



Transplant Essential Data

First Report: 100 Days Post Transplant



Primary Disease Diagnosis: _____

Graft: Auto Allo Syngeneic

Date of This Report: _____ - ____ - ____
YYYY MM DD

PATIENT IDENTIFICATION

Hospital Unique Patient Number: _____

Date of Birth: _____ - ____ - ____
YYYY MM DD

Sex: Male Female

Ethnicity (optional for non-US centers): White/Caucasian
 Black Oriental Other, specify: _____

DISEASE

(complete appropriate disease classification sheet)

Date of initial diagnosis of primary disease: _____ - ____ - ____
YYYY MM DD

TRANSPLANTATION

Date of this transplant: _____ - ____ - ____
YYYY MM DD

Chronological number of this transplant for this patient: _____
 If >1, date of most recent previous transplant: _____ - ____ - ____
YYYY MM DD

If >1, type of most recent previous transplant: Auto Allo

Source of Stem Cells for this transplant (check all that apply):
 Bone marrow Peripheral blood
 Cord blood Other: _____

Donor Type (check one):
 Autologous (self) Syngeneic (monozygotic twin)
 Allogeneic:
 HLA-identical sibling (not monozygotic twin)
 HLA-matched other relative
 HLA-mismatched relative
 HLA-matched unrelated donor
 HLA-mismatched unrelated donor
 Multiple donors

(For allotransplants) donor sex: Male Female

Was the graft manipulated *in vivo* other than for RBC removal or volume reduction? Yes No

Was this transplant part of a planned sequential transplant protocol? Yes No

Additional cell or therapy given (not for relapse)?
(If additional transplant given, submit separate TED form)
 Yes No Unknown

If yes, type of cell(s) (check all that apply):
 Lymphocytes Fibroblasts Dendritic cells
 Mesenchymal Other: _____

If yes, date of first infusion of additional cell therapy (may be the same as transplant date): _____ - ____ - ____
YYYY MM DD

Was Gleevec (STI571, imatinib mesylate) given posttransplant? Yes No Unknown

NA = not applicable, autotransplant

CENTER IDENTIFICATION

Center Identification Code:
 IBMTR/ABMTR: _____
 EBMT: _____

National (specify): _____
 Other (specify): _____

Hospital: _____
 Unit: _____
 Contact person: _____
 Phone #: _____
 Fax #: _____
 Email: _____

BEFORE TRANSPLANTATION

Performance Score Pretransplant:
 Good (KPS ≥80 ~or~ ECOG 0-1 ~or~ Lansky ≥80)
 Poor (KPS <80 ~or~ ECOG 2-4 ~or~ Lansky <80)
 Unknown

Conditioning Regimen: Total Body Irradiation? Yes No
 Non-myeloablative/Reduced Intensity (allo only)? Yes No

AFTER TRANSPLANTATION

Engraftment (Neutrophils >1000/µL)?
 Yes No Unknown
 If yes, date Neutrophils >1000/µL: _____ - ____ - ____
YYYY MM DD

If no, date of latest assessment: _____ - ____ - ____
YYYY MM DD

Maximum Grade of Acute Graft Versus Host Disease (GVHD):
 0 1 2 3 4 Unknown NA

Best disease response to transplant (malignant and pre-malignant diseases only):
 Continued CR CR achieved, date achieved: _____ - ____ - ____
YYYY MM DD

Never in CR posttransplant, date assessed: _____ - ____ - ____
YYYY MM DD

Unknown _____ - ____ - ____
YYYY MM DD

Did the disease for which the patient was transplanted relapse or progress after the transplant?
 Yes No Unknown

Indicate all methods used for the assessment, the date of assessment and whether relapse/progression was detected with that method on the date indicated:
 Molecular
 Date assessed: _____ - ____ - ____
YYYY MM DD

Relapse/progression first detected? Yes No

Cytogenetic
 Date assessed: _____ - ____ - ____
YYYY MM DD

Relapse/progression first detected? Yes No

Hematological/Clinical
 Date assessed: _____ - ____ - ____
YYYY MM DD

Relapse/progression first detected? Yes No

Survival status after transplant:
 Alive Dead Died before transplant
 Date of latest follow-up or death: _____ - ____ - ____
YYYY MM DD

Check here if lost to follow-up

Main cause of death (check one):
 Relapse/Progression/Persistent disease
 Transplantation related causes:
 Rejection/Poor graft function GVHD
 Pulmonary toxicity Cardiac toxicity
 Infection VOD
 Posttransplant lymphoproliferative disorder Other: _____
 Unknown

REGISTRY USE ONLY

Date Received: _____

Retired - Not for Data Submission



**Transplant Essential Data
First Report: 100 Days Post Transplant
Disease Classification Sheet 1**



PATIENT IDENTIFICATION

Hospital Unique Patient Number: _____

CENTER IDENTIFICATION

Center Identification Code:
IBMTR/ABMTR _____

ACUTE LEUKEMIAS

Classification:

Acute Myelogenous Leukemia (AML)

- M0
- M1
- M2
- M3
- M4
- M5
- M6
- M7
- AML NOS
- Transformed from MDS _____
- Other AML, specify: _____

FAB classification: M _____

Acute Lymphoblastic Leukemia (ALL)

- ALL B-lineage
- ALL T-lineage
- Mature B cell (L3)
- T-cell granular lymphocytic leukemia
- Aggressive NK-cell leukemia
- Adult T-cell lymphoma/leukemia (HTLV1+)
- ALL NOS
- Other ALL, specify: _____

Other Acute Leukemias

- Acute undifferentiated
- Acute biphenotypic
- Acute mast cell leukemia
- Other acute leukemia, specify: _____

Complete entire MDS Section on Disease Classification Sheet 2 and remainder of AML Section, except status at transplantation

Was Gleevec (STI571, imatinib mesylate) given for pretransplant therapy? Yes No Unknown
 Was AML caused by prior exposure to therapeutic drugs or radiation? Yes No Unknown

Status at Transplantation:

- Untreated
 - Primary Induction Failure (PIF)
 - CR _____ Number _____ 1st 2nd 3rd or higher
 - Rel _____
- Complete Remission**
- Y N Unk**
- Cytogenetic remission
 - Molecular remission

CHRONIC MYELOGENOUS LEUKEMIA (CML)

Classification:

- CML, Ph+
- CML, Ph-
- CML, NOS

Prior treatment (check all that apply):

- Interferon
- Hydroxyurea (HU)
- Gleevec (STI571, imatinib mesylate)
- Other, specify: _____

Status at Transplantation:

Phase Number For Chronic Phase Only (check all that apply)

- Chronic phase 1st Hematological remission: Yes No, stable phase
- Accelerated phase 2nd Cytogenetic remission: Complete Partial Cytogenetics unknown
- Blast crisis 3rd or higher Molecular (bcr/abl): Present Absent bcr/abl unknown

Other: _____

OTHER LEUKEMIAS

Classification:

- Chronic Lymphocytic Leukemia (CLL), B-cell/ small lymphocytic lymphoma
- CLL, T-cell
- CLL, NOS
- Prolymphocytic Leukemia
 - B-cell
 - T-cell
- Hairy Cell Leukemia
- Other leukemia, specify: _____

Status at Transplantation:

- Untreated
- CR
- PR
- No response/stable
- Progression

CR=complete remission, PR=partial remission, Rel=relapse, CP=chronic phase, AP=accelerated phase, BP=blast phase



**Transplant Essential Data
First Report: 100 Days Post Transplant
Disease Classification Sheet 2**



PATIENT IDENTIFICATION

Hospital Unique Patient Number: _____

CENTER IDENTIFICATION

Center Identification Code:
IBMTR/ABMTR _____

MYELOYDYSPLASTIC OR MYELOPROLIFERATIVE SYNDROMES

Classification:

Myelodysplastic Syndromes (MDS)*

- | | |
|--------------------------|--|
| At diagnosis | At transplantation |
| <input type="checkbox"/> | <input type="checkbox"/> RA |
| <input type="checkbox"/> | <input type="checkbox"/> RARS |
| <input type="checkbox"/> | <input type="checkbox"/> RAEB |
| <input type="checkbox"/> | <input type="checkbox"/> RAEB-t |
| <input type="checkbox"/> | <input type="checkbox"/> AML |
| <input type="checkbox"/> | <input type="checkbox"/> MDS NOS |
| <input type="checkbox"/> | <input type="checkbox"/> Other, specify: _____ |

Myeloproliferative Syndromes (MPS)

- | | |
|--------------------------|--|
| At diagnosis | At transplantation |
| <input type="checkbox"/> | <input type="checkbox"/> Polycythemia vera |
| <input type="checkbox"/> | <input type="checkbox"/> Essential or primary thrombocythemia |
| <input type="checkbox"/> | <input type="checkbox"/> Myelofibrosis with myeloid metaplasia |
| <input type="checkbox"/> | <input type="checkbox"/> Acute myelofibrosis or myelocystic fibrosis |
| <input type="checkbox"/> | <input type="checkbox"/> MPS, NOS |
| <input type="checkbox"/> | <input type="checkbox"/> Other MFS/MPS, specify: _____ |

Other

- | | |
|--------------------------|--|
| At diagnosis | At transplantation |
| <input type="checkbox"/> | <input type="checkbox"/> Chronic myelomonocytic leukaemia (CMML, CMML) |
| <input type="checkbox"/> | <input type="checkbox"/> Juvenile myelomonocytic leukaemia (JMML, JCML, JCMML) |

Status at Transplantation:

- Untreated (Supportive care only)
- Treatment without intent to achieve CR
- Treatment with intent to achieve a CR – CR not achieved
- Treatment with intent to achieve a CR – CR achieved and sustained
- Relapse after CR

Number
<input type="checkbox"/> 1st
<input type="checkbox"/> 2nd
<input type="checkbox"/> 3rd or higher

* If transformed to acute leukemia, report on Disease Classification Sheet 1

ANEMIA/HEMOGLOBINOPATHY

Classification:

- | | |
|---|--|
| <input type="checkbox"/> Acquired Severe Aplastic Anemia (SAA), Idiopathic | <input type="checkbox"/> Diamond-Blackfan anemia (congenital PRCA) |
| <input type="checkbox"/> Acquired SAA, secondary to hepatitis | <input type="checkbox"/> Other constitutional anemia, specify: _____ |
| <input type="checkbox"/> Acquired SAA, secondary to toxin/other drug | <input type="checkbox"/> Thalassemia NOS |
| <input type="checkbox"/> Amegakaryocytosis acquired (not congenital) | <input type="checkbox"/> Sickle cell disease |
| <input type="checkbox"/> Acquired Pure Red Cell Aplasia (PRCA) (not congenital) | <input type="checkbox"/> Other hemoglobinopathy, specify: _____ |
| <input type="checkbox"/> Schwachmann-Diamond | <input type="checkbox"/> Paroxysmal nocturnal hemoglobinuria (PNH) |
| <input type="checkbox"/> Other acquired cytopenic syndrome, specify: _____ | |
| <input type="checkbox"/> Fanconi anemia | |

PLATELET DISORDERS

Classification:

- Congenital amegakaryocytosis/congenital thrombocytopenia
- Glanzmann thrombasthenia
- Other inherited platelet abnormalities, specify: _____

HISTIOCYTIC DISORDERS

Classification:

- Histiocytic disorders, NOS
- Familial erythro/hemophagocytic lymphohistiocytosis (FELH)
- Langerhans Cell Histiocytosis (Histiocytosis-X)
- Hemophagocytosis (reactive or viral associated)
- Malignant histiocytosis
- Other histiocytic disorder, specify: _____

CR=complete remission



**Transplant Essential Data
First Report: 100 Days Post Transplant
Disease Classification Sheet 3**



PATIENT IDENTIFICATION

Hospital Unique Patient Number: _____

CENTER IDENTIFICATION

Center Identification Code: _____

IBMTR/ABMTR _____

LYMPHOMAS

Classification:

Hodgkin Disease

Non-Hodgkin's Lymphoma

B-cell Neoplasms

- Precursor B-lymphoblastic leukemia/lymphoma (precursor B-cell acute lymphoblastic leukemia)
- Lymphoplasmacytic lymphoma
- Splenic marginal zone B-cell lymphoma
- Extranodal marginal zone B-cell lymphoma of MALT type
- Nodal marginal zone B-cell lymphoma (+/- monocytoid B cells)
- Follicular lymphoma
- Mantle cell lymphoma
- Diffuse large B-cell lymphoma
- Burkitt's lymphoma/Burkitt cell leukemia
- Other B-cell lymphoma, specify: _____

T-cell and NK-cell Neoplasms

- Precursor T-lymphoblastic lymphoma/leukemia (precursor T-cell acute lymphoblastic leukemia)
- Extranodal NK/T-cell lymphoma, nasal type
- Enteropathy-type T-cell lymphoma
- Hepatosplenic gamma-delta T-cell lymphoma
- Subcutaneous panniculitis-like T-cell lymphoma
- Mycosis fungoides/Sézary syndrome
- Anaplastic large-cell lymphoma, T/null cell, primary cutaneous type
- Peripheral T-cell lymphoma, NOS
- Angioimmunoblastic T-cell lymphoma
- Anaplastic large-cell lymphoma, T/null cell, primary systemic type
- Other T/NK cell lymphoma, specify: _____

- Grade I
- Grade II
- Grade III
- Unknown

Status at Transplantation:

- Untreated
- Primary refractory (less than partial response to initial therapy)
- CR
 - CR confirmed
 - CR unconfirmed (CRU)
- 1st partial response (PR1)
- Rel

- Number
- 1st
 - 2nd
 - 3rd or higher

Sensitivity to Chemotherapy:

- (within 6 months prior to transplantation)
- Sensitive
 - Resistant
 - Untreated
 - Unknown

* CRU – complete response with persistent scan abnormalities of unknown significance

PLASMA CELL DISORDERS

Classification:

- Multiple myeloma-IgG _____
- Multiple myeloma-IgA _____
- Multiple myeloma-IgD _____
- Multiple myeloma-IgE _____
- Multiple myeloma-Ig light chain only _____
- Multiple myeloma non-secretory _____
- Plasma cell leukemia
- Solitary plasmacytoma (no evidence of myeloma)
- Waldenström macroglobulinemia (IgM)
- Primary Amyloidosis
- Other Plasma Cell Disorder, specify: _____

- SALMON & DURIE**
Stage at Diagnosis
(Multiple Myeloma only)
- 1 and A
 - 2 B
 - 3

Status at Transplantation:

- Untreated
 - CR
 - PR
 - MR
 - Progression/Relapse
 - No response/Stable disease
- Number
- 1st
 - 2nd
 - 3rd or higher

CR=complete remission, PR=partial remission, Rel=relapse, MR=minimal response



**Transplant Essential Data
First Report: 100 Days Post Transplant
Disease Classification Sheet 4**



PATIENT IDENTIFICATION

Hospital Unique Patient Number: _____

CENTER IDENTIFICATION

Center Identification Code: _____

IBMTR/ABMTR _____

BREAST CANCER

Classification:

Breast Cancer

- Inflammatory
- Non-inflammatory

Stage at Diagnosis

- 0
- I
- II
- III
- Inflammatory, no distant metastases
- Metastatic

Status at Transplantation:

- Adjuvant (Stage II, III only)
- Untreated (upfront)
- Primary refractory
- Complete remission (CR)
 - CR confirmed
 - CR unconfirmed (CRU)
- 1st partial response (PR1)
- Relapse
 - Local
 - Metastatic

Number

(complete only for CR or relapse)

- 1st
- 2nd
- 3rd or higher

Sensitivity to Chemotherapy
(complete only for relapse)

- Sensitive
- Resistant
- Untreated
- Unknown

* CRU – complete response with persistent scan abnormalities of unknown significance

OTHER MALIGNANCIES

Classification:

- Head and neck
- Lung cancer, small cell
- Lung cancer, non-small cell
- Lung cancer, NOS
- Thymoma
- Gastric
- Colorectal
- Pancreas
- Hepatobiliary
- Kidney and urinary tract
- Wilm tumor
- Prostate
- Testicular
- External genitalia
- Cervical
- Uterus
- Ewing sarcoma
- Ovary
- Vagina
- Germ cell tumor, extragonadal

- Sarcoma NOS
- Soft tissue sarcoma (include sarcoma PNET)
- Bone sarcoma (excluding Ewing sarcoma) (include sarcoma PNET)
- Rhabdomyosarcoma
- Leiomyosarcoma
- Liposarcoma
- Fibrosarcoma
- Synovial sarcoma
- Hemangiosarcoma
- Lymphangiosarcoma
- Neurogenic sarcoma
- Melanoma
- Central nervous system tumors (include CNS PNET)
- Medulloblastoma
- Neuroblastoma
- Retinoblastoma
- Mediastinal neoplasm,
specify: _____
- Other solid tumor,
specify: _____

Status at Transplantation:

- Untreated (upfront)
- Primary refractory
- Complete remission (CR)
- 1st very good partial response (VGPR1)
- 1st partial response (PR1)
- Relapse
- Adjuvant

Number

(complete only for CR or relapse)

- 1st
- 2nd
- 3rd or higher

Sensitivity to Chemotherapy
(complete only for relapse)

- Sensitive
- Resistant
- Untreated
- Unknown

CR=complete remission, PR=partial remission, VGPR=very good partial response, MR=minimal response



**Transplant Essential Data
First Report: 100 Days Post Transplant
Disease Classification Sheet 5**



PATIENT IDENTIFICATION

Hospital Unique Patient Number: _____

CENTER IDENTIFICATION

Center Identification Code:
IBMTR/ABMTR _____

INHERITED DISORDERS OF METABOLISM

Classification:

- | | |
|--|---|
| <input type="checkbox"/> Osteopetrosis (malignant infantile osteopetrosis) | <input type="checkbox"/> Metachromatic leukodystrophy (MLD) |
| <input type="checkbox"/> Lesch-Nyhan (HGPRT deficiency) | <input type="checkbox"/> Adrenoleukodystrophy (ALD) |
| <input type="checkbox"/> Neuronal ceroid – lipofuscinosis (Batten disease) | <input type="checkbox"/> Krabbe disease (globoid leukodystrophy) |
| <input type="checkbox"/> Mucopolysaccharidosis, NOS | <input type="checkbox"/> Neimann-Pick disease |
| <input type="checkbox"/> Hurler syndrome (IH) | <input type="checkbox"/> I-cell disease |
| <input type="checkbox"/> Scheie syndrome (IS) | <input type="checkbox"/> Wolman disease |
| <input type="checkbox"/> Hunter syndrome (II) | <input type="checkbox"/> Glucose storage disease |
| <input type="checkbox"/> Sanfilippo (III) | <input type="checkbox"/> Polysaccharide hydrolase abnormality, NOS |
| <input type="checkbox"/> Morquio (IV) | <input type="checkbox"/> Aspartyl glucosaminuria |
| <input type="checkbox"/> Maroteaux-Lamy (VI) | <input type="checkbox"/> Fucosidosis |
| <input type="checkbox"/> B-glucuronidase deficiency (VII) | <input type="checkbox"/> Mannosidosis |
| <input type="checkbox"/> Mucopolysaccharidosis (V) | <input type="checkbox"/> Inherited Disorders of Metabolism, NOS |
| <input type="checkbox"/> Mucopolysaccharidosis (V) | <input type="checkbox"/> Other inherited disorders of metabolism,
specify: _____ |
| <input type="checkbox"/> Mucopolysaccharidosis (V) | |
| <input type="checkbox"/> Mucopolysaccharidosis (V) | |
| <input type="checkbox"/> Mucopolysaccharidosis (V) | |
| <input type="checkbox"/> Gaucher disease | |

IMMUNE DEFICIENCIES

Classification:

- ADA deficiency severe combined immune deficiency (SCID)
- Absence of T and B cells SCID
- Absence of T, normal B cell SCID
- Omenn syndrome
- Reticular dysgenesis
- Bare lymphocyte syndrome
- SCID, NOS
- SCID other,
specify: _____
- Ataxia telangiectasia
- HIV infection
- Wiskott Aldrich syndrome
- DiGeorge anomaly
- Chronic granulomatous disease
- Chediak-Higashi syndrome
- Common variable immunodeficiency
- X-linked lymphoproliferative syndrome
- Leukocyte adhesion deficiencies
- Kostmann syndrome-congenital neutropenia
- Neutrophil actin deficiency
- Cartilage hair hypoplasia
- CD40 Ligand deficiency
- Immune Deficiencies, NOS
- Other immune deficiency,
specify: _____

Retired – Not for Data Submission



**Transplant Essential Data
First Report: 100 Days Post Transplant
Disease Classification Sheet 7**



PATIENT IDENTIFICATION	CENTER IDENTIFICATION
Hospital Unique Patient Number: _____	Center Identification Code: IBMTR/ABMTR _____

AUTOIMMUNE DISORDERS

Classification	Involved Organs/Clinical Problem(s) <i>(Check all that apply)</i>	Primary Reason(s) for Transplant <i>(Check all that apply)</i>	Miscellaneous Labs <i>(Check all that apply)</i>
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Other vasculitis

Churg-Strauss

Giant cell arteritis

Takayasu

Behçet's Syndrome

overlap necrotizing arteritis

other vasculitis, specify: _____

Arthritis

<input type="checkbox"/> Rheumatoid arthritis	<input type="checkbox"/> destructive arthritis <input type="checkbox"/> necrotizing vasculitis <input type="checkbox"/> eye (type: _____) <input type="checkbox"/> pulmonary <input type="checkbox"/> extra-articular (specify: _____) <input type="checkbox"/> other, specify: _____	<input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/>
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<input type="checkbox"/> Psoriatic arthritis/psoriasis	<input type="checkbox"/> destructive arthritis <input type="checkbox"/> psoriasis <input type="checkbox"/> other, specify: _____	<input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/>
--	--	--

Juvenile idiopathic arthritis: systemic (Stills disease)

Juvenile idiopathic arthritis: Oligoarticular

Juvenile idiopathic arthritis: Polyarticular

Juvenile idiopathic arthritis: Other, specify: _____

Other, arthritis, specify: _____

Multiple sclerosis

<input type="checkbox"/> Multiple sclerosis (MS)	<input type="checkbox"/> primary progressive <input type="checkbox"/> secondary progressive <input type="checkbox"/> relapsing/remitting <input type="checkbox"/> other specify: _____	<input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/>
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Other Neurological Autoimmune Disease

Myasthenia gravis

Other autoimmune neurological disorder, specify: _____

Hematological Autoimmune Disease

Idiopathic thrombocytopenic purpura (ITP)

Hemolytic anemia

Evan's syndrome

other autoimmune cytopenia, specify: _____

Bowel Disease

Crohn's disease

Ulcerative colitis

Other autoimmune bowel disorder, specify: _____

Retired - Not for Data Submission