



MarkVCID Follow-up Paper Case Report Form Package

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By the MarkVCID Clinical Data, Physiological Data & Cognitive Assessments Subcommittee (Deborah Blacker, MD, ScD, Chair) and Coordinating Center (PI Steven Greenberg, MD, PhD).

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MarkVCID Follow-up Paper CRF Package				
Patient ID:		Patient Initials:		
Visit Date:///	 -	Evaluator Initials:		
Study Visit:				
DEMOCRAPHICS AN	ND DEI ATED EI	EMENTS: FOLLOW-UP		
<u>DEMOGRAFIIICS AI</u>	NO RELATED EL	EMEN 13: FOLLOW-UF		
Date of Collection: /	_/(MM/DD/YYYY)		
1. Sex:				
2. Subject's current marital status	S:			
☐ Married	Never marrie	ed (or marriage was annulled)		
☐Widowed	Living as mar	ried/domestic partner		
Divorced	Unknown			
Separated				
3. What is the subject's living situ	ıation?			
Lives alone				
Lives with one other person	n: a spouse or part	ner		
Lives with one other person	n: a relative, friend	, or roommate		
Lives with caregiver who is	not spouse/partn	er, relative, or friend		
Lives with a group (related	or not related) in	a private residence		
Lives in group home (e.g., a	ssisted living, nurs	sing home, convent)		
Unknown				

MarkVCID Follow-up Paper CRF Package

Patient Initials: _____

Pa	atient ID:	Patient Initials:
Vi	isit Date://	Evaluator Initials:
St	udy Visit:	
	4. What is the subject's level of independence?	
	Able to live independently Requires some assistance with complex activ	ities
	Requires some assistance with basic activitie	
	Completely dependent	
	Unknown	
	5. ZIP Code (first three digits) of subject's primary	residence: Unknown

ati	ent ID:		Pat	tient Initials: _	
	t Date: / / /		Eva	aluator Initials	<u> </u>
tuc	ly Visit:				
	MEDICAL/NEUROLOGICA	AL/PSYCH	IAT	RIC: FOLLOW	-UP
	Date of Collection: / (MM/DD/YYYY)				
Date of Last Study Visit: / (MM/DD/YYYY)			YYY)		
	(To be used to ask patients about medical history since last study visit)			sit)	
	CIGARETTE SMOKING				
		No		Yes	Unknown
	Has the subject smoked since last study visit?				
	If Yes:		•		
	1a. Average number of packs smok 1 cigarette to less than ½ 2 pack to less than 1 pac	pack	sinc	e last study vis	sit:
	\square 1 pack to less than $1½$ pa	acks			
	☐ 1½ packs to less than 2 p	oacks			
	2 packs or more				
	Unknown				
	1b. If the subject has quit smoking s	since last s	tudy	v visit, specify th	ne age at
	which he/she last smoked (i.e., quit	t):	[8-1	110] 🗌 N/A 📋	Unknown

Patient ID:			Patient Initials	:			
/isit Date: / //			Evaluator Initia	als:			
tudy Visit:							
CARDIOVASCULAR DISEAS	CARDIOVASCULAR DISEASE						
Since last study visit, has the No Yes	patient been diagnosed	l wit	h any new cardiova	ascular diseases?			
If yes:							
New Cardiovascular Disease diagnosed since most recent study visit	No		Yes	Not Assessed			
Heart attack/cardiac arrest							
If yes:							
1a. More than one he	eart attack?						
□ No	Yes Uni	kno	wn				
1b. Age at most rece	nt heart attack:		Unknown				
2. Atrial fibrillation							
3. Angioplasty/ endarterectomy/ stent							
4. Cardiac bypass procedure							
Pacemaker and/or defibrillator	5. Pacemaker and/or						
6. Congestive heart failure							
7. Angina							
8. Heart valve replacement or repair							

Patient ID:		F	Patient Initials:		
Visit Date:// Evaluator Initials:					
Study Visit:					
	New Cardiovascular Disease diagnosed since most recent study visit	No	Yes	Not Assessed	
	9. Other cardiovascular disease (specify): (enter 'N/A' if absent)				
	10. Other cardiovascular disease (specify): (enter 'N/A' if absent)				
	11. Other cardiovascular disease (specify): (enter 'N/A' if absent)				

ti	tient ID:			Patient Initials:		
si	t Date:	//		Evaluator Initials:		
uc	ly Visit:					
	CEREBROVAS					
	Since last stud Vascular Event	y visit, has the patient?	been diagnosed Yes	with a Sympto	matic Stroke/Acute	
	New Cerebro	vascular Events dia	gnosed since r	nost recent s	study visit:	
	Event	Age at Event	Type of Sympto Stroke/Acute V Event		Temporally associated with persistent worsening of cognition?	
	Stroke/Acute Vascular Event 1	—— □ Unknown	☐ Ischemic ☐ Hemorrhagic ☐ Stroke type unknown ☐ TIA with clear ischemic mechanism		☐ No ☐ Yes ☐ Unknown	
	Stroke/Acute Vascular Event 2	—— □ Unknown	Ischemic Hemorrhagi Stroke type TIA with cle	unknown ar ischemic	☐ No ☐ Yes ☐ Unknown	
	Stroke/Acute Vascular Event 3	—— □ Unknown	☐ Ischemic ☐ Hemorrhagic ☐ Stroke type unknown ☐ TIA with clear ischemic mechanism ☐ Ischemic ☐ Hemorrhagic ☐ Stroke type unknown ☐ TIA with clear ischemic mechanism		☐ No ☐ Yes ☐ Unknown	
	Stroke/Acute Vascular Event 4	—— □ Unknown			☐ No ☐ Yes ☐ Unknown	
	Stroke/Acute Vascular Event 5	—— □ Unknown	Ischemic Hemorrhagi Stroke type TIA with cle	unknown ar ischemic	☐ No ☐ Yes ☐ Unknown	

		<u> </u>				
ti	ent ID:		Patient Initial	Patient Initials:		
sit	t Date://	Evaluator Init	Evaluator Initials:			
	ly Visit:					
	NEUROLOGIC CONDITIONS					
	Since last study visit, has the p	oatient been diagnose	d with any new net	irologic conditions?		
	New Neurologic Conditions	diagnosed since r	nost recent stud	y visit:		
	Condition	No	Yes	Not Assessed		
	1. Seizures					
	2. Traumatic brain injury (TBI)					
	If TBI yes:	1		1		
	☐ No ☐ Singl ☐ Repe ☐ Unkr	ated/multiple		2)		
	☐ No ☐ Singl ☐ Repe	e ated/multiple	usness (2 3 mmute:	5)		
	Unkr		(as might result from	om military		
	2c. TBI without loss of consciousness (as might result from military detonations or sports injuries)?					
	☐ Singl	e				
	Repe	ated/multiple				
	☐ Unkr	iown				
	2d. Age at most	recent TBI:	Unknown			

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Patient II	atient ID: Patient Initials:				
Visit Date	e://	_ E	valuator Initials	:	
Study Vis	sit:				
MEI	DICAL CONDITIONS				
	Since last study visit, has the patient been diagnosed with any new medical conditions? No Yes				
New	New Medical Conditions diagnosed since most recent study visit:				
	Condition	No	Yes	Not Assessed	
1. Г	Diabetes				
	1a. If yes, which type?Type 1Other type (diabetes insipidus, latent autoimmune diabetes/type 1.5, gestational diabetes)	☐ Type 2 ☐ Unknov			
2. H	Hypertension				
2	2a. If yes, is hypertension treated?	No 🗆	Yes		
3. I	Hypercholesterolemia				
4. E	312 deficiency				
5. 1	Γhyroid disease				
6. <i>A</i>	Arthritis				
	If yes:				
6	6a. Type of arthritis: Rheumatoid Other (specify): Unknown				
		apply): Spine Unknown			
7. I	ncontinence – urinary				
8. I	ncontinence – bowel				
9. S	Sleep apnea				
10.	REM sleep behavior disorder (RBD)				
11.	Hyposomnia/insomnia				

Pati	ent ID:		Patient Initials:		
Visi	t Date:///	_	Evaluator Initi	als:	
Study Visit:					
	PSYCHIATRIC CONDITIONS				
	Since last study visit, has the patient bee conditions? \(\subseteq \text{No} \subseteq \text{Yes} \)	n diagnose	ed with any new psyd	chiatric	
	New Psychiatric Conditions diagnos	ed since	most recent study	visit:	
	Psychiatric Condition	No	Yes	Not Assessed	
	1. Obsessive-compulsive disorder (OCD)				
	2. Developmental neuropsychiatric disorders (e.g., autism spectrum disorder [ASD], attention-deficit hyperactivity disorder [ADHD], dyslexia)				

ati	ent ID:		Patient Initials:		
	t Date://	_	Eva	aluator Initials	<u> </u>
uc	ıdy Visit:				
	FAMILY HIST	ΓORY: FOL	LO'	W-UP	
	Since the last visit, is any new informath history?	tion availabl	e co	oncerning the pa	tient's family
Corrections or new information on previously reported family history: If any previously recorded family history information has been found to be incorrect, corrections to the pertaining data should be made to that previous Family History form. Any newly obtained information (e.g., new mutation information, new reported cases of stroke/TIA or acquired cognitive impairment, new report of autopsy confirmation of diagnoses) should be indicated on this form and should no be submitted as a correction to a previously submitted Family History form.			ncorrect, mily History n, new eport of nd should not		
	Date of Collection: / /		(M	IM/DD/YYYY)	
	FAMILY HISTORY	No		Yes	Unknown
	STROKE/TIA: Is there a family history in a first degree relative of symptomatic stroke or TIA with clear ischemic mechanism?				
	If yes:		1		
	1a. Any cases with onset before age 55?				
	1b. Is there a pattern suggestive of an autosomal dominant family history?				
	2. ACQUIRED COGNITIVE IMPAIRMENT: Is there a family history in a first degree relative of cognitive impairment or dementia or Alzheimer's disease?				

Patio	ent ID:	·	_	Pa	tient Initials:	
		//		Ev	aluator Initials	:
Stud	y Visit	:				
			No		Yes	Unknown
		If yes:				
		2a. Any report of a case in the family with autopsy confirmation of Alzheimer's disease?				
		2b. Any report of cases with autopsy confirmation of another cause of dementia?				
		2c. Any cases with onset before age 65?				
		2d. Is there a pattern suggestive of an autosomal dominant family history?				
<u> </u>	3.	If yes to EITHER autosomal domin following:	ant question	ıs ab	oove (1b, 2d), com	plete the
		3a. Is there a known mutation?				
-	3b. If yes, please indicate which one: PSEN1					
		☐ APP				
		☐ PSEN2				
		☐ CADASIL				
		Other, specify gene if kr	nown:			
		Specify mutation if known:				
		3c. Does this individual carry the n	nutation?			
		\square No \square Yes \square	Unknown			

Pati	ent ID:		Patien	t Initials:	
/isit	t Date://		Evalua	tor Initials: _	
	ly Visit:				
	GENERAL PHYSI	CAL M	EASUR)	ES	
	Were General Physical Measures performe	d?			
	□ No □ Yes				
	If No, please provide the primary re	ason:			
	Physical problem	U Ver	bal refus	al	
	Cognitive/behavior problem	Oth	er probl	em (specify): _	
	Date of Collection: / /	(M	M/DD/Y	YYYY)	
	VITAL SIGNS				
	Measure seated at rest. Take 3 consecutive BF	reading	gs. Avera	ge will be calcı	ılated in EDC.
	1. Blood Pressure Measurement 1:	/_	r	nmHg [Not Done
	Blood Pressure Measurement 2:	/_	r	nmHg [Not Done
	Blood Pressure Measurement 3:	/_	r	nmHg [Not Done
	2. Pulse: beats	s/minu	te	[Not Done
	3. Height: cm in				Not Done
	4. Weight: kg				Not Done
	ADDITIONAL PHYSICAL OBSERVATIONS	5	No	Yes	Unknown
	With or without corrective lenses, is the subject's vision functionally normal?				
	2. With or without a hearing aid(s), is the subject's hearing functionally normal?				

Patient ID:	Patient Initials:
Visit Date://	Evaluator Initials:
Study Visit:	

SHORT PHYSICAL PERFORMANCE BATTERY					
KEY: If the subject cannot complete any of the following exams, please give the reason by entering one of the following codes:					
95 = Physical problem 97 = Other problem	96 = Cognitive/behavior problem 98 = Verbal refusal				
1. Balance Test Score: Side-by-side, semi-tandem, tandem:	[0-4, 95-98]				
2. Gait Speed Test Score:	[0-4, 95-98]				
3. Chair Stand Test Score:	[0-4, 95-98]				

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Patient ID: ______ Patient Initials: ______

Visit Date: ___/___/ Evaluator Initials: ______

Study Visit:

<u>NEUROLOG</u>	SICAL EXAM
Was the Neurological Exam performed? No Yes	
If No, please provide the primary r	eason:
Physical problem	☐ Verbal refusal
Cognitive/behavior problem	Other problem (specify):
Date of Collection: / /	(MM/DD/YYYY)

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Pati	ent ID:	P	Patient Initials:		
	t Date: / /		valuator Initials	:	
	ly Visit:				
	PARKINSONIAN FEATURES				
	Were Parkinsonian signs present? No Yes				
	Parkinsonian Signs: LEFT	No	Yes	Not Assessed	
	1. Resting tremor – arm				
	Slowing of fine motor movements				
	3. Rigidity - arm				
	Parkinsonian Signs: RIGHT	No	Yes	Not Assessed	
	4. Resting tremor – arm				
	5. Slowing of fine motor movements				
	6. Rigidity - arm				
	Parkinsonian Signs	No	Yes	Not Assessed	
	7. Bradykinesia				
	8. Parkinsonian gait disorder				
	9. Postural instability				

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Pati	ent ID:		Patient Initials	:	
	t Date:///		Evaluator Initials:		
	ly Visit:		1		
	-				
	CEREBROVASCULAR FEATU	RES			
	Were neurological signs consi cerebrovascular disease prese			consistent with	
	Findings consistent with stroke / cerebrovascular disease	No	Yes	Not Assessed	
	1. Cortical cognitive deficit (e.g., aphasia, apraxia, neglect)				
	Findings consistent with stroke / cerebrovascular disease: LEFT SIDE OF BODY	No	Yes	Not Assessed	
	2. Lateralized motor weakness				
	3. Lateralized abnormal reflexes (to include pathologically brisk deep tendon reflexes, Babinski signs, others)				
	4. Cortical visual field loss				
	5. Somatosensory loss				
	Findings consistent with stroke / cerebrovascular disease: RIGHT SIDE OF BODY	No	Yes	Not Assessed	
	6. Lateralized motor weakness				
	7. Lateralized abnormal reflexes (to include pathologically brisk deep tendon reflexes, Babinski signs, others)				
	8. Cortical visual field loss				
	Q Comptoconcory loss				

Patient ID:	Patient Initials:
Visit Date://	Evaluator Initials:
Study Visit:	

OTHER FINDINGS		No	Yes	Not Assessed
Patient demonst spontaneous, dis or involuntary cron examination				
2. Is magnetic gait a	apraxia present?			
3. Higher cortical v suggesting poste atrophy (e.g., pro simultagnosia, B syndrome) or ap	rior cortical osopagnosia, alint's			
4. Findings suggest progressive supr (PSP), corticobas other related dis	anuclear palsy sal syndrome, or			
5. Findings suggest muscle wasting, upper motor neulower motor neu	fasciculations, iron and/or			

	1101111101211	<u> </u>	up I	upor or		<u> </u>	
Patient ID:				Pat	Patient Initials:		
	Visit Date:///				luator Init	ials:	
Stu	ly Visit:						
	<u>COGNITIVE DIAGNOSIS</u>						
	Date of Evaluation: /	/		(M	IM/DD/YY	YY)	
	SYNDROMIC DIAGNOSIS						
	Normal Cognition		Imnai	red, Not M	1CI		
	MCI		Deme		101		
	Age of Onset: U	Unkno	wn				
	DDIMADY ETIOLOGICAL	Pres	sent				Mon
	PRIMARY ETIOLOGICAL DIAGNOSES	Pri		Primary	Contribut	ınσ	Non- contributing
		No	Yes			COII	contributing
	1. Alzheimer's disease						
	2. Lewy body disease						
	2a. Parkinson's disease						
	3. Vascular brain injury (based on clinical or imaging evidence)						
	3a. Peri-Ventricular Fazekas Extent Grade	□ 0	1	2	3 U	nknown/	N/A
	3b. Deep Fazekas Extent Grade	□ 0	1	_ 2	3 U	nknown/	N/A
	3c. Deep Fazekas Lesion Count Grade	□ 0	1	2	3 U	nknown/	N/A
		Pres	sent	Primary	Contribut	inσ	Non-
		No	Yes	Tilliary	Contribut	cont	ributing
	4. Traumatic brain injury						
	If Present: 4a. If present, does the subject have symptoms consistent with chronic traumatic encephalopathy?	□ No)	☐ Yes	U	nknown	

MarkVCID Follow-up Paper CRF Package

Patient ID: ______ Patient Initials: ______

Visit Date: ___/ ___/ Evaluator Initials: ______

Study Visit:

	Present		During outre	C + - i + i	Non-	
	No	Yes	Primary	Contributing	contributing	
5. Depression						
If Present:						
5a. Untreated						
☐ Treated with medication and/or counseling						
	Pres	sent	During outer	Cantuilantina	Non-	
	No	Yes	Primary	Contributing	contributing	
6. Cognitive impairment due to alcohol abuse						
If Present: 6a. Current alcohol abuse	□No	_	Yes	Unkno	wn	

Patie	Patient ID:				Patient Initials:			
	Date://			Eval	uator Initials	:		
Study Visit:								
	RELATED ETIOLOGICAL DIAGNOSES	Present	Prima	ary	Contributing	Non- contributing		
	7. Multiple system atrophy							
	8. Frontotemporal lobar degeneration							
	9. Essential tremor							
	10. Down syndrome							
	11. Huntington's disease							
	12. Prion disease (CJD, other)							
	13. Hydrocephalus							
	14. Epilepsy							
	15. CNS neoplasm							
	If present: 15a. ☐ Benign ☐ Malignant							
	16. Human immunodeficiency virus (HIV)							
	17. Bipolar disorder							
	18. Schizophrenia or other psychosis							
	19. Anxiety disorder							
	20. Delirium							
	21. Post-traumatic stress disorder (PTSD)							

Patient ID:			Patient Initials:			
Visit Date: / /]	Eval	uator Initials:	!	_
Study Visit:						
	Present	Prima	ıry	Contributing	Non- contributing	
22. Other psychiatric disease (specify):						
23. Cognitive impairment due to:						
23a. Other neurologic, genetic, or infectious conditions not listed above (specify):						

23b. Other substance

23c. Systemic disease/medical illness

impairment NOS: ____

23d. Medications

23e. Cognitive

abuse

	Markveib renew up rupe	1 CIVI I ackage					
P	Patient ID: Patient Initials:						
V	isit Date:// Evaluator Initials:						
St	tudy Visit:						
	MoCA (MONTREAL COGNITIV	E ASSESSMENT)					
	Was any part of the MoCA administered?						
	□ No □ Yes						
	If No, please provide the primary reason:	Physical problem					
	Cognitive/behavior problem Other problem	n (specify):					
Date of Examination: / / (MM/DD/YYYY)							
	Language of test administration:						
	English						
	☐ Spanish						
	Other (specify):						
	KEY: If the subject cannot complete any of the follow	ying exams, please give the reason by					
	entering one of the following codes:						
		= Cognitive/behavior problem= Verbal refusal					
	1. Visuospatial/ executive — Trails:	[0-1, 95-98]					
	The visuospatialy executive Trans.						
	2. Visuospatial/ executive — Cube:	[0-1, 95-98]					
	3. Visuospatial/ executive — Clock contour:	[0-1, 95-98]					
	4. Visuospatial/ executive — Clock numbers:	[0-1, 95-98]					
	5. Visuospatial/ executive — Clock hands:	[0-1, 95-98]					
	6. Language — Naming:	[0-3, 95-98]					
	7. Memory — Registration (two trials):	[0-10, 95-98]					

Patient ID:	Patient Initials:
Visit Date: / /	Evaluator Initials:
Study Visit:	

8. Attention — Digits:	[0-2, 95-98]
9. Attention — Letter A:	[0-1, 95-98]
10. Attention — Serial 7s:	[0-3, 95-98]
11. Language — Repetition:	[0-2, 95-98]
12. Language — Fluency:	[0-1, 95-98]
13. Abstraction:	[0-2, 95-98]
14. Delayed recall — No cue: (if not completed, enter reason code and skip to question 17)	[0-5, 95-98]
15. Delayed recall — Category cue:	[0-5, 95-98]
16. Delayed recall — Recognition:	[0-5, 95-98]
17. Orientation — Date:	[0-1, 95-98]
18. Orientation — Month:	[0-1, 95-98]
19. Orientation — Year:	[0-1, 95-98]
20. Orientation — Day:	[0-1, 95-98]
21. Orientation — Place:	[0-1, 95-98]
22. Orientation — City:	[0-1, 95-98]

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Pa	Patient ID: Patient Initials:					
	/isit Date: / / Evaluator Initials:					
	udy Visit:					
	NEUROPSYCHOLOGICAL TEST	TING BATTERY				
	Was any part of the Neuropsychological Testing Batt No Yes If No, please provide the primary reason:	_				
	Cognitive/behavior problem Other problem	n (specify):				
	Date of Examination: / / (M	IM/DD/YYYY)				
	Language of test administration: English					
	Spanish					
	Other (enecify)					

	<u>L</u>	<u> </u>					
P	Patient ID: Patient Initials:						
Visit Date:// Evaluator Initials:							
	Study Visit:						
	KEY: If the subject cannot complete any of the follow entering one of the following codes:	ing exams, please give the reason by					
	95 = Physical problem 96	= Cognitive/behavior problem					
	97 = Other problem 98	= Verbal refusal					
	1. Craft Story 21 Recall (Immediate):						
	a) If test not completed, enter reason code and						
	b) Total story units recalled, verbatim scoring:	[0-44]					
	c) Total story units recalled, paraphrase scoring	ng:[0-25]					
	2. Craft Story 21 Recall (Delayed):						
	a) If test not completed, enter reason code and	l skip to question 3a: [95-98]					
	b) Total story units recalled, verbatim scoring:	[0-44]					
	c) Total story units recalled, paraphrase scoring	ng:[0-25]					
	d) Delay time (minutes):	Unknown [0-85]					
	e) Cue ("boy") needed:	☐ No ☐ Yes					
	3. Number Span Test — Forward:	I alin to question (a) [OF 00]					
	a) If test not completed, enter reason code and						
	b) Number of correct trials:	[0-14]					
	c) Longest span forward:	[0, 3-9]					
	4. Number Span Test — Backward:a) If test not completed, enter reason code and	I okin to question 52: [05-08]					
	b) Number of correct trials:	[0-14]					
	c) Longest span backward:	[0, 2-8]					
	5. Category Fluency – Animals:	I alien to associate (c. [OF 00]					
	a) If test not completed, enter reason code and						
	b) Total number of animals named in 60 secon	ids: [0-77]					

Patient ID:	Patient Initials:					
Visit Date://	Evaluator Initials:					
tudy Visit:						
6. Verbal Fluency – Phonemic Tests (words begin a) If test not completed, enter reason code and	o ,					
b) Number of correct F-words generated in 1 m	ninute: [0-40]					
c) Number of F-words repeated in 1 minute:	[0-15]					
d) Number of non-F-words and rule violation e	rrors in 1 minute: [0-15]					
7. Trail Making Test A: a) If test not completed, enter reason code and	skip to question 8a: [95-98]					
b) Total number of seconds to complete (if not	finished by 150 seconds, enter 150)					
	[0-150]					
i. Number of commission errors:	[0-40]					
ii. Number of correct lines:	[0-24]					
8. Trail Making Test B: a) If test not completed, enter reason code and	skip to question 9a: [95-98]					
b) Total number of seconds to complete (if not	finished by 300 seconds, enter 300):					
	[0-300]					
i. Number of commission errors:	[0-40]					
ii. Number of correct lines:	[0-24]					
9. Multilingual Naming Test (MINT): a) If test not completed, enter reason code and	skip to question 10a: [95-98]					
b) Total score (9c+ 9e):	[0-32]					
c) Total correct without any cues (Uncued):	[0-32]					
d) Semantic cues – Number given:	[0-32]					
e) Semantic cues – Number correct with cue:	□ N/A[0-32]					
f) Phonemic cues – Number given:	[0-32]					
g) Phonemic cues – Number correct with cue:	□ N/A [0-32]					

Patient ID:		Patient Initials:	
Visit Date://		Evaluator Initials	:
Study Visit:			
10. Word list learning v a) Name of test:	with immediate/delay/rec HVLT CVLT-SF	ognition: CVLT SEVLT [Spanish]	
	Other (specify):		
b) Total number of v	words on list:		
c) If test not comple	ted, please select reason c	ode:	[95-98]
d) Learning Trial 1:			
e) Learning Trial 2:			
f) Learning Trial 3:			
g) Learning Trial 4:		□ N/A	
h) Learning Trial 5:		□ N/A	
i) Delay duration (i	f multiple options choose l	ongest):	
j) Delayed recall (if	multiple delay options, ch	oose longest):	
k) Recognition hits:			
l) Recognition false	positives:		

Patient ID:			_	Patient Initials:					
		'		Evaluator Initials:					
Study Visit:									
		<u>OR (CLINICAL D</u>	EMENII	IA KATIN	<u>u)</u>				
Was the CDR adi	ministered?								
□ No □	Yes								
If No, ple	ase provide tł	ne primary reason	ı: 🗌 Phys	sical proble	em 🔲 Verbal r	efusal			
Cognitive/be	havior proble	em 🗌 Other pr	oblem (s	necify):					
	navior proble	oener pr	obioiii (b	peeny j					
Date of Evaluation	on: /	/	(MM/DI) /VVVV)					
Date of Evaluation	JII /	/	_ (MM) DL	,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,					
Cartina 1 Chara	Janal CDD								
Section 1: Stand	iara CDR		11.15.41	DIGENER					
Please enter score		0 11	IMPAI	RMENT					
below:	None – 0	Questionable – 0.5	Mil	d – 1	Moderate – 2	Severe – 3			
1. Memory	No memory	Consistent slight		memory	Severe memory	Severe			
	loss, or slight inconsistent	forgetfulness; partial	loss, more		loss; only highly learned	memory loss; only fragments			
	forgetfulness	recollection of	defect int		material	remain			
		events; "benign"	with ever		retained; new				
		forgetfulness	activities		material rapidly lost				
2. Orientation	Fully	Fully oriented	Moderate	difficulty	Severe difficulty	Oriented to			
	oriented	except for slight	with time		with time	person only			
		difficulty with	relationsl	hips; for place at	relationships;				
		time relationships	examinat	•	usually disoriented to				
			have geog	graphic	time, often to				
·			disorienta elsewher		place				
3. Judgment	Solves	Slight impairment	1	e difficulty	Severely	Unable to			
and problem	everyday	in solving	in handlir		impaired in	make			
solving	problems,	problems,	problems		handling	judgments or			
	handles	similarities, and	similariti		problems,	solve problems			
	business and financial	differences	difference judgment	•	similarities, and differences;				
	affairs well;		maintain		social judgment				
	judgment				usually				
	good in				impaired				
	relation to								
	past performance								

	1
Patient ID:	Patient Initials:
Visit Date://	Evaluator Initials:
Study Visit:	

4.	Community affairs	Independent function at usual level in job, shopping, volunteer and social groups	Slight impairment in these activities	Unable to function independently at these activities, although may still be engaged in some; appears normal to casual inspection	No pretense of independent function outside the home; appears well enough to be taken to functions outside the family home	No pretense of independent function outside the home; appears too ill to be taken to functions outside the family home
5.	Home and hobbies	Life at home, hobbies, and intellectual interests well maintained	Life at home, hobbies, and intellectual interests slightly impaired	Mild but definite impairment of function at home; more difficult chores abandoned; more complicated hobbies and interests abandoned	Only simple chores preserved; very restricted interests, poorly maintained	No significant function in the home
6.	Personal care	Fully capable o	of self-care (= 0).	Needs prompting	Requires assistance in dressing, hygiene, keeping of personal effects	Requires much help with personal care; frequent incontinence
8.	8 STANDARD GLOBAL CDR					

	1
Patient ID:	Patient Initials:
Visit Date: / /	Evaluator Initials:
Study Visit:	

Section 2: Supplemental CDR						
Please enter score	IMPAIRMENT					
below:	None – 0	Questionable – 0.5	Mild – 1	Moderate – 2	Severe – 3	
1. Behavior, comportment, and personality	Socially appropriate behavior	Questionable changes in comportment, empathy, appropriateness of actions	Mild but definite changes in behavior	Moderate behavioral changes, affecting interpersonal relationships and interactions in a significant manner	Severe behavioral changes, making interpersonal interactions all unidirectional	
2. Language	No language difficulty, or occasional mild tip-of- the tongue	Consistent mild word-finding difficulties; simplification of word choice; circumlocution; decreased phrase length; and/or mild comprehension difficulties	Moderate word- finding difficulty in speech; cannot name objects in environment; reduced phrase length and/or agrammatical speech and/or reduced comprehension in conversation and reading	Moderate to severe impairments in either speech or comprehension; has difficulty communicating thoughts; writing may be slightly more effective	Severe comprehension deficits; no intelligible speech	

Patient ID: Patient Initials:						
Visit Date://	Date:/ Evaluator Initials:					
Study Visit:						
GDS (GERIATRIC DEPI	GDS (GERIATRIC DEPRESSION SCALE)					
Was the GDS administered?						
□ No □ Yes						
If No, please provide the primary reason:						
Cognitive/behavior problem Other probler	n (specify):					
Date of Evaluation:/ (MM/DD/YYYY)						
	Yes	No	Did not answer			
1. Are you basically satisfied with your life?						
2. Have you dropped many of your activities and interests?						
3. Do you feel that your life is empty?						
4. Do you often get bored?						
5. Are you in good spirits most of the time?						
6. Are you afraid that something bad is going to happen to you?						
7. Do you feel happy most of the time?						
8. Do you often feel helpless?						

Patient ID:		Patient Initials:				
Vis	sit Date://	Evaluator Initials:				
Stı	Study Visit:					
		Yes	No	Did not answer		
9.	Do you prefer to stay at home, rather than going out and doing new things?					
10.	Do you feel you have more problems with memory than most people?					
11.	Do you think it is wonderful to be alive?					
12.	Do you feel pretty worthless the way you are now?					
13.	Do you feel full of energy?					
14.	Do you feel that your situation is hopeless?					
15.	Do you think that most people are better off than you are?					

atient ID:			Patient Initials:			
isit Date://			Evaluator Initials:			
tudy Visit:		L				
	<u>LABORATORY TESTS</u>					
Date of Collection: / (MM/DD/YYYY)						
PHYSIOLOGIC MEASURES If fasting conditions are unknown, mark "not fasting". All tests denoted with * are required. Cholesterol related labs, blood sugar, and homocysteine should be collected under fasting conditions when possible.						
Measure	Fasting		Result			
1. HS-CRP	N/A		mg/L	☐ Not Done		
2. HbA1c	N/A		_ mmol/mol	☐ Not Done		
3. Blood Sugar*	Fasting >8 hours Not fasting		mmol/L	☐ Not Done		
4. Serum cholesterol*	Fasting >8 hours Not fasting		mg/dL	☐ Not Done		
5. HDL cholesterol*	Fasting >8 hours Not fasting		mg/dL	☐ Not Done		
6. LDL cholesterol*	Fasting >8 hours Not fasting		mg/dL	☐ Not Done		
7. Triglycerides*	Fasting >8 hours Not fasting		mg/dL	☐ Not Done		
8. Homocysteine	Fasting >8 hours Not fasting		mg/dL	☐ Not Done		
GENETICS						
Have any genetic tests been performed? If this is a follow up visit, only answer for any new tests performed. No Yes						
If yes:						
APOE genotype:	☐ E2/E2 ☐ E	2/E3	☐ E2/E4			
	☐ E3/E3 ☐ E	3/E4	☐ E4/E4	☐ Not Done		
Has a CWAS been co	ompleted?	Io	□ Voc			

Mai k v CiD Follow-up Faper CKF Fackage							
Pa	atient ID:	Patient Initials:					
V i	isit Date://	Evaluator Initials:					
St	udy Visit:						
	SAMPLE COLLECTION: CSF COLLECTION						
	Status:						
	If not collected, reason not collected:						
	Date CSF Samples Collected:/(MM/DD/YYYY)						
	Time since last meal: hours						
	Time Collected: : (24 hour clock)						
	Collector's Initials: (enter dash if no middle name)						
	Pre-Centrifugation sample: Appearance:	y):					
	Number of 0.25 mL aliquots:						

Patient ID:	Patient Initials:
Visit Date://	Evaluator Initials:
Study Visit:	
Were there any deviations? \(\sum \) No [Yes
If YES, indicate deviations below (select all the Sample not placed on dry ice or in -80°. If selected, please select one of the fole Placed on dry ice or in freezer with Placed on dry ice or in freezer 30-6. Placed on dry ice or in freezer 60+	C freezer immediately after aliquoting lowing: nin 30 minutes of aliquoting 60 minutes after aliquoting
☐ The participant was NOT fasting for a m	inimum of 6 hours prior to collection
Other deviation (specify):	

	<u>1</u>	<u> </u>		
Pa	atient ID:	Patient Initials:		
V	/isit Date:/ Evaluator Initials:			
St	tudy Visit:			
	SAMPLE COLLECTION: PLASM	IA COLLECTION		
	Status:			
	If not collected, reason not collected:			
	Date Plasma Samples Collected: / /	(MM/DD/YYYY)		
	Time since last meal: (hours)			
	Time Collected: : (24 hour clock)			
	Collector's Initials: (enter dash if no mid	ldle name)		
	Number of 0.25 mL plasma aliquots:			
	Number of 1 mL packed cell aliquots for DNA:			
	Temperature of Centrifugation: °C			
	Did plasma remain pink after centrifugation, indi	cating hemolysis?		
	Storage temperature: °C			

Patient ID:	Patient Initials:
Visit Date://	Evaluator Initials:
Study Visit:	
Were there any deviations?] Yes
If YES, indicate deviations below (select all tha Sample tube was not inverted 5-10 times	
☐ Sample not spun within 2 hours of collect If selected, please select one of the follo ☐ Spun 2-3 hours after collection ☐ Spun 3-4 hours after collection ☐ Spun 4+ hours after collection	
☐ Sample not spun at 2000g If selected, please select one of the follo ☐ Spun slower than 2000g ☐ Spun faster than 2000g	wing:
☐ Sample not spun for 10 minutes If selected, please select one of the follo ☐ Spun <10 minutes ☐ Spun >10 minutes	wing:
☐ Sample not placed on dry ice or in -80° C. If selected, please select one of the follo ☐ Placed on dry ice or in freezer within ☐ Placed on dry ice or in freezer 30-60 ☐ Placed on dry ice or in freezer 60+ m	wing: n 30 minutes of aliquoting n minutes after aliquoting
Other deviation (specify):	

	Markveib ronow up rape	1 CM 1 ackage			
Pa	atient ID:	Patient Initials:			
	/isit Date:// Evaluator Initials:				
St	tudy Visit:				
	SAMPLE COLLECTION: SERUI	M COLLECTION			
	Status: Collected Not Collected	-			
	Status. Gonected Not confected				
	If not collected, reason not collected:				
	Date Serum Samples Collected: / /	(MM/DD/YYYY)			
	Time since last meal: (hours)				
	Time Collected: : (24 hour clock)				
	Collector's Initials: (enter dash if no mic	ldle name)			
	Number of 0.25 mL aliquots:				
	Temperature of Centrifugation: °C				
	Did serum remain pink after centrifugation, indic	rating hemolysis?			
	Storage temperature: °C				

Markveib ronow up rupe	i diti i denage
Patient ID:	Patient Initials:
Visit Date://	Evaluator Initials:
Study Visit:	
Were there any deviations?] Yes
If YES, indicate deviations below (select all that After collection, sample not allowed to sit minutes (select all that apply): Sample not kept vertical Sample did not sit for 30-60 minutes If selected, please select one of the Sample sat <30 minutes Sample sat >60 minutes	s after collection
☐ Sample not spun at 2000g If selected, please select one of the follo ☐ Spun slower than 2000g ☐ Spun faster than 2000g	wing:
☐ Sample not spun for 10 minutes If selected, please select one of the follo ☐ Spun <10 minutes ☐ Spun >10 minutes	wing:
☐ Sample not placed on dry ice or in -80° C of the folloout of the following of the	wing: n 30 minutes of aliquoting minutes after aliquoting
Other deviation (specify):	

	Markveib ronow up rape	i citi i acttage
Pá	atient ID:	Patient Initials:
	isit Date://	Evaluator Initials:
	udy Visit:	
	SAMPLE COLLECTION: PLATELET POOR F	PLASMA (PPP) COLLECTION
	Status:	
	If not collected, reason not collected:	
	Date PPP Samples Collected: / /	(MM/DD/YYYY)
	Time Collected: : (24 hour clock)	
	Collector's Initials: (enter dash if no mic	ldle name)
	Time since last meal: hours	
	Number of 0.25 mL aliquots:	
	Did plasma remain pink after centrifugation, indi	cating hemolysis? No Yes
	Storage temperature: °C	

Mai Ky CiD Tollow-up Fape	I Chr Fachage
Patient ID:	Patient Initials:
Visit Date:///	Evaluator Initials:
Study Visit:	
Were there any deviations?] Yes
If YES, indicate deviations below (select all tha Sample tube was not inverted 5-10 times	
Sample not spun within 2 hours of collect If selected, please complete the following Spun hours after collection (roun	ng:
Sample not spun at 500g (first centrifugate of selected, please complete the following Speed sample spun at: g	
Sample not spun for 20 minutes (first cen If selected, please complete the following Duration of spin: min	
Sample not spun at 20C (first centrifugation of spin: C	
Sample not mixed at a 1:1 ratio after first If selected, please complete the following Volume of supernatant (platelet rich plate) Volume of DBS with additives: ml	ng: asma): mL
Sample not spun at 2,200g (second centri If selected, please complete the following Speed sample spun at: g	
Sample not spun for 20 minutes (second of selected, please complete the following Duration of spin: min	

Patient ID:	Patient Initials:
Visit Date:///	Evaluator Initials:
Study Visit:	
Deviations (continued): Sample not spun at 20C (second ce If selected, please complete the formula to the for	
If selected, please select one of the Placed on dry ice or in freezer Placed on dry ice or in freezer	-80° C freezer immediately after aliquoting ne following: r within 30 minutes of aliquoting r 30-60 minutes after aliquoting r 60+ minutes after aliquoting
Other deviation (specify):	

Date of Imaging: ___ / __ _ _ (MM/DD/YYYY)

Other reason: _____

Patient ID:	Patient Initials:				
Visit Date:// Evaluator Initials:					
Study Visit:					
OCTA SCREENING WO	DKCHFFT				
OCTA SCREENING WO	MIXJIILLI				
Date of OCTA Screening: /	(MM/DD/YYYY)				
Exclusion Criteria If the subject answers "yes" to any questions under #1-4 on the subject.	4, please DO NOT perfo	rm OC	TA test	ting	
Criterion			Yes	N/A	
1. Have you ever been diagnosed with any of the following eye diseases?					
1.1. Glaucoma					
1.2. Diabetic Retinopathy					
1.3. <u>Advanced</u> Dry Age-Related Macular Degenera	tion				
1.4. <u>Advanced</u> Wet Age-Related Macular Degeneration					
2. Have you ever had any of the following procedure:	s done?				
2.1. Laser Surgery on either eye for any reason (ex refractive procedures such as LASIK or catarac	S				
2.2. Injections into or around either eye (excluding procedures)					

Patient ID:	Patient Initials:	ent Initials:			
Visit Date:// Evaluator Initials					
Study Visit:					
Criterion		No	Yes	N/A	
3. If you have had your eyes dilated for an exami	nation in the past,				
3.1. Did you have a problem or allergy (excluding blurry vision)? (Mark not applicable if patient has never had their eyes dilated for an eye examination)					
3.2. Were you told not to get dilated again? (Mark not applicable if patient has never had their eyes dilated for an eye examination)					
4. Do you take any prescription eye drops (excluding artificial tears)?					
OCTA Enrollment					
Subject cannot undergo OCTA testing because Subject is enrolled in OCTA testing and agrees not agree to dilation, they are not eligible for enro	to dilation of right eye. If	the sub	ject do	oes	

Patient ID:	Patient Initials:					
Visit Date:///	Evaluator Initials:					
Study Visit:						
OCTA, INITIAL OD ANN	IIIAI EC	NI LOW/ I	ID			
OCTA: INITIAL OR ANN	UAL FU	<u>JLLUW-(</u>	<u> </u>			
Date of OCTA Scans: / / /	(MM	/DD/YYY	YY)			
Right Eye Dilation						
Subject's right eye is topically anesthetized wit		ops Propa	racaine 0.5	5%		
Subject's right eye is dilated with 1-2 drops each	ch of:					
☐ Tropicamide 1%						
☐ Phenylephrine 2.5%						
Other (specify):			hl			
(Note: If subject does not appear well dilated after another drop of each dilating drop)	10 mm	utes it is r	easonable (to administer		
OCTA Scans						
Scan Number		Sig	nal Streng	th		
Right Eye Angiography 3x3 mm Scan 1	8	<u> </u>	<u> </u>	Not Done		
Right Eye Angiography 3x3 mm Scan 2	8	<u> </u>	<u> </u>	Not Done		
Right Eye Angiography 3x3 mm Scan 3	□8	<u> </u>	<u> </u>	☐ Not Done		
Right Eye Angiography 3x3 mm Scan 4	8	<u> </u>	<u> </u>	Not Done		
Right Eye Optic Disc Cube 200x200 Scan 1	8	<u> </u>	<u> </u>	Not Done		
Right Eye Optic Disc Cube 200x200 Scan 2	8	<u> </u>	<u> </u>	Not Done		
Right Eye Optic Disc Cube 200x200 Scan 3	8	<u> </u>	<u> </u>	☐ Not Done		
Right Eye Optic Disc Cube 200x200 Scan 4	8	<u> </u>	<u> </u>	Not Done		

Patient ID:		Patient I	nitials:		
Visit Date: / / /		Evaluator Initials:			
Study Visit:					
Scan Number		S	ignal Streng	gth	
Left Eye Angiography 3x3 mm Scan 1		3	<u> </u>	☐ Not Done	
Left Eye Angiography 3x3 mm Scan 2		3	<u> </u>	☐ Not Done	
Left Eye Angiography 3x3 mm Scan 3		3	<u> </u>	☐ Not Done	
Left Eye Angiography 3x3 mm Scan 4		3	<u> </u>	☐ Not Done	
Left Eye Optic Disc Cube 200x200 Scan 1		3	<u> </u>	☐ Not Done	
Left Eye Optic Disc Cube 200x200 Scan 2		3	<u> </u>	☐ Not Done	
Left Eye Optic Disc Cube 200x200 Scan 3		3	<u> </u>	☐ Not Done	
Left Eye Optic Disc Cube 200x200 Scan 4		3	<u> </u>	☐ Not Done	

Patient ID:	Pat	tient Initia	als:	
Visit Date://	Eva	aluator In	itials:	
Study Visit:				
1. Has the subject seen an eye doctor in the past 5 years?	 lo	Yes	Unknown	
1a. <i>If yes,</i> has the subject released the medical records from this time period?	lo	Yes	Unknown	
2. Does the subject wear glasses or contacts?	lo	Yes	Unknown	
2a. <i>If yes,</i> are they worn to improve reading vision?	lo	Yes	Unknown	
2b. <i>If yes,</i> are they worn to improve distance vision?	lo	Yes	Unknown	
3. Has the subject ever had any of the following?				
3a. Cataract Surgery on Right Eye	lo	Yes	Unknown	
3b. Cataract Surgery on Left Eye	lo	Yes	Unknown	
Same-Day Retest				
Was this the initial OCTA scan?	Ю	Yes		
If this was the initial OCTA scan, was a retest completed on the same day?	Ю	Yes		

Patient ID:	P	Patient Initials:		
Visit Date: / /	Е	Evaluator Initials:		
Study Visit:				
OCTA: TEST/RETES	T _ CA	MEDAV		
OCIA. TEST/RETES)1 – JA	ME DAI		
Date of OCTA Scans: / /	(MN	M/DD/YYY	YY)	
Right Eye Dilation				
Subject's right eye is topically anesthetized wit		rops Propa	racaine 0.5	5%
Subject's right eye is dilated with 1-2 drops each	ch of:			
☐ Tropicamide 1%				
☐ Phenylephrine 2.5%				
Uther (specify):				_
(Note: If subject does not appear well dilated after another drop of each dilating drop)	· 10 mir	iutes it is r	easonable [•]	to administer
OCTA Scans				
Scan Number	Signal Strength			
Right Eye Angiography 3x3 mm Scan 1	□8	<u> </u>	<u> </u>	☐ Not Done
Right Eye Angiography 3x3 mm Scan 2	□8	<u> </u>	<u> </u>	Not Done
Right Eye Angiography 3x3 mm Scan 3	□8	<u> </u>	<u> </u>	Not Done
Right Eye Angiography 3x3 mm Scan 4	□ 8	<u> </u>	<u> </u>	Not Done
Right Eye Optic Disc Cube 200x200 Scan 1	□ 8	<u> </u>	<u> </u>	Not Done
Right Eye Optic Disc Cube 200x200 Scan 2	8	<u> </u>	<u> </u>	Not Done
Right Eye Optic Disc Cube 200x200 Scan 3	8	<u> </u>	<u> </u>	☐ Not Done
Right Eye Optic Disc Cube 200x200 Scan 4	8	<u> </u>	<u> </u>	Not Done

MarkVCID Follow-up Paper CRF Package

Patient ID: _____ Patient Initials: ____ ___

Visit Date: ___/ ___/ Evaluator Initials: ____ ___

Study Visit:

Signal Strength Scan Number 9 Left Eye Angiography 3x3 mm Scan 1 8 Not Done 8 □ 9 $\prod 10$ Left Eye Angiography 3x3 mm Scan 2 Not Done 9 8 □ 10 Left Eye Angiography 3x3 mm Scan 3 Not Done 8 \square 9 $\prod 10$ Left Eye Angiography 3x3 mm Scan 4 Not Done 8 □ 9 Left Eye Optic Disc Cube 200x200 Scan 1 ☐ Not Done □ 9 8 Left Eye Optic Disc Cube 200x200 Scan 2 $\prod 10$ Not Done Left Eye Optic Disc Cube 200x200 Scan 3 8 \square 9 Not Done

Left Eye Optic Disc Cube 200x200 Scan 4

8

□ 9

Not Done

Patient ID:	P	Patient Initials:		
Visit Date: / /	E	Evaluator Initials:		
Study Visit:				
	********	'W 4 4 D 4		
<u>OCTA: TEST/RETEST -</u>	WITH	IN 14 DA	<u>YS</u>	
Date of OCTA Scans: / /	(MI	M/DD/YYY	YY)	
Right Eye Dilation				
Subject's right eye is topically anesthetized wit		rops Propa	racaine 0.5	5%
Subject's right eye is dilated with 1-2 drops each	ch of:			
☐ Tropicamide 1%				
☐ Phenylephrine 2.5%				
Uther (specify):				_
(Note: If subject does not appear well dilated after another drop of each dilating drop)	: 10 mii	nutes it is r	easonable [•]	to administer
OCTA Scans				
Scan Number	Signal Strength			
Right Eye Angiography 3x3 mm Scan 1	□ 8	<u> </u>	<u> </u>	Not Done
Right Eye Angiography 3x3 mm Scan 2	□ 8	<u> </u>	<u> </u>	Not Done
Right Eye Angiography 3x3 mm Scan 3	□ 8	<u> </u>	<u> </u>	Not Done
Right Eye Angiography 3x3 mm Scan 4	□ 8	<u> </u>	<u> </u>	Not Done
Right Eye Optic Disc Cube 200x200 Scan 1	8	<u> </u>	□ 10	Not Done
Right Eye Optic Disc Cube 200x200 Scan 2	8	<u> </u>	□ 10	Not Done
Right Eye Optic Disc Cube 200x200 Scan 3	□ 8	<u> </u>	□ 10	Not Done
Right Eye Optic Disc Cube 200x200 Scan 4	8	<u> </u>	<u> </u>	☐ Not Done

Study Visit:

Scan Number	Signal Strength			
Left Eye Angiography 3x3 mm Scan 1	8	<u> </u>	<u> </u>	☐ Not Done
Left Eye Angiography 3x3 mm Scan 2	□8	<u> </u>	<u> </u>	☐ Not Done
Left Eye Angiography 3x3 mm Scan 3	□8	<u> </u>	□ 10	☐ Not Done
Left Eye Angiography 3x3 mm Scan 4	□8	<u> </u>	<u> </u>	☐ Not Done
Left Eye Optic Disc Cube 200x200 Scan 1	□8	<u> </u>	<u> </u>	☐ Not Done
Left Eye Optic Disc Cube 200x200 Scan 2	□8	<u> </u>	<u> </u>	☐ Not Done
Left Eye Optic Disc Cube 200x200 Scan 3	□8	<u> </u>	<u> </u>	☐ Not Done
Left Eye Optic Disc Cube 200x200 Scan 4	8	<u> </u>	<u> </u>	☐ Not Done