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| Collaborative Research Support Proposal | |
| Primary Investigator Information | |
| Name: Click or tap here to enter text. | |
| Institution:Click or tap here to enter text. | |
| Email:Click or tap here to enter text. | |
| Phone Number:Click or tap here to enter text. | |
| Address:Click or tap here to enter text. | |
| Date of Submission: Click or tap to enter a date. | |
| Study Information | |
| Study Title: Click or tap here to enter text. | |
| Study Sponsor(s) Outside of Illumina:Click or tap here to enter text. | |
| If this study was developed in collaboration with Illumina, who is your primary contact?  Click or tap here to enter text. | |
| Scientific Background/ Rationale and Study Aims | |
| Background Rationale:Click or tap here to enter text. | |
| Study Aims:Click or tap here to enter text. | |
| Project Description and Study Design | |
| Study design (please complete checkboxes as appropriate and add a description of the study design):  Prospective sample collection  Retrospective sample collection  Implementation study  Head-to-head study  Proof of concept  Proof of principle  Biomarker discovery  Click or tap here to enter text. | |
| Study Population (Inclusion/Exclusion Criteria): Click or tap here to enter text. | |
| Sample type(s): Click or tap here to enter text. | |
| Number of samples: Click or tap here to enter text. | |
| Test and Data Analysis Sites: Click or tap here to enter text. | |
| Genetic analysis methodologies: Click or tap here to enter text. | |
| Will results be reported to patients?  Yes  No | |
| Statistical plan: Click or tap here to enter text. | |
| Technology | |
| Check appropriate technology below, and provide any addition details:  Click or tap here to enter text. | |
| Reproductive Health:  VeriSeq NIPT Solution v2.0  Other (specify) Click or tap here to enter text. | Oncology:  TruSight Oncology 500 ctDNA  TruSight Oncology 500  WGS/WTS  Other (specify) Click or tap here to enter text. |
| Infections Disease:  COVIDSeq  Other (specify) Click or tap here to enter text. | Genetic Health:  WGS  WES  WTS  Other (specify) Click or tap here to enter text. |
| Study Aims | |
| Milestones, and any key dates or dependencies for study activation/completion:Click or tap here to enter text. | |
| Endpoints:Click or tap here to enter text. | |
| Deliverables:Click or tap here to enter text.  Conference abstract  Scientific publication  Tool development  Test implementation | |
| Study Timeframe | |
| Estimated study start date:Click or tap to enter a date. | Total Estimated timeline (in months):Click or tap here to enter text. |
| Study Support | |
| Total Study Budget Estimate:Click or tap here to enter text. | |
| **Please attach a copy of the itemized budget to this application.** | |
| Support Requested from Illumina (select all that apply):  Financial, include amountClick or tap here to enter text.  Instrument Loaner, specify instrumentClick or tap here to enter text.  Reagents, specify type and quantityClick or tap here to enter text.  In-house testing supportClick or tap here to enter text.  Bioinformatics support Click or tap here to enter text.  Publication support Click or tap here to enter text.  IRB Guidance Click or tap here to enter text.  Protocol writing support Click or tap here to enter text.  Biostatistics support Click or tap here to enter text.  Research and development support Click or tap here to enter text.  Other, please specify Click or tap here to enter text. | |
| Materials and Support Provided by the PI:  Click or tap here to enter text. | |
| Data Ownership Plan:  Data will not be shared with Illumina  Summary data will be shared with Illumina  De-identified full study data will be shared with Illumina  Click or tap here to enter text. | |

Completed forms should be sent via email to [iResearch@illumina.com](mailto:iResearch@illumina.com) along with:

* Investigator CV
* Study protocol (if applicable)
* Copy of IRB (if applicable)
* Itemized budget
* Any other supporting documents that will aid in the review process

Failure to include all required information may result in delays and support being declined.