

BioFIND Schedule of Activities						
PARKINSON DISEASE (PD) SUBJECTS						
Visit Description	Level #	Baseline/V01 Day 0	V02 Day 14	Telephone Call T01 ⁶	FNL	Unscheduled Visit ⁵
Confidential Subject Identification Log		X				
Written Informed Consent		X				
Screening/Demographics	02	X				
Socio-Economics	04	X				
CTCC Unique ID	06	X				
Inclusion/Exclusion - PD	08	X				
PD Features	10	X				
Primary Diagnosis	12	X				
Medical History (General)	14	X				
Smoking and Alcohol Questionnaire	15	X				
Family History (PD)	16	X				
General Neurological Exam	18	X				
Vital Signs	20	X	X			X
Fasting Status	21	X	X			
Use of PD Medication	22	X	X			
MDS-UPDRS 1	24	X				
MDS-UPDRS 2	24	X				
MDS-UPDRS 3/Hoehn & Yahr	24	X	X ¹			
MDS-UPDRS 4	24	X				
Modified Schwab & England ADL	26	X				
Montreal Cognitive Assessment (MoCA)	30	X				
REM Sleep Disorder Questionnaire	32	X				
DNA Sample	34	X				
Laboratory Procedures	36	X ²	X ⁴			
Saliva and Urine Samples	37		X			
Clinical Labs	38	X ³				
Lumbar Puncture	40		X			
Investigator Signature	42	X	X	X		X
Visit Status	44	X	X	X		X
Adverse Event Log	46		X	X		
Concomitant Medication Log	50	X	X	X		X
Adverse Event Follow-up Log ⁷	54			X		
Conclusion of Study Participation	52				X	

1 MDS-UPDRS part 3 motor only
2 10ml plasma between 1-3 hours after meds
3 CBC, platelet count, PT/PTT
4 Laboratory procedures include PAXgene™ and plasma EDTA purple top
5 Assessments at the discretion of the Investigator and to be recorded in source document
6 Adverse events assessed by phone 7 - 10 days following LP
7 Any AE ongoing at the 7 to 10 day reporting telephone visit should be followed until resolution or stabilization, but not more than 30 days from lumbar puncture.

BioFIND Schedule of Activities						
HEALTHY CONTROL (HC) SUBJECTS						
Visit Description	Level #	Baseline/V01 Day 0	V02 Day 14	Telephone Call T01 ⁶	FNL	Unscheduled Visit ⁵
Confidential Subject Identification Log		X				
Written Informed Consent		X				
Screening/Demographics	02	X				
Socio-Economics	04	X				
CTCC Unique ID	06	X				
Inclusion/Exclusion - HC	09	X				
Primary Diagnosis	12	X				
Medical History (General)	14	X				
Smoking and Alcohol Questionnaire	15	X				
Family History (PD)	16	X				
General Neurological Exam	18	X				
Vital Signs	20	X	X			X
Fasting Status	21	X	X			
MDS-UPDRS 3/Hoehn & Yahr	24	X ¹				
Montreal Cognitive Assessment (MoCA)	30	X				
REM Sleep Disorder Questionnaire	32	X				
DNA Sample	34	X				
Laboratory Procedures	36	X ²	X ⁴			
Saliva and Urine Samples	37		X			
Clinical Labs	38	X ³				
Lumbar Puncture	40		X			
Investigator Signature	42	X	X	X		X
Visit Status	44	X	X	X		X
Adverse Event Log	46		X	X		
Concomitant Medication Log	50	X	X	X		X
Adverse Event Follow-up Log ⁷	54			X		
Conclusion of Study Participation	52				X	

1 MDS-UPDRS part 3 motor only
2 10ml plasma between 1-3 hours after meds
3 CBC, platelet count, PT/PTT
4 Laboratory procedures include PAXgene™ and plasma EDTA purple top
5 Assessments at the discretion of the Investigator and to be recorded in source document
6 Adverse events assessed by phone 7 - 10 days following LP
7 Any AE ongoing at the 7 to 10 day reporting telephone visit should be followed until resolution or stabilization, but not more than 30 days from lumbar puncture.