

**New Jersey Department of Health
Office of Commissioner
INSTITUTIONAL REVIEW BOARD**

APPLICATION TO MODIFY HUMAN SUBJECTS RESEARCH

IRB ID Number	Project Title	
Principal Investigator	Signature	Date Submitted
<p>Type of Modification</p> <p><input type="checkbox"/> Design</p> <p><input type="checkbox"/> Procedure</p> <p><input type="checkbox"/> Methodology</p> <p><input type="checkbox"/> Instrument Type: <input type="checkbox"/> Add <input type="checkbox"/> Delete <input type="checkbox"/> Modify</p> <p><input type="checkbox"/> Study Sites <input type="checkbox"/> Add <input type="checkbox"/> Delete</p>	<p><input type="checkbox"/> Recruitment Materials Type: <input type="checkbox"/> Add <input type="checkbox"/> Delete <input type="checkbox"/> Modify</p> <p><input type="checkbox"/> Personnel <input type="checkbox"/> Add <input type="checkbox"/> Delete</p> <p><input type="checkbox"/> Funding Agency <input type="checkbox"/> Add <input type="checkbox"/> Delete</p> <p><input type="checkbox"/> Consent <input type="checkbox"/> Document <input type="checkbox"/> Process <input type="checkbox"/> Add Translation</p>	<p><input type="checkbox"/> Research Subjects <input type="checkbox"/> Increase Number <input type="checkbox"/> Decrease Number <input type="checkbox"/> Change Inclusion Criteria <input type="checkbox"/> Change Exclusion Criteria <input type="checkbox"/> Add Source <input type="checkbox"/> Delete Source <input type="checkbox"/> Add Vulnerable Population</p> <p><input type="checkbox"/> Other</p>
1. Provide a detailed description of the requested modification.		
2. Provide a justification for the requested modification.		
3. Provide a detailed explanation of how the requested modification a) creates a new risk, if any, or b) affects a previously identified risk. Include a detailed description of the risk (e.g., likelihood, magnitude, duration, etc).		
4. Provide a detailed explanation of how the requested modification a) creates a new benefit, if any, or b) affects a previously identified benefit. Include a detailed description of the benefit (e.g., likelihood, magnitude, duration, etc).		
5. Provide a detailed explanation of how the requested modification will affect currently enrolled research subjects.		
6. Provide a detailed explanation of how current research subjects will be re-consented.		