Selected Adverse Events by SOC and PT - summary (Safety Analysis Set)

Adverse Event	Xanomeline High Dose N=84 n (%)	Xanomeline Low Dose N=84 n (%)	Placebo N=86 n (%)
Any AE	79 (94.0 %)	77 (91.7 %)	69 (80.2 %)
EYE DISORDERS	0 (0.0 %)	0 (0.0 %)	1 (1.2 %)
Eye Allergy	0 (0.0 %)	0 (0.0 %)	1 (1.2 %)
Eye Pruritus	0 (0.0 %)	0 (0.0 %)	1 (1.2 %)
GASTROINTESTINAL DISORDERS	4 (4.8 %)	5 (6.0 %)	10 (11.6 %)
Diarrhoea	4 (4.8 %)	5 (6.0 %)	9 (10.5 %)
Hiatus Hernia	0 (0.0 %)	0 (0.0 %)	1 (1.2 %)
GENERAL DISORDERS AND ADMINISTRATION SITE CONDITIONS	5 (6.0 %)	5 (6.0 %)	1 (1.2 %)
Fatigue	5 (6.0 %)	5 (6.0 %)	1 (1.2 %)

Percentages are based on number of subjects in treatment arm.
N - number of subjects in treatment arm, n - number of subjects in the analysis, SOC - System Organ Class, PT - Prefered Term.