SUBJECT: PAGE CHANGE TO DOE O 443.1B, PROTECTION OF HUMAN RESEARCH SUBJECTS

EXPLANATION OF CHANGE. The purpose of making the page change is to harmonize the definitions in Section 7 of this Order with those in the Federal regulations for the protection of human subjects (10 CFR Part 745). Specifically, the definition of “human subjects research” has been split into “research” and “human subject” and the definitions of “human subject” and “research” have been adopted, verbatim, from 10 CFR Part 745. The definition of “generalizable” has been added, since the determination of whether a project is “research” in 10 CFR Part 745 hinges on whether the work being conducted is generalizable. Additionally, the reference to the “Secretarial Policy Memo on Military or Intelligence-Related Human Subjects Research,” in Section 1, has been deleted, as the requirements specified in that memo were already incorporated into the Order (in March 2011). A number of small corrections and updates to acronyms, references, links, and organization titles have also been made throughout the Order.

1. LOCATIONS OF CHANGES.

<table>
<thead>
<tr>
<th>Page</th>
<th>Paragraph</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Section 1, “Purpose”</td>
</tr>
<tr>
<td>2</td>
<td>Sections 3b, 3c, and 4</td>
</tr>
<tr>
<td>3</td>
<td>Section 4a(2)(d)</td>
</tr>
<tr>
<td>4</td>
<td>Section 4c and 4d(2)</td>
</tr>
<tr>
<td>5</td>
<td>Section 4e</td>
</tr>
<tr>
<td>6</td>
<td>Sections 5a(6), 5b, 5c</td>
</tr>
<tr>
<td>6</td>
<td>Section 5d(4)</td>
</tr>
<tr>
<td>8</td>
<td>Section 5e(1), 5f</td>
</tr>
<tr>
<td>9</td>
<td>Section 5g(10)</td>
</tr>
<tr>
<td>10</td>
<td>Sections 5h, 5h(2)(a)</td>
</tr>
<tr>
<td></td>
<td>5h(2)(b), 6b, 6c, 6d</td>
</tr>
<tr>
<td>11</td>
<td>6e, 6f, 6g, 6h, 6j</td>
</tr>
<tr>
<td>12</td>
<td>Sections 6l, 6r, 7b, 7c</td>
</tr>
<tr>
<td>13</td>
<td>Sections 7g, 7h, 7i</td>
</tr>
<tr>
<td>14</td>
<td>Sections 7j, 7k</td>
</tr>
<tr>
<td>15</td>
<td>7q, 7r, 7s</td>
</tr>
<tr>
<td>CRD 1</td>
<td>Section 1</td>
</tr>
</tbody>
</table>
PROTECTION OF HUMAN RESEARCH SUBJECTS


Cancellation of a directive does not, by itself, modify or otherwise affect any contractual or regulatory obligation to comply with the Order. Contractor Requirements Documents (CRDs) that have been incorporated into a contract remain in effect throughout the term of the contract unless and until the contract or regulatory commitment is modified to either eliminate requirements that are no longer applicable or substitute a new set of requirements.

3. **APPLICABILITY.**

   a. **Departmental Applicability.** Except for exemption in paragraph 3.c., this Order applies to all Departmental elements.

      The Administrator of the National Nuclear Security Administration (NNSA) shall ensure that NNSA employees, contractors, and elements comply with their respective responsibilities under this directive. Nothing in this Order shall be construed to interfere with the NNSA Administrator's authority under Section 3212(d) of Public Law (P.L.) 106-65 to establish Administration-specific policies, unless disapproved by the Secretary.

      In accordance with the responsibilities and authorities assigned by Executive Order 12344, codified at 50 CFR Parts 2406 and 2511, and to ensure consistency throughout the joint Navy/DOE Naval Nuclear Propulsion Program, the Deputy Administrator for Naval Reactors (Director) will implement and oversee requirements and practices pertaining to this Directive for activities under the Director’s cognizance, as deemed appropriate.

   b. **DOE Contractors.** Except for the exemption in paragraph 3.c., the Contractor Requirements Document (CRD), Attachment I, sets forth the requirements of this Order that shall apply to contracts that include the CRD. The CRD shall be included in contracts (i.e., those contracts that include the clause at 48 CFR Part (DEAR) 970.5204-2, Laws, Regulations, and DOE Directives) for the management or operation of a DOE-owned or –leased facility that involves human subjects research (HSR) as defined in paragraph 7.h., and comprehensively explained in Paragraph 4.a., irrespective of the party conducting the HSR under the contract. For all other contracts that involve HSR,
the applicable requirements set forth in this CRD shall be included in the contract terms and conditions as appropriate.

c. **Exemptions for DOE O 443.1B.**

Any requests for partial or full exemptions from the requirements of this Order shall be submitted in writing to the DOE Human Subjects Protection (HSP) Program Manager (and when an NNSA element is involved, the NNSA HSP Program Manager). An exemption may be recommended to the Secretary or the Secretary's designee by the DOE HSP Program Manager (or by the NNSA HSP Program Manager when an NNSA element is involved) after concurrence by the DOE Institutional Official (see paragraph 7f). The basis for granting or denying exemption requests shall be set forth in writing.

**Exemption.** Bonneville Power Administration is exempt from the requirements of DOE O 443.1B.

4. **REQUIREMENTS.** Research using human subjects provides important medical and scientific benefits to individuals and to society. The need for this research does not, however, outweigh the need to protect individual rights and interests. DOE requirements are established in the Federal Policy for the Protection of Human Subjects, 45 CFR Part 46, *Protection of Human Subjects*, in 10 CFR Part 745, DOE’s implementation of Subpart A of 45 CFR Part 46, and in this Order.

a. **Approvals.**

   (1) No HSR conducted with DOE funding, at DOE institutions (regardless of funding source), or by DOE or DOE contractor personnel (regardless of funding source or location conducted), whether done domestically or in an international environment, including classified and proprietary research, shall be initiated without both a Federalwide Assurance (FWA) and approval by the cognizant Institutional Review Board (IRB) in accordance with 10 CFR Part 745.103.

   (2) It is Departmental policy that Human Terrain Mapping (HTM), defined in paragraph 7.k., is managed as HSR and is subject to this Order.

   (a) HTM projects, conducted with DOE funding, at DOE sites/institutions (regardless of funding source), or by DOE or DOE contractor personnel (regardless of funding source or location conducted), whether done domestically or in an international environment, including classified and proprietary research, shall be strictly limited to only those projects involving the analysis and modeling of de-identified data.

   (b) Statements of work for HTM projects shall be submitted to the HSP Program Manager (and when an NNSA element is involved,
the NNSA HSP Program Manager), for DOE Headquarters review and approval prior to initiation. If the project is to be conducted by or for the intelligence community, the Office of Intelligence must also review and approve it prior to initiation. The HSP Program Manager(s) and the Office of Intelligence shall engage the recognized DOE site IRB, and as needed, the principal investigator (PI) and/or sponsor, in clarifying whether the proposed project is HTM and if so, that the data to be used will be de-identified. Additionally, the PI will be asked to provide written verification that only de-identified HTM data (as defined in paragraph 7d) will be used.

(c) The recognized DOE site IRB is the only entity authorized to determine whether the HTM data received by the PI after project initiation meets DOE criteria for de-identification. If the DOE site does not manage or operate its IRB, then the Central DOE IRB shall be the responsible IRB.

(d) All projects funded by other entities, including HTM activities, shall comply with the applicable DOE O 481.1C Admin Chg 2, Strategic Partnership Projects (formerly known as Work for Others (Non-Department of Energy Funded Work)), dated 1-24-05, or DOE O 484.1 Admin Chg 2, Reimbursable Work for the Department of Homeland Security, dated 8-17-06.

(e) If, in the case the sponsor requests assistance in the de-identification of HTM data prior to start of any work on the sponsor’s project and/or re-identification of data following completion of the project, DOE sites may provide such services under a separate contract and/or task order with the sponsor by following the appropriate DOE standard operating procedure approved by the DOE Institutional Official, DOE Office of Science.\(^1\)

b. Solicitations. Any solicitation issued by a DOE element for research involving human subjects shall require compliance with the requirements of this Order, 10 CFR Part 745, and 45 CFR Part 46.

Contracts, Financial Assistance Agreements, and Other Agreements. Any DOE contract, financial assistance agreement, or other agreement involving HSR shall require compliance with the requirements set forth in this Order and/or the CRD

\(^1\) It should be noted that: 1) only limited communications, if needed, may take place between the organization de-identifying and/or re-identifying the sponsor’s data and the organization performing work on the sponsor’s task; b) the identified dataset shall not be shared with the individual who will perform work on the sponsor’s task; and c) the de-identified dataset shall be sent directly by the sponsor to the individual performing work on the sponsor’s task and not by the organization at the DOE site that de-identified it.
associated with this Order (Attachment I), as well as with 10 CFR Part 745 and 45 CFR Part 46. See also CRD (Attachment I).

c. **Notification.** The Human Subjects Protection (HSP) Program Manager (and when an NNSA element is involved, the NNSA HSP Program Manager) shall be:

1. Notified in writing prior to issuance of any new proposal involving HSR, even if it meets the regulatory definition of exempt HSR as outlined in 10 CFR Part 745.101(b), that involves:
   
   a. an institution without an established IRB;
   
   b. a foreign country;
   
   c. a potential for significant controversy (e.g., negative press or reaction from stakeholder or oversight groups);
   
   d. research subjects in a protected class (fetuses, pregnant women, and in vitro fertilization; prisoners; or children); or
   
   e. the generation or use of classified information.

2. The DOE HSP Program Manager (and when an NNSA element is involved, the NNSA HSP Program Manager) shall be notified immediately upon a finding of a suspected or confirmed data breach involving Personally Identifiable Information (PII) in printed or electronic form and reported to the DOE-Cyber Incident Response Capability in accordance with the requirements of DOE O 206.1. The appropriate HSP Program Manager shall also be informed of any corrective actions taken and shall concur on the plan for any remaining corrective actions.

3. The appropriate HSP Program Manager shall be notified in writing within 48 hours, with a description of corrective actions taken, and shall concur on the plan for any remaining corrective actions, following:

   a. unanticipated problems, significant adverse events, and complaints about the research, as well as suspension or termination of IRB approval of research;
   
   b. known or potential incidents of noncompliance with requirements of this Order, 10 CFR Part 745, 45 CFR Part 46.

4. The appropriate HSP Program Manager shall be notified in writing immediately upon the appointment of a new DOE Site IRB Chair or DOE Site Institutional Official.
d. **Reporting.** HSR projects shall be reported annually to the HSR Projects Database (HSRD) in accordance with directions and schedules provided by the appropriate HSP Program Manager.

e. **Protected Classes.** Research involving fetuses, pregnant women, and in vitro fertilization; prisoners; or children shall be conducted in accordance with 45 CFR Part 46 Subparts B, C, and D.

f. **IRB Registration.** Each IRB that is designated by an institution under an assurance of compliance approved for Federalwide use by the Office for Human Research Protections (OHRP) under 45 CFR Part §46.103(a) and that reviews research involving human subjects conducted or supported by the Department of Health and Human Services (HHS) shall be registered with HHS in accordance with 45 CFR Part 46 Subpart E.

5. **RESPONSIBILITIES.**

All DOE employees, contractors, financial assistance recipients, and parties to other DOE agreements share the responsibility to protect the rights and welfare of human research subjects. The Secretary of Energy is responsible for oversight of the conduct of DOE-related human subject research.

a. **Under Secretary for Science and Energy.**

   (1) Monitors implementation of this Order, 10 CFR Part 745, and 45 CFR Part 46, within DOE in accordance with policy established by the Secretary and in consultation with the NNSA, as appropriate.

   (2) Determines what constitutes Departmental-related HSR, in consultation with the NNSA.

   (3) Ensures implementation of human research subject protection measures in accordance with the requirements of this Order, 10 CFR Part 745, and 45 CFR Part 46, in consultation with the NNSA.

   (4) Designates the DOE Institutional Official. For DOE, the Institutional Official is the Associate Director for Biological and Environmental Research, Office of Science.

   (5) Designates the DOE HSP Program Manager. For DOE, the HSP Program Manager resides within the Office of Science’s Office of Biological and Environmental Research.

   (6) Delegates review and approval of statements of work for HTM projects submitted by DOE’s non-NNSA sites to the DOE HSP Program Manager.
b. **Under Secretary for Nuclear Security and Administrator of the National Nuclear Security Administration** designates the NNSA HSP Program Manager and delegates review and approval of statements of work for HTM projects submitted by DOE’s NNSA Sites to the NNSA HSP Program Manager. The NNSA HSP Program manager resides within the Office of the Associate Administrator for Safety, Infrastructure and Operations.

c. **The DOE Institutional Official** is the Senior DOE Official, responsible for overseeing and monitoring Departmental implementation of the requirements of this Order, 10 CFR Part 745, 45 CFR Part 46, as well as Executive Orders, Presidential Memoranda, and other Presidential directives, as applicable, in consultation with the NNSA, as appropriate. The DOE Institutional Official is also responsible for: 1) ensuring the Central DOE Institutional Review Boards (IRBs) comply with applicable Federal and DOE regulations; 2) ensuring the OHRP Federalwide Assurance (FWA) and IRB registration are properly maintained and current; and 3) formally appointing all members of the Central IRBs. The Institutional Official must concur on all requests for partial or full exemptions from the requirements of this Order.

d. **DOE HSP Program Manager.**

(1) Develops procedures for the HSP program in consultation with the NNSA HSP Manager, as appropriate.

(2) Prepares and updates guidance to be followed for obtaining approval for HSR in consultation with the NNSA HSP Manager, as appropriate.

(3) Reviews/approves (or when an NNSA element is involved, reviews and may recommend approval of) local plans to correct any noncompliance or to mitigate adverse study events, ensuring they comply with applicable HSP requirements.

(4) Reviews and approves statements of work for HTM projects submitted by DOE’s non-NNSA sites. Ensures compliance with DOE requirements [see paragraph 4.a.(2)], and, for HTM projects that are Strategic Partnership Projects (SPPs) and Strategic Intelligence Partnership Program (SIPP) projects, coordinates with appropriate Headquarters SPP/SIPP leads prior to approving such statements of work for initiation. Ensures Site Offices and M&O contractors are aware of decisions concerning proposed HTM work.

(5) Provides advice and guidance on evolving Departmental and national bioethics and regulatory issues regarding human research subject protection and helps identify and resolve program/project concerns in consultation with the NNSA HSP Program Manager, as appropriate.
(6) Develops and conducts educational programs on bioethics and human research subjects protection requirements, practices, and procedures relevant to DOE employees, DOE contractor personnel, financial assistance recipients, and the public in consultation with the NNSA HSP Program Manager, as appropriate.

(7) Regularly conducts institutional performance reviews to assess compliance with human research subject protection requirements in consultation with the NNSA HSP Program Manager, as appropriate.

(8) Serves as the Chair of the DOE Human Subjects Working Group and as official DOE representative to groups with bioethics and HSP interests. The NNSA HSP Program Manager shall be invited to attend all such meetings and to co-chair meetings, as appropriate.

(9) Makes recommendations to the Secretary, after concurrence from, and through the Institutional Official, regarding requests for exemptions from the requirements of this Order and satisfies the advance notice and publication requirements of 10 CFR Part 745.101(i) prior to the granting of any exemption (in consultation with the NNSA HSP Program Manager, as appropriate).

(10) Concurs on HSP provisions in interagency agreements, in consultation with the NNSA HSP Program Manager, as appropriate.

(11) Maintains the HSR Projects Database for DOE.

e. NNSA HSP Program Manager.

(1) When an NNSA element is involved, reviews requests for exemptions to requirements of this Order and makes recommendations to the Secretary through the NNSA Administrator after concurrence from the Institutional Official. Ensures that the advance notice and publication requirements of 10 CFR Part 745.101(i) are met prior to the granting of any exemption. Also reviews and approves statements of work for HTM projects submitted by NNSA sites. Ensures compliance with DOE/NNSA requirements and, for SPP/SIPPs HTM projects, coordinates with the NNSA Office of Strategic Partnership Programs prior to approving such projects for initiation. Ensures Site Offices and M&O Contractors are aware of decisions concerning proposed HTM work.

(2) Works with the DOE HSP Program Manager, as outlined in paragraph 5.

f. Office of Intelligence and Counterintelligence.

(1) Reviews and approves, prior to initiation, statements of work for HTM projects received from members of the intelligence community.
(2) Reviews and approves statements of work for non-HTM, intelligence-related HSR prior to initiation.

(3) In these reviews, coordinates with the appropriate HSP Program Manager.

g. Secretarial Officers or their Designees.

(1) Ensure that all proposals for research, studies, tests, surveys, surveillance, or other data collection are reviewed to identify research involving human subjects.

(2) Ensure that any questions or uncertainties regarding the applicability of human research subjects protection requirements to such proposals, and any other issues and concerns regarding the requirements of this Order, are promptly referred to the appropriate HSP Program Manager for resolution.

(3) Ensure that the contracting officer is advised when work statements for proposed agreements include HSR to ensure that the CRD or its requirements (as appropriate) will be applied to HSR conducted with DOE funding, at DOE institutions, or by DOE personnel under agreements other than site/facility management contracts, such as support services contracts, grants, cooperative agreements, work-for-others agreements, and interagency agreements.

(4) Ensure that the contracting officer, after being notified of the affected contracts, incorporates the CRD into the affected contracts by way of the Department of Energy Acquisition Regulations (DEAR) Laws, regulations, and directives clauses included in those contracts. In the case of contracts or other agreements requiring contractor performance of activities covered by the CRD, but which do not contain the Laws, regulations, and DOE directives clause, the contracting officer will work to include the requirements as appropriate.

(5) Ensure their staffs and field elements comply with the requirements of this Order, including the notification requirements in paragraph 4d.

(6) Ensure relevant personnel actively participate in human research subjects protection training and educational programs.

(7) Ensure that self-assessments are periodically conducted to verify compliance with the requirements of this Order.

(8) At their discretion, conduct further review and approve or disapprove research that has been approved by the IRB. (Note: Secretarial Officers or their designees may not approve HSR that has not been approved by an IRB. See 10 CFR Part 745.112.)
(9) Ensure appropriate oversight of the administration of research subjects protection programs of contractors and financial assistance recipients under their cognizance, and other parties to DOE agreements, to ensure compliance with applicable human research subjects protection requirements.

(10) Ensure that the DOE HSP Program Manager and the NNSA HSP Manager are involved in negotiating those portions of interagency agreements that address HSR.

(11) Appoint a point of contact for interacting with the appropriate HSP Program Manager on program-related and/or Department-wide issues.

h. DOE Field/Site Offices.

(1) Ensure contracts and other agreements involving HSR require compliance with the requirements set forth in the CRD associated with this Order (Attachment 1), 10 CFR Part 745, and 45 CFR Part 46.

(2) Ensure that contractors establish and maintain a process for:

(a) Identifying and reporting HTM work according to the requirements of this Order;

(b) Notifying the HSP Program Manager(s) as required in paragraph 1 of this CRD; and

(c) Training relevant personnel in HSP requirements, as required by paragraph 10 of the CRD.

6. REFERENCES.

a. DOE O 206.1, Department of Energy Privacy Program, dated 1-16-09, which ensures compliance with privacy requirements; establishes a Departmental training and awareness program for all DOE Federal and contractor employees to ensure personnel are cognizant of their responsibilities for safeguarding Personally Identifiable Information (PII) and complying with the Privacy Act; and provides Departmental oversight to ensure compliance.

b. DOE P 481.1, DOE's Policy Regarding Laboratories, Plants and Sites Engaging in Strategic Partnership Projects with Other Federal Agencies, Independent Organizations, and the Private Sector, dated 12-17-14, which sets the context in which DOE and its laboratories, plants, and sites should pursue Strategic Partnership Projects (SPPs) with other Federal government agencies, state and local institutions, universities, foreign entities and/or private companies. The Policy is applicable to the DOE laboratories, plants, and sites, and to the DOE programs that own them and facilitate their work.
c. DOE O 481.1C Admin Chg 2, *Strategic Partnership Projects (Formerly Known as Work for Others (Non-Department of Energy Funded Work)),* dated 1-24-05, which establishes the policy, responsibilities, and procedures for authorizing and administering work for non-DOE entities by DOE/National Nuclear Security Administration (NNSA) and/or their respective contractor personnel or the use of DOE/NNSA facilities that is not directly funded by DOE appropriations.

d. DOE M 481.1-1A Chg 1, *Reimbursable Work for Non-Federal Sponsors Process Manual,* dated 1-03-01, provides detailed requirements to supplement DOE O 481.1C, which establishes requirements for the performance of work for non-DOE/non-NNSA entities by DOE/NNSA/contractor personnel and/or the use of DOE/NNSA facilities that is not directly funded by DOE/NNSA appropriations.

e. DOE O 483.1A, *DOE Cooperative Research and Development Agreements,* dated 11-6-13, which provides detailed requirements to supplement DOE O 483.1, DOE Cooperative Research and Development Agreements, dated 1-12-01, which establishes requirements for the performance of technology transfer through the use of Cooperative Research and Development Agreements (CRADAs). DOE O 484.1, *Reimbursable Work for the Department of Homeland Security,* dated 8-17-06. The Order establishes DOE policies and procedures for the acceptance, performance, and administration of reimbursable work directly funded by the Department of Homeland Security.

f. DOE O 484.1 Admin Chg 2, *Reimbursable Work for the Department of Homeland Security,* dated 8-17-06, establishes DOE policies and procedures for the acceptance, performance, and administration of reimbursable work directly funded by the Department of Homeland Security.

g. DOE O 475.2B, *Identifying Classified Information,* dated 10-3-14, establishes the program to identify information classified under the Atomic Energy Act [Restricted Data (RD), Formerly Restricted Data (FRD), and Transclassified Foreign Nuclear Information (TFNI)] or Executive Order (E.O.) 13526 [National Security Information (NSI)], so that it can be protected against unauthorized dissemination in accordance with legal and Departmental requirements.

h. DOE O 471.3, Identifying and Protecting Official Use Only Information, Admin Change 1, dated 1-13-11, which establishes a program within DOE and NNSA to identify certain unclassified controlled information as Official Use Only (OUO) (e.g., PII and proprietary information) and to identify, mark, and protect documents containing such information to ensure it is protected according to legal and Departmental requirements.

i. DOE M 471.3-1, Manual for Identifying and Protecting Official Use Only Information, Admin. Change 1, 1-13-11, which provides detailed requirements to supplement DOE O 471.3. Admin Chg 1 dated 1-13-11.
j. 10 CFR Part 600, Financial Assistance Rules, which provides the policies and procedures for administration and management of all DOE financial assistance activities.

