

R21 Grant Checklist – Forms F

| | |
|---|---|
| Cover Letter | Don't add study or award preferences |
| Abstract/Project Summary | 30 lines maximum |
| Narrative/Relevance | 2-3 sentences |
| Introduction | 1 page (Resubmission/Revision Only) |
| Specific Aims | 1 page |
| Research Strategy | 6 pages |
| References | no limit if applicable |
| Progress Report Publication List | (Renewal Only) |
| Vertebrate Animals | no limit if applicable |
| Select Agent Research | no limit if applicable |
| Multiple PI Leadership Plan | no limit if applicable |
| Consortium Contractual Arrangements | no limit if applicable |
| Letters of Support | no limit |
| Resource Sharing Plan | no limit if applicable |
| Authentication of Key Biological/Chemical Resources ... | 1 page if applicable |
| Appendix | Only if PA instruct to add (survey, questionnaires) |
| Biosketch | 5 page maximum |
| Budget Justification | no limit |
| Facilities and Resources | no limit if applicable |
| PHS Assignment Request Form | Add study or award preference |
| Foreign Justification | If foreign subgrantee on project (no-limit) |

Human Subjects Research Form (mandatory, see HSR Form)

Section 2 – Study Population Characteristics

| | |
|--|------------------------|
| Inclusion of Individuals Across the Lifespan | no limit |
| Inclusion of Women and Minorities | no limit |
| Recruitment and Retention Plan | no limit if applicable |



R21 Grant Checklist – Forms F (Continued)

Study Timeline no limit if applicable
Cumulative Inclusion/Planned Enrollment Report form page if applicable

Section 3- Protection and Monitoring Plans

Protection of Human Subjects no limit
Single IRB plan no limit if applicable
Data Safety Monitoring Plan no limit if applicable
Overall Structure of the Study Team no limit applicable

Section 4 on HSR- (only required if you answered “Yes” to all questions on the Clinical Trial Questionnaire). If you have a Clinical Trial, please provide the documents below:

Study Design 32000 Characters
Statistical Design and Power no limit if applicable
FDA – regulated intervention no limit if applicable
Dissemination Plan no limit if applicable

Section 5 – Clinical Trial Attachments

Subawards documents (Pre-award will request. Please see sub contact information):

- Letter of Intent and 398 face page signed by institutional official
- Detailed Budget and Justification
- Scope of work
- R&R budget form
- Biosketch and profile information for all senior key personnel
- Facilities & Resources
- 5 question certification form for all senior key personnel