Clinic

Institutional Review Board
AAHRPP Accredited

March 4, 2013

, M.D.

RE: FA 07 05 028B: The Cord Blood Center and Hospital Volunteer Umbilical Cord Blood Program

Dear Dr.

Your study renewal application received on February 20, 2013 was reviewed on March 1, 2013 by the convened IRB and **approved for the period of March 24, 2013 to March 23, 2014.**

You are approved to continue this research with the Protocol Version 1.3 dated March 3, 2010, informed consent document V1 dated March 12, 2012, Self Inclusion/Exclusion Requirements, advertisements, Informational Brochures and UCB Collection Material/Infant Data Form.

The stamp-approved Consent form, data form, self inclusion/exclusion requirements form and advertisements are available online under the Stamped Documetns tab.

Written consent is required to document that each person enrlled in this research has been informed about this research and voluntarily agrees to particiapte prior to any involvement in the research.

Any changes or amendments require IRB review and approval prior to implementation. Unanticipated problems including adverse events and deviations are to be reported in accordance with IRB Policy 60: Adverse Events and IRB Policy 70: Unanticipated Problems.

This study may not continue beyond the approved expiration date. To request continuation, submit a renewal application 30 days prior to expiration or a completion report for closure.

Sincerely,

, MS, MHA, CIP
Executive Director, IRB and Human Research Protections

DB:sr

EXPIRATION DATE: March 23, 2014

A signed version of this letter is available online under the Correspondence tab