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: