Human Subject Regulations Decision Charts

September 24, 2004

The Office for Human Research Protections (OHRP) provides the following graphic aids as a guide for institutional review boards (IRBs), investigators, and others who decide if an activity is research involving human subjects that must be reviewed by an IRB under the requirements of the U.S. Department of Health and Human Services (HHS) regulations at 45 CFR part 46. OHRP welcomes comment on these decision charts. The charts address decisions on the following:

- whether an activity is research that must be reviewed by an IRB
- whether the review may be performed by expedited procedures, and
- whether informed consent or its documentation may be waived.

Considerations

The charts are intended to assist IRBs, institutions, and investigators in their decision-making process and should not be used as substitutes for consulting the regulations. OHRP cautions that the full text of applicable regulatory provisions should be considered in making final decisions.

These charts are necessarily generalizations and may not be specific enough for particular situations. Other guidance documents are available related to specific topics, at OHRP Policy Guidance by Topic. OHRP invites inquiries for additional information.

The charts do not address requirements that may be imposed by other organizations, such as the Food and Drug Administration, National Institutes of Health, other sponsors, or state or local governments.

Chart 1: Is an Activity Research Involving Human Subjects?

Chart 2: Is the Human Subjects Research Eligible for Exemption?

Chart 3: Does Exemption 45 CFR 46.101(b)(1) (for Educational Settings) Apply?

Chart 4: Does exemption 45 CFR 46.101(b)(2) or (b)(3) (for Tests, Surveys, Interviews, Public Behavior Observation) Apply?

Chart 5: Does Exemption 45 CFR 46.101(b)(4) (for Existing Data, Documents, Records and Specimens) Apply?

Chart 6: Does Exemption 45 CFR 46.101(b)(5) (for Public Benefit or Service Programs) Apply?

Chart 7: Does Exemption 45 CFR 46.101(b)(6) (for Food Taste and Acceptance Studies) Apply?

Chart 8: May the IRB Review Be Done by Expedited Procedures?

Chart 9: May the IRB Continuing Review Be Done by Expedited Procedures?

Chart 10: May Informed Consent Be Waived or Consent Elements Be Altered under 45 CFR 46.116(d)?

Chart 11: May Documentation of Informed Consent Be Waived Under 45 CFR 46.117(c)?
Chart 1: Is an Activity Research Involving Human Subjects Covered by 45 CFR part 46?

Start here.

Is the activity a systematic investigation designed to develop or contribute to generalizable knowledge? [45 CFR 46.102(d)]

YES → Activity is research. Does the research involve obtaining information about living individuals? [45 CFR 46.102(f)]

NO → Activity is not research, so 45 CFR part 46 does not apply.

YES → The research is not research involving human subjects, and 45 CFR part 46 does not apply.

NO → Does the research involve intervention or interaction with the individuals? [45 CFR 46.102(f)(1), (2)]

NO → Is the information individually identifiable (i.e., the identity of the subject is or may readily be ascertained by the investigator or associated with the information)? [45 CFR 46.102(f)(2)]

NO → The research is not research involving human subjects, and 45 CFR part 46 does not apply.

YES → Activity is research involving human subjects. Is it conducted or supported by HHS? [45 CFR 46.101(a)(1)]

NO → Is the research covered by an applicable OHRP approved assurance created under 45 CFR 46.103?

YES → Unless exempt under 45 CFR 46.101(b), 45 CFR part 46, subpart A requirements apply to the research. As appropriate, subpart B, C, and D requirements also apply.

NO → Go to Chart 2

AND → Other Federal, State and local laws and/or regulations may apply to the activity. [45 CFR 46.101(f)]
Chart 2: Is the Research Involving Human Subjects Eligible for Exemption Under 45 CFR 46.101(b)?

Has HHS prohibited exemption of the human subjects research? (All research involving prisoners, some research involving children.) [Footnote 1 to 45 CFR 46.101(i), 45 CFR 46.401(b)]

**NO**

Will the *only* involvement of human subjects be in one or more of the following categories?

- Research conducted in *established* or *commonly accepted* educational settings, involving *normal education practices*?
  - **YES**
    - Exemption 45 CFR 46.101(b)(1) may apply. Go to Chart 3
  - **AND/OR**

- Research involving the use of *educational tests, survey procedures, interview procedures, or observation of public behavior*?
  - **YES**
    - Exemption 45 CFR 46.101(b)(2) or (b)(3) may apply. Go to Chart 4
  - **AND/OR**

- Research involving collection or study of *existing* data, documents, records, or pathological or diagnostic specimens?
  - **YES**
    - Exemption 45 CFR 46.101(b)(4) may apply. Go to Chart 5
  - **AND/OR**

- Research studying, evaluating, or examining *public benefit or service programs*?
  - **YES**
    - Exemption 45 CFR 46.101(b)(5) may apply. Go to Chart 6
  - **AND/OR**

- Research involving *taste and food quality evaluation or consumer acceptance studies*?
  - **YES**
    - Exemption 45 CFR 46.101(b)(6) may apply. Go to Chart 7
  - **NO**

No exemptions to 45 CFR part 46 apply. Provisions of 45 CFR subpart A apply, and subparts B, C and D also apply if subjects are from covered vulnerable populations.

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**"Only"** means that no non-exempt activities are involved. Research that includes exempt and non-exempt activities is *not* exempt.

If the research involves the use of audio, video, digital or image recordings of subjects, the research does not qualify as exempt according to the UND IRB policies and procedures.
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Chart 5: Does Exemption 45 CFR 46.101(b)(4) (for Existing Data Documents and Specimens) Apply?

From Chart 2

Does the research involve only the collection or study of existing data, documents, records, pathological specimens, or diagnostic specimens? *
("Existing" means existing before the research is proposed to an institutional official or the IRB to determine whether the research is exempt.)

YES

Are these sources publicly available?

YES

Research is exempt under 45 CFR 46.101(b)(4) from 45 CFR part 46 requirements.

NO

NO

Will information be recorded by the investigator in such a manner that the subjects cannot be identified, directly or through identifiers linked to the subjects?

YES

Research is not exempt under 45 CFR 46.101(b)(4) from 45 CFR part 46 requirements.

NO

Go to Chart 8

* Note: See OHRP guidance on research use of stored data or tissues and on stem cells at http://www.hhs.gov/ohrp/policy/index.html#tissues and #stem, and on coded data or specimens at #cocied for further information on those topics.

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If the research involves the use of audio, video, digital or image recordings of subjects, the research does not qualify as exempt according to the UND IRB policies and procedures.
Chart 6: Does Exemption 45 CFR 46.101(b)(5) (for Public Benefit or Service Programs) Apply?

From Chart 2

Is the research or demonstration project conducted or approved by the Department or Agency Head?

YES

Does the research or demonstration project involve only the study, evaluation, or examination of:

Public benefit or service programs;

YES

Procedures for obtaining benefits or services under public benefit or service programs;

YES

Research is exempt under 45 CFR 46.101(b)(5) from all 45 CFR part 46 requirements.*

NO

Possible changes in or alternatives to public benefit or service programs or to procedures for obtaining benefits or services under public benefit or service programs;

YES

Possible changes in methods or levels of payment for benefits or services under those public benefit or service programs?

YES

Research is not exempt under 45 CFR 46.101(b)(5).

NO

Go to Chart 8

NO

* Note: See OHRP guidance on exemptions at http://www.hhs.gov/ohrp/policyindex.html#exempt for further description of requirements for this exemption.

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If the research involves the use of audio, video, digital or image recordings of subjects, the research does not qualify as exempt according to the UND IRB policies and procedures.
Chart 7: Does Exemption 45 CFR 46.101(b)(6) (for Food Taste and Acceptance Studies) Apply?

From Chart 2

Does the research involve only a taste and food quality evaluation or a food consumer acceptance study?

YES

Are wholesome foods without additives consumed?

YES → Research is exempt under 45 CFR 46.101(b)(6) from all 45 CFR part 46 requirements.

NO

Is food consumed that contains a food ingredient, agricultural chemical, or environmental contaminant at or below the level found to be safe by the Food and Drug Administration or approved by the Environmental Protection Agency or the Food Safety and Inspection Service of the U.S. Department of Agriculture?

YES

NO → Research is not exempt under 45 CFR 46.101(b)(6).

Go to Chart 8

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If the research involves the use of audio, video, digital or image recordings of subjects, the research does not qualify as exempt according to the UND IRB policies and procedures.
Chart 8: May the IRB Review Be Done by Expedited Procedures Under 45 CFR 46.110?*

* Note: See expedited review categories and OHRP guidance on the use of expedited review procedures at [http://www.hhs.gov/ohrp/policy/index.htm#expedited](http://www.hhs.gov/ohrp/policy/index.htm#expedited) for further information on expedited review.

From Chart 2, 3, 4, 5, 6, or 7

Has the research been *previously reviewed* and approved by the IRB?

YES

Is the review a continuing review? [45 CFR 46.109(d)]

NO

Does the research present *no more than minimal risk* to human subjects? and does the research involve *only procedures included in categories 1 through 7* on the list of categories of research that may be reviewed through an expedited review procedure? [45 CFR 46.110(b)(1)]

YES

Is the research classified? [Paragraph (D) of Categories of Research That May Be Reviewed By an IRB through an Expedited Review Procedure.]

YES

Review by convened IRB is required.

NO

Are measures in place to make risks no more than minimal?

YES

Go to Chart 10

NO

Could identification of subjects put them at risk of criminal or civil liability, or be socially or economically damaging? [Paragraph (C) of Categories.]

YES

NO

Go to Chart 9

Research is eligible for IRB review through expedited procedures. Agency head may restrict, suspend, terminate or choose not to authorize an institution’s or IRB’s use of the expedited review procedure. [45 CFR 46.110(d)]

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Chart 9: Can Continuing Review be Done by Expedited Procedures Under 45 CFR 46.110?

From Chart 8

Has the research been previously reviewed and approved by the IRB using expedited procedures?

- **YES**
  - Have conditions changed such that the research is no longer eligible for expedited review (e.g., protocol change, or experience shows research to be of greater than minimal risk)?
    - **YES**
      - Review by convened IRB is required.
    - **NO**
      - Go to Chart 10

- **NO**
  - Have conditions changed to make the research eligible for expedited review under the applicability criteria and categories 1 through 7 on the list of categories that may be reviewed by expedited procedures (e.g., research is within those categories and experience confirms research to be of no greater than minimal risk)?
    - **YES**
      - Research is eligible for IRB review through expedited procedures.
    - **NO**
      - Category 8
        - (a) For this site: Is the research permanently closed to enrollment of new subjects?
          - **AND**
            - Have all subjects completed all research-related interventions?
              - **AND**
                - Does the research at this site remain active only for long-term follow-up of subjects?
                  - **YES**
                    - Category 9
                      - (c) Are the remaining research activities at this site limited to data analysis?
                        - **YES**
                          - Is the research conducted under an IND or IDE?
                            - **YES**
                              - NO
                        - **NO**
                          - NO
                    - (b) Have no subjects been enrolled at this site?
                      - **AND**
                        - Have no additional risks been identified anywhere?
                          - **YES**
                            - NO
                          - **NO**
                            - NO
                - **NO**
                  - NO
          - **NO**
            - NO
        - **NO**
          - NO
    - **NO**
      - NO

*Note: See expedited review categories, OHRP guidance on the use of expedited review procedures and on continuing review at [http://www.hhs.gov/ohrp/policy/index.html#expedited](http://www.hhs.gov/ohrp/policy/index.html#expedited) and [http://www.hhs.gov/ohrp/policy/index.html#continuing](http://www.hhs.gov/ohrp/policy/index.html#continuing) for further information on expedited review.
Chart 10: Can Informed Consent Be Waived or Consent Elements Be Altered Under 45 CFR 46.116(c) or (d)??

**(Note: If subjects include children to whom 45 CFR part 46, subpart D applies, an alternative provision for waiver of parental permission might apply. [See 45 CFR 46.408(c)])

From Chart 8 or 9

Will the research or demonstration project be **conducted by or subject to** the approval of **state or local government officials**? [45 CFR 46.116(c)(1)]

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NO
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Will the research involve greater than minimal risk, as defined in Section 46.102(i)? [45 CFR 46.116(d)(1)]

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NO
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Is it **practicable** to conduct the research without the waiver or alteration? [45 CFR 46.116(d)(3)]

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NO
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Will waiving or altering the informed consent adversely affect the subjects’ rights and welfare? [45 CFR 46.116(d)(2)]

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NO
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Will pertinent information be **provided** to subjects later, if appropriate? [45 CFR 46.116(d)(4)]

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YES
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Waiver of informed consent or alteration of consent elements is allowed if IRB documents these findings and approves waiver or alteration.

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NO
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Is it **practicable** to conduct the research without the waiver or alteration? [45 CFR 46.116(c)(2)]

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YES
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No waiver of informed consent or alteration of consent elements is allowed.*

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NO
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Is the project designed to study, evaluate, or otherwise examine: (i) Public benefit of service programs, (ii) procedures for obtaining benefits or services under those programs; (iii) possible changes in or alternatives to those programs or procedures; or (iv) possible changes in methods or levels of payment for benefits or services under those programs? [45 CFR 46.116(c)(1)]

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NO
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Go to Chart 11

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NO
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If informed consent is not waived entirely

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NO
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* Note: See OHRP guidance on informed consent requirements in emergency research at http://www.hhs.gov/ohrp/policy/index.html#emergency for further information on emergency research informed consent waiver.

This flowchart does not pertain to research that is FDA regulated. See SOP 702 and SOP 703 for consent requirements for FDA regulated research.
Chart 11: Can Documentation of Informed Consent Be Waived Under 45 CFR 46.117(c)?

From Chart 10

Would the consent document be the only record linking the subject and the research and would the principal risk be potential harm resulting from a breach of confidentiality? [45 CFR 46.117(c)(1)]

NO

Does the research present no more than minimal risk and involve no procedures for which written consent is normally required outside the research context? [45 CFR 46.117(c)(2)]

YES

IRB may waive the requirement for a signed consent form for some or all subjects.

NO

IRB may require investigator to provide subjects with a written statement regarding the research. [45 CFR 46.117(c)]

AND

If IRB Allows Waiver of Documentation Under 45 CFR 46.117(c)(1)

Investigator will ask each subject if he or she wants documentation linking the subject with the research. [45 CFR 46.117(c)(1)]

Subject’s wishes will govern whether informed consent is documented. [45 CFR 46.117(c)(1)]

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