k. 10 CFR Part 602, Epidemiology and Other Health Studies Financial Assistance Program, which sets forth the policies and procedures applicable to the award and administration of financial assistance agreements and cooperative agreements for health-related research, education/training, conferences, communication, and related activities.

l. 10 CFR Part 605, Office of Science (formerly Office of Energy Research) Financial Assistance Program, as explained at www.er.doe.gov/grants/605.asp, which provides policies and procedures for the administration and management of basic and applied research financial award agreements awarded by the Office of Science.

m. 10 CFR Part 745, Protection of Human Subjects, which set Federal requirements for DOE for the protection of human subjects involved in research activities.

n. 10 CFR Part 1008, Records Maintained on Individuals (Privacy Act) which establishes the procedures to implement the Privacy Act of 1974 (PL. 93-579, 5 U.S.C. 552a) within DOE of Energy.

o. 45 CFR Part 46, Protection of Human Subjects, Subparts B, C, and D, which sets out DOE prescribed DHHS requirements for protected classes of human research subjects and Subpart E for IRB registration.


q. The Freedom of Information Act, 5 USC Section 552, as amended, which establishes the right of citizens to request information from Federal agencies and establishes a framework of procedures to implement this right.

r. DOE Policy Memorandum on Research Involving Intentional Modification of the Human Environment, 4-25-13.

7. DEFINITIONS.

a. Appropriate HSP Program Manager. The DOE HSP Program Manager (and when an NNSA element is involved, the NNSA HSP Program Manager).

b. Assurance. The written documentation, satisfactory to the Secretary of Energy, required from the prospective performing institution, that ensures institutional compliance with and implementation of DOE and Department of Health and Human Services (DHHS) regulations for the protection of human research
subjects. The only documentation currently meeting this requirement is a Federalwide Assurance (FWA). See: http://ohrp.cit.nih.gov/efile/FwaStart.aspx.

c. **Adverse Event.** Any unfavorable medical occurrence in a human subject, including any abnormal sign (for example, abnormal physical exam or laboratory finding), symptom, or disease, temporally associated with the subjects participation in the research, whether or not considered related to the subject’s participation in the research. A significant adverse event is an adverse event that is unexpected and substantively impacts the human subjects.

d. **De-identified Data.** A data set that has no, or limited, identifiers and for which a person with current knowledge of generally accepted scientific principles determines that the risk that the information could be used, alone or in combination with other reasonably available information, by an anticipated recipient, to identify an individual who is a subject of the information, has been reduced to the extent practicable. A graded approach must be used in balancing de-identification of the datasets and the usability of the dataset to accomplish the needed research.

e. **DOE Human Subjects Protection Program Manager (DOE HSP Program Manager).** The individual designated by the Under Secretary for Science and Energy to oversee the non-NNSA components of DOE’s Human Subjects Protection Program.

f. **DOE Institutional Official.** The Senior DOE Official responsible for overseeing and monitoring Departmental implementation of the requirements of 45 CFR Part 46, 10 CFR 745, Protection of Human Subjects, and this Order, in consultation with NNSA, as appropriate.

g. **DOE HSR Projects Database (HSRD).** An unclassified compilation of summary information, which is available on the website at: https://www3.orau.gov/hsrdreport, updated annually, on every HSR project funded by DOE, conducted at DOE institutions or facilities, or performed with DOE or contractor personnel.

h. **Generalizable.** Information/research findings that can be applied to populations or situations beyond that studied.

i. **Human Subject.** A living individual about whom an investigator (whether professional or student) conducting research obtains

(1) Data through intervention or interaction with the individual, or

(2) Identifiable private information.

*Intervention* includes both physical procedures by which data are gathered (for example, venipuncture) and manipulations of the subject or the subjects’s
environment that are performed for research purposes. *Interaction* includes communication or interpersonal contact between investigator and subject. *Private information* includes information about behavior that occurs in a context in which an individual can reasonably expect that no observation or recording is taking place, and information which has been provided for specific purposes by an individual and which the individual can reasonably expect will not be made public (for example, a medical record).

