

ELEMENTS OF INFORMED CONSENT

Checkboxes to be completed by reviewers

<u>Elements of Informed Consent:</u>	<u>Yes</u>	<u>No</u>	<u>N/A</u>
(1) Purpose of Research:			
A statement that the study involves research	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
An explanation of the purposes of the research	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
Expected duration of the subject's participation	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
A description of the procedures to be followed	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
Probability of random assignment to each treatment	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
Identification of any procedures that are experimental	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
(2) Risks and Discomforts:			
A description of any reasonably foreseeable risks or discomforts to the subject	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
(3) Benefits:			
A description of any benefits to the subject or to others, which may reasonably be expected from the research	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
(4) Alternatives:			
A disclosure of appropriate alternative procedures or courses of treatment, if any, that might be advantageous to the subject	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
(5) Confidentiality:			
A statement describing the extent, if any, to which confidentiality of records identifying the subject will be maintained and, if relevant, that other agencies might inspect the records	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
(6) Compensation for Injury:			
For research involving more than minimal risk, an explanation as to whether any compensation and an explanation as to whether any medical treatments are available if injury occurs and, if so, what they consist of, or where further information may be obtained	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
(7) Research Questions:			
An explanation of whom to contact for answers to pertinent questions about the research and research subjects' rights, and whom to contact in the event of a research-related injury to the subject	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
(8) Voluntary Participation:			
A statement that participation is voluntary, that refusal to participate will involve no penalty or loss of benefits to which the subject is otherwise entitled, and that the subject may discontinue participation at any time without penalty or loss of benefits to which the subject is otherwise entitled	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Additional Elements of Informed Consent (When Appropriate):

- (1) A statement that the particular treatment or procedure may involve risks to the subject (or to the embryo or fetus, if the subject is or may become pregnant) which are currently unforeseeable.
 - (2) Anticipated circumstances under which the subject's participation may be terminated by the investigator without regard to the subject's consent.
 - (3) Any additional costs to the subject that may result from participation in the research.
 - (4) The consequences of a subject's decision to withdraw from the research and procedures for orderly termination of participation by the subject.
 - (5) A statement that significant new findings developed during the course of the research which may relate to the subject's willingness to continue participation will be provided to the subject.
 - (6) The approximate number of subjects involved in the study.
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