	afnlp_	_class							
_goldstd	I	Pos.	Neg	.	Total					
Abnormal	-+	614		+ 8	632					
Normal	I	57	9.	7	154					
Total	-+ 	671	11	+ 5	786					
True abnorr	mal dia	gnosis	defined a	as bina	ary_adj	_goldst	.d = 1			
						[9]	95% Confid	lence Inter	val]	
Prevalence				Pr ( <i>1</i>		80%	77%	83.1%		
Sensitivity	 У			Pr (+ 7		97.2%	95.5%	98.3%		
Specificity	Y			Pr(- 1	1)	63%	54.8%	70.6%		
ROC area			(Sens. + :	Spec.)/	/2	.801	.762	.839		
Likelihood	ratio	(+)	Pr(+ A),	/Pr(+ N	1) 	2.62	2.14	3.23		
Likelihood	ratio	(-)	Pr(- A)	/Pr(- 1	1)	.0452	.0282	.0724		
Odds ratio			LR(·	+)/LR(-	-)	58	32.9	102		
Positive p	redicti	.ve val	ıe	Pr(Al+	+)	91.5%	89.1%	93.5%		
Negative p	redicti	ve val	le	Pr(N -	-)	84.3%	76.4%	90.5%		
COMPREHE	ENSIV	e mod	EL							
Logistic regression				Numb	er of obs	; =	786			
						LR c	:hi2(47)	=	459.16	
						Prob	> chi2	=	0.0000	
Log likelih	hood =	-159.2	6033			Pseu	ido R2	=	0.5904	
binary_a	adj_gol	.dstd	Coe:	f. St	ed. Err		z P>  z	:  [95%	Conf. Interva	 al]

++							
afnlp_mrn_predict	2.2063	.5049817	4.37	0.000	1.216554	3.196046	
agegrp1	.0328202	.0130682	2.51	0.012	.0072069	.0584335	
sex	.006181	.3288827	0.02	0.985	6384173	.6507793	
I							
race_categ							
Black	-2.187707	1.514649	-1.44	0.149	-5.156364	.7809499	
Asian	.2546778	3.239757	0.08	0.937	-6.095129	6.604485	
Other/missing	656952	.6036018	-1.09	0.276	-1.83999	.5260858	
I							
hispanic	.1670156	.2225115	0.75	0.453	269099	.6031302	
I							
index_pay1_categ							
2	.5163838	.7883102	0.66	0.512	-1.028676	2.061443	
3	2421059	.4610076	-0.53	0.599	-1.145664	.6614523	
4	.9414504	.8906946	1.06	0.291	804279	2.68718	
5	2.136887	1.231747	1.73	0.083	2772917	4.551066	
I							
inpatientdx	1369238	.4172727	-0.33	0.743	9547633	.6809157	
countoutpatient_afib	.1403936	.0740849	1.90	0.058	0048101	.2855974	
afibicd_primary	-1.262697	.3451367	-3.66	0.000	-1.939153	5862419	
index_year	0405696	.075878	-0.53	0.593	1892877	.1081485	
amidiag_all	.561467	.6321116	0.89	0.374	6774491	1.800383	
caddiag_all	0842321	.3797421	-0.22	0.824	8285129	.6600487	
valvediag_all	0823799	.3639308	-0.23	0.821	7956711	.6309113	
chfdiag_all	113876	.4480115	-0.25	0.799	9919625	.7642104	
pvddiag_all	6556172	.4073686	-1.61	0.108	-1.454045	.1428106	
cvddiag_all	-1.12224	.3907799	-2.87	0.004	-1.888155	3563258	
dementiadiag_all	.0200449	.5326683	0.04	0.970	-1.023966	1.064056	
pulmdzdiag_all	3854925	.3913458	-0.99	0.325	-1.152516	.3815311	
rheumdiag_all	.5394939	.8438259	0.64	0.523	-1.114374	2.193362	
ulcerdiag_all	.2380188	.8312635	0.29	0.775	-1.391228	1.867265	
liverdiag_all	4841053	.4199376	-1.15	0.249	-1.307168	.3389572	
dmdiag_all	0962111	.3521609	-0.27	0.785	7864338	.5940115	
renaldiag_all	2154304	.5936916	-0.36	0.717	-1.379045	.9481838	

ckddiag_all   1.626106	.6297206	2.58	0.010	.391876	2.860335			
lymphdiag_all   .5713589	1.101511	0.52	0.604	-1.587562	2.73028			
pulmhtndiag_all   .4727443	.4709309	1.00	0.315	4502633	1.395752			
htndiag_all  5871352	.3913627	-1.50	0.134	-1.354192	.1799215			
thyroiddiag_all   .012911	.4038323	0.03	0.974	7785859	.8044078			
coagdiag_all   .4569073	.5011965	0.91	0.362	5254199	1.439234			
elecdiag_all  613357	.4176761	-1.47	0.142	-1.431987	.2052732			
anemiadiag_all   .2884396	.4747701	0.61	0.543	6420927	1.218972			
cancerdiag_all   .1483555	.38792	0.38	0.702	6119538	.9086648			
dialysis_icdproc_all  9619088	1.086018	-0.89	0.376	-3.090466	1.166648			
valve_icdproc_all   -1.106743	.9657247	-1.15	0.252	-2.999529	.7860425			
cabg_icdproc_all   1.961944	1.187286	1.65	0.098	3650924	4.288981			
pci_icdproc_all  623917	1.42916	-0.44	0.662	-3.425019	2.177185			
angio_icdproc_all  3559314	.8904829	-0.40	0.689	-2.101246	1.389383			
ppm_defib_icdproc_all   .0118631	1.02624	0.01	0.991	-1.99953	2.023257			
dccv_icdcpt_binary  0867684	1.027261	-0.08	0.933	-2.100162	1.926625			
ablate_cpt_binary   -1.658924	1.220243	-1.36	0.174	-4.050557	.7327092			
ecg_afib   2.881581	.6455071	4.46	0.000	1.616411	4.146752			
q_afnlp_total   1.182809	.2549187	4.64	0.000	.683178	1.682441			
_cons   78.62454	152.8337	0.51	0.607	-220.9241	378.1731			
Note: 0 failures and 14 successes c	Note: 0 failures and 14 successes completely determined.							
Logistic model for binary_adj_golds	td							
number of observations = 786								
area under ROC curve = 0.9504								
Empirical cutpoint estimation								
Method:	Liu	Liu						
Reference variable:	binary_a	<pre>binary_adj_goldstd (0=neg, 1=pos)</pre>						
Classification variable:	comprehe	comprehensive_lr						
Empirical optimal cutpoint:	.7892637	.7892637						
Sensitivity at cutpoint:	0.90	0.90						
Specificity at cutpoint:	0.87	0.87						
Area under ROC curve at cutpoint:	Area under ROC curve at cutpoint: 0.89							
1								

(591 real changes made) Detailed report of sensitivity and specificity \_\_\_\_\_ Correctly Cutpoint Sensitivity Specificity Classified LR+ LR-\_\_\_\_\_ 90.35% 87.01% 89.69% 6.9568 0.1109 ( >= 1 )\_\_\_\_\_ ROC -Asymptotic Normal--Obs Area Std. Err. [95% Conf. Interval] \_\_\_\_\_ 786 0.8868 0.0148 0.85779 0.91582 binary\_adj | comprehensive\_class goldstd | Total Pos. Neg. | Abnormal | 571 61 | 632 20 134 | Normal | 154 Total | 591 195 | 786 True abnormal diagnosis defined as binary\_adj\_goldstd = 1 [95% Confidence Interval] \_\_\_\_\_ Pr(A) 80% 77% 83.1% Prevalence \_\_\_\_\_ Sensitivity Pr(+|A) 90.3% 87.8% 92.5% Pr(-|N) 87% 80.7% 91.9% Specificity (Sens. + Spec.)/2 .887 .858 .916 ROC area Likelihood ratio (+) Pr(+|A)/Pr(+|N) 6.96 4.62 10.5 Likelihood ratio (-) Pr(-|A)/Pr(-|N) .111 .0867 .142 LR(+)/LR(-) 62.7 36.7 107 Odds ratio

Positive predictive value	Pr(A +)	96.6%	94.8%	97.9%
Negative predictive value	Pr(N -)	68.7%	61.7%	75.2%