Eligibility

Applicants must...

• be a MD, DO, or PhD
• hold a full-time clinical or research position where the research will be conducted
• be an Active, Candidate or International member of ASPS or obtain sponsorship from an Active ASPS member at your institution
  o This is a requirement if you are currently in your training
  o For multiple PIs, a sponsor is required for only the applying PI

Grant Description:

The goals of this grant program are to accelerate the translation of scientific discoveries and technical developments into practical solutions that improve human health through innovation funding, and to encourage collaborative, transdisciplinary work to accelerate the translation of medical discoveries into improved health. This mechanism provides funding support for early-stage ideas to help researchers complete proof-of-concept studies and develop prototypes that address unmet clinical needs with a robust value proposition. A further goal of this early-stage funding is to enable investigators to gather data that enables future funding, information dissemination and most importantly, momentum that helps to move these novel solutions from the idea stage to clinical use. Note: Transdisciplinary collaborations are encouraged but are not a requirement for funding.

This seed grant(s) is awarded in the areas of medical technology, therapeutics, diagnostics, population health sciences and community engagement. Emphasis will be placed on technologies that have the potential to advance rapidly into patient care through development by venture funded startups, licensing to established companies, or other pathways. Examples include, but are not limited to, prototype device development, preclinical studies, and pilot clinical studies.

Applications will be evaluated based upon the importance of the study question, soundness of study design, demonstration of study feasibility through preliminary/pilot data, the quality of the investigator team, and use of appropriate statistical and analytic methods.

Award Amount: up to $50,000
• Project Duration: Up to 12 months; Note: Projects must be completed and all expenses incurred within that 12-month timeframe; no cost extensions are not permitted.

Earliest Start Date: July 1, 2019
Award Announcements: May 2019
Reporting: Progress reports (technical and financial) are due at six (6) months and twelve (12) months. Details on reporting requirements will be sent to the PI once awarded.

The PSF is dedicated to fostering the growth of research in Plastic Surgery. In order to ensure The PSF is successful in building a diverse, committed, inclusive, and self-sustaining plastic surgery research community, applications may also be evaluated on whether the PI is currently funded by The PSF, the PI has an open no cost extension by The PSF, or whether there are multiple grant submissions from the same lab at the same institution.

The PI or their Sponsor’s donation history to the PSF may also be considered in the review of the application.
**Online Application:** All Applications must be submitted online through proposalCENTRAL. (proposalcentral.altum.com)

**Deadline:** Grants must be submitted in proposalCENTRAL by 11:59 P.M. (EASTERN) on or before the deadline. NO late submissions will be accepted. Corrections of oversights/errors discovered after the deadline will not be allowed.

**Multiple Submissions:** Projects will be awarded only one PSF grant per annual grant cycle. Applicants may submit more than one grant application ONLY if they are scientifically different, but only one research project may be funded.

**Grant Writing Module:** All applicants who have not been previously funded by The PSF must complete The PSF’s Grant Writing Module. For ASPS Members, log on to the PSEN Research and Technology page here: http://www.psenetwork.org/research-technology and complete the free Grant Writing module by Ergun Kocak, MD. For Non-ASPS Members click here: (http://www.thepsf.org/Documents/GrantWriting/story.html). Only the Applicant needs to complete the Module. Upon completion, print the final screen showing your completion of the Module or sign the “Grant Writing Module Certificate” found in proposalCENTRAL templates. Upload either document into the Research Plan & Other Attachments section in proposalCENTRAL.

**Templates:** Budget & Budget Justification, Biosketch, Other Support, Resources and Grant Writing Module Certificate templates are provided on the Research Plan & Other Attachments section in proposalCENTRAL.

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**Deliverables Check List with Page Limits**

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**Information Entered Directly into proposalCENTRAL**

- Project Summary (2,500 characters max, including spaces)
- Impact Statement (800 characters max, including spaces)
- Biography (1,500 characters max, including spaces)

**Documents to Upload- Page Limits will be Strictly Enforced**

- Signed Face Page
- Cover Letter for Resubmissions
- Budget
  - Budget Justification 2 PAGES
  - Biographical Sketch 5 PAGES each
- Other Support
- Resources
- Research Plan:
  - Specific Aims 1 PAGE
  - Research Strategy 12 PAGES
  - Human Subjects 1 PAGE
  - Vertebrate Animals 1 PAGE
  - Literature Cited NO LIMIT
  - Leadership Plan 1/2 PAGE
  - Consortium/Contracts
- IRB /IACUC Approval Documentation* (Each project must submit an IRB or APLAC approval number, or provide confirmation of non-human/non-animal subjects determination within 90 days of award notification)
Sponsor Letter*  
Letters of Support*  
Grant Writing Module*  
Appendix*  

NO LIMIT  
(Upload any/all images as “Appendix” items)

* Submit if applicable – all other documentation is required

Formatting Requirements:
- The minimum acceptable font size is 11 (Arial or Helvetica; no condensed fonts).
- The maximum number of lines per inch is 6; DO NOT PACK LINES BY SETTING LINE SPACING AT “EXACTLY”.
- Use at least one-half inch margins (top, bottom, left and right) for all pages, including continuation pages.

### Key Personnel

**Documentation is required from the following personnel**

<table>
<thead>
<tr>
<th>Personnel</th>
<th>Role Description</th>
<th>Biosketch</th>
<th>Other Support</th>
<th>Letters of Support</th>
<th>Sponsor Letter</th>
</tr>
</thead>
<tbody>
<tr>
<td>PI (Applicant)</td>
<td>Applies for and writes the grant. Oversees and directs the research.</td>
<td>✓</td>
<td>✓</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Multiple-PI “Co-PI”</td>
<td>Shares equal responsibility with PI. Does not need to be from the same institution as the PI.</td>
<td>✓</td>
<td>✓</td>
<td>✓</td>
<td></td>
</tr>
<tr>
<td>Co-Investigator</td>
<td>Highly involved in the scientific development or execution of the project. Contributes measurable effort. Does not need to be from the same institution as the PI.</td>
<td>✓</td>
<td>✓</td>
<td>✓</td>
<td></td>
</tr>
<tr>
<td>Collaborator</td>
<td>Moderately involved in the scientific development or execution of the project. Contributes measurable effort. Typically from the same institution as the PI.</td>
<td>✓</td>
<td></td>
<td></td>
<td>✓</td>
</tr>
<tr>
<td>Sponsor</td>
<td>Required for non-ASPS applicants and those in training. Can also serve another role on the grant but isn’t required to.</td>
<td>✓</td>
<td>*</td>
<td>*</td>
<td>✓</td>
</tr>
<tr>
<td>Other</td>
<td>Research Fellows, Research Assistants, Technicians, Paid Consultants, etc.</td>
<td>*</td>
<td>*</td>
<td>*</td>
<td></td>
</tr>
</tbody>
</table>

*Not required unless also serving in another role, example: Co-Investigator or Collaborator, then follow those guidelines.

### Application Deliverables

Enter the following information directly into the Project Summary section of your application in proposalCENTRAL.

**Project Summary:** This is the summary description of your research project. In language suitable for the public, please describe the project’s broad, long-term objectives and its specific aims. Describe concisely the study design and methods, as well as the rationale, and techniques to be used to achieve the aims.

**Impact Statement:** In language suitable for the general public, describe the potential of real, clinical impact this research is likely to have on the practice of plastic surgery.

**Biography:** In language suitable for the general public, please include a professional biography.
Upload these deliverables in the Research Plan & Other Attachments section of your application in proposalCENTRAL.

Face /Signature Page: After completing your application in proposalCENTRAL, you will need to print your Face Page (Signature page) to obtain your institutional signatures. Once you obtain all signatures, please convert your Face Page to PDF and upload to your application in proposalCENTRAL. Please plan accordingly as obtaining the appropriate signatures may take time. Your grant submission will not be reviewed without institutional sign off.

Budget: Complete all sections of the budget template following these limitations:
- Budgets can NOT include: indirect costs, administrative costs, travel to meetings (presentations), and publications costs. Grant funds will not be used to purchase capital equipment* costing more than 5% of the total budget, intellectual property services, or food.
  - *Capital Equipment is defined as any asset having a useful life of more than one year. Examples include but are not limited to (laptop computers, iPads, software, cameras, cryogenic systems, incubators, dry boxes, cell counters, etc.)
- Budgets can NOT include salary support for PI, Co-PI, Co-Investigators, or Collaborators.
- Budgets CAN include salary for investigator and/or research personnel, travel (if project related) and supplies.

Budget Justification (2 PAGES): Please clarify and describe the purpose and need for each item listed on the Budget page, i.e., Personnel, Consultant Costs, Supplies, etc. Under Personnel, roles for each person should match what you’ve entered into proposalCENTRAL. For a study that requires more resources than this grant provides, please describe any institutional or other support you have.

Biographical Sketch (5 PAGES): Submit an NIH biosketch for those directly involved with the project (see chart above for required personnel). Expand the space for educational training if necessary. If you are omitting publications due to space limitations, include the statement: “The following publications were selected from among a total of ____ (#).” List selected ongoing or completed (during the last three years) research projects. Begin with the projects that are most relevant to the research proposed in this application. Include the project number, dates, source of funds, project title, your role in the project and briefly indicate the overall goals of the research project. Do not list award amounts or percent effort in projects. See example provided within the proposalCENTRAL Templates page.

Other Support: Please provide information on all active or pending support from any source (see chart above for required personnel). Include the project number, dates, source of funds, project title, award amounts, percent of effort in months, and briefly indicate the overall goals of the research project. Indicate and explain any scientific or budgetary overlap between funding, and other overarching projects. For individuals with no active or pending support, please indicate NONE. See example provided within the proposalCENTRAL Templates page.

Resources: Limit the description of resources available to those identified on the form.

Upload each section of the Research Plan as separate PDFs into proposalCENTRAL even if you are indicating N/A.

Research Plan (A - G)

A. Specific Aims (1 PAGE): This section should include a brief introductory paragraph. The introduction should give a brief overview of the project and state its significance and central hypothesis. Each Specific Aim shall be comprised of a title, rationale and hypothesis.

B. Research Strategy (12 PAGES): Organize the Research Strategy in the specified order and using the instructions provided below. Start each section with the appropriate section heading - Significance, Innovation, Approach. Cite published experimental details in this section and provide the full reference in the Literature Cited section. Tables and Figures should be included within the text of this section.

Significance
• Explain the importance of the problem or critical barrier to progress or gap in knowledge in the field that the proposed project addresses.
• Explain how the proposed project will improve scientific knowledge, technical capability, and/or clinical practice in one or more broad fields.
• Describe how the concepts, methods, technologies, treatments, services, or preventative interventions that drive this field will be changed if the proposed aims are achieved.

Innovation
• Explain how the application challenges current research or clinical practice paradigms.
• Describe any novel theoretical concepts, approaches or methodologies, instrumentation, or interventions to be developed or used, and any advantage over existing methodologies, instrumentation, or interventions.

Approach
• Discuss the PI’s preliminary studies, data, and/or experience pertinent to this application.
• Provide detailed plans for analysis including statistical methods, control and experimental groups, as well as expected outcomes.
• Describe the overall strategy, methodology, and analyses to be used to accomplish the specific aims of the project. Include how the data will be collected, analyzed, and interpreted as well as any resource sharing plans as appropriate. Please make sure to include discussion of your sample size justification and your power calculations, if applicable.
• Cite relevant publications to support your approach.
• Discuss potential problems, alternative strategies, and benchmarks for success anticipated to achieve the aims. Provide detailed pitfalls and alternative strategies. Consider and discuss how negative data will be used or interpreted.
• If the project is in the early stages of development, describe any strategy to establish feasibility, and address the management of any high risk aspects of the proposed work.
• Point out any procedures, situations, or materials that may be hazardous to personnel and precautions to be exercised.
• State how the findings from this study will inform the next stage of research.
• State expected outcomes and any potential limitations/obstacles to obtaining results.

C. Human Subjects (1 PAGE): If applicable, briefly summarize the use of human subjects, including specific references to adherence of accepted standards, and documentation of the project’s review by an appropriate IRB or hospital ethics board. This includes Protection of Human Subjects, Inclusion of Women and Minorities, Inclusion of Children, and Targeted/Planned Enrollment Tables for Race and Ethnicity. Do not include the entire IRB application. In addition, if available include copies of all Approval Letters from the appropriate IRB Board(s), including BioSafety and/or Radiation Safety (if applicable). Indicate N/A if you are not using Human Subjects within your research.

D. Vertebrate Animals (1 PAGE): If applicable, briefly summarize the use of animals in scientific research, including specific reference to adherence to accepted standards (e.g. NIH publication No. 86 23). In addition, if available upload the documentation of the project’s review by the appropriate institutional committee, including BioSafety and/or Radiation Safety. Indicate N/A if you are not using Vertebrate Animals within your research.

