Nonprofit Funder	Terms stated Non-negotiable	Export Controls/Sanctions	Conflict of Interest	Research Integrtity/Misconduct	Indemnification	Human/Animal Compliance	Deliverables/Progress Reports/Milestones	Data Sharing	Public Access
Adelson Medical Research Foundation							Financial reports as requested, including confirmation of compliance with the budget. Research progress reports as requested. Equipment purchased must be made available to other foundation funded researchers (not institution specific)		
ALEX LEMONADE STAND FDTN	Not listed in materials available on website	Not specified	Not specified	Not specified	Please submit a copy of your institution's current proof of liability insurance either a Certificate of Insurance listing Alex's Lemonade Stand Foundation as an additional insured or a Letter of Indemnification if your institution is self-insured.	Grantees must submit IRB or IAUC assurances either at time of application or before funding is released.	Annual research progress report which must be reviews by the Scientific Advisory Board and reccommended for continuation of funding	Not specified	Not specified
ALS Therapy Alliance							Final financial report within 60 days of expiration. Progress reports every 6 months and final report.		
Alzheimer's Association	Not listed in materials available on website	Not specified	Only listed in context of review	Not listed in materials available on website	Not listed in materials available on website	IRB and animal welfare assurances must be submitted within 90 days after receipt of award notification.	Annual progress reports, final report within 90 days of frant's end date. Some grants may have six month milestones/updates.	Not specified	Not specified
American Association of Cancer Research	No, require acceptance of grant agreement terms at time of application	Not specified	For some grants, tobacco industry funding is considered conflict of interest (for fellowships mentors can't be accepted any funding from the tobacco industry).	Not specified aside from inclusion in indemnification	e Institution hereby indemnifies and holds harmless AACR for any and all claims, liabilities, losses, and expenses (including attorneys' fees) to the extent arising from or caused by any of Institution's negligent, reckless, or intentionally wrongful act or omission, including without limitation research misconduct, undisclosed conflict of interest or professional malpractice, or fraud or other misconduct in applying for or expending Grant Funds or in carrying out, or reporting on, the Project.	PI and their institution responsible for obtaining assurances/certifications from all team member institutions (for SU2C grants). Human subjects must have IRB approval and consent form as well as IND/IDE where applicable, and report serious and adverse events to the IRB, FDA, and SU2C. Animal research shall comply with applicable NIH and Public Health Service Animal Welfare Policy, including IACUC documentation.	Milestones and deliverables pre-defined in agreement, progress reports could be up to twice a year, and a monthly clinical trial report (for SU2C grants). Grant funds may be withheld if reports are not filed in a timely manner or if milestones and deliverables are not being satisfactorily pursued and acheived.	Only requiring making data available to AACR (at AACR's expense) or regulatory authorities.	Not specified
American Asthma Foundation		not specified	HHS compliant policy required	HHS compliant policy required	not specified	IRB approval ust be provided (NIH OHRPguidelines). IACUC also as applies	Must attend annual meeting. Annual progress report. Annual expenditure reports		
American Cancer Society	Intellectual property terms non-negotiable. Others not specified in materials online.	Not specified.	Adhere to USDHHS guidelines, report new conflicts to ACS	ACS adheres to federal guidelines for scientific misconduct. Allegations must be brought forth to ACS program director overseeing grant. Through chain of command for applicants, reviewers and grantees.	Not speficied.	Industry standards. IACUC and/or IRB approval before funding.	Final expenditures due <90 days of end, annual progress reports and final progress report <60 days.	Not specified.	Not specified.
American Diabetes Association	Not listed in materials available on website	Grantee must comply with applicable United States economic sanctions, anti-terrorism laws, andanti-money laundering laws, including but not limited to the USA PATRIOT Act, the lawsadministered by the Office of Foreign Assets Control (OFAC) of the U.S. Department of the Treasury ,Executive Order 13224 and any local laws that apply in the jurisdiction in which the Grantee isoperating. Grantee certifies that Grantee takes reasonable steps to ensure that Grant Funds are notultimately distributed to any person or entity designated as a Specially Designated National by OFAC(a list of which is online at treasury.gov/resource-center/sanctions/SDN-List/Pages/default.aspx) andthat Grantee takes reasonable steps to ensure that it does not support terrorist activity and/orterrorism-related training or money laundering. This provision must be included in all relevantsubcontracts when applicable.			To the extent permitted by applicable law, Grantee and Grantee Institution (each an "Indemnifying Party") will defend, indemnify and hold harmless the Association, including its regents, directors, officers, employees and agents (collectively, the "Indemnified Parties"), from and against any and all losses, claims, liabilities, damages and costs of whatever kind and nature, including reasonable attorney fees and legal costs, for death or injury of any person and for loss or damage to any property, occurring or claimed to occur as a result of the negligence or willful misconduct of the Indemnifying Party, including its regents, directors, officers, employees, faculty, students and agents, or the failure of the Indemnifying Party to perform its obligations under the Grant; providing, however, the Indemnifying Party shall not be obligated to defend, indemnify and hold harmless any Indemnified Party to the extent any such losses, claims, liabilities, damages, and costs are the result of the negligence or willful misconduct of an Indemnified Party or the failure of an Indemnified Party to perform any obligation under the Grant. This provision shall survive expiration or termination of the Grant.	Animals: must meet standards of PHS or equivalent, and verify IACUC assurances before final approval of grant funding and annually throughout grant period. Humans: meet HHS and NIH guidelines, IRB certification before final approval of grant funding, must adhere to the International Conference on Harmonization for Good Clinical Practice and all relevant host country standards.	Annual scientific report and financial reconciliation, final report within 60 days after grant end, agree to annual follow-up assessments for five years (in response to Association request)	Not specified	Not specified
American Heart Association	Yes	no	Compliance with PHS regulations.	AHA cn determine if institutional policy is acceptable. If so, will follow norml institutional process. Consult with AHA general counsel.		IRB/IACUC approval. Conform to NIH guidance , includinf IBC review if required	annual scientific report within the 30 days prior to end date. Final report must contain list of published or accepted publications	Data required for independent verification of research results must be deposited in AHA approved repository	Must be deposited in Pubmed within 12 months of publication
American Lung Association	Not specified in materials on website	Not specified.	No funding from tobacco industry, compliance with USDHHS financial conflcits of interest 42 C.F.R. pt. 50, subpt. F to the same extent as each would were the research funded by the National Institutes of Health.	Not specified, aside from indemnification and animal/human subject testing notices	ALA will not assume responsibility for and the grantee institution will indemnify and hold the American Lung Association harmless from any lawsuit, claim, judgment, damages, awards, or malpractice arising from research or investigations related to an award as a condition of accepting this award.	Applicants/ awardees must comply with the PublicHealth Service Policy on Human Care and Use of Laboratory Animals and the National Research Council Guide for the Care and Use of Laboratory Animals to the same extent as each would were the research funded by the National Institutes of Health (must have IACUC on file prior to start). Humab subects and/or human stem cells must comply with the provisions of the UnitedStates Department of Health and Human Services 45 C.F.R. pt. 46 to the same extent as each would were theresearch funded by the National Institutes of Health. IRB must be provided prior to start of award.	Financial disbursement report at conclusion. Information on progress reports unspecified.	Not specified.	Not specified
Bill and Melinda Gates Foundation	Terms are non-negotiable	Anti-Terrorism provision--Grantee will not use funds to support activities prohibited by US laws relating to combating terrorism, with persons on the List of Specially Designated Nationals, or in or with countries against which the US maintains comprehensive sanctions, including paying or reimbursing the expenses of persons from such countries or territories, unless such activities are authorized by the US governemnet and approved by the Foundation	not specified	Not specified	If the Project involves clinical trials, trials involving human subjects, post-approval studies, field trials involving genetically modified organisms, experimental medicine, or the provision of medical/health services ("Indemnified Activities"), grantee institution will indemnify, defend, and hold harmless the Foundation against anything resulting from grantee institution's actions/omissions, etc, relating to any Indemnified Activities. Grantee institution's indemnification obligations are limited to the extent permitted or precluded under applicable federal, state or local laws, including federal or state tort claims acts, the Federal Anti-Deficiency Act, state governmental immunity acts, or state constitutions.	Compliance with all applicable regulatory standards	Specified milestone/targets, and interim and final progress reports	Foundation needs to be able to use data as per its Global Access Commitment.	BMGF Open Access Policy: https://www.gatesfoundation.org/How-We-Work/General-Information/Open-Access-Policy
BRAIN & BEHAVIORAL RES FD		not specified	Not specified	not specified	not specified	Comply with HHS guidelines for animal, human subjects and rDNA	final financial report within 90 days of termination. Interim progress report at 6 months, second one year, final report 90 days after grant end.		

Chan Zuckerberg Initiative							**Reports, Communications, Books and Records** a. Reports. At least ten (10) business days prior to each anniversary described in Sections 1.b.i.b, 1.b.i.c, 1.b.i.d, and 1.b.i.e, the University shall provide to CZF an interim detailed written report (each such report, an "Interim Report") describing the use of any Grant Payments (i) during the period beginning on the Initial Funding Date and ending on the fifteenth (15th) day of the month immediately preceding such anniversary; and (ii) not otherwise described in a prior Interim Report. No later than sixty (60) days after the end of the Term, the University shall provide to CZF a final detailed written report describing the use of the Grant Payments (i) during the period beginning on the Initial Funding Date and ending on the last date of the Term; and (ii) not otherwise described in a prior Interim Report (such report, the "Final Report" and together with the Interim Reports, the		dissemination of the results of the projects it supports and expects that the Project Director and/or grantee Institution will actively seek to disseminate, on a national basis, project results, through conference presentations, publication in professional journals, and through release to the media. If grant results have not been disseminated within one yar of termination of the grant, and no plans are in place for doing so, the Fund shall reserve the right to develop and execute a plan for such dissemination.
Commonwealth Fund	not specified	not specified	not specified	silent	HHS compliant review IRB	according to provided schedule	foundation retains license to use data at no additional cost		
CYSTIC FIBROSIS FDTN	No. CFF negotiates standard terms for all awards to each institution/system.	Institution must warrant it will not violate US export control laws	Institution must warrant it has established COI policies that prevent it and its employees from using their positions for personal gain, and institution will report actual COIs as they related to CFF-funded research to CFF within 10 business days of confirmation of an actual COI.	Institution must have policies and procedures in place for the avoidance and reporting of scientific misconduct. Institution must report alleged or actual scientific misconduct within 10 days of institution becoming aware of the misconduct, or alleged misconduct.	CFF and its affiliates assume no responsibility for activities performed by PI under this award. Subject to applicable state law, institution agrees to indemnify and hold harmless CFF and its trustees officers, employees, agents, and affiliates against all actions, claims, demands, costs, liabilities, and expenses (including reasonable attorney's fees) arising out of the activities associated with the award.	Institution warrants it will obtain all necessary approval for human subjects and animal research. Copies of the approvals will be provided to CFF prior to initiating research, and changes and other documentation will be submitted to CFF.	Quarterly and final financial reports; annual progress reports. Detailed instructions on both are included in the terms and conditions.	Not specified	CFF expects PI to publish in relevant peer-reviewed journals, and when data from the CFF National Patient Registry is used for analysis and interpreted in a manuscript, CFF reserves teh right to request revisions to any such analysis or interpretation.
David and Lucile Packard Foundation	No.	Not specified	Not specified	Not specified	Not specified	Not specified	Reporting, as specified in award	Not specified	Not specified

Doris Duke Charitable Foundation	No. Language indicates that if a subsequent grant is awarded where there are overlapping scientific objectives "they will work towards an appropriate resolutions".	Not specified	not specified in general guidelines and instructions, but has a separate 6 page COI policy document.	not specified	not specified	fund research that uses non-human animals. All research conducted involving human subjects must comply with appropriate federal, state and local regulations pertaining to the use of human subjects in research, including all requirements of the United States Department of Health and Human Services. No part of the grant may be used to support any research involving human subjects that does not have the approval of the appropriate Institutional Review Boards (IRB). Appropriate IRB approvals must be in place before research involving human subjects commences (with copies of IRB approvals submitted to DDCF) and must thereafter be maintained and in effect throughout the research project. All federal, state and local regulations and policies governing the conduct of research must be followed without exception, including, but not limited to those relating to human subject protection, radiation and environmental health and safety. The grantee institution is responsible for ensuring that the research conducted as part of this grant is in compliance with all relevant federal, state and local policies and regulations. DDCF must be notified immediately if there are any regulatory issues, protocol violations or policy changes that impact the ability of the grantee to conduct the research as part of this grant.	Progress and financial reports must be completed using the foundation's web-based application and reporting system. A link to the online report form that includes a template for the financial report and additional instructions will be emailed to the grantee approximately two months in advance of the due date. Please refer to your grant agreement for the specific date requirements of your progress/financial reports. Generally, progress and financial reports are due on an annual basis. Progress reports should detail the scientific progress made, as well as provide updated IRB approval status, budgetary, expense and other support information. A separate financial report gives a detailed comparison of actual expenses to the approved line-item budget, as well as an explanation for any overages and/or unspent funds and a budget for the following year, when appropriate. The financial report should be signed by the appropriate financial officer of the grantee institution.	DDCF reserves the right to include information relating to the grants that it funds in DDCF's periodic reports, newsletters or news releases or in any other materials issued by or on behalf of DDCF. DDCF is a member of the Health Research Alliance (HRA) and has agreed to deposit grant information in a database of privately funded grants. The following grant information will be uploaded into this database: investigator name, degree(s), clinical specialty, Open Researcher and Contributer ID (ORCID), institution and institutional information, project title, abstract, grant start date and duration, and grant amount. The Health Research Alliance (HRA) aggregates these data for its member organizations and for periodic publication of findings. View more information on HRA reporter and use of these data by HRA members here .	The foundation requires that any publication based on a DDCF-funded research project must be made freely available and downloadable online in a timely manner and with as few restrictions as possible, in order to ensure that DDCF-funded research can be accessed, read and built upon. Access to PMC is made available to the grantee through DDCF's membership in the Health Research Alliance (HRA) (a national consortium of non-governmental, nonprofit funders of biomedical research and training) and DDCF's registration of data about its clinical research grants in the HRA reporter database. DDCF will provide detailed instructions for depositing documents in PMC to grantees upon DDCF's receipt of a fully executed grant agreement.
Ellison Medical Foundation	The awardee institution may not independently make changes to the agreement received from the foundation.	Not specified	Some conflicts listed for specific programs (senior scholars can't apply twice, new scholars can't receive a "conflicting award")	The Association expects investigators and institutions to follow the National Institutes of Health (NIH) guidelines on research conduct in "proposing, performing, or reviewing research, or in reporting research results" (U.S. Code of Federal Regulations, title 42, §93.103). Grantee Institution is expected to have established procedures in place for addressing research misconduct and is charged with fostering research integrity through enforcement of its Institutional policies. The Association's Panel on Ethical Scientific Programs (ESP) works on behalf of Association scientific programs and publications to objectively and efficiently investigate cases of potential or perceived misconduct. Grant data and research records, including the original versions of manuscripts, figures and other files and supporting materials, must be retained for a period of seven (7) years and must be produced for review by ESP, if requested. Refer to the Association's Policies and Procedures Related to Scientific Ethics and Integrity in Research Grants ("Misconduct Policy") for further details	The Ellison Medical Foundation shall be indemnified by the awardee institution against any and all claims arising out of the project to the extent allowed by law.	When relevant the foundation will require IRB, IACUC, and documentation that the PI has completed trainin on the protection of human research participants before an award can be made. Annual updates must be submitted with progress reports.	For multiyear awards, funding for renewal years is contingent upon submission of an acceptable annual progress report. For all awards, the awardee institution must provide final scientific progress and financial stewardship reports within 60 days of the termination of the award. Failure to submit final scientific and financial reports will render the institution ineligible for consideration of future funding by the Ellison Medical Foundation	The Ellison Medical Foundation specifically desires that all research results shall be made available to the public as promptly as possible in keeping with the foundation's commitment to free and open publication of research results to both validate findings and to allow other scientists to further advance work in the field. The foundation recognizes and endorses the obligation of awardee institutions to take steps to bring the practical applications of its research forward to public use and benefit	See data sharing.
FORD FDTN		organization will not promote or engage in violence, terrorism, bigotry or the destruction of any State, nor will it make subgrants to any entity that engages in these activities. It is understood that all grant activities shall be performed in accordance with regulations promulgated by the Office of Foreign Asset Controls of the U.S. Department of Treasury ("OFAC") and that no Foundation funds shall be used to provide funds or services to any person or entity listed on either the Specially Designated Nationals List or the Consolidated Sanctions List published periodically by OFAC. It is further understood that no Foundation grant funds shall be used for any prohibited activity in any countries under sanctions by the U.S. government, including prohibited travel to and from those countries, or for the unauthorized provision of funds or services to any person, entity or organization from those countries.	not specified	not specified	not specified	not specified	Final written report including project narrative and financial accounting, according to line item categories of the submitted grant udget, which includes a statement by the responsible financial officer of your organization certifying the accuracy of the report.		At the end of the grant period the Foundation also shall be furnished a copy of any publication, audie or video program, film, or other media product produced by your organization under this grant for archival and/or research purposes. The Foundation shall have the right to make and disseminate additional copies of any such grant product. In addition, your organization hereby grants to the Foundation a license to disseminate on the Foundation's website any product produced by your organization under this grant.
Gordon and Betty Moore Foundation	No.	not specified	not specified	not specified	Not specified	not specified	Interim and final progress reports; detailed outcome table included with award	Grantee must make the data developed in whole or in part with any Grant funds publicly available as soon as possible at no cost, or, when justified, at a reasonable cost	Moore Foundation's Open Access Policy requires that a final (post-print) version of all peer-reviewed articles produced as a result of research supported, either in whole or in part, by the Foundation's funding, be made publicly and freely available (open access) within 12 months of publication.

JDRF	Not specified in online materials.	The Grantee Institution agrees to comply with applicable United States economic sanctions, anti-terrorism laws, and anti-money laundering laws, including, but not limited to, the USA PATRIOT Act , the laws administered by the United States Treasury Department's Office of Foreign Assets Control, Executive Order 13224, and any state or local laws that apply in the jurisdiction in which the Grantee Institution is operating. This provision must be included in allrelevant subcontracts/subawards, when applicable.	Notify JDRF of conclicts.	Agreeing to terms acknowledges acceptance of policies related to misconduct. Any misconduct (scientific or financial) must be reported to JDRF immediately. Follows NIH guidelines.		Per federal government guidelines (NIH).	Annual reports due 30 days prior to next funding period. Final report due 75 days after end (includes final expenditures).	Not specified.	It is a condition of JDRF funding that the final peer-reviewed manuscripts be made available in the PubMed Central online archive immediately upon acceptance of journal publication. Final manuscript within 12 months of publication. Electronic version immiately upon acceptance.
John Templeton Foundation		Throughout the term of this agreement, Grantor shall comply with all appliacble laws, regulations, rules decrees nd orders, whether federal, state, local, domestic or foreign, including but not limited to U.S. Treasury Department's OFAC. Compliance described in the OFAC Compliance section below and IRS rules and regulations relating to tax witholding and reporting for payments to non - U.S> persons. OFAC COmpliance. Grantee represents that (i) it does not knowingly support (including through subgrants), employ or do business with, directly or indirectly, individuals, entities, or groups that are subject to OFAC sanctions, or with individuals, entities or groups known to Grantee to support terrorism or to have violated OFAC sanctions and (ii) it has not changed its name in the last 5 years except as previously disclosed in writing to the Grantor..... (also includes inspection provision).	not specified	not specified	Each party shall be responsible for its negligent acts or omissions and the negligent acts of its employees, officers, or directors, to the extent allowed by law.	not specified	6 year retention of all grant financial records. Reporting (Progress and financial) on predetermined schedule in appendix based on proposal milestones.	not specified	not specified
Kresge Foundation		your knowledge, your organization, members of your governing body, your staff, and any consultants/contractors for your project do not advocate, plan, sponsor, commit, threaten to commit, or support terrorism. By your acceptance of this grant, you agree to provide us with information required for us to comply with Exevcutive Order 13224, the USA Patriot Act, and other applicable laws, administrative rules, and Executive Orders. By accepting this grant, you further agree that all dfunds, including subawards to subrecipients, will be used in complianc with all applicable anti-terrorist financing and asset control laws, regulations, rules, and executive orders. In this regard, you agree to take reasonable steps to ensure that no person or entity expected to receive funds in conection with this Grant Agreement is designated on (a) the Annex to Executive Order No. 13224, as amended or supplemented from time to time, or (b) the Lists of Specially Designated Nationals or Blocked Persons mainained by the Office of Foreign Asset Controls of the U.S. Department of Treasury. Finally, you certify that you will not provide material support or resources to an individual or entity that advocates, plans, sponsors, engages in, or has engaged in, terrorist activity, or that has been so designated, and will immediately cease such support if an entity is so designated after the date of the Grant Agreement.	not specified	not specifies	none	none	Annual reports including project narrative and financial report. Financial record retention 5 years.	none	none
Laura and John Arnold Foundation		not specified	not specified	not specified	[redacted] directors, founders, employees and agents and each of their affiliates (collectivelty the "Foundation Parties"), from and against, any liability, damage, loss or expense (including reasonable attorneys' fees and expenses of litigation) incurred or imposed upon the Foundation Parties in connection with any claims, suits, actions, demands or judgements, arising out of or related to (a) any act or omissionof Grantee, is employees or agents in applying for or accepting the Grant; (b) the expending of the Grant funds furnihed pursuant to this agreement; or (c) the carrying outof any programs or projects funded by the Grant. The Foundation shall not be liable for any losses, damages, claims or other liabilities arising out of Grantees activities. It is expressly understood tht the Foundation, by making the Grant and entering into this Agreement, has no obligation to provide othr or additional support to Grantee	not specified	annual progress reports and final report including financial reporting(detailed expenditures). Financial record retention for 3 years after the end of the award.	not specified	not specified
Lymphoma Research Foundation	No, some terms may be negotiated	Not specified	Not specified	Grantees agree to abide by all applicable federal standards defining integrity and misconduct in research.	The Sponsoring Institution and the LRF Grantee shall, jointly and severally, indemnify and defend LRF and hold it harmless against any and all liabilities, claims, and demands that relate to the research of the LRF Grantee or the LRF Grant, including, but not limited to those for personal injury, property damage, or malpractice. (Note: this term is negotiable for institutions which can't indemnify or prefer to have the institution as the sole party)	Projects involve human/animal subjects must meet or exceed standards for Federal Government funding. Certifications from institution IRB/Animal Review Committess must be provided to LRF before funds are released (if not uploaded with application).	Varies slightly but most programs one yearly written progress report, including financials, on LRF template, to be reviewed by LRF's grant oversight committee. Deadline a minimum of 60 days past grant anniversary date. Funding may be held if report is not received in time and again if oversight committee needs further information to approve the report.	"Unique research resources" produced by LRF funding (including data, models, computer programs, etc. should be made available to other researchers when "resonably requested to do so" - unique biological information should be submitted to appropriate data banks.	Researchers should submit or have submitted final manuscript (including postprint) to Pub Med Central, to be made publicly available no later than 12 months after official publication date.

Paul Allen Frontiers Group	Not listed in materials available on wet	Not listed in materials available on website	Yes, all investigators must complete training program (and retate every 4 years) or whithin 30 days if the policy changes, disclose significan financial interests, subrecipients can rely on their own policy but must confirm that their policy complies with regulations and report all FCOIs to Allen within 45 days of discovery	Not listed in materials available on website	Not listed in materials available on website	Not listed in materials available on website	Not listed in materials available on website	Can't find the actual policy but clearly encourage if not require data sharing as part of its funding	We pride ourselves on our open science approach, publishing data online as soon as they pass our quality standards, and often before we publish on it ourselves.
PCORI	Do appear to accept attachments or addendums to their contract, unclear what form those take.	Recipient shall comply with all applicable federal, state, and local laws, regulations, and requirements of any applicable jurisdiction. (Does not otherwise specify.)	Required to list conflicts of interest in grant contract and include a management plan for any COI for recipient, PI, or Key personnel.	Recipient required to have its own polies and procedures including with respect to data privacy and is expected to enforce those guidelines. Required to repoart any findings of research/financial misconduct to PCORI within 30 days of the conclusion of an investigation (related to PCORI funded reseaarch). Must be notified in writing with nature of violation, corrective actions that will be taken and timeline for those actions. PCORI reserves the right to take their own corrective action or terminate the Contract.	To the extent permitted under applicable law, recipient agrees to indemnify PCORI (and all agents) including reasonable attorney's fees and other expenses incurred by PCORI on account of any willful or negligent act or omission of the Recipient (or its agents). Recipient's obligation to indemnify shall be limited to the extent that recipient is afforded sovereign immunity under applicable laws.	Human subjects shall have and maintain up-to-date IRB approval and must provide PCORI with copies of the approval documentation. (Doesn't specify animal, but PCORI does not really cover bench/translational research.)	Milestones set through agreement between PCORI and recipient, Recipient to submit Interim progress reports (schedule set in contract), including technical and non technical sections, a draft final research report for peer review, then revise and submit final taking peer review comments into account. Reporting on the status of recruitment, subcontracting, and IRB oversight also specified in contract, as well as other specialized reports where applicable. Final report to be submitted no later than 60 days after ending of grant term.	Recipient shall develop and maintain a plan that addresses data management and data sharing of Research Project data in a manner that is appropriate for the nature of the Research Project and the types of Research Project data, and that is consistent with applicable privacy, confidentiality, and other legal requirements.	Recipient shall ensure that an electonic copy of the final peer-reviewed manuscript of any research finding are submitted to PubMedCentral.
Pew Charitable Trust	No. PCT open to negotiating standard terms for all awards to the institution.	Not specified	Not specified	Not specified	Grantee shall indemnify Pew from all claims, etc. arising from or relating to our breach of any provision of the agreement, our performance under each project, and intentional misconduct. Also, if any federal, state or local government or administrative agency determines that our personnel acted as a Pew employee while conducting work under an agreement, we indemnify Pew from any liabilities related to such determination.	Not specified	Narrative and financial reports, as specified for each project	Not specified	Not specified
Prostate Cancer Foundation	I tried to call to obtain info. Then resorted to email getting the following response: We don't have these policies stated on our website and we currently don't have any open RFAs. Using the link I sent you, you will be able to download previous RFAs which may contain some of the information you're seeking. Other than that, we don't have policy statements in a format that we make public or share upon request.	not specified online or in RFAs	Not specified online or in RFAs	not specified	not specified	An IRB-approved protocol ready for activation is required for all human-related studies, including specimen acquisition. In addition, documentation of the availability of the experimental agent to be studied is required.	not specified semiannual progress reports and finanacial reports	Not specified.	Not specified
Rita Allen Foundation	no	none	none	none	none	none	none	none	none
Robert Wood Johnson Foundation	No.	not specified	not specified	Not specified	Not specified	Grantee must certify that research will be conducted in compliance with 45CFRPart46, and related guidance	Annual and final narrative reports	Foundation retains a license to republish data, or use/license to other to use data for Foundation's charitable, educational and scientific research purposes.	not specified
SLOAN FDTN	no	not specified	not specified	not specified	not specified	not specified	Final written and financial report	not specified	not specified
ST BALDRICKS FOUNDATION	Not specified	not specified	Not specified	It is the responsibility of the institution to immediately report to SBF if it has a reasonable good faith basis to believe there has been Misconduct, and to report any Misconduct or change in the funded researcher's employment status with the institution, including administrative leave, which may occur during the term of any award that is pertinent related to the work described in the grant application. Failure to abide by the terms above, or any other SBF policies and procedures in connection to the application and/or grant, may result in SBF suspending grant funding or canceling the grant, to be decided by SBF in its sole discretion. If a grant is discontinued, the PI shall receive a 30-day written notice. Any funds unspent at the time the notice is issued shall be returned to SBF.	Identification with the Foundation shall also be made in any news released about the fellowship or the Fellow's research project by the public relations department or its equivalent at the sponsoring institution.	All human subject research or research on human tissue which is supported by St. Baldrick's must comply with the regulations applicable to that supported by the National Institutes of Health. As part of the application, the applicant will be asked to submit documentation of approval of the study by his or her Institutional Review Board (IRB), along with the Institution's Human Subjects Assurance Number. Approval is not required at the application stage but will be required prior to issuing funds. If the grantee's research plan changes after the award to include human subjects or tissue, the grantee must submit proper documentation of IRB approval to St. Baldrick's. The Foundation adheres to the most current guidelines applicable to the care and treatment of animals used in laboratory work as outlined by the National Institutes of Health. As part of the application, the Fellow must submit a statement that the institution meets and adheres to these policies. Failure to notify the Foundation of compliance with these guidelines or the improper use of laboratory animals may result in termination of the fellowship.	The Fellow must submit reports of his/her annual research progress online via ProposalCENTRAL by the dates specified in ProposalCENTRAL. In addition to the scientific report, this includes a lay report written for the general public. The progress report shall be accompanied by an evaluation report from the Mentor directly responsible for the Fellow's work. A brief Interim Update verifying receipt of funds and usage is due after the first three (3) months of each grant year. These reports must be completed using the templates available on ProposalCENTRAL. Progress reports that are more than thirty (30) days late will impact the Primary Investigator's consideration and release of future awards. These reports shall be reviewed by the Foundation to evaluate the research progress of each Fellow. The Foundation reserves the right to terminate any fellowship if it determines that there has been inadequate research progress or a failure to adhere to the original proposal submitted with the application.	Not specified.	Not specified

The Leukemia Lymphoma Society	Unclear, some guidelines state that "by	Not specified	Not specified	Sponsoring Institution will enforce Grantee following Institution's research misconduct policies, and represents that such policies are at least as rigorous as those followed by the NIH.	LLS assumes no responsibility for any of the activities of the Sponsored Research, including any acts or omissions of Grantee. Sponsoring Institution will indemnify, defend and hold LLS harmless from any and all claims, damages, costs and expenses that may arise as a result of the Sponsored Research and the activities of the Grantee in connection with this Agreement unless caused by the willful misconduct or gross negligence of LLS and to the fullest extent authorized under the Constitution and laws of Sponsoring Institution's state, if applicable.	Sponsoring Institution will ensure that Grantee obtains prior written approval from the Sponsoring Institution's Institutional Review Board (or equivalent institutional authority) ("IRB") for the protection of human/animal subjects before undertaking any form of human/animal subject research. An original executed copy of this approval must be submitted to LLS within ten (10) days after such approval is obtained. With respect to research projects that do not deal with human subject research, Sponsoring Institution must furnish to LLS a letter executed simultaneously with this Agreement stating that: "The research project does not involve the use of human subjects or human tissue." (For animals allowed to just state in the application that no animals will be used.)	Varies depending on program in general appears to be yearly until end of grant, and then 60 days following end of grant. Eac report must include an updated lay summary reflecting progress since original application. Report submitted via LLS template through their online portal.	Not specified	Not specified
The National Pancreas Foundation	No, grantee must sign terms at time of application	Not specified	Not specified	Signing attestation statement includes "the investigator's commitment that the research will be conducted under the supervision of an Institution at which appropriate standards and oversights are working."	Not specifed	The principal investigator's institution will be responsible for certifying to the Foundation that the research is conducting in accordance with current medical research standards and that is in compliance with the current guidelines of the U.S. Department of Health and Human Resources regarding Vertebrate Animals, Recombinant DNA, Research Misconduct, and Financial Conflict of Interest. The institution will further verify that any research involving human subjects or vertebrate animals has been approved by the necessary review boards and committees, in accordance with existing laws and regulations.	Grants are one-year only, final financial report and written summary of research results listing all publications.	Not specified	Not specified
The Simons Foundation		not specified	not specifies	not specified	not specified	Follows NIH guidelines for animal and humans . Approvals in place at start of award.	progress reports due 60 days following end of funding year. Carry forward and NCE requests must be made 30 days prior to end of period. Final invention sttement must be submitted within 60 days of end date. Financial statements due 60 days after funding period (annual) No revisions <$500, over $500 only if grant still active.	It is essential that instituions and Pis share renewable reagents and data developed using Simons Foundation funds with other qualified investigators. Pis will be required to have renewable reagents and data-sharing plan in place prior to receiving a grnat. At the Foundation's discretion, the PI may be required to provide the foundation with an electronic copy of all properly deidentified research-generated data prior to the end of the grant. In all reasonable cases, the foundation will assume financial responsibility for costs associated with the data transfer.	The Simons Foundation encourages Pis to post preprints on recognized servers, such as arXiv or bioRxiv, in parallel with (or even before) submission to peer reviewed journal. The Simons Foundation also encourages Pis to publish under Open Access licenses. All preprints and Open Access publication resulting from projects supported by the Simons Foundation must be submitted through proposalCENTRAL with the nnual and Final Progress Report web forms.
V FOUNDATION	no	none	Must have written guidelines	Must have written guidelines. It is the responsibility of the institution and the grantee to inform TVF immediately, in writing, of any misconduct finfding resulting from an investigation into the conduct of an investigator whose work is supported by TVF, and to keep TVF informed in a timely manner of the progress and outcome of such investigation. A finding by the institution that the investigator was guilty of fraud is sufficient grounds to terminate TVF's support of this project.	TVF does not assume responsibility for activities supported by the grant and shall not be liable for any activities conducted by the grantee upon receipt of the grant funds. The grantee agrees to indemnify and hold harmelss, to the extent permitted by state law, TVF and its members, directors, employees and agents against and from all liabilities and expenses (including reasonable attorney's fees and court costs) which are threatened, amay be imposed upon, incurred by, or asserted against such persons relating to the grant funds and any research project conducted by the grantee or research results developed by grantee.	NIH/NCI guidelines and IRB approval for human subjects, IACUC approval animal use; clinical must have IND approval	annual ptrogress report including scientific progress report, publications, follow-on grant funding, lay summary, financial report comparing actual to budget with explanations for >10% variance	no	Yes through HRA initiative: Pubmed ce
William and Flora Hewlett Foundation		**Anti-Terrorism**. You agree to use Grant Funds in compliance with all applicable antiterrorist financing and asset control laws, regulations, and executive orders, including without limitation Title 31 Chapter V of the U.S. Code of Federal Regulations. You will not use Grant Funds to (a) provide resources or support to parties associated with terrorism and the terrorist-related lists promulgated by the U.S. Government, including the List of Specially Designated Nationals (www.treasury.gov/sdn); or (b) support transactions or dealings in countries against which the U.S. maintains a comprehensive embargo (currently, Cuba, Iran, (North) Sudan, Syria, North Korea, or the Crimea Region of Ukraine), unless such transactions or dealings are fully authorized by the U.S. government under applicable law and specifically approved by the Foundation in its sole discretion.							