PPMI

1 3 2 **CONCLUSION OF STUDY PARTICIPATION** 7 4

SUBJECT ID | | | | | VISIT NO | F | N | L |

INITIALS | | | | SITE NO | | | | VISIT DATE | | | MM | | | DD | | | | | YYYY |

2. Did the subject complete the study? (00 = No, 01 = Yes) 2. | | |

If subject prematurely withdrew: 4. | | |

4. What was the primary reason for withdrawal:
   01 = Adverse Event (complete AE Log)
   02 = Lost to Follow-up
   03 = Subject withdrew consent (specify in 4a)
   04 = Pregnancy
   05 = Protocol violation
   06 = Death of subject
   07 = Investigator decision (specify in 4a)

   09 = Clinical Monitor decision (specify in 4a)
   10 = Sponsor decision (specify in 4a)
   11 = Primary Care Physician decision (specify in 4a)
   12 = Informant/Caregiver decision (specify in 4a)
   13 = Institutionalized
   14 = Inability to continue giving consent
   15 = Other (specify in 4a)

4a. Specify: _____

5. Date of premature withdrawal:
   (Date investigator deemed the subject would no longer participate in the study)

5. | | | MM | | | DD | | | | | YYYY |