

June 9, 2011

To Whom It May Concern,

On February 2, 1998, the National Marrow Donor Program® (NMDP) submitted an IND application to the U.S. Food and Drug Administration (FDA): "*Centralized Cord Blood Registry to Facilitate Allogeneic, Unrelated Donor Cord Blood Transplantation.*" The FDA assigned the IND number BB-IND-7555. The FDA accepted the IND which included the protocol of the same name: "*Centralized Cord Blood Registry to Facilitate Allogeneic, Unrelated Donor Cord Blood Transplantation, A Phase 2 Clinical Trial Protocol.*"

On February 4, 2011, the NMDP submitted a new protocol under the same IND: "*Protocol 10-CBA: A Multicenter Access and Distribution Protocol for Unlicensed Cryopreserved Cord Blood Units (CBUs) for Transplantation in Pediatric and Adult Patients with Hematologic Malignancies and Other Indications.*" The FDA accepted that protocol under BB-IND-7555.

I can be reached by phone at (612) 884-8703 or by email at lphillip@nmdp.org if you have questions.

Kind Regards,



Lisa Phillips Johnson, MT(ASCP)SBB, CQA(ASQ) CQM-OE
Director, Regulatory Compliance