Listing of Subjects With Series Adverse Events
ASaT

Trial Number: xxxx

Subject ID	Rel Day of Onset	Aderse Event	Duration	Intensity	Serious	Related	Action Taken	Outcome
Placebo								
Trial Number = CDISCPILOT01, Site Number = 701, Subject ID = 01-701-1015, Gender = F, Race = WHITE, AGE = 63 Years								
01-701-1015	2	APPLICATION SITE ERYTHEMA	NA	MILD	N	PROBABLE		NOT RECOVERED/NOT RESOLVED
		APPLICATION SITE PRURITUS	NA	MILD	N	PROBABLE		NOT RECOVERED/NOT RESOLVED
	8	DIARRHOEA	3 DAY	MILD	N	REMOTE		RECOVERED/RESOLVED
Trial Number = CDISCPILOT01, Site Number = 701, Subject ID = 01-701-1023, Gender = M, Race = WHITE, AGE = 64 Years								
01-701-1023	3	ERYTHEMA	24 DAY	MILD	N	POSSIBLE		NOT RECOVERED/NOT RESOLVED
		ERYTHEMA	NA	MODERATE	N	PROBABLE		NOT RECOVERED/NOT RESOLVED

Listing of Subjects With Series Adverse Events
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Trial Number: xxxx

Subject ID	Rel Day of Onset	Aderse Event	Duration	Intensity	Serious	Related	Action Taken	Outcome
Placebo								
Trial Number = CDISCPILOT01, Site Number = 701, Subject ID = 01-701-1023, Gender = M, Race = WHITE, AGE = 64 Years								
01-701-1023	3	ERYTHEMA	24 DAY	MILD	N	POSSIBLE		RECOVERED/RESOLVED
	22	ATRIOVENTRICULAR BLOCK SECOND DEGREE	NA	MILD	N	POSSIBLE		NOT RECOVERED/NOT RESOLVED
Trial Number = CDISCPILOT01, Site Number = 701, Subject ID = 01-701-1047, Gender = F, Race = WHITE, AGE = 85 Years								
01-701-1047	1	HIATUS HERNIA	1 DAY	MODERATE	N	NONE		NOT RECOVERED/NOT RESOLVED
		HIATUS HERNIA	1 DAY	MODERATE	N	NONE		RECOVERED/RESOLVED
	23	UPPER RESPIRATORY TRACT INFECTION	NA	MILD	N	NONE		NOT RECOVERED/NOT RESOLVED
	27	BUNDLE BRANCH BLOCK LEFT	NA	MILD	N	NONE		NOT RECOVERED/NOT RESOLVED

Listing of Subjects With Series Adverse Events
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Trial Number: xxxx

Subject ID	Rel Day of Onset	Aderse Event	Duration	Intensity	Serious	Related	Action Taken	Outcome
Placebo								
Trial Number = CDISCPILOT01, Site Number = 701, Subject ID = 01-701-1118, Gender = M, Race = WHITE, AGE = 52 Years								
01-701-1118		COUGH	NA	MILD	N	NONE		NOT RECOVERED/NOT RESOLVED
Trial Number = CDISCPILOT01, Site Number = 701, Subject ID = 01-701-1130, Gender = M, Race = WHITE, AGE = 84 Years								
01-701-1130	23	URINARY TRACT INFECTION	8 DAY	MILD	N	NONE		NOT RECOVERED/NOT RESOLVED
		PYREXIA	1 DAY	MILD	N	NONE		RECOVERED/RESOLVED
		URINARY TRACT INFECTION	8 DAY	MILD	N	NONE		RECOVERED/RESOLVED
	84	EYE SWELLING	1 DAY	MILD	N	NONE		RECOVERED/RESOLVED
		EYE ALLERGY	1 DAY	MILD	N	NONE		RECOVERED/RESOLVED
	97	EYE PRURITUS	NA	MILD	N	REMOTE		NOT RECOVERED/NOT RESOLVED
		PRURITUS	NA	MILD	N	REMOTE		NOT RECOVERED/NOT RESOLVED

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Trial Number: xxxx

Subject ID	Rel Day of Onset	Aderse Event	Duration	Intensity	Serious	Related	Action Taken	Outcome
Placebo								
Trial Number = CDISCPILOT01, Site Number = 701, Subject ID = 01-701-1130, Gender = M, Race = WHITE, AGE = 84 Years								
01-701-1130	97	NASAL CONGESTION	NA	MILD	N	REMOTE		NOT RECOVERED/NOT RESOLVED
Trial Number = CDISCPILOT01, Site Number = 701, Subject ID = 01-701-1153, Gender = F, Race = WHITE, AGE = 79 Years								
01-701-1153	21	INCREASED APPETITE	25 DAY	MILD	N	POSSIBLE		NOT RECOVERED/NOT RESOLVED
		INCREASED APPETITE	25 DAY	MILD	N	POSSIBLE		RECOVERED/RESOLVED
Xanomeline High Dose								
Trial Number = CDISCPILOT01, Site Number = 701, Subject ID = 01-701-1028, Gender = M, Race = WHITE, AGE = 71 Years								
01-701-1028	3	APPLICATION SITE ERYTHEMA	NA	MILD	N	POSSIBLE		NOT RECOVERED/NOT RESOLVED
	21	APPLICATION SITE PRURITUS	NA	MILD	N	PROBABLE		NOT RECOVERED/NOT RESOLVED

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Trial Number: xxxx

Subject ID	Rel Day of Onset	Aderse Event	Duration	Intensity	Serious	Related	Action Taken	Outcome
Xanomeline High Dose								
Trial Number = CDISCPILOT01, Site Number = 701, Subject ID = 01-701-1034, Gender = F, Race = WHITE, AGE = 77 Years								
01-701-1034	58	APPLICATION SITE PRURITUS	NA	MILD	N	PROBABLE		NOT RECOVERED/NOT RESOLVED
	125	FATIGUE	NA	MILD	N	POSSIBLE		NOT RECOVERED/NOT RESOLVED
Trial Number = CDISCPILOT01, Site Number = 701, Subject ID = 01-701-1133, Gender = F, Race = WHITE, AGE = 81 Years								
01-701-1133	61	APPLICATION SITE ERYTHEMA	NA	MILD	N	PROBABLE		NOT RECOVERED/NOT RESOLVED
		APPLICATION SITE PRURITUS	NA	MILD	N	PROBABLE		NOT RECOVERED/NOT RESOLVED
		APPLICATION SITE ERYTHEMA	NA	MODERATE	N	PROBABLE		NOT RECOVERED/NOT RESOLVED
		APPLICATION SITE PRURITUS	NA	MODERATE	N	PROBABLE		NOT RECOVERED/NOT RESOLVED

Listing of Subjects With Series Adverse Events
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Trial Number: xxxx

Subject ID	Rel Day of Onset	Aderse Event	Duration	Intensity	Serious	Related	Action Taken	Outcome
Xanomeline High Dose								
Trial Number = CDISCPILOT01, Site Number = 701, Subject ID = 01-701-1146, Gender = F, Race = WHITE, AGE = 75 Years								
01-701-1146	-4	RASH	18 DAY	MILD	N	NONE		NOT RECOVERED/NOT RESOLVED
		RASH	18 DAY	MILD	N	NONE		RECOVERED/RESOLVED
	13	APPLICATION SITE IRRITATION	2 DAY	MILD	N	PROBABLE		RECOVERED/RESOLVED
	15	FATIGUE	NA	MILD	N	POSSIBLE		NOT RECOVERED/NOT RESOLVED
	22	APPLICATION SITE ERYTHEMA	NA	MILD	N	PROBABLE		NOT RECOVERED/NOT RESOLVED
		APPLICATION SITE PRURITUS	NA	MILD	N	PROBABLE		NOT RECOVERED/NOT RESOLVED

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Trial Number: xxxx

Subject ID	Rel Day of Onset	Aderse Event	Duration	Intensity	Serious	Related	Action Taken	Outcome
Xanomeline High Dose								
Trial Number = CDISCPILOT01, Site Number = 701, Subject ID = 01-701-1146, Gender = F, Race = WHITE, AGE = 75 Years								
01-701-1146	22	APPLICATION SITE ERYTHEMA	NA	MODERATE	N	PROBABLE		NOT RECOVERED/NOT RESOLVED
		APPLICATION SITE PRURITUS	NA	MODERATE	N	PROBABLE		NOT RECOVERED/NOT RESOLVED
	34	APPLICATION SITE PAIN	NA	MODERATE	N	PROBABLE		NOT RECOVERED/NOT RESOLVED
	38	APPLICATION SITE VESICLES	NA	MODERATE	N	PROBABLE		NOT RECOVERED/NOT RESOLVED
		APPLICATION SITE IRRITATION	NA	MODERATE	N	PROBABLE		NOT RECOVERED/NOT RESOLVED
Trial Number = CDISCPILOT01, Site Number = 701, Subject ID = 01-701-1148, Gender = M, Race = WHITE, AGE = 57 Years								
01-701-1148	-569	DYSPEPSIA	NA	MODERATE	N	NONE		NOT RECOVERED/NOT RESOLVED

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Trial Number: xxxx

Subject ID	Rel Day of Onset	Aderse Event	Duration	Intensity	Serious	Related	Action Taken	Outcome
Xanomeline High Dose								
Trial Number = CDISCPILOT01, Site Number = 701, Subject ID = 01-701-1148, Gender = M, Race = WHITE, AGE = 57 Years								
01-701-1148	-25	DEPRESSED MOOD	NA	MODERATE	N	NONE		NOT RECOVERED/NOT RESOLVED
	3	APPLICATION SITE ERYTHEMA	NA	MILD	N	PROBABLE		NOT RECOVERED/NOT RESOLVED
		APPLICATION SITE PRURITUS	NA	MILD	N	PROBABLE		NOT RECOVERED/NOT RESOLVED
	51	LOWER RESPIRATORY TRACT INFECTION	30 DAY	MODERATE	N	NONE		NOT RECOVERED/NOT RESOLVED
		LOWER RESPIRATORY TRACT INFECTION	30 DAY	MODERATE	N	NONE		RECOVERED/RESOLVED
	115	FLANK PAIN	3 DAY	MODERATE	N	NONE		RECOVERED/RESOLVED
	117	CALCULUS URETHRAL	1 DAY	MODERATE	N	NONE		RECOVERED/RESOLVED
	134	EPISTAXIS	1 DAY	MILD	N	NONE		RECOVERED/RESOLVED

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Trial Number: xxxx

Subject ID	Rel Day of Onset	Aderse Event	Duration	Intensity	Serious	Related	Action Taken	Outcome
Xanomeline High Dose								
Trial Number = CDISCPILOT01, Site Number = 701, Subject ID = 01-701-1148, Gender = M, Race = WHITE, AGE = 57 Years								
01-701-1148	174	ACTINIC KERATOSIS	NA	MILD	N	NONE		NOT RECOVERED/NOT RESOLVED
Trial Number = CDISCPILOT01, Site Number = 701, Subject ID = 01-701-1180, Gender = M, Race = WHITE, AGE = 56 Years								
01-701-1180	1	VOMITING	1 DAY	MILD	N	NONE		NOT RECOVERED/NOT RESOLVED
		VOMITING	1 DAY	MILD	N	NONE		RECOVERED/RESOLVED
	4	MICTURITION URGENCY	33 DAY	MILD	N	POSSIBLE		NOT RECOVERED/NOT RESOLVED
		MICTURITION URGENCY	33 DAY	MILD	N	POSSIBLE		RECOVERED/RESOLVED
	23	NASOPHARYNGITIS	2 DAY	MILD	N	NONE		RECOVERED/RESOLVED
	36	APPLICATION SITE VESICLES	NA	MODERATE	N	POSSIBLE		NOT RECOVERED/NOT RESOLVED

