

# **Research Authorization Form (RAF)**

*To be completed prior to submission of an IRB/IACUC application. Please see instruction form for an explanation of each section.* 

# **RAF Submission Date:**

# Section 1: General project information

- a. Principal Investigator (First, Last):
- b. Full project title (and abbreviation, if applicable). Project title <u>must</u> match IACUC or IRBNet submission:
- c. Brief project overview:

d.	Type of submission:	O New (	С	Renewal		t		
	Previous RAF # (if renewal or amendment):							
e.	Type of research:	O Animal (	С	Human	Other (desc	ribe):		
f.	Study classification:	Survey	0	bservational	Clinical tria	I		
		Other (desci	ribe): _					
	i. Clinical trial phase Phase 0 Phase 1 Phase 2 classification (check all that apply): Phase 3 Phase 4							
			11030 3		•			
Section 2: Funding information								
		deral grant (e.g. N	IH, DO	O(D	ther non-profit g	rant (e.g. AAO, Fight for Sight)		
	PI	discretionary fund	ls	R	equest for SIVR fi	unding		
	Industry: sponsor initiated Industry: investigator initiated							
	Ot	ner (please list):						
b.	Name of funding agency:							
C.	Type of grant (e.g., R01, K2	3, New Investigato	or Awa	rd):				
d.	Total project period (MM/Y	'Y - MM/YY):						
e.	Funds requested i. Year one: Direc ii. All years: Direc		Indir Indir	ect \$ ect \$	Total \$ Total \$			
f.	iii. N/A (1 year or l F&A Rate:	ess):						
Section	a 3: Shared resources							
a.	Will additional space or inst	itutional resource	s be re	equired?	ONO	OYes (describe below)		

- b. Will any additional equipment be borrowed or donated?
- Yes (complete below) i. Will any be borrowed from the college (UEC, CVRC, other labs)? Indicate name and location below.

ii. Will any be borrowed/loaned from an outside entity? Indicate name and provider below.

c. Will the study require additional faculty/staff release time?



No

Yes (describe below)

## Section 4: Study personnel and compliance

a. List all key personnel and their corresponding Role in the study.

Please enter the information required per column. If any changes were made on the annual FCOI disclosure after November 1<sup>st</sup>, kindly enter the last date that it was updated.

#### By marking as completed, the Principal Investigator confirms that all listed personnel have an up-to-date and accurate Annual Financial Conflicts of Interest (FCOI) for the current Fiscal year, and that they have completed all required CITI Training Courses.

Name (First, Last,	Role in study	Comple	eted the Annual	Completed	Expiration Dates for required
Degrees)	(PI,	FCOI disclosure for		CITI Training	CITI Training
	investigator,	current FY			
	coordinator,	(note date of completion)			
	etc.)				
					Conflicts of Interest:
					<ul> <li>Group 1 All Investigators &amp;</li> </ul>
					Key Personnel:
					<ul> <li>Responsible Conduct of</li> </ul>
					Research (RCR):
					<ul> <li>Conflicts of Interest:</li> </ul>
					<ul> <li>Group 1 All Investigators &amp;</li> </ul>
					Key Personnel:
					<ul> <li>Responsible Conduct of</li> </ul>
					Research (RCR):

Conflicts of Interest:
<ul> <li>Group 1 All Investigators &amp;</li> </ul>
Key Personnel:
<ul> <li>Responsible Conduct of</li> </ul>
Research (RCR):
Conflicts of Interest:
<ul> <li>Group 1 All Investigators &amp;</li> </ul>
Key Personnel:
<ul> <li>Responsible Conduct of</li> </ul>
Research (RCR):
<ul> <li>Conflicts of Interest:</li> </ul>
<ul> <li>Group 1 All Investigators &amp;</li> </ul>
Key Personnel:
Responsible Conduct of
Research (RCR):
Conflicts of Interest:
Group 1 All Investigators &
Key Personnel:
Responsible Conduct of
Research (RCR):
Conflicts of Interest:
Group 1 All Investigators &
Key Personnel:
Responsible Conduct of
Research (RCR):
Conflicts of Interest:
<ul> <li>Group 1 All Investigators &amp;</li> </ul>
Key Personnel:
<ul> <li>Responsible Conduct of</li> </ul>
Research (RCR):
Conflicts of Interest:
Group 1 All Investigators &     Kay Personnely
Key Personnel:
Responsible Conduct of
Research (RCR):
Conflicts of Interest:
Group 1 All Investigators &
Key Personnel:
Responsible Conduct of
Research (RCR):
<ul> <li>Conflicts of Interest:</li> </ul>
<ul> <li>Group 1 All Investigators &amp;</li> </ul>
Key Personnel:
<ul> <li>Responsible Conduct of</li> </ul>

