Participants with Severe Adverse Events (Incidence \geq 0% in One or More Treatment Groups) (ASaT Population)

]	Placebo		Xanomeline Low Dose		Xanomeline High Dose		Total	
	n	(%)	n	(%)	n	(%)	n	(%)	
Participants in population	86		84		84		254		
with one or more severe AE	7	(8.14)	16	(19.05)	8	(9.52)	31	(12.20)	
with no severe AE	79	(91.86)	68	(80.95)	76	(90.48)	223	(87.80)	
cardiac disorders	3	(3.5)	0	(0.0)	1	(1.2)	4	(1.6)	
atrial fibrillation	0	(0.0)	0	(0.0)	1	(1.2)	1	(0.4)	
atrioventricular block second degree	1	(1.2)	0	(0.0)	0	(0.0)	1	(0.4)	
myocardial infarction	2	(2.3)	0	(0.0)	0	(0.0)	2	(8.0)	
gastrointestinal disorders	0	(0.0)	0	(0.0)	2	(2.4)	2	(0.8)	
gastrointestinal haemorrhage	0	(0.0)	0	(0.0)	1	(1.2)	1	(0.4)	
nausea	0	(0.0)	0	(0.0)	1	(1.2)	1	(0.4)	
general disorders and administration site conditions	0	(0.0)	7	(8.3)	0	(0.0)	7	(2.8)	
application site dermatitis	0	(0.0)	1	(1.2)	0	(0.0)	1	(0.4)	
application site erythema	0	(0.0)	2	(2.4)	0	(0.0)	2	(8.0)	
application site irritation	0	(0.0)	3	(3.6)	0	(0.0)	3	(1.2)	
application site pruritus	0	(0.0)	1	(1.2)	0	(0.0)	1	(0.4)	

Participants with Severe Adverse Events (Incidence \geq 0% in One or More Treatment Groups) (ASaT Population)

	Placebo		Xanomeline Low Dose		Xanomeline High Dose		Total	
	n	(%)	n	(%)	n	(%)	n	(%)
general disorders and administration site conditions	0	(0.0)	7	(8.3)	0	(0.0)	7	(2.8)
application site warmth	0	(0.0)	1	(1.2)	0	(0.0)	1	(0.4)
sudden death	0	(0.0)	1	(1.2)	0	(0.0)	1	(0.4)
infections and infestations	0	(0.0)	1	(1.2)	0	(0.0)	1	(0.4)
nasopharyngitis	0	(0.0)	1	(1.2)	0	(0.0)	1	(0.4)
injury, poisoning and procedural complications	1	(1.2)	0	(0.0)	1	(1.2)	2	(8.0)
hip fracture	1	(1.2)	0	(0.0)	1	(1.2)	2	(8.0)
musculoskeletal and connective tissue disorders	1	(1.2)	1	(1.2)	0	(0.0)	2	(8.0)
arthritis	1	(1.2)	0	(0.0)	0	(0.0)	1	(0.4)
muscle spasms	0	(0.0)	1	(1.2)	0	(0.0)	1	(0.4)
neoplasms benign, malignant and unspecified (incl cysts and polyps)	0	(0.0)	1	(1.2)	1	(1.2)	2	(8.0)
colon cancer	0	(0.0)	1	(1.2)	0	(0.0)	1	(0.4)
prostate cancer	0	(0.0)	0	(0.0)	1	(1.2)	1	(0.4)
nervous system disorders	0	(0.0)	3	(3.6)	4	(4.8)	7	(2.8)
dizziness	0	(0.0)	0	(0.0)	1	(1.2)	1	(0.4)
headache	0	(0.0)	1	(1.2)	0	(0.0)	1	(0.4)
partial seizures with secondary generalisation	0	(0.0)	0	(0.0)	1	(1.2)	1	(0.4)
stupor	0	(0.0)	1	(1.2)	0	(0.0)	1	(0.4)

Participants with Severe Adverse Events (Incidence \geq 0% in One or More Treatment Groups) (ASaT Population)

n 0	(%)	n	(0/)				
0		111	(%)	n	(%)	n	(%)
	(0.0)	3	(3.6)	4	(4.8)	7	(2.8)
0	(0.0)	2	(2.4)	1	(1.2)	3	(1.2)
0	(0.0)	1	(1.2)	1	(1.2)	2	(8.0)
1	(1.2)	1	(1.2)	0	(0.0)	2	(0.8)
0	(0.0)	1	(1.2)	0	(0.0)	1	(0.4)
1	(1.2)	0	(0.0)	0	(0.0)	1	(0.4)
1	(1.2)	0	(0.0)	0	(0.0)	1	(0.4)
1	(1.2)	0	(0.0)	0	(0.0)	1	(0.4)
0	(0.0)	4	(4.8)	1	(1.2)	5	(2.0)
0	(0.0)	1	(1.2)	0	(0.0)	1	(0.4)
0	(0.0)	1	(1.2)	0	(0.0)	1	(0.4)
0	(0.0)	1	(1.2)	1	(1.2)	2	(8.0)
0	(0.0)	1	(1.2)	0	(0.0)	1	(0.4)
	1 0 1 1 1 0 0 0	0 (0.0) 1 (1.2) 0 (0.0) 1 (1.2) 1 (1.2) 1 (1.2) 0 (0.0) 0 (0.0) 0 (0.0) 0 (0.0)	0 (0.0) 1 1 (1.2) 1 0 (0.0) 1 1 (1.2) 0 1 (1.2) 0 1 (1.2) 0 0 (0.0) 4 0 (0.0) 1 0 (0.0) 1 0 (0.0) 1 0 (0.0) 1	0 (0.0) 1 (1.2) 1 (1.2) 1 (1.2) 0 (0.0) 1 (1.2) 1 (1.2) 0 (0.0) 1 (1.2) 0 (0.0) 1 (1.2) 0 (0.0) 0 (0.0) 4 (4.8) 0 (0.0) 1 (1.2) 0 (0.0) 1 (1.2) 0 (0.0) 1 (1.2)	0 (0.0) 1 (1.2) 1 1 (1.2) 1 (1.2) 0 0 (0.0) 1 (1.2) 0 1 (1.2) 0 (0.0) 0 1 (1.2) 0 (0.0) 0 1 (1.2) 0 (0.0) 0 0 (0.0) 4 (4.8) 1 0 (0.0) 1 (1.2) 0 0 (0.0) 1 (1.2) 0 0 (0.0) 1 (1.2) 1	0 (0.0) 1 (1.2) 1 (1.2) 1 (1.2) 1 (1.2) 0 (0.0) 0 (0.0) 1 (1.2) 0 (0.0) 1 (1.2) 0 (0.0) 0 (0.0) 1 (1.2) 0 (0.0) 0 (0.0) 1 (1.2) 0 (0.0) 0 (0.0) 0 (0.0) 4 (4.8) 1 (1.2) 0 (0.0) 1 (1.2) 0 (0.0) 0 (0.0) 1 (1.2) 0 (0.0) 0 (0.0) 1 (1.2) 0 (0.0) 0 (0.0) 1 (1.2) 1 (1.2)	0 (0.0) 1 (1.2) 1 (1.2) 2 1 (1.2) 1 (1.2) 0 (0.0) 2 0 (0.0) 1 (1.2) 0 (0.0) 1 1 (1.2) 0 (0.0) 0 (0.0) 1 1 (1.2) 0 (0.0) 0 (0.0) 1 1 (1.2) 0 (0.0) 0 (0.0) 1 0 (0.0) 4 (4.8) 1 (1.2) 5 0 (0.0) 1 (1.2) 0 (0.0) 1 0 (0.0) 1 (1.2) 0 (0.0) 1 0 (0.0) 1 (1.2) 0 (0.0) 1 0 (0.0) 1 (1.2) 1 (1.2) 2

This is footnote 1 This is footnote 2

Source: [Study MK9999P001: adam-adae]