Listing of Participants With Serious Adverse Events
Weeks 0 to 12
All Participants as Treated

USUBJID	ASTDY	Adverse Event	Duration	Intensity	Serious	Related	Action Taken	Outcome
Low Dose								
Subject ID = 01-718-1170, Gender = F, Race = WHITE, AGE = 80 Years, TRT = Xanomeline Low Dose								
01-718-1170	27	SYNCOPE	2 Day	SEVERE	Y	Probable	None	Resolved
High Dose								
Subject ID = 01-709-1424, Gender = M, Race = WHITE, AGE = 77 Years, TRT = Xanomeline High Dose								

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High Dose								
Subject ID = 01-709-1424, Gender = M, Race = WHITE, AGE = 77 Years, TRT = Xanomeline High Dose								
01-709-1424	5	SYNCOPE	1 Day	MODERATE	Y	Possible	None	Resolved
Subject ID = 01-718-1371, Gender = F, Race = WHITE, AGE = 69 Years, TRT = Xanomeline High Dose								
01-718-1371	38	PARTIAL SEIZURES WITH SECONDARY GENERALISATION	4 Day	SEVERE	Y	None	None	Resolved

Related: Investigator-assessed relationship of the adverse event to study medication. Y = RELATED, N = NOT RELATED
Action Taken: Discontinued = DRUG WITHDRAWN, Interrupted = DRUG INTERRUPTED, Reduced = DOSE REDUCED, Increased = DOSE INCREASED, None = DOSE NOT CHANGED, N/A = NOT APPLICABLE.
Outcome: Resolved = RECOVERED/RESOLVED, Resolving = RECOVERING/RESOLVING, Sequelae = RECOVERED/RESOLVED WITH SEQUELAE, Not resolved = NOT RECOVERED/NOT RESOLVED.
Adverse event terms are from MedDRA Version 25.0.

Source: [CDISCpilot: adam-adsl; adae]