

**SAN JUAN DE DIOS EDUCATIONAL FOUNDATION, INC.  
INSTITUTIONAL REVIEW BOARD**

**PROTOCOL ASSESSMENT FORM (SJDEFIIRB FORM 2.3)**

<b>PROTOCOL NUMBER:</b>	<b>IRB REFERENCE NUMBER:</b>				
<b>PROTOCOL TITLE:</b>					
<b>PRINCIPAL INVESTIGATOR:</b>					
<i>To the Principal Investigator: Please indicate the page number of the relevant items in this section applicable to your protocol.</i>	<i>To the Reviewer: Please assess the appropriateness of the contents of the various sections, as outlined in this assessment form, and propose revisions as deemed necessary. You may put your comments in the space provided, or alternatively, an electronic version of this form is available upon request, to facilitate encoding of comments.</i>				
<b>A. The protocol contains the following:</b>	<b>Pages</b>	<b>YES</b>	<b>NO</b>	<b>NA</b>	<b>Comments</b>
<b>1. Background of Study</b>		[ ]	[ ]	[ ]	
<b>2. Significance of Study</b>		[ ]	[ ]	[ ]	
<b>3. Rationale of Study</b>		[ ]	[ ]	[ ]	
<b>4. Literature Review</b>		[ ]	[ ]	[ ]	
a. Results of animal/human studies		[ ]	[ ]	[ ]	
b. Known risks of procedures		[ ]	[ ]	[ ]	
c. Known benefits of procedures		[ ]	[ ]	[ ]	
d. Known adverse effects of drugs/procedures		[ ]	[ ]	[ ]	
<b>5. Objectives of Study</b>		[ ]	[ ]	[ ]	
a. Primary objective		[ ]	[ ]	[ ]	
b. Secondary objectives		[ ]	[ ]	[ ]	
<b>6. Statement of risks of the project</b>		[ ]	[ ]	[ ]	
a. To study participants		[ ]	[ ]	[ ]	
b. To community		[ ]	[ ]	[ ]	
<b>7. Possible adverse events (AE)</b>		[ ]	[ ]	[ ]	
<b>8. Statement of benefits of the project</b>		[ ]	[ ]	[ ]	
a. To study participants		[ ]	[ ]	[ ]	
b. To community		[ ]	[ ]	[ ]	
<b>9. Recruitment of participants</b>		[ ]	[ ]	[ ]	
a. Recruitment of procedures		[ ]	[ ]	[ ]	
b. Inclusion/Exclusion Criteria		[ ]	[ ]	[ ]	
<b>10. Methods</b>		[ ]	[ ]	[ ]	
a. Type of study design		[ ]	[ ]	[ ]	
b. Setting for project		[ ]	[ ]	[ ]	
c. Duration of project		[ ]	[ ]	[ ]	
d. Procedures to be done		[ ]	[ ]	[ ]	
<b>11. Informed Consent</b>		[ ]	[ ]	[ ]	

	Pages	Yes	No	NA	Comments
<b>12. Total Site Budget</b>		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
<b>13. Curriculum Vitae of Investigators</b>					
a. Complete name, titles, institutional affiliations of Principal Investigator		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
b. Names of co-workers		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
c. Job description of co-workers		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
d. Responsibilities of each co-worker		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
e. Contract with sponsor		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
<b>14. Project Sponsors</b>					
a. Complete Name		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
b. Address		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
c. Name of contact person/s		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
d. Telephone number of contact person/s		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
e. Statement of sponsor's interest/co-authorship		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
<b>15. Elements of Informed Consent</b>					
15.1 Conflict of Interest					
Full disclosure of potential sources of conflict of interest involving any of the authors or the granting agency		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
15.2 Privacy and confidentiality of Health Information					
a. Full disclosure of publication rights		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
b. Amount and method of reimbursement of trial related expenses of the study participants		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
15.3 Vulnerable subjects involved in the study					
a. A description of who may solicit consent, how, and when it will be done		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
b. A description of who may give consent (involving minors and not legally competent to give consent)		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
15.4 Risks					
a. Provision for management of adverse reactions		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
b. Interim analysis and provisional or mandatory cessation guidelines in case harmful effects are demonstrated during the study		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
c. Non-material compensation to participant such as health education or other creative benefits, where no clear, direct benefit from the project will be received by the participant		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	

	Pages	Yes	No	NA	Comments
d. Guarantee of medical care/ indemnification of study participants in case of trial-related injuries, which shall not be subject to previous waiver		[ ]	[ ]	[ ]	
e. Level of risk					
i. Low		[ ]	[ ]	[ ]	
ii. Medium		[ ]	[ ]	[ ]	
iii. High		[ ]	[ ]	[ ]	

Please use this space for additional explanation/comments like use of contraception in SJDEFI which is a Catholic institution, other socially-sensitive issues, and funding sources (should not be tobacco industry-related)

**B. COMMENTS:**

Approval

Modifications required prior to approval

Minor Modifications

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Major Modifications

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Disapproval

Reason/s for disapproval:

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\_\_\_\_\_

\_\_\_\_\_  
Signature above printed name

\_\_\_\_\_  
Date

**C. FINAL ACTION:**

APPROVED

DISAPPROVED

\_\_\_\_\_  
Signature above printed name

\_\_\_\_\_  
Date