**POSE - End of study reporting form**

|  |  |
| --- | --- |
| Centre number |  |
| Institution Name |  |
| Recruiting period | From (dd/Mmm/YYYY) to (dd/Mmm/YYYY) |
|  | # of patients |
| 1-Screened patients | Total # of all patients (meeting the protocol inclusion criteria) who had anaesthesia during the period of recruitment |  |
| 2-Included patients | Total # of patients corresponding to inclusion criteria that you succeed to capture  |  |
| 3-Screened failure patients | Total # of patients corresponding to inclusion criteria that you failed to capture (logistics problem, consent mandatory but not signed, patient refused study participation etc.) |  |
| Please confirm collaborators from your centre with substantial involvement in the study.Each participating centre including at least 1 patient can designate one investigator to be mentioned as a collaborator within the POSE Study Group in the publication. Please add the names of the collaborators/ authors according to the number of enrolled patients in the table below.  |
| # of patients recruited and included | # of collaborators per centre | **Last Name** | **First Name** | **E-mail address** |
| 1 | 1stCollaborator |  |  |  |
| 25 | 2ndCollaborator |  |  |  |
| 48 | 3rdCollaborator |  |  |  |
| 71 | *1st Co-Author* |  |  |  |
| 72 | 4thCollaborator |  |  |  |
| 96 | 5thCollaborator |  |  |  |
| 120 | 6thCollaborator |  |  |  |
| 143 | *2nd Co-Author* |  |  |  |
| 143 | 7thCollaborator |  |  |  |
| 167 | 8thCollaborator |  |  |  |
| 191 | 9thCollaborator |  |  |  |
| 214 | *3rd Co-Author* |  |  |  |
| 215 | 10thCollaborator |  |  |  |
| 238 | 11thCollaborator |  |  |  |
| .... | ..... |  |  |  |

Last and first name Site Local Investigator: Date (dd/Mmm/YYYY): Signature of Site Local Investigator: