Summary of Adverse Events by Treatment Group
Safety Analysis Set

Adverse Events	Placebo (N=86)	Xanomeline High Dose (N=84)	Xanomeline Low Dose (N=84)
ABDOMINAL DISCOMFORT	0	1	0
ABDOMINAL PAIN	1	2	3
ACROCHORDON EXCISION	0	1	0
ACTINIC KERATOSIS	0	1	0
AGITATION	2	1	2
ALCOHOL USE	0	1	0
ALLERGIC GRANULOMATOUS ANGIITIS	0	1	0
ALOPECIA	1	0	0
AMNESIA	0	2	0
ANXIETY	2	0	4
APPLICATION SITE BLEEDING	0	0	1
APPLICATION SITE DERMATITIS	9	12	15
APPLICATION SITE DESQUAMATION	0	0	1
APPLICATION SITE DISCHARGE	0	1	0
APPLICATION SITE DISCOLOURATION	0	0	1

Adverse events are coded using MedDRA version 25.0.
Events are sorted alphabetically by preferred term.

Source: ADAE dataset, Data cutoff: 01JAN2023