Submit Designated Review

1. * I do NOT have a conflicting interest: ☐

2. * Determination:

<table>
<thead>
<tr>
<th>Name</th>
<th>Related Worksheet</th>
</tr>
</thead>
<tbody>
<tr>
<td>Approved</td>
<td>HRP-314 - Worksheet - Criteria for Approval and Additional Considerations</td>
</tr>
<tr>
<td>Modifications Required to Secure &quot;Approved&quot;</td>
<td>HRP-314 - Worksheet - Criteria for Approval and Additional Considerations</td>
</tr>
<tr>
<td>Not Human Research</td>
<td>HRP-310 - Worksheet - Human Research Determination</td>
</tr>
<tr>
<td>Modifications Required to Secure &quot;Not Human Research&quot;</td>
<td>HRP-310 - Worksheet - Human Research Determination</td>
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<tr>
<td>Human Research, Not Engaged</td>
<td>HRP-311 - Worksheet - Engagement Determination</td>
</tr>
<tr>
<td>Modifications Required to Secure &quot;Human Research, Not Engaged&quot;</td>
<td>HRP-311 - Worksheet - Engagement Determination</td>
</tr>
</tbody>
</table>

Clear

3. Review level: (select one if "Approved" or "Modifications Required to Secure 'Approved'"

<table>
<thead>
<tr>
<th>Name</th>
<th>Related Worksheet</th>
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<tbody>
<tr>
<td>Exempt</td>
<td>HRP-312 - Worksheet - Exemption Determination</td>
</tr>
<tr>
<td>Expedited</td>
<td>HRP-313 - Worksheet - Eligibility for Review Using the Expedited Procedure</td>
</tr>
</tbody>
</table>

Clear

If exempt, indicate the categories (see HRP-312 for full regulatory criteria):
(check all that apply)

- [ ] (1) Educational settings
- [ ] (2) Tests, surveys, interviews, or observation
- [ ] (3) Public officials or protected information
- [ ] (4) Data, documents, or specimens
- [ ] (5) Demonstration projects
- [ ] (6) Taste and food quality
- [ ] Other

If expedited, indicate the categories (see HRP-313 for full regulatory criteria):
(check all that apply)

- [ ] (1)(a) Drug studies
- [ ] (1)(b) Device studies
- [ ] (2)(a) Blood samples from healthy, non-pregnant adults
- [ ] (2)(b) Blood samples from others
- [ ] (3) Noninvasive biological specimens
- [ ] (4) Noninvasive procedures
- [ ] (5) Data, documents, records, or specimens
(6) Voice, video, digital, or image recordings
(7) Behavioral research/social science methods
(8)(a) Long-term follow-up
(8)(b) No subjects enrolled
(8)(c) Data analysis
(9) Convened IRB determined minimal risk
Other

4. Dates:

Last day of approval period (Use 11/5/2015 if 12-month approval period):

5. If modifications are required, enter them below:

6. Notes:

7. Supporting documents: (attach any relevant checklists completed as part of the review)

Add

Name
There are no items to display

8. * Are you ready to submit this review?

   Yes  No  Clear