



UNIVERSITY OF COLORADO
HOSPITAL
Far Beyond The Ordinary™

HRRC CONDITIONAL APPROVAL FOR WIRB STUDIES

COMIRB #
Protocol Title:
Study Sponsor:
Principal Investigator:
Principal Investigator's Department:
Research Coordinator: Phone:

The study listed above is conditionally approved by the HRRC based on the documents submitted.

Final HRRC approval requires WIRB approval, an approved study budget and a signed copy of the Clinical Trial Agreement.

Any amendments to the Protocol received after the initial submission to the HRRC that alter the Consent Form or change the study budget must be sent to the HRRC.

Please note COMIRB will be unable to process your submission to WIRB without this form.

UCH APPROVAL

HRRC CHAIRPERSON APPROVAL: _____
(Signature)

(Date)