Appendix A: Data collection

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| --- | --- | --- | --- |
| Procedures | Enrolment | Post allocation ( trial) | Close out |
| V1 | Diary | T1 | T2 | T3 | V2 |
| Eligibility assessment | √ |  |  |  |  |  |
| Informed consent +/- assent | √ |  |  |  |  |  |
| Baseline assessment | √ |  |  |  |  |  |
| Randomization | √ |  |  |  |  |  |
| Allocation of study medications | √ |  |  |  |  |  |
| Adverse events assessment |  |  | √ | √ | √ | √ |
| NRS score | √ |  | √ | √ | √ | √ |
| The Self-Rating Anxiety Scale (SAS) | √ |  |  |  |  | √ |
| The Self-Rating Depression Scale (SDS) | √ |  |  |  |  | √ |
| The EuroQol (EQ-5d) | √ |  |  |  |  | √ |
| Satisfaction with the medication (0-10) |  |  |  |  |  | √ |
| The quality and duration of sleep | √ | √ |  |  |  |  |
| Number of study medications used |  |  |  |  |  | √ |
| Duration that analgesics were taken |  |  |  |  |  | √ |

V1 = Visit 1; V2 = Visit 2 (14 days after randomization); T1 = telephone follow-up 1 (1 day after randomization); T2 = telephone follow-up 2 (3 days after randomization); T3 = telephone follow-up 3 (7 days after randomization).

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