• I agree this dataset is intended for public access and use.
• I agree that downloading of datasets implies agreement to Terms and Conditions.
• I agree to the citation policy: All publications or presentations of these data shall acknowledge CIBMTR as a data source.
• I agree to apply the following CIBMTR citation policy to the research paper: This dataset was collected by the Center for International Blood and Marrow Transplant Research (CIBMTR) which is supported primarily by the Public Health Service U24CA076518 from the National Cancer Institute; the National Heart, Lung, and Blood Institute; the National Institute of Allergy and Infectious Diseases; 75R60222C00011 from the Health Resources and Services Administration; N00014-23-1-2057 and N00014-24-1-2507 from the Office of Naval Research; NMDP; and the Medical College of Wisconsin.
• I agree to not sell information derived from the data unless express written permission has been received from CIBMTR.
• Ensure protection of the data. If accessing the data from a remote location on a time-sharing Network, computer system, or local area network with any statistical package, you will not share with any other individual(s).
• I agree that the data are private and confidential and that you will have in place and shall maintain administrative, technical, procedural, and physical safeguards sufficient to protect the confidentiality of the data, including preventing unauthorized access and use.
• I agree to not use or permit others to use or to link the data to identify individuals, or to combine the data with other patient-level information.
• I agree to not make copies of the data. Agree to neither release nor permit others to release the files or data therein to any person (including media and subcontractors) except with the written approval of CIBMTR in order to accomplish the study goals.
• Certify that you are responsible for ensuring that only staff with a “need to know” shall access the data and that any support staff assigned to this project and having access to these data will likewise follow these provisions.
• I agree that all data will be destroyed upon completion of the approved use.
• I agree to comply with all applicable laws and regulation with regard to use of the data, including but not limited to appropriate IRB oversight consistent with the uses of the data where applicable. Any publication deriving from these data must contain a statement confirming the study was conducted in accordance with all applicable human subjects’ protections laws and regulations.
• I acknowledge that European Union patient-level data is redacted. For publications that include EU patient data the publicly available dataset will not contain the same patient population as the publication, and therefore users may not be able to fully replicate the findings from the publication.
• I acknowledge that supplemental data may not be included and will depend on the funds used to acquire the data and the contractual agreements in place for those funds and data.