

## Submit Pre-Review



### 1. Regulatory oversight: (check all that apply)

- DOD (Department of Defense)
- DOE (Department of Energy)
- DOJ (Department of Justice)
- ED (Department of Education)
- EPA (Environmental Protection Agency)
- FDA (Food and Drug Administration)
- HHS (Department of Health and Human Services)
- ICH GCP (International Center for Harmonization of Good Clinical Practice)
- VA (Department of Veterans Affairs)
- Other Federal Agency

### 2. Special determinations and waivers: (check all that apply)

Determination	Related Checklist
<input type="checkbox"/> Children	<a href="#">HRP-416 - Checklist - Research Involving Children</a>
<input type="checkbox"/> Children who are wards of the state	<a href="#">HRP-416 - Checklist - Research Involving Children</a>
<input type="checkbox"/> Cognitively impaired adults	<a href="#">HRP-417 - Checklist - Research Involving Cognitively Impaired Adults</a>
<input type="checkbox"/> Neonates of uncertain viability	<a href="#">HRP-414 - Checklist - Research Involving Neonates of Uncertain Viability</a>
<input type="checkbox"/> Nonsignificant risk device	<a href="#">HRP-418 - Checklist - Non-Significant Risk Device</a>
<input type="checkbox"/> Non-viable neonates	<a href="#">HRP-413 - Checklist - Research Involving Non-Viable Neonates</a>
<input type="checkbox"/> Pregnant women	<a href="#">HRP-412 - Checklist - Research Involving Pregnant Women</a>
<input type="checkbox"/> Prisoners	<a href="#">HRP-415 - Checklist - Research Involving Prisoners</a>
<input type="checkbox"/> Students / Employees	
<input type="checkbox"/> Waiver of consent documentation	<a href="#">HRP-411 - Checklist - Waiver of Written Documentation of the Consent Process</a>
<input type="checkbox"/> Waiver of consent for emergency research	<a href="#">HRP-419 - Checklist - Waiver of the Consent Process for Emergency Research</a>
<input type="checkbox"/> Waiver of HIPAA authorization	<a href="#">HRP-441 - Checklist - HIPAA Waiver of Authorization</a>
<input type="checkbox"/> Waiver/alteration of the consent process	<a href="#">HRP-410 - Checklist - Waiver or Alteration of Consent Process</a>

### 3. \* Risk level:

- Greater than minimal risk
  - No greater than minimal risk
- Clear

### 4. \* Type of research: (check all that apply)

- Biomedical
- Social Behavioral

**5. Incomplete Information and/or Missing materials:**

**6. Review Considerations:** 

**7. Add supporting documents:**

Name

There are no items to display

**8. \* Are you ready to submit this pre-review?** 

Yes  No [Clear](#)

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