## **Table 14.2.3.1** Selected Adverse Events by SOC and PT - summary (Safety Analysis Set)

|  | Xanomeline<br>High Dose<br>N=84 n (%) | Xanomeline<br>High Dose<br>N=84 n (%) | Placebo<br>N=86 n<br>(%) |
|--|---------------------------------------|---------------------------------------|--------------------------|
| Any AE   | 79 ( 94.0 %)                          | 77 ( 91.7 %)                          | 69 ( 80.2 %)             |
| GASTROINTESTINAL DISORDERS                           | 4 ( 4.8 %)                            | 5 ( 6.0 %)                            | 9 ( 10.5 %)              |
| Diarrhoea  | 4 ( 4.8 %)                            | 5 ( 6.0 %)                            | 9 ( 10.5 %)              |
| 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