Prodromal Schedule of Activities (Years 6+)

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	Visit Number	R12	V13	R13	V14	R14	V15	R15	V16	R16	V17	R17	V18	R18	V19	R19	Annual	Remote	^H Event Driven Modified Visit
Assessment	**Timepoint	66 mths	72 (Y6)	78 mths	84 (Y7)	90 mths	96 (Y8)	102 mths	108 (Y9)	114 mths	120 (Y10)	126 mths	132 (Y11)	138 mths	144 (Y12)	150 mths	156+ (Y13+)	162 mths+	
Consent Activities																		•	
Documentation of Informed Consent		As Needed																	
Continuing Consent			X		X		X		X		X		X		X		X		
Consent to share contact information				•	•	•			A	As Neede	d			•	•	•	•		
Research Proxy Designation									A	As Neede	ed								
Informed Consent Tracking Log		As Needed																	
General Activities																			
myPPMI Registration ^S		As Needed																	
Vital Signs + Height and Weight			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	
Conclusion of Study Participation		As Needed																	
Neurological/Motor Assessments																			
Participant Motor Function Questio	onnaire		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		
Initiation of Dopaminergic Therapy		X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	
MDS-UPDRS Part Ia			I		I		I		I		I		I		I		I		
MDS-UPDRS Part III Treatment Determination/Motor Exam/Hoehn & Yahr ^{d,c}			I		I		I		I		I		I		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	
MDS-UPDRS Repeat Part III/Hoehn & Yahr ^{d,e}			I		I		I		I		I		I		I		I		
MDS-UPDRS Part IV ^d			I		I		I		I		I		I		I		I		
Modified Schwab & England ADL			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 Research Diagnosis			I		I		I		I		I		I		I		I		
Clinical Global Impression (CGI)			I		I		I		I		I		I		I		I		
Clinical Diagnosis		X	Х	Х	X	Х	X	X	X	Х	Х	X	X	Х	Х	X	X	X	
Non-Motor Assessments																			
REM Sleep Behavior Disorder Scre	ening Questionnaire		P		P		Р		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		
Participant Global Impression (PGI))		P		P		P		P		P		P		P		P		
PDAQ-27			P		P		P		P		P		P		P		P		
Neuro QoL			P		P		P		P		P		P		P		P		
Cognitive Assessments																			
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-Revi	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		
Modified Semantic Fluency (Anima	ils 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		

Prodromal Schedule of Activities (Years 6+)

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Assessment	**Timepoint	66 mths	72 (Y6)	78 mths	84 (Y7)	90 mths	96 (Y8)	102 mths	108 (Y9)	114 mths	120 (Y10)	126 mths	132 (Y11)	138 mths	144 (Y12)	150 mths	156+ (Y13+)	162 mths+	
Modified Boston Naming Test*			X		X		X		X		Х		X		X		X		
Cognitive Change			P		P		P		P		P		P		P		P		
Cognitive Categorization			I		I		I		I		I		I		I		I		
Neuropsychological Assessments																			
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		
Clinical and Biological Samples		•		•		•	•		•						•		•		
Research Biosamples (blood + urine)			X		X		X		X		Х		X		X		Х		
Lumbar puncture					Х				X				X				X^L		
Skin biopsy					X														
Imaging Activities																			
Pregnancy Test (prior to tracer injection), if application	ble		X				X				X								
Dopamine Imaging			X				X				X								
MRI			X				X				X								
Safety and General Health																			
*Adverse Events			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	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			As Needed																
LEDD Concomitant Medication Log			As Needed																
Procedure for PD Log	rocedure for PD Log As Needed																		
Report of Pregnancy									A	As Neede	ed								
I = Investigator (or trained designee) completed assessment																		

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

As needed assessments can be located under the Event Driven category in EDC

 $P = Participant \ completed \ assessment$

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

 $[\]label{eq:definition} d = Investigator \ or \ Coordinator \ may \ complete \ treatment \ and \ timing \ information.$

 $e = If \ the \ participant \ is \ on \ levodopa, \ dopamine \ agonists, \ or \ has \ had \ DBS, \ the \ MDS-UPDRS \ Part \ III \ should \ be \ performed \ in \ the \ OFF \ and \ ON \ state.$

H= see protocol section 11 for modification of visit schedule due to New Clinical Diagnosis, Need for PD Therapy or withdrawal from study

L= Cadence should continue to follow every 2 years.

S= Site to inform about myPPMI and assist with registration, if not yet done.

^{*}Completed on paper source first, and then scores entered in to EDC.

^{**}Window of +45 days either side of Target Visit Date

 $^{\#\} Adverse\ events\ collected\ only\ day\ of\ and\ 2-3\ business\ days\ post\ Dopamine\ Imaging,\ LP\ and\ skin\ biopsy\ per\ protocol.$