**Sedana Medical Research Grant - Application form**

|  |  |
| --- | --- |
| **Principal Investigator** | Name, title, academic/clinical affiliation |
| **Co-Investigator(s)\*** | Name, title, academic/clinical affiliation |
| **Site(s)** | Institution(s) where the study will take place |
| **Title of study** | Title |
| **Background and rationale** | Background and medical/scientific rationale, hypothesis and clinical significance |
| **Patient population** | Patient inclusion and exclusion criteria  |
| **Intervention** | Planned treatment, dose range/duration, administration route |
| **Comparison\*** | Control intervention or comparator |
| **Outcomes** | Primary endpoint and secondary endpoint(s) |
| **Study design** | Design of the study (e.g., randomized, blinded, non-interventional, multicenter) |
| **Sample size** | Number of patients and rationale or power calculation  |
| **Timelines** | Anticipated study duration |
| **Statistics** | Short description of the methods for data analysis |
| **Resources and feasibility** | Description of setting (ICU type, patient availability, facilities) |
| **Research coordinator** | Name and contact details (unless PI/co-investigator has this role) |
| **Requested support from Sedana Medical**  | Specify support required (e.g., funding, devices, services)  |
| **Request for data access\*** | State if access to data from a Sedana-sponsored study is required; if yes, specify which data in table/appendix |
| **Ethical considerations and approvals\*** | Status of ethical/IRB application (e.g. not yet filed, approval pending, approval code available)  |
| **Database registration\*** | Status of study plan registration in public database (e.g. clinicaltrials.gov) |

*\*if applicable*