

Society of Surgical Oncology Clinical Investigator Award in Breast Cancer Research

Funded by Susan G. Komen for the Cure[®]

Purpose

- To promote patient-oriented breast cancer research conducted by surgical oncologists in clinical and translational science.

Eligibility

- Applicants must be surgical oncologists within 10 years of completion of training and be full SSO members.
- Applicants must commit to at least 25% effort for this award.

Terms of the Award

- The award will be funded for two years at \$100,000 (\$50,000 per year) beginning October 1, 2009.
- The award is given to the sponsoring institution and may be used for partial support of applicant's salary, research fellows or staff support, tuition, travel (no more than \$2,000 per year) and/or supplies. No additional funds are available for paying indirect costs.
- A brief year-one progress report will be required by October 31, 2010. The SSO reserves the right to withhold the second year of support in the event of unsatisfactory progress.
- A final report (not to exceed 4 pages) will be required before October 31, 2011. This should include a summary of the project and abstracts and publications acknowledged as supported by this award.
- A brief 5-minute presentation will be required at the 2012 Annual Meeting.

Application Procedure

The SSO envisions a proposal in which the applicant plays a central role in the conduct of a specific clinical breast cancer research project. This might include a leadership role in a clinical trial involving prevention, diagnosis, or therapy; cancer outcomes research; survivorship research; or a translational research project related to a prospective clinical trial. The clinical research focus must be hypothesis-driven and must have a direct patient-oriented focus. Any clinical trial may be investigator-initiated, industry-driven or organized by a cooperative group.

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While the applicant need not be the principal investigator of the trial, nor the lead institutional investigator, how participation in the trial will serve to enhance their research experience must be clearly articulated. It is anticipated that applicants will be early to mid-career and are seeking extramural support to further their goals of establishing an independent career in breast cancer research. The potential and track record of the applicant, the training plan and environment, and the scientific merit of the clinical trial will comprise the review criteria.

The application MUST include the following items in order:

- A. Cover page. The cover page must include the following information: the title of the proposal, name of applicant, applicant's position, institution and contact information, and the name and contact information for the individual authorized to act for the applicant if the award is made.
- B. Curriculum Vitae: A three-page biosketch (including other support) in the standard NIH format.
- C. Supporting Letters. A letter from the applicant's institution is required and should include a review of the proposal and summary of the applicant's qualifications. It should also describe the facilities and support available to complete the project. The letter should be from the institution's cancer center director, chair of surgery or dean of research. Other supporting letters (no more than 3) may be included.
- D. Research Proposal: This should include the following items: *(limited to eight pages, single spaced, 12 pt font, one inch margin)*.
 1. Abstract: The abstract should describe the clinical trial, the hypothesis being tested, the role of the applicant and how this will serve as a valuable training experience.
 2. Personal statement: The applicant should briefly describe their past accomplishments and how this award mechanism will serve their career goals.
 3. Research plan: This section should focus on the hypothesis being tested, the exact role of the applicant, any didactic course work, and a description of interaction with mentors and colleagues who may also submit supporting letters.
 4. Training environment: This should include a brief description of the clinical cancer trials infrastructure at the applicant's institution, with a specific reference on how it may be used to advantage in this training mechanism.
 5. Literature cited. (not included in 8 page proposal limit)
- E. Budget: In no more than 1 page, briefly describe proposed budget utilization and justification. This should include any required indirect costs that would reduce the actual amount of direct research dollars available to the applicant. The reviewers will take this into account and may not recommend awards when the indirect cost rate is substantial.
- F. Appendix. Clinical protocol, informed consent form, IRB approval notice. **Not to exceed five (5) pages in length.**

The application deadline is July 1, 2009

The complete application must be assembled as a single pdf file and sent electronically to heatherheller@surgonc.org by midnight July 1, 2009.

Applications received after this date, or those that do not conform to the instructions will not be considered. Applicants will be informed of the outcome in September.

PLEASE NOTE: Funding of grants will be determined by scientific peer review process and consistent with any restrictions required by the grantor.