

# Society of Urologic Nurses and Associates Young Investigators Program

# **Application for Clinical Research Support**

Date:	
Name:	
Address:	
Phone #:	Fax #:
E-mail URL:	
SUNA Membership #:	Preferred contact route:
I am aware that acceptance of this research scholar sharing results via podium presentation at a future publication in UNJ with acknowledgement for SUNA	SUNA meeting, submitting the study and results for
My signature also signifies that there is no conflict of gain from any commercial product associated with t	of interest or attachment of this research with personal this research project.
Signature & date	
Title of Research Project:	

## **Research Project Abstract:**

(Brief summary of research focus/objective, targeted participants, and research methods/data collection approaches to be used; 300 word limit.)

## **Society of Urologic Nurses and Associates**

## **Young Investigators Program**

### **Application for Clinical Research Support**

#### **Directions related to the application process**

- 1. Identify a clinical practice issue or concern which warrants research investigation. Develop a preliminary research study proposal. Secure IRB guidelines at the targeted involved institution(s) and follow their instructions related to proposal title, consent forms, etc. Set default drivers to standard margins (1 or 1.5 inch all around), double space text, easy to read font type (not script), font size of 10 or11, and logical pagination.
- 2. Complete the Application for Clinical Research Support cover sheet.
- 3. Develop the <u>Description of Research Project</u> section using the directions provided. Submit this section in an anonymous format; identify pages with only a running head of no more than 5 key words (i.e., "A research study exploring the urinary infection following with renal stenosis" would have a running head of "stenosis & infection").
- 4. Complete an <u>Investigator Biographical Sketch</u> for each person significantly involved in the conduct of the research. Sketches are not required for data collection assistants if their involvement is limited to data collection only.
- 5. Submit the materials as one unit to the SUNA National Office, East Holly Ave, Box 56, Pitman, NJ 08071. Materials should include:

Application for Clinical Research Support, Description of Research Project, Investigator Biographical Sketches

Applications may be emailed to <a href="mailto:suna@ajj.com">suna@ajj.com</a> as a PDF file.

Faxed versions will **not** be accepted.

#### Funding

Upon acceptance of the application, funds will be released in the following manner: 25% upon receipt of the agreement; 25% upon receipt of an interim report, 25% upon receipt of final report and 25% upon presentation at a SUNA national conference and submission of the study and results for publication in UNJ.

#### **Description of Research Project**

#### **Directions**

#### **Purpose of the study**

In a few sentences, briefly describe the purpose of the study. May be stated as a problem statement, hypothesis or study question.

#### Significance of the study

Briefly describe the significance of this problem to or within urologic nursing practice. This section is limited to  $\frac{1}{2}$  page.

#### **Background of the problem**

What is the issue or concern to be addressed? Why is it a practice problem? How does your study contribute to what has already been done (AKA extending evidence-based practice)? This section is limited to 1 page.

#### **Methods**

Describe how you will accomplish the objectives. Include a description of the setting, criteria for subject/participant selection and sample size, and your method for subject/participant recruitment. If variables will be manipulated or measured, describe that process also. Finally, describe data collection and analysis processes. Attach a copy of the instrument that will be used in data collection. This section is limited to 3 pages.

#### **Ethical conduction**

Describe how voluntary and informed consent is assured. In what ways will anonymity and confidentiality be addressed? How will you minimize hazards, discomforts and negative effects of your research? Append the informed consent form and documentation of Institutional Review Board approval to this section of your application packet.

#### **Study Budget**

Provide a budget associated with the conduction of the study. Include supplies, equipment, special staff, salary and other expenses needed to complete the study. Justification for budget items are required. A project timeline must also be included.

#### References

Include full citations only for references cited in the application. Do not include references supporting the academic side of the proposal.

#### **Appendices**

As indicated above, attach the informed consent form, instruments to be used in data collection and IRB approval documentation.

# **Biographical Sketch**

Name:	
Professional Title:	
Role in research study: Primary investigator Other (explain)	Associate Investigator

Academic and experiential qualifications for this study: