

Registry Use Only Sequence Number: Date Received:	OMB No: 0915-0310 Expiration Date: 10/31/2022 <p>Public Burden Statement: The purpose of the data collection is to fulfill the legislative mandate to establish and maintain a standardized database of allogeneic marrow and cord blood transplants performed in the United States or using a donor from the United States. The data collected also meets the C.W. Bill Young Cell Transplantation Program requirements to provide relevant scientific information not containing individually identifiable information available to the public in the form of summaries and data sets. An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. The OMB control number for this information collection is 0915-0310 and it is valid until 10/31/2022. This information collection is voluntary under The Stem Cell Therapeutic and Research Act of 2005, Public Law (Pub. L.) 109-129, as amended by the Stem Cell Therapeutic and Research Reauthorization Act of 2010, Public Law 111-264 (the Act) and the Stem Cell Therapeutic and Research Reauthorization Act of 2015, Public Law 114-104. Public reporting burden for this collection of information is estimated to average 0.85 hours per response when collected at 100 days post-transplant, 0.85 hours per response when collected at 6 months post-transplant, 0.65 hours per response when collected at 1 and 2 years post-transplant, and 0.52 hours per response annually thereafter, including the time for reviewing instructions, searching existing data sources, and completing and reviewing the collection of information. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden, to HRSA Reports Clearance Officer, 5600 Fishers Lane, Room 14N136B, Rockville, Maryland, 20857 or paperwork@hrsa.gov.</p>
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CIBMTR Center Number: _____ CIBMTR Research ID: _____ Event date: _____ - _____ - _____ YYYY MM DD Visit: <input type="checkbox"/> 100 day <input type="checkbox"/> 6 months <input type="checkbox"/> 1 year <input type="checkbox"/> 2 years <input type="checkbox"/> >2 years, Specify: _____

Organ failure (not due to GVHD or infection)

- Liver failure (not VOD) – **Go to question 5**
- Venous-occlusive disease (VOD) / sinusoidal obstruction syndrome (SOS) – **Go to question 5**
- Cardiac failure – **Go to question 5**
- Pulmonary failure– **Go to question 5**
- Central nervous system (CNS) failure – **Go to question 5**
- Renal failure – **Go to question 5**
- Gastrointestinal (GI) failure (not liver) – **Go to question 5**
- Multiple organ failure – **Go to question 4**
- Other organ failure – **Go to question 4**

Malignancy

- New malignancy (post-HCT or post-cellular therapy) – **Go to question 5**
- Prior malignancy (malignancy initially diagnosed prior to HCT or cellular therapy, other than the malignancy for which the HCT or cellular therapy was performed) – **Go to question 5**

Hemorrhage

- Pulmonary hemorrhage – **Go to question 5**
- Diffuse alveolar hemorrhage (DAH) – **Go to question 5**
- Intracranial hemorrhage – **Go to question 5**
- Gastrointestinal hemorrhage – **Go to question 5**
- Hemorrhagic cystitis – **Go to question 5**
- Other hemorrhage – **Go to question 4**

Vascular

- Thromboembolic – **Go to question 5**
- Disseminated intravascular coagulation (DIC) – **Go to question 5**
- Thrombotic microangiopathy (TMA) (Thrombotic thrombocytopenic purpura (TTP)/Hemolytic Uremic Syndrome (HUS))– **Go to question 5**
- Other vascular - **Go to question 4**

Other

- Accidental death – **Go to question 5**
- Suicide – **Go to question 5**
- Other cause - **Go to question 4**

4. Specify: _____

5. Contributing cause of death (check all that apply)
- Recurrence / persistence / progression of disease for which the HCT or cellular therapy was performed – **Go to question 7**
 - Acute GVHD – **Go to question 7**
 - Chronic GVHD – **Go to question 7**
 - Graft rejection or failure – **Go to question 7**
 - Cytokine release syndrome – **Go to question 7**

Infection

- Infection, organism not identified – **Go to question 7**
- Bacterial infection – **Go to question 7**
- Fungal infection – **Go to question 7**
- Viral infection – **Go to question 7**
- COVID-19 (SARS-CoV-2) – **Go to question 7**
- Protozoal infection – **Go to question 7**
- Other infection – **Go to question 6**

Pulmonary

- Idiopathic pneumonia syndrome (IPS) – **Go to question 7**
- Pneumonitis due to Cytomegalovirus (CMV) – **Go to question 7**
- Pneumonitis due to other virus – **Go to question 7**
- Other pulmonary syndrome (excluding pulmonary hemorrhage) – **Go to question 6**
- Diffuse alveolar damage (without hemorrhage) – **Go to question 7**
- Acute respiratory distress syndrome (ARDS) (other than IPS) – **Go to question 7**

Organ failure (not due to GVHD or infection)

- Liver failure (not VOD) – **Go to question 7**
- Venous-occlusive disease (VOD) / sinusoidal obstruction syndrome (SOS) – **Go to question 7**
- Cardiac failure – **Go to question 7**
- Pulmonary failure – **Go to question 7**
- Central nervous system (CNS) failure – **Go to question 7**
- Renal failure – **Go to question 7**
- Gastrointestinal (GI) failure (not liver) – **Go to question 7**
- Multiple organ failure – **Go to question 6**
- Other organ failure – **Go to question 6**

CIBMTR Center Number: _____ CIBMTR Research ID: _____

- II - Rash on > 50% of skin, bilirubin 2-3 mg/dL, or diarrhea 500 – 1000 mL/day or persistent nausea or vomiting
- III - Bilirubin 3-15 mg/dL, or gut stage 2-4 diarrhea > 1000 mL/day or severe abdominal pain with or without ileus
- IV - Generalized erythroderma with bullous formation, or bilirubin >15 mg/dL
- Not applicable (acute GVHD present but grade is not applicable)

List the stage for each organ at diagnosis of acute GVHD:

23. Skin:
- Stage 0 – no rash, no rash attributable to acute GVHD
 - Stage 1 – maculopapular rash, < 25% of body surface
 - Stage 2 – maculopapular rash, 25–50% of body surface
 - Stage 3 – generalized erythroderma, > 50% of body surface
 - Stage 4 – generalized erythroderma with bullae formation and/or desquamation
24. Lower intestinal tract: (use mL/day for adult recipients and mL/kg/day for pediatric recipients)
- Stage 0 – no diarrhea, no diarrhea attributable to acute GVHD / diarrhea < 500 mL/day (adult), or < 10 mL/kg/day (pediatric)
 - Stage 1 – diarrhea 500 - 1000 mL/day (adult), or 10 - 19.9 mL/kg/day (pediatric)
 - Stage 2 – diarrhea 1001 - 1500 mL/day (adult), or 20 - 30 mL/kg/day (pediatric)
 - Stage 3 – diarrhea > 1500 mL/day (adult), or > 30 mL/kg/day (pediatric)
 - Stage 4 – severe abdominal pain, with or without ileus, and/or grossly bloody stool
25. Upper intestinal tract:
- Stage 0 – no persistent nausea or vomiting
 - Stage 1 – persistent nausea or vomiting
26. Liver:
- Stage 0 – No liver acute GVHD / bilirubin < 2.0 mg/dL (< 34 µmol/L)
 - Stage 1 – bilirubin 2.0–3.0 mg/dL (34–52 µmol/L)
 - Stage 2 – bilirubin 3.1–6.0 mg/dL (53–103 µmol/L)
 - Stage 3 – bilirubin 6.1–15.0 mg/dL (104–256 µmol/L)
 - Stage 4 – bilirubin > 15.0 mg/dL (> 256 µmol/L)
27. Other site(s) involved with acute GVHD

CIBMTR Center Number: _____ CIBMTR Research ID: _____

- Yes – **Go to question 28**
- No – **Go to question 29**

28. Specify other site(s): _____

Specify the maximum overall grade and organ staging of acute GVHD since the date of last report

29. Maximum overall grade of acute GVHD:

- I - Rash on \leq 50% of skin, no liver or gut involvement
- II - Rash on $>$ 50% of skin, bilirubin 2-3 mg/dL, or diarrhea 500 – 1000 mL/day or persistent nausea or vomiting
- III - Bilirubin 3-15 mg/dL, or gut stage 2-4 diarrhea $>$ 1000 mL/day or severe abdominal pain with or without ileus
- IV - Generalized erythroderma with bullous formation, or bilirubin $>$ 15 mg/dL
- Not applicable (acute GVHD present but cannot be graded)

30. Date maximum overall grade of acute GVHD: _____ - _____ - _____

31. Skin:

- Stage 0 – no rash, no rash attributable to acute GVHD
- Stage 1 – maculopapular rash, $<$ 25% of body surface
- Stage 2 – maculopapular rash, 25–50% of body surface
- Stage 3 – generalized erythroderma, $>$ 50% of body surface
- Stage 4 – generalized erythroderma with bullae formation and/or desquamation

32. Lower intestinal tract: (use mL/day for adult recipients and mL/kg/day for pediatric recipients)

- Stage 0 – no diarrhea, no diarrhea attributable to acute GVHD / diarrhea $<$ 500 mL/day (adult), or $<$ 10 mL/kg/day (pediatric)
- Stage 1 – diarrhea 500 - 1000 mL/day (adult), or 10 - 19.9 mL/kg/day (pediatric)
- Stage 2 – diarrhea 1001 - 1500 mL/day (adult), or 20 - 30 mL/kg/day (pediatric)
- Stage 3 – diarrhea $>$ 1500 mL/day (adult), or $>$ 30 mL/kg/day (pediatric)
- Stage 4 – severe abdominal pain, with or without ileus, and/or grossly bloody stool

33. Upper intestinal tract:

- Stage 0 – no persistent nausea or vomiting
- Stage 1 – persistent nausea or vomiting

34. Liver:
- Stage 0 – No liver acute GVHD / bilirubin < 2.0 mg/dL (< 34 µmol/L)
 - Stage 1 – bilirubin 2.0–3.0 mg/dL (34–52 µmol/L)
 - Stage 2 – bilirubin 3.1–6.0 mg/dL (53–103 µmol/L)
 - Stage 3 – bilirubin 6.1–15.0 mg/dL (104–256 µmol/L)
 - Stage 4 – bilirubin > 15.0 mg/dL (> 256 µmol/L)

35. Other site(s) involved with acute GVHD
- Yes – **Go to question 36**
 - No – **Go to question 37**

36. Specify other site(s): _____

37. Did chronic GVHD develop since the date of last report?
- Yes – **Go to questions 38**
 - No - **Go to question 39**
 - Unknown – **Go to question 39**

38. Date of chronic GVHD diagnosis: _____ Date estimated – **Go to questions 40**

YYYY MM DD

39. Did chronic GVHD persist since the date of last report?
- Yes – **Go to questions 40**
 - No - **Go to question 43**
 - Unknown – **Go to question 43**

Specify the maximum grade of chronic GVHD since the date of last report:

40. Maximum grade of chronic GVHD: (according to best clinical judgment)
- Mild
 - Moderate
 - Severe
 - Unknown

CIBMTR Center Number: _____

CIBMTR Research ID: _____

Copy and complete questions 53-59 to report each new malignancy diagnosed since the date of last report. The submission of a pathology report or other supportive documentation for each reported new malignancy is strongly recommended.

53. Specify the new malignancy:

- Acute myeloid leukemia (AML / ANLL) – **Go to question 56**
- Other leukemia – **Go to question 56**
- Myelodysplastic syndrome (MDS) – **Go to question 56**
- Myeloproliferative neoplasm (MPN) – **Go to question 56**
- Myelodysplasia / myeloproliferative neoplasm (MDS / MPN)– **Go to question 56**
- Hodgkin lymphoma – **Go to question 55**
- Non-Hodgkin lymphoma – **Go to question 55**
- Post-transplant lymphoproliferative disorder (PTLD)– **Go to question 55**
- Clonal cytogenetic abnormality without leukemia or MDS – **Go to question 56**
- Uncontrolled proliferation of donor cells without malignant transformation – **Go to question 56**
- Breast cancer – **Go to question 56**
- Central nervous system (CNS) malignancy (e.g. glioblastoma, astrocytoma) – **Go to question 56**
- Gastrointestinal malignancy (e.g. colon, rectum, stomach, pancreas, intestine) – **Go to question 56**
- Genitourinary malignancy (e.g. kidney, bladder, ovary, testicle, genitalia, uterus, cervix) – **Go to question 56**
- Lung cancer – **Go to question 56**
- Melanoma – **Go to question 56**
- Basal cell skin malignancy – **Go to question 56**
- Squamous cell skin malignancy – **Go to question 56**
- Oropharyngeal cancer (e.g. tongue, buccal mucosa) – **Go to question 56**
- Sarcoma – **Go to question 56**
- Thyroid cancer – **Go to question 56**
- Other new malignancy – **Go to question 54**

54. Specify other new malignancy: _____ - **Go to question 56**

55. Is the tumor EBV positive?

- Yes
- No

56. Date of diagnosis: _____

CIBMTR Center Number: _____ CIBMTR Research ID: _____

YYYY MM DD

57. Was documentation submitted to the CIBMTR? (e.g. pathology / autopsy report or other documentation)
- Yes
 - No
58. Was the new malignancy donor / cell product derived?
- Yes – **Go to question 59**
 - No – **Go to question 59**
 - Not done – **Go to question 60**
59. Was documentation submitted to the CIBMTR? (e.g. cell origin evaluation (VNTR, cytogenetics, FISH))
- Yes
 - No

Chimerism Studies (Cord Blood Units, Beta Thalassemia, and Sickle Cell Disease Only)

This section relates to chimerism studies from allogeneic HCTs using cord blood units or for recipients whose primary disease is beta thalassemia or sickle cell disease. If this was an autologous HCT, or an allogeneic HCT using a bone marrow or PBSC product, or a different primary disease, continue to disease assessment.

