**Data to collect for adverse events substudy**

**Trial basics**

ID,

Comparator drugs

Primary outcome measure – surrogate outcome only, yes/no

Age cut-offs

Phase

(these are mainly relevant to the matching process)

**Baseline characteristics**

Mean age + SD

Percentage male/ female

Mean BMI +SD

(These are minimum criteria to try to get some representativeness of the population and were the ones I found easiest to access consistently)

**Adverse events**

Number of participants randomised

Number of participants completing trial

Total follow up time(weeks)

Number of events/patients with AE ( this is what was more accessible than number of AEs) -

Number of events/patients with SAEs

Number of deaths

( We already have all the information above for the clinical trials carried out in older people exclusively and 23 of the trials carried out in adults of all ages)