**Intervention extraction form[[1]](#footnote-1)**

|  |  |  |  |  |
| --- | --- | --- | --- | --- |
| **Label** | **Type** | **Parameter** | | |
| Description of intervention | | | | |
| Description of intervention | text |  |  | |
| Dosage | number/ single choice for unit |  | □ Milligram  □ Milliliter  □ Other  □ Unclear  □ Not applicable | |
| Schedule | number/ single choice for unit |  | □ Per minute  □ Per hour  □ Per day  □ Per week  □ Per month  □ Per year  □ Fixed schedule  □ Unclear  □ Not applicable | |
| Route of administration | single choice | □ Per os  □ Intramuscular  □ Intravenous  □ Subcutaneous  □ Intraarticular  □ Intrathecal  □ Intradermal  □ Superficial (skin)  □ Intraosseal  □ Intraarterial  □ Other  □ Unclear  □ Not applicable | | |
| Maximum treatment duration/administration | number/ single choice for unit |  | □ Minute(s)  □ Hour(s)  □ Day(s)  □ Week(s)  □ Month(s)  □ Year(s)  □ Administration(s)  □ Unclear |  |
| Average treatment duration/administration | number/ single choice for unit |  | □ Minute(s)  □ Hour(s)  □ Day(s)  □ Week(s)  □ Month(s)  □ Year(s)  □ Administration(s)  □ Unclear |  |
| Maximum follow-up Maximum study duration/administration | number/ single choice for unit |  | □ Minute(s)  □ Hour(s)  □ Day(s)  □ Week(s)  □ Month(s)  □ Year(s)  □ Administration(s)  □ Unclear | |
| Average study duration/administration | number/ single choice for unit |  | □ Minute(s)  □ Hour(s)  □ Day(s)  □ Week(s)  □ Month(s)  □ Year(s)  □ Administration(s)  □ Unclear | |
| Blinding of participants | | | | |
| Description by investigator(s) | text |  | | |
| Risk of bias | single choice | □ Low risk  □ High risk  □ Unclear risk  □ Unclear risk – probably low  □ Unclear risk – probably high | | |
| Rationale for choice | text |  | | |
| Blinding of health care providers | | | | |
| Description by investigator(s) | text |  | | |
| Risk of bias | single choice | □ Low risk  □ High risk  □ Unclear risk  □ Unclear risk – probably low  □ Unclear risk – probably high | | |
| Rationale for choice | text |  | | |
| Participants | | | | |
| Number of allocated participants (to intervention) | number |  | | |
| General comments on the intervention | | | | |
| Comment | text |  | | |

1. To be completed for each trial arm/group i.e. including control [↑](#footnote-ref-1)