**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 by order of importance.Please mention site local coordinator as #1.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. For each further 25 included, followed up and documented patients a further collaborator can be mentioned in this table. Enrollment of ≥ 75 patients entitles to designate 1 Co-Author in addition to the collaborators.  |
| # of patients recruited | # of collaborators per centre | **Last Name** | **First Name** | **E-mail address** |
| 1 | 1 |  |  |  |
| 26 | 2 |  |  |  |
| 51 | 3 |  |  |  |
| 75 | *1 Co-Author* |  |  |  |
| 76 | 4 |  |  |  |
| 101 | 5 |  |  |  |
| 126 | 6 |  |  |  |
| .... | .... |  |  |  |

Last and first name Site Local Investigator:

Date (dd/Mmm/YYYY):

Signature of Site Local Investigator:

**Return this form to POSE study team: akowark@ukaachen.de**