Publication Instructions:

BMT CTN Protocol 0201
- Primary and Ancillary Studies –

The requirements below are for publications and presentations made pursuant to the primary BMT CTN Study 0201 and the following correlative studies:
Patient Quality of Life
Immunophenotyping, Tetramer, TREC and T-Cell Response Assays
Immunoglobulin, Various Antibody & IL-2/IL-7 Levels
Donor HSC Graft Characterization ASH presentation

See below for Donor Quality of Life

For other ancillary studies, please contact the e-mail below.

Acknowledgment Requirements:

For all Publications, Reports, etc.

"Support for this study was provided to the Blood and Marrow Transplant Clinical Trials Network by grant #U10HL069294 from the National Heart, Lung, and Blood Institute and the National Cancer Institute. The Department of the Navy, Office of Navy Research, and the National Marrow Donor Program also provided support for the BMT CTN 0201 study. Enrollment support was provided by DKMS Germany. Any views, opinions, findings, conclusions or recommendations expressed in this material are those of the author(s) and do not reflect the views or the official policy or position of the above mentioned parties."

For all Presentations, Symposiums, etc.

"Support for this study was provided to the Blood and Marrow Transplant Clinical Trials Network by grant #U10HL069294 from the National Heart, Lung, and Blood Institute and the National Cancer Institute, the Health Resources and Services Administration Contract No. HHSH234200637020C, the Department of the Navy, Office of Naval Research, and the National Marrow Donor Program."

Additional Language for Publications on the Donor Quality of Life Ancillary Study:

"Contributions to the Donor Quality of Life ancillary study were made by The Health Resources and Services Administration Contract No. HHSH234200637020C to the National Marrow Donor Program. The authors of this _____[publication/report] alone are responsible for reporting and interpreting the data and all or part of the data used to compile this _____ [publication/report] were collected pursuant to a contract with the Department of Health and Human Services, Health Resources and Services Administration, Healthcare Systems Bureau, Division of Transportation."

* If HRSA staff substantially contributed in the analysis or in the writing of the report or publication, those staff MUST be included as authors.

For correlative studies utilizing biospecimens and/or clinical data obtained from the NHLBI Biologic Specimen and Data Repository Information Coordinating Center, include the following as applicable:

If the Research Plan involves collaboration with Study Investigators, acknowledge the source of the data by including language similar to the following either in the acknowledgement or in the text of the manuscript:

"This manuscript was prepared using BMT CTN 0201 Research Materials obtained from the NHLBI."

If the Research Plan does not involve collaboration with Study investigators, or if the Study has ended, use the following language:

"This manuscript was prepared using BMT CTN 0201 Research Materials obtained from the NHLBI Biologic Specimen and Data Repository Information Coordinating Center and does not necessarily reflect the opinions or views of the BMT CTN 0201 or the NHLBI."

Manuscripts and abstracts resulting from the Research Plan should not use the name of the Study in the title of the manuscript/abstract unless the title clearly denotes the source of the Research Materials as being from the NHLBI Biologic Specimen and Data Repository Information Coordinating Center.

The following language should be inserted in the acknowledgements for all BMT CTN Studies that utilize CIBMTR data:
The CIBMTR registry is supported primarily by the U24-CA76518 from the National Cancer Institute, the National Heart, Lung, and Blood Institute, and the National Institute of Allergy and Infectious Diseases and from HHSH234200637015C (HRSA/DHHS) to the Center for International Blood and Marrow Transplant Research.

Review Requirements:

For all Publications, Reports, etc.

BMT CTN: Abstracts, presentations, and proposed publications relating to data obtained from BMT CTN protocols or to activities of BMT CTN Committees or Protocol Teams are to be submitted to the DCC for review.

HRSA and ONR: One copy of each publication (or paper planned for publication) must be provided to the Office of Naval Research (ONR) and, if regarding the Donor Quality of Life Ancillary Study, to the Health Resources and Services Administration (HRSA). **** Review submissions to HRSA and ONR should be submitted to the email below.**

Questions regarding the above requirements may be directed to the Data Coordinating Center, NMDP Contracts & Procurement Department at NMDP_BMTCTN_Pub@nmdp.org.