

NIMH Data Archive

Data Use Certification

Last updated: May 2020

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Introduction

The National Institute of Mental Health (NIMH) Data Archive (NDA) is an NIH-funded collaborative resource that contains human subjects research data from multiple research data repositories.

The NIMH Data Archive Data Use Certification (DUC) is used to request access to shared research data in a data repository within the NIMH Data Archive. Shared data are available with either an Institutional sponsorship or an Individual sponsorship. All data access requests require acceptance of the Data Use Terms and Conditions contained in this DUC. (See the *NIMH Data Archive Recipient Information and Certifications* form in this document for available data and associated sponsorship types.)

- Institutional sponsorship requires Recipients to be affiliated with an NIH recognized institution (foreign or domestic), based upon registration in the NIH's eRA Commons system, with an active Federal Wide Assurance (FWA) issued by the Department of Health and Human Services, Office for Human Research Protections (OHRP). The signature of an Authorized Institutional Business Official is also required on this DUC.
- Individual sponsorship may be requested by a Recipient without the need for sponsorship by or affiliation with an NIH recognized institution and, therefore, the signature of an Authorized Institutional Business Official or an active institutional FWA is not required.

A Data Access Committee(s) (DAC) will objectively review a data access request sponsored by an Institution. Individual sponsorships do not require DAC review. To submit data to the NIMH Data Archive, the NIMH Data Archive Data Submission Agreement (DSA) must be completed, which is a separate document.

The NIMH Data Archive (NDA)

The National Institutes of Health (NIH) and NIMH have developed a data infrastructure to store the collection of data from participants in research studies, regardless of the source of funding. The extensive information collected by these studies is harmonized and subsequently stored in one of several data repositories within the NIMH Data Archive (NDA) data infrastructure, providing a rare and valuable scientific resource. A current list of all NDA data repositories and links to their websites is available at <https://nda.nih.gov/about/about-us.html>.

The NIH and NIMH seek to encourage the use of these resources to achieve rapid scientific progress. Moreover, NIMH has made data sharing a requirement for all clinical research it funds (see [NOT-MH-19-033](#)). In order to take full advantage of such resources and maximize their research value, it is important that data are made **broadly available**, on appropriate terms and conditions, to the largest possible number of qualified investigators in a timely manner.

Data collected by the Submitters have been stripped of all individual identifiers, but the unique and intrinsically personal nature of genomics data, brain imaging, and other derivative data of which are included in these repositories, combined with the recent increase in the accessibility of conducting genotype and other sequence analyses (in terms of technological capacity and cost), has altered the framework through which "identifyability" can be defined. To protect and assure the confidentiality and privacy of all participants, the Recipient who is granted access to these data is expected to adhere to the specifications of this DUC. Failure to do so could result in denial of further access to data.

Data Use Terms and Conditions

I request access to shared data from the NIMH Data Archive for the purpose of scientific investigation, scholarship or teaching, or other forms of research and research development as described in the following NIMH Data Archive Data Use Certification (DUC). I, and any Other Recipients listed in this DUC, agree to the following terms:

1. Research Data Use Statement

Generally, these data will be used by the Recipient in connection with the purpose indicated and described in the *Research Data Use Statement* on the DUC. Recipients are encouraged to explore shared data in the NIMH Data Archive for a variety of purposes including secondary analysis, hypothesis generation, and replication regardless of whether said exploration leads to analysis in support of a question beyond the scope of the originally identified purpose described in the *Research Data Use Statement*.

2. Non-transferability of Agreement

This DUC is not transferable. If a Recipient changes institution and wishes to retain access to the NIMH Data Archive, a new DUC is required.

3. Non-Identification of Subjects

Recipients agree that data will not be used to establish the individual identities of any of the study participants from whom data were obtained (or their relatives) and/or contact the individual study participant, except as permitted by law (e.g., in connection with a separately negotiated collaboration with the original research team or the enrollment of the consented subject in the Recipient's study). Recipients agree to not publish or disseminate any derived data that could aid in the re-identification of any of the study participants (or their relatives). Recipients agree to notify the NIH at NDAHelp@mail.nih.gov as soon as possible if, upon use of NIMH Data Archive data, identifying information is discovered.

4. Use of the NIH Global Unique Identifier (GUID)

The Global Unique Identifier (GUID) is a computer-generated alphanumeric code that is unique to each research participant. The GUID allows the NIMH Data Archive to link together all submitted information on a single participant, giving researchers access to information even if the data were collected at different locations or through different studies. If Recipients access data on individuals for whom they, themselves, have previously submitted data to the NIMH Data Archive, Recipients may gain access to more data about an individual participant than they, themselves, collected. Consequently, these research activities may be considered "human subjects research" within the scope of 45 C.F.R. 46. Recipients must comply with the requirements contained in 45 C.F.R. 46, as applicable, which may require Institutional Review Board (IRB) approval of the Research Data Use Statement.

5. Data Disclaimers

Recipients acknowledge that the NIH does not and cannot warrant the results that may be obtained by using any data or data analysis tools included in the NIMH Data Archive. The NIH disclaims all warranties as to the accuracy of the data in the NIMH Data Archive or the performance or fitness of the data or data analysis tools for any particular purpose.

6. Data Access for Research

Data and Supporting Documentation in the NIMH Data Archive are eligible for access by qualified researchers, pursuant to the terms set forth in this DUC. Recipients acknowledge that other researchers have access to the data and that downloading, and duplication of research is a distinct possibility,

thereby decreasing subject data protections. Raw or nearly raw research data files (e.g. fastq, bam, MRI, and EEG recordings) are made available for just in time computation, regardless of where the computational resources may reside. Therefore, data copied shall not be persisted (i.e., stored) beyond the time necessary for computation and shall be expunged once computation has been completed. Recipients are encouraged to utilize the NIMH Data Archive computational capabilities described at <https://nda.nih.gov/tools/nda-tools.html#cloud>.

7. Supporting Documentation

Recipients agree to review the supporting information, materials, and documentation (“Supporting Documentation”) for the data accessed in the NIMH Data Archive to enable efficient use of the submitted data by Recipients unfamiliar with the data or the research project. Examples of supporting documentation include:

- Research protocol(s)
- Questionnaire(s)
- Study manuals

8. Sharing of a NIMH Data Archive Study/Acknowledgements

Recipients agree to create and share an NIMH Data Archive Study (<https://nda.nih.gov/get/manuscript-preparation.html>) for each publication, computational pipeline, or other public disclosure of results from the analysis of data accessed in the NIMH Data Archive, whether reporting positive or negative results, thereby linking it to the underlying data. Recipients agree to create the NIMH Data Archive Study when a manuscript is submitted for review and share the Study when the publication is released. Recipients agree to acknowledge the appropriate NIMH Data Archive data repository and the relevant Digital Object Identifier(s) (DOI), which will be created by NIMH Data Archive staff, in any and all oral and written presentations, disclosures, and publications (including abstracts, as space allows) resulting from any and all analyses of data, whether or not the Recipient is collaborating with Submitter(s). The oral or written presentation, disclosure, or publication should include an acknowledgement statement, which includes a disclaimer of NIH endorsement, as appropriate. Acknowledgements specific to each NDA data repository are maintained at <https://nda.nih.gov/get/manuscript-preparation.html>.

If the Research Project involves collaboration with Submitters or NIH staff (as indicated in the DUC), then Recipient will acknowledge Submitters or NIH staff as co-authors, if appropriate, on any presentation, disclosure, or publication.

9. No Distribution of Data

Recipients agree to retain control over data from the NIMH Data Archive, and further agree not to transfer or sell data, with or without charge, in any form, to any other entity or any individual or to distribute the data to anyone other than the Other Recipients listed on this DUC who also agree to the terms in this DUC. This includes any data derived from the data in the NIMH Data Archive if the associated GUID is distributed with that derived data or if the derived data can aid in the re-identification of a research participant.

10. Non-Governmental Endorsement; Liability

Recipients agree not to claim, infer, or imply endorsement of the research project described in the *Research Data Use Statement*, the entity, or personnel conducting the research project or any resulting commercial product(s) by the United States Government, the Department of Health & Human Services, the National Institutes of Health, or the National Institute of Mental Health. The United States Government assumes no liability except to the extent provided under the Federal Tort Claims Act (28

U.S.C. § 2671-2680).

11. Recipient's Compliance with Institutional Requirements

Recipients with Institutional sponsorship acknowledge that access, if provided, is for research that is approved by the Institution with which they are affiliated, which must be operating under an active Federal Wide Assurance (FWA) issued by the Department of Health & Human Services, Office for Human Research Protections (OHRP). Furthermore, Recipients agree to comply with all applicable rules for the protection of human subjects, which may include Department of Health and Human Services regulations at 45 C.F.R. Part 46, and other federal and state laws for the use of this data. Recipients agree to report promptly to the NIH any unanticipated problems involving risks to subjects or others. This DUC is made in addition to, and does not supersede, any of Recipient's institutional policies or any local, State, and/or Federal laws and regulations that provide additional protections for human subjects.

12. Recipient's Permission to Post Information Publicly

Recipient agrees to permit the NIMH Data Archive to publicly summarize the Recipient's research use of data along with the Recipient's name and organizational/institutional affiliation.

13. Privacy Act Notification

Recipients agree that information collected by the NIH from a Recipient, as part of the DUC, may be made public in part or in whole for tracking and reporting purposes. This Privacy Act Notification is provided pursuant to Public Law 93-579, Privacy Act of 1974, 5 U.S.C. Section 552a. Authority for the collection of the information requested below from Recipients comes from the authorities regarding the establishment of the National Institutes of Health, its general authority to conduct and fund research and to provide training assistance, and its general authority to maintain records in connection with these and its other functions (42 U.S.C. 203, 241, 289l-1 and 44 U.S.C. 3101), and Sections 301 and 493 of the Public Health Service Act. These records will be maintained in accordance with the Privacy Act System of Record Notice 09-25-0156

([https://oma.od.nih.gov/forms/Privacy%20Documents/Documents/Privacy%20Act%20Systems%20of%20Records%20Notices%20\(SORNs\)%205-1-15.pdf](https://oma.od.nih.gov/forms/Privacy%20Documents/Documents/Privacy%20Act%20Systems%20of%20Records%20Notices%20(SORNs)%205-1-15.pdf)) covering "Records of Participants in Programs and Respondents in Surveys Used to Evaluate Programs of the Public Health Service, HHS/PHS/NIH/OD." The primary uses of this information are to document, track, monitor, and evaluate the use of NIMH Data Archive datasets, as well as to notify interested Recipients of updates, corrections or other changes to the database.

The Federal Privacy Act protects the confidentiality of some NIH records. The NIH will use the information collected for the purposes described above. In addition, the Act allows the release of some information in the Recipient's records without the Recipient's permission; for example, if it is requested by members of Congress or other authorized individuals. The information requested in this DUC is voluntary, but necessary for obtaining access to data in the NIMH Data Archive.

14. Security

Recipients acknowledge that the data being made available were made available for researcher use with the expectation that the data will be protected in a manner consistent with security best practices. Such practices include, but are not limited to, the following:

- Accounts and passwords will not be shared.
- Data are protected from anonymous access. Any data transferred or stored outside of the NIMH Data Archive will be protected using standard encryption protocols and/or strong password protection.
- When finished using the data, the data will be expunged, as permitted by law.

15. Annual Update/Research Use Reporting

Recipients will provide to NDAHelp@mail.nih.gov an annual summary of research accomplishments from using the NIMH Data Archive and agree to create and share an NIMH Data Archive Study for each public disclosure of results pursuant to the Sharing of an NIMH Data Archive Study/Acknowledgements term in this DUC. The NIH encourages Recipients who publish manuscripts based on a combination of data from the NIMH Data Archive data, data derived from NDA data, and data collected independent of the NIMH Data Archive to consider submitting the complete analyzed dataset to the NIMH DataArchive.

16. Amendments

Amendments to this DUC must be in writing and signed by authorized representatives of all parties.

17. Termination

Either party may terminate this DUC, without cause, provided 30 days' advanced written notice to the other party. Recipients agree to immediately report violations of this agreement to the appropriate NIMH Data Archive Data Access Committee. Additionally, the NIH may terminate this agreement with 5 days' advanced written notice if the NIH determines, in its sole discretion, that a Recipient has committed a material breach of this DUC. The NIH may, in its sole discretion, provide a Recipient with 30 days' advanced written notice to remedy a breach before termination.

18. Term and Access Period

Recipients are granted permission to access requested and approved data from the NIMH Data Archive for a period of one year and this DUC will automatically terminate at that time. Data access may be renewed upon certification of a new DUC.

19. Accurate Representations

Recipients expressly certify that the contents of any statements made or reflected in this document are truthful and accurate.

Burden Disclosure Statement

Public reporting burden for this collection of information is estimated to vary from 15 min to 1.5 hours per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing the collection of information. **An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number.** Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden, to: NIH, Project Clearance Branch, 6705 Rockledge Drive, MSC 7974, Bethesda, MD 20892-7974, ATTN: PRA (0925-0667). Do not return the completed form to this address.

NIMH Data Archive Recipient Information and Certifications

1. Access Request:

Application Type			Permission Group	Recipient Sponsor*
NEW		RENEWAL	NIMH Data Archive (NDA)	Institutional
NEW		RENEWAL	Adolescent Brain Cognitive Development Study (ABCD)	Institutional
NEW		RENEWAL	Connectome Coordination Facility (CCF)	Institutional
NEW		RENEWAL	Osteoarthritis Initiative (OAI)	Individual
NEW		RENEWAL	NIAAA Data Archive (NIAAA _{DA})	Institutional
NEW		RENEWAL		

**Institutional sponsorship requires the signature of an Authorized Institutional Business Official and an active Federal Wide Assurance (FWA) number in the Signatures section below. See the "Introduction" section on page 1 for more information.*

See <https://ndaa.nih.gov/about/about-us.html> and https://ndaa.nih.gov/user/dashboard/data_permissions.html for a current list of all NDA Data Repositories and Permission Groups.

2. Lead Recipient:

First Name: _____ Last Name: _____ Degree: _____

Institution: _____

City: _____ **State/Province:** _____ **Country:** _____

Phone: _____ E-mail Address: _____

3. Research Data Use Statement: Describe the purpose of the scientific investigation, scholarship or teaching, or other form of research and research development for which you are requesting access to the NIMH Data Archive. If you are requesting access to a controlled access permission group, this statement must demonstrate adherence to the consent-based data use limitations.

4. Renewal Applicants Only:

Has a publication, computational pipeline, or other public disclosure of results from the analysis of data accessed in the NIMH Data Archive resulted from a Recipient's previous access period? Yes No

If Yes, has an NDA Study been created? Yes List the NDA Study number(s): _____
 No* List the PubMed ID(s) or citation(s): _____

* See 8. Sharing of a NIMH Data Archive Study/Acknowledgements above. Contact the NDA Help Desk (NDAHelp@mail.nih.gov) to create an NDA Study.

5. Other Recipient(s): List all individuals who will access, use, or analyze the data regardless of position title or data use role. Use additional sheets as needed.

First Name: _____ Last Name: _____ Degree: _____

Institution: _____

City: _____ State/Province: _____ Country: _____

Phone: _____ E-mail Address: _____

First Name: _____ Last Name: _____ Degree: _____

Institution: _____

City: _____ State/Province: _____ Country: _____

Phone: _____ E-mail Address: _____

First Name: _____ Last Name: _____ Degree: _____

Institution: _____

City: _____ State/Province: _____ Country: _____

Phone: _____ E-mail Address: _____

First Name: _____ Last Name: _____ Degree: _____

Institution: _____

City: _____ State/Province: _____ Country: _____

Phone: _____ E-mail Address: _____

First Name: _____ Last Name: _____ Degree: _____

Institution: _____

City: _____ State/Province: _____ Country: _____

Phone: _____ E-mail Address: _____

- 6. Authorized Institutional Business Official:** Requests to access data requiring an Institutional sponsorship must list an individual with an “SO” role as defined in the NIH eRA Commons - <https://commons.era.nih.gov/commons>

Name: _____ Email Address: _____

- 7. Signatures:** By signing and dating this DUC to request access to data in the NIMH Data Archive, I and my Institutional Official (*if required*) certify that we will abide by the Data Use Terms and Conditions defined in this DUC. I further acknowledge that I have shared this document with any Other Recipients who will participate in the use of data from the NIMH Data Archive. My Institutional Business Official (*if required*) also acknowledges that they have shared this document with appropriate institutional organizations.

Lead Recipient Signature

Date

Authorized Institutional Business Official Signature (*if required*)

Date

Inquiries and requests to access data in the NIMH Data Archive should be sent to:

Office of Technology Development and Coordination (OTDC), Program Director
National Institute of Mental Health | National Institutes of Health
6001 Executive Boulevard, Room 8125, MSC 9640 Bethesda, MD 20892-9640
Telephone: 301-443-3265 | Email: NDAHelp@mail.nih.gov