



# The EBMT activity survey report 2017: a focus on allogeneic HCT for nonmalignant indications and on the use of non-HCT cell therapies

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## Abstract

Hematopoietic cell transplantation (HCT) is widely used for acquired and congenital disorders of the hematopoietic system. Number of transplants done in Europe and associated countries continues to rise with 45,418 HCT in 41,100 patients [(17,155 allogeneic (42%) and 23,945 autologous (58%)] reported by 683 centers in 50 countries in 2017. Main indications were myeloid malignancies 10,147 (25%; 96% allogeneic), lymphoid malignancies 26,488 (64%; 19% allogeneic), solid tumors 1,607 (3.9%; 2% allogeneic), and nonmalignant disorders 2,667 (7%; 81% allogeneic). Trends in donor choice seen before continue, with growing numbers of haploidentical HCT and decreasing use of cord blood. Of interest is that after many years of continued growth, the number of patients receiving an allogeneic HCT for marrow failure is decreasing slightly ( $p < 0.001$ ). Such a change may be explained by the use of thrombopoietin analogs in aplastic anemia patients. Other nonmalignant indications, however continue to grow, most importantly HCT for hemoglobinopathies by 36%, equally for thalassemias and sickle cell disease. Non-HCT cell therapies have increased by 28% since 2015 and genetically modified T cells is type of cell therapy with the fastest growth. These annual reports reflect current activity and trends and are useful for health-care planning.

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## Introduction

Hematopoietic cell transplantation (HCT) is an established procedure for many inherited or acquired disorders of the hematopoietic system, whether benign or neoplastic,

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including those of the immune system, and as enzyme replacement in metabolic disorders [1–4]. The activity survey of the European Society of Blood and Marrow Transplantation (EBMT), describing the status of HCT in Europe and affiliated countries, has become an instrument to observe trends and to monitor changes in technology [5–14]. The survey using a standardized structure captures the numbers of HCT from highly committed participating teams, divided by indication, donor type, and stem cell source. More recently, the survey has included information on novel cell therapies with hematopoietic stem cells for non-hematopoietic use, and the use of non-hematopoietic stem and progenitor cells. This coincides with the interest of the World Health Organization (WHO; [www.who.org](http://www.who.org)) in cell and tissue transplants, and further stresses the need for adequate and timely information [15]. The analysis of the survey data spanning 27 years and amassing data on almost 708,000 transplants in over 630,000 patients has shown a continued and constant increase in the annual numbers of HCT and transplant rates for both allogeneic and autologous HCT. This report is based on the 2017 survey data. In addition to transplant rates and indications, it focuses on more recent changes possibly related to development of alternative therapies.

## Patients and methods

### Data collection and validation

Participating teams were invited to report their data for 2017 as listed in Table 1. The survey allows the possibility to report additional information on the numbers of subsequent transplants performed because of relapse, rejection, or those that are part of a planned sequential transplant protocol.

Additional information on the numbers of donor lymphocyte infusions (DLIs), reduced intensity HCT, and the numbers of pediatric HCT is also collected. Quality control measures included several independent systems: confirmation of validity of the entered data by the reporting team, selective comparison of the survey data with MED-A data sets in the EBMT Registry database, and crosschecking with the National Registries.

### Teams

A total of 720 centers from 50 countries were contacted for the 2017 survey (40 European and 10 affiliated countries); of which 683 teams reported. This corresponds to a 95% return rate and includes 83% EBMT members and 17% non-EBMT members. Thirty-seven active teams failed to report in 2017. Reporting teams are listed in the online appendix in alphabetical order by country, city, and EBMT

center code, with their reported numbers of first and total HCT, and of first allogeneic and autologous HCT as supplementary material. The WHO regional office definitions were used to classify countries as European or non-European. Nine neighboring non-European countries participated in the 2017 EBMT survey: Algeria, Iraq, Iran, Israel, Jordan, Lebanon, Saudi Arabia, South Africa, and Tunisia. Their data, 3101 HCT in 2948 patients, from 30 actively transplanting teams make up 6.9% of the total data set and are included in all analyses.

### Patient and transplant numbers

Wherever appropriate, patient numbers corresponding to the number of patients receiving a first transplant, and transplant numbers reflecting the total number of transplants performed are listed. The term sibling donor includes human leukocyte antigen (HLA) identical siblings and twins but not siblings with HLA mismatches. Unrelated donor transplants include HCT from matched or mismatched unrelated donors with peripheral blood and marrow as a stem cell source but not cord blood HCT. Haploidentical transplants are being described as any family member with 2 or more loci mismatch within the loci HLA-A, -B, -C, -DRB1, and -DQB1 in graft versus host (GvH) and/or host versus graft direction. Other family member donors are those related donors that are mismatched to a lesser degree than a full haplotype. Additional non-first transplants may include multiple transplants defined as subsequent transplants within a planned double or triple autologous or allogeneic transplant protocol, and re-transplants (autologous or allogeneic) defined as unplanned HCT for rejection or relapse after a previous HCT.

### Transplant rates

Transplant rates, defined as the total number of HCT per 10 million inhabitants, were computed for each country without adjustments for patients who crossed borders and received their HCT in a foreign country. Population numbers for the European countries in 2017 were obtained from Eurostats (<http://epp.eurostat.ec.europa.eu>) and from the US census bureau database for the non-European countries (<https://www.census.gov/data-tools>).

