1. Indicates that participant/subject prematurely discontinued study intervention:

[ ]  Yes [ ]  No [ ]  Unknown

1. Primary reason participant/subject discontinued study intervention (choose all that apply):

[ ]  Adverse event

[ ]  Other clinical decision (e.g., investigator decision, primary care provider decision, etc.) OR other reason specified by the protocol (i.e., institutionalization, pregnancy, etc.)

[ ]  Death

[ ]  Participant's/Subject's decision (e.g., unwilling/unable to commit time and/or resources, moved from area, etc.)

[ ]  Lost to follow-up

[ ]  Other, specify: