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**Boston  
Children's  
Hospital**

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Applicability Boston Children's  
Hospital- Policies  
& Procedures

## Review of Research Involving Vulnerable Populations Not Covered by HHS Subparts B, C, and D Policy/Procedure

### Internal Approval

SVP, Research

EVP & Chief Scientific Officer, Research

### Scope

This policy will address and provide guidance concerning vulnerable populations not covered by U.S. Department of Health & Human Services (HHS) 45 CFR 46 subparts B, C, and D.

### Policy Statements

Boston Children's Hospital applies U.S. Department of Health & Human Services (HHS) 45 CFR 46 subparts B, C, and D to federally funded research and research under the jurisdiction of the U.S. Food & Drug Administration. Institutional Review Board (IRB) policies for these populations are covered separately, see:

- Pregnant Women, Human Fetuses, and Neonates
- Incarcerated Youth and Prisoners
- Children
- Children in State Custody

Adults with questionable decision-making capabilities are addressed in a separate IRB policy: Research Involving Individuals with Decisional Impairment.

Other groups such as the socially, educationally, and economically disadvantaged, elderly, and terminally

ill do not have specific regulatory protections. However, when research involves one of these other groups the IRB will evaluate the need for additional protections on a protocol-by-protocol basis.

When the Boston Children's Hospital IRB reviews research involving a vulnerable category of subjects, it will include one or more individuals qualified to represent that group, either through personal experience or experience working with the populations. In addition, the IRB will ascertain that additional safeguards to protect the rights and welfare of any vulnerable subjects will be included in the research.

## Procedures

For research involving participants who are vulnerable to coercion and/or undue influence and who can give consent, the IRB will apply the following additional criteria:

1. The inclusion of the vulnerable population is acceptable because either:
  - a. The inclusion of the vulnerable population is likely because of the setting of the research, and the setting is not designed to target vulnerable participants; or
  - b. The research is designed for a disease or condition relevant to the vulnerable population under study.
2. The research does not target vulnerable participants as a matter of convenience.
3. The recruitment process includes additional safeguards to minimize coercion and undue influence.
4. The IRB will consider the nature of the risks, the type of vulnerability, and the nature and level of anticipated benefit in addition to the availability of alternatives.
5. The consent process includes additional safeguards to minimize coercion and undue influence.
6. The financial payment (if any) to participants is not coercive or unduly influential.

## Related Content

- Department of Health and Human Services Regulations (HHS 45 CFR 46)
  - Subpart B, C, D
- IRB Policies
  - Pregnant Women, Human Fetuses, and Neonates
  - Incarcerated Youth and Prisoners
  - Children
  - Children in State Custody
  - Research Involving Individuals with Decisional Impairment.

## Approval Signatures

Step Description

Approver

Date

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## Applicability

Boston Children's Hospital- Policies & Procedures

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