**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: