## RUSH UNIVERSITY

RUSH MEDICAL COLLEGE • COLLEGE OF NURSING • COLLEGE OF HEALTH SCIENCES • THE GRADUATE COLLEGE

## Rush Research Portal **Grant Walkthrough Sponsored Programs Administration**

**Updated June 2020** 

#### Grant Module of the RRP

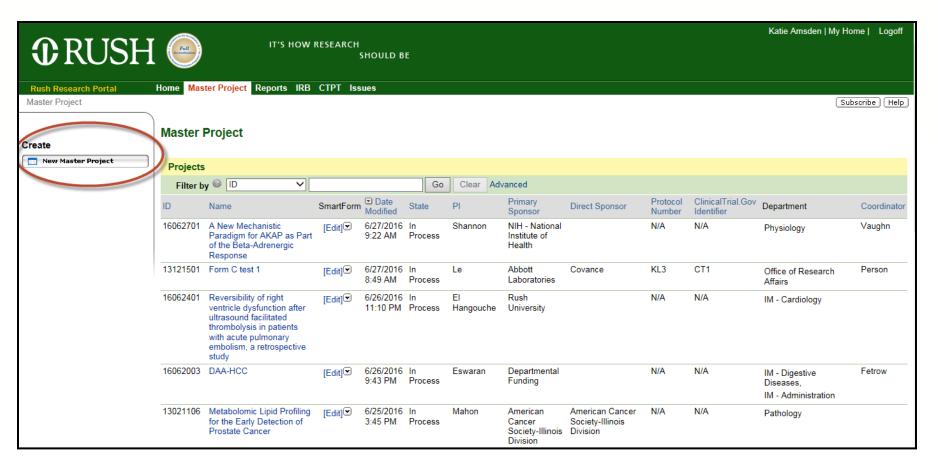


#### What is the purpose of the Grant module in the Rush Research Portal?

- Create, route and submit to Sponsored Programs for review/submission:
  - Federal and non-federal grant applications
  - Grant progress report requests
  - Grant transfer documents
  - Letter of intent requests

#### **Grant Walkthrough**

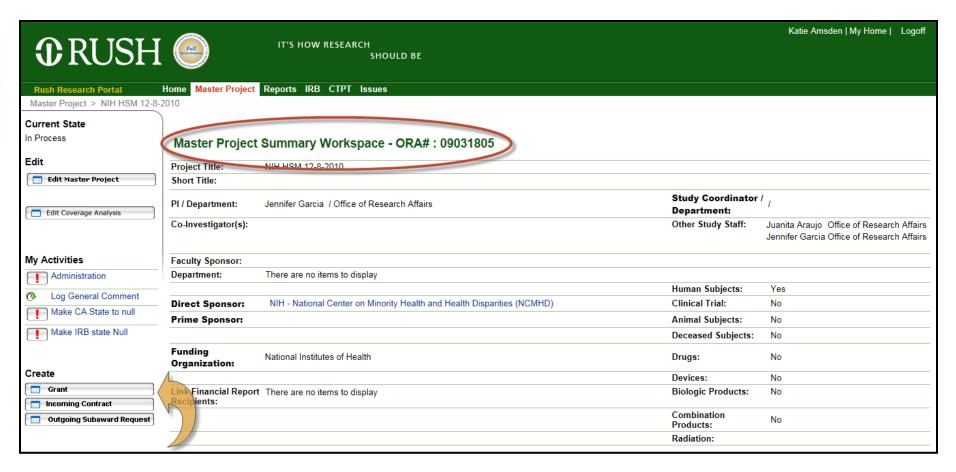
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 Start by searching for an existing master project, or if the project is new, create a new master project by clicking on the "Master Project" tab, then click on "New Master Project" under "Create" on the left side of the screen.

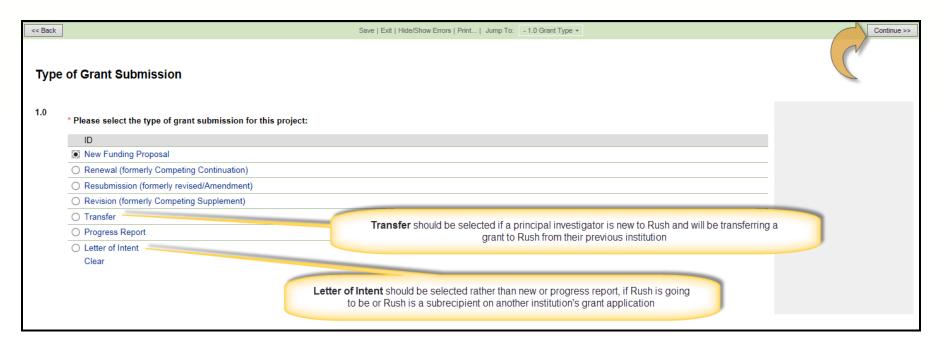
#### **Grant Walkthrough**





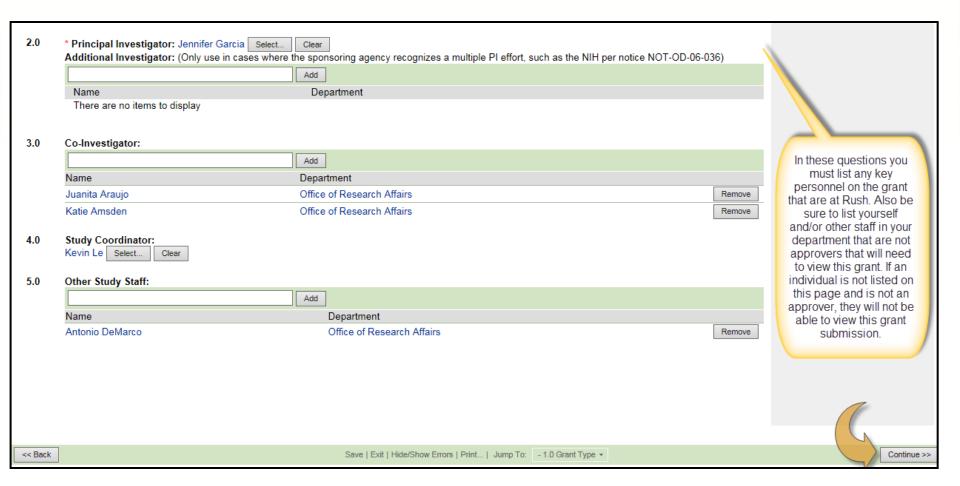
 When you are in the Master Project Summary Workspace, you will have an activity along the left side of the page to create a grant. Click on the "Grant" button.





- This is the question on the first page of the grant smartforms, the selection you make here determines the subsequent smartform pages, thus, it is imperative that you make an accurate selection. See the NIH page for types of grant submissions:: https://grants.nih.gov/grants/how-to-apply-application-guide/prepare-to-apply-and-register/type-of-applications.htm
- Complete each smartform page and choose continue to save and move to the next page.



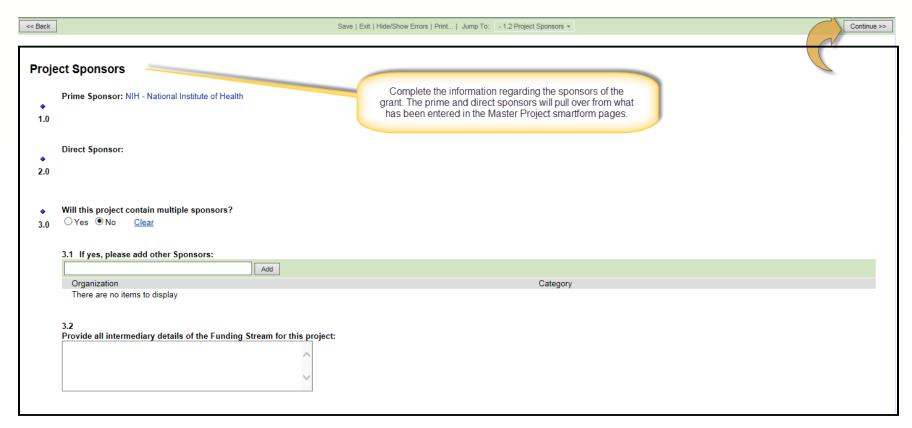


• These are the next questions on the first grant smartform page. Please be sure to add all key personnel and other necessary department staff to this page.

#### **1** RUSH UNIVERSITY

<< Back	Save   Exit   Hide/Show Errors   Print   Jump To: -1.1 General Informa	ation •	Continue >>
Fund	ding Proposal General Information		
1.1	* Proposal Title: (Note 200 Character limit for NIH)    Diffice of Research Affairs Test		
1.2	information will help SPA to	on requested for your grant, this o have the information necessary to ess your grant.	
1.3	* Enter Number Of Budget Period(s): 2		
1.4	* Application Submission Deadline: 7/5/2016 Please pay attention to the due date you progrant being submitted by another institution, the date the prime institution has requested letter of intent by.	please provide	
1.5	PI (listed on the Master Project): Jennifer Garcia  Grant Administrator: Katie Amsden Select Clear	By default, the grains set to the person this grant, unless the grant.	who created
1.6	Please enter members of the Study Staff:  NOTE: This will NOT populate SF424. The users listed on this page, along with the PI, will have READ and EDIT access to this funding land.  Name  Department  There are no items to display	In order to receive regarding this gran identify the people receive these notifi using the activity c.  E-Mail Distribution	t, you must who should ications by alled 'Edit
1.7	* Link Financial Report Recipients:  Individuals in this role will redaily post-award financial refer the grant/fund activity or		nancial reports activity on the
	Name Department	Link dashboard, inc award summary re	
	Juanita Araujo Office of Research Affairs	Remove labor and non-labor transaction and co	r costs
	Katie Amsden Office of Research Affairs	Remove reports, and an ope	
	Jennifer Garcia Office of Research Affairs	Remove report.	

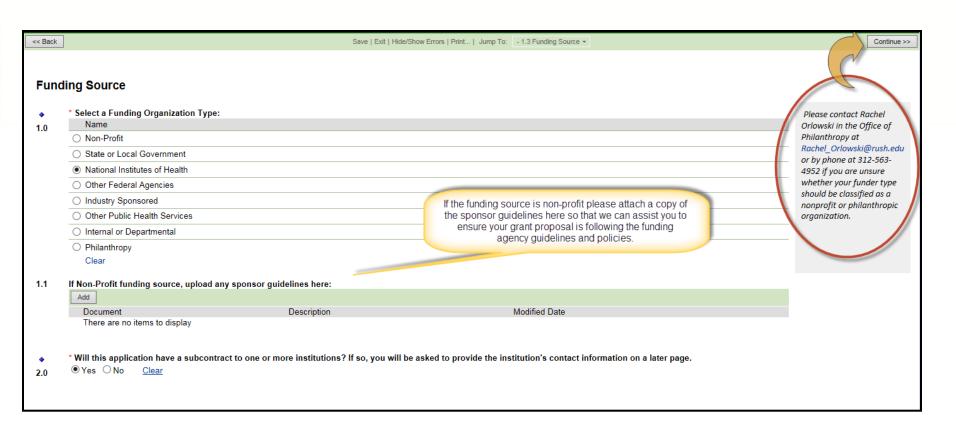




 Prime Sponsor is the original funding source. For example, if Rush University Medical Center applies for a grant to the National Insitutes of Health (NIH) than NIH is the Prime Sponsor because they administer the funds.

**Direct Sponsor** is the organization directly receiving the funds from the prime sponsor and who directly applied for the funding.

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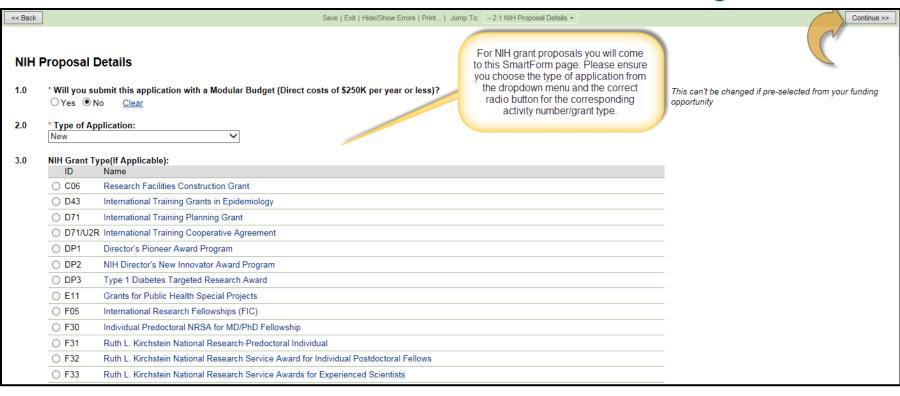


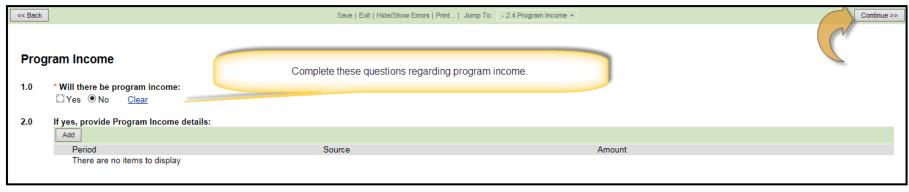


<< Back		Save   Exit   Hide/Show Errors	s   Print   Jump To: - 2.0 Federal Grant Information •	Continue >>
Fede	eral Grant Information	question pertains to li	funded you will come to this SmartForm page. This first mited submission opportunities, this is very important. If s about this, please contact a member of the SPA team.	
1.0	* Are the number of applications from this program announcement/sponsor limited?   Yes ONo Clear			
	If you answered Yes, please reach out to a member of the Sponsored Programs team to discuss.			
2.0	Why is this important? If we sub How do I determine if the progra announcement, specifically the elig	mit more proposals than a prog m announcement/sponsor ha pibility section to determine if a i ber below and click the FIND be	utton. This system will search for your number and information regarding	t review.
	* Opportunity ID (PA or RFA Nu	mber):		For Further
	CFDA Number:		Find	Information: www.grants.gov
	Competition ID:			
			Enter the Program Announcement or Request for Appl	
3.0	If NIH Funding Source, Is the PI	an NIH new investigator?	in the corresponding box, click "Find" and select the rate to the correct opportunity.	idio button next

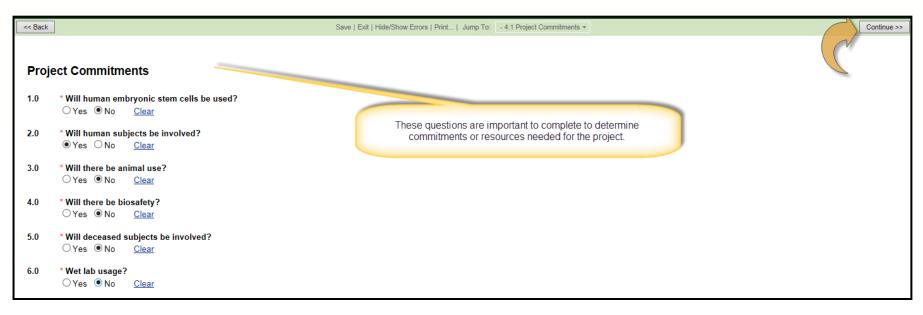
• If the grant application will use the SF424 package, this page will ensure that the correct package of forms is created for you to complete for this particular grant opportunity. Make sure to select the right version of the forms package.

#### TRUSH UNIVERSITY





#### TRUSH UNIVERSITY



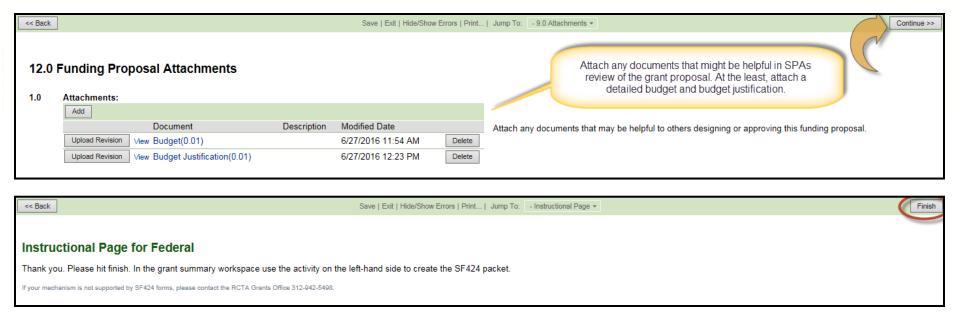






• The information will auto-populate based on what is completed on the "General Information" page of the Grant smartforms.

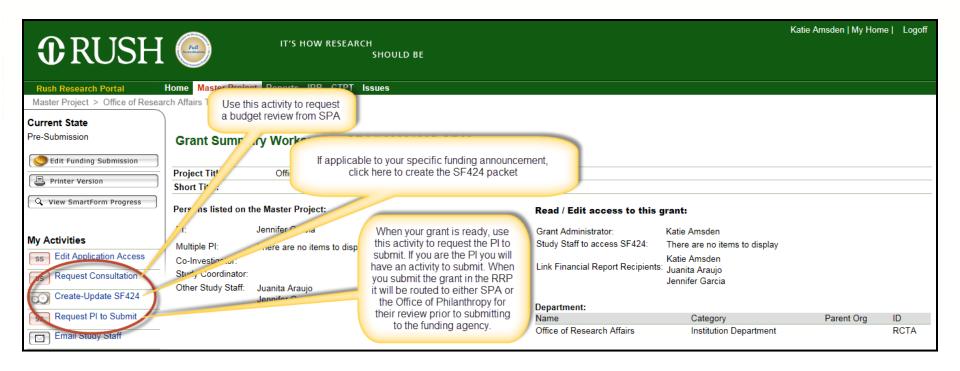




- For Letter of Intent (LOI) requests, attach a budget on the Rush budget template, budget justification and scope of work for SPA review.
- Once you have completed the SmartForm pages and click "Finish", you will be taken back to the Grant Summary Workspace.

