

“How I Did It” Extramural Funding Applications

Saieesh Rao

4/10/2025

Awards I applied for and secured

- **F32 Ruth Lillian Kirschstein National Research Service Award from the Agency for Healthcare Research and Quality (AHRQ F32)**
 - \$174,497 over two years
- **American College of Surgeons Resident Research Scholarship (ACS)**
 - \$60,000 over two years
- **Association for Academic Surgery / AAS Foundation Trainee Research Fellowship Award (AAS)**
 - \$30,000 over one year

- Accepts applications in April, August, and December
- Allows 1 resubmission attempt if rejected
- Number of awards limited by federal budget allocation
- No interview

- One submission deadline annually in September
- 5 awards per year
- No interview

- One submission deadline annually in August
- 4 awards per year, one per research category
- Has an interview

Disclaimers

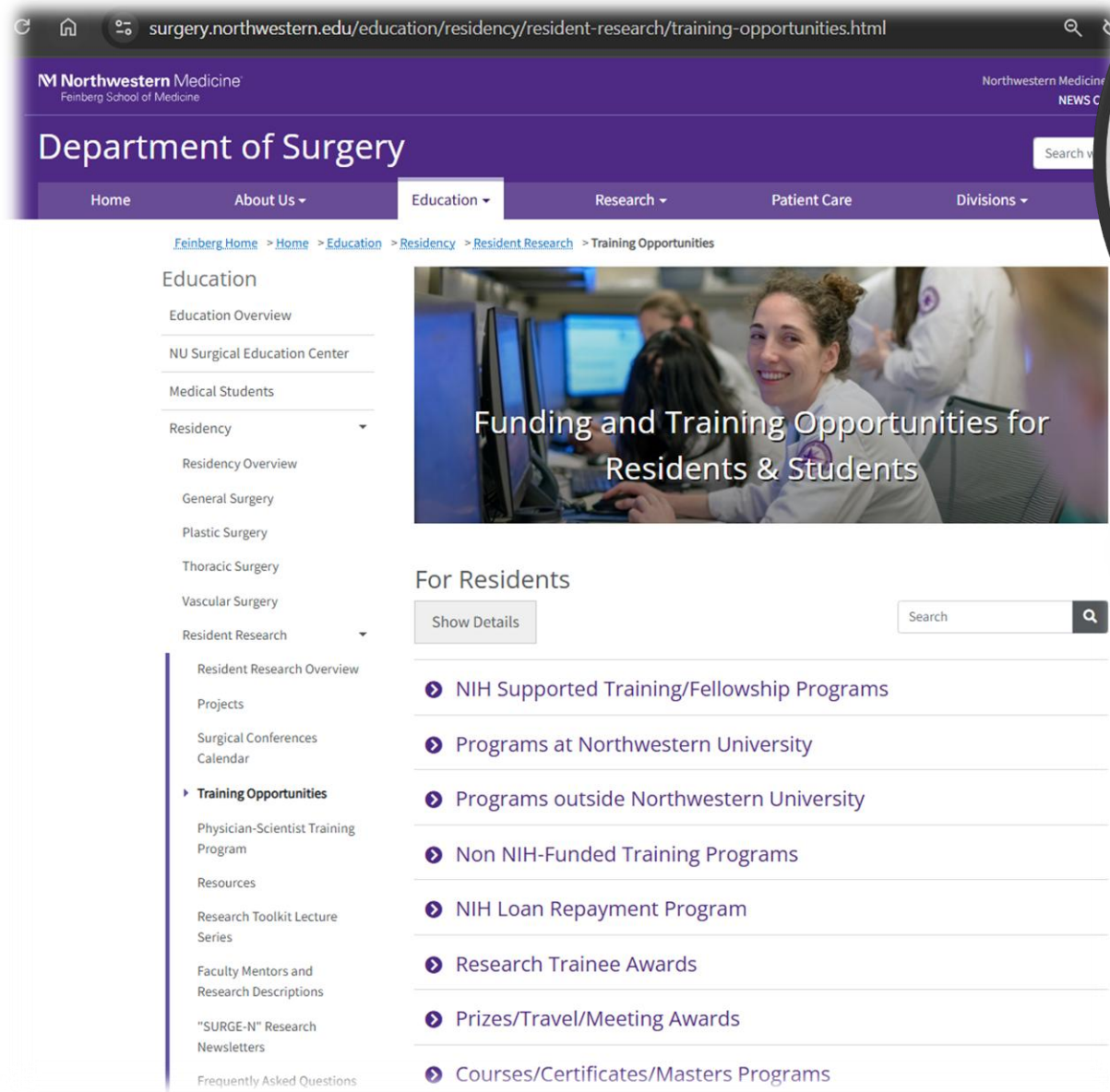
- This presentation represents one person's perspective
- This presentation combines anecdote, objective fact, and opinion
- I will make strong recommendations based on my experience, YMMV
- The eventual success of the applications is in larger part due to support from mentors and assistance from previous applicants
- The federal funding environment is in flux and may make my strategy less applicable in the future

The Most Important Thing

TIMELINE

- Timeline, timeline, timeline
- If you are very serious about getting **individual award** funding, start thinking about it at end of **intern year**
 - “The best time to start was back then... the next best time is now”
 - April PGY2 is still doable!
- Other training grants (T32, R38) available to Northwestern residents are applied for in **third year**
 - **These are your backups, not the goal**
- What does the timeline look like, actually?
- What is the strategy to maximize your efficiency applying for awards?

How to Identify Awards



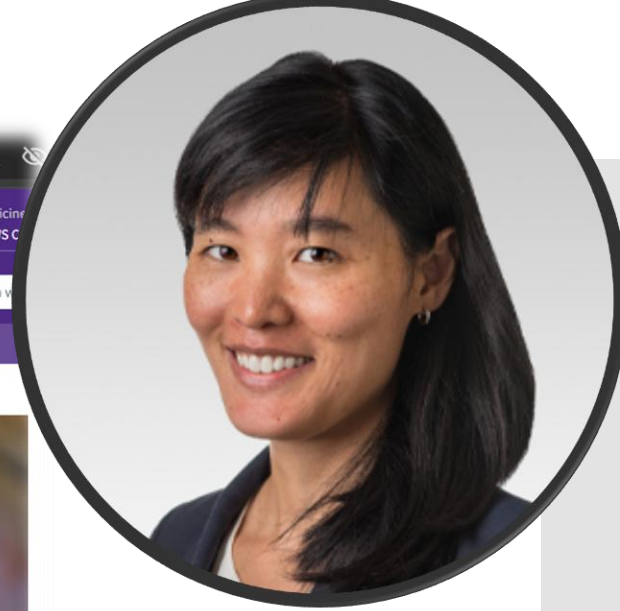
The screenshot shows a web browser at the URL surgery.northwestern.edu/education/residency/resident-research/training-opportunities.html. The page header includes the Northwestern Medicine logo and the Department of Surgery name. A navigation menu is visible with options: Home, About Us, Education (selected), Research, Patient Care, and Divisions. A breadcrumb trail reads: [Feinberg Home](#) > [Home](#) > [Education](#) > [Residency](#) > [Resident Research](#) > [Training Opportunities](#).

The left sidebar contains a navigation menu under the heading "Education":

- Education Overview
- NU Surgical Education Center
- Medical Students
- Residency
 - Residency Overview
 - General Surgery
 - Plastic Surgery
 - Thoracic Surgery
 - Vascular Surgery
- Resident Research
 - Resident Research Overview
 - Projects
 - Surgical Conferences Calendar
 - Training Opportunities**
 - Physician-Scientist Training Program
 - Resources
 - Research Toolkit Lecture Series
 - Faculty Mentors and Research Descriptions
 - "SURGE-N" Research Newsletters
 - Frequently Asked Questions

The main content area features a banner image of a smiling woman in a lab coat with the text "Funding and Training Opportunities for Residents & Students". Below the banner is a section titled "For Residents" with a search bar and a "Show Details" button. A list of training opportunities is displayed, each with a right-pointing arrow icon:

- NIH Supported Training/Fellowship Programs
- Programs at Northwestern University
- Programs outside Northwestern University
- Non NIH-Funded Training Programs
- NIH Loan Repayment Program
- Research Trainee Awards
- Prizes/Travel/Meeting Awards
- Courses/Certificates/Masters Programs



List of Awards on the Website

NIH Supported Training/Fellowship Programs

- NIH Ruth L. Kirschstein National Research Service Award (NRSA) Individual Postdoctoral Fellowship (F32)

Programs at Northwestern University

- Chicago Kidney Urology Hematology network FOR city-Wide reseArch tRaining and career Development (Chicago KUJ FORWARD) (Deadline - Rolling basis until slots are filled)
- NQUIRES NCI Resident Research Opportunity in Surgical Oncology (PDF) (NIH/NCI R38) (Deadline 9/29/23)
- NQUIRES Resident Research Opportunity in Surgical Oncology (PDF) (NIH/NCI T32) (Deadline 9/29/23)
- Northwestern University Minority Health Disparities Research Training Program (NIH/NIMDH T37)
- TL1 - Multidisciplinary Training Program in Child and Adolescent Health
- Vascular Surgery Scientist Training Program (NIH/NHLBI T32) (Latest application date 3/1/24)
- Transplant Surgery Scientist Training Program
- NU School of Professional Studies Advanced Graduate Certificates (Including Health Informatics and Artificial Intelligence)
- NU-THRIVE: Postdoctoral T32 Fellowship in Translational Science, HIV, and Sexual and Gender Minority Health
- The University of Chicago and Northwestern University Postdoctoral Health Services Research Program Postdoctoral Fellowship
- Robert J. Havey, MD Institute for Global Health

Programs outside Northwestern University

- The Advanced Immunobiology T32 Training Program (AITP) at Duke University (Deadline 2/1/24)
- Training of Academic Surgical Oncologists T32 Training Program at MD Anderson Cancer Center
- T32 NIH Training Grant Position at Boston University (DOCX)
- University of Arizona Aerospace Medicine and Surgery Fellowship

Non NIH-Funded Training Programs

- American College of Surgeon Clinical Scholars in Residence
- American College of Surgeons Scholarships
- National Numbered Surgical Education Trials Group Research Fellowship (2 years)
- Howard Hughes Medical Institute Hanna H. Gray Fellows Program (Deadline 12/1/23; Internal Deadline 11/27/23)

NIH Loan Repayment Program

- NIH Loan Repayment Program (Deadline 11/21/24)

Research Trainee Awards

- Steven J. Stryker, M.D., Gastrointestinal Surgery Research and Education Endowment (Deadline 3/1/24)
- Association of Academic Surgeons (AAS)/AAS Foundation Trainee Research Awards (in areas of basic science, clinical, and education) (Deadline 8/21/23; internal deadline 8/9/23)
- AATS David J. Sugarbaker Surgical Resident Investigator Award (Deadline 12/1/23; internal deadline 11/20/23)
- AATS Foundation Programs (Search "Residents" for full list)(Deadline 12/1/23; internal deadline 11/20/23)
- American Association for Cancer Research (many opportunities)
- American College of Surgeons Resident Research Scholarships (Deadline 9/15/23; internal deadline 9/5/23)
- American Heart Association Postdoctoral Fellowship (Deadline 9/7/23; internal deadline 8/28/23)
- American Society of Transplant Surgeons Jon Fryer Resident Research Scholarship (Deadline TBA)
- American Society of Transplant Surgeons Veloxis Fellowship in Transplantation (Deadline TBA)
- The MacLean Center for Clinical Medical Ethics at the University of Chicago (Deadline 1/15/24; internal deadline 1/4/24)
- Nina Starr Braunwald Research Award (Thoracic Surgery Foundation) (Deadline 9/15/23; internal deadline 9/5/23)
- Society of American Gastrointestinal and Endoscopic Surgeons (Deadline 10/1/23; internal deadline 9/19/23)
- Society of University Surgeons Resident Research Scholar Award (Deadline 6/28/24; internal deadline 6/18/24)
- Surgical Outcomes Club Michael Zinner HSR Fellowship (Deadline May 2024)
- Thoracic Surgery Foundation Resident Research Fellowship (Deadline 9/15/23; internal deadline 9/5/23)
- VA Chief Resident in Quality and Patient Safety Program (CRQS)
- Vascular and Endovascular Surgical Society/Medtronic Vascular Resident Research Award (Deadline 10/6/23; internal deadline 9/22/23)

Prizes/Travel/Meeting Awards

- 2023 AMA Research Challenge
- AOA Postgraduate Fellowship (for 2K) (Deadline 5/28/24; Internal deadline is 5/28/24)
- Midwestern Vascular Surgical Society travel scholarship
- Midwestern Vascular Surgical Society Trainee Awards
- The Society of Asian Academic Surgeons Foundation Academic Surgical Congress Research Award (Deadline 8/6/23; internal deadline 7/26/23)

Others not shown:

- ASCRS (colorectal) Resident Research Fellowship
- Am Assoc Thoracic Surgery
- Plastic Surgery Foundation
- Probably more...

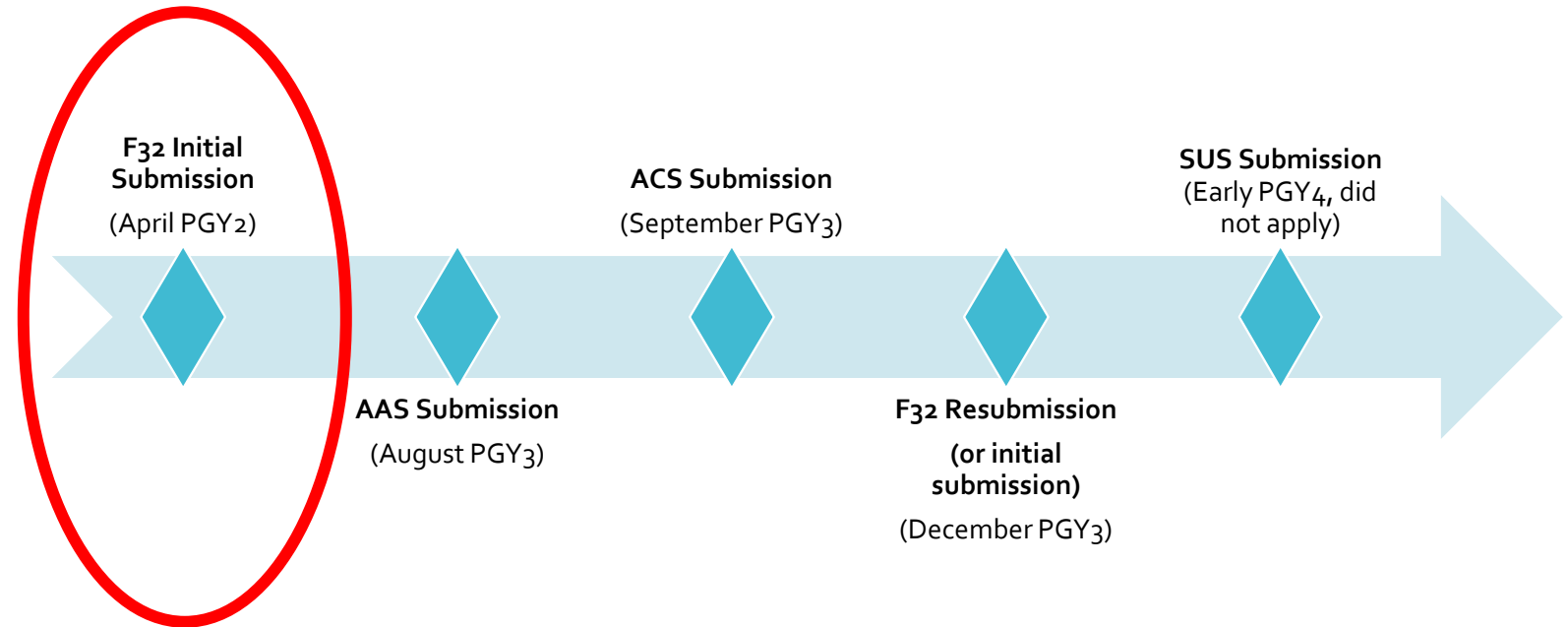
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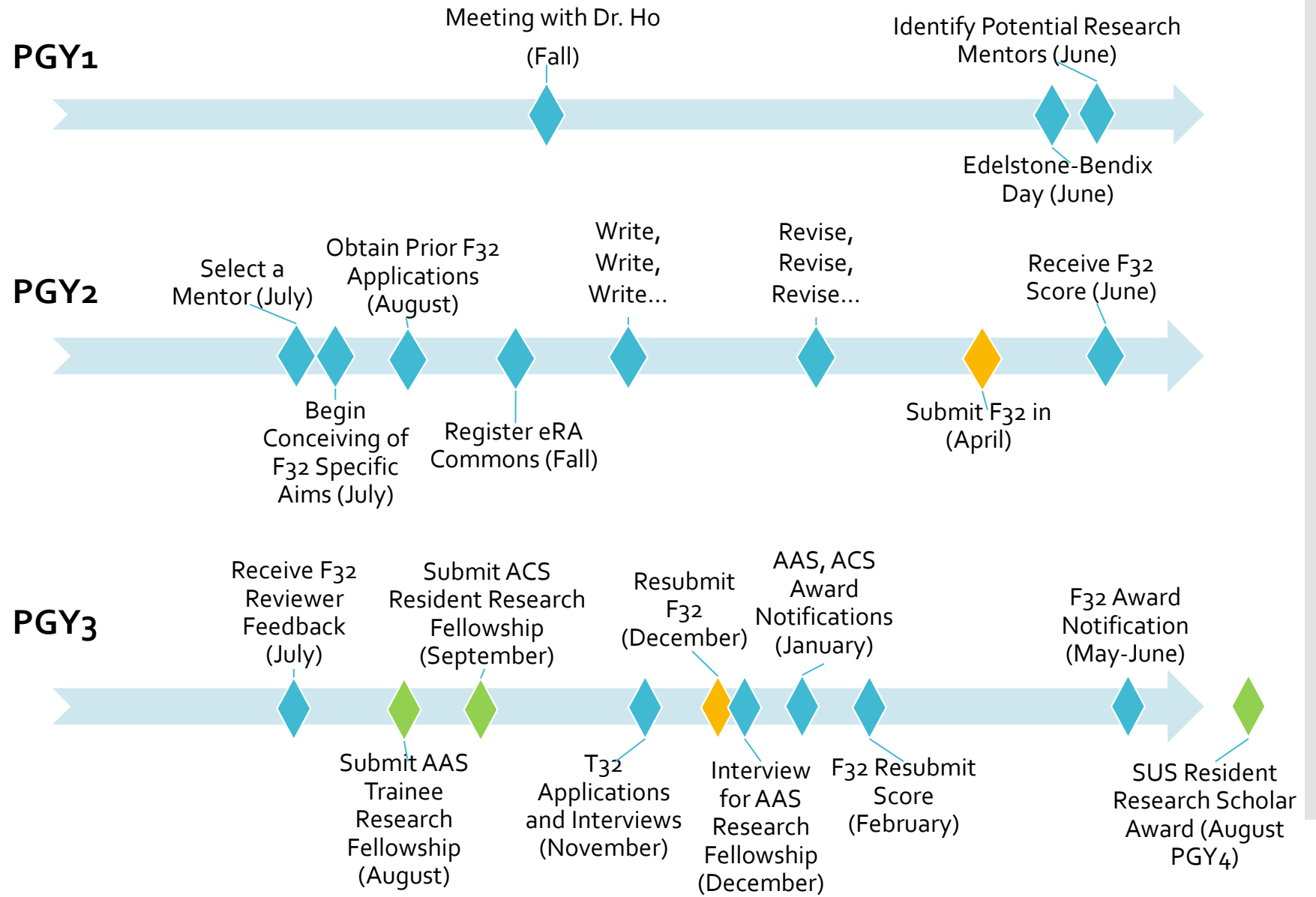
Courses/Certificates/Masters Programs

- ACS Leadership and Advocacy Summit (April)
- ACS Residents as Teachers and Leaders (March)
- Artificial Intelligence at Northwestern
- FSM Center for Education in Health Sciences Master of Science in HSOR
- Institute of Image-Guided Surgery (IHU Strasburg) Diploma Program
- McGaw Medical Education Clinical Scholars Program (Programs in Bioethics, Global Health, Health Equity and Advocacy, and Medical Education)
- NU Center for Leadership Fellowship in Leadership
- NUCATS Master of Science in Clinical Investigation
- School of Professional Studies Artificial Intelligence Certificate Program

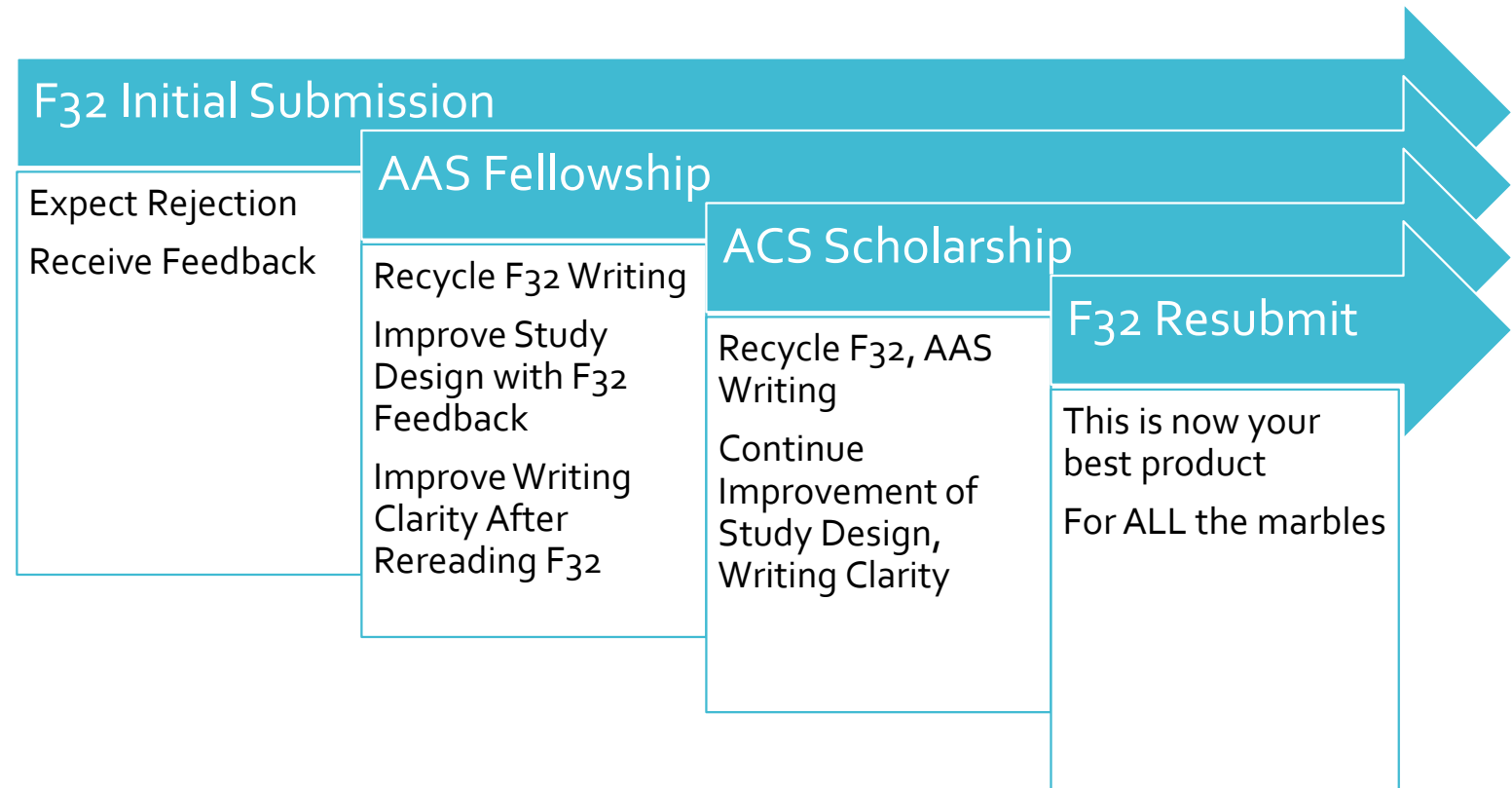
The Timeline



The Timeline



The Strategy: Make the most of the opportunity to resubmit your F32



- The latest you can submit an F32 in time to start research years with funding is December PGY₃
- Working backwards gives us the April PGY₂ deadline for your initial F32 submission, so you can resubmit in December PGY₃
- This conveniently provides the window to recycle the initial F32 submission for other award applications at the start of PGY₃
- Not shown: Society of University Surgeons (SUS) Resident Research Scholar Award applications are due in **August PGY₄** if you don't get any of these funds

Comparison of Award Submission Requirements

The society awards can be written entirely with F32 materials

Remember to modify letters, cover pages for specific awards

	F32	ACS	AAS
Cover Letter	1 page	1 page introducing the applicant, career objectives, training plan	1 page
Introduction	1 page (resubmission only)		
Project Summary/Abstract	1 page summarizing specific aims, training plan, and environment		1 page
Project Narrative	2-3 sentence summary		
Specific Aims	1 page	1 page	5 pages
Research Strategy	6 pages	3 pages	
References Bibliography	No Limit	1 page	
Background & Goals of Fellowship Training	6 pages		
Sponsor / Co-Sponsor Statements	6 pages		
Letters of Support	6 pages		
Letters of Reference	At least 3, no more than 5	1 chair letter, 1 research PI letter	1 chair letter, 1 research PI letter
CV / Biosketch	Applicant, Sponsors, Co-Sponsors	No	Applicant, Research PI
Facilities & Other Resources	No Limit		
Equipment	No Limit		
Human Subjects Protections	8 subcategories, <1 page each, however extensive PHS Human Subjects and Clinical Trials Form		
Respective Contributions	1 page		
Selection of Sponsor and Institution	1 page		
Responsible Conduct of Research	1 page		
Institutional Environment and Commitment to Training	2 pages		
Budget and Justification	No Limit	1 page	1 page

Using Peer Materials

- Jessie shared her final F32 submission materials as well as those of prior applicants (successful and not)
- Much of the language was recyclable
- There were differences between basic science vs. health services research
 - Particularly regarding facilities, equipment, selection of institution given my focus was different
- Dr. Ho has collected these in a central repository for internal use

Share Copy link Add shortcut to My files Download

My files > F32 resources

Name	Modified	Modified By	File size
Reiter F32 documents	January 17, 2023	Reiter, Audra	0 items
1F32HL162378-01-Summary Statement.pdf	October 23, 2022	Ho, Jessie	139 KB
FINAL-Activities Planned-JWH.docx	October 23, 2022	Ho, Jessie	22.8 KB
FINAL-Assembled Proposal v2.pdf	October 23, 2022	Ho, Jessie	2.61 MB
FINAL-Authentication of Key Biological-J...	October 23, 2022	Ho, Jessie	14.8 KB
FINAL-biosketch-JWH.docx	October 23, 2022	Ho, Jessie	48.1 KB
FINAL-Budget_justification Jessie Ho F32...	October 23, 2022	Ho, Jessie	23.2 KB
FINAL-Concurrent Support-JWH.docx	October 23, 2022	Ho, Jessie	12.4 KB
FINAL-Cover Letter-JWH.pdf	October 23, 2022	Ho, Jessie	33.3 KB
FINAL-Cover Letter-JWH-3.pdf	October 23, 2022	Ho, Jessie	70.0 KB
FINAL-Doctoral Dissertation and Researc...	April 1, 2023	Sanchez, Joseph	24.1 KB
FINAL-Equipment-JWH.docx	October 23, 2022	Ho, Jessie	14.9 KB
FINAL-Facilities and Other Resources-JW...	October 23, 2022	Ho, Jessie	24.2 KB
FINAL-Institutional Environment and Com...	October 23, 2022	Ho, Jessie	21.1 KB
FINAL-Keller-biosketch.docx	October 23, 2022	Ho, Jessie	40.5 KB
FINAL-Project Narrative-JWH.docx	October 23, 2022	Ho, Jessie	12.7 KB
FINAL-Project Summary-JWH.docx	October 23, 2022	Ho, Jessie	16.4 KB
FINAL-Resource Sharing Plan.docx	October 23, 2022	Ho, Jessie	12.5 KB

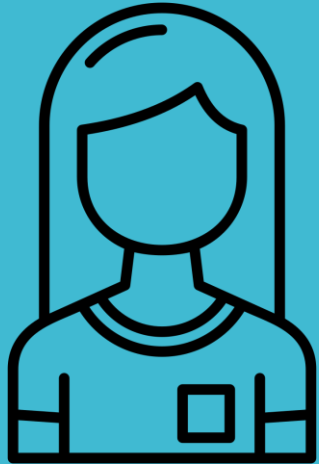
Biggest Challenges for F32

- **Obvious:**
 - Clinical residency remains time consuming and the priority
- **Less Obvious:**
 - The long timeline in mid-PGY2 is deceptive
 - It is NOT a lot of time and it is NOT too early
 - YOU are the primary driver of the proposal
 - You collaborate with your mentor, but YOU are inventing the Specific Aims and Research Strategy
 - This sometimes means making up the research strategy as you go
 - Easy to get stuck or feel you are in quicksand because the amount of work is paralyzing
 - YOU likely will ghostwrite all sponsor, co-sponsor, and reference letters
 - Iterative revisions can be demoralizing
 - See-saw been nitpicking during some weeks and major revisions of research strategy during other weeks
 - The proposal needs multiple sponsors with different expertise
 - You need to find sponsors and obtain their consent; your mentor can help assemble a sponsorship team
 - The training plan is more important than the actual project
 - F32 is a training grant, and the project is a vehicle for training you in research
 - Your proposal will be rejected for a stellar project and inadequate training plan

My Experience

- The **initial F32 submission** required effort which felt overly demanding as a clinical resident
- Despite starting early, it was easy to defer working on non-essential grant writing in favor of clinical residency expectations
- Having prior F32 applications as a reference led to a false sense of security
- **AAS and ACS** applications were significantly easier having already put the thought into the F32 and recycling much of the language
- Adapting the F32 application to the reduced page requirements for AAS and ACS was still laborious
- The **F32 resubmission** was started over a month in advance and still required excessive time wordsmithing, formatting, and crafting a new training plan from scratch
- Input from my mentor and co-sponsors greatly improved the application, but at the same time created more work from incorporating their feedback
- That said, the amount of funding has made research life much easier given increased funding for conference travel, analyst support, etc.
- My experience does not have to be yours
- Future grant applications now appear less daunting – I have muscle memory on how to put together a grant and know what level of effort is necessary, so I can plan better





Testimonial

Anonymous PGY5 #1

- *“I think my year was honestly a little weird with [finding research funding] ...because **we all were so late in finding opportunities and missed a lot of deadlines**”*
- *“I think at the time I wish I knew **1.** What options there were, and what the due dates were **2.** Exactly how to access the grant advising services at NM and if we HAD to use them (since usually means needs to submit [weeks] earlier) and **3.** How many to apply for”*
- *“I think what happened to me was that **I had a sense I wanted to apply for an F32 and the loan forgiveness but didn’t have a project identified in time and couldn’t pull myself together in 3 months**”*
- *“There was sorta like a sense like, you should do this, and really no guidance except just winging it. Like **I wish I had a timeline** but I feel like since then Dr. Ho has shared some of that stuff so hopefully that’s not a problem anymore”*

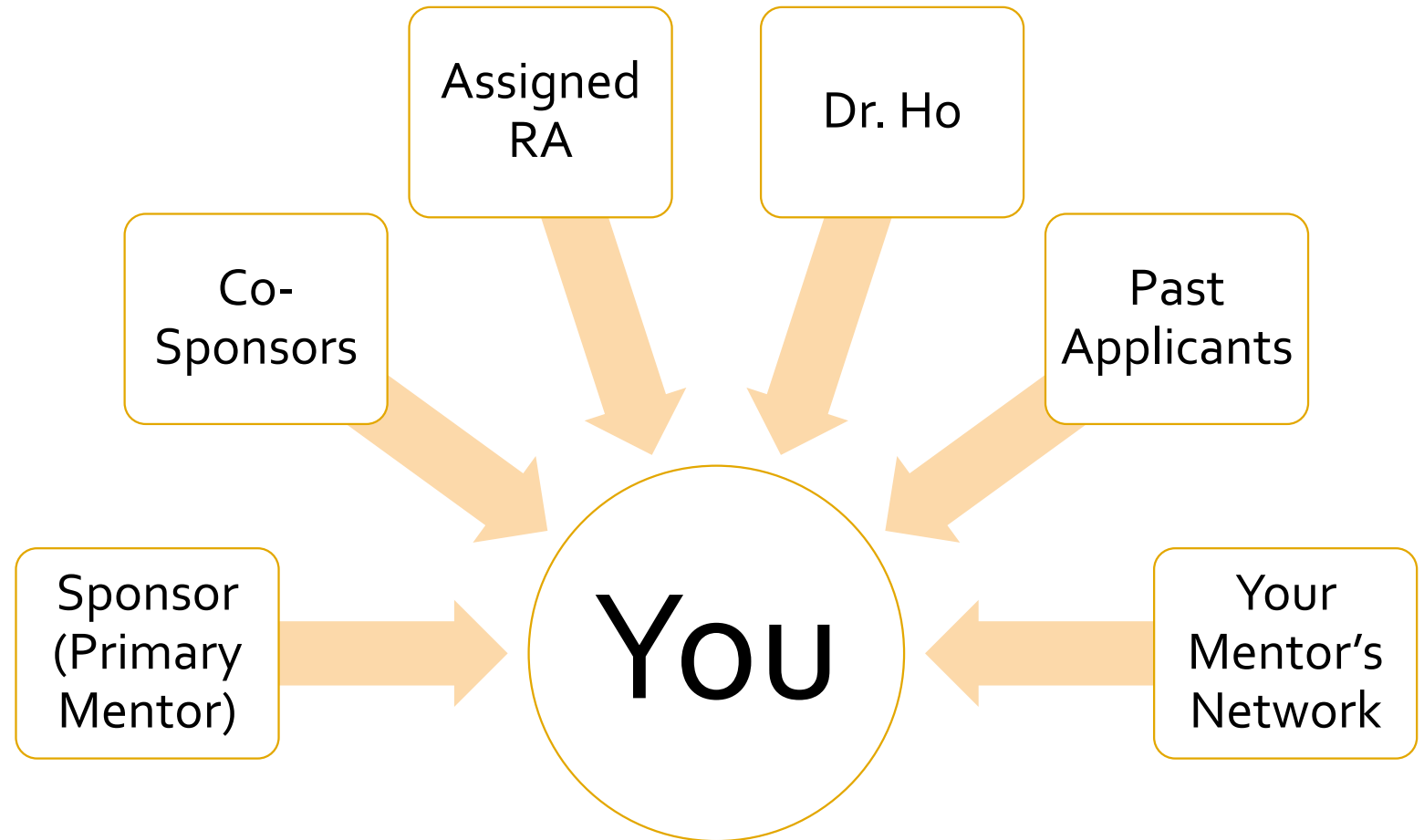


Testimonial

Anonymous PGY5 #2

- *“... I would’ve been more successful **if I had started sooner and had acquired a mentor sooner also.**”*
- *“You **can’t really write one of these things within 3 months like I tried to do.**”*

You Have Support



Finding a Mentor

- Ideally someone with a track record of supporting research residents
- Should be someone you envision yourself getting along with
- Ask prior research residents for their experiences with their mentors
 - Biggest complaint: lack of mentor responsiveness
- **Two approaches:**
 - Choose mentor for their subspecialty:
 - Pros: they can introduce you to colleagues at conferences, you demonstrate interest in the field, you will have much deeper exposure to field which shows on the interview trail
 - Cons: if you change your subspecialty interest (not uncommon), these benefits evaporate somewhat, you might be doing projects you are not really enthused about (clinical research when you want basic science, etc.)
 - Choose mentor because of their research methods:
 - Pros: what you learn will be applicable no matter your future subspecialty, you can brand yourself as a methods expert (“qualitative methods”, “basic science”, “health economics” etc.) and how that relates to your career goals
 - Cons: less opportunity to network with people in your chosen field, you will have to do some outside work to demonstrate interest if important for your eventual subspecialty (publications, conferences, etc.)
- Faculty at conferences insist productivity and project ownership >> subspecialty focus, but there’s likely a balance

Finding a Mentor

- My advice: pick the mentor for their **research methods** because you will be happy regardless
 - Double whammy if you change specialty interest AND don't like basic science / statistics / qualitative interviews, etc.
 - Methods are 90% of your time (what your day-to-day looks like)
 - You can always be a methods expert and apply them to projects in your new field
- People change their minds during research years
 - Research year experiences will convince people for or against certain specialties
 - Research year lifestyle, family events, etc. also play a role
- You can pick up additional projects / mentors during research years, but you need to find a mentor early to start applying for grants


RA Checklist & Internal Deadlines

- Your **research assistant (RA)** is a dedicated expert in grant submissions employed by Northwestern University
- Your RA will be the same as that of your primary mentor (sponsor)
- They will create your account on **eRA Commons** and make a checklist of submission materials for your specific funding opportunity
- They will demand all administrative submission materials to be given **two weeks early** before the funder's submission deadline
 - One week early** for the remainder (scientific content like specific aims, research strategy, goals and training)
 - This is practically a big deal for your timeline as time runs short


Document Type	Details	Responsible	Status	Internal Deadline	Received
Budget (Draft)	Confirm all Key Personnel, Subcontracts, & large line items Composed of stipends, tuition and fees, and institutional allowance	PI		11/3/2023	
Letters of Reference	At least 3, no more than 5. Letters should be from individuals not directly involved in the application (not sponsors/co-sponsors), but who are familiar with the applicant's qualifications, training, and interests. Reference letters are submitted directly through the eRA Commons and do not use Grants.gov. This process requires that the referee be provided information including (a) the PI's (candidate's) eRA Commons user name, (b) the PI's first and last name as they appear on the PI's eRA Commons account, and (c) the number assigned to this Funding Opportunity Announcement. Instructions for Referees are also found at: https://grants.nih.gov/grants/how-to-apply-application-guide/submission-process/reference-letters.htm http://grants.nih.gov/grants/funding/424/Referee_instructions_Mentored_Career_Awards.doc	PI		12/7/2023	Referees Must Submit Thru eRA Commons Directly prior to the deadline.
KEY PERSONNEL & SIGNIFICANT CONTRIBUTORS BIOSKETCHES					
Principal Investigator		PI/RA		11/10/2023	
Sponsor	Anne Stey	PI/RA		11/10/2023	
Co-Sponsor	Allen Heinemann	PI/RA		11/10/2023	
Co-Sponsor	Tara Lagu	PI/RA		11/10/2023	
Co-Sponsor	Alexander Lundberg	PI/RA		5/18/2016	
Co-Sponsor	Adin-Cristian Andrei	PI/RA		5/18/2016	
BUDGET					
Budget (Final)	Include applicable stipend amount requested for the initial period of support and the number of months, actual tuition fees, and the standard institutional allowance	PI/RA		11/13/2023	
Budget Justification	Justify each line item	PI/RA		11/13/2023	
OTHER ADMINISTRATIVE DOCUMENTS					
Cover Letter	Individual fellowship applicants must include a cover letter that contains a list of referees (including name, departmental affiliation, and institution).	PI		11/17/2023	done
Introduction	Required - application must include an introduction addressing issues raised in the previous critique (Summary Statement). Limit: 1 pg	PI		11/17/2023	done
Project Summary/Abstract	In addition to summarizing the research project to be conducted under the fellowship award, describe the fellowship training plan and the environment in which the research training will take place. The Limit: 30 lines of text	PI		11/17/2023	done
Project Narrative	Limit: 2-3 Sentences, lay language	PI		11/17/2023	done
Specific Aims	Limit: 1 pg	PI		11/17/2023	done
Facilities & Other Resources	Description of institutional facilities and resources available to the fellowship applicant	PI/RA		11/17/2023	done
Equipment		PI	Not applicable	11/17/2023	na
Sponsor Co-Sponsor Statements	6 page limit.	PI/RA		11/17/2023	done
Letters of Support	Attachments may be provided (if applicable) by collaborators, consultants, advisors, etc. 6 page limit	PI/RA		11/17/2023	na
HUMAN SUBJECTS:					
Protection of Human Subjects		PI		11/17/2023	done
Data and Safety Monitoring Plan		PI		11/17/2023	done
Inclusion of Individuals across the Lifespan		PI		11/17/2023	done
Inclusion of Women and Minorities		PI		11/17/2023	done
Recruitment and Retention Plan		PI		11/17/2023	done
Study Timeline		PI		11/17/2023	done
Section 4 of HS Form	Only required if clinical trial	PI		11/17/2023	done
Inclusion Enrollment Report		PI		11/17/2023	done
Respective Contributions	1 page	PI		11/20/2023	done
Selection of Sponsor and Institution	1 page	PI		11/20/2023	done
Responsible Conduct of Research	1 page	PI		11/20/2023	done
Alt Phone Number/Degree Sought/Field of Training	Please review 'PHS Fellowship Supplemental Form' for details	PI		11/20/2023	
Background & Goals of Fellowship and Training	6 page limit. Consists of a) Doctoral Dissertation and Research Experience; b) Training Goals and Objectives; and c) Activities Planned Under this Award	PI		11/20/2023	done
Institutional Environment & Commitment to Training	2 pages	PI		11/20/2023	done
APPLICATION (Science portion)					
Research Strategy	Limit: 6 pgs Should address the significance of the proposed studies, including the background leading to the present application; and the approach to provide experimental support of the proposed hypothesis.	PI		11/29/2023	done
References Bibliography	Bibliography of all referenced cited	PI		11/29/2023	done
Appendix - Required Materials	Maximum of 10 PDF attachments allowed. Do not use the Appendix to circumvent page limits	PI		11/29/2023	na
Fellow/Mentor review of proposal PDFs	Fellow and mentor team to review full proposal PDFs to ensure everything looks accurate and as expected prior to SR's submission			11/29/2023-11/30/2023	
Internal review of full application	SR to review and send notes; RA and PI to resolve any errors so NU approval can be obtained. SR will then submit to grants.gov and PI will receive a confirmation with the grant number. PI to check grants.gov after submission to confirm everything looks as expected.			12/1/2023-12/8/2023	

eRA Commons

- Website used by HHS for submission of NIH, AHRQ grants
- Account must be created by grants manager at university (your RA)
- All communication and updates are through eRA Commons portal
- Your RA will be the one to actually submit, the account is mostly for communications from the funding agency


Login with Login.gov ?


Login with eRA Credentials ?
Username:


Password:
 

(For External Users Only)
[Forgot Password/Unlock Account?](#)

Login with Federated Account ?

Login with PIV/CAC
 Login using Smart Card

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on your fede

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the eRA Servi

ALERT: eRA's sch

Note: Improved
additional details

Note: Users with
into a single eRA
account to supp

Note: eRA posts



Status: PI Search

Recent/Pending eSubmissions

- Applications that require action (e.g., to view errors/warnings) prior to submission completion
- Applications that are available to view (during two business day correction window) prior to submission completion
- Applications that have been rejected by Signing Official



List of Applications/Awards

- Funded Awards
- Successfully submitted applications, both paper and electronic
- Review assignment status, review results, summary statements, and Notices of Award
- Other Commons features (e.g., Just In Time, eSNAP, Closeout, Financial Status Report) for previously submitted applications/awards



Search by Grants.gov Tracking Num

Enter the Grants.gov Tracking Number into the following box for easy access to a specific award application

Contacts

Administration:

Scientific Review Officer (SRO)

Name: Kenney, Nicholas J.

Phone: 301-427-1869

Email:
Nicholas.Kenney@ahrq.hhs.gov

Administration:

Grants Management Specialist (GMS)

Name: Caponiti, Anna

Phone: (301) 427-1402

Email:
anna.caponiti@ahrq.hhs.gov

Administration:

Program Official (PO)

Name: Chanlongbutra, Amornrat

Phone: 301-427-1542

Status Information ?

Filter



Expand All

Collapse All

Print

1F32HS029776-01A1

Status Fellowship awarded.	Project Title Long Term Outcomes and Cost-Implications of Inequitable Access to AcuteInpatient Rehabilitation	
PI Name Rao, Saieesh	NIH Appl. ID 10996951	Application ID 1F32HS029776-01A1

▼ Status

Status Fellowship awarded.	Last Status Update Date 06/27/2024	
PI Name Rao, Saieesh	Institution Name NORTHWESTERN UNIVERSITY AT CHICAGO	NIH Appl. ID 10996951
	School Name FEINBERG SCHOOL OF MEDICINE	
	School Category SCHOOLS OF MEDICINE	
	Division Name NONE	
	Department Name	

Priority Score

Found in eRA Commons shortly after the study section meeting date

- The score your application receives is the “priority score” or “impact score”
- Range from 10-90, lower scores are better
- About half of applications are not scored at all after initial review (“ND” – not discussed)
- Initial submission was not funded with a score of 40
- Resubmission was funded with a score of 20
- Percentile based on score was not calculated for my application, but that is an equivalent metric

Review

Application

Award Document Number:
FHS029776A

FSR Accepted Code: N

Snap Indicator Code:

Impact Score:
20

Percentile:

Early Stage Investigator Eligible:

New Investigator Eligible:

Eligible for FFATA Reporting:

Yes

Study Section

Scientific Review Group:
HCRT

Council Meeting Date (YYYY/MM):
2024/05

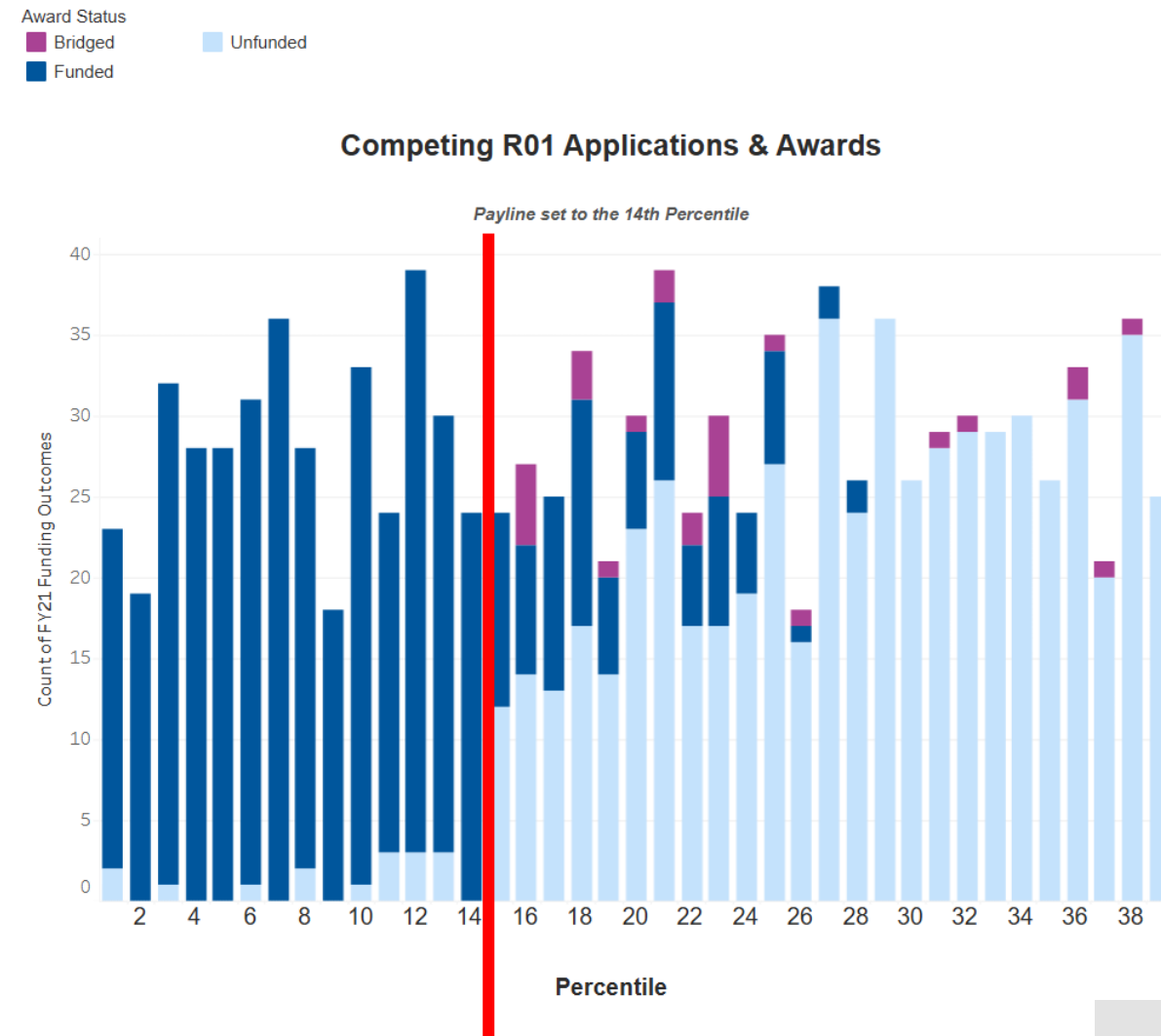
Meeting Date:
02/15/2024

Advisory Council (AC)

Overall Impact or Criterion Strength	Score	Descriptor
High	1	Exceptional
	2	Outstanding
	3	Excellent
Medium	4	Very Good
	5	Good
	6	Satisfactory
Low	7	Fair
	8	Marginal
	9	Poor
Other Designations for Final Outcome		
AB	Abstention	
CF	Conflict of Interest	
DF	Deferred	
ND	Not Discussed	
NP	Not Present	
NR	Not Recommended for Further Consideration	

Paylines

- Your priority score or percentile is used to compare your application to others
- The payline is the **conservative estimate** of the percentile / score below which your grant will be funded
- Varies cycle to cycle
- Not a sure thing (see graph)



Benefits of applying

- Even if you don't secure funding:
 - You will have a clear outline of your research strategy
 - You will have identified pitfalls and know what to troubleshoot (data access, IRB approval, institutional red tape)
 - You will be ready to start research years running
 - You will have better rapport with your mentor
 - You will have firsthand experience writing grants, so your first won't be as an attending
 - You can still report your priority score on your CV even if you miss the payline



Look & Feel of F32 Specific Aims, Research Strategy

Original and Revised Submissions

Specific Aims

Spinal cord injury (SCI) is a widespread and debilitating condition affecting an estimated 294,000 patients in the United States.¹ Sequelae range from impaired sensorimotor function to complete paralysis. SCI also has detrimental effects on employment, social reintegration, and healthcare system utilization.² Importantly, neurologic losses are mitigated by a post-acute care strategy which emphasizes intensive physical therapy, frequent nursing care, and regular physician visits.^{3,4} These activities typically take place at an acute inpatient rehabilitation facility (IRF). IRF care benefits SCI patients who have 0.6 times decreased odds of mortality, 9.4 times increased odds of independent living at one year, and improved neurologic outcomes compared to those not admitted to an IRF.³ It is thereby imperative that SCI patients are triaged to IRFs for post-acute care.

Despite the benefits of IRF care for SCI patients, our group has shown that publicly insured SCI patients are less likely to receive care at IRFs compared to privately insured patients, even after controlling for age, injury severity, and comorbidities.⁵ The reasons for this finding are unclear, as are its long term clinical and fiscal implications. Given that IRF care improves neurologic outcomes, increased access to IRF treatment may improve reduce total long-term healthcare system costs. However, there is a critical literature gap assessing whether such an investment would be cost-effective. Understanding the long-term healthcare costs for patients admitted versus not admitted to an IRF can determine whether upfront investment in IRF care reduces long-term healthcare system costs while improving outcomes and care quality metrics for these patients.

Our overarching goal is to inform healthcare policy at state and national levels by forecasting the clinical and economic implications of expanded care access. Here, our objective is to determine the cost-effectiveness of expanded IRF access in reducing morbidity and healthcare utilization in the SCI patient population. We hypothesize expanded IRF access for publicly-insured SCI patients will reduce long-term morbidity and healthcare resource utilization at readily acceptable cost-effectiveness thresholds. We are prepared to undertake the proposed research because we have extensive experience in clinical care, health economics, economic forecasting, and data analytics using publicly available administrative datasets. We will ensure our findings are relevant and informative to policy makers. We have the following specific aims:

Aim #1: Understand long-term harm associated with SCI patients who received IRF care versus those who did not. We will use the California Department of Health Care Access and Information (HCAI) dataset to identify SCI patients between 2015-2018 and examine the incidence of SCI-associated complications as enumerated in the AHRQ Quality Indicators and obtained via ICD-10 medical coding data.

H1: Rates of SCI-associated complications, such as DVTs, pressure ulcers, UTIs, etc. will be higher among patients who undergo rehabilitation at an IRF than those who do not at one year after their index injury after controlling for clinical risk factors such as age, ~~Elixhauser~~ comorbidity index, and New Injury Severity Score.

Aim #2: Quantify long-term rates of hospital resource use among SCI patients who received IRF care versus those who did not. We will use the California Department of Health Care Access and Information (HCAI) dataset to identify SCI patients between 2015-2018. We will examine individual patient costs over one year following initial injury, as well as individual costs associated with IRF versus non-IRF care.

H2: Total healthcare resource utilization, including ED visits, rehospitalization, etc. will be higher among patients who undergo rehabilitation at an IRF than those who do not at one year after their index injury after controlling for clinical risk factors such as age, ~~Elixhauser~~ comorbidity index, and New Injury Severity Score.

Aim #3: Forecast cost-savings associated with redirecting patients to appropriate post-acute care rehabilitation. This may involve escalation or de-escalation of care. We will apply care costs from patients cared for at IRFs to matched patients who did not receive IRF care based on aforementioned risk factors.

H3: Upfront investment in IRF care for SCI patients would reduce long term global cost to the healthcare system at one year, primarily by improving functional outcomes in the post-acute care period and reducing the incidence of SCI-associated complications.

SPECIFIC AIMS

Traumatic spinal cord injury (SCI) is a widespread and debilitating condition affecting an estimated 294,000 patients in the United States.¹ SCI is sudden, unanticipated, and afflicts young to middle aged people in the prime of their lives. Sequelae range from impaired sensorimotor function to complete paralysis. SCI also has detrimental impact on employment, social reintegration, and healthcare system utilization.² Importantly, neurologic losses are mitigated by a post-acute care strategy which emphasizes intensive physical therapy, frequent nursing care, and regular physician visits.^{3,4} These activities take place at an acute inpatient rehabilitation facility (IRF). SCI patients sent to IRF following initial injury have 0.6 times decreased odds of mortality, and 9.4 times increased odds of independent living at one year versus those not sent to an IRF.³

Despite the benefits of IRF care for SCI patients, our group has shown that publicly insured SCI patients are less likely to receive care at IRFs compared to privately insured patients, even after controlling for age, injury severity, and comorbidities.⁵ The long-term clinical and cost implications are unknown. Given that IRF care improves neurologic outcomes, increased access to IRF treatment may reduce total long-term SCI complications thus reducing healthcare costs. However, there is a critical literature gap assessing whether this investment would be cost-effective long-term. Quantifying the long-term implications for patients admitted versus not admitted to an IRF can determine whether upfront investment in IRF care reduces complications while reducing long-term healthcare costs for injured patients.

Our overarching goal is to inform healthcare policy at state and national levels by forecasting the clinical and economic implications of expanded care access. The objective of this application is to determine if expanded IRF access reduces complications and cost among SCI patients. We hypothesize expanded IRF access for publicly-insured SCI patients will reduce long-term complications at acceptable cost-effectiveness thresholds. We have extensive experience in clinical care, health economics, economic forecasting, and data analytics using all-payer claims datasets. We will ensure our findings are relevant and informative to policy makers. We have the following specific aims:

Aim 1: Understand long-term complications and unplanned healthcare use associated with SCI patients who did not receive IRF post-acute care versus those who did. We will use California Department of Health Care Access and Information (HCAI) data to identify traumatic SCI patients aged 18-64 between 2015-2018. We will examine the incidence of SCI-associated complications as indicated by Agency for Healthcare Research & Quality (AHRQ) Prevention Quality Indicators. We will compare complications and unplanned healthcare use among patients sent to IRFs, skilled nursing facilities (SNF), and home using multivariable mixed-effects logistic regression models.

H1: SCI-associated complication rates (e.g., venous thromboembolism, urinary tract infection) will be lower in patients treated at an IRF than those who were not, one year after index injury controlling for patient variables (e.g., age, ~~Elixhauser~~ Comorbidity Index, injury intent, Spine Abbreviated Injury Score).

Aim 2: Quantify long-term healthcare costs associated with SCI patients who received IRF care versus those who did not. We will use the California Department of Health HCAI data to identify traumatic SCI patients aged 18-65 between 2015-2018. We will examine individual patient emergency department, inpatient, and post-acute care costs over one year after initial injury. We will compare costs associated with IRF versus sub-acute nursing facility care using a hierarchical log-transformed linear regression.

H2: Healthcare cost, including emergency department, inpatient, and post-acute care costs, will be lower in patients treated at an IRF than those who are not at one year after index injury controlling for patient variables.

Aim 3: Forecast cost-savings associated with expanded access for SCI patients to IRF for post-acute care rehabilitation. We will derive and validate a machine learning model for predicting SCI patients' discharge disposition using New York State Emergency Department and Inpatient Discharge Databases from AHRQ's Healthcare Cost and Utilization Project (HCUP) 2015-2018. We will apply this model to California patients, predicting their counterfactual rate of IRF utilization, and combine several methods including propensity matching, proportional hazards, and Monte Carlo simulation to calculate subsequent cost-savings.

H3: Upfront investment in access to IRF care for SCI patients reduces long term healthcare cost at one year.

Research Strategy

1. Significance

This year in the United States, approximately 17,900 patients will suffer from new spinal cord injury (SCIs).¹ These injuries often occur in the setting of trauma and are initially diagnosed and treated in the acute inpatient setting. While the acute care in the initial hospitalization following injury is critical to preservation of neurologic capabilities, ultimate mitigation of neurologic deficits and maximal recovery depends on the quality of rehabilitation provided in the post-acute care setting. In fact, access to quality rehabilitation, under the guidance of experts trained in the rehabilitation of SCI patients, is one of the most important factors in reducing patient mortality and improving subsequent functional status.^{3,4}

Expert rehabilitation tailored to the care of SCI patients is provided at sites known as inpatient rehabilitation facilities (IRFs). Patients are discharged from hospitals to IRFs to complete their course of rehabilitation after their acute hospital needs are met. IRFs provide a space in which patients undergo at least three hours of daily physical therapy and are frequently visited by physicians trained in neurology and rehabilitation.⁶ IRF care is associated with 0.6 decreased odds of 1-year mortality and 9.4 greater odds of returning home to live independently.³ Additionally, patients rehabilitated at IRFs have improved long-term functional outcomes such as increased physical mobility and self-care function compared to those discharged to skilled nursing facilities (SNFs).⁴ Given the well described benefits of IRF care for the near-term benefits and functional recovery among SCI patients, it is no surprise that some jurisdictions enforce the provision of IRF care for SCI patients. Regulations in New York state demand that SCI patients be triaged to facilities which can provide IRF-level care; this results in appropriate transfers of SCI patients to higher levels of care for completion of rehabilitation treatment.⁷

Most other states have no such regulations. This allows for the possibility of SCI patients forgoing necessary rehabilitation care and incurring preventable morbidity such as pressure ulcers, urosepsis, deep venous thrombosis (DVT), and otherwise not returning to their highest potential level of function. Previous work by our group examining patients in California, a state without as stringent regulations regarding SCI care as New York, demonstrates that thousands of patients who experience traumatic neurologic injuries, including SCI but also including traumatic brain injury (TBI), are not triaged to IRF level care in the post-acute period. From the available clinical registry data, it is extremely unclear why these patients are not triaged to IRF care. Despite controlling for clinical factors such as patient age, comorbidities, and severity of injury, the most predictive factor distinguishing patients not admitted to IRF care was public insurance status.⁵ Identification of this factor suggests that a health policy intervention at state or federal levels may mitigate the disparity in IRF access for publicly insured patients.

Such an intervention would be justified by data documenting reduction of long-term SCI-associated morbidity by IRF care at acceptable cost-effectiveness thresholds. However, there is little data documenting reduction of long-term harms and healthcare costs following IRF care. Our group, having already laid the preliminary groundwork by studying short-term costs associated with IRF care, is expertly poised to assess the long-term healthcare system costs and clinical outcomes associated with access to acute inpatient rehabilitation for SCI patients.

2. Innovation

Over the past several years, the team led by Dr. Stey has led research efforts to improve the cost and quality of healthcare in the United States. This work recognizes that drivers of healthcare costs include not only decisions made regarding care in the acute care hospital setting that most physicians are familiar with, but also decisions regarding care in the post-acute care setting as well. There is increasing evidence in the literature that much preventable patient harm and increased healthcare system cost occurs due to inadequate care in the post-acute care setting. For example, understanding that patients undergoing curative cancer surgery remain at increased risk for venous thromboembolism (VTE) after surgery, including post-discharge, informed interventions to reduce this risk.⁹⁻¹⁰ We propose to extend the analogy to rehabilitation of SCI patients, arguing that the quality of post-acute care rehabilitation not only influences clinical metrics such as mortality and functional status, as previously described, but also influences long-term healthcare system utilization and

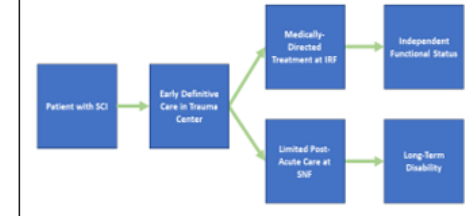
RESEARCH STRATEGY

A. SIGNIFICANCE

A.1. Spinal cord injuries (SCI) are a common and sudden cause of disability in young, working aged people where the degree of disability can be mitigated through access to highly specialized healthcare immediately after injury. This year in the United States, approximately 17,900 patients will suffer a new SCI.¹ These injuries occur from falls, vehicle collisions, sports accidents, and self-inflicted or interpersonal violence.^{1,6} SCI patients are initially treated in specialized hospitals known as trauma centers which provide early definitive care such as neurosurgical decompression^{7,8} and mean arterial pressure elevation following injury to promote neurologic healing.⁸⁻¹¹ However, functional recovery depends on rehabilitation in the first year after injury.^{3,6,11}

A.2. Expert rehabilitation tailored to the care of SCI patients occurs at inpatient rehabilitation facilities (IRFs) which improve functional recovery while reducing long-term complications and mortality. Patients are sent for rehabilitation after initial hospitalization (**Figure 1**). Timely rehabilitation is critical because neurologic adaptation is highest one year after injury.⁶ Patients at IRFs do at least three hours of daily physical

Figure 1: Independence after SCI can be regained with rehabilitation at IRF



therapy and are cared by multidisciplinary teams with physicians trained in neurology and rehabilitation.¹² IRF care is associated with 40% decreased odds of mortality and nine-fold greater odds of independent living at one year compared to skilled nursing facility care (SNFs).³⁴

A.3. Preventable complications occur when SCI patients forgo expert rehabilitation at IRFs. Examples include urinary tract infections from indwelling catheters preventable with bladder training protocols,^{8,13-18} ventilator-associated pneumonias preventable with SCI-specific ventilator weaning protocols,¹⁹⁻²⁷ and pressure ulcers²⁸⁻³¹ and venous thromboembolism³²⁻³⁸ from immobility. These complications particularly occur when patients are discharged to SNF.³ SNFs provide only one hour of rehabilitation daily and lack SCI protocols or multidisciplinary teams. SNFs offer heterogenous care and precipitate functional decline among the disabled.³⁹

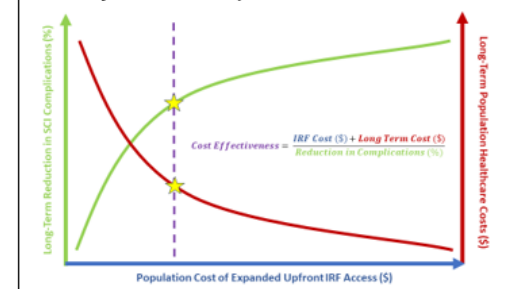
A.4. IRF beds are extremely limited which has led to biased patient selection for IRF. IRF access is limited by insurance coverage,^{40,41} race,⁴²⁻⁴⁶ social support,⁴⁷ and proximity.⁴⁶ Previous work by our group demonstrated that thousands of neurotrauma patients in California are never sent to IRF after hospitalization. That study was motivated by our team's clinical experience working in California that even young, healthy SCI patients were referred to SNF rather than IRF because of anticipated issues with insurance approval. Our study confirmed an inverse association between public insurance and with odds of IRF care despite controlling for age, comorbidities, and injury severity when compared to private insurance.⁵

A.5. SCI is different from other indications for acute IRF because SCI occurs in young, working aged people who decades of life and potential economic productivity ahead of them. This imposes large long-term costs to public payers. Despite being only 14.8% of the population, disabled people account for 72% of Medicaid costs and 54% of Medicare costs, having higher per-capita expenditures than the non-disabled.⁴⁸⁻⁵⁰ Furthermore, SCI patients are often primary earners for families and have potentially decades of economic productivity if neurologic function is regained.^{48,51-53}

A.6. Some jurisdictions have health policies which mandate provision of IRF care for SCI patients. New York state law requires that hospitals discharge SCI patients to IRFs for high quality rehabilitation.³⁹ However, most states lack regulations. Patients not sent to IRFs go to SNFs where intensive rehabilitative care is not possible.^{3,5} Our team quantified that in California it would take \$364 million annually to expand IRF access to all neurotrauma patients.⁵ While large, this cost must be viewed in the context of 1) reducing SCI complication rates, 2) reducing emergency and inpatient encounters, and 3) expected resulting cost savings.

A.7. There is a critical literature gap in whether the upfront cost of IRF care is offset by long-term

Figure 2: Calculate the ideal investment in IRF access to maximally reduce SCI complications for the least total cost



Specific Aim 1: Understand long-term harm associated with SCI patients who received IRF care versus those who did not. We will use the California Department of Health Care Access and Information (HCAI) dataset to

identify SCI patients between 2015-2018 and examine the incidence of SCI-associated complications as enumerated in the AHRQ Quality Indicators and obtained via ICD-10 medical coding data.

Rationale for Aim 1:

Previous literature and our own clinical experience demonstrate that SCI patients experience severe complications requiring hospitalization or emergency treatment as a result of their disability. Incidence of these complications, such as pressure ulcers, urosepsis, and DVT, are critical healthcare quality measures assessed by AHRQ. To establish whether expanded access to IRF care is economically merited for SCI patients, one must establish baseline empirical rates of these complications in both the IRF and non-IRF treated populations.

Experimental Approach

To assess the efficacy of IRF care in promoting health and functioning among SCI patients, we will examine AHRQ Prevention Quality Indicators (PQI) Composite Measures in the SCI population for one-year following their acute injury. PQI Composite Measures include rates of inpatient admission for diabetes complications, hypertension, heart failure, community acquired pneumonia, urinary tract infections, and Asthma/COPD as well as rates of lower extremity amputation.¹² We will also track rates of SCI-associated complications not specifically enumerated in the AHRQ PQI Composite Measures, such as incidence of DVTs and urinary retention. Given that a minority of patients will likely experience recurrent complications whereas others may experience none, rates will be determined by calculating the both the number of unique patients who experience complications (so that each patient is only counted once) as well as the total incidence of complications (in which each patient may count more than once).

The HCAI dataset documents every inpatient admission and emergency department encounter in the state of California. To limit our analysis to SCI patients, we will filter for those patients whose records contain an International Statistical Classification of Diseases and Related Health Problems, version 10 (ICD-10) code corresponding to SCI. These codes are S14, S24, and S34, corresponding to injuries of the cervical, thoracic, and lumbosacral spina cord, respectively. Given that ICD-10 codes may be persistent well after the period of a patient's initial injury (to document a history of SCI), index presentations of SCI will be identified by the presence of a corresponding E-code, and ICD-10 code documenting external traumatic circumstances responsible for the patient's presentation. Patient records will be linked across inpatient and ED encounters to identify the index presentation of a patient's spinal cord injury. The time period under examination will be between 2015 and 2018; this period was chosen for ease of analysis given a full conversion from ICD-9 to ICD-10 coding conventions by this time. During record linkage, each patient with an identified index admission associated with traumatic SCI (as indicated by an E-code) will have their initial post-acute care disposition recorded (IRF versus SNF, for example) for follow-up analysis.

Preliminary analysis demonstrates that 22,946 emergency room records in the 2015-2018 study period document a patient encounter with an active or historical diagnosis of SCI. Similarly, 22,328 inpatient encounters involving an active or historical diagnosis are found during the same time period. Once records are linked for individual SCI patients across inpatient and emergency room encounters, we will use several multinomial mixed-effects logistic regression models to evaluate the association between patient demographics (insurance status, race, ethnicity, median household income, and post-acute care disposition after index hospitalization) and morbidity as defined by readmission, emergency room encounters, the incidence of the AHRQ PQI Composite Measures, and incidence of SCI-associated complications as enumerated earlier.

Completion of this aim will result in knowledge of baseline incidences of unexpected healthcare utilization and long-term SCI-associated morbidity. Demographic and clinical factors which indicate increased long-term SCI associated morbidity will be identified as well.

C.2. Aim 1: Understand long-term complications and unplanned healthcare

0.976] 0.951] IRF = Inpatient Rehabilitation Facility, SNF = Skilled Nursing Facility, NISS = New Injury Severity Score

use associated with SCI patients who did not receive IRF post-acute care versus those who did. We will use the California Department of Health Care Access and Information (HCAI) data to identify traumatic SCI patients aged 18-64 between 2015-2018. We will examine the incidence of SCI-associated complications as indicated by Agency for Healthcare Research & Quality (AHRQ) Prevention Quality Indicators. We will compare complications and unplanned healthcare use among patients sent to IRFs vs. sub-acute nursing facilities using multivariable mixed-effects logistic regression models.

C.2.a. Rationale for Aim 1: SCI patients experience frequent unplanned encounters with the healthcare system due to complications of their injury. Incidence of these complications, such as pressure ulcers, urosepsis, and DVTs are healthcare quality metrics. This aim will describe baseline incidence of unexpected healthcare use and complications and help establish whether expanded IRF access may reduce complications.

C.2.b. Preliminary Data: 22,946 emergency room records not resulting in inpatient admission in the 2015-2018 period document a patient encounter with an active or historical diagnosis of SCI. Similarly, 22,328 inpatient encounters involving an active or historical diagnosis are found during the same period. Given an anticipated complications rate of 25%,^{35,61} this gives us 80% power to detect a 5.1% change in complications.⁶²

C.2.c. Data Sources:

C.2.c.1. California Department of Health Care Access and Information (HCAI) Patient Discharge Data and Emergency Department Data for years 2015-2018

C.2.c.2. Center for Medicare & Medicaid Services (CMS) Inpatient Rehabilitation Facility Quality Reporting Program - Provider Data for years 2015-2018

C.2.c.3. Center for Medicare & Medicaid Services (CMS) Skilled Nursing Facility Quality Reporting Program - Provider Data for years 2015-2018

C.2.d. Inclusion/Exclusion Criteria: Patients between the ages of 18 and 64 who present to a California hospital with a new diagnosis of spinal cord injury indicated by International Classification of Disease (ICD)-10 diagnosis codes of S14, S24, or S34 will be included. Restricting patient age to non-senior adults reduces the likelihood of poor pre-injury functional status and re-admissions from non-SCI related causes. Patients discharged from acute care hospitalization to long-term care facility and 'other facilities' will be excluded (<20% of discharges) because limited claims data preclude determination of drivers of discharge to these settings. The study period (2015-2018) was chosen given full conversion from ICD-9 to ICD-10 codes by this time.

C.2.e. Primary Predictor of Interest: Post-acute care facility following hospitalization for SCI using the 'dispo' variable. This will be confirmed with patient record linkage numbers where discharge date of hospitalization is equal to admission date of post-acute care. Additional covariates that will be controlled include age, sex, race, median income of zip code, Elixhauser Comorbidity Score,⁶³ Abbreviated Injury Severity Score for Spinal Injury, New Injury Severity Score,⁶⁴ insurance status and distance from hospital to nearest IRF and SNF.⁶⁵ The "icdpcr" package⁶⁶ will be used to calculate injury severity scores⁶⁷, and the "comorbidity" package⁶⁸ will be used to calculate Elixhauser comorbidity scores.⁶⁹

C.2.f. Outcome Measures:

C.2.f.1. Rates of SCI complications such as pressure ulcers, urosepsis, and deep venous thrombosis at one year following injury as measured by AHRQ Prevention Quality Indicators.⁵⁷

C.2.f.2. Rates of emergency department and inpatient encounters at one year following injury.

C.2.f.3. Rates of emergency department and inpatient encounters with diagnoses enumerated in the AHRQ prevention quality indicators at one year following injury.

C.2.f.4. **Statistical Analysis:** Each patient will have their initial post-acute care facility recorded (IRF, SNF, home). All subsequent encounters for each patient (grouped by patient record linkage number) within one year of initial injury will be tabulated. Complication rates will be determined using both the number of unique patients with complications and the total incidence of complications. Multivariable mixed-effects logistic and Poisson regression models will evaluate the association between patient post-acute care (IRF, SNF, home) and future complications, ED encounters, readmission, and AHRQ Prevention Quality Indicators. Model predictors are in C.2.e. **k-fold cross validation** will ensure model reliability.^{77,78} All analyses will be in R.

C.2.g. Potential Problems and Alternative Approaches: First, despite rigorous exclusion criteria and covariates to control for injury severity and comorbidity, unmeasured variable bias may still be present. Instrumental variable analysis will be used to address this bias;⁷⁰ distance between acute care hospital and nearest IRF will be used as the instrumental variable to draw causal inference between IRF post-acute care and SCI-associated preventable complications rates. Second, selection bias may be present in post-acute care facility selection. Inverse probability treatment weighting (IPTW) will control for selection bias in post-acute care. These weights will create a pseudopopulation with confounders equally distributed and included in a second logistic regression modeling SCI-associated complication rates associated with IRF, SNF, and home.

C.2.h. Timeline: We expect to use four months (by Nov '23) while also data cleaning and in coursework.

Specific Aim 2: Quantify long-term rates of hospital resource use among SCI patients who received IRF care versus those who did not. We will use the California Department of Health Care Access and Information (HCAI) dataset to identify SCI patients between 2015-2018. We will examine individual patient costs over one year following initial injury, as well as individual costs associated with IRF versus non-IRF care.

Rationale for Aim 2:

The second arm to establishing the cost-effectiveness of IRFs, after tabulating incidence of SCI-associated morbidity, is to tabulate health-care system costs incurred by SCI-patients over the long term. Our group has previously demonstrated that short-term costs for SCI patients associated with IRF care substantial and unlikely to be offset by short term reductions in morbidity. However, reduced healthcare utilization over a longer time horizon as a result of IRF care may make IRF care appear more cost-effective. While both Aims 1 and 2 are necessary for a cost-effectiveness calculation, they are functionally independent and may be completed separately from each other.

Experimental Approach

The experimental approach in Aim 2 largely mirrors that of Aim 1 in terms of the data set used, data filtering performed, and linkage of patient records. For our estimation of healthcare system costs, the HCAI dataset contains charges placed during individual emergency department encounters and inpatient admissions. However, charges placed by healthcare entities are rarely equivalent to the healthcare system costs, as healthcare entities routinely overbill for services and are rarely paid the full amount of their charges.

To estimate healthcare system costs for SCI-related healthcare encounters, we will utilize cost-to-charge ratios (CCRs) to convert charge data present in HCAI to healthcare system costs. CCRs are a fundamental tool in econometrics and AHRQ research methodology; charges are converted to costs simply by multiplying the charge by the CCR.¹³ Calculation of CCRs involves examination of individual hospital and facility balance sheets; namely, each hospital's CCR is calculated by subtracting the hospital's total operating revenue from total operating expenses, and dividing that difference by the hospital's gross patient's revenue.¹⁴ Fortunately, this information is made available at a granular level by HCAI. Our preliminary work has calculated a CCR for all hospitals in California, with a median CCR of 0.27. This means that for every dollar charged to the California healthcare system, the median healthcare system cost is \$0.27. Charges for each encounter at each facility will be converted to costs using each hospital's unique CCR.

Given the prospective nature of possible policy interventions stemming from this work, all healthcare costs will be adjusted for inflation to U.S. dollar amounts in the year that the work is to be published. To do this, we will follow standard econometric procedures adjusting for inflation using the Market Basket Wage index as previously done in our prior work.⁵

C.3. Aim 2: Quantifying long-term healthcare costs associated with SCI patients who received IRF care versus those who did not. We will use the California Department of Health HCAI data to identify traumatic SCI patients aged 18-64 between 2015-2018. We will examine patients' emergency department, inpatient, and post-acute care costs up to one year after initial injury. We will compare costs associated with IRF versus sub-acute nursing facility care using a hierarchical log-transformed linear regression.

C.3.a. Rationale for Aim 2: Our group previously showed that short-term costs for SCI patients associated with IRF care are higher than SNF. However, reduced healthcare utilization after IRF may make IRF care more cost-effective (Figure 2). While both Aims 1 and 2 are necessary for a cost-effectiveness calculation, they are functionally independent and may be completed separately from each other.

C.3.b. Preliminary Data: Using HCAI data for years 2015-2018, median total cost of acute and post-acute care was \$129,000 (Q1-Q3, \$72,500-\$217,000) for patients sent to IRFs compared to \$53,100 (Q1-Q3, \$27,900-\$154,000) for patients sent to SNFs. Total median adjusted cost difference was \$18,461 (95%CI [\$5,908–\$38,064]) more for patients discharged to IRF versus SNF. Median adjusted cost-per-day at IRF was \$1,045 (95%CI [\$752-\$2,399]) more than for SNF, suggesting higher intensity care. Given the above, we have 80% power to detect a change of \$1458 in total median adjusted cost between SNF and IRF cohorts.⁷¹

C.3.c. Data Source:

C.3.c.1. California Department of Health Care Access and Information (HCAI) Patient Discharge Data and Emergency Department Data for years 2015-2018.

C.3.c.2. California Department of Health Care Access and Information (HCAI) Hospital Annual Financial Disclosure Reports for years 2015-2018.

C.3.c.3. Center for Medicare & Medicaid Services (CMS) Inpatient Rehabilitation Facility Quality Reporting Program – Provider Data for years 2015-2018.

C.3.c.4. Center for Medicare & Medicaid Services (CMS) Skilled Nursing Facility Quality Reporting Program – Provider Data for years 2015-2018.

C.3.c.5. Bureau of Economic Analysis Personal Consumption Expenditures (PCE) Index for 2015-2018.⁷²

C.3.d. Inclusion/Exclusion Criteria: See C.2.d.

C.3.e. Primary Predictors: See C.2.e.

C.3.f. Outcome Measures:

C.3.f.1. Log-cost of initial hospitalization, post-acute care (IRF, SNF, home), and all emergency and inpatient encounters within one year, with subgroup analysis for encounters due to SCI complications.

C.3.g. Statistical Analysis: Encounters and charges will be grouped by record linkage number (C.2.g).

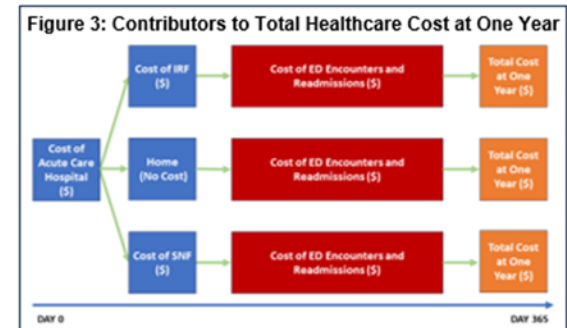
C.3.g.1. Cost to charge conversion: Charges will be derived from emergency department, inpatient, and post-acute care (IRF, SNF, home) encounters (Figure 3). To estimate costs from hospital charges, we utilize cost-to-charge ratios (CCRs). CCRs are a fundamental tool in econometrics and AHRQ methodology. Charges are multiplied by the CCR to yield costs.⁷³ Calculation of CCRs uses individual hospital and facility balance sheets; namely, each facility's CCR is calculated by subtracting total operating revenue from total operating expenses, and dividing that difference by gross patient's revenue.⁷⁴ This information available at a granular through HCAI. Preliminary data shows a median CCR of 0.27 for all hospitals in California; for every dollar charged in California, the median healthcare system cost is \$0.27. Charges for each encounter at each facility for each year will be converted to costs using each facility's unique CCR.

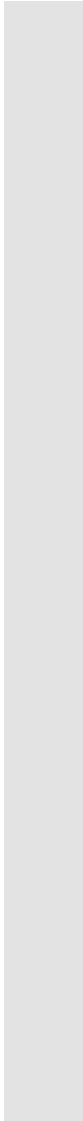

C.3.g.2. Adjust for inflation: costs will be adjusted for inflation to the year of publication using the PCE Price Index⁷² as suggested by AHRQ's Medical Expenditure Panel Survey (MEPS).^{75,76}

C.3.g.3. Cost modeling: The association of predictor variables with long term healthcare costs will be analyzed using hierarchical multivariable log-transformed linear regression. **k-fold cross validation** with starting k=10 as suggested by the literature will be used to ensure performance and reliability of the model.^{77,78}

C.3.h. Potential Problems and Alternative Approaches: Cost data are highly skewed which may bias cost predictions. We will winsorize extreme outliers of CCR and costs to the 1% and 99% percentile and repeat analyses with and without winsorization. CCR are missing in up to 5% of facilities per year. We will use multiple imputations with the 'mice' package in R⁷⁹ to predict missing CCRs based on facility characteristics.

C.3.i. Timeline: We expect four months to complete this aim (Nov '23 – Feb '24) while also in coursework.





Look & Feel of F32 Background & Goals of Fellowship Training

Revised Submission

CANDIDATE'S BACKGROUND

A. Doctoral Dissertation & Research Experience

2011-2012 **Title:** Association of Mast Cells with Colon Cancer Progress
Role: Student Researcher, Northwestern University
PI: Khashavarsha Khazaie, PhD DSc, Department of Immunology Chicago, IL (now at Mayo Clinic, AZ)

Project Summary: My first exposure to research was in the lab a examined mast cell density in human colonic tissue and its corre from normal tissue (euplasia) to dysplastic and neoplastic tissue well as immunohistochemistry to identify mast cells in tissue. Ma using ImageJ. I found that increased mast cell density was signi histology, suggesting mast cell-driven inflammation as a possibl

Skills Developed: I was introduced to laboratory research and pr skills in data collection, wet lab protocols, and statistics. I preser form at the Illinois Math & Science Academy's IMSAloquium Res

2014-2017 **Title:** Investigating Structure-Function Relationships in Cata
Role: Student Researcher, University of Chicago
PI: Joseph Piccirilli, PhD

Project Summary: Ribonucleic acids (RNA) are responsible for c gene transfer. Capable of both catalysis and heredity, RNA is int expanded known RNA biochemistry by analyzing the catalytic ac Using both experiment and computation, our group discovered a cation in the RNA's active site and published in the prestigious N

Skills Developed: Building on wet lab methods from the Khazaie synthesis, HPLC, gel electrophoresis, and kinetic studies with ra my intellectual independence, owning my experiments and pres

2015 **Title:** Investigating the Oncogenic Potential of Mutations in)
Role: Student Researcher, Memorial Sloan Kettering Cancer Ce
PI: Omar Abdel-Wahab, MD, Human Oncology and Pathogenes Kettering Cancer Center, New York City, NY

Project Summary: The Abdel-Wahab lab at MSKCC uses funcn epigenetic drivers of leukemias and lymphomas. Having studied Lab, I was intrigued by Dr. Abdel-Wahab's lab investigation of sy translation of the XPO1 gene. These errors altered XPO1's activ p53 from the nucleus. By engineering leukemic cell lines with mi XPO1 knockdown with shRNAs inhibits cell proliferation, sugges

Skills Developed: My time in the Abdel-Wahab lab was the caps drove tumorigenesis. In addition to gaining familiarity with viral tr microscopy, I also assisted with mouse models and scaling proje

2017-2019 **Title:** Development of a Machine Learning Model for Rapid D
Role: Medical Student Researcher, University of Chicago Pritzke
PI: Matthew Churnek, MD MPH PhD, University of Chicago Pritz IL (now at University of Wisconsin – Madison)

Project Summary: Delays in care timeliness lead to morbidity, m costs for acutely ill patients. While several early warning scores patients, these scores do not point to the reversible causes of cl deficiency, we developed a machine learning model to accuratel among high-risk inpatients. Models were trained on over one the

clinical deterioration. Results demonstrated 91% sensitivity in de detecting sepsis, and over 70% in detecting volume overload, hy arrhythmia based on vital sign trends alone. I presented at the 2 (ATS) International Conference, and it was the basis for Dr. Chu

Skills Developed: This was a departure from my prior wet-lab re Stata, and Python, and gained familiarity with machine learning forest models. These skills have been continuously useful in my

CAREER GOALS AND OBJECTIVES

B. Goals for Fellowship and Training

My **long-term** goal is to be an independently funded investigator focused quality of surgical care by *planning* and *predicting* the long-term effects of policy state, and national levels. To this end, I hope to exploit the large-scale increase i to predict these outcomes with big data and simulation methodologies. My prima rigorous simulation of *counterfactual data* (that is, data which does not yet exist i interventions) from existing datasets using state-of-the-art machine learning and career development award will fill gaps in my knowledge of health policy and ski machine learning that limit my ability to effectively answer outstanding questions

The **short-term** objectives of this F32 proposal are to address training g: an early career independent investigator. Since policy interventions typically can implementation using conventional randomized controlled trial approaches, meti experiments are used to qualitatively predict the effects of interventions in an un propose to extend this methodology by using natural experiments in one populat treatment effects in a second population of interest. Doing this requires me to ac **aims:** quantitative methods in (1) **Healthcare Quality** to understand the current d Healthcare System and quality measurement structures; (2) **Health Economics** t inferences (3) **Machine Learning** to use non-parametric approaches to answer h **Scientific Writing and Dissemination of Research** to ensure sustained research c This proposal is an experiential application of training in the above four learning aims map one-to-one with my specific aims. I have selected my sponsors and co program, and research environment with attention to their ability to help me achi

CAREER DEVELOPMENT PLAN AND TRAINING ACT

C. Activities Planned under Award

This proposal involves gaining new knowledge and skills to examine the rehabilitation on the long-term clinical and fiscal outcomes of spinal cord injury (S **research aims** are to (1) Determine the effect of access to post-acute care on th complications and healthcare encounters in a large, longitudinal; (2) Determine t disposition on the healthcare costs of SCI patients in that same patient registry; i and econometrics to predict the effect of an intervention increasing access to inp patients' long-term clinical and fiscal outcomes. As a resident in general surgery and regularly sees SCI-associated complications, I have highly relevant clinical e study. I additionally have prior experience with machine learning, supercomputin However, I have no formal training, and I have identified four learning aims to ac improving health systems and policy. I have identified **coursework in the four l** **quality, health economics, machine learning, and scientific writing and dissemin** During the first year of training, I will complete an in-person **Master's in Health Research** (HSOR) at Northwestern University addressing three of the four learn **Machine Learning coursework** over two years, providing formal training in all **three research aims is expected to take 75% FTE, whereas coursework is e** **travel, conferences, and mentorship meetings are expected to take the rem**

TABLE 1: CAREER DEVELOPMENT TIMELINE AND MENTORSHIP

Components of Career Development Plan	
Healthcare Quality	Federal Policy Making and Healthcare Reform (HSR 470) Northwestern University Ethical Issues in Health Services Research (HSR 460) Northwestern University Mentorship with Tara Lagu, MD MPH PharmD and Anne Stey, MD MSc
Health Economics (Learning Aim 2)	Health Economics & Healthcare Financing (HSR 433) Northwestern University Topics in Health Services Research: Methods and Measurement (HSR 433) Northwestern Univer Applied Quantitative Methods & Analysis for Researchers (HSR 456) Northwestern University Main and Advanced Causal Inference Workshop (August 2024) Northwestern University Mentorship with Alexander Lundberg, PhD Mentorship with Adin-Cristian Andrei, PhD
Machine Learning (Learning Aim 3)	Statistical Horizons Machine Learning for Estimating Causal Effects 3-Day Remote Seminar Statistical Horizons Longitudinal Data Analysis Using R 3-Day Remote Seminar

TABLE 2: Primary and Co-Mentors, Scientific Advisors, and Statistics Advisor

Scientist	Role	Mentoring Contribution	Meetings
Anne Stey, MD MSc (Assistant Professor of Surgery – Trauma Surgery)	Primary Sponsor	<ul style="list-style-type: none"> Provide research expertise in design and experimental methods for relating to econometrics and data analysis Mentorship for a career in health policy and economics as a surgeon-scientist General oversight of career development and research deliverables 	<ul style="list-style-type: none"> Weekly virtual or in-person check-in Weekly research group meetings
Allen Heinemann, PhD (Professor of Physical Medicine and Rehabilitation, Emergency Medicine, and Medical Social Sciences)	Co-Sponsor	<ul style="list-style-type: none"> Director of the Center for Rehabilitation Outcomes Research at Shirley Ryan Ability Lab, the #1 rehabilitation hospital in U.S. Provide expertise in the practical care of spinal cord injury patients and metrics for evaluating the success of rehabilitation Advisor for scientific writing regarding rehabilitation outcomes and preparation for conference presentations 	<ul style="list-style-type: none"> Monthly meeting concurrent with weekly research group meeting Ad-hoc meetings
Tara Lagu, MD MPH (Director, Institute for Public Health and Medicine – Center for Health Services & Outcomes Research; Professor of Medicine and Medical Social Sciences)	Co-Sponsor	<ul style="list-style-type: none"> Co-investigator with Dr. Stey and Dr. Heinemann on project Mentorship in Health Services and Public Health research Provide expertise in disability care based on past work on that subject 	<ul style="list-style-type: none"> Monthly meeting concurrent with weekly research group meeting Ad-hoc meetings
Alexander Lundberg, PhD (Assistant Professor of Emergency Medicine)	Co-Sponsor	<ul style="list-style-type: none"> Member of the Buehler Center for Health Policy and Economics Provide additional research expertise in health economics, econometrics, and health law 	<ul style="list-style-type: none"> Weekly research group meetings
Adin-Cristian Andrei, PhD (Professor of Preventative Medicine and Biostatistics)	Co-Sponsor	<ul style="list-style-type: none"> Expertise in machine learning, computationally intensive methods, propensity score methods for causal inference, and survival analysis Mentorship in statistical design and choice of computational methods 	<ul style="list-style-type: none"> Bimonthly meeting concurrent with weekly research group meeting Ad-hoc meetings

LEARNING AIMS

1. Healthcare Quality

1.1. Mentorship (Anne Stey, MD, MSc, Tara Lagu, MD, MPH, PharmD): Dr. Lagu is a Professor of Medicine at Northwestern University and a renowned health policy researcher. She studies health disparities among marginalized groups, particularly those with disabilities, and has practical experience outside the academy, including two years at the Centers for Medicare and Medicaid Services developing hospital quality metrics. She has mentored over 30 trainees and a dozen junior faculty in obtaining career development awards.

1.2. Structured Meetings: I will attend monthly research group meetings with Dr. Lagu as well as ad-hoc remote meetings as necessary. They will support Research Aim 1 and throughout.

1.3. Coursework: I will complete Federal Policy Making and Healthcare Reform (HSR 470) and Ethical Issues in Health Services Research (HSR 460) for the Master's.

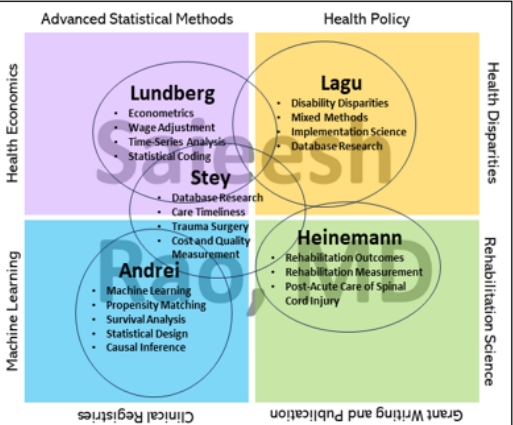


Figure 1. My sponsor and co-sponsors are chosen for their expertise regarding my four learning aims and additional content are expertise.

2. Health Economics

2.1. Mentorship (Alexander Lundberg, PhD): Research Aims 2 and 3 provide the opportunity for formal training in health economics. Dr. Lundberg is an Assistant Professor and applied microeconomist at Northwestern University. He has taught econometrics and computer programming at both the graduate and undergraduate levels, and he received the 2021 Teaching Excellence Award at Northwestern University for his graduate course in biostatistics with Stata. He also has direct experience coding in many of the domains for this project (e.g., cost-to-charge ratios in administrative datasets, comorbidity and injury severity indexes, Monte Carlo simulations, medical diagnosis codes, wage index adjustments).

2.2. Structured Meetings: I will attend weekly research group meetings with Dr. Lundberg as well as ad-hoc remote meetings as necessary. He will support Research Aims 2 and 3 and throughout.



Look & Feel of AAS and ACS Applications



Edge Cases

Dealing with Multiple Funding Sources



Awards Unable to be Combined

Declined AAS Award in favor of ACS

- An application for the ACS Resident Research Scholarship may be submitted even if comparable application to other organizations has been made.
- If the recipient is submitting, submitted, and/or offered a scholarship, fellowship, or research award from another extramural organization, it is the responsibility of the recipient to contact the College's Scholarships Administrator. Those applicants receiving other extramural awards will have to choose between the ACS award and the other awarding body. Intramural awards are allowed (e.g., departmental support, institutional training grant, institutional career development award)
- Applicants who have already earned extramural research funding for their research period, irrespective of funding source or scientific overlap, are not eligible for this scholarship.



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The winner will receive \$30,000 for one year to be used for salary support and/or direct-cost expenditures incurred in the conduct of the proposed research project. The project must be completed within the 1 year funding period. **In the event that investigator is awarded another extramural award, the investigator will be required to accept only one source of funding.** No indirect costs will be covered. The award winners will be acknowledged at the AAS/SUS Awards Ceremony during the 2025 Academic Surgical Congress.

Office of Sponsored Research

Cannot accept awards
funding over 1.0 FTE salary

Had to resubmit budget to
ACS without salary support



Rao, Saieesh

To: Moore, Heather



Wed 3/20/2024 3:17 PM



General



Rao - ACS Budget Template v2.xlsx
17 KB



Hi Heather,

Thanks for responding so promptly to the other thread!

Attached is the revised budget I submitted to ACS. It is not yet confirmed yet by ACS but hopefully can be used for OSR purposes to get things rolling. There is no salary support in the revised budget. I'll keep you posted with any changes or updates!

Best,
Saieesh