E. Literature Cited: Please list all references in order of occurrence of their first mention in your proposal, in number or superscripted form.

F. Leadership Plan (1/2 PAGE): For applications proposing multiple PIs, or the Applicant has an ASPS Sponsor, a leadership plan is required. The governance and organizational structure should be described, including communication plans and procedures for resolving conflicts. The shared administrative, technical and scientific responsibilities for the project or program should be delineated for the PIs. Indicate N/A if you are not required to submit a leadership plan.
G. **Consortium and Contractual Agreements**: If applicable, describe all research relationships required for this project carefully. Indicate N/A if this is not required.

**IRB/IACUC**: Institutional IRB/IACUC approval letters must be on file in The PSF Executive Offices within ninety (90) days of written notification of the Award. If the approval is not on file within the ninety (90) day time frame, the Award will be rescinded by The PSF. All annual renewals of IRB/IACUC approvals must be sent to The PSF Executive Offices within thirty (30) days of receiving such renewal.

**Accurate Compliance with Institutional Regulatory, Financial and Conflict-of-Interest Policies**

Awardees must be in compliance with all aspects of their institutional policies for human subjects and animal research (IRB/APLAC) approvals; conflict-of-interest (COI) disclosure forms; and NIH/federal policies for allowable and allocable costs.

- For all research that involves live or dead vertebrate animals, an animal use protocol must be reviewed and approved by APLAC prior to commencement of the project.
- Project protocols should be submitted to their institutional IRB as soon as possible after notification of the grant award is received.
- Once investigators are notified of IRB approval/exemption, the study protocol ID# and date of approval/exemption should be submitted via proposalCENTRAL.
- If required, project investigators and their research staff must complete HIPAA (Health Insurance Portability and Accountability Act), and CITI (Collaborative Institutional Training Initiative) training.

**Sponsor Letter**: If the applicant is in training, an original letter from your ASPS Sponsor must be submitted (see chart above for requirements). This letter must verify that the Applicant will be present and fully available to carry out the proposed work during the allotted time period. Applicants, for which this eligibility requirement is not met, will not have their application reviewed.

**Letters of Support**: All Co-PIs, Co-Investigators and Collaborators must submit an original Letter of Support for their involvement in your research project (see chart above for required personnel). Letters of Support are addressed to the applicant and should describe the Co-PI/Co-Investigator/Collaborator’s credibility, intended contribution, role, commitment, and provide support for the work being proposed. If the Sponsor is also serving as a Co-PI, Co-Investigator or Collaborator, then the Sponsor Letter and Letter of Support can be combined. The Sponsor should indicate that they are also serving as a Co-PI, Co-Investigator or Collaborator for the project. A separate letter from each Co-PI/Co-Investigator/Collaborator is required.

**Resubmissions**: In addition to resubmitting all required application documents, resubmissions must be accompanied by a cover letter that summarizes the substantial additions, deletions or changes to the application. If you were provided a summary statement from your previous review, please also include a point-by-point discussion of the issues raised in your summary statement critiques. Please upload your Cover Letter in the Research Plan & Other Attachments section of your application in proposalCENTRAL. Note: Only those resubmitting a grant application must upload a Cover Letter.

**No-cost extensions**: Projects must be completed and all expenses incurred within that 12-month timeframe; no cost extensions are not permitted.

**Additional Funding**: The Principal Investigator may apply for additional funding. Those who want to expand upon their project at the end of the original time period will need to re-apply for additional funding.

**Transferring Institutions**: This grant is NOT transferable to another individual within the Institution and NOT transferable to another institution OR to operating funds. If the Principal Investigator or Sponsor leaves the Institution, the Principal Investigator, Sponsor and the Institution must notify The PSF within thirty (30) days and all unused funds MUST be returned.
Budget Changes: A certain degree of latitude to re-budget within and between budget categories to meet unanticipated needs is allowable. If your expenses fall outside of your original approved budget categories by a degree of more than 10% of your total award, prior written approval will be required, along with a revised budget and a justification for why the changes are necessary to complete your project. In addition, inclusion of any budget category that was not included in the original approved budget must also be approved by The PSF. All reallocation requests are subject to the current grant guidelines and must be approved by The PSF.