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These materials, such as flyers, advertisements, and letters, are an important part of a research study, and are considered to be the beginning of the informed consent process. While neither the Office of Human

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Research Protection
(OHRP) nor the Food
and Drug
Administration (FDA)
address subject
recruitment in their
regulations, both require
that an IRB review
methods and materials
used to recruit study
subjects.

*Subject Recruitment
Materials:*

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Understanding the ...

Materials used in the recruitment of potential study participants such as advertisements, flyers, and letters are an important and integral part of a research study. They are often the only way for a potential subject to learn about a new clinical trial.

Subject Recruitment

Page 13/36

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Materials

Understanding The ...

Subject Recruitment Materials Understanding

The Oftentimes, the only way for potential research subjects to learn about a new clinical trial is through recruitment materials. These materials, such as flyers, advertisements, and letters, are an important part of a

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research study, and are considered to be the beginning of the informed consent process.

*Subject Recruitment
Materials*

*Understanding The
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The consider the text of
the recruitment material
and its placement. These*

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tools are considered to be part of the subject selection and the informed consent process for the study.

The type of recruiting tool should be specified (e.g., posted flier, website notice, newspaper ad, radio, newsletter from support groups, social

Subject Recruitment

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*Understanding The
Advertisements and
recruitment materials
for human research
subjects (posters, flyers,
newspaper/magazine
ads, scripts for
radio/TV, electronic
mail, or solicitations
from outside sources)*

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are considered an extension of the informed consent and subject selection processes.

Subject Selection, Recruitment And Compensation

Advertisements and recruitment materials for human research subjects (posters, flyers, newspaper/magazine

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ads, scripts for radio/TV, electronic mail, or solicitations from outside sources) are considered an extension of the informed consent and subject selection processes.

*Subject Selection,
Recruitment, and
Compensation*
Recruiting Study

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Subjects - Information Sheet. FDA requires that an Institutional Review Board (IRB) review and have authority to approve, require modifications in, or disapprove all research ...

Recruiting Study

Subjects | FDA

The JHM IRB will consider the text of the

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recruitment material and its placement. These tools are considered to be part of the subject selection and the informed consent process for the study. The type of recruiting tool should be specified (e.g., posted flier, website notice, newspaper ad, radio, newsletter from support groups, social

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(networking site ad, etc.)

*JHM - Recruiting Study
Subjects*

- b. Reviewing subject recruitment materials and strategies.
- c. Assuring that all applicable institutional policies and federal regulations related to research with human subjects are followed.
- d. Reviewing manuscripts

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prior to submission for
publication. e.

Protecting the rights and
welfare of human
subjects.

*CITI Online Program
Flashcards | Quizlet*

All recruitment
materials must have
tracking information
such as an item number,
version number or, at a
minimum, a date. It

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should state the purpose of the study. It should be written in simple language, comparable in reading level to the consent document. It must be quite clear that it is for research or for an investigation.

Recruitment Materials / Advertising

Advertisements, flyers,
information sheets,

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notices, internet postings and/or media are used to recruit subjects. The text of these needs to be included within or as an attachment to.

Recruitment Methods & Materials Page 2 of 8
AAHRPP Elements
II.3.C, II.3.D, II.3.D,
II.4.A, III.1.E. the
webIRB application.

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*Recruitment Methods
and Tools - Research
Administration*

The Subject

Identification and
Recruitment section of
the initial review
application asks study
teams to provide
information about the
methods used to identify
and recruit subjects, and
the overall recruitment
plan. When does the

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Subject Identification
and Recruitment section
display? The Subject
Identification and

Recruitment section is
the eighth section of the
initial review
application and will
automatically display
for most application
types when study teams
intend to interact or ...

Completing the Subject
Page 31/36

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Identification and Recruitment Section

Recruitment to primary care-based studies may occur within the consultation or in dedicated research clinics. For practical and logistic reasons, some ...

Understanding the origins of the ethical safeguards surrounding research using human subjects may, however,

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allow primary care
researchers to challenge
the appropriateness of
these safeguards ...

The

*Ethical issues regarding
recruitment to research
studies ...*

To assist the subject in
understanding the
material, the eIC may
use interactive
electronic-based
technology, which may

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include diagrams,
images, graphics,
videos, and narration.

The eIC should be
appropriate for the
intended audience,
taking into
consideration the
subject's age, language,
and comprehension
level.

*Use of Electronic
Informed Consent:*
Page 34/36

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Questions and Answers

...
Materials

Recruitment of
Research Participants

This guidance document defines the standards and parameters for utilization of advertisements and other recruitment materials in research studies. The main purpose of such materials is to inform

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and invite potential
subjects to participate in
a research study.

Understanding The

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