



**Translational Research Program  
Guidelines & Instructions**

**Effective dates:  
May 1, 2018 – June 30, 2019**

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## Key Points:

- It is highly recommended to access the LLS Research Portal at <https://lls.fluxx.io> to begin the application process well in advance of any deadlines.
- It is recommended that final submissions at each stage (letter of intent/full application) be completed the day prior to the deadline. No aspects of the application, except regulatory approvals, will be accepted past the deadline.
- All components of the application must be present in the order indicated in these guidelines.
- All formatting must adhere to the policy stated in these guidelines.
- Completion of several steps in the process initiates emails sent from the LLS Research Portal. LLS staff may also send emails during the application process. Spam filters should be monitored for these emails.
- Contact [researchprograms@lls.org](mailto:researchprograms@lls.org) if expected emails are not received by the times indicated in these guidelines.
- The deadlines stated in the [TRP Key Dates](#) section are strictly enforced. No exceptions are made to this policy.
- Please do not attach documents to the application that are not specifically called for such as papers in press or published papers. The application could be administratively triaged if this rule is violated.

## General Information

### About The Leukemia & Lymphoma Society

The Leukemia & Lymphoma Society, Inc. (LLS) is a national voluntary health agency dedicated to the conquest of hematologic malignancies and relevant premalignant conditions. LLS supports research, patient aid, community service programs, advocacy, and public and professional education.

### Description of Awards

The formation of the Translational Research Program (TRP) was to foster collaboration between basic and clinical scientists with the intent of enhancing the transfer of basic research findings to clinical usefulness.

Applications are sought proposing novel approaches to the prevention, diagnosis or treatment of hematological malignancies and related pre-malignant conditions. Proposals should be based on molecular, cellular or integrated systems findings and be conceptually innovative. The application should have a clear plan for the eventual clinical translation of the studies proposed and the results expected. This feature will be an important consideration of the review process.

This program is intended to provide support over an initial three-year period. Two additional years may be available through the competitive, peer-reviewed TRP Renewal process to solidify progress made in the initial award and further support a clinical trial. To be considered for a TRP renewal award, a clinical protocol for a Phase I or Phase II clinical trial based on the initial Translational Research Program grant must be submitted to the institution's IRB for approval and the work must be a direct result of the funded TRP award (see Renewal Guidelines and Instructions for detailed information).

### Maximum TRP Award Duration & Value

**\*Please note: The TRP award amount you are given will reflect the amount you request in the budget section of your application. Any requests to increase funding must be in writing to LLS and are subject to the availability of funds.**

<i>Duration</i>	<b>Maximum Annual Direct Costs</b>	<b>Maximum Annual Indirect Costs</b>	<b>Maximum Annual Total Costs</b>	<b>Maximum Total for 3 Years</b>
3 years	\$180,018.00	\$19,982.00	\$200,000.00	\$600,000.00

## Who Can Apply

### Citizenship

LLS welcomes applications from both US citizens and non-citizens, as well as applicants who are performing research outside the US. Applicants must be appointed to a not-for-profit institution at the time the funding commences.

This year, LLS and the Snowdome Foundation (an Australian-based not-for-profit organization supporting blood cancer research) have joined forces to provide funding for up to two TRPs that will support investigators who conduct research in Australia. The awardee may be Australian or an Australian researcher working with others outside Australia, or an Australian and an investigator outside Australia applying jointly (“co-PI”).

### Degree

Applicants must hold a PhD, MD, DVM or equivalent degree.

### Sponsoring Institution’s Acceptance of Contract Terms

Applicants who are offered a TRP Award will be sent a contract. The current contract is found on LLS’s website, ([www.lls.org/research/translational-research-program](http://www.lls.org/research/translational-research-program)). Currently, the NIH does not accept LLS’s contract terms.

### Leadership and Staffing

A Principal Investigator may only submit ONE application per application cycle and cannot serve as a Principal Investigator OR Co-Principal Investigator on more than ONE application per cycle. A Co-Investigator (also known as Collaborator) CAN serve as Co-Investigator on more than one application with no limit. A Principal Investigator or Co-Principal Investigator can serve as a Collaborator on other applications. Members of the peer review committee cannot apply for a TRP award. They may apply for a renewal award if they are not serving on the renewal committee since these are awarded independently of the regular TRP awards.

The Application will require one Principal Investigator who is responsible for the preparation and submission of the proposal including budget, the conduct of the research programs and adherence with all stipulations made by LLS in this document, the Policies & Procedures document, and in the grant contract, if funded. Co-investigators are allowed on multiple applications; however, one individual is to be designated as the Principal Investigator and this individual is limited to one application only.

An Applicant may only submit one new application and/or one Renewal application per award cycle.

### Relevance

The proposed research should be clinically directed or clinically translatable in hematologic malignancies that is intended to develop innovative approaches to treatment, diagnosis, or prevention. Projects currently funded by LLS can be viewed under the Grant Finder section of the LLS website at <http://www.lls.org/grant-finder>.

## Review Process & Applicant Notification

### Review Process of LOI

Letters of Intent for the traditional TRP RFP topics are reviewed and approved by LLS at the time of submission. Once the LOI has been reviewed, the Applicant will be notified via an automated email as to whether or not they have been invited to submit a Full Application. The LOI is non-competitive and is for eligibility purposes only. If invited for Full Application submission, the Applicant will immediately have access to this section in Fluxx. If you have not received an email regarding your LOI approval within five business days, contact [researchprograms@lls.org](mailto:researchprograms@lls.org).

The deadline to submit all Full Applications is October 31, 2018 at 3 PM ET. Full Applications will only be accepted via the LLS Research Portal (<https://lls.fluxx.io>). The submission deadlines will be strictly enforced. Please note that all times are Eastern Time (ET). If any date falls on a weekend or a U.S. holiday, the deadline becomes the following business day.

### TRP Key Dates

	Date	Time
Call For Proposals	May 2018	
Letter of Intent due	August 31, 2018	3:00 PM ET
Full Application due	October 31, 2018	3:00 PM ET
Panel Review Meetings	March 2019	
Award Notification	May 2019	
LLS's receipt of signed contract*	June 1, 2019	
Award Start Date	July 1, 2019	

\*Current contract Terms & Conditions can be found on the [TRP webpage](#).

**It is highly recommended that submissions are done the day prior to the deadline.** Internet traffic may be slow near the deadline, which may result in difficulties in submission. In addition, LLS's response time to questions may be delayed by the high volume received near the deadline. Therefore, it is imperative that any questions be posed to LLS as far ahead as possible. The LLS Research Portal automatically shuts down submissions after the deadline has passed.

## Review Criteria

An application will be judged on these criteria:

- The probability of an advance in prevention, diagnosis or treatment in the near-term
- The conceptual basis upon which the proposal rests
- The novelty of the concept and strategy
- Thoughtful and clear presentation
- The overall plan for bringing the research findings to clinical application
- Experience, background and qualifications of investigators
- Adequacy of resources and environment (facilities, access to patient samples if needed, data management and data analysis, etc.)
- Adequacy of provisions for protection of human subjects

### Review Process of Full Applications

Full Applications will be reviewed after the October 31<sup>st</sup> submission deadline by the TRP Subcommittee of the Medical & Scientific Affairs Committee. If an application does not meet the program goals, scope or guidelines, it will be administratively disqualified. Applications are assigned an initial score by the primary and secondary reviewers. Only applications that fall above a scoring level determined by the committee chair will be discussed in detail for final ranking by the entire committee.

Once ranked, priority scores and funding recommendations of the TRP Subcommittee will be presented to the Medical & Scientific Affairs Committee and Mission Oversight Committee for final determination of awardees. The Mission Oversight Committee will determine the number of awards funded, based on scientific merit and the budget approved by LLS's National Board of Directors.

TRP applications will be rank-ordered based on their Overall Priority Score (10-90; which reflects the average of all the reviewers' priority scores multiplied by ten).

Any Applicant selected for funding will be notified within 45 days of the funding decision. Please do not call or email LLS to determine whether the application has been received, when it will be reviewed or the results of the review. Please check Fluxx for the status of your application. All priority scores are confidential in that they are available to LLS's Medical & Scientific Affairs Committee, LLS's Mission Oversight Committee, its Research Subcommittee and administrative personnel only. Feedback may only be provided for applications discussed by the full review committee.

**The LLS will continue to pursue proposals in several specific research areas that it considers “high unmet need.”**

### **General Instructions for Applying**

All submissions must use the LLS Research Portal powered by Fluxx at <https://lls.fluxx.io>. It is recommended that you familiarize yourself with this portal well in advance of any deadlines.

#### Registration

Both the Applicant and Sponsoring Institution must be registered in Fluxx. If you have applied to LLS in the past, you do not need to create a new registration. Simply log in, or contact [researchprograms@lls.org](mailto:researchprograms@lls.org) to reset your password. Once updated, the Applicant can begin the LOI. Email [researchprograms@lls.org](mailto:researchprograms@lls.org) for assistance creating a new account in Fluxx if you do not already have one. Only LLS staff members have administrative permission to create new accounts.

#### Institutional Designation

Applicants should create their profile from the standpoint of where they will perform their research described in the application. The Applicant must indicate the name of the Sponsoring Institution as well as the name of the signing officials for that institution. To register a new institution, contact [researchprograms@lls.org](mailto:researchprograms@lls.org).

#### Data Entry

Both the LOI and the full application may be accessed and changed multiple times as needed prior to the submission deadline. However, neither the LOI nor the full application can be changed once the deadline has passed or the application has been finally submitted. Moreover, some fields may not be modified in the full application following submission of the LOI.

#### Forms and Format

Applicants will provide information on the Fluxx website at the LOI phase; there is no other template necessary at this phase. For the full application phase, a template will be provided in Fluxx. All information must be typed in English using commonly accepted grammar and punctuation. Some information will be captured when Applicants populate fields on Fluxx. Fields in bold are required. All Applicants must use single-spaced text and Times New Roman, size 12 font. Margins are preset in the template and must remain as set. The Applicant's name should be typed in the upper right corner of each page of the template. Failure to use the provided template or to adhere to font size, spacing, margins and/or page limitations will result in the disqualification of the application. In addition, character limitations must be adhered to.

#### Contacting LLS

Questions that are not clarified in this document, the FAQ on [the TRP webpage](#), or Fluxx should be addressed to [researchprograms@lls.org](mailto:researchprograms@lls.org).



## Beginning an Application

Under “Information” in the left navigation bar, click “Translational Research Program.”

Click “Apply to TRP!” and you will be directed to the Letter of Intent form.

Follow the instructions for each web form field. Bold font indicates required information.

Character limitations include spaces. Character and other length limitations are strictly enforced on the web form and the uploaded project description template. If character limits are not adhered to, the application may be triaged.

You may save your work and return to it at any time by clicking “Save.” Clicking “Submit” will lock your application and prevent further modification at that stage. Contact [researchprograms@lls.org](mailto:researchprograms@lls.org) if you submit in error (must be before the deadline).

Once your letter of intent is approved, you will receive an automated email from Fluxx. Consider that these emails may end up in your spam filter. If selected to submit a Full Application, log back in and click “New or Pending” under “Requests” to continue with your application.

Download and complete the project description template, including all required signatures, and upload to the gray “Supporting Documentation” section of the web form. Margins are preset and must not be changed. Text must be written single spaced in Times New Roman size 12 font.

**Only one PDF file is accepted in this section (Project Description Template combined with biosketch(es)),** so delete any other documents uploaded during the process.

Click “Submit” to formally submit your application to LLS.

## Specific Instructions for Applying

### Letter of Intent

Each Applicant must submit the LOI by **August 31<sup>st</sup> at 3:00pm ET** via the LLS Research Portal (<https://lls.fluxx.io>) or the following business day if this date falls on a weekend or a U.S. holiday. The Applicant should carefully craft the information requested in the LOI as this information is automatically populated into the full application and is subject to the Changes clause listed below.

The LOI for the traditional TRP RFP topics will be evaluated by staff on a rolling basis. If the LOI is approved, the Applicant will be notified by an automated email from Fluxx stating that he/she may proceed to the Full Application phase. Applicants may also check the status of their LOI on Fluxx.

### Completing the LOI

#### **Organization Information**

**Sponsor Institution:** Indicate the name of the institution where the research will be performed. If this institution is not listed, please contact [researchprograms@lls.org](mailto:researchprograms@lls.org).

**Principal Investigator:** The Principal Investigator is the Applicant.

**Institutional Signing Official (ISO):** The ISO is the institutional representative responsible for the signing and agreeing to the accuracy of the application and the terms of the award, should the application be selected for funding.

**Financial Officer:** The Financial Officer is the institutional representative responsible for the financial administration of externally-funded research.

**Additional Access (Admin/Assistant):** Access may be given to personnel to assist in the application process. This is the institutional representative responsible for the day-to-day administration of externally-funded research (or the Research Administrator).

**Technology/Transfer Official (TTO):** The TTO is the institutional representative responsible for overseeing Intellectual Property.

**Zip Code of Sponsor Institution:** Enter the zip code of the Sponsoring Institution if located within the United States. If not located within the U.S., this can be left blank.

#### **Project or Program Information**

If you are applying to the Renewal program, check the corresponding box in the drop-down menu, provide the previous TRP grant number, then refer to the Renewal Guidelines to complete your application.

### **Request Proposal Information**

If your proposed research falls within a topic listed, please choose “Yes” from the selections. Otherwise, choose “No.” Choosing “No” for all topics does not disqualify your application from review. The LLS seeks proposals responsive to the requests for proposals, but will also consider other exceptional proposals with the near-term potential of clinical translation. *Applicants with research proposals that are responsive to the RFP should indicate this on the title page of their Full Application.*

#### General Topics of particular interest:

1. **Personalized medicine approach for cancer treatment.** Advances in cancer care have significantly improved lives of patients with hematologic diseases such as CLL, CML, Hodgkin Lymphomas, and ALL. LLS believes that, with time, complete cures can be achieved for certain diseases or subtypes of diseases. Therefore, LLS will continue to support research that may revolutionize cancer care for any hematologic disease through the use of state of the art technologies for molecular profiling, early disease detection, prognostic/predictive biomarkers, development of liquid biopsy technology and novel target identification, and patient selection.
2. **Development of novel therapies and/or novel therapeutic strategies including those that target mutational and epigenetic events within the microenvironment. LLS is especially interested in novel immunotherapy approaches. Such therapies can be applicable to any hematologic malignancies but emphasis is warranted in the following areas:**
  - a. Aggressive subtypes of Non-Hodgkin Lymphoma including but not limited to DLBCL, tFL, PTCL, and ALCL
  - b. Indolent lymphoma including but not limited to CLL, FL, WM (therapies with the potential to provide significant extension of lives of patients or total disease control in defined subtypes)
  - c. Myeloid disorders including MPN/MDS/AML as well as lymphoid disorders such as ALL
  - d. Multiple Myeloma and pre-emergent conditions
3. **Improvements of safety and efficacy of stem cell transplantation.**
4. **Long-term outcome assessment following the therapies.**
5. **Progress in understanding neoplastic stem cell growth and differentiation as well as cancer cell/microenvironmental interactions especially with translation to novel therapies.**
6. **Prevention of disease by detection and/or treatment of pre-blood cancerous states, smoldering low grade disease, or relapse using state-of-the-art methodologies and new therapies.**

### Special Topics of Interest

- 1. LLS-Snowdome Foundation Translational Research Program.** The Snowdome Foundation is an Australian-based not-for-profit organization whose mission is to accelerate the next-generation treatments for Australian blood cancer patients to help them live longer, better lives. To enhance our common goal, the LLS and Snowdome Foundation will jointly fund up to two meritorious TRP applications focused on blood cancer research from investigators working in Australia, Australian investigators working in other countries, or to Australian and non-Australian researchers jointly applying as co-PIs. While any area of blood cancer research will be considered, the organizations especially seek work focused on multiple myeloma. Applications must be submitted to the LLS TRP program and will be evaluated within the general pool of TRP applications. In addition, applications will be jointly reviewed by both foundations to ensure they meet the funding objectives of both organizations, and scientific progress of each awarded TRP will be evaluated by both organizations on an annual basis. LLS will administer the grant program.
- 2. LLS expansion in pediatric research.** LLS recognizes that new precision medicine and immunotherapies are needed to improve outcomes for pediatric blood cancers. The goal is to develop curative therapies that have reduced long-term complications compared to current cytotoxic therapies. Research may focus on pediatric leukemias, lymphomas, as well as other pediatric blood cancers such as Langerhan's Cell Histiocytosis. We encourage research applications attempting to justify and explore novel therapies for pediatric blood cancers, especially those that uniquely target mutations found in pediatric cancers. The grant applications in pediatrics will be evaluated by a separate section within the TRP program to ensure appropriate evaluation by pediatric cancer experts and attention to programmatic goals. This special call for TRPs is part of a larger effort to build the LLS portfolio of research for pediatrics. Suggested topics for research are provided in a separate description of LLS' expansion in pediatric research, which can be downloaded from [www.lls.org/research/translational-research-program](http://www.lls.org/research/translational-research-program).
- 3. Outcomes research based on digital records.** LLS recognizes the value and feasibility of examining both short- and long-term outcomes for blood cancer patients by using readily available genomic profiles together with electronic health care records. We encourage grant applications using electronic records to examine real-world outcomes, which may include long-term studies (of a retrospective nature) to define better treatments outcomes. Genomic analyses of large data sets to identify patients who may selectively respond to existing or new therapies are encouraged.

## Grant Information

Project Title: Provide a title adhering to the 100 character limitation (which includes spaces).

Project Summary: Provide a summary adhering to the 500 character limitation (which includes spaces). Charts and graphs cannot be included in the project summary section of Fluxx.

Scientific Abstract: Briefly describe the proposed research in 3,000 characters (including spaces) or less using technical language. Once the LOI has been submitted, the scientific abstract may not change. Greek characters or symbols must not be used.

Lay Abstract: Using lay language, clearly state the proposed research in 3,000 characters (including spaces) or less. Once the LOI has been submitted, the lay abstract may not change. Greek characters or symbols must not be used.

Amount Requested: The total amount, including both direct and indirect costs, cannot exceed \$200,000.00/year. Enter the total amount of funding requested over the life of the grant (Maximum \$600,000.00). The amount requested on Fluxx should match the budget section of the full application template.

Proposed Start Date: The start date for all TRP grants is July 1 in the year the award is made (i.e. if an award is made to your application in May 2019, the start date will be July 1, 2019).

Proposed End Date: The end date for all TRP grants is June 30 three years after the year the award is made (i.e. if an award is made to your application in May 2019, the end date will be June 30, 2022).

Previous Submission: Indicate whether you have previously submitted this proposal (or one similar) to LLS and indicate the date of any prior submission.

### Key Personnel or Collaborators Information

New collaborator or key personnel contacts may be added to the collaborator section by typing the name(s) into the boxes. These include Co-Principal Investigators and Co-Investigators. *This section helps LLS identify conflicts with reviewer assignments.*

***If you plan to submit an application or serve as Co-Principal Investigator on an application, you will not be eligible to serve on the program's review panel this cycle.***

### Save and Review

Validation will automatically occur after clicking the "Save" button. Validation is a safety measure for the Applicant to ensure that all required fields are complete. If required fields are empty or incomplete, the system notifies the Applicant of fields that require information.

After clicking "Save" you will be directed to review your LOI. Please ensure all information is accurate, and then click the "Submit" button to submit your LOI to LLS.

### Submission of the LOI

Each Applicant must submit the LOI by **August 31<sup>st</sup> at 3:00 pm Eastern Time** via Fluxx or the following business day if this date falls on a weekend or a U.S. holiday. After clicking the

“Submit” button, the Applicant will receive an email from Fluxx stating that the LOI was successfully submitted. **If you did not receive the confirmatory email from Fluxx within five business days of LOI submission, please e-mail [researchprograms@lls.org](mailto:researchprograms@lls.org).**

Signatures of the Applicant and Sponsoring Institution Officials are not required for submission of the LOI.

### Changes

Information collected in the LOI will automatically populate fields in the full application. Once submitted, changes may only be made after receiving prior approval from LLS. The Applicant must email LLS ([researchprograms@lls.org](mailto:researchprograms@lls.org)) requesting any change and identifying the elements to be changed. Any changes made without the prior approval of LLS may result in the disqualification of the application.

## **Full Application**

Each Applicant must submit a full application by **October 31<sup>st</sup> at 3:00 pm Eastern Time** via Fluxx or the following business day if this date falls on a weekend or a U.S. holiday. Some sections of the full application will be automatically captured electronically on Fluxx from the LOI. Other pieces of information will be captured in the application template that must be downloaded, completed and then uploaded by the Applicant. The Applicant may not modify any information provided in the submitted LOI as this is subject to the Changes clause listed above and may result in disqualification of the application.

## **Completing the Full Application**

### Project or Supporting Documentation

Log onto the LLS Research Portal (<https://lls.fluxx.io>), click “New or Pending” under “Requests” on the left, click on your application, and then click “Edit.”

Download and complete the Project Description/Budget Template. The Project Description is limited to 7 pages.

***The completed Project Description/Budget Template and all appendices, including biosketch(es), must be uploaded as one single PDF file. Failure to submit as a single PDF may result in disqualification of your application.***

### Project Description/Budget Template

The Project Description/Budget template includes 4 fields, as follows: a) Project Description b) Other Research Support c) Budget d) Budget Justification.

Each Project Description is limited to 7 pages and should be presented in the following sequence:

- Title and Specific Aims (approximately 0.25 pages)
- Scientific and Clinical Significance of the Work (approximately 1.0 page)
- Previous Studies/Preliminary Data (approximately 2.5 pages)
- Research Methods (approximately 0.75 pages)
- Interaction with Other Investigators (approximately 0.5 page)
- Resources and Environment (Major lab items or facilities) (approximately 1.0 page)
- References Cited (approximately 1.0 page)

### Budget

The Budget and Budget Justification should provide itemized detail for each major category for all years of the project. The budget can be summarized in Year One and extrapolated for the remaining three years. All totals and subtotals should be completed on the form.

The maximum annual total cost (direct and indirect) cannot exceed \$200,000.00 per year. The aggregate costs over three (3) years cannot exceed \$600,000.00.

Permissible Direct Costs include the following with the specified limitations:

- Personnel Expenses including salary, wage, or stipend with fringe benefits. In total, no more than forty percent (40%) of the direct costs may be requested for the salary and fringe benefit expenses of professional staff with a post-graduate degree (i.e. M.D., Ph.D., D.V.M.) regardless of function or role. This restriction does not apply to technical staff (lab assistants, nurses, etc.).

- Supplies & Materials requests should be itemized by category.
- Equipment Purchase requests must identify each item of equipment with an acquisition cost of more than \$500.
- Travel Expense requests cannot exceed \$1000 per year of the award.
- Other Direct Cost requests can include patient care costs.

Permissible Indirect Costs (often referred to as Institutional Overhead, IDC, M&A, G&A or pooled costs) are those costs incurred for common or joint objectives that cannot be readily identified with a particular project (general maintenance, utilities, library, etc.) as defined in Office of Management and Budget Circular A-21. Indirect costs are limited to (11.1%) of total direct costs. For Sponsoring Institutions that do not choose to use these funds for indirect costs, LLS allows the funds to be applied to the Grantee's/Principal Investigator's stipend or fringe benefits cost.

Impermissible Costs include membership dues, tuition, books, journals and publication costs.

#### Biosketch(es) and Other Support

Use the NIH Biosketch format and include Other Support. ERA Commons user name is not required. The Other Support section must contain all current and pending support from any source. The section of the Biosketch containing the Research Support section may be used, but all current and pending support must be included; completed support should not be included. As per the NIH format, the goals of each grant must be stated. **In addition, specific aims must be listed for current and pending grants that may overlap or appear to overlap with the LLS application.** This includes any grants or portions of grants submitted to any organization, including LLS.

If funding decisions about potentially overlapping, pending grants become available following submission of an LLS application, LLS must be notified within five business days of the applicant's receipt of that information.

LLS recognizes that some investigations, particularly those involving clinical trials, may require multiple funding sources, so overlap of specific aims with another grant may be appropriate and acceptable. The need for and details about such overlap should be clearly explained in the application. However, LLS will consider an applicant's other current grant support in its funding decisions. This may result in LLS funding only part of the grant or none of the grant, depending on the level of overlap. In addition, LLS reserves the right to adjust the level of funding of an awarded grant should another overlapping or potentially overlapping grant that was pending at the time of grant submission be awarded to the applicant.

Failure to abide by LLS's rules on disclosure of current or pending support may jeopardize the funding of the current grant application and may affect future LLS funding decisions.

#### Signature Page

All applications must be signed by the Principal Investigator (and Co-Principal Investigator, if applicable) and Institutional Signing Official.

#### Upload the Project Template to Supporting Documentation

Upload the completed template and biosketch(es) as one single PDF to the gray Supporting



Documentation section by clicking the green plus sign. Choose “Project Description/Application” from the drop down menu before uploading.

Organization Assurances

The Applicant must complete the organization assurances section. The following provides an overview:

Human Subjects: The Applicant must indicate if human materials or subjects will be involved in the proposed research. The status (approved, pending or exempt) of Institutional Review Board (IRB, or equivalent oversight entity) approval must be provided. The Human Subject Assurance Number (OHRP) must be included. If the research project has received IRB approval, the date must be provided and documentation must be uploaded as the Human Investigation Statement. The application may be submitted with IRB approval pending. However, an award will not be made without documented IRB approval if it was pending at the time of application submission. It is recommended that the Applicant notify LLS before the March review date if the IRB status has changed. If a project is exempt from IRB review, the certificate of exemption must be uploaded as the Human Investigation Statement.

Laboratory Animals: The Applicant must indicate if laboratory animals will be involved in the proposed research. The status and date of Institutional Animal Care and Use Committee (IACUC, or equivalent oversight entity) approval must be provided. The Animal Welfare Assurance number must be included. Documentation of Sponsoring Institutional approval must be uploaded as the Laboratory Animal Statement. The application may be submitted with IACUC approval pending. However, an award will not be made without documented IACUC approval if it was pending at the time of application submission. It is recommended that the Applicant notify LLS before the March review if the IACUC status has changed.

Recombinant DNA: The Applicant must indicate if the proposed research involves the use of recombinant DNA. Documentation of Sponsoring Institutional approval must be uploaded with the application.

Biohazard Statement: The Applicant must indicate if the proposed research involves the use of biohazards. If the Applicant indicates affirmatively, then an institutional statement of assurances regarding potential biohazards and safeguards must be uploaded as the Biohazard Statement.

Clinical Protocol Appendix (if applicable)

Provide a one page summary and a link to the [clinicaltrials.gov](http://clinicaltrials.gov) website for any clinical protocol essential to the proposed research. Include IRB approval date, IRB compliance number and effective dates of approval. Projects for which IRB approval is pending must include a statement to that effect. The Applicant should notify LLS of IRB approval prior to the March grant review. The applicant must provide information **if a trial is receiving funding from a sponsor** specifically how much money is to be received and what the funds will be used for.

Budgeting Information

Enter the budgeting information as required on the web form fields.

Applicant Assurance

Check the box to accept the terms as stated on the web form field.

Save and Review

Validation will automatically occur after clicking the “Save” button. Validation is a safety measure for the Applicant to ensure that all required fields are complete. If required fields are empty or incomplete, the system notifies the Applicant of fields that require information.

Submission & Confirmation

After clicking “Save” you will be directed to review your application. Please ensure all information is accurate, and then click the “Submit” button to submit your application to LLS.

**Once an application is submitted, only LLS staff can delete files. If you need a file deleted, contact [researchprograms@lls.org](mailto:researchprograms@lls.org) for assistance.**

**If you plan to withdraw your application at any time during the application cycle, please inform LLS staff of your decision by writing to [researchprograms@lls.org](mailto:researchprograms@lls.org).**