



NIH FORMS VERSION I
RESEARCH PROPOSAL CHECKLIST
Updated June 10, 2026

The following checklist is designed for most National Institutes of Health research (e.g., R01, R21, etc.) proposals. The checklist is designed to assist PIs in responding to a NIH funding opportunity announcement (FOA). Kindly note that particular FOAs (Fellowships, STTR/SBIR, Training, etc.) have specific requirements that may not be included in this checklist. ***This document is designed only to serve as a project management tool. It does NOT replace the detailed information available within the FOA, and the funding agency's forms, instructions, and review criteria.*** For any questions, please refer to the FOA, contact your program officer, or contact your [ORA Contract Administrator](#).

****When submitting NIH proposals, complete in [ASSIST](#)****

FORMATTING BASICS

- Font Size: 11 pt or larger
- Recommended Fonts: Arial, Georgia, Helvetica, Palatino Linotype
- No headers or footers
- No page numbers
- Use section headings
- File names are 50 characters or less (no special characters: "&", "*", "%", "#", or "/" in file name)
- Margins: Minimum of ½ inch margins on all sides
- Hyperlinks: Typically limited to citing relevant publications in biosketches and publication lists only. See [NOT-OD-20-174](#) and the [NIH Format Attachments page](#).

****This list is not exhaustive. All formatting requirements are here:**

<https://grants.nih.gov/grants/how-to-apply-application-guide/format-and-write/format-attachments.htm>

RESOURCES

[SF424 Application Guide](#)

[Standard Due Dates](#)

[Page Limits](#)

[UMD Standard Institutional Information](#)

✓	Proposal Component	Notes	SF424 Guide Reference
	SF424 (R&R) Form	Title: limited to 200 characters, including spaces and punctuation EIN: 1520710851-A1 If you check the "Changed/Corrected Application" box, then "Field 4.c Previous Grants.gov Tracking ID" is required	G.200
	PHS 398 Cover Page Supplement		G.210
	RR Other Project Information	▪ Human Subject Assurance Number: IRB FWA: 00005856 ☒ Need help determining whether your application includes human subjects? Check out the NIH Human Subjects Research website for information, including a Decision Tool to help make the determination. ☐ Justification Requirement: If your application proposes using human specimens or data but you mark "No" to the "Are Human Subjects Involved?" question, you must provide a clear justification explaining why this does not constitute human subjects research ▪ Animal Welfare Assurance Number: IACUC OLAW: D16-00172 ▪ International Collaborators: Include a "Foreign Justification" attachment in Field 12 - Other Attachments. ▪ Project Summary/Abstract: 30 lines of text ▪ Project Narrative: limited to three (3) sentences ▪ Bibliography/Reference Cited: Must include PubMed Central or NIHMS reference #, if applicable (when applicant (PI) is the author or co-author of the article) If a PMCID is not yet available because the journal is submitting articles directly to PMC, you should indicate "PMC Journal - In Process"	G.220 G.220.7 G.220.8 G.220.9 G.220.10 G.220.11 G.220.12

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		through established repositories, etc., including an amount for each category and a brief explanation. The recommended length of the justification should be no more than half a page.	
PHS 398 Research Plan	▪ Introduction: For resubmission or revision applications only, unless otherwise specified in the NOFO. Limit one page. ▪ Specific Aims: State concisely the goals of the proposed research and summarize the expected outcome(s), including the impact that the results of the proposed research will have on the research field(s) involved. Limit one page (check NIH Table of Page Limits), unless otherwise specified in the NOFO. ▪ Research Strategy: Page limit varies by project type. Consult the NIH Table of Page Limits. Unless otherwise specified in the NOFO, start each section with the appropriate headings in the following order: Significance, Innovation, and Approach (includes Preliminary Studies for New Applications and Progress Report for Renewal/Revision Applications). ▪ Progress Report Publication List: Only for renewal applications. ▪ Vertebrate Animals: Required if vertebrate animals are involved, or if you answered "Yes" to the question #2 "Are Vertebrate Animals Used?" on the Other Project Information Form. See Section G.400.5 for additional guidance. ▪ Select Agent Research: Required if select agents are involved. See section G.400.6 for additional guidance. ▪ Multiple PI/PD Leadership Plan: Required for multi-PD/PI projects. For background information, see the NIH Multiple Principal Investigators page. See G.400.7 for Renewal and Resubmission applications. ▪ Consortium/Contractual Arrangement: Required if subrecipients are included. ▪ Letters of Support: Attach all appropriate letters of support, including any letters necessary to demonstrate the support of consortium participants and collaborators such as Senior/Key Personnel and Other Significant Contributors included in the grant application. ▪ Resource Sharing Plan: This section should include the following. The Data Management and Sharing and Genomic Data sharing Plans are now included in Other Plans (see below). ☐ Sharing Model Organisms – Required when proposing to develop model organisms regardless of amount requested. ☐ Research Tools: Required when Research Tools / resources have been developed with NIH funds. See also NIH GPS 8.2.3. ▪ Other Plans: (Data Management and Sharing Plan) Recommended not to exceed two pages. Sample format available on Data Management and Sharing Plan Format Page. Refer to the list of NIH activity codes subject to the DMS Policy and your NOFO to determine if your application is required to provide an attachment and address a Data Management and Sharing (DMS) Plan. Applicants proposing to conduct research that will generate scientific data are subject to the NIH Data Management and Sharing Policy and must attach a DMS Plan. ☐ Genomic Data Sharing – Include as a part of the DMS Plan when seeking funding for research that generates large-scale human or non-human genomic data. See more information on the NIH Genomic Data Sharing (GDS) Policy and applicability of the GDS policy. ▪ Authentication of Key Biological and/or Chemical Resources: Suggested maximum 1 page; briefly describe methods to ensure the identity and validity of key biological and/or chemical resources used in the proposed studies. ▪ Appendix: Include if applicable; check NOFO for specific instructions. <u>Do not use to circumvent page limits</u>. See NOT-OD-17-098 and G.400.13 regarding allowable appendix materials. Max 10 PDF attachments allowed; if more than 10 are needed, combine	G.400.1 G.400.2 G.400.3 G.400.4 G.400.5 G.400.6 G.400.7 G.400.8 G.400.9 G.400.10 G.400.11 G.400.12 G.400.13	

		remaining info into attachment #10.	
	PHS Human Subjects and Clinical Trials Information Form	- For detailed information, visit the NIH Human Subjects Research website. Use of Human Specimens and/or Data: Required regardless of response to question #1 "Are Human Subjects Involved?" on the RR Other Project Information Form. Delayed Onset Study: Required only when human subjects research is anticipated within the period of award but definite plans cannot be described in the application. Study Record and Attachments: Required for any project involving Human Subjects and/or Clinical Trials.	G.500
	PHS Assignment Request Form	Optional.	G.600