

Quick Guide for a Compelling Research Strategy

Suggested section lengths are based on a 12-page research strategy. For detailed guidance, see the [NIH application guide](#) and [SF424](#).

THE ENTIRE RESEARCH STRATEGY

- Read the RFP to learn what is important to the funder**
 - The proposal should address the goals and scope stated in the RFP
- Use formatting to enhance readability**
 - Highlight key information with bold or underline (use sparingly)
 - Place figures and tables close to the relevant text
 - Check spelling and grammar
- Number and label the sections to help reviewers find important information**
 - Sections and subsections should be in a logical order
 - Use consistent numbering and formatting throughout
 - Leverage section numbering to reduce repetition (e.g., “To determine the epigenetic effects of sleep deprivation in adults, we will compare histone modifications in serum obtained from researchers 3 days before, 1 day after, and 7 days after a grant deadline, using the methods described in [Aim 1.2.2](#).”)
 - Check all numbering and references to other sections before submitting
- Follow [NIH formatting guidelines](#) and [page limits](#)**
 - Margins are at least ½ inch
 - Font size in main text is 11 pt or higher
- Check the PDF for accuracy and legibility before submitting**

SIGNIFICANCE (usually 1.5-2 pages)

- Establish the importance of the problem that the project addresses**
 - Usually the opening paragraph
 - Provide context of the “big picture” problem
 - Introduce the main research question
 - Explain impacts to public health, individual health, the economy, and/or scientific progress
 - Support claims with statistics and citations
- Describe the scientific premise of the project**
 - 2-3 paragraphs
 - Summarize background information that is directly relevant to the research question
 - Do not review the entire field. Instead, build a strong, focused rationale for the hypothesis. How does each bit of information support the hypothesis?
 - Published literature and key preliminary data
 - Consider strengths and weaknesses of supporting data
 - State the hypothesis and goals of the project
 - The project should directly address the hypothesis and main problem presented
- Explain the likely short-term impacts of the project for scientific knowledge, technical capability, and/or clinical practice (2-3 sentences)**

- **Indicate how the completed project will change the field**
 - Usually concludes the significance section
 - Concepts, methods, technologies, treatments, services, or preventive interventions, etc.
 - Longer-term possibilities can be included, but should not be over-stated

INNOVATION (usually ½ to 1 page)

- **Innovation can take several forms**
 - Conceptual innovations - challenge a paradigm or provide an important knowledge leap
 - Technical innovations – develop or new or substantially improved way of doing things
 - Resource innovations – establish and provide proof of concept for a crucial new resource
 - May be relevant for research and/or clinical practice
 - Consider impact beyond your own field
- **Explain the limitations of current knowledge and tools**
- **Describe how the project will address these limitations**
- **Explain the advantages of the innovation over existing options**
 - Cost savings, improved efficiency, new capabilities, better safety or accuracy, important shifts in knowledge

APPROACH (usually ~9 pages)

- **Open with the overall strategy for addressing the central hypothesis**
 - Summarize the general approach, including key resources and methodologies across aims
 - Explain how the aims work together so the whole is greater than the sum of the parts
 - Flowcharts and diagrams can be helpful
- **Present preliminary data that support the central hypothesis and/or experimental feasibility**
 - Can be in one combined section and/or within each aim
- **Usually arranged into 2-4 specific aims**
 - Aims should be related, but not dependent on one another
 - Experiments should provide valuable information even if the hypothesis is incorrect
- **Provide enough detail to show the experimental design is justified, robust, and unbiased**
 - Study populations, model systems, key reagents
 - Replicates (technical and biological), controls, comparison groups, biological variables (e.g., sex)
 - Methods
 - Data quality control, data processing, power analyses, and statistical design and interpretation
 - Indicate potential hazards and relevant precautions
- **Discuss potential problems and alternative strategies**
 - Technical challenges can be addressed with alternative methods or data demonstrating feasibility
 - Are there potential challenges in obtaining key resources or recruiting participants?
 - What if the underlying assumptions and/or the stated hypotheses are incorrect?
 - Provide a realistic way to mitigate any problem that is mentioned
- **Briefly describe one or two concrete future directions that can build on this study**

Date: 06/28/2019

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Funding: NIGMS award U54GM104934