

# **Bone Marrow Transplant**

**Adjudication Guideline** 

**Rule Category:** Medical

**Approved by:** Daman

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& Research

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### 1. Abstract

#### 1.1 For Members

A bone marrow transplant is a medical treatment that replaces bone marrow with healthy cells. The replacement cells can either come from your own body or from a donor.

A bone marrow transplant is also called a stem cell transplant or, more specifically, a hematopoietic stem cell transplant. Transplantation can be used to treat certain types of cancer, such as leukaemia, myeloma, and lymphoma, and other blood and immune system diseases that affect the bone marrow.

#### 1.2 For Medical Professionals

A stem cell or bone marrow transplant replaces damaged blood cells with healthy ones. It can be used to treat conditions affecting the blood cells, such as leukaemia and lymphoma.

## 2. Scope

The scope of this adjudication rule is to highlight the medical criteria, patient eligibility criteria and coverage details for Bone marrow transplant procedures for plans administered by Daman, subject to policy terms and conditions.

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## 3. Adjudication Policy

## 3.1 Eligibility / Coverage Criteria

## **Indications:**

#### Adults:

Malignant disorder	Non-malignant disorders
<ul> <li>Acute Myeloid Leukemia</li> <li>Acute Promyelocytic         Anemia     </li> <li>Acute Lymphoblastic         Leukemia     </li> <li>Chronic Myeloid Leukemia</li> <li>Myelodysplastic Syndrome</li> <li>Myelofibrosis and         Myelofibrosis and         Myelofibrotic disease     </li> <li>Plasma cell disorders</li> <li>Myeloma</li> <li>Light chain amyloidosis</li> <li>POEMS Syndrome</li> <li>Replase after autologous         transplant     </li> <li>Hodgkin's Lymphoma</li> <li>High Grade B Cell         Lymphoma     </li> <li>Primary Nervous system         lymphoma     </li> <li>Lymphoma</li> <li>Waldenström         macroglobinemia     </li> <li>Germ cell tumors and         Ewing's Sarcoma     </li> </ul>	<ul> <li>Severe Aplastic Anemia</li> <li>Sickle cell disease</li> <li>Hemophagocytic disorders</li> <li>Multiple and Systemic         Sclerosis</li> <li>Wiskott Aldrich Syndrome</li> </ul>

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### **Paediatric:**

Malignant disorder	Non-malignant disorders
<ul> <li>Acute Lymphoblastic Leukemia</li> <li>Chronic Myeloid Leukemia</li> <li>Myelodysplastic Syndrome</li> <li>T cell Non-Hodgkin's lymphoma</li> <li>Burkitt's Lymphoma</li> <li>Hodgkin's Lymphoma</li> <li>Ewing's Sarcoma</li> <li>Neuroblastoma</li> <li>Wilms Tumor</li> <li>Osteosarcoma</li> <li>Medulloblastoma</li> <li>Other Malignant Brain tumors</li> </ul>	<ul> <li>Severe Aplastic Anemia</li> <li>Sickle cell disease</li> <li>Thalassemia</li> <li>Hemophagocytic disorders</li> </ul>

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#### **Timings for referral:**

#### **Adult Leukemias and Myelodysplasia:**

<u>Acute Lymphoblastic Leukemia (ALL)</u>: Adult defined as greater than or equal to 40 years. High-resolution HLA typing is recommended at diagnosis for all patients. HSCT consultation should take place early after initial diagnosis for all patients with ALL, including:

- Primary induction failure
- Measurable (also known as minimal) residual disease after initial therapy.
- First relapse
- CR1
- CR2 and beyond, if not previously evaluated

<u>Myelodysplastic Syndromes (MDS)</u>: High-resolution HLA typing is recommended at diagnosis for all patients. Any intermediate or high IPSS or IPSS-R score Any MDS with poor prognostic features, including:

- Treatment-related MDS
- Refractory cytopenias
- Adverse cytogenetics and molecular features
- Transfusion dependence
- Failure of hypomethylating agents or chemotherapy
- Moderate to severe marrow fibrosis

#### Chronic Myeloid Leukemia (CML):

- Inadequate hematologic or cytogenetic/molecular response to tyrosine kinase inhibitor (TKI) therapies
- Disease progression
- Intolerance to TKI therapies
- Accelerated phase.
- Blast crisis (myeloid or lymphoid)
- T315l mutation

<u>Myeloproliferative Neoplasms (MPN):</u> High-resolution HLA typing is recommended at diagnosis for all patients. Intermediate- or high-risk disease, including:

- High-risk cytogenetics
- Poor initial response or at progression

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#### Myelofibrosis (MF):

- DIPSS Intermediate-2 (INT-2) and high-risk disease
- DIPSS Intermediate-1 (INT-1) with low platelet counts, refractory, red blood cell transfusion dependent, circulating blast cells > 2%, complex cytogenetics.
- High risk driver mutations (ASXL1, EZH2, TET2, IDH1, IDH2, SRSF2, and TP53) or triple negative (lack of a driver mutation such as JAK2, MPL, or CALR) should be considered in decision making.

#### Chronic Lymphocytic Leukemia (CLL):

Resistance or intolerance to BTK inhibitors and/or BCL2 inhibitors

### **Pediatric Acute Leukemias and Myelodysplasia:**

<u>Acute Myeloid Leukemia (AML):</u> High-resolution HLA typing is recommended at diagnosis for all patients. Early after initial diagnosis, all patients with AML including:

- Age < 2 years at diagnosis
- Primary induction failure
- Measurable (also known as minimal) residual disease after initial therapy.
- CR1 except favorable risk AML [defined as:t (8;21) (q22;q22.1); RUNX1- RUNX1T1, inv(16)(p13.1q22) or t(16;16)(p13.1;q22); CBFB-MYH11, mutated NPM1 without FLT3-ITD or with FLT3-ITD low, biallelic mutated CEBPA]
- Monosomy 5 or 7
- Treatment-related leukemia
- First relapse
- CR2 and beyond, if not previously evaluated

#### Acute Lymphoblastic Leukemia (ALL) (age < 15 years):

- Infant at diagnosis, unfavorable genetics, age < 3 months with any White Blood Cell Count (WBC), or < 6 months with WBC > 300,000 at presentation
- Primary induction failure
- Presence of measurable (also known as minimal) residual disease after initial therapy
- High/very high-risk CR1, including: o Philadelphia chromosome positive slow-TKI responders or with Ikaros Zinc Finger 1 (IKZF1) deletions; Philadelphia-like o Intrachromosomal amplification of chromosome 21 (iAMP21) o 11q23 rearrangement
- First relapse
- CR2 and beyond, if not previously evaluated
- Chimeric Antigen Receptor Therapy (CAR-T)



<u>Acute Lymphoblastic Leukemia (ALL):</u> (adolescent and young adults aged 15-39 years) High-resolution HLA typing is recommended at diagnosis for all patients.

- Primary induction failure
- Presence of measurable (also known minimal) residual disease after initial therapy
- High/very high-risk CR1, including: o Philadelphia chromosome positive or Philadelphia-like o iAMP21 o 11q23 rearrangement o B-cell with poor-risk cytogenetics
- First relapse
- CR2 and beyond, if not previously evaluated

Myelodysplastic Syndromes (MDS)

At diagnosis for all subtypes

Juvenile Myelomonocytic Leukemia (JMML)

At diagnosis

#### **Plasma Cell Disorders:**

- 1- Multiple Myeloma
  - At diagnosis
  - At progression and/or relapse
- 2- Light Chain Amyloidosis
  - At diagnosis
  - At progression and/or relapse
- 3- POEMS Syndrome (Osteosclerotic Myeloma)
  - At diagnosis

#### A. Lymphomas:

- Hodgkin Lymphoma
- Non-Hodgkin Lymphoma

#### **B.** Other Malignant Diseases

- Germ Cell Tumors
- Neuroblastoma
- Ewing Family of Tumors

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Medulloblastoma

#### **C. Non-Malignant Disorders**

- Immune Deficiency Diseases
- Inherited Metabolic Disorders
- Hemoglobinopathies
  - Sickle Cell Disease
  - o Transfusion-Dependent Thalassemia
- Hemophagocytic Lymphohistiocytosis (HLH)
- Severe Aplastic Anemia and Other Marrow Failure Syndromes
- Systemic Sclerosis
- Multiple Sclerosis (MS)

### 3.2 Requirements for Coverage

- The service codes for Bundle codes 22-01, 22-04, 22-05 and 22-08 are reported with Encounter type = 1, Bundle codes 22-02, 22-03, 22-06 and 22-07 are reported with Encounter type = 3
- Pre-authorization is required for all service codes and excluded medication mentioned within this adjudication at the start of the treatment.
- Drugs Plerixafor and defibrotide, or an equivalent should be administered under strict medical supervision at the medical facility, with SRVC code for "Short stay observation" as per medical necessity when requested between the bundles.
- Providers shall only claim the rate set for the respective service code and any excluded services.
- In line with DOH Cirular 106/2023, Department of Health has designated Abu Dhabi Stem Cell Center & Yas Clinic Khalifa City as the Center of Excellence (COE) for HSCT in the Emirate of Abu Dhabi
- The added inidication are to be followed as per "HSCT Indications and Timing of Referral for Adult and Pediatric" Guidelines published at DoH Website with the Reference No. DOH/GD/HFS/HSCT ITRAP/V1/2024
- For the services that are included in the service code providers are required to report the proper codes as activity line but keep charges at a value of zero as a prerequisite for reimbursement. Excluded services such as drugs/labs and other activities are defined in BMT reimbursement packages.

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### 3.3 Non-Coverage

- Missing services/benefits Reporting activity items included in each bundle is a
  prerequisite for payment. The claim has to be submitted after completing the
  bundle to allow reporting all expected and performed services.
- The BMT bundle codes are eligible to be billed for Thiqa and ABM policies.

## 3.4 Payment and Coding Rules

• Please apply regulator payment rules and regulations and relevant coding manuals for ICD, CPT, etc. Kindly code the ICD-10 and the CPT codes to the highest level of specificity.

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## **BUNDLES**

Code	<b>Code Description</b>	<b>Details</b>
22-01	Bundled reimbursement for Bone Marrow Pre- transplantation work-up (Autologous)	The bundle reimbursement for Bone Marrow pretransplantation work-up includes all procedures necessary for the pre-transplant work-up, extensive examination, Laboratory testing, Radiological and imaging analysis, Multidisciplinary team consultation.  Excluded Services from this bundle payment are:  • Medications plerixafor and defibrotide, or an equivalent, will be reimbursed in accordance with FDA label indication and require prior authorization.  • Any additional cost pertaining to complications (excluding Potentially Preventable Complications of BMT transplant procedure).  • List of CPT codes, see appendix 1, will be reimbursed outside the bundle based on medical necessity.
22-02	Bundled reimbursement for Preparation (Autologous)	<ul> <li>The bundle reimbursement for Bone Marrow preparation includes all procedures necessary for the preparation, Evaluation and Management, laboratory testing and radiological analysis,</li> <li>Mobilization and Apheresis procedures and patient specific conditioning protocol.</li> <li>Excluded Activities:         <ul> <li>Medications plerixafor and defibrotide, or an equivalent, will be reimbursed in accordance with FDA label indication and require prior authorization</li> <li>Any additional cost pertaining to complications (excluding Potentially Preventable Complications of BMT transplant procedure).</li> </ul> </li> </ul>
22-03	Bundled reimbursement for bone marrow transplant (Autologous)	The bundle reimbursement for Bone Marrow transplantation includes all inpatient procedures necessary for the Bone Marrow Transplantation to the day of discharge.  Excluded Activities:

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	1	
22-04	Bundled reimbursement for post-transplant follow-up (Autologous)	<ul> <li>Medications plerixafor and defibrotide, or an equivalent, will be reimbursed in accordance with FDA label indication and require prior authorization</li> <li>Any additional cost pertaining to complications (excluding Potentially Preventable Complications of BMT transplant procedure).</li> <li>The bundle reimbursement for Bone Marrow post-transplant follow-up includes all procedures necessary for the post-transplant follow-up (four months from discharge date), Evaluation and Management, laboratory testing and radiological analysis, medication up to 7 days, vaccination cost and cryopreservation for 6 months.</li> </ul>
		<ul> <li>Excluded Activities:</li> <li>Medications plerixafor and defibrotide, or an equivalent, will be reimbursed in accordance with FDA label indication and require prior authorization</li> <li>Any additional cost pertaining to complications (excluding Potentially Preventable Complications of BMT transplant procedure).</li> </ul>
22-05	Bundled reimbursement for Pre-transplantation work-up (Allogenic)	The bundle reimbursement for Bone Marrow Pretransplant work-up includes all procedures necessary for the pre-transplant work-up (Donor and recipient), extensive examination prior to transplantation, laboratory testing, radiological analysis, and multidisciplinary team consultation. Excluded Activities:  • Medications plerixafor and defibrotide, or an equivalent, will be reimbursed in accordance with FDA label indication and require prior authorization  • Any additional cost pertaining to complications (excluding Potentially Preventable Complications of BMT transplant procedure).
22-06	Bundled reimbursement for Preparation (Allogenic)	The bundle reimbursement for Bone Marrow preparation includes all procedures necessary for the preparation, Evaluation and Management, laboratory testing and radiological analysis, Mobilization and Apheresis procedures and patient specific conditioning protocol.  Excluded Activities:  Medications plerixafor and defibrotide, or an equivalent, will be reimbursed in accordance with FDA label indication and require prior authorization  Any additional cost pertaining to complications (excluding Potentially Preventable Complications of BMT transplant procedure).

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22-07	Bundled reimbursement for bone marrow transplant (Allogenic)	The bundle reimbursement for Bone Marrow transplantation includes all inpatient procedures necessary for the Bone Marrow Transplantation to the day of discharge.  Excluded Activities:  Medications plerixafor and defibrotide, or an equivalent, will be reimbursed in accordance with FDA label indication and require prior authorization  Any additional cost pertaining to complications (excluding Potentially Preventable Complications of BMT transplant procedure).
22-08	Bundled reimbursement for post-transplant follow-up (Allogenic)	The bundle reimbursement for Bone Marrow post- transplant includes all procedures necessary for the post-transplant follow-up (four months from discharge date), Evaluation and Management, laboratory testing and radiological analysis, discharge medication up to 7 days, vaccination cost and cryopreservation for 6 months. Excluded Activities:  Medications plerixafor and defibrotide, or an equivalent, will be reimbursed in accordance with FDA label indication and require prior authorization Any additional cost pertaining to complications (excluding Potentially Preventable Complications of BMT transplant procedure).

## 4. Denial Codes

Code	Code Description
CODE-012	Encounter type inconsistent with service(s) / diagnosis
MNEC-005	Service/supply may be appropriate, but too frequent
AUTH-001	Prior approval is required and was not obtained
PRCE-002	Payment is included in the allowance for another service
CLAI-016	Incorrect billing regime

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## 5. Appendices

## **5.1** Reference

## **Appendix 1: Excluded codes from the bundle:**

A maria side s	Codo	On a swin bin in
ACTIVITY	Code	Jescription
	88182	Flow cytometry, cell cycle or DNA analysis
		Flow cytometry, cell surface, cytoplasmic, or nuclear
	88184	marker, technical component only; first marker
		Flow cytometry, cell surface, cytoplasmic, or nuclear
		marker, technical component only; each additional
		marker (List separately in addition to code for the first
	88185	marker)
	88187	Flow cytometry, interpretation; 2 to 8 markers
FLOW	88188	Flow cytometry, interpretation; 9 to 15 markers
CYTOMETRY	88189	Flow cytometry, interpretation; 16 or more markers
		Flow cytometry, cell surface, cytoplasmic, or nuclear
	88184	marker, technical component only; first marker
		Flow cytometry, cell surface, cytoplasmic, or nuclear
		marker, technical component only; each additional
	88185	marker (List separately in addition to code for the first
DURACLONE T	x7	marker) ,
REG	88187	Flow cytometry, interpretation; 2 to 8 markers
		Flow cytometry, cell surface, cytoplasmic, or nuclear
	88184	marker, technical component only; first marker
MAXPAR		Flow cytometry, cell surface, cytoplasmic, or nuclear
DIRECT		marker, technical component only; each additional
IMMUNE	88185	marker (List separately in addition to code for the first
PROFILING	x29	marker)
ASSAY	88189	Flow cytometry, interpretation; 16 or more markers
MINIMAL		
RESIDUAL		
DISEASE	Code will	depend on target gene and methodology used.
STEM CELL		
KIT	86367	
TCR	86356	
ALFA/BETA	x2	
CD 10		Cell enumeration using immunologic selection and
CD 19		identification in fluid specimen (e.g., circulating tumor
SELECTION	86152	cells in blood);

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	86153	Cell enumeration using immunologic selection and identification in fluid specimen (e.g., circulating tumor cells in blood); physician interpretation and report, when required
	86152	Cell enumeration using immunologic selection and identification in fluid specimen (e.g., circulating tumor cells in blood);
CD 34+ SELECTION	86153	Cell enumeration using immunologic selection and identification in fluid specimen (e.g., circulating tumor cells in blood); physician interpretation and report, when required
BUSULFAN TEST	80375	
	81267	Chimerism (engraftment) analysis, post transplantation specimen (e.g., hematopoietic stem cell), includes comparison to previously performed baseline analyses; without cell selection
CHIMERISM	81268	Chimerism (engraftment) analysis, post transplantation specimen (e.g., hematopoietic stem cell), includes comparison to previously performed baseline analyses; with cell selection (eg, CD3, CD33), each cell typ

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## • Excluded Paediatric BMT services:

Mhala	1	
Whole Genome		
		Conomo (og unovalginod syndromo), coguence analysis
Sequencin		Genome (eg, unexplained syndrome); sequence analysis constitutional or heritable disorder or Whole Genome
g for	01/25	
· ·	81425	Sequencing for Donor
		Conomo (og unovalained constitutional or heritable disorder
•	01/175	
g for Dollor	01423	
Danal	96920	, , , , , , , , , , , , , , , , , , , ,
	00030	
	Q6Q21	
(FNA)	00031	,
Donor	86833	
	00032	
•		, , , , , , , , , , , , , , , , , , , ,
	86833	
(DSA)	00055	Serii quantitative paner (eg, titer), TEA class II
		Magnetic resonance (eg. proton) imaging abdomen; without
	74181	
	74101	
MRI T2*	75557	
	73337	
•		
	76377	
		process sections of an analysis and monitoring
,		
	81403	Molecular pathology procedure, Level
Cama alice sul	C	· · · · · · · · · · · · · · · · · · ·
_		a person accompanying a registered inpatient insured, of
oldy	Code 26	any age that is critically ill, or (2) parent accompanying a
		child under 10 years of age
Recipient Whole Genome Sequencin g for Donor  Panel- Reactive Antibodies (PRA)  Donor- Specific Antibodies (DSA)  MRI T2* for Liver and Heart in patients with iron overload RBC Genotyping in selected patients  Caregiver's Stay	81425 86830 86831 86832 86833 74181 75557 76377 81403 Service Code 26	any age that is critically ill, or (2) parent accompanying a

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- https://www.doh.gov.ae/-/media/2EEB17C27F4C48598C9F9328F415DF3B.ashx
- https://www.doh.gov.ae/-/media/Feature/shafifya/Prices/Adjudication-Rules/Addendum-36-to-DOH-claims-Adjudication-Rules -Bone-Marrow-Transaplant.ashx
- https://www.uptodate.com/contents/determining-eligibility-for-allogeneichematopoietic-celltransplantation?search=bone%20marrow%20transplant&source=search\_result &selectedTitle=2~150&usage type=default&display rank=2
- https://www.dynamed.com/procedure/hematopoietic-stem-cell-transplantationhsct-considerations
- https://www.cancer.org/cancer/managing-cancer/treatment-types/stem-celltransplant/process.html
- Relapse after Allogeneic Stem Cell Transplantation of Acute Myelogenous Leukemia and Myelodysplastic Syndrome and the Importance of Second Cellular Therapy - Transplantation and Cellular Therapy, Official Publication of the American Society for Transplantation and Cellular Therapy (astctjournal.org)

## 5.2 Revision History

Date	Change(s)
28.05.2024	Creation of Adjudication Guideline-External Instruction Template.

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