|  |
| --- |
| **Project identification** |
| Project name/-description |  |
| Principal Investigator (Project Manager) |   |
| Entity or person responsible for research’s representative |  |
| Identification in the continuous record of project activities |  |

|  |  |  |
| --- | --- | --- |
| **Task/duties** | **Task carried out by:** | **Further information** |
| Assessment of soundness/ justifiability including privacy protection |  |  |
| Registration of the project in the FORSKPRO;UiO’s system for continuous overview |  |  |
| Day-to-day execution of research tasks \* |  |  |
| Applications for internal and external approvals |  |  |
| Communication with legal authorities |  |  |
|  |  |  |
| Processing (handling) and storage of research data and other project documentation |  |  |
| Follow-up of the project funding |  |  |
| Final and other reports to REK |  |  |
| Final reports and other reports to any other authorities |  |  |
| Any other tasks: |  |  |
|  |  |  |
|  |  |  |

\* Clinical trials of medicinal products or medical devices are documented in separate forms, cf

Procedure Description 3, Annex 3.2 Form, the Principal Tester’s (Principal Investigator’s) delegation of tasks

Procedure Description 4, Annex 4.2 Form, the Principal Tester’s (Principal Investigator’s) delegation of tasks

The project was presented at the Department -/ unit level \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

 dd.mm.yyyy

Principal Investigator*’*s signature \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_