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| --- | --- |
| I. Project Title | |
|  | **Date:** |
| **II. Project Group Investigators (Please list names and institutes of all investigators)**  *(for external investigator- it is required to include at least 1 ILCCO/TRICL investigator in the Project Group)* | |
| |  |  |  |  | | --- | --- | --- | --- | | **Author/Co-author Name** | **Institution** | **Phone Number** | **Email** | |  |  |  |  | |  |  |  |  | |  |  |  |  | |  |  |  |  | |  |  |  |  | |  |  |  |  | | |
| III. Background | |
|  | |
| IV. Specific Aims | |
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| **V. Methods** | |
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| **VI. Materials or variables needed from the study PIs (Please separate out the required vs optional components)** | |
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| **VII. Time line** | |
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| **VIII. Funding Sources and Declaration of Conflict of Interests.** *To ensure full transparency and to protect collaborating study PIs, ILCCO/TRICL requires the Project Leaders to disclose any circumstances that could give rise to a potential conflict of interests related to the proposed project activity in particular, or to lung cancer and/or tobacco products in general, including but not limited to funding sources, employment and consulting, board membership and investment interests within the last 5 years.* | |
|  | |
| **IX. Other remarks** (e.g. **publication** plan, etc) | |
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| **X. Scope/Focus of your proposal (see definition below)** | |
| Scope: 1=Genome-Wide, 2=Gen-Sets, 3=Targeted Regions, 4=Specific Pathway.  Focus: 1=Risk, 2=Prognosis, 3=Both Risk and Prognosis, 4=Method, 5=Other Phenotype, 6=Interactions | |