Private information must be individually identifiable (i.e., the identity of the subject is or may readily be ascertained by the investigator or associated with the information) in order for obtaining the information to constitute research involving human subjects.

j. Human Subjects Research (HSR). Any activity meeting the definitions of both: 1) research; and 2) human subject, as defined in this section.

k. Human Terrain Mapping. Research and data gathering activities primarily conducted for military or intelligence purposes to understand the “human terrain,”—the social, ethnographic, cultural, and political elements of the people among whom the U.S. Armed Forces are operating and/or in countries prone to political instability. This work includes observations, questionnaires, and interviews of groups of individuals, as well as modeling and analysis of collected data, and may become the basis for U.S. military actions in such locations. In addition to Human Terrain Mapping (HTM), such activities are often referred to as human social culture behavior (HSCB) studies. It is DOE policy that HTM activities will be managed as HSR.

l. HTM Data. Data collected or used as part of HTM efforts, as described above, as well as any auxiliary data on the same group(s) of individuals.

m. NNSA Human Subjects Protection Designee (NNSA HSP Program Manager). The individual appointed by the NNSA Administrator to oversee the Human Subjects Protection Program for NNSA elements.

n. Institution. Any public or private entity or agency (including Federal, State, and other agencies). This term refers to laboratories and other facilities managed by DOE, DOE contractors, or DOE financial assistance recipients.

o. Institutional Review Board (IRB). A committee or board established by an institution that performs initial and continuing reviews of research involving human subjects, and is registered with the Office for Human Research Protections (OHRP) and designated on an FWA.

p. Personally Identifiable Information. Any information collected or maintained about an individual, including but not limited to, education, financial transactions, medical history and criminal or employment history, and
information that can be used to distinguish or trace an individual’s identity, such as his/her name, Social Security number, date and place of birth, mother’s maiden name, biometric data, and any other personal information that is linked or linkable to a specific individual.

q. Research. A systematic investigation, including research development, testing and evaluation, designed to contribute to generalizable knowledge. Activities which meet this definition constitute research for purposes of this policy, whether or not they are conducted or supported under a program which is considered research for other purposes. For example, some demonstration and service programs may include research activities.

r. Strategic Intelligence Partnership Program (SIPP): SIPP, formerly the Intelligence Work for Others (IWFO) program, is the mechanism by which DOE provides highly-specialized scientific and technical services and products to non-DOE Intelligence Community (IC) and other agencies for intelligence and intelligence-related activities carried out under unique IC authorities held by DOE’s Office of Intelligence and Counterintelligence and sponsoring IC agencies.

s. Strategic Partnership Projects (SPPs): SPP, formerly the Work for Others (WFO) program, is the mechanism by which non-DOE entities fund DOE/NNSA and/or their contractors or use of DOE/NNSA facilities for work that is not directly funded by DOE/NNSA appropriations.

t. Unanticipated Problem. In general, to be classified as an unanticipated problem, any incident, experience, or outcome should meet all three of the following criteria:

1. Unexpected (in terms of nature, severity, or frequency) given (a) the research procedures that are described in the protocol-related documents, such as the IRB-approved research protocol and informed consent document; and (b) the characteristics of the subject population being studied

2. Related or possibly related to participation in the research (possibly related means there is a reasonable possibility that the incident, experience, or outcome may have been caused by the procedures involved in the research)

3. Likely to place subjects or others at greater risk of harm (including physical, psychological, economic, or social harm) than was previously known or recognized.

8. CONTACT. Questions regarding this Order should be addressed to the DOE Program Manager, HSP Program, Office of Science, Office of Biological and Environmental Research, telephone 301-903-3213, or the NNSA HSP Program Manager, as
BY ORDER OF THE SECRETARY OF ENERGY:

DANIEL B. PONEMAN
Deputy Secretary
CONTRACTOR REQUIREMENTS DOCUMENT
DOE O 443.1B, PROTECTION OF HUMAN SUBJECTS

Regardless of the performer of the work, the contractor is responsible for compliance with the requirements of this Contractor Requirements Document (CRD).

The contractor is responsible for flowing down the requirements of this CRD to subcontracts at any tier to the extent necessary to ensure the contractor’s compliance with the requirements.

Note: Throughout this CRD, the term "Human Subjects Protection Program Manager (HSP Program Manager)" refers either to the DOE HSP Program Manager or to the NNSA HSP Program Manager except where otherwise noted.

As directed by the contracting officer, the contractor shall—

1. Ensure notification of the DOE HSP Program Manager (and, when an NNSA element is involved, the NNSA HSP Program Manager):
   a. Prior to initiation of any new HSR project, even if it meets the regulatory definition of exempt HSR as outlined in 10 CFR Part 745.101(b), involving:
      (1) an institution without an established Institutional Review Board (IRB);
      (2) a foreign country;
      (3) the potential for significant controversy (e.g., negative press or reaction from stakeholder or oversight groups);
      (4) research subjects in a protected class (fetuses, pregnant women, and in vitro fertilization; prisoners; or children); or
      (5) the generation or use of classified information.
   b. Within 48 hours of the following, and, provide a description of corrective actions taken immediately following the incident, as well as corrective actions to be taken for concurrence by the appropriate HSP Program Manager:
      (1) any significant adverse events, unanticipated problems, and complaints about the research,
      (2) any suspension or termination of IRB approval of research;
      (3) any significant non-compliance with HSP Program procedures or other requirements, which shall be reported to the IRB for evaluation for further action with the appropriate HSP Program Manager;
   c. Immediately, of a finding of a suspected or confirmed data breach involving PII in printed or electronic form and to the DOE-Cyber Incident Response Capability
immediately, in accordance with the requirements of the CRD associated with DOE O 206.1, and provide a description of any corrective actions taken within 48 hours and a description of corrective actions to be taken for concurrence by the appropriate HSP Program Manager.

d. Upon appointment of a new DOE Site IRB Chair or DOE Site Institutional Official.

