Summary of Adverse Events
Weeks 0 to 12
All Participants as Treated

	Placebo		Xanomeline Low Dose		Xanomeline High Dose		Total	
	n	(%)	n	(%)	n	(%)	n	(%)
Participants in population	86		84		84		254	
with one or more adverse events	69	(80.2)	77	(91.7)	79	(94.0)	225	(88.6)
with drug-related[a] adverse events	44	(51.2)	73	(86.9)	70	(83.3)	187	(73.6)
with serious adverse events	0	(0.0)	1	(1.2)	2	(2.4)	3	(1.2)

[a]Determined by the investigator to be related to the drug.

Source: [CDISCpilot: adam-adsl; adae]