### Analysis

Wherever appropriate, the absolute numbers of transplanted patients, transplants, or transplant rates are shown for specific countries, indications, or transplant techniques. Myeloid malignancies include acute myeloid leukemia (AML), myelodysplastic or myelodysplastic/myeloproliferative neoplasm (MDS or MD/MPN overlap), myeloproliferative

**Table 1** Numbers of HCT in Europe 2017 by indication, donor type, and stem cell source

Transplant activity 2017																			
No. of patients																			
	Allogeneic																		
	Family							Unrelated											
	HLA-id		Twin		Haplo $\geq 2$ MM			Other family			BM		BM+		Total				
	BM	PB	Cord	All	BM	PB	Cord	BM	PB	Cord	BM	PB	Cord	BM	Only	PB	Cord	Allo	Auto
<b>Myeloid malignancies</b>	380	2532	0	11	319	946	10	85	1	527	4804	157	7	368	0	9772	375	10,147	
Acute myeloid leukemia	267	1802	0	7	235	716	7	70	1	350	3101	120	7	353	0	6676	360	7036	
First complete remission	195	1123		7	116	330	5	40		218	1655	64	4	289		3753	293	4046	
Not first complete remission	50	474			80	270	1	20	1	96	919	39	3	56		1950	59	2009	
AML therapy-related	8	53			14	46		3		15	130	8		2		277	2	279	
AML from MDS/MPN	14	152			25	70	1	7		21	397	9		6		696	6	702	
Chronic myeloid leukemia	35	84	0	0	6	22	0	2	0	25	161	0	0	0	0	335	0	335	
Chronic phase	22	35			2	6		2		14	55					136	0	136	
Not chronic phase	13	49			4	16				11	106					199	0	199	
MDS or MD/MPN overlap	66	478		4	64	169	3	9		132	1097	34		5		2056	5	2061	
MPN	12	168			14	39		4		20	445	3		10		705	10	715	
<b>Lymphoid malignancies</b>	297	1434	6	7	193	579	11	47	1	408	1963	69	29	21,444	0	5015	21,473	26,488	
Acute lymphatic leukemia	245	731	6	3	89	285	9	38	1	340	876	58	5	85	0	2681	90	2771	
First complete remission	139	516	2	1	49	130	7	30		158	590	30	4	78		1652	82	1734	
Not first complete remission	106	215	4	2	40	155	2	8	1	182	286	28	1	7		1029	8	1037	
Chronic lymphocytic leukemia	4	50			6	20		3		5	141	1		9		230	9	239	
Plasma cell disorders—MM	5	122	1	1	12	17		2		8	192	1	4	12,353		360	12,357	12,717	
Plasma cell disorders—other	6					2				1	16			335		25	335	360	
Hodgkin lymphoma	16	131	0	1	30	99		1		11	153	2	11	2141		444	2152	2596	
Non-Hodgkin lymphoma	27	394		2	56	156	2	3		43	585	7	9	6521		1275	6530	7805	
<b>Solid tumors</b>	3	1	0	0	10	20	0	0	0	0	2	0	48	1521	2	36	1571	1607	
Neuroblastoma	2				8	14				2			22	513	2	26	537	563	
Soft tissue sarcoma/Ewing	1				1	6							4	225		8	229	237	
Germinal tumors													2	370		0	372	372	
Breast cancer														13		0	13	13	
Other solid tumors	1				1								20	400		2	420	422	
<b>Nonmalignant disorders</b>	668	246	27	3	96	167	61	61	2	450	325	67	15	479	0	2173	494	2667	

**Table 1** (continued)

Transplant activity 2017																
No. of patients																
	Allogeneic										Autologous				Total	
	Family					Unrelated					BM	Only	PB	BM+	Cord	Allo
HLA-id	BM	PB	Twin		Haplo $\geq$ 2MM		Other family		BM	PB						
			Cord	All	BM	PB	BM	PB			BM	PB	BM	PB	Cord	
Bone marrow failure—SAA	176	107	3	28	32	3	6	3	6	120	100	8	1	583	1	584
Bone marrow failure—other	54	23	3	8	14	7	10	62	35	3	3	219	0	219	0	219
Thalassemia	178	57	15	5	18	22	15	1	58	49	2	2	4	420	6	426
Sickle cell disease	110	41	6	11	14	6	2	19	5	1	1	215	1	215	1	216
Primary immune deficiencies	125	10	3	33	81	15	19	144	102	21	6	7	554	13	567	
Inh. disorders of metabolism	23	3	8	6	7	7	7	44	29	32	5	5	159	10	169	
Autoimmune disease	2	5	3	2	1	2	1	3	5	1	462	23	463	486		
Others	28	17	11	11	20	5	2	31	39	6	32	159	32	191		
<b>Total patients</b>	1376	4230	33	21	629	1732	87	195	4	1416	7133	299	99	23,844	2	17,155
Re/additional transplants	44	181	79	249	2	17	1	76	446	31	13	3178	1	1126	3192	4318
<b>Total transplants</b>	1420	4411	33	21	708	1981	89	212	5	1492	7579	330	112	27,022	3	18,281

*HCT* hematopoietic stem cell transplantation, *HLA-id* human leukocyte antigen identical, *BM* bone marrow, *MM* multiple myeloma, *AML* acute myeloid leukemia, *MDS* myelodysplastic syndrome, *MD/MPN* myelodysplastic/myeloproliferative neoplasm, *SAA* severe aplastic anemia, *PB* peripheral blood

neoplasm (MPN), and chronic myeloid leukemia (CML). Lymphoid malignancies include acute lymphocytic leukemia (ALL), chronic lymphocytic leukemia (CLL), Hodgkin lymphoma (HL), non-Hodgkin lymphoma (NHL), and plasma cell disorders (PCD including multiple myeloma (MM) and others). The nonmalignant disorders include bone marrow failure (BMF including severe aplastic anemia (SAA) and others), thalassemia and sickle cell disease (HG), primary immune disease, inherited disease of metabolism, and autoimmune disease (AID). Others include histiocytosis and rare disorders not included in the above. BMF syndromes were grouped until 2004, at which time they were separated into SAA and other marrow failure. Trends shown over time include changes in absolute number of patients transplanted from 1990 to 2017. We use graphical representation to indicate changes over time. We consider an increase or decrease of  $\pm 5\%$  as a non-relevant change. To confirm trends we used SPSS to automatically fit the best exponentially smoothed, autoregressive integrated moving average model without any further pre-specification. To detect possible deviations from trends, we calculated the observed and predicted counts as well as the 95% confidence intervals (CIs).

## Results

### Participating teams in 2017

Of the 683 teams, 430 (63%) performed both allogeneic and autologous transplants; 236 (34%) restricted their activity to autologous HCT, and 17 (2%) to allogeneic transplants only. Of the 683 active centers, 131 (19%) centers performed transplants on both adult and pediatric patients. An additional 115 (17%) centers were dedicated pediatric transplant centers and 437 (64%) centers performed transplants on adults only. Thirty-seven teams failed to report in 2017, which when compared to previously reported data by these teams, accounts for approximately 630 non-reported HCT.

### Numbers of patients and transplants

In 2017, 45,418 transplants were reported in 41,100 patients (first transplant); of these, 18,281 HCT (40%) were allogeneic and 27,137 (60%) autologous (Table 1). When compared with 2016 the total number of transplants increased by 4.1% (3.6% allogeneic HCT and 4.4% autologous HCT) [16]. In patients receiving their first transplant in 2017, the increase was 3.9% for allogeneic HCT and 5.0% for autologous HCT. In addition, there were 4318 second or subsequent transplants, 1126 being allogeneic, mainly to treat relapse or graft failure and 3192

autologous, the majority of which were most likely part of multiple transplant procedures such as either tandem procedures, or as salvage autologous transplants for plasma cell disorders. Furthermore, 986 of the allogeneic HCTs were reported as being given after a previous autologous HCT, and were mainly for lymphoma or plasma cell disorders.

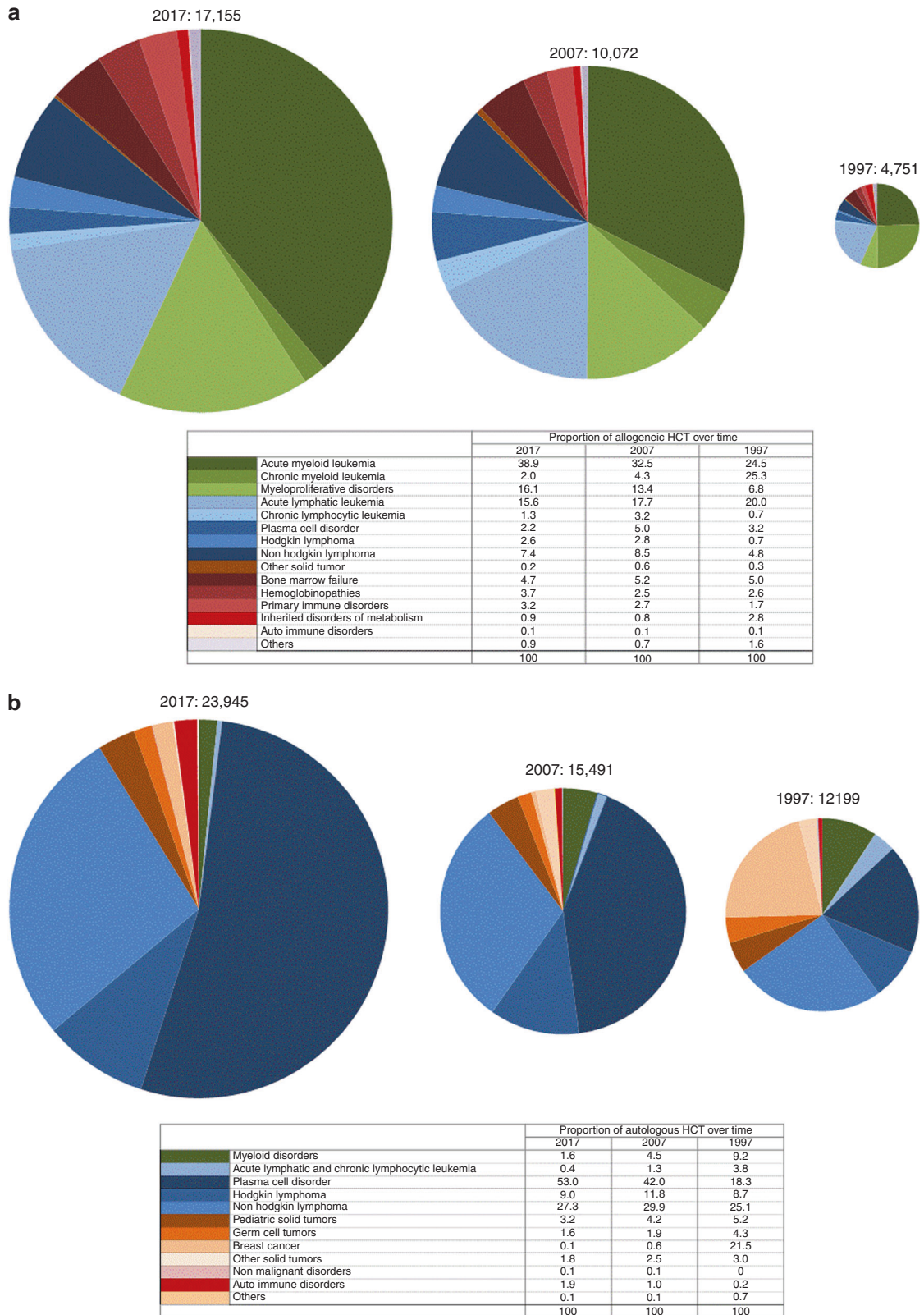
The total number of patients transplanted under the age of 18 in both dedicated and joint adult-pediatric units was 5056 (3725 allogeneic and 1331 autologous). This is an increase of 7.8% (5.1% allogeneic and 16.2% autologous) when compared to 2016, where 4690 HCT (3545 allogeneic and 1145 autologous) were reported. From the 5056 patients, 3983 transplants in 3505 patients (2701 allogeneic, 77%; and 804 autologous, 23%) were performed in dedicated pediatric centers. Due to limitations in the design of the survey, indication for HCT can only be observed in the dedicated pediatric centers. The main indications were myeloid malignancies 588 (17%; 99% allogeneic), lymphoid malignancies 851 (24%; 86% allogeneic), solid tumors 696 (20%; 4% allogeneic), and nonmalignant disorders 1370 (39%; 99% allogeneic). When compared to 2016, increases in allogeneic HCT were seen in sickle cell disease (55.8%), thalassemia (40.4%), ALL (12.8%), and AML (9.6%). For autologous HCT the main increase was seen in HL (30%) and pediatric solid tumors (17.5%).

### Indications and trends

Indications for HCT in 2017 are listed in detail in Table 1. Main indications for HCT were myeloid malignancies (AML, CML, MDS or MD/MPN overlap, and MPN): 10,147 (25% of total; 96% of which were allogeneic); lymphoid malignancies (ALL, CLL, HL, NHL, and PCD): 26,488 (64%; 19% allogeneic); solid tumors: 1607 (4%; 2% allogeneic); nonmalignant disorders: 2667 (7%; 81% allogeneic), and others: 191 (0.5%). As seen in previous years, the majority of HCT for lymphoid malignancies were autologous, while most transplants for myeloid malignancies were performed using stem cells from allogeneic donors. Autologous HCT for nonmalignant disorders predominantly include patients with autoimmune disorders.

Figure 1a, b shows distribution of disease indications for allogeneic (Fig. 1a) and autologous (Fig. 1b) HCT as a pie graph. For comparative analyses, the 2017 data are juxtaposed to the analogous data from the years 2007 and 1997 using the size of the pie graph to represent the absolute number of transplanted patients and the pie slices for proportions of specific indications [9, 17]. The 2017 data are used to depict the pie graph with the 100% size for allogeneic and autologous HCT respectively, pie size for 2007 and 1997 are proportionally smaller.

As shown (Fig. 1a), the number of patients treated with an allogeneic HCT has increased from 4751 to 17,155 over



**Fig. 1** Relative proportion of disease indications for hematopoietic cell transplantation (HCT) in Europe between the years 1997, 2007, and 2017. **a** Relative proportion of allogeneic HCT. **b** Relative proportion of autologous HCT

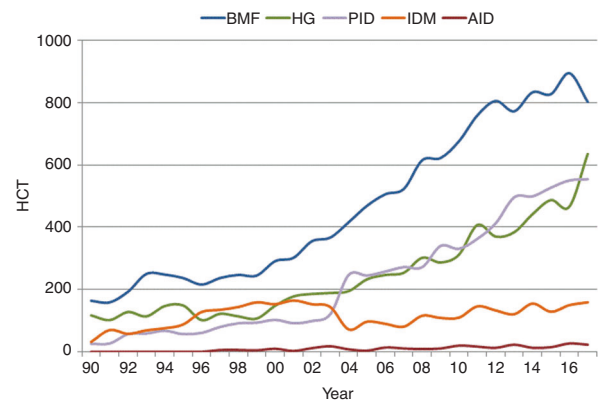
20 years. For allogeneic HCT, AML is the most frequent indication (39%), of these 56% were for patients in CR1, 29% for patients with more advanced disease, and 15% for patients with transformed AML, either therapy-related or from MDS/MPN. The 20-year trend confirms the decline in HCT for CML from 25.3 to 1.9%, whereas AML, MDS, and MPN have increased substantially. The proportion of ALL transplants has decreased only slightly, other lymphoid malignancies have rather increased. Compared to 2016, there were increases in allogeneic HCT for AML by 6.3%, and MDS and MPN by 9.5% and 9.0%, respectively. HL increased by 13.8%, and thalassemia and sickle cell disease by 27.7% and 56.9%, respectively. Decreases were seen for CML by 12.8%, PCD by 16.9%, CLL by 16.4%, and BMF by 10.3%.

In Fig. 1b for autologous HCT, the number of patients has increased from 12,199 to 23,945 (an increase of 196% over 20 years as compared to allogeneic HCT where the increase is 360% over 20 years). In autologous HCT, PCD is the most frequent indication (53%), of these 51.5% were for MM. Compared to 2016, there were increases in autologous HCT for AID by 31.9%, neuroblastoma by 12.1%, soft tissue and Ewing sarcoma by 20.5%, and MM by 7.0%. In autologous HCT, myeloma has become the dominant indication increasing from 18 to 53% over the past 20 years. Proportions of HCTs for HL and NHL have remained stable. Autologous HCT for AML has decreased from 13 to 1.72%. Already well described is the change in autologous HCT for solid tumor where the decrease in use of this technology in breast cancer dominates the graphical representation.

Within allogeneic HCT 7166 were performed using non-meloablative conditioning in 2017. This comprises 39% of all allogeneic HCT, and has remained stable over the last 9 years. European maps depicting transplant rates by country are provided in the supplementary figures section (supplementary Figure 1a for allogeneic, 1b for autologous)

## Nonmalignant disorders

Figure 2 depicts allogeneic HCT numbers for nonmalignant disorders. Since 2016, an increase in numbers of HCT for hemoglobinopathies (sickle cell disease by 56.9% and thalassemia by 27.7%) and inherited disorders of metabolism by 6.0% has been observed with stable numbers for primary immunodeficiencies and AID. The increase for sickle cell disease HCT is clearly (>95% CI for the prediction model) significant, for thalassemia of borderline significance. A decline in the numbers of HCT is seen for BMF by 10.3%. This decline is still borderline within the lower confidence limits of the prediction model but after many years of increasing HCT numbers for marrow failure this is the first time that numbers actually do decline. This is likely due to

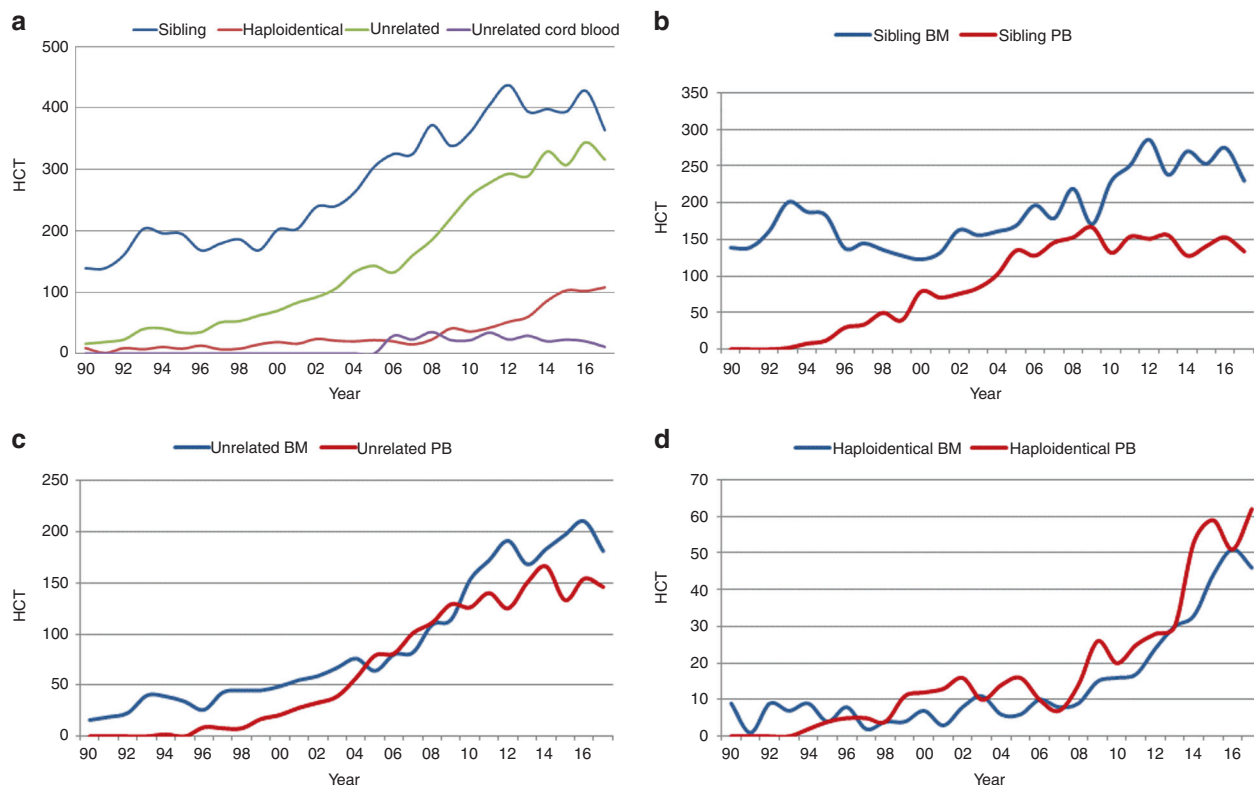


**Fig. 2** Trends in the absolute numbers of allogeneic hematopoietic cell transplantation (HCT) for nonmalignant disorders in Europe 1990–2017

the use of thrombopoietin analogs recently approved to treat refractory aplastic anemia.

Autologous HCT (not shown in the figure) is predominantly used for the treatment of AIDs, where a continuous increase has been seen for the last 12 years. Compared to 2016, the increase is 32%. This recent increase is however mainly due to one non-EBMT member center in Eastern Europe reporting large numbers of autologous HCT for multiple sclerosis.

We now focus on marrow failure, where Fig. 3a shows recent changes in practice for BMF: types of donors used indicate that the use of sibling and unrelated donor transplants are decreasing by 15% and 8% respectively. Whilst the numbers of haploidentical HCT for marrow failure remain stable, the number of cord blood transplants performed is very low. In marrow failure, the use of marrow is generally recommended over the use of peripheral blood as a stem cell source, mainly because of the absence of benefit of potentially higher alloreactivity conferred by the higher number of T lymphocytes in the peripheral blood graft [18, 19]. Figure 3b shows the changes in stem cell source for sibling donor HCT, 3c for unrelated donor HCT, and 3d for haploidentical HCT. Marrow is the preferred source of stem cells in sibling donor HCT but is less so in unrelated donor HCT. In haploidentical HCT for marrow failure, slightly more peripheral blood HCT is performed when compared to marrow. Marrow failure syndromes are rare diseases and the experience of teams doing these transplants is evidently limited. Table 2 shows the number of transplants for BMF by type of donor and the number of teams performing these transplants. Of note, e.g., in haploidentical HCT the 108 patients receiving transplants were treated in 64 different centers, the median number of haploidentical HCT per team is 1 (range 1–9). Over 100 patients received haploidentical HCTs for marrow failure in spite of a very limited number of published series [20–23] in the field. European maps depicting transplant rates by country



**Fig. 3** Changes in donor choice and stem cell source for bone marrow failure in Europe 1990–2017. **a** Donor choice. **b** Human leukocyte antigen identical sibling hematopoietic cell transplantation (HCT) by

bone marrow (BM) or peripheral blood (PB). **c** Unrelated HCT by BM or PB. **d** Haploidentical HCT by BM or PB

**Table 2** BMF data by center in 2017

	HCT, <i>N</i>	Teams, <i>N</i>	Median and range
Sibling	363	168	1 (1–20)
Unrelated	317	163	1 (1–8)
Haploidentical	108	64	1 (1–9)
Cord blood	14	14	1 (1)
Total	802		

BMF bone marrow failure, HCT hematopoietic cell transplantation

are provided in the Supplementary Figures section. Supplementary Figure 1a shows allogeneic transplant rates per country for all indications. Whereas the map depicting allogeneic HCT reflects income class of countries, wealthier countries usually having higher rates than middle-income countries, transplant rates for marrow failure look somewhat different. The countries with the highest reported rates for marrow failure transplants include Lithuania, Norway, Ireland, Turkey, and Israel.

## Cell therapy

Table 3 shows cell therapies performed in EBMT centers in 2017. There were 2825 patients receiving DLIs, which is a

slight decrease of 1.9% since 2016 where 2879 DLIs were reported. The majority of DLIs were given for relapse (1284) and graft enhancement (661). Other reasons being for residual disease (422) and “per protocol” (458). A total of 1202 patients received other forms of cell therapy, most commonly mesenchymal stromal cells ( $n = 557$ ; 91% allogeneic), mainly to treat GvH disease (GvHD). Expanded/selected T lymphocytes ( $n = 179$ ; 100% allogeneic) and genetically modified T cells ( $n = 151$ ; 50% allogeneic) were the second most frequently reported cell therapy. The area with the highest growth is genetically modified T cells. The main indications for cell therapies (excluding DLI) being GvHD or graft enhancement ( $n = 688$ ; 95% allogeneic), malignancy ( $n = 253$ ; 50% allogeneic), infections ( $n = 124$ ; 100% allogeneic), AID ( $n = 61$ ; 43% allogeneic), and genetic disease ( $n = 14$ ; 7% allogeneic). The number of cell therapies have increased by 27.6% since 2015, when they were first introduced into the survey (2015, HCT = 942; and 2017, HCT = 1202) [24, 25].

## Discussion

The EBMT activity survey has been conducted annually since 1990 [7]. The 2010 survey reported for the first time



**Table 3** Non-HCT cellular therapies using manipulated cells in 2017

Number of patients	DLI	MSC		NK cells		Selected/ expanded T cells or CIK		TREGS		Geneti- cally modified T cells		Dendritic cells		Expanded CD34+ cells		Geneti- cally modified CD34+ cells		Other		
		Allo	Auto	Allo	Auto	Allo	Auto	Allo	Auto	Allo	Auto	Allo	Auto	Allo	Auto	Allo	Auto	Allo	Auto	
GvHD		413	5			8		36											13	
Graft enhancement	661	55	18	1		14		5		57		4	6		1			44	8	
Autoimmune disease		6	14	19									1						21	
Genetic disease		1															13			
Infection		3				113													8	
Malignancy				6		42		24	8	16	78	5	35	2				32	5	
DLI for residual disease	422																			
DLI for relapse	1284																			
DLI per protocol	458																			
Regenerative medicine		31	11	1		2								2				4	11	
Total		2825	509	48	27	0	179	0	65	8	73	78	5	39	11	0	1	13	101	45

more than 30,000 patients transplanted in a given year, >40,000 transplants in 2014, and >45,000 in 2017. As before, autologous HCT continues to expand (4.4%) and at a higher rate than allogeneic HCT (3.6%). In allogeneic HCT, some indications continue to increase but not others. Impressive changes are observed in nonmalignant disorders, there were considerable increases of allogeneic HCT for hemoglobinopathies, including the thalassemias and sickle cell disease. The majority of which is with matched sibling donors and among sibling donor HCT, 73% is with marrow as a stem cell source. For the first time we describe a lower number of transplants for marrow failure. Such a decrease coincides with the approval of the thrombopoietin mimetic eltrombopag for relapsed refractory aplastic anemia [21]. Next to aplastic anemia, decreased use of allogeneic HCT was also seen in CLL and PCD by almost 20%, which is best explained by the competition of new drugs being developed in these fields. Therefore, it seems that the use of allogeneic HCT is modified by availability of drugs to treat the disease. Furthermore, such shifts in use of transplant technology occur rather rapidly.

In autologous HCT, the only striking increase is in HCT for AID, which has increased by almost a third since 2016, mostly related to autologous HCT to treat multiple sclerosis [26].

We have generated pie graphs showing number of transplants as pie size and proportion of indications by segment over the last 20 years. Of interest is that some of the proportions have remained stable over the years, e.g., allogeneic HCT for ALL. While, as described before, CML has decreased dramatically as a main indication. It appears to have been replaced in much greater numbers, but in

similar proportions, by MDS and more recently MPN and AML. Whether this shift in indications is biased by the availability of treatment resources, or follows some other logic, is difficult to discern.

For marrow failure, interesting aspects include the resolute resistance in the use of peripheral blood over marrow. This is seen more so with unrelated donor HCT than with sibling donor transplantation, and may possibly be related to the unwillingness of some unrelated donors to undergo marrow harvest. The use of haploidentical HCT for marrow failure is not substantiated by much available data, and the reason for this use of technology may be due to the general popularity of haploidentical HCT in recent years.

The section on cell therapies shows the gradual increasing use of mesenchymal stromal cells, most commonly to treat GvHD. There is a large increase in the use of gene modified T cells from 36 in 2016 to 151 in 2017. Whether these numbers reflect all patients receiving gene-modified T cells in Europe during the year is not entirely clear, as some centers may not report patients treated in clinical trials. It is our goal to truly reflect the activity of this rapidly growing field within Europe. As CAR-T cells [27] have become commercially available in 2018, it is most important that EBMT centers continue the well-established practice of transparently sharing data on activity of advanced therapy medicinal products manufactured from hematopoietic cells used and on the outcome of patients. These data reflect the ability of the EBMT to capture data not only on HCT but also on non-HCT cell therapies in this field of rapidly emerging treatment options.

The annual activity survey of the EBMT reflects current activity and trends in the field of transplant technology. It is valuable for the dissemination of the most recent information on indications, donor, and stem cell usage, which can ultimately be beneficial in health care planning.

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## Compliance with ethical standards

**Conflict of interest** The authors declare that they have no conflict of interest.

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## References

- Copelan EA. Hematopoietic stem-cell transplantation. *N Engl J Med.* 2006;354:1813–26.
- Appelbaum FR. Hematopoietic-cell transplantation at 50. *N Engl J Med.* 2007;357:1472–5.
- Sureda A, Bader P, Cesaro S, Dreger P, Duarte RF, Dufour C, et al. Indications for allo- and auto-SCT for haematological diseases, solid tumours and immune disorders: current practice in Europe, 2015. *Bone Marrow Transplant.* 2015;50:1037–56.
- Gratwohl A, Baldomero H, Aljurf M, Pasquini MC, Bouzas LF, Yoshimi A, et al. Hematopoietic stem cell transplantation: a global perspective. *JAMA.* 2010;303:1617–24.
- Gratwohl A, Pasquini MC, Aljurf M, Atsuta Y, Baldomero H, Foeken L, et al. One million haemopoietic stem-cell transplants: a retrospective observational study. *Lancet Haematol.* 2015;ume 2 (No. 3):e91–100.
- Gratwohl A, Baldomero H, Schwendener A, Gratwohl M, Apperley J, Frauendorfer K, et al. The EBMT activity survey 2008 impact of team size, team density and new trends. *Bone Marrow Transplant.* 2011;46:174–91.
- Gratwohl A. Bone marrow transplantation activity in Europe 1990. Report from the European Group for Bone Marrow Transplantation (EBMT). *Bone Marrow Transplant.* 1991;8:197–201.
- Gratwohl A, Baldomero H, Horisberger B, Schmid C, Passweg J, Urbano-Ispizua A. Accreditation Committee of the European Group for Blood and Marrow Transplantation (EBMT). Current trends in haematopoietic stem cell transplantation in Europe. *Blood.* 2002;100:2374–86.
- Gratwohl A, Baldomero H, Schwendener A, Rocha V, Apperley J, Frauendorfer K, et al. The EBMT activity survey 2007 with focus on allogeneic HSCT for AML and novel cellular therapies. *Bone Marrow Transplant.* 2009;43:275–91.
- Gratwohl A, Schwendener A, Baldomero H, Gratwohl M, Apperley J, Niederwieser D, et al. Changes in use of hematopoietic stem cell transplantation; a model for diffusion of medical technology. *Haematologica.* 2010;95:637–43.
- Passweg JR, Baldomero H, Peters C, Gaspar HB, Cesaro S, Dreger P, et al. Hematopoietic SCT in Europe: data and trends in 2012 with special consideration of pediatric transplantation. *Bone Marrow Transplant.* 2014;49:744–50.
- Passweg JR, Baldomero H, Bader P, Bonini C, Cesaro S, Dreger P, et al. Hematopoietic stem cell transplantation in Europe 2014: more than 40 000 transplants annually. *Bone Marrow Transplant.* 2016;51:786–92.
- Passweg JR, Baldomero H, Bader P, Bonini C, Duarte RF, Dufour C, et al. Use of haploidentical stem cell transplantation continues to increase; the 2015 European Society for Blood and Marrow Transplant activity survey report. *Bone Marrow Transplant.* 2017;52:811–7.
- Niederwieser D, Baldomero H, Szer J, Gratwohl M, Aljurf M, Atsuta Y, et al. Hematopoietic stem cell transplantation activity worldwide in 2012 and a SWOT analysis of the Worldwide Network for Blood and Marrow Transplantation Group including the global survey. *Bone Marrow Transplant.* 2016;51:778–85.
- World Health Organisation, WHO (<http://www.who.int/topics/transplantation/en/>).
- Passweg JR, Baldomero H, Bader P, Basak GW, Bonini C, Duarte R, et al. Is the use of unrelated donor transplantation leveling off in Europe? The 2016 European Society for Blood and Marrow Transplant activity survey report. *Bone Marrow Transplant.* 2018;53:1139–48.
- Gratwohl A, Passweg J, Baldomero H, Hermans J. Blood and marrow transplantation activity in Europe 1997. European Group for Blood and Marrow Transplantation (EBMT). *Bone Marrow Transplant.* 1999;3:231–45.
- Schrezenmeier H, Passweg JR, Marsh JC, Bacigalupo A, Brede-son CN, Bullorsky E, et al. Worse outcome and more chronic GVHD with peripheral blood progenitor cells than bone marrow in HLA-matched sibling donor transplants for young patients with severe acquired aplastic anemia. *Blood.* 2007;110:1397–400.
- Eapen M, Le Rademacher J, Antin JH, Champlin RE, Carreras J, Fay J, et al. Effect of stem cell source on outcomes after unrelated donor transplantation in severe aplastic anemia. *Blood.* 2011;118:2618–21.
- Clay J, Kulasekararaj AG, Potter V, Grimaldi F, McLoman D, Raj K, et al. Nonmyeloablative peripheral blood haploidentical stem cell transplantation for refractory severe aplastic anemia. *Bone Marrow Transplant.* 2014;20:1711–6.
- Young NS. Aplastic anemia. *N Engl J Med.* 2018;379:1643–56.

22. Esteves I, Bonfim C, Pasquini R, Funke V, Pereira NF, Rocha V, et al. Haploidentical BMT and post-transplant Cy for severe aplastic anemia: a multicenter retrospective study. *Bone Marrow Transplant.* 2015;50:685–9.
23. DeZern AE, Zahurak M, Symons H, Cooke K, Jones RJ, Brodsky RA. Alternative donor transplantation with high-dose post-transplantation cyclophosphamide for refractory severe aplastic anemia. *Biol Blood Marrow Transplant.* 2017;23:498–504.
24. Bonini C, Mondino A. Adoptive T-cell therapy for cancer: the era of engineered T cells. *Eur J Immunol.* 2015;45:2457–69.
25. Tolar J, Le Blanc K, Keating A, Blazar BR. Concise review: hitting the right spot with mesenchymal stromal cells. *Stem Cells.* 2010;28:1446–55.
26. Snowden JA, Badoglio M, Labopin M, Giebel S, McGrath E, Marjanovic Z. Evolution, trends, outcomes, and economics of hematopoietic stem cell transplantation in severe autoimmune diseases. *Blood Adv.* 2017;1:2742–55.
27. Maude SL, Frey N, Shaw PA, Aplenc R, Barrett DM, Bunin NJ, et al. Chimeric antigen receptor T cells for sustained remissions in leukemia. *N Engl J Med.* 2014;371:1507–17.