



Trends in Use and Outcomes of Autologous and Allogeneic Hematopoietic Cell Transplantation among Racial/Ethnic Groups in the U.S.

WHY?

There has been an increase in the number of autologous (your own cells) and allogeneic (cells from donor) hematopoietic cell transplants (HCT) worldwide and improvement in outcomes after HCT for most hematologic disorders over time. Racial/ethnic disparities in access and

outcomes of HCT are well documented, and there have been efforts made to mitigate these disparities. This research looked at whether there have been improvements in the utilization and outcomes after HCT among various U.S. racial/ethnic groups over time.

EQUAL OUTCOMES FOR ALL. ACCESS TO CARE FOR ALL.



WHAT?

This observational study evaluated utilization rates and outcomes after HCT comparing non-Hispanic whites and racial/ethnic groups in the U.S. Provider reported race/ethnic groups were divided into non-Hispanic white, African American, Hispanic and other (Asian, Pacific Islander, Native American).



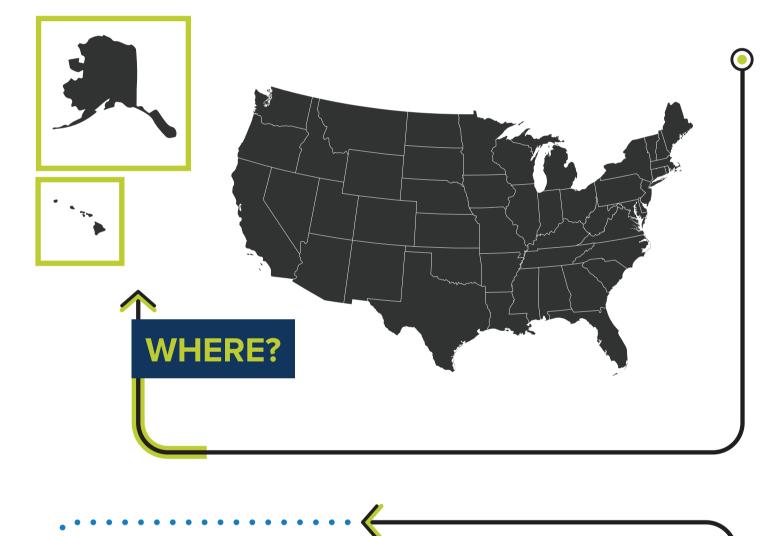
WHO?

80,080 adults who had their first autologous HCT and 60,412 adult and pediatric patients who had their first allogeneic HCT to treat a blood cancer between 2009-2018.

	Disease	Transplant Type	Patient Population (Age)
	Non-Hodgkin lymphoma (NHL)	Autologous	Adult
	Hodgkin disease (HD)	Autologous	Adult
	Multiple myeloma	Autologous	Adult
	Acute myeloid leukemia (AML)	Allogeneic	Pediatric + Adult
	Acute lymphoblastic leukemia (ALL)	Allogeneic	Pediatric + Adult
	Lymphoma, including chronic lymphocytic leukemia (CLL)	Allogeneic	Pediatric + Adult
	Myelodysplastic syndrome (MDS)/ myeloproliferative disease (MPD)	Allogeneic	Pediatric + Adult

WHEN?

2009-2018



significantly improved for all racial/ethnic groups over time. In pediatric patients, overall survival after allogeneic HCT is significantly lower for African Americans and those who were in the

Increased utilization of autologous and allogeneic HCT from 2009-2018 in African

American, Hispanic and other races.

on this challenge.

IMPACT/FINDINGS:

non-Hispanic Whites, with Hispanics having similar overall survival to non-Hispanic White patients.

"other" category (Asian, Pacific Islander, Native American) compared to

Overall survival after autologous HCT and allogeneic HCT in adults has

Although there have been improvements in HCT utilization and outcomes over time, there is still a gap in outcome for pediatric patients undergoing allogeneic HCT who are African American or were in the "other" category (Asian, Pacific Islander, Native American). There is a need to strengthen efforts to improve access to and outcomes of HCT. It is the responsibility of all stakeholders to take

National Marrow Donor Program® (NMDP)/Be The Match® is committed to

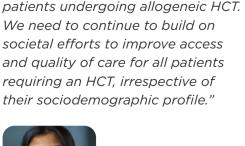
achieving equal opportunity and outcomes for all patients.

research and philanthropic funding to get there.

We're committed to making sure all stem cell transplant patients have equal access to transplant and equal outcomes. That's not the case today, and we won't rest until that goal is achieved. We need more

FROM THE EXPERTS

adults from different racial/ethnic groups. Unfortunately, disparities in outcomes persist in pediatric



Our study shows encouraging

gap between outcomes of

autologous and allogeneic

results in terms of narrowing the

hematopoietic cell transplantation in

2018. Survival has improved for most HCT patients, however there is still room for improvement in some subgroups. Hopefully our study helps focus attention on patients who need additional study to improve their survival after HCT." Theresa Hahn, PhD, MS Professor of Oncology Clinical Epidemiologist Roswell Park Comprehensive

Cancer Center

The good news is that utilization of

both autologous and allogeneic HCT has increased across all races and

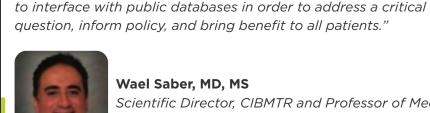
Hispanic ethnicity between 2009 and

Mayo Clinic

Bone and Marrow

Transplant Physician

Nandita Khera, MD, MPH



The CIBMTR leveraged its infrastructure and used modern tools

Medical College of Wisconsin

Wael Saber, MD, MS Scientific Director, CIBMTR and Professor of Medicine,

Division of Hematology/Oncology, Department of Medicine