2. Ensure that research involving human subjects, regardless of source of funding, is conducted in accordance with applicable requirements. (See also 10 CFR Part 745 and 45 CFR Part 46).¹

3. Ensure that contractor-issued solicitations or proposals for research, studies, tests, surveys, surveillance, or other data collection are reviewed to identify research involving human subjects and that any resulting agreements include the substance of the requirements in this CRD.

4. Ensure that no research involving human subjects, regardless of funding source, is initiated without prior IRB approval under the terms of an approved assurance covering the research.

5. Ensure that any Human Terrain Mapping (HTM) work complies with DOE requirements specified in this CRD, namely:

a. HTM projects, conducted with DOE funding, at DOE institutions (regardless of funding source), or by DOE contractor personnel (regardless of funding source or location of work conducted), whether done domestically or in an international environment, and including classified and proprietary research, shall be strictly limited to only those projects involving the analysis and modeling of de-identified data.

b. Documented process and procedures shall be developed to ensure that: 1) statements of work for HTM projects are submitted to the Site Office, for information, and to the appropriate HSP Program Manager (and when an NNSA element is involved, the NNSA HSP Program Manager), for DOE Headquarters review and approval, prior to initiation, and 2) relevant M&O personnel are trained in HTM requirements. If the project is to be conducted by or for the intelligence community, the Office of Intelligence must also review and approve it prior to project initiation. The HSP Program Manager(s) and the Office of Intelligence shall engage the recognized DOE site IRB and, if needed, the principal investigator (PI) and/or sponsor, in clarifying whether the proposed project is HTM, and if so, that the data to be used will be de-identified. Additionally, the PI will provide written verification that only de-identified data (as defined in paragraph 7d of this Order) will be used.

¹ Ensure that research is reviewed at intervals appropriate to the degree of risk, but not less than once per year, to assess the risk to test subjects and to assure the risk is reasonable in relation to anticipated benefits.
c. For Headquarters approved projects, the recognized DOE site IRB is the only entity authorized to determine whether the HTM data received by the PI after project initiation meets DOE criteria for de-identification. If the DOE site does not have an internal IRB, the Central DOE IRB will be the responsible IRB.

d. If, in the case the sponsor requests assistance in the de-identification of HTM data prior to start of any work on the sponsor’s project and/or re-identification of data following completion of the project, DOE sites may provide such services under a separate contract and/or task order with the sponsor by following the appropriate DOE standard operating procedure approved by the DOE Institutional Official, DOE Office of Science.

6. Submit an application for a Federalwide Assurance (FWA) to the Office of Human Research Protections (OHRP) with Department of Health and Human Services (DHHS) and, once approved by DHHS, maintain this FWA covering proposed and ongoing HSR and provide a copy to the appropriate HSP Program Manager. The Secretary of Energy uses the approved FWA as appropriate written documentation from DOE Sites committing to institutional compliance with and implementation of DOE and DHHS regulations for the protection of human research subjects. See http://ohrp.nih.gov/efile/FwaStart.aspx and/or contact the DOE HSP Program Manager, Office of Science, Office of Biological and Environmental Research, telephone 301-903-3213, or the NNSA HSP Program Manager, as appropriate.

7. Periodically conduct self-assessments to ensure compliance with the HSP Program procedures and other requirements.

8. Prepare and submit an annual report for the HSR Projects Database in accordance with directions and schedules provided by the appropriate HSP Program Manager.

9. Submit requests for exemptions from these requirements in writing through the contracting officer to the appropriate HSP Program Manager.

10. Ensure relevant personnel actively participate in HSP training and educational programs.

---

2 It should be noted that: a) only limited communications, if needed, may take place between the organization de-identifying and/or re-identifying the sponsor’s data and the organization performing work on the sponsor’s task; b) the identified data set shall not be shared with the individual who will perform work on the sponsor’s task; and c) the de-identified dataset shall be sent directly by the sponsor to the individual performing work on the sponsor’s task and not by the organization at the DOE site that de-identified it.