Adverse Events Summary by Treatment

USUBJID	TRTA	AEDECOD	AESEV	AESER	AEREL
01-701-1015	Placebo	APPLICATION SITE ERYTHEMA	MILD	N	PROBABLE
01-701-1015	Placebo	APPLICATION SITE PRURITUS	MILD	N	PROBABLE
01-701-1015	Placebo	DIARRHOEA	MILD	N	REMOTE
01-701-1023	Placebo	ERYTHEMA	MILD	N	POSSIBLE
01-701-1023	Placebo	ERYTHEMA	MODERATE	N	PROBABLE
01-701-1023	Placebo	ATRIOVENTRICULAR BLOCK SECOND DEGREE	MILD	N	POSSIBLE
01-701-1023	Placebo	ERYTHEMA	MILD	N	POSSIBLE
01-701-1028	Xanomeline High Dose	APPLICATION SITE ERYTHEMA	MILD	N	POSSIBLE
01-701-1028	Xanomeline High Dose	APPLICATION SITE PRURITUS	MILD	N	PROBABLE
01-701-1034	Xanomeline High Dose	APPLICATION SITE PRURITUS	MILD	N	PROBABLE
01-701-1034	Xanomeline High Dose	FATIGUE	MILD	N	POSSIBLE
01-701-1047	Placebo	HIATUS HERNIA	MODERATE	N	NONE
01-701-1047	Placebo	HIATUS HERNIA	MODERATE	N	NONE

USUBJID	TRTA	AEDECOD	AESEV	AESER	AEREL
01-701-1047	Placebo	UPPER RESPIRATORY TRACT INFECTION	MILD	N	NONE
01-701-1047	Placebo	BUNDLE BRANCH BLOCK LEFT	MILD	N	NONE
01-701-1097	Xanomeline Low Dose	ERYTHEMA	MILD	N	POSSIBLE
01-701-1097	Xanomeline Low Dose	PRURITUS GENERALISED	MODERATE	N	POSSIBLE
01-701-1097	Xanomeline Low Dose	APPLICATION SITE VESICLES	MILD	N	PROBABLE
01-701-1097	Xanomeline Low Dose	APPLICATION SITE PRURITUS	MILD	N	POSSIBLE
01-701-1097	Xanomeline Low Dose	PRURITUS GENERALISED	MODERATE	N	POSSIBLE
01-701-1097	Xanomeline Low Dose	PRURITUS GENERALISED	MODERATE	N	POSSIBLE
01-701-1097	Xanomeline Low Dose	PRURITUS GENERALISED	MODERATE	N	POSSIBLE
01-701-1097	Xanomeline Low Dose	PHARYNGOLARYNGEAL PAIN	MILD	N	NONE
01-701-1097	Xanomeline Low Dose	NASAL CONGESTION	MILD	N	NONE
01-701-1097	Xanomeline Low Dose	APPLICATION SITE PRURITUS	MODERATE	N	POSSIBLE
01-701-1111	Xanomeline Low Dose	ERYTHEMA	MILD	N	NONE

USUBJID	TRTA	AEDECOD	AESEV	AESER	AEREL
01-701-1111	Xanomeline Low Dose	PRURITUS	MILD	N	NONE
01-701-1111	Xanomeline Low Dose	LOCALISED INFECTION	MODERATE	N	NONE
01-701-1111	Xanomeline Low Dose	ERYTHEMA	MILD	N	NONE
01-701-1111	Xanomeline Low Dose	PRURITUS	MILD	N	NONE

Abbreviations: USUBJID=Subject ID, TRTA=Treatment, AEDECOD=Adverse Event, AESEV=Severity, AESER=Serious, AEREL=Related

Source: ADAE Dataset from Clinical Trial Database