

**[ACRONYM] PROTOCOL SYNOPSIS (SAMPLE)**  
**(Prepared by the [SG NAME] with the support of [SPONSOR])**

Protocol Number	Provide the protocol number identifier.
Protocol Title	Provide the title from the protocol.
Acronym/Title	Provide the acronym and title upon which the acronym is based
Clinical Phase	Specify Phase I, Phase II, Phase III, or Phase IV.
Investigators	Specify Parkinson Study Group (PSG), Huntington Study Group (HSG), or other Investigator(s).
Study Centers	Specify the number of study sites.
Study Period	Specify the length of the study.
Study Objective	Provide a brief description of the study objective from the protocol.
Study Population	Provide a brief description of the study population.
Study Design	Provide a brief summation of the design of the study using information regarding number of sites, blinding, dosage escalation, length of study, treatment course, patient groups, etc.
Number of Subjects	Provide number of subjects required for the study.
Main Eligibility/ Exclusion Criteria	Specify the subject diagnosis and all Main Inclusion/ Exclusion Criteria
Route and Dosage Form	Specify the route (oral, IV, etc.) and dosage form (tablets, caplets, placebo, etc.).
Dosage	Specify dosage (i.e., 150 mg QD).
Duration of Treatment	Specify the length of treatment course.
Primary Outcome Measure(s)	Provide a brief description of any Primary Outcome(s) to be studied.
Secondary Outcome Measure	Provide a brief description of any Secondary Outcome(s) to be studied.
Sample Size Considerations	Briefly describe any Sample Size issues or considerations.

**SCHEDULE OF ACTIVITIES (SAMPLE)**

	<b>Screening</b>	<b>Baseline</b>	<b>Maintenance Phase</b>				<b>Post-Drug Evaluation</b>
	<b>Visit SC Week-2 (-14d)</b>	<b>Visit BL Week 0 (Day 0)</b>	<b>Visit 1 (10d+3d)</b>	<b>Visit 2 (10d+3d)</b>	<b>Visit 3 (10d+3d)</b>	<b>Visit 4 (10d+3d)</b>	<b>Visit 5 (10d+3d)</b>
<b>Written Informed Consent</b>							
<b>Subject Entry Number</b>							
<b>Eligibility Criteria</b>							
<b>Med/Neuro History</b>							
<b>Physical Exam</b>							
<b>Hoehn &amp; Yahr Scale</b>							
<b>UPDRS II-IV</b>							
<b>CGI</b>							
<b>MMSE</b>							
<b>Beck Depression Inventory</b>							
<b>Vital Signs</b>							
<b>ECG-12 LEAD</b>							
<b>Standard Clinical Labs*</b>							
<b>Pregnancy Test</b>							
<b>Adverse Experience Information</b>							
<b>Concomitant Drug Therapy</b>							
<b>Instruct and Supply Home Diary Card</b>							
<b>Review Home Diary Card</b>							
<b>Dispense Study Drug</b>							
<b>Check Drug Compliance</b>							

\*Specific analyses may be listed here at the bottom of the page.