## Section 5: Animal subjects research information

- a. Please attach NIH style budget page (<u>https://grants.nih.gov/grants/funding/phs398/fp4.pdf</u>)
- b. Species:
- c. Strain(s):
- d. Number of animals for each species/strain:

#### Section 6: Human subjects research information

- a. Population
  - i. Number of subjects to be enrolled:

ii. Will minors (under age 18) be enrolled?

iii. Study entry criteria (inclusion/exclusion):

b. Procedures and treatment

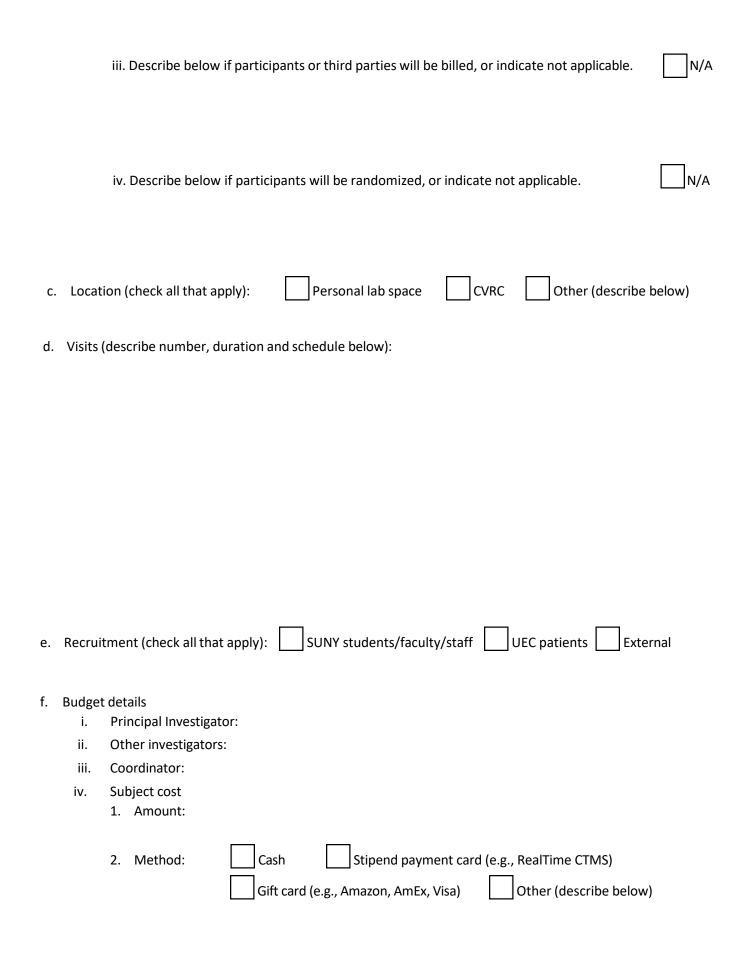
i. Tests/equipment to be used:

ii. Describe below any clinical treatments to be provided, or indicate not applicable.



Yes

No



v.	Start-up/closeout/admin costs. Indicate if not applicable.	N/A
vi.	Advertising costs. Indicate if not applicable.	N/A
vii.	Other costs (supplies, travel, publication, etc.). Indicate if not applicable.	N/A

# g. Other/Notes

## Section 7: Certification

I certify that the information provided in this form is accurate and complete and that I will abide by federal, state, College, and, Research Foundation guidelines and regulations while conducting this research.

**PI Signature:** 

Date:

# SECTION BELOW FOR ADMINISTRATIVE USE ONLY

Does this project require a Conflict of Interest (COI) management plan for any of the listed study team members? (If applicable, please provide details in the Notes section below and include the relevant plan(s) for the IRB to review.)

Yes No

Notes:

RAF #:

Approval date:

**Approver's Signature:**