SUPPLEMENTAL CONTENT 3: NUMBER OF ADVERSE EVENTS (AE) REPORTED PER TRIAL, CATEGORISED BY THE PRESENCE OR ABSENCE OF AN AE ASSESSMENT METHOD (N=117)

- Reported ≥1 AE
  - n=40
  - n=22

- Reported 0 AE
  - n=15
  - n=31

- Results missing
  - n=9

- Trials with a known AE assessment method (n=64/117)
- Trials with an unknown AE assessment method (n=53/117)