SUPPLEMENTAL MATERIAL

- KiCS-AF Registry Coder's Data Dictionary
- The list of ethical review board which approved the study
- Supplemental Table 1: Definitions of Cardiovascular death
- Supplemental Table 2: The number of patients who refused to participate in the KiCS-AF registry at each participating center

		1. Patient Demographics
Seq. # 1101	Name:	System ID
	Coding instruction:	Indicate the patient's ID in the KiCS-AF registry (auto)
	Target Value :	At the time of registration
Seq. # 1104	Name:	Date of Birth
	Coding instruction:	Indicate the patient's date of birth
	Target Value :	At the time of registration
Seq. # 1105	Name:	Sex
	Coding instruction:	Indicate the patient's sex at birth
	Target Value :	Status at the initial visit
	Selection:	1. Male
		2. Female
	Supporting Definition:	(none)
Seq. # 1106	Name:	Encounter Date
	Coding instruction:	Indicate the date of the patient encounter or visit to the MD office
	Target Value:	At the time of registration
Seq. # 1107	Name:	Registration Date
	Coding instruction:	Indicate the patient's registration date to the KiCS-AF registry
	Target Value:	At the time of registration
Seq. # 1108	Name:	Referral from Emergency Department
	Coding instruction:	Indicate if the patient referral from emergency department
	Target Value:	At the time of registration
	Selection:	1. No (planned outpatient)
		2. Yes
Seq. # 1109	Name:	Symptoms at Encounter
	Coding instruction:	Indicate the patient's symptoms related to atrial fibrillation on
		encounter
	Target Value :	At the time of initial visit (Multiple-choice)
	Selection:	1. Palpitation
		2. Dyspnea (at rest, on exertion)
		3. Difficulty in activities
		4. Dizziness
		5. Fatigue
		6. Chest pain
		7. Syncope
0		8. No symptoms related with atrial fibrillation
Seq. # 1110	Name:	Diagnosed AF with medical check-up/screening

i i		Indicate if the patient has diagnosed atrial fibrillation with medical
	Coding instruction:	check-up/screening (eg. human dry dock)
	Tanget Value	At the time of initial visit
	Selection :	
	Selection.	2. Yes
Seq. # 1111	Name:	
Seq. # 1111		Indicate the patient's height in centimeters (cm).
	_	At the time of initial visit
Seq. # 1112	Name:	
Seq. # 1112		Indicate the patient's weight
	_	At the time of initial visit
Seq. # 1113	G	Systolic Blood Pressure
Seq. # 1113		Indicate the patient's systolic blood pressure
	_	At the time of initial visit
Seq. # 1114		Diastolic Blood Pressure
Seq. # 1114		Indicate the patient's diastolic blood pressure
	G	At the time of initial visit
Seq. # 1115	G	Heart Rate
Seq. π 1113		Indicate the patient's heart rate
	_	At the time of initial visit
Seq. # 1116	G	Heart Rhythm
Seq. # 1110		Indicate the patient's heart rhythm
	_	At the time of initial visit
	G	1. Sinus Rhythm
		2. Atrial Fibrillation
		3. Atrial Flutter
		4. Others
Seq. # 1117	Name:	
•	Coding instruction:	Indicate the findings of the patient's electrocardiogram
	G	At the time of initial visit (Multiple-choice)
		1. Left ventricular hypertrophy
		2. Left bundle branch block
		3. Right bundle branch block
		4. 1st degree atrioventricular block
		5. Nothing detected
Seq. # 1118	Name:	-
		Indicate the patient's QTc time (meec) in the electrocardiogram
	_	At the time of initial visit
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Seq. # 1119	Nama ·	QRS Time in Electrocardiogram
Seq. # 1117		Indicate the patient's QRS Time (meec) in the electrocardiogram
	_	At the time of initial visit
	Target value:	2. Medication
Seq. # 1201		Aspirin Prescribed
	_	Indicate if the patient had Aspirin prescribed
		At the time of initial visit
	Selection:	
C // 1202	Name	2. Yes
Seq. # 1202		Clopidogrel Prescribed
		Indicate if the patient had Clopidogrel prescribed
		At the time of initial visit
	Selection:	
Com # 1202	Nama	2. Yes
Seq. # 1203		Other Antiplatelet Agents Prescribed
	_	Indicate if the patient had other Antiplatelet agents prescribed
	Selection:	At the time of initial visit
	Selection:	2. Yes
Seq. # 1204	Name :	Warfarin Prescribed
Seq. # 1204		Indicate if the patient had warfarin prescribed
	_	At the time of initial visit
	Selection:	
	Selection .	2. Yes
		If chose "Yes", indicate dose (mg)
Seq. # 1205	Name:	
20 4 1200		Indicate if the patient had dabigatran prescribed
	G	At the time of initial visit
	Selection :	
		2. Yes
		If chose "Yes", indicate the dose (mg)
	Selection:	
		2. 220 mg
		3. 300 mg
Seq. # 1206	Name:	Rivaroxaban Prescribed
	Coding instruction :	Indicate if the patient had rivaroxaban prescribed
		At the time of initial visit
	Selection:	
1	ı	'

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		2. Yes
		If chose "Yes", indicate the dose (mg)
	Selection:	
		2. 15 mg
Seq. # 1207		Apixaban Prescribed
Coo	ding instruction:	Indicate if the patient had apixaban prescribed
	Target Value :	At the time of initial visit
	Selection:	1. No
		2. Yes
		If chose "Yes", indicate the dose (mg)
	Selection:	1. 5 mg
		2. 10 mg
Seq. # 1208	Name:	Edoxaban Prescribed
Coc	ding instruction:	Indicate if the patient had edoxaban prescribed
	Target Value:	At the time of initial visit
	Selection:	1. No
		2. Yes
		If chose "Yes", indicate the dose (mg)
	Selection:	1. 30 mg
		2. 60 mg
Seq. # 1209	Name:	Initiation Date of Oral Anticoagulants
Coo	ding instruction:	Indicate the documented date of starting anticoagulants
	Target Value :	At the time of initial visit
Seq. # 1210	Name:	ACE inhibitors/ ARBs Prescribed
Coo	ding instruction:	Indicate if the patient had ACE inhibitors or ARBs prescribed
	Target Value :	At the time of initial visit
	Selection:	1. No
		2. Yes
Seq. # 1211	Name:	Beta Blockers Prescribed
Coo	ding instruction:	Indicate if the patient had beta blockers prescribed
	Target Value:	At the time of initial visit
	Selection:	1. No
		2. Yes
Seq. # 1212	Name:	Calcium Antagonist Prescribed
Coo	ding instruction:	Indicate if the patient had calcium antagonist prescribed
	Target Value:	At the time of initial visit
	Selection:	1. No
		2. Yes

Seq. # 1213	Name:	Digoxin Prescribed
1		Indicate if the patient had digoxin prescribed
	Target Value :	At the time of initial visit
	Selection :	
		2. Yes
Seq. # 1214	Name:	Diuretics Prescribed
	Coding instruction:	Indicate if the patient had diuretics prescribed
	Target Value :	At the time of initial visit
	Selection:	1. No
		2. Yes
Seq. # 1215	Name:	Statin Prescribed
	Coding instruction:	Indicate if the patient had statins prescribed
	Target Value :	At the time of initial visit
	Selection:	1. No
		2. Yes
Seq. # 1216	Name:	Proton Pump Inhibitor Prescribed
	Coding instruction:	Indicate if the patient had proton pump inhibiters prescribed
	9	At the time of initial visit
	Selection:	
		2. Yes
Seq. # 1217		NSAIDs Prescribed
	_	Indicate if the patient had NSAIDs prescribed
	_	At the time of initial visit
	Selection:	
G #1310	N T	2. Yes
Seq. # 1218	Name:	Antihyperuricemic Agents Prescribed
	_	Indicate if the patient had antihyperuricemic agents prescribed
	Selection :	At the time of initial visit
	Selection.	2. Yes
Seq. # 1219	Name:	
Seq. # 1217		Indicate if the patient had antiarrhythmic drugs prescribed
	_	At the time of initial visit
	Selection:	
	~00000011	2. Yes
Seq. # 1220	Name:	Vaughan-Williams Ia Drugs Prescribed
•		Indicate if the patient had Vaughan-Williams type Ia drugs prescribed
	_	At the time of initial visit
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	Selection :	1. procainamide
		2. quinidine
		3. sibenzolin
		4. disopyramide
Seq. # 1221	Name:	Vaughan-Williams Ib Drugs Prescribed
_	Coding instruction :	Indicate if the patient had Vaughan-Williams type Ib drugs prescribed
	Target Value :	At the time of initial visit
	Selection :	1. aprindine
		2. mexiletine
Seq. # 1222	Name:	Vaughan-Williams Ic Drugs Prescribed
	Coding instruction:	Indicate if the patient has Vaughan-Williams type Ic drugs prescribed
	Target Value :	At the time of initial visit
	Selection:	1. pilsicainide
		2. flecainide
		3. propafenone
Seq. # 1223	Name:	Vaughan-Williams III Drugs Prescribed
	Coding instruction:	Indicate if the patient had Vaughan-Williams type III drugs prescribed
	Target Value :	At the time of initial visit
	Selection:	1. sotalol
		2. amiodarone
Seq. # 1224	Name:	Vaughan-Williams IV Drugs Prescribed
	Coding instruction:	Indicate if the patient has Vaughan-Williams type IV drugs prescribed
	Tangat Value	At the time of initial visit
	rarget value:	
	Selection :	
	G	
Seq. # 1301	G	1. bepridil
Seq. # 1301	Selection:	1. bepridil 3. Past history about AF Type of AF
Seq. # 1301	Selection : Name :	1. bepridil 3. Past history about AF Type of AF Indicate the type of the patient's Atrial Fibrillation
Seq. # 1301	Selection : Name : Coding instruction :	1. bepridil 3. Past history about AF Type of AF Indicate the type of the patient's Atrial Fibrillation
Seq. # 1301	Selection : Name : Coding instruction : Target Value :	1. bepridil 3. Past history about AF Type of AF Indicate the type of the patient's Atrial Fibrillation At the time of initial visit
Seq. # 1301	Selection : Name : Coding instruction : Target Value :	1. bepridil 3. Past history about AF Type of AF Indicate the type of the patient's Atrial Fibrillation At the time of initial visit 1. Initial
Seq. # 1301	Selection : Name : Coding instruction : Target Value :	1. bepridil 3. Past history about AF Type of AF Indicate the type of the patient's Atrial Fibrillation At the time of initial visit 1. Initial 2. Paroxysmal
Seq. # 1301	Selection : Name : Coding instruction : Target Value :	1. bepridil 3. Past history about AF Type of AF Indicate the type of the patient's Atrial Fibrillation At the time of initial visit 1. Initial 2. Paroxysmal 3. Persistent
Seq. # 1301 Seq. # 1302	Selection : Name : Coding instruction : Target Value : Selection :	1. bepridil 3. Past history about AF Type of AF Indicate the type of the patient's Atrial Fibrillation At the time of initial visit 1. Initial 2. Paroxysmal 3. Persistent 4. Permanent
	Selection : Name : Coding instruction : Target Value : Selection :	1. bepridil 3. Past history about AF Type of AF Indicate the type of the patient's Atrial Fibrillation At the time of initial visit 1. Initial 2. Paroxysmal 3. Persistent 4. Permanent 5. Unknown
	Selection : Name : Coding instruction : Target Value : Selection :	1. bepridil 3. Past history about AF Type of AF Indicate the type of the patient's Atrial Fibrillation At the time of initial visit 1. Initial 2. Paroxysmal 3. Persistent 4. Permanent 5. Unknown Diagnosis Date of AF
	Selection : Name : Coding instruction : Target Value : Selection :	1. bepridil 3. Past history about AF Type of AF Indicate the type of the patient's Atrial Fibrillation At the time of initial visit 1. Initial 2. Paroxysmal 3. Persistent 4. Permanent 5. Unknown Diagnosis Date of AF Indicate the documented date of diagnosis of atrial fibrillation. If no

Seq. # 1303	Name :	Reversible Factors of AF
		Indicate if the patient's AF is due to a transient and/or reversible
	Coding instruction:	cause.
	Target Value :	At the time of initial visit
	Selection:	1. None
		2. Infectious diseases (ex. Pneumonia)
		3. Active hyperthyroidism
		4. Cardiac surgery within 3 months
		5. Pregnancy
		6. Other reversible cause
Seq. # 1304	Name:	AF management strategy - Rate Control/Rhythm Control
	Coding instruction:	Indicate the management strategy for the patient's Atrial Fibrillation
	Target Value:	At the time of initial visit
	Selection:	1. Rate control
		2. Rhythm control
		Rate control is the attempted control of ventricular rate with no
	Supporting Definition:	commitment to restore or maintain sinus rhythm. Rhythm control is
		the attempted restoration and/or maintenance of sinus rhythm.
Seq. # 1305	Name:	History of defibrillation
	Coding instruction:	Indicate if the patient has undergone defibrillation. If yes, indicate all
	Coung mstruction.	dates the patient received a defibrillation.
	Target Value:	Any occurrence between birth and the initial visit
	Selection:	1. No
		2. Non-pharmacological
		3. Pharmacological
		4. Both non-pharmacological and pharmacological
Seq. # 1306	Name:	History of Catheter Ablation
	Coding instruction:	Indicate if the patient has undergone catheter ablation
	Target Value :	Any occurrence between birth and the initial visit
	Selection:	1. No
		2. Yes
Seq. # 1307	Name:	History of Surgical Ablation
	Coding instruction:	Indicate if the patient has undergone surgical ablation (including
	coming mon action .	"maze" surgery)
	Target Value :	Any occurrence between birth and the initial visit
	Selection:	1. No
		2. Yes
Seq. # 1308	Name:	History of AV Node Ablation and/or Pacemaker Implantation

Coding instruction: Indicate if the patient has undergone AV node ablation or	pacemaker
implantation	
Target Value: Any occurrence between birth and the initial visit	
Selection: 1. No	
2. Yes	
eq. # 1309 Name: History of Left Atrial Appendage Closure	
Coding instruction: Indicate if the patient has undergone left atrial appendage	closure
Target Value: Any occurrence between birth and the initial visit	
Selection: 1. No	
2. Yes	
4. Past medical history	
eq. # 1401 Name: Heart Failure	
Coding instruction: Indicate if the patient has been diagnosed with heart failure	e (HF).
Target Value: Any occurrence between birth and the initial visit	
Selection: 1. No	
2. Yes (Hospitalized within 1 year)	
3. Yes (Not hospitalized)	
4. Yes (Details unknown)	
New York Heart Association Functional Classification for	or Heart
Name: Failure (if diagnosed with heart failure)	
Indicate the patient's NYHA class if he/she was diagnosed	with heart
Coding instruction: failure	
Target Value: At the time of initial visit	
Selection: 1. I	
2. II	
3. III	
4. IV	
Supporting Definition: (none)	
Name: Cardiomyopathy	
Coding instruction: Indicate if the patient has been diagnosed with cardiomyop	oathy
Coding instruction: Indicate if the patient has been diagnosed with cardiomyop Target Value: Any occurrence between birth and the initial visit	oathy
	oathy
Target Value: Any occurrence between birth and the initial visit	oathy
Target Value: Any occurrence between birth and the initial visit Selection: 1. No	oathy
Target Value: Any occurrence between birth and the initial visit Selection: 1. No 2. Yes (dilated cardiomyopathy)	oathy
Target Value: Any occurrence between birth and the initial visit Selection: 1. No 2. Yes (dilated cardiomyopathy) 3. Yes (hypertrophic cardiomyopathy)	oathy
Target Value: Any occurrence between birth and the initial visit Selection: 1. No 2. Yes (dilated cardiomyopathy) 3. Yes (hypertrophic cardiomyopathy) 4. Yes (ischemic cardiomyopathy)	

	Selection :	1 No
	Selection .	2. Yes
	Supporting Definition:	
Seq. # 1405	Name:	
, and the second	Coding instruction:	Indicate if the patient has been implanted cardiac devices.
	O	Any occurrence between birth and the initial visit
	Selection :	•
		2. Yes (permanent pacemaker)
		3. Yes (implantable cardiac defibrillator)
		4. Yes (cardiac resynchronization therapy)
Seq. # 1406	Name:	Coronary Artery Disease
	~	Indicate if the patient has been diagnosed with coronary artery
	Coding instruction:	disease.
	Target Value :	Any occurrence between birth and the initial visit
	Selection:	1. No
		2. Yes
	C	Documentation of CAD includes (but not limited to) angina pectoris,
	Supporting Definition:	ischemic heart disease, or coronary artery disease.
Seq. # 1407	Name:	Myocardial Infarction
	Coding instruction:	Indicate if the patient has been diagnosed with myocardial infarction.
	Target Value :	Any occurrence between birth and the initial visit
	Selection:	1. No
		2. Yes
Seq. # 1408	Name:	Percutaneous Coronary Intervention
	Coding instruction:	Indicate if the patient has undergone percutaneous coronary
	Coung mstruction.	intervention (PCI).
	Target Value :	Any occurrence between birth and the initial visit
	Selection:	1. No
		2. Yes
Seq. # 1409	Name:	Type of Stent
	Coding instruction:	Indicate the used devices during PCI
	Target Value :	Any occurrence between birth and the initial visit
	Selection:	1. Bare Metal Stent
		2. Drug-Eluting Stent
		3. Plain Old Balloon Angioplasty
		4. Unknown
Seq. # 1410	Name:	Coronary Artery Bypass Grafting

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	Coding instruction:	Indicate if the patient has undergone coronary artery bypass grafting
		(CABG).
	Target Value :	Any occurrence between birth and the initial visit
	Selection:	1. No
		2. Yes
Seq. # 1411	Name:	Valvular Heart Disease
	Coding instruction:	Indicate if the patient has been diagnosed with rheumatic mitral
	County instruction.	stenosis
	Target Value :	Any occurrence between birth and the initial visit
	Selection:	1. No
		2. Yes
Seq. # 1412	Name:	Valve Surgery or Percutaneous Valvuloplasty
	Coding instruction :	Indicate if the patient has undergone valve surgery or percutaneous valvuloplasty
	Target Value :	
	Selection :	•
		2. Yes (aortic valve)
		3. Yes (mitral valve)
		Documentation of percutaneous valvuloplasty includes (but not
		limited to) percutaneous transvenous mitral commissurotomy, balloon
	Supporting Definition:	aortic valvuloplasty, transcatheter aortic valve replacement, or mitral
		clip.
Seq. # 1413	Name:	Congenital heart disease
		Indicate if the patient has been diagnosed with congenital heart
	Coding instruction:	disease
	Target Value :	The value on registration date
	Selection:	1. No
		2. Yes
Seq. # 1414	Name:	Pericarditis or Myocarditis
	Coding instruction :	Indicate if the patient has been diagnosed with pericarditis or
	T 437.1	myocarditis
		Any occurrence between birth and the initial visit
	Selection:	
Com # 445	% .T	2. Yes
Seq. # 1415		Hypertension
		Indicate if the patient has been diagnosed with hypertension
		Any occurrence between birth and the initial visit
	Selection :	1. No

		2 V
		2. Yes
	Supporting Definition:	Hypertension is defined as documentation of SBP>=140mmHg,
0 111116	N T	DBP>=90mmHg or taking any antihypertensive drugs
Seq. # 1416	Name:	Dyslipidemia Liting is the state of the sta
	Coding instruction:	Indicate if the patient has been diagnosed with dyslipidemia
	Target Value :	•
	Selection:	1. No
		2. Yes
	Supporting Definition:	Dyslipidemia is defined as documentation of LDL>=140mg/dl,
		fasting TG>=150mg/dl or any cholesterol-lowering drugs
Seq. # 1417	Name:	
	_	Indicate if the patient has been diagnosed with diabetes mellitus
	Target Value :	Any occurrence between birth and the initial visit
	Selection:	1. No
		2. Yes (diet therapy)
		3. Yes (drug therapy)
		4. Yes (insulin therapy)
		Diabetes mellitus is defined as documentation of fasting BS
	Supporting Definition:	>=126mg/dl, random>=200mg/dl of blood glucose or HbA1c>=6.5,
		or taking any diabetes drugs
Seq. # 1418	Name:	Prior Stroke or TIA
	Coding instruction :	Indicate if the patient has been diagnosed with symptomatic stroke or
	Coung msu ucuon .	transit ischemic attack (TIA)
	Target Value :	Any occurrence between birth and the initial visit
	Selection:	1. No
		2. Yes (stroke)
		3. Yes (TIA)
	C	Stroke is defined as a new, sudden, focal neurologic deficit that
	Supporting Definition:	persists beyond 24 hours (vs. TIA: reversible within 24 hours).
Seq. # 1419	Name:	Cerebral Hemorrhage
	Coding instruction:	Indicate if the patient has been diagnosed with cerebral hemorrhage
	Target Value :	Any occurrence between birth and the initial visit
	Selection:	1. No
		2. Yes
Seq. # 1420	Name:	Gastrointestinal Bleeding
	~ .	Indicate if the patient has been diagnosed with class 3 or more
	Coding instruction:	gastrointestinal bleeding on BARC criteria
	Target Value :	Any occurrence between birth and the initial visit
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	Selection:	
		2. Yes (within 6 months)
		3. Yes (over 6 months ago)
Seq. # 1421	Name:	Peripheral Vascular Disease
	Coding instruction:	Indicate if the patient has been diagnosed with peripheral vascular disease
	Target Value :	Any occurrence between birth and the initial visit
	Selection:	1. No
		2. Yes
Seq. # 1422	Name:	Chronic Obstructive Pulmonary Disease
		Indicate if the patient has undergone chronic obstructive pulmonary
	Coding instruction:	disease (COPD).
	Target Value :	Any occurrence between birth and the initial visit
	Selection:	1. No
		2. Yes
Seq. # 1423	Name:	Sleep Apnea
	Coding instruction:	Indicate if the patient has been diagnosed with sleep apnea.
	Target Value :	Any occurrence between birth and the initial visit
	Selection:	1. No
		2. Yes (not using continuous positive airway pressure)
		3. Yes (using continuous positive airway pressure)
Seq. # 1424	Name:	Hyperthyroidism
	Coding instruction:	Indicate if the patient has been diagnosed with hyperthyroidism.
	Target Value :	Any occurrence between birth and the initial visit
	Selection:	1. No
		2. Yes
Seq. # 1425	Name:	Dialysis
	Cally and and and an a	Indicate if the patient is undergoing dialysis, including hemodialysis
	Coding instruction:	and peritoneal dialysis
	Target Value :	Any occurrence between birth and the initial visit
	Selection:	1. No
		2. Yes
Seq. # 1426	Name:	Malignant Tumor
	Coding instruction	Indicate if the patient has been diagnosed with malignant tumor that
	Coding instruction:	defines the patient's long-term prognosis
	Target Value :	Any occurrence between birth and the initial visit
	Selection:	1. No
		2. Yes
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Seq. # 1427	Name:	Smoking				
	Coding instruction :	Indicate if the patient is a smoker currently or quit within the past 12				
	Coung mstruction.	months.				
	Target Value:	Any occurrence between birth and the initial visit				
	Selection:	1. No				
		2. Yes				
Seq. # 1428	Name:	Drug Addiction				
	Coding instruction :	Indicate if the patient has been diagnosed and treated or hospitalized with drug addiction				
	Target Value :	Any occurrence between birth and the initial visit				
	Selection:	-				
		2. Yes				
Seq. # 1429						
1	Coding instruction:	Indicate if the patient drinks more than 160g of pure alcohol/week.				
		Any occurrence between birth and the initial visit				
	Selection :	•				
		2. Yes				
Seq. # 1430	Name:	Risk Assessment for Stroke on Medical Record				
_	Coding instruction:	Indicate if the patient's CHADS2 score is listed on the medical record.				
	Target Value :	At the time of initial visit				
	Selection:	1. No				
		2. Yes				
		5. Examination				
Seq. # 1501	Name:	Transthoracic Echocardiography				
	Coding instruction:	Indicate if the patient has undergone transthoracic echocardiography				
	Coung instruction:	within 6 months before and after the actual date of registration.				
	Target Value :	Any occurrence within 6 months before and after the date of registration.				
	Selection:	1. No				
	Selection.	2. Yes				
Seq. # 1502	Name:	Rhythm on transthoracic Echocardiography				
Seq. II 1202	Coding instruction :	Indicate the patient's heart rhythm on examination				
	Target Value :	The value within 6 months before and after the date of registration.				
	Selection:	Sinus rhythm				
		2. Atrial fibrillation				
		3. Others				
Seq. # 1503	Name:	Left Ventricular Ejection Fraction (LVEF) Percent				
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	Coding instruction:	Indicate the most recent documented left ventricular quantitative
		assessment
	Target Value :	The value within 6 months before and after the date of registration.
Seq. # 1504	Name:	Left Ventricular End Diastolic Diameter (LVEDD)
	Coding instruction:	Indicate the patient's left ventricular end diastolic diameter in
	_	centimeters.
	Target Value :	The value within 6 months before and after the date of registration.
Seq. # 1505	Name:	Left Ventricular End Systolic Diameter (LVESD)
	Coding instruction :	Indicate the patient's left ventricular end systolic diameter in
		centimeters.
	Target Value :	The value within 6 months before and after the date of registration.
Seq. # 1506	Name:	Left Atrium Diameter (LA diameter)
	Coding instruction:	Indicate the patient's left atrium diameter in centimeters.
	Target Value :	The value within 6 months before and after the date of registration.
Seq. # 1507	Name:	E-wave Velocity
	Coding instruction:	Indicate the patient's e-wave velocity in centimeters per second.
	Target Value :	The value within 6 months before and after the date of registration.
Seq. # 1508	Name:	A-wave Velocity
	Coding instruction:	Indicate the patient's a-wave velocity in centimeters per second.
	Target Value:	The value within 6 months before and after the date of registration.
Seq. # 1509	Name:	Deceleration Time of E-wave
	Coding instruction:	Indicate the patient's deceleration time of e-wave in milliseconds.
	Target Value:	The value within 6 months before and after the date of registration.
Seq. # 1510	Name:	Early Diastolic Filling Velocity
	Coding instruction :	Indicate the patient's early diastolic filling velocity (either IVS or LW
	Couning instruction.	basal) in centimeters per second.
	Target Value:	The value within 6 months before and after the date of registration.
Seq. # 1511	Name:	Aortic Stenosis
	Coding instruction:	Indicate if the patient has been diagnosed with aortic stenosis
	Target Value:	The value within 6 months before and after the date of registration.
	Selection:	1. Normal / trace
		2. Mild
		3. Moderate
		4. Severe
Seq. # 1512	Name:	Aortic Regurgitation
	Coding instruction:	Indicate if the patient has been diagnosed with aortic regurgitation
	Target Value :	The value within 6 months before and after the date of registration.
	Selection:	1. Normal / trace
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		2. Mild
		3. Moderate
		4. Severe
Seq. # 1513	Name:	Mitral Stenosis
	Coding instruction:	Indicate if the patient has been diagnosed with mitral stenosis
	Target Value :	The value within 6 months before and after the date of registration.
	Selection:	1. Normal / trace
		2. Mild
		3. Moderate
		4. Severe
Seq. # 1514	Name:	Mitral Regurgitation
	Coding instruction :	Indicate if the patient has been diagnosed with mitral regurgitation
	Target Value :	The value within 6 months before and after the date of registration.
	Selection:	1. Normal / trace
		2. Mild
		3. Moderate
		4. Severe
Seq. # 1515	Name:	Tricuspid Regurgitation
	Coding instruction :	Indicate if the patient has been diagnosed with tricuspid regurgitation
	Target Value :	The value within 6 months before and after the date of registration.
	Selection:	1. Normal / trace
		2. Mild
		3. Moderate
		4. Severe
Seq. # 1516	Name:	Transesophageal Echocardiography
	Coding instruction :	Indicate if the patient has undergone transesophageal
	ouring morraction.	echocardiography within 6 months.
	Target Value :	Any occurrence within 6 months before and after the date of
	imger value v	registration.
	Selection:	1. No
		2. Yes
Seq. # 1517	Name:	Rhythm on Transesophageal Echocardiography
	Coding instruction:	Indicate the patient's heart rhythm on examination
	_	The value within 6 months before and after the date of registration.
	Selection:	1. Sinus rhythm
		2. Atrial fibrillation
		3. Others
Seq. # 1518	Name:	Thrombus in Left Atrium or Left Atrial Appendage

		Indicate if the patient has thrombus in left atrium or left atrial
	Coding instruction:	appendage during TEE
		Any occurrence within 6 months before and after the date of
	Target Value :	registration.
	Selection:	1. No
		2. Yes
		3. Unknown
Seq. # 1519	Name:	Spontaneous Echo Contrast
	Coding instruction:	Indicate if the patient has spontaneous echo contrast.
	Target Value :	The value within 6 months before and after the date of registration.
	Selection:	1. No
		2. Yes
Seq. # 1520	Name:	Left Atrial Appendage Velocity
	Coding instruction	Indicate the patient's left atrial appendage velocity in centimeters per
	Coding instruction:	second.
	Target Value:	The value within 6 months before and after the date of registration.
Seq. # 1521	Name:	Blood Test
	Coding instruction:	Indicate if the patient has undergone blood tests within 3 months.
	Target Value :	Any occurrence within 3 months before and after the date of
	ranget value.	registration.
	Selection:	1. No
		2. Yes
Seq. # 1522	Name:	Hemoglobin
		Indicate the patient's hemoglobin (Hb) in grams per deciliter
	Target Value :	The value within 3 months before and after the date of registration.
Seq. # 1523	Name:	Blood Urea Nitrogen
	Coding instruction:	Indicate the patient's blood urea nitrogen in milligrams per deciliter.
	Target Value :	The value within 3 months before and after the date of registration.
Seq. # 1524	Name:	Creatinine
	Coding instruction:	Indicate the patient's creatinine in milligrams per deciliter.
	Target Value :	The value within 3 months before and after the date of registration.
Seq. # 1525	Name:	Total Bilirubin
	Coding instruction:	Indicate the most recent documented total bilirubin in milligrams per deciliter.
	Target Value :	The value within 3 months before and after the date of registration.
Seq. # 1526	Name:	Uric Acid
	Coding instruction :	Indicate the most recent documented uric acid in milligrams per
	-	deciliter.

	_	The value within 3 months before and after the date of registration.
Seq. # 1527	Name :	
	G	Indicate the most recent documented AST in Units per liter.
	Target Value :	The value within 3 months before and after the date of registration.
Seq. # 1528	Name:	ALT
	Coding instruction:	Indicate the most recent documented ALT in Units per liter.
	Target Value :	The value within 3 months before and after the date of registration.
Seq. # 1529	Name:	CRP
	Coding instruction:	Indicate the most recent documented CRP in milligrams per deciliter.
	Target Value :	The value within 3 months before and after the date of registration.
Seq. # 1530	Name:	PT-INR #1
	Coding instruction :	Indicate the patient's 1st PT-INR.
	Target Value :	The value within 3 months before and after the date of registration.
Seq. # 1531	Name :	PT-INR #2
	Coding instruction :	Indicate the patient's 2nd PT-INR.
	Target Value :	The value within 3 months before and after the date of registration.
Seq. # 1532	Name:	PT-INR #3
	Coding instruction :	Indicate the patient's 3rd PT-INR.
	Target Value :	The value within 3 months before and after the date of registration.
Seq. # 1533	Name :	PT-INR #4
	Coding instruction :	Indicate the patient's 4th PT-INR.
	Target Value :	The value within 3 months before and after the date of registration.
Seq. # 1534	Name:	PT-INR #5
	Coding instruction :	Indicate the patient's 5th PT-INR.
	Target Value :	The value within 3 months before and after the date of registration.
Seq. # 1535	Name:	APTT
	Coding instruction :	Indicate the most recent documented APTT in seconds.
	Target Value :	The value within 3 months before and after the date of registration.
Seq. # 1536	Name:	BNP
_	Coding instruction :	Indicate the most recent documented BNP measure in picograms per milliliter.
	Target Value :	The value within 3 months before and after the date of registration.
Seq. # 1537	Name :	D-dimer
		Indicate the most recent documented D-dimer measure in micrograms
	Coding instruction:	per milliliter
	Target Value :	The value within 3 months before and after the date of registration.

List of the name of committee which approved the KiCS-AF study

	Institution (Japanese)	Institution (English)	Name of committee (Japanese)	Name of committee (English)
1	慶應義塾大学医学部	Keio University, School of medicine	倫理委員会	Independent Ethics Committee
2	独立行政法人国立病院機構 東京医療センター	National Hospital Organization, Tokyo Medical Center	倫理委員会	Independent Ethics Committee
3	独立行政法人国立病院機構 埼玉病院	National Hospital Organization, Saitama Hospital	倫理委員会	Independent Ethics Committee
4	日野市立病院	Hino Municipal Hospital	倫理委員会	Independent Ethics Committee
5	さいたま市民病院	Saitama City Hospital	倫理委員会	Independent Ethics Committee
6	東京歯科大学市川総合病院	Tokyo Dental College, Ichikawa General Hospital	倫理審査委員会	Independent Ethics Committee
7	横浜市民病院	Yokohama Municipal Citizen's Hospital	倫理委員会	Independent Ethics Committee
8	立川病院	Tachikawa Hospital	倫理委員会	Independent Ethics Committee
9	北里大学北里研究所病院	Kitasato University, Kitasato Institute Hospital	研究倫理委員会	Independent Ethics Committee
10	済生会宇都宮病院	Saiseikai Utsunomiya Hospital	倫理委員会	Independent Ethics Committee
11	けいゆう病院	Keiyu Hospital	倫理委員会	Independent Ethics Committee

Supplemental Table 1. Definitions of the Cardiovascular death

Cardiovascular Death:	Death by any cardiovascular mechanism (arrhythmia, sudden death, heart		
Acute Myocardial	failure, stroke, pulmonary embolus, pulmonary artery disease) within 30 d after		
Infarction (MI)	an acute MI, related to the immediate consequences of the MI, such as		
	progressive HF or recalcitrant arrhythmia. There may be assessable		
	(attributable) mechanisms of cardiovascular death during this time period, but		
	for simplicity, if the cardiovascular death occurs within 30 d of an acute MI, it		
	will be considered a death due to MI.		
Cardiovascular Death:	eath: SCD was defined as unexpected and otherwise unexplained death in a		
Sudden Cardiac Death	h previously stable patient or death from documented or presumed cardiac		
(SCD)	arrhythmia without a clear non-cardiovascular cause, including patients who		
	were comatose and then died after attempted resuscitation. Patients who die		
	and had been out of contact for more than 24 hours were classified as		
	undetermined cause of death.		
Cardiovascular Death: Death associated with clinically worsening symptoms and/or sig			
Heart Failure (HF)	regardless of HF etiology		
Cardiovascular Death: Death after a stroke that is either a direct consequence of the str			
Stroke	complication of the stroke.		
Cardiovascular Death:	Cardiovascular death not included in the above categories but with specific,		
Other	known cause (e.g., pulmonary embolism, aortic dissection, aortic aneurysm or		
	peripheral arterial disease)		

Above definitions are accordance with the 2014 ACC/AHA Key Data Elements and Definitions for Cardiovascular Endpoint Events in Clinical Trials*

Hicks KA, Tcheng JE, Bozkurt B, Chaitman BR, Cutlip DE, Farb A, et al. 2014 ACC/AHA Key Data Elements and Definitions for Cardiovascular Endpoint Events in Clinical Trials: A Report of the American College of Cardiology/American Heart Association Task Force on Clinical Data Standards (Writing Committee to Develop Cardiovascular Endpoints Data Standards). *J Am Coll Cardiol* 2015; **66**: 403-469.

Supplemental Table 2.

The number of patients who refused to participate in the KiCS-AF registry at each participating center

	Number of patients who	Number of patients who participate	Refusal
Participating center	refused to participate	in the KiCS-AF registry	rate,%
Overall	42	1,322	3.1
Keio University Hospital	14	484	2.8
National Hospital Organization, Saitama Hospital	14	333	4.2
Hino Municipal Hospital	9	224	4.0
Tokyo Dental College, Ichikawa General Hospital	1	194	0.5
Saiseikai Utsunomiya Hospital	4	87	4.5

^{*}The number of patients who refused to participate was only available for limited number of the participating center