

## Checklist: The NIH Proposal at-a-glance

A MaineHealth Member

NOTE: See Detailed Instructions for further guidance on each section

TITLE: A short, informative statement that includes the study design and the sample
<b>SENIOR/KEY PERSONNEL</b> : Lists members of the research team who will have direct responsibilities for conduct of the research and who will contribute to the scientific direction
<b>OTHER SIGNIFICANT CONTRIBUTORS:</b> Scientists with particular expertise who have agreed to consult on the project but are not giving measurable time
BIOSKETCHES: Needed for all senior,/key personnel and Other Significant Contributors
LETTERS OF SUPPORT: Needed from consultants and collaborators not listed on budget and from Other Significant Contributors
ABSTRACT: Briefly describes project, including significance, goals, hypotheses, specific aims, methods (30 lines max)
NARRATIVE/RELEVANCE: Describes public health relevance in lay terms (2-3 sentences)
<b>FACILITIES/OTHER RESOURCES:</b> Describes office, lab, computer, animal facilities, core facilities, clinical resources, and scientific environment (including collaborative opportunities, seminars) conducive to success
EQUIPMENT: Describes major equipment available and to be used for project
BUDGET JUSTIFICATION: Justifies tasks of personnel; justifies supplies, travel, other items if non-modular.
SPECIFIC AIMS: States purpose, goals, hypotheses and numbered specific aims (1 page)
RESEARCH STRATEGY (12 pgs or R01; 6 pgs for R21): Must include Significance, Innovation and Approach.
REFERENCES CITED: Full bibliography of works cited in proposal including all author names
RESOURCE SHARING PLAN: Describes how you will share data with scientific community
<b>AUTHENTICATION OF KEY BIOLOGICAL OR CHEMICAL RESOURCES</b> : Explains how you will authenticate the reliability of biological and chemical specimens used (1 page)
ADDITIONAL SECTIONS (IF APPLICABLE)
HUMAN SUBJECTS:  o If using deidentified specimens only, justify this determination
o If claiming exemption, justify the exemption claim
<ul> <li>If non-exempt human subjects;</li> <li>Section 1: Basic info</li> <li>Section 2: Study Population Characteristics (includes Inclusion Enrollment Report – not required for exemption 4)</li> <li>Section 3: Protection and Monitoring Plans</li> <li>Section 4: Protocol Synopsis (Clinical Trials Only)</li> </ul>
VERTEBRATE ANIMALS: Discusses proposed use, justification of use and minimization of pain
INTRODUCTION: (re submissions only): 1 page response to critique of the original grant cycle
PROGRESS REPORT (renewals only): List of publications from previous grant cycle
MULTIPLE P.I. PLAN: Description of management organization if multiple Principal Investigators on proposal.
CONSORTIUM ARRANGEMENTS: Description of role of subcontracting institution and how it contributes to project.

NOTE: If you are including a subcontract with another institution, MMC needs a budget, budget justification, and signed face page from that institution (Grants and Contracts will coordinate this requirement).