#### THE STRONG HEART STUDY VII CARDIOVASCULAR DISEASE IN AMERICAN INDIANS

### MORBIDITY SURVEY – DECISION

ID n					
Date	Date of this event:      month / day / year				
A.	DIA	GNOSIS (enter appropriate code number):		_	
	01.	Definite non-fatal myocardial infarction			
	1b.	Probable non-fatal myocardial infarction			
	02.	Possible non-fatal myocardial infarction			
	03.	Definite non-fatal stroke			
	04.	Possible non-fatal stroke			
	06.	Definite CHD			
	07.	Possible CHD (those with some, but not all, criteria or with for definite CHD)	equivocal criteria		
	08.	TIA			
	09.	Other CVD, specify:			
	10.	Non–CVD, specify:			
	11.	ESRD (dialysis or transplant):			
	12.	Heart Failure (Please fill out the HF PROCEDURE F	ORM)		
B.	Crite	ria used:			
1.	MYC	CARDIAL INFARCTION (Please check all applicable cri	teria)		
ı	A. Def	inite MI		<u>  </u>	
	1.	Evolving diagnostic ECG*, or		<u>  </u>	
	2.	Diagnostic biomarkers (2 x ULN)*			
	B. Pr	obable MI		<u>  </u>	
	1.	Positive ECG findings plus cardiac symptoms or signs with available biomarkers, or	nout	<u>  </u>	
The S		Positive ECG findings plus equivocal biomarkers eart Study VII - 12/30/2014 Page 1 of 6		 Morbidity Decision Forn	

C.	Po	ssible MI	
	1.	Equivocal biomarkers plus nonspecific ECG findings, or	
	2.	Equivocal biomarkers plus cardiac symptoms or signs, or	<u>  </u>
	3.	Missing biomarkers plus positive ECG	LI
· For F	CG a	and cardiac biomarker definition, please refer to: SHS VI Manual, Section 2.3.	
COM	/IEN	TS:	
2	ет	ROKE	
<b>4.</b>			
A.	Def	inite non-fatal stroke	ll
	1.	Stroke of unknown type etiology: Definite stroke of unknown etiology when CT or MRI not done. Information is inadequate to diagnose ischemic (infarction), intracerebral hemorrhage, or subarachnoid hemorrhage.	
	2.	Definite ischemic stroke: CT or MRI scan within 14 days of onset of a focal neurological deficit lasting more than 24 hours with evidence of brain infarction (mottled cerebral pattern or decreased density in a defined vascular territory), no intraparenchymal or subarachnoid hemorrhage by CT/MRI, (or lumbar puncture if done). A nonvascular etiology must be absent.	
	3.	Definite primary intracerebral hemorrhage: Focal neurological deficit lasting more than 24 hours. Confirmation of intraparenchymal hemorrhage in a compatible location, not caused by trauma, with CT/MRI scan within 14 days of stroke.	<u>  </u>
	4.	Subarachnoid hemorrhage: Sudden onset of a headache, neck stiffness, loss of consciousness. There may be a focal neurological deficit, but neck stiffness is more prominent. Blood in the subarachnoid or intraventricular space by CT/MRI - not caused by trauma.	
	5.	Non-fatal stroke after cardiovascular invasive interventions:  Stroke associated with the intervention within 30 days of cardiovascular surgery, or within 7 days of cardiac catheterization, arrhythmia ablation, angioplasty, atherectomy, stent deployment or other invasive coronary or peripheral vascular interventions.	
	6.	Non-fatal stroke post non-cardiovascular surgery: Stroke occurring within 30 days of non-cardiovascular surgery.	<u>  </u>
В.	Po	ssible non-fatal stroke	
	a.	History or rapid onset (approximately 48 hours from onset to time of admission or maximum acute neurologic deficit) of localizing neurologic deficit and/or change in state of consciousness, and	
	1b	. Documentation of localizing neurologic deficit by unequivocal physician or laboratory finding within 6 weeks of onset with 24 hours duration of objective physician findings, or	
	2a	. Discharge diagnosis with consistent primary or secondary codes (ICD-9-CM codes 431, 432, 434, 436, 437), and	<u>  </u>

	other disease pro	nequivocal physician or laboratory findings of any    cess or event causing focal brain deficit or coma other than or hemorrhage according to hospital records.
C.	Ischemic stroke subty	pe classification (complete for cases of definite ischemic stroke).
	[ ] 1.	Large-artery atherosclerosis: Clinical and brain imaging findings of either significant (>50%) stenosis or occlusion of a major brain artery or branch cortical artery, presumably due to atherosclerosis, and clinical findings of cerebral cortical impairment (aphasia, neglect, restricted motor involvement, etc.) or brain stem or cerebellar dysfunction. A history of intermittent claudication, transient ischemic attacks (TIAs) in the same vascular territory, a carotid bruit, or diminished pulses helps support the clinical diagnosis. Cortical or cerebellar lesions and brain stem or subcortical hemispheric infarcts greater than 1.5 cm in diameter on CT or MRI are considered to be of potential large-artery atherosclerotic origin. Supportive evidence by duplex imaging or arteriography of a stenosis of greater than 50% of an appropriate intracranial or extracranial artery is needed. Diagnostic studies should exclude potential sources of cardiogenic embolism. The diagnosis of stroke secondary to large- artery atherosclerosis cannot be made if duplex or arteriographic studies are normal or show only minimal changes.
		*Probable    *Possible
	[ ] 2.	Cardioembolism: Patients with arterial occlusions presumably due to an embolus arising in the heart. Cardiac sources are divided into high-risk and medium-risk groups based on the evidence of their relative propensities for embolism. At least one cardiac source for an embolus must be identified for a possible or probable diagnosis of cardioembolic stroke. Clinical and brain imaging findings are similar to those described for large-artery atherosclerosis. Evidence of a previous TIA or stroke in more than one vascular territory or systemic embolism supports a clinical diagnosis of cardiogenic stroke. Potential large-artery atherosclerotic sources of thrombosis or embolism should be eliminated. A stroke in a patient with a medium-risk cardiac source of embolism and no other cause of stroke is classified as a possible cardioembolic stroke.
		*Probable    *Possible
	[ ] 3.	Small-artery occlusion (lacune): Patients whose strokes are often labeled as lacunar infarcts in other classifications. The patient should have one of the traditional clinical lacunar syndromes and should not have evidence of cerebral cortical dysfunction (aphasia, neglect, restricted motor involvement, etc.). A history of diabetes mellitus or hypertension supports the clinical diagnosis. The patient should also have a normal CT/MRI examination or a relevant brain stem or subcortical hemispheric lesion with a diameter of less than 1.5 cm demonstrated. Potential cardiac sources for embolism should be absent, and evaluation of the large extracranial arteries should not demonstrate a stenosis of greater than 50% in an ipsilateral artery.
		*Probable    *Possible
		* A <b>probable</b> diagnosis is made if the clinical findings, neuroimaging data, and results of diagnostic studies are consistent with one subtype and other

etiologies have been excluded. A possible diagnosis is made when the

				clinical findings and neuroimaging data suggest a specif other studies are not done.	ic subtype but
		]	] 4.	Acute stroke of other determined etiology: Patients with stroke, such as non atherosclerotic vasculopathies, hyp or hematologic disorders. Patients in this group should or MRI findings of an acute ischemic stroke, regardless Diagnostic studies such as blood tests or arteriography these unusual causes of stroke. Cardiac sources of emlarge-artery atherosclerosis should be excluded by other	ercoagulable states, have clinical and CT of the size or location. should reveal one of bolism and
		[	] 5.	Stroke of undetermined etiology: In several instances, the cannot be determined with any degree of confidence. So no likely etiology determined despite an extensive evaluation was cursory. This cate patients with two or more potential causes of stroke so the unable to make a final diagnosis. For example, a patient cardiac source of embolism who also has another possil identified would be classified as having a stroke of underexamples would be a patient who has atrial fibrillation are of 50%, or the patient with a traditional lacunar syndromic carotid stenosis of 50%.	Some patients will have ation. In others, no egory also includes hat the physician is t with a medium-risk ple cause of stroke termined etiology. Other an ipsilateral stenosis
СО	MMEN	ITS: _			
		_			
		_			
3.	DEF	INITE	CORONA	RY HEART DISEASE (CHD)	
	a.	Card	iac cath pr	roven coronary artery disease (1 or more vessels ≥ 50% ste	nosis), <b>or</b>
	b.	PTC	A,		<u>  </u>
	C.	Coro	nary artery	y bypass grafting, <b>or</b>	<u>  </u>
	d1.	Abno	ormal stres	s ECG, and	<u>  </u>
	d.2.	Abno	ormal imag	ing, <b>or</b>	<u></u>
	e.	Posit	tive functio	nal test of ischemia (such as treadmill)	<u>  </u>
СО	MMEN	ITS: _			
		_			
		_			

### 4. HEART FAILURE (if yes, fill out Heart Failure form)

Two major criteria or one major and two minor criteria:

	a.	Major criteria  [ ] i. Paroxysmal nocturnal dyspnea or Orthopnea  [ ] ii. Neck vein distention  [ ] iii. Rales  [ ] iv. Cardiomegaly  [ ] v. Acute pulmonary edema  [ ] vi. S3 gallop  [ ] vii. Increased venous pressure >16cm water  [ ] viii. Circulation time ≥ 25 seconds  [ ] ix. Hepatojugular reflux
	b.	Minor criteria  [ ] i. Ankle edema  [ ] ii. Night cough  [ ] iii. Dyspnea on exertion  [ ] iv. Hepatomegaly  [ ] v. Pleural effusion  [ ] vi. Vital capacity reduced by one-third from maximum  [ ] vii. Tachycardia (rate of ≥ 120/min.)
	C.	Major or minor criteria [ ] i. Weight loss > 4.5kg in 5 days in response to treatment  AND
	d.	[ ] No known non-cardiac process leading to fluid overload such as renal failure
COMMENTS:  5. OTHER		ATAL CARDIOVASCULAR DISEASE
	<ul><li>a.</li><li>b.</li><li>c.</li><li>d.</li><li>e.</li><li>f.</li><li>g.</li><li>h.</li><li>i.</li><li>j.</li><li>k.</li><li>l.</li><li>m.</li></ul>	Purposely left blank – CHF moved to #4 above CHF secondary to ESRD (diagnosis = 10) Cardiomyopathy Valvular Heart Disease Left Ventricular Hypertrophy Atrial Fibrillation Non-coronary heart surgery or carotid or other vascular surgery (does not include procedures for PVD) Pacemaker implantation Positive non-coronary angiography (does not include procedures for PVD) Arrhythmia Angina pectoris (Class 2 chest pain, or relieved by nitroglycerides; diagnosis = 07) PVD (either peripheral arterial surgical procedures, angiogram or amputation) Aortic aneurysm

If there was coronary or peripheral vascular procedure done, fill out CVD Test Procedures form or Peripheral Vascular Procedure form.			
COMMENTS:			
ADMINISTRATIVE INFORMATION:			
Reviewer code:			
Review date:	month / day / year		

If you have any comments on this case, please use the space below:

#### THE STRONG HEART STUDY VII CARDIOVASCULAR DISEASE IN AMERICAN INDIANS

### **MORBIDITY SURVEY**

#### **Cardiovascular Test Procedures Abstract**

ID nur	mber:		<u> </u>	_	_  _
1.	WAS CATHETERIZATION/ANGIOGRAM DONE? Yes   1 No (Go to Q18)		Yes, but no re	eport   3	
2.	If YES, When?		_ /  month	_  /   day	 year
3.	Where:  Hospital/Clinic		City/Stat	e	
Was /	Any Vessel ≥ 50% Stenotic in	Yes	No	Uncertain	Unknown
4.	Left Main:	1	2	8	9
5.	Left anterior descending:	1	2	8	9
6.	Right coronary:	1	2	8	<u> </u> 9
7.	Circumflex artery:	1	2	8	<u> </u> 9
8.	Ejection Fraction (%):			<u> </u>	
	777= normal, % not specified 888=ab 999=unknown/no response	normal, %	not specified		
9.	Left Ventricular Function: Normal   1	A	ssessed, resu	lts not specifi	ed   3
	Depressed   2	N	ot assessed (	(Go to Q17)	9
10.	Was Akinetic Wall Observed?				
	Yes   1 No ( <b>Go to Q15)</b>   2	Uncert	ain   8	Unkn	own  9
		Yes	No	Uncertain	Unknown
11.	Anterior:	1	2	8	<u> </u>  9
12.	Inferior:	1	2	8	<u> </u> 9
13.	Apex:	1	2	8	<u> </u> 9
14.	Diffuse:	1	2	<u> </u>  8	<b></b>
The Stro	ong Heart Study VII Page 31 Cardiac Proce	dure Form		The District of Children The District of Children TRB APPROVED	IRB APPROVAL DATE: 04/13/20

The Strong Heart Study VII Version Date: 12/30/2014

)20

Findi	ing of Valvular Function:	Yes	No	Uncertain	Unknown
15.	Mitral regurgitation:	<u> </u>	<u> </u> 1   <u> </u> 2	8	9
16.	Aortic regurgitation:		<u> </u> 1   <u> </u> 2	8	<u> </u> 9
17.	Was Angioplasty performed?		<u> </u> 1   <u> </u> 2	8	<u> </u> 9
18.	WAS COMPUTED TOMOGRAPHIC	CALCIUM SCORIN	IG DONE?		
	Yes   1 N	o <b>(Go to Q22)</b>   2	2	Yes, but no re	port   3
19.	If YES, When?		_ / _ month	/   day	 year
20.	Where:		C:4x/S	· toto	
04	·		City/S	i i i	1 1
21.	Agotston score:	T DONES			
22.	WAS TREADMILL EXERCISE TES				
	Yes   1 N	o <b>(Go to Q29)</b>   2	2	Yes, but no re	port   3
23.	If YES, When?		_ / _ month	/   day	 year
24.	Where:  Hospital/Clinic		City/S	State	
25.	Treadmill ECG:				
	Normal   1 Borderline   2	Abnormal   3	Inconclusive	8 No rep	oort   9
26.	Maximum heart rate (beats/minute):		999=no report	: <u> </u>	
27.	Maximum systolic blood pressure (m	ımHg):	999=no report	: <u> </u>	
28.	Treadmill time (round to nearest who	ole number minute):	99=no report		
29.	WAS THALLIUM TEST, OR OTHER	R NUCLEAR IMAGE	TEST DONE?		
	Yes   1 N	o <b>(Go to Q34)</b>   2	2	Yes, but no rep	oort   3
30.	If YES, When?		_ / _ month	/   day	 year
31.	Where:		Oir /6	· toto	
32.	Hospital/Clinic  What Stress: Exercise   1 Aden	osine   2 Dobuta	City/S	her Drug 4  Place Rough Parkeyer Parkeyer  Res AppPROVED	IRB NUMBER: 10188 IRB APPROVAL DATE: 04/13/20

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33.	l est results:	Positive   1	Negative   2	Equivocal   3	No report   9
ADMI	NISTRATIVE IN	FORMATION:			
34.	Reviewer code	)			
35.	Review date:			/  month	/     day year

## THE STRONG HEART STUDY VII CARDIOVASCULAR DISEASE IN AMERICAN INDIANS

## MORBIDITY SURVEY PERIPHERAL VASCULAR PROCEDURES/REVASCULARIZATION ABSTRACT

ID nu	ımber: 	_  _	_				
1.	Was	periph	eral angiogram (ICD-	9 procedure o	code 88.48)	done?	
	`	Yes	_ 1 No   2 <b>(G</b>	o to Q2) Yes,	but no repor	t   9	
	a.	If yes	s: Contrast angiogram	MR :	angiogram	CT an	giogram
	b.	If yes	s, when?		L_	—ı——ı ı——ı—ı ı–	 day year
	C.	Whe	re:				
	d.	Was	any vessel ≥ 50% ster	notic?			
		i.	Aorta:	Yes   1	No   2	Uncertain  8	Unknown   9
			If yes, which side?	Right	Left	Both	
		ii.	Iliac:	Yes   1	No   2	Uncertain   8	Unknown   9
			If yes, which side?	Right	Left	Both	
		iii.	Femoral:	Yes   1	No   2	Uncertain   8	Unknown   9
			If yes, which side?	Right	Left	Both	
		iv.	Popliteal or lower:	Yes   1	No   2	Uncertain   8	Unknown   9
			If yes, which side?	Right	Left	Both	
		V.	Carotid stenosis	Yes   1	No   2	Uncertain   8	Unknown   9
			If yes, which side?	Right	Left	Both	
	e.	Was	there evidence of previ	ious revascula	rization? Y	es   1	No   2
2.	Was	peripho	eral angioplasty or su	rgical revasc	ularization d	one?	
			angioplasty   1 -9 procedure code 39.	.50)		scularization   3 cedure code 39.2	

No |\_\_\_|2 **(Go to Q3)** Yes, but no report |\_\_\_|9

	a.	If yes, when?         _ /  /  /  /           month       day       year
	b.	Where:
3.	Was	amputation (ICD-9 procedure codes 84.10 – 84.19) performed?
		Yes   1 No   2 <b>(Go to Q4.)</b> Yes, but no report   9
	a.	If yes, which side? Right    Left    Both
	b.	Which part?
		Upper body, Arm=1, Hand=2, Finger=3,
		Lower body, Above knee=1, Below knee=2    Foot=3, Toe(s)=4
	b.	When:
	C.	Where:
4.	Was	carotid angioplasty/stenting done?
		Yes   1 No   2 <b>(Go to Q5.)</b> Yes, but no report   9
	a.	If yes, which side? Right    Left    Both
	b.	If yes, when?   _ /  / _ /  month day year
	C.	Where:
5.	Was	carotid endarterectomy done?
		Yes   1 No   2 <b>(Go to end.)</b> Yes, but no report   9
	a.	If yes, which side? Right    Left    Both
	b.	When:   _ /  / _ _   month day year
	C.	Where:
<b>ADN</b> 5.		ATIVE INFORMATION:  ewer code:
6.	Revie	ew date:   _ / _ _

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IRB APPROVAL DATE: 04/13/2020

**Instructions**: The same procedures used for the ongoing surveillance in each center should be used, including evaluation of clinic charts and/or use of the IHS computerized records as well as direct contact with participants when necessary.

The purpose of this study is to derive an estimate of the proportion of participants who have undergone diagnostic or therapeutic procedures documenting definite lower extremity peripheral arterial disease since the Phase III SHS examination, and the proportion thereof for whom the necessary records are still available. Therefore, medical records for hospitalizations or outpatient encounters dealing with the diagnostic or procedural codes listed below and occurring since 1 January 1998 should be requested and reports of the procedures of interest should be obtained. Earlier events that correspond to the same procedures should be noted but charts need not be abstracted.

The following diagnostic codes should be identified:

For Peripheral Angiograms: ICD-9 procedure code **88.48** For Peripheral Angioplasty: ICD-9 procedure code **39.50** 

For Peripheral Surgical Revascularization: ICD-9 procedure codes 39.25 and 39.29

For Amputation: ICD-9 procedure codes **84.10-84.19** For Carotid Endarterectomy: ICD-9 procedure code **38.12** 

For Angioplasty: ICD-9 procedure code **00.61** For Stenting: ICD-9 procedure code **00.45** 



## THE STRONG HEART STUDY VII CARDIOVASCULAR DISEASE IN AMERICAN INDIANS

#### **HEART FAILURE PROCEDURES**

SH	S ID:   _
	ATRIAL FIBRILLATION AT TIME OF HF?         Yes     1         No     2         Unknown     9
B.	WHICH IMAGING STUDY WAS PERFORMED DURING THIS ADMISSION? Please check ALL that were done. If more than one imaging study was done in the same admission, please use one of these forms for EACH IMAGING STUDY to record the results of that study.
	1 Echocardiogram
	2 Nuclear Imaging
	3 Invasive Angiogram
	4 CT Angiogram
	5 MRI Angiogram
	6 Other, Specify:
	7 Not sure, no results found in chart
	8 None
lf r	ot sure or none, skip to Q8.
1.	Name of test:
2.	Date of test:   _ /  /  _     month day year
3.	Facility name:
	City/State:
4.	Ejection fraction: Measured:   % Estimated:   %
	If % not stated, 777 = normal, or range $\geq$ 50% 888 = abnormal, or range $<$ 50% 999 = unknown/no response
5.	Ejection fraction interpretation:  Normal     1 Depressed     2 NR     9
6.	Segmental wall motion abnormalities?         Yes    1         No    2         NR    9
	If yes, degree of abnormality: Mild    1 Moderate    2 Severe    3 Unknown    9
7.	Transmitral time: E Velocity:cm/sec A Velocity:cm/sec Peak E/A Ratio:

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					Н	FP
Decel. Time:	_msec	IVRT:	Septal E':	Peak S':	Septal A':	

SH	HS ID:   _ _ _		
8.	Valvular disease?	Yes    1 No    2 Unknown    9  If No or Unknown, go to Q9.	
	If Yes,	ii ito or omalomi, go to cor	
	a. Mitral regurgitation/insufficiency:  1+    1	4+    4 Unknown    9	
	b. Mitral stenosis: Mild    1 Mo	oderate    2 Severe    3 Unknown	9
	c. Aortic regurgitation/insufficiency:		
	1+   1 2+   2 3+   3	4+    4 Unknown    9	
	d. Aortic stenosis: Mild    1 Mo	oderate    2 Severe    3 Unknown	9
	e. Tricuspid regurgitation:  1+    1	4+    4 Unknown    9	
9.	Right ventricular systolic pressure/PA systolic pr If not stated, 777 = normal 888 = abnormal 999 = u		.]
C.	B-TYPE NATRIURETIC PEPTIDE (BT-BNP):	pg/ml. Upper Limit of Normal: pg/ml	
	N-TYPE NATRIURETIC PEPTIDE (NT-BNP):		
D.	CARDIOMYOPATHY DIAGNOSIS: Ischemic:	Non-Ischemic: Hypertrophic:	
	Valvular disea	ase: Acute MI: NR    9	
	No cardiomyo	opathy	
Re	eviewer Code:     Re	eview Date:  _ /  _ /  /   year	_

#### THE STRONG HEART STUDY VII

### CARDIOVASCULAR DISEASE IN AMERICAN

#### INDIANS MORTALITY SURVEY - FINAL DECISION

ID nur	mber:				
Date o	of death:	month / day / year		Age at o	death:
A.	Cause of	death, choose from the list	t below:		
	Cause of	death:			
	Contribu	tory cause of death 1:			
	Contribu	tory cause of death 2:			
	1a 02 03 04 05 06 07	I = Definite myocardial infar a = Probable myocardial infar 2 = Definite sudden death d B = Definite coronary heart of 4 = Possible coronary heart 5 = Definite stroke 6 = Possible stroke 7 = Definite congestive hear 8 = Possible congestive hear 9 = Other cardiovascular dis	arction ue to coronary disease disease rt failure art failure		
If is N	on-CVD de	eath, choose one from the	e following lis	t and complete the evidenc	e code:
				Evidence Code: (up to 3 Codes)	
22 = U 23 = U 24 = 0 25 = F 26 = E 27 = 0 28 = S 29 = H 30 = N 31 = E 32 = S 33 = H 88 = 0	Unintentional Chronic observations and allied Cheumonia Diabetes me Chronic live Suicide Homicide ar Nephritis, ne ESRD Septicemia HIV/AIDS Other, speci	ite: ————————————————————————————————————	s/all other hrosis	01 = Pathology Report 02 = Clinical Diagnosis only 03 = Pulmonary function te 04 = Blood glucose test 05 = Abnormal liver functio 06 = Abnormal kidney func 07 = Positive culture (blood 08 = Positive antibody test 09 = Positive blood test (ar 10 = Autopsy 11 = Police/Coroner's inves 12 = Other medical records Specify:	est In tests Ition test Id or sputum) Iny type) Istigation Is evidence
Was tl	he death al	cohol related?	Yes   1	No   2	Unknown   9

Criteria used for the cause of death: (Please check the appropriate boxes.)										
01.	Definite fatal myocardial infarction									
	[ ] 1(a)	Definite MI within 4 weeks of death by criteria: Yes No								
			2  2							
	OR	2. Diagnostic dicinations (2 x OLIV)	<b>-</b>							
	[ ] 1(b)	Acute MI diagnosed by autopsy								
	AND									
	[ ] 2.	No known non-atherosclerotic or noncardiac-atherosclerotic condition that was probably lethal according to death certificate, autopsy report, hospital records, or physician records.								
1a.	Probable fa	atal MI								
	[ ] 1.	Death within 28 days of hospital admission, cases defined as:								
		Yes No	)							
	1a.	Positive ECG findings plus cardiac symptoms or signs Without biomarkers, or   1	2							
	1b.	Positive ECG findings plus equivocal biomarkers   1	2							
		OR								
	[ ] 2.	Death within 6 hours of hospital admission with cardiac   1  _	2							
		symptoms and/or signs. Other confirmatory data								
		(biomarkers, ECG) are absent or non-diagnostic.								
* For	ECG and card	liac biomarker definitions, please refer to: SHS VI Manual, Section 2.3.								
02.	Definite suc	dden death due to CHD								
	[ ] 1.	Death witnessed as occurring within 1 hour after the onset of cardiac symptoms (prolonged cardiac pain, shortness of breath, fainting) or within hour after the subject was last seen without symptoms.	า 1							
	AND									
	[ ] 2.	No documentation of acute MI within 4 weeks prior to death.								
	AND									
	[ ] 3.	No known non-atherosclerotic or noncardiac-atherosclerotic process that was probably lethal according to death certificate, autopsy report, hospital records or physician report.								

B.

03.	Definite fatal CHD					
	[ [ [	] 1. ] 2. ] 3. ] 4.	Death certificate with consistent underlying or immediate causes, <b>AND</b> No documentation of definite acute MI within 4 weeks prior to death, <b>AND</b> Criteria for sudden death not met (above), <b>AND</b> No known non-atherosclerotic or noncardiac-atherosclerotic process or event that was probably lethal according to death certificate, autopsy report, hospital records, or physician records,			
	ΑN	ND				
	[	] 5(a)	Previous history of MI according to relative, physician, or hospital records, <b>OR</b>			
	[	] 5(b)	Autopsy reporting severe atherosclerotic-coronary artery disease or old MI without acute MI (50% proximal narrowing of two major vessels or 75% proximal narrowing of one more vessel, if anatomic details given.), <i>OR</i>			
	]	] 5(c)	Death occurring greater than 1 and less than or equal to 24 hours after the onset of severe cardiac symptoms or after subject was last seen without symptoms (without meeting criteria for Probable MI), <b>OR</b>			
	[	] 5(d)	Angiogram reporting severe (≥ 50% narrowing) atherosclerotic coronary artery disease, <i>OR</i>			
	[	] 5(e)	Other positive physical signs or lab findings.			
04.	Po	ssible fatal	CHD			
	[	] 1.	No documentation by criteria of definite acute MI within 4 weeks prior to death,  AND			
	[	] 2.	No documentation by criteria of definite sudden death,  AND			
	[	] 3.	No documentation by criteria of definite fatal CHD,  AND			
	[	] 4.	Death certificate with consistent underlying or immediate cause, <b>AND</b>			
	[	] 5.	No known non-atherosclerotic or noncardiac-atherosclerotic process that was probably lethal according to death certificate, autopsy report, hospital records, or physician records.			
05.	De	efinite fatal	stroke (also complete 6.1, 6.2 and Supplemental Form)			
	[	] 1a.	Cerebral infarction or hemorrhage diagnosed at autopsy,  AND			
	[	] 1b.	No other known disease process or event such as brain tumor, subdural hematoma, metabolic disorder or peripheral lesion that could cause focal neurologic deficit, with or without coma, according to death certificate, autopsy, hospital records, or physician records.			

OR

	[	] 2a.	History of rapid onset (approximately 48 hours from onset to time to admission or maximum acute neurologic deficit) of focal neurologic deficit with or without change in state of consciousness,  AND
	[	] 2b.	Focal neurologic deficit within 6 weeks of death documented by unequivocal physician or laboratory findings with 24 hours duration of objective physician findings,  AND
	[	] 2c.	No other known disease process or event such as brain tumor, subdural hematoma, metabolic disorder, or peripheral lesion that could cause focal neurologic deficit, with or without coma, according to death certificate, autopsy, hospital records, or physician records,
06.	Po	ossible (Und	documented) fatal stroke
	[	] 1.	Death certificate consistent with underlying or immediate cause (ICD-9, code 431 – 437), but neither autopsy evidence nor adequate pre-terminal documentation of the event, <i>AND</i>
	[	] 2.	No evidence at autopsy examination of the brain, if performed, of any disease process that could cause focal neurologic signs that would not be connected with cerebral infarction or hemorrhage.  OR
	[	] 3.	Focal neurological deficit and death within 24 hours, without MRI or other diagnostic image.
	<u>St</u>	roke subtyp	e classification (complete for cases of definite fatal stroke).
	[	] 1.	Stroke of unknown type etiology: Definite stroke of unknown etiology when CT or MRI not done. Information is inadequate to diagnose ischemic
	]	] 2.	(infarction), intracerebral hemorrhage, or subarachnoid hemorrhage.  Definite ischemic stroke: CT or MRI scan within 14 days of onset of a focal neurological deficit lasting more than 24 hours with evidence of brain infarction (mottled cerebral pattern or decreased density in a defined vascular territory), no intraparenchymal or subarachnoid hemorrhage by
	[	] 3.	CT/MRI. A nonvascular etiology must be absent.  Definite primary intracerebral hemorrhage: Focal neurological deficit lasting more than 24 hours. Confirmation of intraparenchymal hemorrhage in a compatible location, not caused by trauma, with CT/MRI scan within 14 days of stroke.
	[	] 4.	Subarachnoid hemorrhage: Sudden onset of a headache, neck stiffness, loss of consciousness. There may be a focal neurological deficit, but neck stiffness is more prominent. Blood in the subarachnoid or
	[	] 5.	intraventricular space by CT/MRI, not caused by trauma.  Non-fatal stroke after cardiovascular invasive interventions:  Stroke associated with the intervention within 30 days of cardiovascular surgery, or within 7 days of cardiac catheterization, arrhythmia ablation, angioplasty, atherectomy, stent deployment or other invasive coronary or
	[	] 6.	peripheral vascular interventions.  Non-fatal stroke post non-cardiovascular surgery: Stroke occurring within 30 days of non-cardiovascular surgery.

]	] 1.	Large-artery atherosclerosis: Clinical and brain imaging findings of either significant (>50%) stenosis or occlusion of a major brain artery or branch cortical artery, presumably due to atherosclerosis, and clinical findings of cerebral cortical impairment (aphasia, neglect, restricted motor involvement, etc.) or brain stem or cerebellar dysfunction. A history of intermittent claudication, transient ischemic attacks (TIAs) in the same vascular territory, a carotid bruit, or diminished pulses helps support the clinical diagnosis. Cortical or cerebellar lesions and brain stem or subcortical hemispheric infarcts greater than 1.5 cm in diameter on CT or MRI are considered to be of potential large-artery atherosclerotic origin. Supportive evidence by duplex imaging or arteriography of a stenosis of greater than 50% of an appropriate intracranial or extracranial artery is needed. Diagnostic studies should exclude potential sources of cardiogenic embolism. The diagnosis of stroke secondary to large- artery atherosclerosis cannot be made if duplex or arteriographic studies are normal or show only minimal changes.
		*Probable
[	] 2.	Cardioembolism: Patients with arterial occlusions presumably due to an embolus arising in the heart. Cardiac sources are divided into high-risk and medium-risk groups based on the evidence of their relative propensities for embolism. At least one cardiac source for an embolus must be identified for a possible or probable diagnosis of cardioembolic stroke. Clinical and brain imaging findings are similar to those described for large-artery atherosclerosis. Evidence of a previous TIA or stroke in more than one vascular territory or systemic embolism supports a clinical diagnosis of cardiogenic stroke. Potential large-artery atherosclerotic sources of thrombosis or embolism should be eliminated. A stroke in a patient with a medium-risk cardiac source of embolism and no other cause of stroke is classified as a possible cardioembolic stroke.
		*Probable    *Possible
[	] 3.	Small-artery occlusion (lacune): Patients whose strokes are often labeled as lacunar infarcts in other classifications. The patient should have one of the traditional clinical lacunar syndromes and should not have evidence of cerebral cortical dysfunction (aphasia, neglect, restricted motor involvement, etc.). A history of diabetes mellitus or hypertension supports the clinical diagnosis. The patient should also have a normal CT/MRI examination or a relevant brain stem or subcortical hemispheric lesion with a diameter of less than 1.5 cm demonstrated. Potential cardiac sources for embolism should be absent, and evaluation of the large extracranial arteries should not demonstrate a stenosis of greater than 50% in an ipsilateral artery.
		*Probable
		* A <b>probable</b> diagnosis is made if the clinical findings, neuroimaging data, and results of diagnostic studies are consistent with one subtype and other etiologies have been excluded. A <b>possible</b> diagnosis is made when the clinical findings and neuroimaging data suggest a specific subtype but other studies are not done.

Ischemic stroke subtype classification (complete for cases of definite ischemic stroke).

[	] 4.	Acute stroke of other determined etiology: Patients with rare causes of stroke, such as non atherosclerotic vasculopathies, hypercoagulable states, or hematologic disorders. Patients in this group should have clinical and CT or MRI findings of an acute ischemic stroke, regardless of the size or location. Diagnostic studies such as blood tests or arteriography should reveal one of these unusual causes of stroke. Cardiac sources of embolism and large-artery atherosclerosis should be excluded by other studies.
[	] 5.	Stroke of undetermined etiology: In several instances, the cause of a stroke cannot be determined with any degree of confidence. Some patients will have no likely etiology determined despite an extensive evaluation. In others, no cause is found but the evaluation was cursory. This category also includes patients with two or more potential causes of stroke so that the physician is unable to make a final diagnosis. For example, a patient with a medium-risk cardiac source of embolism who also has another possible cause of stroke identified would be classified as having a stroke of undetermined etiology. Other examples would be a patient who has atrial fibrillation and an ipsilateral stenosis of 50%, or the patient with a traditional lacunar syndrome and an ipsilateral carotid stenosis of 50%.
D	efinite fat	al congestive heart failure (Please fill out the HF PROCEDURE FORM)
<u>T</u> \	wo major	criteria or one major and two minor criteria:
a.	[ [ [ [	j i. Paroxysmal nocturnal dyspnea or Orthopnea ] ii. Neck vein distention ] iii. Rales ] iv. Cardiomegaly ] v. Acute pulmonary edema ] vi. S3 gallop ] vii. Increased venous pressure >16cm water ] viii. Circulation time ≥ 25 seconds ] ix. Hepatojugular reflux
b.	[ [ [ [	or criteria  i. Ankle edema  jii. Night cough  jiii. Dyspnea on exertion  jiv. Hepatomegaly  v. Pleural effusion  vi. Vital capacity reduced by one-third from maximum  vii. Tachycardia (rate of ≥ 120/min.)
c.		jor or minor criteria  1 i. Weight loss > 4.5kg in 5 days in response to treatment

07.

**AND** 

[ ]

d.

No known non-cardiac process leading to fluid overload such as renal failure

	08. Possible fatal congestive heart failure							
		[		ificate or medical records with neither autopsy evidence nor				
	09.	Oth	er fatal cardio	/ascular diseases				
		[	] i. Death cert	ificate or medical records with	consistent underly	ing (	or immediate	
	Cause. Check that applies.							
		[		th certificates are the only sou D 10: I00 to I09, I11, I13, I20 to				4
	ICD -	9	ICD - 10	Disease				
	390-39	92	100-102	Acute rheumatic fever		[	]	
	393-39	98	105-109	Chronic rheumatic heart dis	ease	[	]	
	402		l11	Hypertensive heart disease		[	]	
	404-40	05		Hypertensive disease		[	]	
	410-4	14	120-125	Ischemic heart disease		[	]	
	415-417 420-429 429.2 431-437			Diseases of pulmonary circ	ulation	[	]	
				Other forms of heart disease			]	
				Cardiovascular disease, unspecified		[	]	
				Cerebrovascular disease			]	
	799			III-defined or unknown		[	]	
			l13	Hypertensive heart and rena	al disease	[	]	
			127	Other pulmonary heart dise	ase	[	]	
			130-152	Other forms of heart diseas	е	[	]	
	443.9		173.9	Peripheral vascular disease	)	[	]	
Comm	nent:							
			INFORMATI					
Rev	viewer c	ode:						
Rev	riew dat	e:		-	month / day / year			
Revie	wer:		er Use Only					
Fii	rst revie	ew	1 Se	cond review   2 Stroke	e review   3	1	Adjudication   9	

If you have any comment on this case, please use the space below:					

## THE STRONG HEART STUDY VII CARDIOVASCULAR DISEASE IN AMERICAN INDIANS

# MORTALITY SURVEY INFORMANT INTERVIEW

ID r	number:		
A.	DECEDENT (Completed by study center	staff prior to interview.)	
1.	Name:		
	Last First		Middle
2.	Date of death:	<u> </u>  /  <u> </u> month day	/    year
B.	RECORD OF CALLS or HOME VISIT TO	COMPLETE INTERVIEW	
			Interview Completed
	DATE TIME (mo/day/yr) (24 hr clock)		1=Yes 2=No 9=Refused
	1)		
	2)		
C.	Person Providing Information (Complete	d by study center staff prior to	interview.)
3.	a. Name: Last First		 Middle
4.	Before we get started, could you please tell m	what was your relationship to the	ne deceased?
	You are the	of the decease	ed.
5.	What did the patient die from?		
_			
6.	Were you present when he/she died?		
	Yes   1 (Go to Q8)	No   2 Unknow	wn   9

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7.	If no, how long befor	e he/she died	did you last se	e him/her	?		
	1 hour or less 24 hours or le		1   2	More Unkn	than 24 hours own	3   9	
8.	Do you know of anyo	one else who r	may have been	n present a	at about the tim	e of his/her d	eath?
	,	Yes   1	No	<u> </u> 2	Unknown	_ <b> </b> 9	
	If yes can you give Contact information	1					
the forces the consorted health physical verbar Question persorted the consorted health physical persorted health physical person health p	ease describe the everiollowing conditions: colousness, sudden weath, health on the day lectan and will help to be atim and ask pertinerations: Are you aware on have the illness? We type and how long prior	chest pain, sakness, slurred he/she died, abetter understant questions of any illnesse vas the individ	shortness of d speech, etc. and of the dea and the cause when appropries the individu	breath, a Please t ath itself. of your le riate attac al had pri	agitation, sudd ell me what yo This informatio oved one's dea or to death?	len collapse u know of his on will be re- ath. <i>(Recor</i> sheet if need If yes – how	or loss of her general viewed by a standard summary led) Probing long did the

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befor him/h <i>alrea</i> <i>answ</i>	next set of questions deal specifically with the last episode of pain or discomfort that occurred the his/her death. This is defined as starting at the time you noticed discomfort that caused her to stop or change what he/she was doing. NOTE TO INTERVIEWERS: If the informant has dy answered these questions in the description of circumstances, just fill out the correct ter(s) as noted below. Respect the informant's wishes about continuing the interview and answers to as many of the following questions as possible.
10.	Did his/her last episode of pain or discomfort specifically involve the chest?  Yes   1 No   2 Unknown   9
11.	Did he/she experience pain or discomfort in his/her chest, left arm or shoulder or jaw either just before death or within 3 days (72 hours) of death?  Yes   1
12.	Did he/she take nitroglycerine because of this last episode of pain or discomfort?  Yes    1 No    2 Unknown    9
13.	Did he/she take any other medicine for chest discomfort prior to death? Yes No If yes what?
14. 15.	How long was it from the beginning of his/her last episode of pain or discomfort to the time he/she stopped breathing on his/her own? (use the shortest interval known to be true)  5 minutes or less   1
10.	
	a. If yes, what year did he/she start dialysis?
	b. How many times per week did he/she receive dialysis?
	c. Did he/she stop dialysis before death? Yes No Unknown
	If yes, how long before death?      /  /  /    days months years
16.	Within 3 days of death, or just before he/she died, did any of the following symptoms begin for the first time or did the patient complain of any of these symptoms:
	Yes No Unknown

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	a.	Shortness of breath?		1
	b.	Dizziness?		1   2   9
	C.	Palpitations (pounding in the		1   2   9
	d.	Marked or increased fatigue,	tiredness, or weakness?	1   2   9
	e.	Headache?		1   2   9
	f.	Sweating?		1   2   9
	g.	Paralysis?		1   2   9
	h. :	Loss of speech?	ation or abdominal discountant	1   2   9
	i. i		stion or abdominal discomfort?	· · · · · · · · · · · · · · · · · · ·
	J. k.	nausea or vomiting? Other? specify:		1
	Ν.	Other? specify.		1   2   9
		· · · · · · · · · · · · · · · · · · ·	ons are about his/her medica	
		<u>Please provide</u>	as much information as pos	<u>sible</u>
17.		e his/her final illness, had he/s	ne ever had pains in the chest	from heart disease, for example,
	arigine	Yes   1	No   2( <b>If no, go to Q20?</b> )	) Unknown   9
18.	Did he	e/she ever take nitroglycerin fo	r this pain?	
	D.G.110	Yes    1	No    2	Unknown    9
		<u> </u>	<u> </u>	<u>,                                     </u>
19.	Any o	ther medications such as aspi Yes   1	rin, tums or other antacids?  No   2	Unknown   9
00	D:			
20.	illness		ollowing medical condition or	procedures before his/her final
				Yes No Unknown
	a.	heart attack?		
	b.	stroke?		
	C.	heart failure?	part condition	1    2    9
	d.	any other heart disease or he lf yes, specify:	eart condition	1   2   9
	e.	coronary bypass surgery (CA	ABBAGE)	1    2
	<u> </u>  9	, ,, , , , , , , , , , , , , , , , , ,	,	,,
	f.	coronary angioplasty (balloo	• . • ,	1   2   9
	g.	insertion of pace maker (defi	brillator)	1   2   9
	h.	any other heart surgery?		1   2   9
		The west few assetions are	abaut bia/bay baalth in the v	
		The next lew questions are	about his/her health in the <u>y</u>	eai prior to death
21.	Was h	ne/she hospitalized or taken to	a clinic	Yes No Unknown
		year prior to death?		1   2   9
		month prior to death?		1 2 9
		7 days prior to death?		1   2   9
22	\\/a==	any haanitalizations for hazart	attack or chest pain?	
22.	vvere	any hospitalizations for heart a Yes    1 No	attack or cnest pain?  2 Unknown    9	
		169   1 140  _	2 OHKHOWH   9	

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23.	Was a hospitalization for heart surgery?	Yes   1 No   2 Unknown   9
24.	What was the date of the <u>last</u> hospital admission? (If unknown, draw two lines across the boxes)  If the information in questions 25	month day year 5- 28 is already known to you, skip to Q29.

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25.	Can you tell me the name and location of the hospital? <i>(If unknown, check the box.)</i> a. Name:			
	b. Address:			
	City/town:			
	State-Zip:			
26.	Was he/she seen by a physician anytime in the year prior to death?  Yes   1 No   2 Unknown  9			
27.	Can you tell me the name and address of this physician or healthcare facility?     IHS only			
	a. Name:			
	b. Address:			
	City/town:			
	State-Zip: ————————————————————————————————————			
28.	Can you tell me the name and address of his/her usual physician?  If same as Q27, check here.			
	a. Name:			
	b. Address:			
	City/town:			
	State-Zip:			
29.	Now, think back to about <u>one month</u> before he/she died. At that time, was he/she sick or ill; were his/her activities limited, or was he/she normally active for the most part?			
	Sick/ill/limited activities   1 Normally active   2 Unknown   9			
30.	Was he/she being cared for at a nursing home or at another place at the time of death?  Yes, nursing home, specify   1			
	next few questions are concerned specifically with emergency medical care he/she may have ved just prior to or at the time of death.			
31.	Was he/she taken to a hospital/clinic in the week before his/her death? Yes   1   2 No			

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32.	If Yes, could you tell me the name and location of this facility:			
	a.	Name:		
	b.	Address:		
		City/town:		
		State-Zip:		
33.		re someone else whom we could contact, who might know more about the circumstances nding his/her death or his/her usual state of health?		
		Yes   1 No   2 Unknown   9 (If Yes, complete the front of the second Informant Interview)		
34.	Did info	ormant provide consent to gather further information?  Yes   1 No   2 Not applicable   3  (If Yes, ask the informant to sign the consent form for us to review the decedent's medical records)		
35.	How re	eliable was the participant in completing the questionnaire?		
Very r	eliable	1 Reliable   2 Unreliable   3 Very unreliable   4 Uncertain   5		
<b>ADMII</b> 36.	_	TIVE INFORMATION: ewer code:		
37.		ew date:		
		month day year		

## THE STRONG HEART STUDY VII CARDIOVASCULAR DISEASE IN AMERICAN INDIANS

# **SUPPLEMENTAL STROKE FORM - Mortality and Morbidity Surveys** (Complete for mortality codes 5 or 6 and morbidity codes 3, 4 or 8)

ID number:			
Date of this event:		/  / _ Month day	 year
A.	ISCHEMIC STROKE LOCATION		YES NO
1.	Right hemisphere		1   <u></u>  2
2.	Left hemisphere		1   2
3.	Basilar		1 <u></u>  2
4.	Hemispheric and Basilar		1   2
5.	Unknown		1   2
B.	BRAIN IMAGING		
6.	HEAD CT	Yes	<u> </u>  1
		No (go to Q 7)	2
		Yes, but no report	3
	6.1 If yes, timing of Head CT	<48 h since symptom onset	1
		≥48 h since symptom onset	2
		Unknown	3
7.	BRAIN MRI	Yes	1
		No (go to Q 8)	   2
		Yes, but no report	<u> </u> 3
C.	NEUROVASCULAR IMAGING		
8.	CAROTID DUPLEX	Yes	<u> </u>  1
		No (go to Q 9)	2
		Yes, but no report	IRB NUMBER: 1/3/188

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IRB NUMBER: 1888
IRB APPROVAL DATE: 04/13/2020

9.	TRANSCRANIAL DOPPLER (TCD)	Yes	<u> </u>  1
		No, (go to Q 10)	2
		Yes, but no report	3
10.	MAGNETIC RESONANCE ANGIOGR	APHY (MRA) Yes	<u> </u> 1
		No (go to Q 11)	2
		Yes, but no report	3
11.	CT ANGIOGRAPHY	Yes	<u> </u>  1
		No (go to Q 12)	2
		Yes, but no report	3
12.	ANGIOGRAPHY	Yes	<u> </u> 1
		No, (go to Q 13)	2
		Yes, but no report	3
D.	STROKE DEFICIT		
13.	MODIFIED RANKIN SCALE (Code Maximal Severity Within 7 Days		(0-6)
	<ul> <li>1 = no significant disability despite syn</li> <li>2 = slight disability: unable to carry out without assistance</li> <li>3 = moderate disability: requiring some</li> <li>4 = moderately severe disability: unable bodily needs without assistance</li> <li>5 = severe disability: bedridden, incontent</li> <li>6 = death</li> <li>9 = information insufficient for coding</li> </ul>	all previous activities but able to look help, but able to walk without assistence, and ur	ok after own affairs stance nable to attend to own
E.	STROKE TREATMENT		
14.	Intravenous thrombolysis	Yes	<u>       </u> 1
		No	2
15.	Presentation within 3 hours from symp	tom onset Yes	<u> </u> 1
		No	2
F.	BRAIN EXAMINATION AT AUTOPSY	Yes	1
		No	RB NUMBER 10188
The St	rong Heart Study VII Page 5	3Supplemental Stroke	IRB APPROVAL DATE: 04/13/2020

	Yes, but no report	<u> </u> 3
ADMINISTRATIVE INFORMATION: Reviewer code:		
Review date:	_ /  / _ Month day	_    year

If you have any comments on this case, please use the space below: