

## Parkinson's Disease and PD Genetic Schedule of Activities (Years 0 - 5)

Visit Number	Screening	Baseline (BL)	V02	V04	V05	V06	R06	V08	R08	V10	R10	V12	Unsched	<sup>b</sup> Transition Activities
Assessment	**Timepoint	-60 days	0	6 mths	12 (Y1) mths	18 (Y2) mths	30 mths	36 (Y3) mths	42 mths	48 (Y4) mths	54 mths	60 (Y5) mths	--	--
<b>Consent Activities</b>														
Documentation of Informed Consent		I	As Needed											I
Continuing Consent				X		X		X		X		X		
Research Proxy Designation		I	As Needed											X
Consent to share contact information		X	As Needed											X
Informed Consent Tracking Log		X	As Needed											X
<b>General Activities</b>														
Demographics		X												X
Family History		X												X
Socio-Economics		X												X
Physical Examination		X												
Vital Signs (Height and Weight BL + Annually)		X	X	X	X	X	X	X	X	X	X	X	X	X
Review Inclusion/Exclusion Criteria		I	I											
Visit Status		X	X	X	X	X	X	X	X	X	X	X	X	X
Screen Fail		As Needed												As Needed
Conclusion of Study Participation			As Needed											
<b>Neurological/Motor Assessments</b>														
Participant Motor Function Questionnaire			P		P		P		P		P		P	
Freezing and Falls			X		X		X		X		X		X	
PD Diagnosis History		I												
Neurological Examination		I			I		I		I		I		I	I
MDS-UPDRS ON/OFF Determination & Dosing				X	X	X	X	X <sup>c</sup>	X	X <sup>c</sup>	X	X <sup>c</sup>	X	
MDS-UPDRS Part Ia, Part III and Hoehn & Yahr			I	I	I	I	I	<sup>a</sup> I	I	<sup>a</sup> I	I	<sup>a</sup> I	I	
MDS-UPDRS Part Ib and Part II			P	P	P	P	P	P	P	P	P	P	P	
Modified Schwab & England ADL			I	I	I	I	I	I	I	I	I	I	I	
MDS-UPDRS Part IV				I <sup>d</sup>	I <sup>d</sup>	I <sup>d</sup>	I <sup>d</sup>	I <sup>d</sup>	I <sup>d</sup>	I <sup>d</sup>	I <sup>d</sup>	I <sup>d</sup>	I <sup>d</sup>	
MDS-UPDRS Repeat Part III/Hoehn & Yahr				I <sup>d</sup>	I <sup>d</sup>	I <sup>d</sup>		I <sup>d</sup>		I <sup>d</sup>		I <sup>d</sup>		
Features of Parkinsonism			I	I	I	I	I	I	I	I	I	I	I	
Other Clinical Features			I	I	I	I	I	I	I	I	I	I	I	
Primary Clinical Diagnosis			I	I	I	I	I	I	I	I	I	I	I	
<b>Non-Motor Assessments</b>														
Olfactory Testing (UPSIT)			P											
REM Sleep Behavior Disorder Screening Questionnaire			P		P		P		P		P		P	
Epworth Sleepiness Scale			P		P		P		P		P		P	
SCOPA-AUT			P		P		P		P		P		P	
Neuro QoL			P		P		P		P		P		P	
<b>Cognitive Assessments</b>														
Montreal Cognitive Assessment*		X			X		X		X		X		X	
Clock Drawing*		X			X		X		X		X		X	
Lexical Fluency*			X		X		X		X		X		X	
Hopkins Verbal Learning Test-Revised*			X		X		X		X		X		X	
Benton Judgment of Line Orientation*			X		X		X		X		X		X	
Semantic Fluency (Animals only)*			X		X		X		X		X		X	
Letter Number Sequencing*			X		X		X		X		X		X	
Symbol Digit Modalities Test*			X		X		X		X		X		X	
Trail Making Test (A and B)*			X		X		X		X		X		X	
Modified Boston Naming Test*			X		X		X		X		X		X	
Cognitive Change			P	P	P	P	P	P	P	P	P	P	P	
Cognitive Categorization			I		I		I		I		I		I	
<b>Neuropsychological Assessments</b>														
State-Trait Anxiety Inventory for Adults			P		P		P		P		P		P	
Geriatric Depression Scale			P		P		P		P		P		P	

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Visit Number	Screening	Baseline (BL)	V02	V04	V05	V06	R06	V08	R08	V10	R10	V12	Unsched	<sup>b</sup> Transition Activities	
Assessment	Timepoint	-60 days	0	6 mths	12 (Y1) mths	18 mths	24 (Y2) mths	30 mths	36 (Y3) mths	42 mths	48 (Y4) mths	54 mths	60 (Y5)	--	--
QUIP		P		P			P		P		P		P		
<b>Clinical and Biological Samples</b>															
Clinical Lab blood sample		X											X		
Research samples (blood + urine)			X	X	X	X	X		X		X		X		
Lumbar puncture			X		X		X		X		X		X		
Skin biopsy <sup>f</sup>			X				X				X				X <sup>c</sup>
<b>Imaging Activities</b>															
Pregnancy Test (prior to DaTscan injection), if applicable		X			X		X				X				
DaTscan Imaging		X			X		X				X				
MRI			X		X		X				X				
<b>Safety and General Health</b>															
<sup>†</sup> Adverse Events		X	X		X		X		X		X		X	X	
Adverse Event Telephone Assessment		X	X		X		X		X		X		X		
Current Medical Conditions Review		X	X	X	X	X	X	X	X	X	X	X	X	X	
Concomitant Medication Review		X	X	X	X	X	X	X	X	X	X	X	X	X	
LEDD Concomitant Medication Log		X	X	X	X	X	X	X	X	X	X	X	X	X	
Participation in Other Studies		X	X	X	X	X	X	X	X	X	X	X	X	X	X
Surgery for PD Log		As Needed													
Report of Pregnancy		As Needed													

<sup>†</sup> = Investigator completed assessment

P = Participant completed assessment

X = Investigator or Coordinator completed assessment (or as otherwise delegated)

ROX Visits are conducted remotely (e.g., video, audio)

a = rigidity and postural stability will not be assessed for Remote visits; Part III and Hoehn & Yahr not done if phone/audio only

b = Transition Activities completed for all previously enrolled participants transitioning into new database at first visit only

c = Previously enrolled participants transitioning to new database may be asked to have skin biopsy. If not done at first visit, may be conducted at a subsequent in person visit.

d = Only complete once participant has initiated dopaminergic medication/DBS for treating the symptoms of PD

e = Completed to record timing of the single MDS-UPDRS assessment during remote visits. If participant is on medications for treating PD, the preferred state for the MDS-UPDRS remote assessment is ON.

f = Skin biopsy will be conducted at participating sites.

\*Completed on paper source first, and then scores entered in EDC

\*\*Window of +45 days either side of Target Visit Date

#Adverse events collected only day of and 2-3 days post DaTscan, LP and skin biopsy per protocol.

## Parkinson's Disease and PD Genetic Schedule of Activities (Years 6-13)

Visit Number	R12	V13	R13	V14	R14	V15	R15	V16	R16	V17	R17	V18	R18	V19	R19	V20	Unscheduled	<sup>b</sup> Transition Activities	
Assessment	**Timepoint	66 mths	72 (Y6)	78 mths	84 (Y7)	90 mths	96 (Y8)	102 mths	108 (Y9)	114 mth	120 (Y10)	126 mths	132 (Y11)	138 mths	144 (Y12)	150 mths	156 (Y13)	---	---
<b>Consent Activities</b>																			
Documentation of Informed Consent	As Needed																	I	
Continuing Consent		X		X		X		X		X		X		X		X			
Consent to share contact information	As Needed																		
Research Proxy Designation	As Needed (I)																		
Informed Consent Tracking Log	As Needed																		
<b>General Activities</b>																			
Demographics																			X
Family History																			X
Socio-Economics																			X
Vital Signs + Height and Weight		X		X		X		X		X		X		X		X		X	X
Visit Status	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X
Screen Fail																			As Needed
Conclusion of Study Participation	As Needed																		
<b>Neurological/Motor Assessments</b>																			
Participant Motor Function Questionnaire		P		P		P		P		P		P		P		P			
Freezing and Falls		X		X		X		X		X		X		X		X			
Neurological Examination		I		I		I		I		I		I		I		I		I	
MDS-UPDRS Part Ia, Part III and Hoehn & Yahr	<sup>a</sup> I	I	<sup>a</sup> I	I	<sup>a</sup> I	I	<sup>a</sup> I	I	<sup>a</sup> I	I	<sup>a</sup> I	I	<sup>a</sup> I	I	<sup>a</sup> I	I	<sup>a</sup> I	I	
MDS-UPDRS Part Ib and Part II	P	P	P	P	P	P	P	P	P	P	P	P	P	P	P	P	P	P	
Modified Schwab & England ADL	I	I	I	I	I	I	I	I	I	I	I	I	I	I	I	I	I	I	
MDS-UPDRS Part IV	I <sup>d</sup>	I <sup>d</sup>	I <sup>d</sup>	I <sup>d</sup>	I <sup>d</sup>	I <sup>d</sup>	I <sup>d</sup>	I <sup>d</sup>	I <sup>d</sup>	I <sup>d</sup>	I <sup>d</sup>	I <sup>d</sup>	I <sup>d</sup>	I <sup>d</sup>	I <sup>d</sup>	I <sup>d</sup>	I <sup>d</sup>	I <sup>d</sup>	
MDS-UPDRS ON/OFF Determination & Dosing	X <sup>c</sup>	X	X <sup>c</sup>	X	X <sup>c</sup>	X	X <sup>c</sup>	X	X <sup>c</sup>	X	X <sup>c</sup>	X	X <sup>c</sup>	X	X <sup>c</sup>	X	X <sup>c</sup>	X	
MDS-UPDRS Repeat Part III/Hoehn & Yahr		I <sup>d</sup>		I <sup>d</sup>		I <sup>d</sup>		I <sup>d</sup>		I <sup>d</sup>		I <sup>d</sup>		I <sup>d</sup>		I <sup>d</sup>		I <sup>d</sup>	
Features of Parkinsonism		I	I	I	I	I	I	I	I	I	I	I	I	I	I	I	I	I	
Other Clinical Features		I	I	I	I	I	I	I	I	I	I	I	I	I	I	I	I	I	
Primary Clinical Diagnosis	I	I	I	I	I	I	I	I	I	I	I	I	I	I	I	I	I	I	
<b>Non-Motor Assessments</b>																			
REM Sleep Behavior Disorder Screening Questionnaire		P		P		P		P		P		P		P		P			
Epworth Sleepiness Scale		P		P		P		P		P		P		P		P			
SCOPA-AUT		P		P		P		P		P		P		P		P			
Neuro QoL		P		P		P		P		P		P		P		P			
<b>Cognitive Assessments</b>																			
Montreal Cognitive Assessment*		X		X		X		X		X		X		X		X			
Clock Drawing*		X		X		X		X		X		X		X		X			
Lexical Fluency*		X		X		X		X		X		X		X		X			
Hopkins Verbal Learning Test-Revised*		X		X		X		X		X		X		X		X			
Benton Judgment of Line Orientation*		X		X		X		X		X		X		X		X			
Semantic Fluency (Animals only)*		X		X		X		X		X		X		X		X			
Letter Number Sequencing*		X		X		X		X		X		X		X		X			
Symbol Digit Modalities Test*		X		X		X		X		X		X		X		X			
Trail Making Test (A and B)*		X		X		X		X		X		X		X		X			
Modified Boston Naming Test*		X		X		X		X		X		X		X		X			
Cognitive Change		P		P		P		P		P		P		P		P			
Cognitive Categorization		I		I		I		I		I		I		I		I			

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Visit Number		R12	V13	R13	V14	R14	V15	R15	V16	R16	V17	R17	V18	R18	V19	R19	V20	Unsched	<sup>b</sup> Transition Activities
Assessment	**Timepoint	66 mths	72 (Y6)	78 mths	84 (Y7)	90 mths	96 (Y8)	102 mths	108 (Y9)	114 mth	120 (Y10)	126 mths	132 (Y11)	138 mths	144 (Y12)	150 mths	156 (Y13)	---	---
<b>Neuropsychological Assessments</b>																			
State-Trait Anxiety Inventory for Adults			P		P		P		P		P		P		P		P		
Geriatric Depression Scale			P		P		P		P		P		P		P		P		
QUIP			P		P		P		P		P		P		P		P		
<b>Clinical and Biological Samples</b>																			
Clinical Lab blood sample																		X	
Research samples (blood + urine)			X		X		X		X		X		X		X		X		
Lumbar puncture					X				X				X				X		
Skin biopsy <sup>f</sup>																			X <sup>c</sup>
<b>Safety and General Health</b>																			
<sup>#</sup> Adverse Events					X				X				X				X	X	
Adverse Event Telephone Assessment					X				X				X				X		
Current Medical Conditions Review		X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X
Concomitant Medication Review		X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X
Participation in Other Studies		X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X
LEDD Concomitant Medication Log		As Needed																	
Surgery for PD Log		As Needed																	
Report of Pregnancy		As Needed																	

I = Investigator completed assessment

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\*\*Window of +45 days either side of Target Visit Date

#Adverse events collected only day of and 2-3 days post LP and skin biopsy per protocol.

## Prodromal Schedule of Activities (Years 0 - 5)

### Prodromal Schedule of Activities (Years 0 - 5)

Visit Number	SC (DAT)	BL (Clinic)	R01	V04	R04	V06	R06	V08	R08	V10	R10	V12	Unsched	<sup>b</sup> Transition Activities	
Assessment	**Timepoint	-60 days	0	6 mths (Y1)	12 mths (Y2)	18 mths (Y3)	24 mths (Y4)	30 mths (Y5)	36 mths (Y6)	42 mths (Y7)	48 mths (Y8)	54 mths (Y9)	60 mths (Y10)	--	--
<b>Consent Activities</b>															
Documentation of Screening DaTscan Consent		I													
Documentation of Informed Consent			I	As Needed											I
Continuing Consent				X		X		X		X		X			
Research Proxy Designation			I	As Needed											X
Consent to share contact information			X	As Needed											X
Informed Consent Tracking Log		X	X	As Needed											
<b>Pre-Screening Activities</b>															
Prodromal History		X													
Olfactory Testing (UPSIT)		p <sup>e</sup>													
<b>General Activities</b>															
Demographics			X												X
Family History			X												X
Socio-Economics			X												X
Physical Examination			X												
Vital Signs (Height and Weight BL + Annually)			X		X		X		X		X		X	X	
Review Inclusion/Exclusion Criteria		I	I												
Visit Status		X	X	X	X	X	X	X	X	X	X	X	X	X	
Screen Fail			As Needed												As Needed
Conclusion of Study Participation				As Needed											
<b>Neurological/Motor Assessments</b>															
Participant Motor Function Questionnaire			P		P		P		P		P		P		
Freezing and Falls			X		X		X		X		X		X		
Neurological Examination			I		I		I		I		I		I	I	
MDS-UPDRS ON/OFF Determination & Dosing				X <sup>c</sup>	X	X <sup>c</sup>	X	X <sup>c</sup>	X	X <sup>c</sup>	X	X <sup>c</sup>	X		
MDS-UPDRS Part Ia, Part III and Hoehn & Yahr			I	<sup>a</sup> I	I	<sup>a</sup> I	I	<sup>a</sup> I	I	<sup>a</sup> I	I	<sup>a</sup> I	I		
MDS-UPDRS Part Ib and Part II			P	P	P	P	P	P	P	P	P	P	P		
Modified Schwab & England ADL			I	I	I	I	I	I	I	I	I	I	I		
MDS-UPDRS Part IV				I <sup>d</sup>	I <sup>d</sup>	I <sup>d</sup>	I <sup>d</sup>	I <sup>d</sup>	I <sup>d</sup>	I <sup>d</sup>	I <sup>d</sup>	I <sup>d</sup>	I <sup>d</sup>		
MDS-UPDRS Repeat Part III/Hoehn & Yahr					I <sup>d</sup>		I <sup>d</sup>		I <sup>d</sup>		I <sup>d</sup>		I <sup>d</sup>		
Features of Parkinsonism			I	I	I	I	I	I	I	I	I	I	I		
Other Clinical Features			I	I	I	I	I	I	I	I	I	I	I		
Primary Clinical Diagnosis			I	I	I	I	I	I	I	I	I	I	I		
<b>Non-Motor Assessments</b>															
Olfactory Testing (UPSIT)							P								
REM Sleep Behavior Disorder Screening Questionnaire			P		P		P		P		P		P		
Epworth Sleepiness Scale			P		P		P		P		P		P		
SCOPA-AUT			P		P		P		P		P		P		
Neuro QoL			P		P		P		P		P		P		
<b>Cognitive Assessments</b>															
Montreal Cognitive Assessment*			X		X		X		X		X		X		
Clock Drawing*			X		X		X		X		X		X		
Lexical Fluency*			X		X		X		X		X		X		
Hopkins Verbal Learning Test-Revised*			X		X		X		X		X		X		
Benton Judgment of Line Orientation*			X		X		X		X		X		X		
Semantic Fluency (Animals only)*			X		X		X		X		X		X		

## Prodromal Schedule of Activities (Years 0 - 5)

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Letter Number Sequencing*			X		X		X		X		X		X		
Symbol Digit Modalities Test*			X		X		X		X		X		X		
Trail Making Test (A and B)*			X		X		X		X		X		X		
Modified Boston Naming Test*			X		X		X		X		X		X		
Cognitive Change			P		P		P		P		P		P		
Cognitive Categorization			I		I		I		I		I		I		
<b>Neuropsychological Assessments</b>															
State-Trait Anxiety Inventory for Adults			P		P		P		P		P		P		
Geriatric Depression Scale			P		P		P		P		P		P		
QUIP			P		P		P		P		P		P		
<b>Clinical and Biological Samples</b>															
Clinical Lab blood sample			X											X	
Research samples (blood + urine)			X		X		X		X		X		X		
Lumbar puncture			X		X		X		X		X		X		
Skin biopsy <sup>f</sup>			X				X				X				X <sup>c</sup>
<b>Imaging Activities</b>															
Pregnancy Test (prior to DaTscan injection), if applicable		X			X		X				X				
DaTscan Imaging		X			X		X				X				
MRI			X		X		X				X				
<b>Safety and General Health</b>															
# Adverse Events		X	X		X		X		X		X		X	X	
Adverse Event Telephone Assessment		X	X		X		X		X		X		X		
Current Medical Conditions Review			X	X	X	X	X	X	X	X	X	X	X	X	
Concomitant Medication Review		X	X	X	X	X	X	X	X	X	X	X	X	X	
Participation in Other Studies		X	X	X	X	X	X	X	X	X	X	X	X	X	X
LEDD Concomitant Medication Log				As Needed											
Surgery for PD Log				As Needed											
Report of Pregnancy				As Needed											

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f = Skin biopsy will be conducted at participating sites.

g = Performed for sites recruiting participants not referred from Screening Core

\*Completed on paper source first, and then scores entered in EDC

\*\*Window of +45 days either side of Target Visit Date

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<b>Consent Activities</b>																		
Documentation of Informed Consent	As Needed																	I
Continuing Consent		X		X		X		X		X		X		X		X		
Consent to share contact information	As Needed																	
Research Proxy Designation	As Needed (I)																	
Informed Consent Tracking Log	As Needed																	
<b>General Activities</b>																		
Demographics																		X
Family History																		X
Socio-Economics																		X
Vital Signs + Height and Weight		X		X		X		X		X		X		X		X	X	
Visit Status	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	
Screen Fail																		As Needed
Conclusion of Study Participation	As Needed																	
<b>Neurological/Motor Assessments</b>																		
Participant Motor Function Questionnaire		P		P		P		P		P		P		P		P		
Freezing and Falls		X		X		X		X		X		X		X		X		
Neurological Examination		I		I		I		I		I		I		I		I	I	
MDS-UPDRS Part Ia, Part III and Hoehn & Yahr	<sup>a</sup> I	I	<sup>a</sup> I	I	<sup>a</sup> I	I	<sup>a</sup> I	I	<sup>a</sup> I	I	<sup>a</sup> I	I	<sup>a</sup> I	I	<sup>a</sup> I	I		
MDS-UPDRS Part Ib and Part II	P	P	P	P	P	P	P	P	P	P	P	P	P	P	P	P		
Modified Schwab & England ADL	I	I	I	I	I	I	I	I	I	I	I	I	I	I	I	I		
MDS-UPDRS Part IV	I <sup>d</sup>	I <sup>d</sup>	I <sup>d</sup>	I <sup>d</sup>	I <sup>d</sup>	I <sup>d</sup>	I <sup>d</sup>	I <sup>d</sup>	I <sup>d</sup>	I <sup>d</sup>	I <sup>d</sup>	I <sup>d</sup>	I <sup>d</sup>	I <sup>d</sup>	I <sup>d</sup>	I <sup>d</sup>		
MDS-UPDRS ON/OFF Determination & Dosing	X <sup>c</sup>	X	X <sup>c</sup>	X	X <sup>c</sup>	X	X <sup>c</sup>	X	X <sup>c</sup>	X	X <sup>c</sup>	X	X <sup>c</sup>	X	X <sup>c</sup>	X	X <sup>c</sup>	
MDS-UPDRS Repeat Part III/Hoehn & Yahr		I <sup>d</sup>		I <sup>d</sup>		I <sup>d</sup>		I <sup>d</sup>		I <sup>d</sup>		I <sup>d</sup>		I <sup>d</sup>		I <sup>d</sup>		
Features of Parkinsonism		I	I	I	I	I	I	I	I	I	I	I	I	I	I	I		
Other Clinical Features		I	I	I	I	I	I	I	I	I	I	I	I	I	I	I		
Primary Clinical Diagnosis	I	I	I	I	I	I	I	I	I	I	I	I	I	I	I	I		
<b>Non-Motor Assessments</b>																		
REM Sleep Behavior Disorder Screening Questionnaire		P		P		P		P		P		P		P		P		
Epworth Sleepiness Scale		P		P		P		P		P		P		P		P		
SCOPA-AUT		P		P		P		P		P		P		P		P		
Neuro QoL		P		P		P		P		P		P		P		P		
<b>Cognitive Assessments</b>																		
Montreal Cognitive Assessment*		X		X		X		X		X		X		X		X		
Clock Drawing*		X		X		X		X		X		X		X		X		
Lexical Fluency*		X		X		X		X		X		X		X		X		
Hopkins Verbal Learning Test-Revised*		X		X		X		X		X		X		X		X		
Benton Judgment of Line Orientation*		X		X		X		X		X		X		X		X		
Semantic Fluency (Animals only)*		X		X		X		X		X		X		X		X		
Letter Number Sequencing*		X		X		X		X		X		X		X		X		
Symbol Digit Modalities Test*		X		X		X		X		X		X		X		X		
Trail Making Test (A and B)*		X		X		X		X		X		X		X		X		
Modified Boston Naming Test*		X		X		X		X		X		X		X		X		
Cognitive Change		P		P		P		P		P		P		P		P		
Cognitive Categorization		I		I		I		I		I		I		I		I		

## Prodromal Schedule of Activities (Years 6-13)

Visit Number	R12	V13	R13	V14	R14	V15	R15	V16	R16	V17	R17	V18	R18	V19	R19	V20	Unsched	<sup>b</sup> Transition Activities
Assessment	**Timepoint 66 mths	72 (Y6)	78 mths	84 (Y7)	90 mths	96 (Y8)	102 mths	108 (Y9)	114 mth	120 (Y10)	126 mths	132 (Y11)	138 mths	144 (Y12)	150 mths	156 (Y13)	---	---
<b>Neuropsychological Assessments</b>																		
State-Trait Anxiety Inventory for Adults		P		P		P		P		P		P		P		P		
Geriatric Depression Scale		P		P		P		P		P		P		P		P		
QUIP		P		P		P		P		P		P		P		P		
<b>Clinical and Biological Samples</b>																		
Clinical Lab blood sample																	X	
Research samples (blood + urine)		X		X		X		X		X		X		X		X		
Lumbar puncture				X				X				X				X		
Skin biopsy <sup>f</sup>																		X <sup>c</sup>
<b>Safety and General Health</b>																		
# Adverse Events				X				X				X				X	X	
Adverse Event Telephone Assessment				X				X				X				X		
Current Medical Conditions Review	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	
Concomitant Medication Review	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	
Participation in Other Studies	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X
LEDD Concomitant Medication Log	As Needed																	
Surgery for PD Log	As Needed																	
Report of Pregnancy	As Needed																	

I = Investigator completed assessment

P = Participant completed assessment

ROX Visits are conducted remotely (e.g., video, audio)

a = rigidity and postural stability will not be assessed for Remote visits; Part III and Hoehn & Yahr not done if phone/audio only

b = Transition Activities completed for all previously enrolled participants transitioning into new database at first visit only

c = Previously enrolled participants transitioning to new database may be asked to have skin biopsy. If not done at first visit, may be conducted at a subsequent in person visit.

d = Only complete once participant has initiated dopaminergic medication/DBS for treating the symptoms of PD

e = Completed to record timing of the single MDS-UPDRS assessment during remote visits. If participant is on medications for treating PD, the preferred state for the MDS-UPDRS remote assessment is ON.

f = Skin biopsy will be conducted at participating sites

\*Completed on paper source first, and then scores entered in to EDC.

\*\*Window of +45 days either side of Target Visit Date

#Adverse events collected only day of and 2-3 days post LP and skin biopsy per protocol.



## Healthy Control Schedule of Activities (Years 0 - 5)

Visit Number	Screening	Baseline (BL)	V02	V04	V05	V06	R06	V08	R08	V10	R10	V12	Unsched	<sup>b</sup> Transition Activities	
Assessment	**Timepoint	-60 days	0	6 mths	12 (Y1)	18 mths	24 (Y2)	30 mths	36 (Y3)	42 mths	48 (Y4)	54 mths	60 (Y5)	--	--
<b>Consent Activities</b>															
Documentation of Informed Consent	I	As Needed												I	
Continuing Consent				X		X		X		X		X			
Research Proxy Designation	X	As Needed (I)													
Consent to share contact information	X	As Needed												X	
Informed Consent Tracking Log	X	As Needed												X	
<b>General Activities</b>															
Demographics	X														X
Family History	X														X
Socio-Economics	X														X
Physical Examination	X														
Vital Signs (Height and Weight BL + Annually)	X	X	X	X	X	X		X		X		X	X		
Review Inclusion/Exclusion Criteria	I	I													
Visit Status	X	X	X	X	X	X	X	X	X	X	X	X	X	X	
Screen Fail		As Needed													As Needed
Conclusion of Study Participation		As Needed													
<b>Neurological/Motor Assessments</b>															
Participant Motor Function Questionnaire		P		P		P		P		P		P			
Freezing and Falls		X		X		X		X		X		X			
Neurological Examination	I			I		I		I		I		I	I		
MDS-UPDRS Part Ia, Part III and Hoehn & Yahr		I	I	I	I	I	<sup>a</sup> I	I	<sup>a</sup> I	I	<sup>a</sup> I	I	<sup>a</sup> I		
MDS-UPDRS Part Ib and Part II		P	P	P	P	P	P	P	P	P	P	P	P		
Modified Schwab & England ADL		I	I	I	I	I	I	I	I	I	I	I	I		
Features of Parkinsonism		I	I	I	I	I	I	I	I	I	I	I	I		
Other Clinical Features		I	I	I	I	I	I	I	I	I	I	I	I		
Primary Clinical Diagnosis		I	I	I	I	I	I	I	I	I	I	I	I		
<b>Non-Motor Assessments</b>															
Olfactory Testing (UPSIT)		P													
REM Sleep Behavior Disorder Screening Questionnaire		P		P		P		P		P		P			
Epworth Sleepiness Scale		P		P		P		P		P		P			
SCOPA-AUT		P		P		P		P		P		P			
Neuro QoL		P		P		P		P		P		P			
<b>Cognitive Assessments</b>															
Montreal Cognitive Assessment*	X			X		X		X		X		X			
Clock Drawing*	X			X		X		X		X		X			
Lexical Fluency*		X		X		X		X		X		X			
Hopkins Verbal Learning Test-Revised*		X		X		X		X		X		X			
Benton Judgment of Line Orientation*		X		X		X		X		X		X			
Semantic Fluency (Animals only)*		X		X		X		X		X		X			

## Healthy Control Schedule of Activities (Years 0 - 5)

Visit Number		Screening	Baseline (BL)	V02	V04	V05	V06	R06	V08	R08	V10	R10	V12	Unscheduled	<sup>b</sup> Transition Activities
Assessment	**Timepoint	-60 days	0	6 mths	12 (Y1)	18 mths	24 (Y2)	30 mths	36 (Y3)	42 mths	48 (Y4)	54 mths	60 (Y5)	---	---
Letter Number Sequencing*			X		X		X		X		X		X		
Symbol Digit Modalities Test*			X		X		X		X		X		X		
Trail Making Test (A and B)*			X		X		X		X		X		X		
Modified Boston Naming Test*			X		X		X		X		X		X		
Cognitive Change			P	P	P	P	P		P		P		P		
Cognitive Categorization			I		I		I		I		I		I		
<b>Neuropsychological Assessments</b>															
State-Trait Anxiety Inventory for Adults			P		P		P		P		P		P		
Geriatric Depression Scale			P		P		P		P		P		P		
QUIP			P		P		P		P		P		P		
<b>Clinical and Biological Samples</b>															
Clinical Lab blood sample		X												X	
Research samples (blood + urine)			X	X	X	X	X		X		X		X		
Lumbar puncture			X		X		X		X		X		X		
Skin biopsy <sup>d</sup>			X				X				X				X <sup>c</sup>
<b>Imaging Activities</b>															
Pregnancy Test (prior to DaTscan injection), if applicable		X													
DaTscan Imaging		X													
MRI			X												
<b>Safety and General Health</b>															
# Adverse Events		X	X		X		X		X		X		X	X	
Adverse Event Telephone Assessment		X	X		X		X		X		X		X		
Current Medical Conditions Review		X	X	X	X	X	X	X	X	X	X	X	X	X	
Concomitant Medication Review		X	X	X	X	X	X	X	X	X	X	X	X	X	
Participation in Other Studies		X	X	X	X	X	X	X	X	X	X	X	X	X	X
Report of Pregnancy		As Needed													

I = Investigator completed assessment

P = Participant completed assessment

X = Investigator or Coordinator completed assessment (or as otherwise delegated)

R0X Visits are conducted remotely (e.g., video, audio)

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c = Previously enrolled participants transitioning to new database may be asked to have a skin biopsy. If not done at first visit, may be conducted at a subsequent in person visit.

d = Skin biopsy will be conducted at participating sites.

\*Completed on paper source first, and then scores entered in EDC.

\*\*Window of +45 days either side of Target Visit Date

#Adverse events collected only day of and 2-3 days post DaTscan, LP and skin biopsy per protocol.

## Healthy Control Schedule of Activities (Years 6-13)

Visit Number	R12	V13	R13	V14	R14	V15	R15	V16	R16	V17	R17	V18	R18	V19	R19	V20	Unsched	<sup>b</sup> Transition Activities	
Assessment	**Timepoint	66 mths	72 (Y6)	78 mths	84 (Y7)	90 mths	96 (Y8)	102 mths	108 (Y9)	114 mth	120 (Y10)	126 mths	132 (Y11)	138 mths	144 (Y12)	150 mths	156 (Y13)	---	---
<b>Consent Activities</b>																			
Documentation of Informed Consent		As Needed																	I
Continuing Consent			X		X		X		X		X		X		X		X		
Consent to share contact information		As Needed																	
Research Proxy Designation		As Needed (I)																	
Informed Consent Tracking Log		As Needed																	
<b>General Activities</b>																			
Demographics																			X
Family History																			X
Socio-Economics																			X
Vital Signs + Height and Weight			X		X		X		X		X		X		X		X		X
Visit Status		X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X
Screen Fail																			As Needed
Conclusion of Study Participation		As Needed																	
<b>Neurological/Motor Assessments</b>																			
Participant Motor Function Questionnaire			P		P		P		P		P		P		P		P		
Freezing and Falls			X		X		X		X		X		X		X		X		
Neurological Examination			I		I		I		I		I		I		I		I		I
MDS-UPDRS Part Ia, Part III and Hoehn & Yahr		<sup>a</sup> I	I	<sup>a</sup> I	I	<sup>a</sup> I	I	<sup>a</sup> I	I	<sup>a</sup> I	I	<sup>a</sup> I	I	<sup>a</sup> I	I	<sup>a</sup> I	I		
MDS-UPDRS Part Ib and Part II		P	P	P	P	P	P	P	P	P	P	P	P	P	P	P	P		
Modified Schwab & England ADL		I	I	I	I	I	I	I	I	I	I	I	I	I	I	I	I		
Features of Parkinsonism			I		I		I		I		I		I		I		I		
Other Clinical Features			I		I		I		I		I		I		I		I		
Primary Clinical Diagnosis		I	I	I	I	I	I	I	I	I	I	I	I	I	I	I	I		
<b>Non-Motor Assessments</b>																			
REM Sleep Behavior Disorder Screening Questionnaire			P		P		P		P		P		P		P		P		
Epworth Sleepiness Scale			P		P		P		P		P		P		P		P		
SCOPA-AUT			P		P		P		P		P		P		P		P		
Neuro QoL			P		P		P		P		P		P		P		P		
<b>Cognitive Assessments</b>																			
Montreal Cognitive Assessment*			X		X		X		X		X		X		X		X		
Clock Drawing*			X		X		X		X		X		X		X		X		
Lexical Fluency*			X		X		X		X		X		X		X		X		
Hopkins Verbal Learning Test-Revised*			X		X		X		X		X		X		X		X		
Benton Judgment of Line Orientation*			X		X		X		X		X		X		X		X		
Semantic Fluency (Animals only)*			X		X		X		X		X		X		X		X		
Letter Number Sequencing*			X		X		X		X		X		X		X		X		
Symbol Digit Modalities Test*			X		X		X		X		X		X		X		X		
Trail Making Test (A and B)*			X		X		X		X		X		X		X		X		
Modified Boston Naming Test*			X		X		X		X		X		X		X		X		
Cognitive Change			P		P		P		P		P		P		P		P		
Cognitive Categorization			I		I		I		I		I		I		I		I		
<b>Neuropsychological Assessments</b>																			
State-Trait Anxiety Inventory for Adults			P		P		P		P		P		P		P		P		

## Healthy Control Schedule of Activities (Years 6-13)

Visit Number	R12	V13	R13	V14	R14	V15	R15	V16	R16	V17	R17	V18	R18	V19	R19	V20	Unsched	<sup>b</sup> Transition Activities
Assessment	**Timepoint 66 mths	72 (Y6)	78 mths	84 (Y7)	90 mths	96 (Y8)	102 mths	108 (Y9)	114 mth	120 (Y10)	126 mths	132 (Y11)	138 mths	144 (Y12)	150 mths	156 (Y13)	---	---
Geriatric Depression Scale		P		P		P		P		P		P		P		P		
QUIP		P		P		P		P		P		P		P		P		
<b>Clinical and Biological Samples</b>																		
Clinical Lab blood sample																		X
Research samples (blood + urine)		X		X		X		X		X		X		X		X		
Lumbar puncture				X				X				X				X		
Skin biopsy <sup>d</sup>																		X <sup>c</sup>
<b>Safety and General Health</b>																		
#Adverse Events				X				X				X				X	X	
Adverse Event Telephone Assessment				X				X				X				X		
Current Medical Conditions Review	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	
Concomitant Medication Review	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	
Participation in Other Studies	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X
Report of Pregnancy	As Needed																	

I = Investigator completed assessment

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#Adverse events collected only day of and 2-3 days post LP and skin biopsy per protocol.