60. Were chimerism studies performed since the date of last report?
- Yes – **Go to question 61**
 - No – **Go to question 80**
61. Was documentation submitted to the CIBMTR? (e.g. chimerism laboratory reports)
- Yes
 - No
62. Were chimerism studies assessed for more than one donor / multiple donors?
- Yes
 - No

Provide date(s), method(s) and other information for all chimerism studies performed since the date of last report.

CIBMTR Center Number: _____ CIBMTR Research ID: _____

- Total mononuclear cells (lymphs & monos) – **Go to question 78**
- T-cells (includes CD3+, CD4+, and/or CD8+) – **Go to question 78**
- B-cells (includes CD19+ or CD20+) – **Go to question 78**
- Granulocytes (includes CD33+ myeloid cells) – **Go to question 78**
- NK cells (CD56+) – **Go to question 78**
- Other – **Go to question 75**

75. Specify: _____

76. Total cells examined: _____

77. Number of donor cells: _____ - **Go to question 80**

78. Were donor cells detected?

- Yes - **Go to question 79**
- No – **Go to question 80**

79. Percent donor cells: _____ %

Copy questions 63 – 79 if needed for multiple chimerism studies.

Disease Assessment at the Time of Best Response to HCT

80. Compared to the disease status prior to the preparative regimen, what was the best response to HCT since the date of the last report? (Include response to any therapy given for post-HCT maintenance or consolidation, but exclude any therapy given for relapsed, persistent, or progressive disease)

- Continued complete remission (CCR) - **For patients transplanted in CR- Go to question 103**
- Complete remission (CR) - **Go to question 82**
- Not in complete remission - **Go to question 81**
- Not evaluated - **Go to question 103**

81. Specify disease status if not in complete remission:

- Disease detected - **Go to question 84**
- No disease detected but incomplete evaluation to establish CR - **Go to question 84**

82. Was the date of best response previously reported?

CIBMTR Center Number: _____ CIBMTR Research ID: _____

- Ibrutinib
- Imatinib mesylate (Gleevec)
- Ixazomib
- Lenalidomide (Revlimid)
- Lestaurtinib
- Midostaurin
- Nilotinib (AMN107, Tasigna)
- Nivolumab
- Pembrolizumab
- Pomalidomide
- Quizartinib
- Rituximab (Rituxan, MabThera)
- Sorafenib
- Sunitinib
- Thalidomide (Thalomid)
- Other systemic therapy- **Go to question 106**

106. Specify other systemic therapy: _____

107. Specify other therapy: _____

Relapse or Progression Post-HCT

Report if the recipient has experienced a clinical/hematologic relapse or progression post-HCT. If the relapse or progression was detected in a previous reporting period indicate that and continue on. If the first clinical/hematologic relapse occurred since the date of last report, indicate the date it was first detected in this reporting period.

108. Did the recipient experience a clinical/hematologic relapse or progression post-HCT?

- Yes - **Go to question 109**
- No - **Go to question 111**

109. Was the date of the first clinical/hematologic relapse or progression previously reported?

- Yes - **Go to question 119 (only valid >day 100)**

CIBMTR Center Number: _____

CIBMTR Research ID: _____

- Bortezomib (Velcade)
- Bosutinib
- Carfilzomib
- Chemotherapy
- Dasatinib (Sprycel)
- Decitabine (Dacogen)
- Gemtuzumab (Mylotarg, anti-CD33)
- Gilteritinib
- Ibrutinib
- Imatinib mesylate (Gleevec)
- Ixazomib
- Lenalidomide (Revlimid)
- Lestaurtinib
- Midostaurin
- Nilotinib (AMN107, Tasigna)
- Nivolumab
- Pembrolizumab
- Pomalidomide
- Quizartinib
- Rituximab (Rituxan, MabThera)
- Sorafenib
- Sunitinib
- Thalidomide (Thalomid)
- Other systemic therapy- **Go to question 117**

117. Specify other systemic therapy: _____

118. Specify other therapy: _____

Current Disease Status

119. What is the current disease status?

- Complete remission (CR) - **Go to question 121**

