National Marrow Donor Program® Institutional Review Board
IRB Registration: IRB00001253 Assurance: FWA00000441

Notice of Action

Date: July 13, 2020 **Study Number:** IRB-1991-0002

Meeting Date: Expedited Review

Principal Investigator: Stephen Spellman

Study Title: *Protocol for a Research Sample Repository for Hematopoietic Cell Transplantation, Other Cellular Therapies and Marrow Toxic Injuries*

Protocol Version: 12.0

Number of participants approved: Unlimited

TYPE OF REVIEW:
AMENDMENT. Documents reviewed are listed below:

- 2020 Repository Record of Revisions (Misc/Other)
- Repository DONOR Adult Parent Consent v18.0 (Consent Form)
- Repository DONOR Adult Parent Consent v18.0_tracked (Consent Form)
- Repository DONOR Match Algorithm Consent v16.0 (Consent Form)
- Repository DONOR Match Algorithm Consent v16.0_tracked (Consent Form)
- Repository DONOR Minor Related Assent 12-17 v13.0 (Consent Form)
- Repository DONOR Minor Related Assent 12-17 v13.0_tracked (Consent Form)
- Repository DONOR Minor Related Assent 7-11 v13.0 (Consent Form)
- Repository DONOR Minor Related Assent 7-11 v13.0_tracked (Consent Form)
- Repository MARROW TOXIC INJURY Adult Parent Consent v14.0 (Consent Form)
- Repository MARROW TOXIC INJURY Adult Parent Consent v14.0_tracked (Consent Form)
- Repository MARROW TOXIC INJURY Assent 12-17 v12.0.doc (Consent Form)
- Repository MARROW TOXIC INJURY Assent 12-17 v12.0_tracked.doc (Consent Form)
- Repository RECIPIENT Adult Parent Consent v19.0 (Consent Form)
- Repository RECIPIENT Adult Parent Consent v19.0_tracked (Consent Form)
- Repository RECIPIENT Minor Assent 12-17 v14.0.doc (Consent Form)
- Repository RECIPIENT Minor Assent 12-17 v14.0_tracked.doc (Consent Form)
- Repository SECONDARY PRIMARY MALIGNANCY Adult Parent Consent v3.0.docx (Consent Form)
- Repository SECONDARY PRIMARY MALIGNANCY Adult Parent Consent v3.0_tracked.docx (Consent Form)

STATUS:
APPROVED

Notice of Action

Amendment Approved as of: July 13, 2020

- All projects must be reviewed for continuation of work. No modification may be made in the protocol or in the wording of the IRB approved consent(s) without the prior approval of the IRB.
- For donors not covered by the NMDP IRB, additional IRB approval from participating donor centers will be required.
- Donor/recipient consent forms are attached, if applicable.
- The Principal Investigator is responsible for reviewing the information in the email that accompanies this Notice of Action.

Reconsent (if applicable)**:** The NMDP IRB determined that re-consent of study subjects is NOT required. Standard practice has not changed, this amendment is primarily clarification of standard practice.

Authorized signature:

Electronically signed by Brian Lindberg, J.D. on 7/13/2020 2:09:38 PM