#### **Grant Walkthrough**

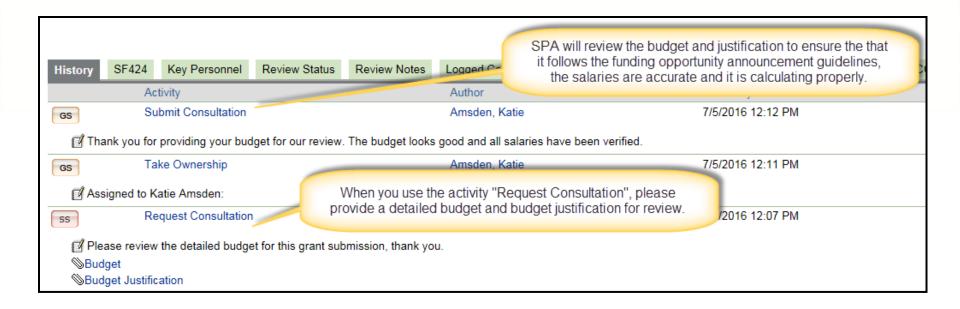




- In the Grant Summary Workspace, you can:
  - Request Consultation
  - Create-Update SF424
  - Submit Application/Request PI to Submit
  - Withdraw Grant (if the state is still in pre-submission)

### Grant Walkthrough – Request Consultation





### Grant Walkthrough – Create SF424

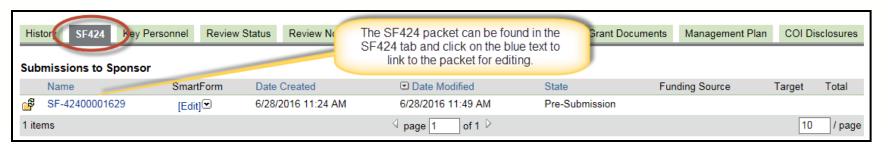


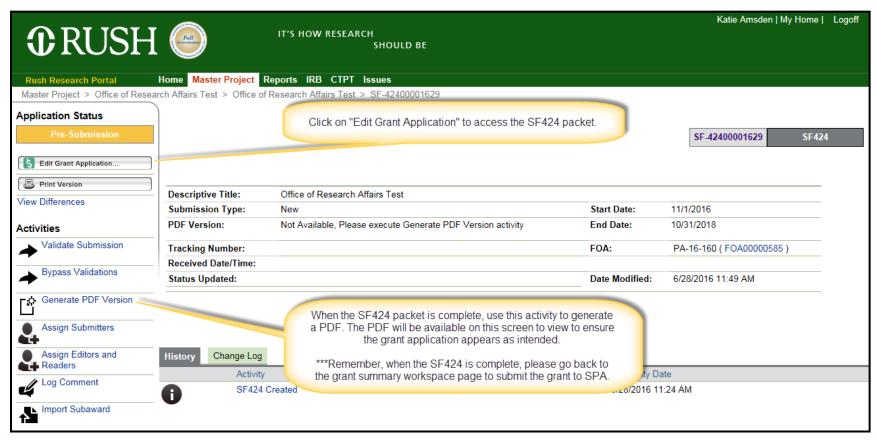
Create SF424 (Al	rtoMap)		
Special note: Th re-generate your	le button will generate your \$F424 for you l \$F424 again, you must contact the researc	ased on the information that you have completed in your forms. Once you generate your 424, to reduce any inforamtion lost office at 312-542-5438 and ask for a grant specialist.	, this button will no longer be available. If for some reason you need to
To continue and	create your \$F424, please select the OK bu	ton at the bottom of this screen. If you want to return to your forms, please click cancel.	
ET .	Action	Status	
₩ Research & R	elated Senior/Key Person Profile (Expanded) \	2.0	
F PHS Human	Subjects and Clinical Trials Information V2.0		
Research & R	elated Project/Performance Site Location(s) V.	.0	
Research And	Related Other Project Information V1.4		
₩ PHS 398 Res	earch Plan V4.0		
₩ SF424 (R&R)	V2.0		
PHS 398 Cov	er Page Supplement V5.0		
Research & R	elated Budget V1.4		
□ PHS 398 Mod	ular Budget V1.2		
E Research & R	elated Subaward Budget Attachment(s) Form	YR 30 ATT V1.4	
PHS Assignm	ent Request Form V3.0		
			OK Cancel

• When the "Create/Update SF424" button is pressed the system will pull up the application forms associated with the program announcement. Click the checkbox next to each page that should be included in the application. Press "OK" once all forms are generated.

### Grant Walkthrough – Create SF424

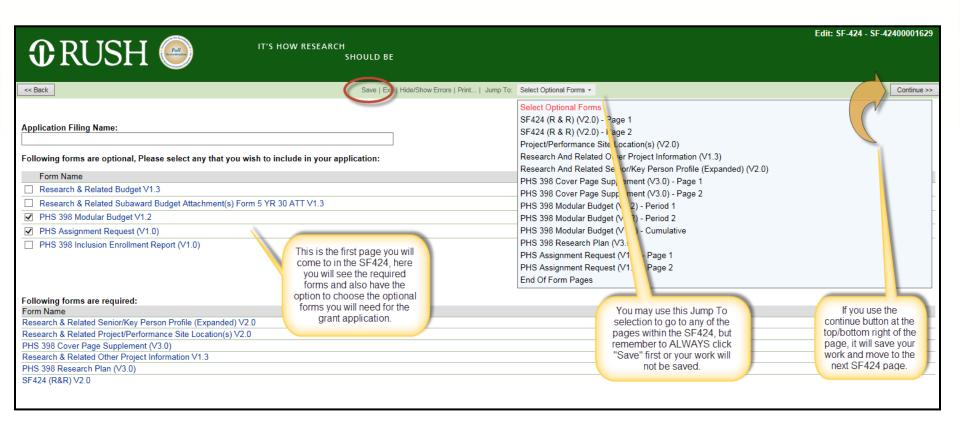
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#### Grant Walkthrough – SF424

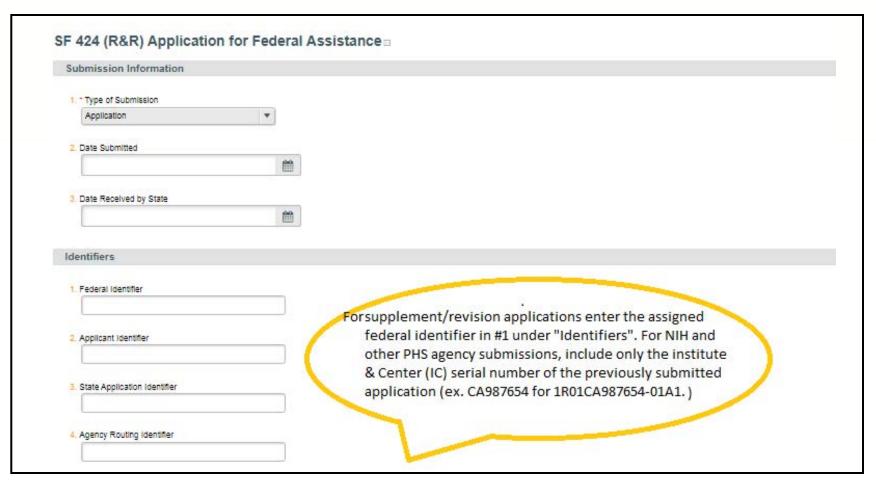
#### TRUSH UNIVERSITY



 This is the first page you will come to in the SF424. For specific instructions on completing sections of the application, please refer to the PA/RFA and the NIH Application guide linked in the PA/RFA.

## Grant Walkthrough – SF424 page 1

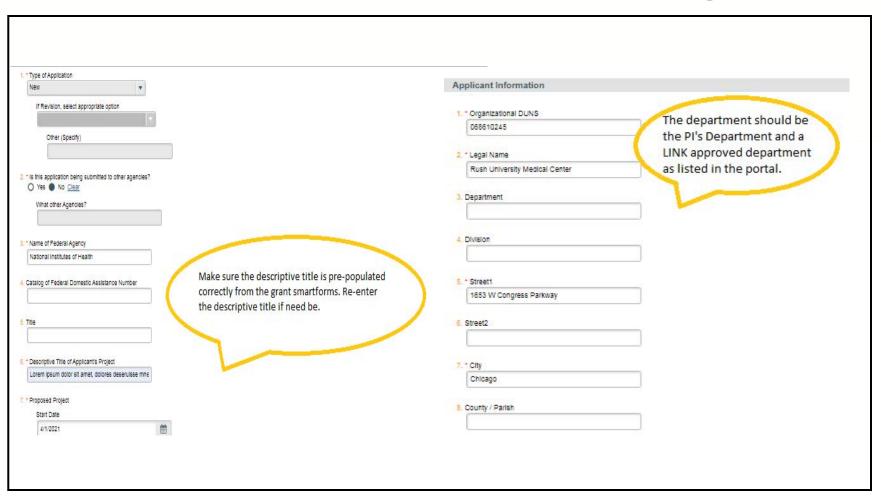




 Most of page 1 of the SF424 will be pre-populated with the information you have filled in from the grant smartforms. Please be sure to review the page for accuracy and enter any missing information.

## Grant Walkthrough – SF424 page 1

#### **1** RUSH UNIVERSITY



## Grant Walkthrough – SF424 page 1

#### **1** RUSH UNIVERSITY

1.	Total Federal Funds Requested		
			Review/Certification/SFLLL
2 .	Total Non-Federal Funds		Is application subject to review by state executive order 12372 process?  Program is not covered by E.O. 12372  Date
3.	Total Federal & Non-Federal Funds		
	\$0.00		<ol><li>By signing this application, I certify (1) to the statements contained in the list of certifications, or fraudulent statements or claims may see</li></ol>
4.	Estimated Program Income	Enter the total project	☐ - Lagree Check this box
	\$0.00	costs for the entire project period including indirects in #1 and #3, found on the last pg of	SFLLL (disclosure of Lobbying Activities) or other Explanatory Documentation [None]
		the budget.	

## Grant Walkthrough – SF424 Project/Performance Site(s)



Zip/Postal Code: 60612 Project/ Performance Site Congressional District: IL-007  Project/Performance Site Location(s) 1-299  Add  Organization Name  There are no items to display	Research & Related Project/Performance Site Location(s)				
Organization Name: Rush University Medical Center  DUNS Number: 068610245  *Street 1: 1653 W Congress Parkway  *City: Chicago  State: IL: Illinois  Zip/Postal Code: 60612  Project/Performance Site Location(s) 1-299  Add  Organization Name  There are no items to display	Project/Performance Site Primary Location				
DUNS Number: 068610245  *Street 1: 1653 W Congress Parkway  *City: Chicago  State:   L:   Illinois					
*Street 1: 1653 W Congress Parkway  *City: Chicago  State:   L:   Illinois	nter				
*City: Chicago  State: IL: Illinois  Province: *Country: USA: UNITED STATES  Zip/Postal Code: 60612  Project/ Performance Site Congressional District: IL-007  Project/Performance Site Location(s) 1-299  Add  Organization Name Street1  City ZIP Code There are no items to display					
State: IL: Illinois	Street 2:				
Project/ Performance Site Congressional District: IL-007  Project/Performance Site Location(s) 1-299  Add  Organization Name There are no items to display	County:				
Project/Performance Site Location(s) 1-299  Add  Organization Name Street1 City ZIP Code There are no items to display	➤ Province: *Country: USA: UNITED STATES	~			
Organization Name Street1 City ZIP Code There are no items to display	Zip/Postal Code: 60612 Project/ Performance Site Congressional District: IL-007				
Organization Name Street1 City ZIP Code There are no items to display	Project/Performance Site Location(s) 1 200				
There are no items to display					
	Street1 City 7IP Code				
	Olicott Olicy Zil Odde				
Indicate the primary site where the work will be performed. If a portion of the project will be performed at any other site(s), identity the site location(s) by clicking add and indicating the other site(s).	Oily Zii ood	. If a			

• Please do not check the box that you are submitting this application as an individual. This application is being submitted by Rush University Medical Center.

## Grant Walkthrough – SF424 Other Project Info



RESEARCH & RELATED Other Project Information	
If no, is the IRB review Pending?  Oyes No Clear IRB Approval Date:  5/20/2014  Description Accuracy Number:  100000483	Project Information used for all grant s. This form includes on the use of human d vertebrate animals,
2. *Are Vertebrate Animals Used? Yes No Clear  2.a. If YES to Vertebrate Animals  Is the IACUC review Pending? Yes No Clear  See Yes No Clear  Yes No Clear  See Yes No Clear	s fields to upload an , project narrative, , equipment lists, and es descriptions.
3. *Is proprietary/privileged information included in ∴ the application?   Yes  No  Clear	
4.a. *Does this Project have an Actual or Potential Impact - positive or negative - on the environment?  4.b. If yes, please explain:  4.c. If this project has an actual or potential impact on the environment, has an exemption been authorized or an environmental assessment (EA) or environmental impact statement (EIS) been performed?  4.d. If yes, please explain:	
5. *Is the research performance site designated, or eligible to be designated, as a historic place?  5.a. If yes, please explain:  Clear	
6. *Does this project involve activities outside the United States or partnerships with international collaborators? Oyes No Clear 6.a. If yes, identify countries:  6.b. Optional Explanation:	
7. Project Summary/Abstract: [None] Add 8. Project Narrative: [None] Add 9. Bibliography & References Cited: [None] Add 10. Facilities & Other Resources: [None] Add 11. Equipment: [None] Add 12. Other Attachments:	IIH application guide rants/how-to-apply-

## Grant Walkthrough – SF424 Key Personnel

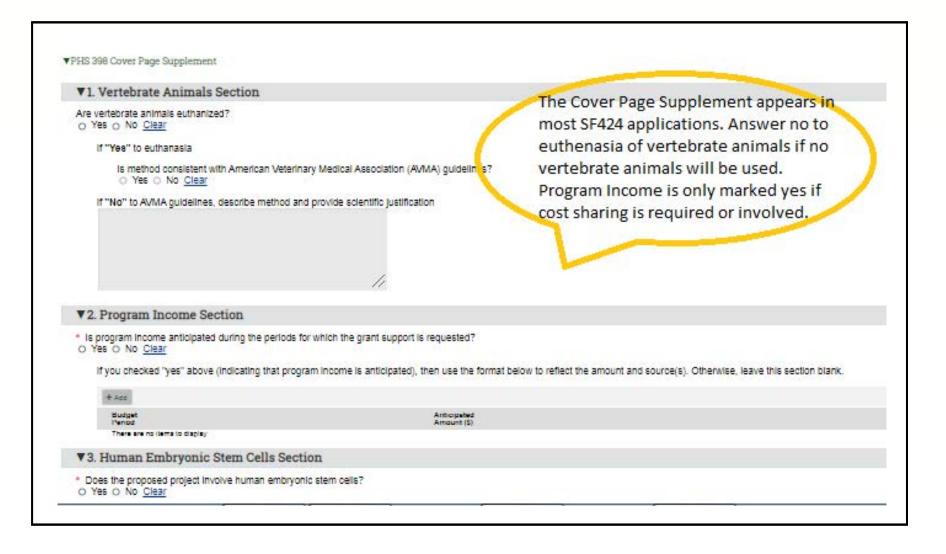


	Research & Related Senior/Key Person (with	Expanded) Profile	The information for	
	the PD/PI needs to			
Prefix: "First Name: Jennifer Position/Title: Director, Sponsored Programs Ad Department:	Middle Name: "Last I Organization: Division:	Name: Garcia Suffix	match exactly to what is listed on page 1 of the SF424.	
*Street 1: County/Parish:	Street 2:	Province:		
*City:	State:			
*Country:	Fach PC	D/PI must be registered in the eRA C	commons and must	
*Phone Number: Fax Nu		ned the PI Role in that system. Each		
Credential, e.g., agency login:	their	respective eRA Commons ID in the	Credential field.	
*Project Role: PD/PI	Other Project Role Category:			
Degree Type: Post Doctoral Post Doctoral Scholar  Post Doctoral Scholar	Choose the	correct project role for each of the ke	ey personnel.	
Other Professional Attach Biogra Attach Currer Support: Consultant Co-Investigator	DDOFILE(S) Senior/Key Decree	1.00		
Co-Investigator PROFILE(S) Senior/Key Person 1-99 Other (Specify)				
		Edit Display	Order	
Add				
Last Name First Name Dep There are no items to display	artment Title Project Role	Project Role Category		
There are no nems to display				
ADDITIONAL PROFILE(S)				
ADDITIONAL SENIOR/KEY PERSON PROFILE [None] Add Additional Biographical Sketch(es) [None] Add				
Additional Current and Pending Support(s) [None] Add				

• Only add a senior/key person profile for consultants or a postdoc if they meet the definition of senior/key.

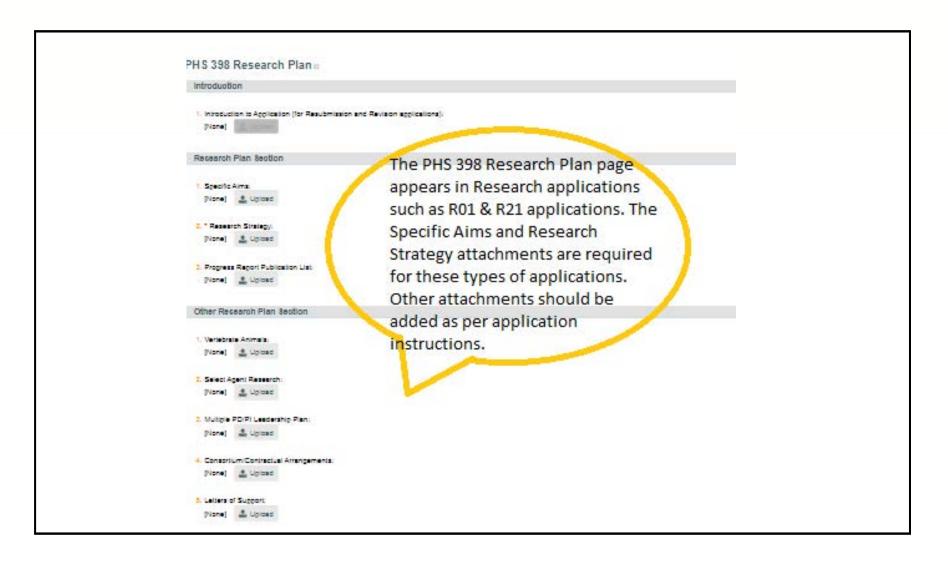
## Grant Walkthrough – SF424 - PHS 398 Cover Page Supplement





#### Grant Walkthrough – SF424 - PHS 398 Research Plan

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# Grant Walkthrough – SF424 - PHS 398 Human Subjects and Clinical Trials Form

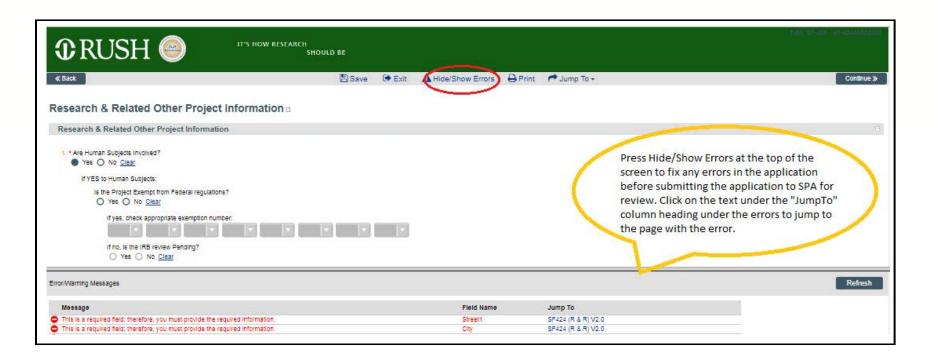


Use of Human Specimens and/or Data  * Does any of the proposed research in the application involve human specimens and/or data?  O Yes O No Clear  Provide an explanation for any use of human specimens and/or data not considered to be human subjects research.  In One Lapses  V Research & Related Other Project Information  Please complete the human subjects section of the Research & Related Other Project Information form prior to completing this form.  The following items are taken from the Research & Related Other Project Information form and displayed here for your reference. Any changes to these fields must be this form.  Are Human Subjects involved?  Yes  Is the Project Exempt from Federal regulations?  Exemption number:  V PHS Human Subjects and Clinical Trials Information  If No to Human Subjects  Add a record for each proposed Human Subject sudy by selecting "Add New Study" or "Add New Delayed Onset Study" as appropriate. Delayed onset studies are: Delayed Onset Studies. For delayed onset studies, you will provide the study name and a justification for omission of human subjects study information.  Add a study record and complete each relevant section  Other Requested Information  Add a study record and complete each relevant section  per the application instructions if the research has been identified as Human Subject Research.  Also human subject to the proposed Human Subject subject indentified as Human Subject Research.	
Use of Human Specimens and/or Data  * Does any of the proposed research in the application involve human specimens and/or data?  O'Yes O'No Clear  Provide an explanation for any use of human specimens and/or data not considered to be human subjects research.  Income  * Does any of the proposed research in the application involve human specimens and/or data.  Provide an explanation for any use of human specimens and/or data not considered to be human subjects research.  * Provide an explanation for any use of human specimens and/or data not considered to be human subjects research.  * Presearch & Related Other Project Information  Please complete the human subjects section of the Research & Related Other Project Information form and displayed here for your reference. Any changes to these fields must be in this form.  Are Human Subjects involved?  Yes Is the Project Exempt from Federal regulations?  Exemption number:  * PHS Human Subjects and Clinical Trials Information  If No to Human Subjects  Add a record for each proposed Human Subjects and Clinical Trials information for an and a pustification for omission of human subjects study information.  Other Requested Information  Cher Requested Information  Add a study record and complete each proposed Human Subject subject in study or 'Add New Delayed Onset Study' as appropriate. Delayed onset studies are:  Delayed Onset Study as appropriate. Delayed onset studies are:  Other Requested Information  Add a study record and complete each relevant section  [None] * Jumes*  Delayed Onset Study Research has been identified as Human Subject Research.  Also human subjects  Study Record(e)	will be conducted that involves human
Numan specime Provide an explanation for any use of human specimens and/or data not considered to be human subjects research.    Note   Logical   Logical   Provided	d/or data that is not classified as human
Provide an explanation for any use of human specimens and/or data not considered to be human subjects research. the subject that   Or, human specimens   Or, human subjects   Or, human subjects   Or, human subjects   Order   Or	ch provide an explanation. For example, lens that only contain information about
Please complete the human subjects section of the Research & Related Other Project Information form prior to completing this form.  The following items are taken from the Research & Related Other Project Information form and displayed here for your reference. Any changes to these fields must be in this form.  Are Human Subjects involved? Yes  Is the Project Exempt from Federal regulations?  Exemption number:  **PHS Human Subjects and Clinical Trials Information**  If No to Human Subjects  Skip the rest of the PHS Human Subjects and Clinical Trials information.  If Yes to Human Subjects  Add a record for each proposed Human Subject acudy by selecting 'Add New Study' or 'Add New Delayed Onset Study' as appropriate. Delayed onset studies are: Delayed Onset Studies. For delayed onset studies, you will provide the study name and a justification for omission of human subjects study information.  Other Requested Information  None]	at has been de-identified to all study staff. crimens taken from deceased individuals.
The following items are taken from the Research & Related Other Project Information form and displayed here for your reference. Any changes to these fields must be a this form.  Are Human Subjects involved? Yes Is the Project Exempt from Federal regulations? Exemption number:  **PHS Human Subjects and Clinical Trials Information  If No to Human Subjects Skip the rest of the PHS Human Subjects and Clinical Trials information.  If Yee to Human Subjects  Add a record for each proposed Human Subject and by selecting 'Add New Study' or 'Add New Delayed Onset Study' as appropriate. Delayed onset studies are: Delayed Onset Studies. For delayed onset studies, you will provide the study name and a justification for omission of human subjects study informs on.  Other Requested Information  Add a study record and complete each relevant section  [None]	
Are Human Subjects involved? Yes Is the Project Exempt from Federal regulations? Exemption number:   PHS Human Subjects and Clinical Trials Information  If No to Human Subjects and Subjects and Clinical Trials Information  If Yes to Human Subjects  Skip the rest of the PHS Human Subjects and Clinical Trials information.  If Yes to Human Subjects  Add a record for each proposed Human Subject acudy by selecting 'Add New Study' or 'Add New Delayed Onset Study' as appropriate. Delayed onset studies are: Delayed Onset Studies. For delayed onset studies, you will provide the study name and a justification for omission of human subjects study information.  Other Requested Information  (None)	
Y PHS Human Subjects and Clinical Trials Information  If No to Human Subjects and Clinical Trials Information  If No to Human Subjects  Skip the rest of the PHS Human Subjects and Clinical Trials Information  If Yes to Human Subjects  Add a record for each proposed Human Subject and y by selecting 'Add New Study' or 'Add New Delayed Onset Study' as appropriate. Delayed onset studies are 'Delayed Onset Studies. For delayed onset studies, you will provide the study name and a justification for omission of human subjects study information.  Other Requested Information  (None)	made on the Research & Related Other Project Information form and may impact the data items you are required to complete on
▼ PHS Human Subjects and Clinical Trials Information  If No to Human Subjects  Skip the rest of the PHS Human Subjects and Clinical Trials information. Trials  If Yee to Human Subjects  Add a record for each proposed Human Subject and by selecting 'Add New Study or 'Add New Delayed Onset Study' as appropriate. Delayed onset studies are:  Delayed Onset Studies. For delayed onset studies, you will provide the study name and a justification for omission of human subjects study informs on.  Other Requested Information  (None)	
▼ PHS Human Subjects and Clinical Trials Information  If No to Human Subjects  Skip the rest of the PHS Human Subjects and Clinical Trials information. Town.  If Yee to Human Subject  Add a record for each proposed Human Subject acudy by selecting "Add New Study" or "Add New Delayed Onset Study" as appropriate. Delayed onset studies are: Delayed Onset Studies. For delayed onset studies, you will provide the study name and a justification for omission of human subjects study informs on.  Other Requested Information  Add a study record and complete each relevant section  [None]	
If No to Human Subjects  Skip the rest of the PHS Human Subjects and Clinical Trials information 7  If Yee to Human Subjects  Add a record for each proposed Human Subject at Judy by selecting 'Add New Study' or 'Add New Delayed Onset Study' as appropriate. Delayed onset studies are:  Delayed Onset Studies. For delayed onset studies, you will provide the study name and a justification for omission of human subjects study information.  Other Requested Information  Add a study record and complete each relevant section  [None]	
Skip the rest of the PHS Human Subjects and Clinical Trials Information Town.  If Yes to Human Subjects  Add a record for each proposed Human Subject and by selecting "Add New Study" or "Add New Delayed Onset Study" as appropriate. Delayed Unset Studies. For delayed onset studies are:  Delayed Onset Studies. For delayed onset studies, you will provide the study name and a justification for omission of human subjects study information.  Other Requested Information  [None]	
If Yee to Human Subjects  Add a record for each proposed Human Subject and y by selecting 'Add New Study' or 'Add New Delayed Onset Study' as appropriate. Delayed onset studies are 'Delayed Onset Studies. For delayed onset studies, you will provide the study name and a justification for omission of human subjects study information.  Other Requested Information  [None]	
Add a record for each proposed Human Subject. Judy by selecting 'Add New Study' or 'Add New Delayed Onset Study' as appropriate. Delayed onset studies. For delayed onset studies. For delayed onset studies, you will provide the study name and a justification for omission of human subjects study information.  Add a study record and complete each relevant section per the application instructions if the research has been identified as Human Subject Research.  Study Record(e)  Study Inter Study Inter These see or Jens to space.	
Other Requested Information  [None]	
Description	those for which there is no well-defined plan for human subject involvement at the time of submission, per agency policies on
Study Record(e) been identified as Human Subject Research.  Alsoh human subject study records using unique flenames.  + Acc  Short Shody little  These are no lettle to dopey	
Study Record(8)  Aliach human subject study records using unique flanames:  + Acc  Short Shudy finite  There are no sems to depay	
+ Acc Shody little Study little Study little Study little	
Short Shudy Ritle There are no lears to copiey	
There are no Jerra to capay	
	Display Order
Delayed Chaet Study(rea)	
+ Acc	
Studylitie Anticipated Clinical Irad? There are no lama to daptay	

For a guide to determine if the research meets the definition of Human Suject Research see the link: https://grants.nih.gov/policy/humansubjects/hs-decision.htm

#### Grant Walkthrough – SF424

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 When finished completing the application and all errors have been fixed, save and close out of the SF424 and press "Request PI Submit" in the Grant Workspace.

Once the PI submits the application it's routed to the Department Chair for approval and then routed to SPA for review.

## Grant Walkthrough – SF424



- This walkthrough is to serve as a guide and is a brief overview of the SF424 packet used for federal grant submissions
- SF424 pages may differ depending on the specific program announcement or funding opportunity
- For specific instructions on SF424 requirements please refer to the specific PA or RFA and the NIH application guide linked in the PA/RFA.
- If you have additional questions regarding grant submissions please reach out to a member of the Sponsored Programs Administration team in the Office of Research Affairs, contact info is on the following page

## **Sponsored Programs Administration Contacts**

TRUSH UNIVERSITY

#### Jennifer Garcia, BS, CRA

Director, Sponsored Programs Administration

Phone: (312) 942-3554

Jennifer\_Garcia@rush.edu

#### Yvonne Harris, MPA, CRA

Senior Grant and Subaward Specialist

Phone: (312) 942-1990

Yvonne\_Harris@rush.edu

#### Lorraine Gibson, BA, CRA

**Grant and Subaward Specialist** 

Phone: (312) 942-2411

Lorraine\_Gibson@rush.edu

#### Jennifer Stadler, BA, CRA

**Grant and Subaward Specialist** 

Phone: (312) 563-1989

Jennifer a Stadler@rush.edu