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Trial Number: xxxx

Subject ID	Rel Day of Onset	Aderse Event	Duration	Intensity	Serious	Related	Action Taken	Outcome
Xanomeline High Dose								
Trial Number = CDISCPILOT01, Site Number = 701, Subject ID = 01-701-1180, Gender = M, Race = WHITE, AGE = 56 Years								
01-701-1180	36	APPLICATION SITE PRURITUS	NA	MODERATE	N	POSSIBLE		NOT RECOVERED/NOT RESOLVED
		APPLICATION SITE ERYTHEMA	NA	MODERATE	N	POSSIBLE		NOT RECOVERED/NOT RESOLVED
		HEADACHE	NA	MODERATE	N	NONE		NOT RECOVERED/NOT RESOLVED
Trial Number = CDISCPILOT01, Site Number = 701, Subject ID = 01-701-1181, Gender = F, Race = WHITE, AGE = 79 Years								
01-701-1181	5	AGITATION	1 DAY	MODERATE	N	NONE		RECOVERED/RESOLVED
Xanomeline Low Dose								
Trial Number = CDISCPILOT01, Site Number = 701, Subject ID = 01-701-1097, Gender = M, Race = WHITE, AGE = 68 Years								
01-701-1097	3	ERYTHEMA	NA	MILD	N	POSSIBLE		NOT RECOVERED/NOT RESOLVED
	51	PRURITUS GENERALISED	1 DAY	MODERATE	N	POSSIBLE		RECOVERED/RESOLVED

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Trial Number: xxxx

Subject ID	Rel Day of Onset	Aderse Event	Duration	Intensity	Serious	Related	Action Taken	Outcome
Xanomeline Low Dose								
Trial Number = CDISCPILOT01, Site Number = 701, Subject ID = 01-701-1097, Gender = M, Race = WHITE, AGE = 68 Years								
01-701-1097	51	APPLICATION SITE VESICLES	3 DAY	MILD	N	PROBABLE		RECOVERED/RESOLVED
	52	APPLICATION SITE PRURITUS	NA	MILD	N	POSSIBLE		NOT RECOVERED/NOT RESOLVED
		APPLICATION SITE PRURITUS	NA	MODERATE	N	POSSIBLE		NOT RECOVERED/NOT RESOLVED
	80	PRURITUS GENERALISED	1 DAY	MODERATE	N	POSSIBLE		RECOVERED/RESOLVED
	90	PRURITUS GENERALISED	1 DAY	MODERATE	N	POSSIBLE		RECOVERED/RESOLVED
	109	PRURITUS GENERALISED	2 DAY	MODERATE	N	POSSIBLE		RECOVERED/RESOLVED

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Trial Number: xxxx

Subject ID	Rel Day of Onset	Aderse Event	Duration	Intensity	Serious	Related	Action Taken	Outcome
Xanomeline Low Dose								
Trial Number = CDISCPILOT01, Site Number = 701, Subject ID = 01-701-1097, Gender = M, Race = WHITE, AGE = 68 Years								
01-701-1097	109	PHARYNGOLARYNGEAL PAIN	4 DAY	MILD	N	NONE		RECOVERED/RESOLVED
		NASAL CONGESTION	4 DAY	MILD	N	NONE		RECOVERED/RESOLVED
Trial Number = CDISCPILOT01, Site Number = 701, Subject ID = 01-701-1111, Gender = F, Race = WHITE, AGE = 81 Years								
01-701-1111	-61	LOCALISED INFECTION	NA	MODERATE	N	NONE		NOT RECOVERED/NOT RESOLVED
	-5	ERYTHEMA	6 DAY	MILD	N	NONE		NOT RECOVERED/NOT RESOLVED
		PRURITUS	6 DAY	MILD	N	NONE		NOT RECOVERED/NOT RESOLVED
		ERYTHEMA	6 DAY	MILD	N	NONE		RECOVERED/RESOLVED
		PRURITUS	6 DAY	MILD	N	NONE		RECOVERED/RESOLVED
	1	MICTURITION URGENCY	NA	MILD	N	NONE		NOT RECOVERED/NOT RESOLVED

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Trial Number: xxxx

Subject ID	Rel Day of Onset	Aderse Event	Duration	Intensity	Serious	Related	Action Taken	Outcome
Xanomeline Low Dose								
Trial Number = CDISCPILOT01, Site Number = 701, Subject ID = 01-701-1111, Gender = F, Race = WHITE, AGE = 81 Years								
01-701-1111	7	ARTHRALGIA	NA	MODERATE	N	NONE		NOT RECOVERED/NOT RESOLVED
		CELLULITIS	NA	MODERATE	N	NONE		NOT RECOVERED/NOT RESOLVED
Trial Number = CDISCPILOT01, Site Number = 701, Subject ID = 01-701-1115, Gender = M, Race = WHITE, AGE = 84 Years								
01-701-1115	3	APPLICATION SITE ERYTHEMA	NA	MILD	N	PROBABLE		NOT RECOVERED/NOT RESOLVED
		APPLICATION SITE PRURITUS	NA	MILD	N	PROBABLE		NOT RECOVERED/NOT RESOLVED
		APPLICATION SITE ERYTHEMA	NA	MODERATE	N	PROBABLE		NOT RECOVERED/NOT RESOLVED
		APPLICATION SITE PRURITUS	NA	MODERATE	N	PROBABLE		NOT RECOVERED/NOT RESOLVED

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Trial Number: xxxx

Subject ID	Rel Day of Onset	Aderse Event	Duration	Intensity	Serious	Related	Action Taken	Outcome
Xanomeline Low Dose								
Trial Number = CDISCPILOT01, Site Number = 701, Subject ID = 01-701-1115, Gender = M, Race = WHITE, AGE = 84 Years								
01-701-1115	27	APPLICATION SITE IRRITATION	1 DAY	MILD	N	POSSIBLE		NOT RECOVERED/NOT RESOLVED
		APPLICATION SITE IRRITATION	1 DAY	MILD	N	POSSIBLE		RECOVERED/RESOLVED
	46	FATIGUE	NA	MODERATE	N	REMOTE		NOT RECOVERED/NOT RESOLVED
	51	APPLICATION SITE IRRITATION	NA	MODERATE	N	PROBABLE		NOT RECOVERED/NOT RESOLVED
		APPLICATION SITE DERMATITIS	NA	MODERATE	N	PROBABLE		NOT RECOVERED/NOT RESOLVED
Trial Number = CDISCPILOT01, Site Number = 701, Subject ID = 01-701-1188, Gender = M, Race = WHITE, AGE = 71 Years								
01-701-1188	2	ERYTHEMA	NA	MILD	N	PROBABLE		NOT RECOVERED/NOT RESOLVED

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Trial Number: xxxx

Subject ID	Rel Day of Onset	Aderse Event	Duration	Intensity	Serious	Related	Action Taken	Outcome
Xanomeline Low Dose								
Trial Number = CDISCPILOT01, Site Number = 701, Subject ID = 01-701-1188, Gender = M, Race = WHITE, AGE = 71 Years								
01-701-1188	2	PRURITUS	NA	MILD	N	PROBABLE		NOT RECOVERED/NOT RESOLVED
		ERYTHEMA	NA	MODERATE	N	PROBABLE		NOT RECOVERED/NOT RESOLVED
		PRURITUS	NA	MODERATE	N	PROBABLE		NOT RECOVERED/NOT RESOLVED
	18	URTICARIA	NA	MILD	N	PROBABLE		NOT RECOVERED/NOT RESOLVED
		URTICARIA	NA	MODERATE	N	PROBABLE		NOT RECOVERED/NOT RESOLVED
	35	APPLICATION SITE URTICARIA	NA	MILD	N	PROBABLE		NOT RECOVERED/NOT RESOLVED

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Trial Number: xxxx

Subject ID	Rel Day of Onset	Aderse Event	Duration	Intensity	Serious	Related	Action Taken	Outcome
Xanomeline Low Dose								
Trial Number = CDISCPILOT01, Site Number = 701, Subject ID = 01-701-1188, Gender = M, Race = WHITE, AGE = 71 Years								
01-701-1188	35	URTICARIA	NA	MILD	N	PROBABLE		NOT RECOVERED/NOT RESOLVED
Trial Number = CDISCPILOT01, Site Number = 701, Subject ID = 01-701-1192, Gender = F, Race = WHITE, AGE = 80 Years								
01-701-1192	-782	COUGH	NA	MODERATE	N	NONE		NOT RECOVERED/NOT RESOLVED
	13	NASAL MUCOSA BIOPSY	1 DAY	MILD	N	NONE		RECOVERED/RESOLVED
	17	SECRETION DISCHARGE	39 DAY	MILD	N	NONE		NOT RECOVERED/NOT RESOLVED
		ERYTHEMA	39 DAY	MILD	N	NONE		NOT RECOVERED/NOT RESOLVED
This is footnote 1 This is footnote 2								

Source: [Study MK9999P001: adam-adae]