

following items will be considered in the determination of scientific merit and priority score:

1. **Intervention.** Is the potential efficacy or effectiveness of the proposed program within the target population or community theoretically justified and supported with epidemiological, methodological, and behavioral research? How feasible is the implementation of the efficacy or effectiveness study as proposed?

2. **Dissemination.** Are there plans for the dissemination of findings and the sharing and release of data? Are these plans well articulated? Do the plans include provision for disseminating findings to stakeholders outside of academia?

Protection of Human Subjects from Research Risks: Does the application adequately address the requirements of title 45 CFR part 46 for the protection of human subjects? Not scored; however, an application can be disapproved if the research risks are sufficiently serious and protection against risks is so inadequate as to make the entire application unacceptable.

Inclusion of Women and Minorities in Research: Does the application adequately address the CDC Policy requirements regarding the inclusion of woman, ethnic, and racial groups in the proposed research? This includes: (1) The proposed plan for the inclusion of both sexes and racial and ethnic minority populations for appropriate representation; (2) The proposed justification when representation is limited or absent; (3) A statement as to whether the design of the study is adequate to measure differences when warranted; and (4) A statement as to whether the plans for recruitment and outreach for study participants include the process of establishing partnerships with community(ies) and recognition of mutual benefits.

Inclusion of Children as Participants in Research Involving Human Subjects: The NIH maintains a policy that children (*i.e.*, individuals under the age of 21) must be included in all human subjects research, conducted or supported by the NIH, unless there are scientific and ethical reasons not to include them. This policy applies to all initial (Type 1) applications submitted for receipt dates after October 1, 1998.

All investigators proposing research involving human subjects should read the "NIH Policy and Guidelines" on the inclusion of children as participants in research involving human subjects that is available at <http://grants.nih.gov/grants/funding/children/children.htm>.

Budget: The reasonableness of the proposed budget and the requested

period of support in relation to the proposed research.

2. Review and Selection Process

Applications will be reviewed for completeness by the Procurement and Grants Office (PGO) and for responsiveness of the eligibility information by the NCIPC. Incomplete applications and applications that are non-responsive to the eligibility criteria will not advance through the review process. You will be notified that the application did not meet submission requirements.

Applications that are complete and responsive to the PA will be subjected to a preliminary evaluation (streamline review) by a peer review committee, the Initial Review Group (IRG), convened by NCIPC, to determine if the application is of sufficient technical and scientific merit to warrant further review by the IRG. CDC will withdraw from further consideration applications judged to be noncompetitive and promptly notify the principal investigator or program director and the official signing for the applicant organization. Those applications judged to be competitive will be further evaluated by a dual review process.

1. The primary review will be a peer review conducted by the IRG. All applications will be reviewed for scientific merit in accordance with the review criteria listed above. Applications will be assigned a priority score based on the National Institutes of Health (NIH) scoring system of 100–500 points.

2. The secondary review will be conducted by the Science and Program Review Subcommittee (SPRS) of NCIPC's Advisory Committee for Injury Prevention and Control (ACIPC). The ACIPC Federal agency experts will be invited to attend the secondary review, and will receive modified briefing books (*i.e.*, abstracts, strengths and weaknesses from summary statements, and project officer's briefing materials). ACIPC Federal agency experts will be encouraged to participate in deliberations when applications address overlapping areas of research interest, so that unwarranted duplication in federally funded research can be avoided and special subject area expertise can be shared. The NCIPC Division Associate Directors for Science (ADS) or their designees will attend the secondary review in a similar capacity as the ACIPC Federal agency experts to assure that research priorities of the announcement are understood and to provide background regarding current research activities. Only SPRS members will vote on funding recommendations,

and their recommendations will be carried to the entire ACIPC for voting by the ACIPC members in closed session. If any further review is needed by the ACIPC, regarding the recommendations of the SPRS, the factors considered will be the same as those considered by the SPRS.

The committee's responsibility is to develop funding recommendations for the NCIPC Director based on the results of the primary review, the relevance and balance of proposed research relative to the NCIPC programs and priorities, and to assure that unwarranted duplication of federally-funded research does not occur. The secondary review committee has the latitude to recommend to the NCIPC Director, to reach over better-ranked proposals in order to assure maximal impact and balance of proposed research. The factors to be considered will include:

a. The results of the primary review including the application's priority score as the primary factor in the selection process.

b. The relevance and balance of proposed research relative to the NCIPC programs and priorities.

c. The significance of the proposed activities in relation to the priorities and objectives stated in "Healthy People 2010," the Institute of Medicine report, "Reducing the Burden of Injury," and the "CDC Injury Research Agenda."

All awards will be determined by the Director of the NCIPC based on priority scores assigned to applications by the IRG, recommendations by the secondary review committee, *e.g.*, ACIPC, consultation with NCIPC senior staff, and the availability of funds.

VI. Award Administration Information

1. Award Notices

Successful applicants will receive a Notice of Grant Award (NGA) from the CDC Procurement and Grants Office. The NGA shall be the only binding, authorizing document between the recipient and CDC. The NGA will be signed by an authorized Grants Management Officer, and mailed to the recipient fiscal officer identified in the application.

2. Administrative and National Policy Requirements

45 CFR Part 74 or 92

For more information on the Code of Federal Regulations, see the National Archives and Records Administration at the following Internet address: <http://www.access.gpo.gov/nara/cfr/cfr-table-search.html>.

The following additional requirements apply to this project: