

Adverse Event Count by Treatment Group

| Adverse Events | Placebo | Xanomeline High Dose | Xanomeline Low Dose |
|--|---------|----------------------|---------------------|
| ABDOMINAL PAIN | 1 | 2 | 3 |
| AGITATION | 2 | 1 | 2 |
| ALOPECIA | 1 | 0 | 0 |
| ANXIETY | 2 | 0 | 4 |
| APPLICATION SITE DERMATITIS | 9 | 12 | 15 |
| APPLICATION SITE ERYTHEMA | 3 | 23 | 20 |
| Adverse events are coded according to MedDRA version 23.0 adam-adae | | | |