

GUIDE FOR ASSIGNED REVIEWERS' PRELIMINARY COMMENTS ON MENTORED RESEARCH SCIENTIST DEVELOPMENT AWARD (K01) APPLICATIONS

PA NUMBER: PA-06-001

Complete details at: <http://grants.nih.gov/grants/guide/pa-files/PA-06-001.html>

The purpose of the Mentored Research Scientist Development Award (K01) is to provide support and “protected time” (three, four, or five years) for an intensive, supervised career development experience in the biomedical, behavioral, or clinical sciences leading to research independence.

General considerations when reviewing Mentored Research Scientist Development Award applications:

- Candidates must be U.S. citizens or non-citizen nationals or an individual lawfully admitted for permanent residence that hold a research or health-professional doctoral degree or its equivalent and can commit a minimum of 75% of full-time professional effort conducting research and relevant career development activities specified in the application.
- The candidate must demonstrate and justify the need for a three, four, or five-year period of additional supervised research experience. Planning, direction, and execution of the proposed career development program and research project will be the responsibility of the candidate and his/her mentor.
- The proposed career development experience must be in a research area new to the applicant and/or one in which an additional supervised research experience will substantially augment the research capabilities of the applicant.

Note: Although most of the NIH Institutes and Centers (ICs) use K01 awards to support career development experiences that lead to independence, characteristics of ideal candidates may vary. For example, some of the ICs reserve this award for individuals who propose to train in a completely new field or for individuals who have had a hiatus in their careers because of illness or pressing family circumstances. Other ICs reserve the K01 for faculty from underrepresented groups or faculty at minority serving institutions who may want to enhance their research skills and knowledge through a period of supervised training at a research center. Reviewers are strongly encouraged to contact their study section's Scientific Review Administrator and discuss any special review considerations for K01 applications they are preparing to review.

Review Criteria

The goals of NIH-supported career development programs are to help ensure that diverse pools of highly trained scientists are available in adequate numbers and in appropriate research areas to address the Nation's biomedical, behavioral, and clinical research needs. The scientific review group will address and consider each of these criteria in assigning the application's overall score, weighting them as appropriate for each application.

- Candidate
- Career Development Plan
- Research Plan

- Training in the Responsible Conduct of Research
- Appropriateness of and statements by former Mentor, Co-Mentor(s), Consultant(s), and Collaborator(s)
- Environment and Institutional Commitment to the Candidate

For revised applications, also comment briefly on whether the application is improved, the same, or worse. In addition, provide a one-sentence summary of your evaluation at the end of each section. After considering all of the review criteria, briefly summarize the strengths and weaknesses of the application and recommend an overall level of merit in a section titled Summary and Recommendation (see below). Please note that your comments will be used essentially unedited in the final summary statement sent to the candidate.

The application does not need to be strong in all categories to receive a high priority score. These criteria are listed in logical order and not in order of priority.

Candidate

- Potential to develop as an independent and productive researcher
- Quality of the candidate's research, academic and (if relevant) clinical record;
- Commitment to meeting the program objectives to become an independent researcher;
- Quality of the letters of reference from three well-established scientists evaluating the candidate's potential to pursue an independent health-related research scientist career;
- Letters of reference submitted by mentor(s)/ co-mentor(s) will be considered independent of and in addition to of the three required reference letters.

Career Development Plan

Likelihood that the plan will contribute substantially to the scientific development of the candidate leading to scientific independence, based on:

- Appropriateness of the content, scope, phasing, and duration of the career development plan when considered in the context of prior training/research experience and the stated training and research objectives for achieving research independence;
- Plans for monitoring and evaluating the candidate's research and career development progress.

Research Plan

Reviewers recognize that an individual with limited research experience is less likely to be able to prepare a research plan with the breadth and depth of that submitted by a more experienced investigator. Nevertheless, a fundamentally sound research plan must be provided.

- Scientific and technical merit of the research question, design and methodology;
- Relevance of the proposed research to the candidate's career objectives;
- Appropriateness of the research plan to the stage of research development and as a vehicle for developing the research skills described in the career development plan.

Training in the Responsible Conduct of Research

- Quality and appropriateness of the proposed training in the responsible conduct of research.

Statements by Mentor/Co-Mentor(s), Consultant(s), and Collaborator(s)

- Appropriateness of the mentor's research qualifications in the area of the proposed research;
- Quality and extent of the mentor's proposed role in providing guidance and advice to the candidate;
- Previous experience in fostering the development of independent investigators;
- History of research productivity and peer-reviewed support;
- Adequacy of active/pending support for the proposed research project; and
- Strength of the mentor's statement.

Environment and Institutional Commitment to the Candidate

- Clear commitment of the institution to ensure that a minimum of 75% of the candidate's effort will be devoted directly to research, with the remaining percent effort being devoted to activities related to the successful development of a research career including clinical responsibilities;
- Strength of the institutional commitment to the career development of the candidate;
- Adequacy of research facilities and training opportunities, including faculty capable of productive collaboration with the candidate; and
- Quality and relevance of the environment for the scientific and professional development of the candidate, and
- Assurance that the institution intends for the candidate to be an integral part of its research program.

Summary and Recommendation

In one paragraph, briefly summarize the most important points of the Critique, addressing the strengths and weaknesses of the application in terms of the six review criteria. An application does not need to be strong in all categories to receive a good rating. Each scored application will receive a numerical rating that will reflect your opinion of its merit. The numerical rating is based on a scale from 1.0 for the most meritorious to 5.0 for the least meritorious with increments of 0.1 unit. Reviewers should score the "average" application they customarily review in their Scientific Review Group with a score of 3.0. This practice is designed to have 3.0 be the median.

Additional Review Criteria:

In addition to the above criteria, the following items will continue to be considered in the determination of scientific merit and the priority score and should be addressed in the critique.

Protection of Human Subjects from Research Risks: Evaluate the application with reference to the following criteria: risk to subjects, adequacy of protection against risks, potential benefit to the subjects and to others, importance of the knowledge to be gained. (If the applicant fails to address **all** of these elements, notify the SRA immediately to determine if the application should be withdrawn.) If all of the criteria are adequately addressed, and there are no concerns. Write "Acceptable Risks and/or Adequate Protections." A brief

explanation is advisable. If one or more criteria are inadequately addressed, write, "Unacceptable Risks and/or Inadequate Protections" and document the actual or potential issues that create the human subjects concern. If the application indicates that the proposed human subjects research is exempt from coverage by the regulations, determine if adequate justification is provided. If the claimed exemption is not justified, indicate "Unacceptable" and explain why you reached this conclusion. Also, if a clinical trial is proposed, evaluate the Data and Safety Monitoring Plan. (If the plan is absent, notify the SRA immediately to determine if the application should be withdrawn.) Indicate if the plan is "Acceptable" or "Unacceptable", and, if unacceptable, explain why it is unacceptable.

Inclusion of Women Plan:

Inclusion of Minorities Plan:

Inclusion of Children Plan:

Public Law 103-43 requires that women and minorities must be included in all NIH-supported clinical research projects involving human subjects unless a clear and compelling rationale establishes that inclusion is inappropriate with respect to the health of the subjects or the purpose of the research. NIH requires that children (individuals under the age of 21) of all ages be involved in all human subjects research supported by the NIH unless there are scientific or ethical reasons for excluding them. Each project involving human subjects must be assigned a code using the categories "1" to "5" below. Category 5 for minority representation in the project means that only foreign subjects are in the study population (no U.S. subjects). If the study uses both then use codes 1 thru 4. Examine whether the minority and gender characteristics of the sample are scientifically acceptable, consistent with the aims of the project, and comply with NIH policy. For each category, determine if the proposed subject recruitment targets are "A" (acceptable) or "U" (unacceptable). If you rate the sample as "U", consider this feature a weakness in the research design and reflect it in the overall score. Explain the reasons for the recommended codes; this is particularly critical for any item coded "U".

| Category | Gender (G) | Minority (M) | Children (C) |
|-----------------|-------------------|---------------------------------|------------------------------------|
| 1 | Both Genders | Minority & non-minority | Children & adults |
| 2 | Only Women | Only minority | Only children |
| 3 | Only Men | Only non-minority | No children included |
| 4 | Gender Unknown | Minority representation unknown | Representation of children unknown |
| 5 | | Only Foreign Subjects | |

NOTE: To the degree that acceptability or unacceptability affects the investigator's approach to the proposed research, such comments should appear under "Approach" in the five major review criteria above, and should be factored into the score as appropriate.

Vertebrate Animals: Express any comments or concerns about the appropriateness of the responses to the five required points, especially whether the procedures will be limited to those that are unavoidable in the conduct of scientifically sound research.

Biohazards: Note any materials or procedures that are potentially hazardous to research personnel and indicate whether the protection proposed will be adequate.

OTHER CONSIDERATIONS: These comments are useful to NIH but should not influence your overall score.

Administrative Note: (e.g., There is potential overcommitment and/or scientific overlap with other existing grants and/or pending applications.)

Data Sharing Plan: Applications requesting more than \$500,000 direct costs in any year of the proposed research are expected to include a data sharing plan in their application. Certain Program Announcements may request a data sharing plan for all applications regardless of the amount of direct costs. Assess the reasonableness of the data sharing plan or the rationale for not sharing research data.

Model Organism Sharing Plan: The NIH policy on sharing of model organisms for biomedical research was announced in the May 7, 2004 issue of the NIH Guide (<http://grants.nih.gov/grants/guide/notice-files/NOT-OD-04-042.html>). Starting with the October 1, 2004 receipt date, all new and competing-renewal NIH grant applications that plan to produce model organisms will be expected to include a sharing plan. Unlike the NIH Data Sharing Policy, the submission of a model organism sharing plan is NOT subject to a cost threshold of \$500,000 or more in direct costs in any one year, and is expected to be included in all applications where the development of model organisms is anticipated.

Budget: Evaluate the direct costs only. Do not focus on detail. For all years, determine whether all categories of the budget are appropriate and justified. Provide a rationale for each suggested modification in amount or duration of support.

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