**Franciscan Children’s**

## RESEARCH APPLICATION

## FOR IRB REVIEW

**Study Title (including any nicknames): \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_**

**Principal Investigator(s) (Name, FC department, and Contact Information):**

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**Co-investigator(s) (Name and FC department):**

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**Department Director of PI (Name and Signature Required):**

**Departmental Director of Co-Investigator(s) (Name and Signature(s) Required): \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_**

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Date of Submission:

This form is to be used when submitting a research application to the Institutional Review Board at FC. *Incomplete applications will not be reviewed*. Please complete the outline below to describe your proposed study. Please address each section using additional space as needed.

Applications should be submitted to the IRB Administrator, [SaiSankeerth.Madakasira@franciscanchildrens.org](mailto:SaiSankeerth.Madakasira@franciscanchildrens.org)

1. Study Summary (please include the purpose of the study; study hypotheses; a brief description of the participants, study design, and methods; anticipated results; and plans for disseminating the findings)

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**2. Study Benefits (**State benefits to the research participant, the medical/scientific community, and to the general public’s health and welfare).

**3. Study Risks (**List the possibilities for harm to the participants as a result of their participation in the research, including discomforts, hazards, or inconveniences and indicate what measures will be taken to prevent or to minimize these risks).

**4. Confidentiality** (State what steps will be taken to maintain confidentiality of data anonymity/privacy of participants).

**5. Contact with Participants**

a. Does this research involve any non-affiliated person (someone who is not an employee or staff member of FC) who will recruit, treat, observe, provide information to participants, or otherwise participate in this research? ( )no ( ) yes (If “yes”, please complete Appendix A)

b. Who will supervise their conduct as it pertains to the research participants?

c. Describe their involvement with the participants/research study.

d. Address the following issues with respect to the professional qualifications of each non-affiliated person involved in the research study. (May attach CV) *CITI training certificate must accompany application*

1. Degrees achieved (identify type and institutions)

2. Licenses held (all states)

3. Hospital/health care organization affiliations

**6. Study Participants**

a. Will the study involve children as research participants? ( ) yes ( ) no

b. Will the participant be legally competent to consent or refuse to participate?

( ) yes ( ) no (If “no” who will consent?): \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

c. Will all participants be English speaking?

( )yes ( ) no (If no, what provisions have been made for non-English speaking participants?

d. Will participants be hospitalized for this study? ( ) yes ( ) no

1. If hospitalized for medical reasons, will participation in this study increase the length of stay? ( ) yes ( ) no

2. If outpatients, will extra visits be required for this study over and above those required for the participant’s clinical care? ( ) yes ( ) no

1. If the answer to (1) and (2) above is **yes**, state the anticipated source of payment for hospitalization, visits, and tests required solely for purposes of this study:

e. Will participants incur charges, specifically as a consequence of participation?

( )yes (if “yes”, state specifically what they will be charged for) ( )no

f. Will there be payment to participants?

( )yes (state terms of payment, including amount, who will receive it, what form it will take, formula for pro-rating, if any, and when it will be paid) ( )no

g. Outline methods for recruitment including identifying and contacting potential participants. Also, state follow-up contact procedures, if any, to be used. (Please attach samples of any letters, advertisements, etc.)

h. Outline the process by which the participant’s consent will be sought. (State the length of time between the conclusion of the consent process and the beginning of the participant’s participation, and who will conduct the consent discussion.)

i. State specific criteria for participant eligibility for this study (e.g. “normal kidney function” is not an acceptable inclusion criterion but instead should be listed as, “Creatinine X times the upper limit of hospital normal”).

j. State specific criteria for participant exclusion for this study.

**7. Study Components**

a.Controlled Substances in Schedule II ( ) yes ( ) no

b. Investigational Drug(s) ( ) yes ( ) no

(if **yes, indicate** IND# and submit copy of investigator’s brochure) **IND#\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_**

c. FDA Approved Drug Used in Accordance with FDA Labeling ( ) yes (x ) no

d. Use of an Approved Drug ( ) yes ( ) no

e. Review of Medical Records or other existing data (x ) yes ( ) no

f. Audio Recordings ( ) yes ( ) no

g. Visual Recordings and/or photographs ( ) yes ( ) no

h. Telephone Interviews and/or Personal Interviews (please attach copy) ( ) yes ( ) no

i. Questionnaires and/or Psychological or Similar Tests (please attach copy) ( ) yes ( x ) no

j. On-line/Internet-based Questionnaires or Survey (s) ( ) yes ( ) no

k. Handling of Blood and/or Bodily Fluids (please provide documentation that all study staff have been instructed in the relevant OSHA regulations). N/A

**8. Research Support**

a. ( ) Outside funding-specify: \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

b**.** ( ) Internal (FHC) funding-specify: \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

**c.** ( ) None

(If you checked a or b please complete Appendix B)

1. **Information about Possible Conflicts of Interest**

Do any of the researchers have a conflict of interest (financial, investigator / provider, or other) related to the proposed research?

( ) yes ( ) no

If yes, each researcher having a conflict of interest must be identified and the conflict described in the above application and any conflict must also be described in the consent form for the participants.

**I, the undersigned, accept the responsibility for assuring that this study will be conducted in accordance with all applicable federal and state laws and regulations and the policies of Franciscan Children’s Hospital,** **with regard to the protection of human participants participating in this study.**

I agree to request and obtain approval from the Institutional Review Board prior to making any changes in this study as described in the application. I also agree to promptly notify the IRB of any emergent problems that may arise in the course of the study, including untoward results or unanticipated side effects.

Signature of Principal Investigator(s): \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

#### Franciscan Children’s Hospital

#### IRB Application

#### APPENDIX A - Review by Other Institutions/Institutional Reliance

**Study Title:** \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

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**Principal Investigator(s):** \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_ \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

I. Has this proposal been reviewed by an Institutional Review Board/Human Participants Committee at another institution?

( ) no ( ) yes ( ) N/A-FC ( ) to be submitted

Submission Date:

Pending Approval:

Approval Date:

Deferral Date:

A. Name of institution:

B. If known, application ID # at other institution:

C. Principal Investigator for other submission:

D. Is a Reliance agreement being sought? \_\_\_\_\_\_Yes \_\_\_\_\_No

If yes, please provide detail(s) of request.

**Franciscan Children’s Hospital**

**IRB Application**

**APPENDIX B - Study Funding**

**Study Title:** \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

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**Principal Investigator(s):** \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_ \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

I. Funds for expected support for the proposed research

### Have been awarded ( ) no ( ) yes

Have been applied for ( ) no ( ) yes

Will be applied for ( ) no ( ) yes

II. Funds will be administered through the business office at:

( ) FC

( ) Other-please specify: \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

II. Grant/Contact Number:

1. Period of support (start-end dates):
2. Amount of support for each year: