Double unrelated umbilical cord blood versus HLA-haploidentical bone marrow transplantation (BMT CTN 1101)

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On behalf of the Blood and Marrow Transplant Clinical Trials Network

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Conflict of interest

Research funding: GamidaCell, Magenta, Astex, FateTherapeutics

Advisory Board: AlloVir

Hypothesis

As compared to dUCB, haplo-BM would have a 15% higher 2-year PFS.

- Null hypothesis: no difference in 2-year PFS between treatment arms
- Target sample size of 205 per group to maintain type I error of 5% while providing 80% statistical power for a two-sided test to detect a 15% increase in 2-year PFS in haplo-BM arm

Study Design

- Phase III, randomized trial of RIC: dUCB versus haplo-BM
- Hematologic malignancy (acute leukemia / lymphoma)
- Primary endpoint: progression-free survival at 2 years
- Intent-to-treat analysis from time of randomization

Accrual

Closed for Accrual June 2018 18 to 70 years Acute leukemia or lymphoma 18 to 70 years Both dUCB and haplo-BM donors Acute leukemia or lymphoma Both dUCB and haplo-BM donors n=368 n=368 **Randomization Stratified Center** 33 centers **dUCB** haplo-BM No transplant n=11 No transplant n=15 n=186 n=183 Relapse n= 1 Relapse n= 1 Transplant Transplant Death in relapse n= 7 Death in relapse n= 9 n=175 n=168 Withdrew consent n= 3 Withdrew consent n= 4 Death in remission n= 0 Death in remission n= 1 dUCB n= 172 haplo-BM n=154 dUCB n=11 haplo-BM n= 1 Other n=2Other n=3

Open for Accrual June 2012

Characteristics: Acute Leukemia

	dUCB N = 186	haplo-BM N = 182
Disease status		
1 st complete remission	74%	85%
2 nd complete remission	26%	15%
≥3 rd complete remission		<1%
Cytogenetic risk		
Favorable	13%	15%
Intermediate	46%	41%
Poor	32%	33%
Not reported	10%	12%

Characteristics: Lymphoma

	dUCB N = 186	haplo-BM N = 182
Disease status		
Complete response	39%	32%
Partial response	48%	57%
Follicular or other non-Hodgkin	14%	11%

Graft Characteristics

	dUCB N=175	haplo-BM N=167
TNC infused, median,	2.95 (1.85 – 4.32) x	2.68 (1.87 – 3.63) x
IQR	10 ⁷ /kg	10 ⁸ /kg
CD34 infused, median,	1.30 (0.70 – 2.30) x	2.87 (1.44 – 3.86) x
IQR	10 ⁵ /kg	10 ⁶ /kg
CD3 infused, median,	5.50 (1.90 – 8.20) x	2.96 (2.24 – 4.28) x
IQR	10 ⁶ /kg	10 ⁷ /kg

Results by Intention-to-treat

from the time of randomization:

- Progression-free-survival
- Treatment-related mortality
- Relapse/Progression
- Overall survival

by treatment arm:

- Neutrophil recovery
- Platelet recovery
- Acute GVHD
- Chronic GVHD

Primary Endpoint – Progression-Free Survival at 2 yrs.



Intention-to- treat	dUCB	haplo-BM
Number of events	117	104
PFS at 2 yrs.	35%	41%
95% Conf. interval	28%-42%	34%-48%
Median follow-up	25 months	25 months
Range	20-26 months	23-25 months

Δ=6.1% (95%Cl -5% to 17%)

Multivariate Analysis: Progression-free Survival

	Hazard Ratio (95% confidence interval)	P-value
Donor		0.060
haplo-BM	1.00	
dUCB	1.30 (0.99 – 1.70)	0.060
Adjusted for transplant center		
Donor		
Haplo-BM	1.00	
dUCB	1.27 (0.92-1.75)	0.162

Multivariate Analysis: Progression-free Survival

	Hazard Ratio (95% confidence interval)	P-value
Disease Risk		0.029
Acute leukemia, CR 1	1.00	
Acute leukemia, CR 2 and CR 3	0.91 (0.62 – 1.34)	0.64
Lymphoma, CR	0.58 (0.34 – 1.01)	0.053
Lymphoma, PR	1.51 (1.04 – 2.20)	0.032
Follicular lymphoma	0.82 (0.38 – 1.78)	0.621
Age, >59 years at randomization	1.00 (0.76 – 1.32)	0.979
Performance score, 90 – 100	1.05 (0.79 – 1.40)	0.742

Cumulative Incidence of Relapse at 2 yrs.



Cumulative Incidence of TRM at 2 yrs.



Overall Survival at 2 yrs.



Intention-to- treat	dUCB	haplo-BM
Number of events	117	104
OS at 2 yrs.	46%	57%
95% Conf. interval	38%-53%	49%-64%
Main COD	dUCB	haplo-BM
Primary disease	55	46
Infections	16	13
Organ failure	14	8

Neutrophil Recovery by Treatment Arm



By treatment arm	dUCB	haplo-BM
Number of patients	175	167
Anc > 500	95%	99%
95% Conf. interval	90%-97%	94%-100%
Median in days	15 (r: 4-69)	17 (r:1-87)

Platelet Recovery > 20K by Treatment Arm



Grades II-IV Acute GVHD at day +180 by Treatment Arm



By treatment arm	dUCB	haplo-BM
Number of patients	175	167
aGVHD at 180 days	35%	28%
95% Conf. interval	28%-42%	22%-35%

Grades III-IV	9%	7%	
95% Conf. interval	5%-13%	4%-12%	

Chronic GVHD at 2 yrs. by Treatment Arm



By treatment arm	dUCB	haplo- BM
Number of patients	175	167
cGVHD at 2 yrs.	22%	26%
95% Conf. interval	16%-29%	20%-33%



No significant difference in PFS at 2-years between dUCB and haplo-BM with or without adjustment for transplant center.

Neutrophil recovery was faster in dUCB but higher incidence by day +60 in haplo-BM.

TRM was lower and survival was higher in haplo-BM.

There were no significant differences in relapse, grade II-IV or III-IV acute GVHD, and chronic GVHD.

Conclusion

Among the 368 patients randomized on this study, no significant difference was observed in the 2-year PFS between the dUCB and haplo-BM arms suggesting both donor types extend access to transplantation.

Although the trial did not record the expected 15% difference in 2-year PFS between treatment arms in adults with hematologic malignancy, lower NRM and higher overall survival favor haplo-BM transplantation.

However Taking a closer look into the data...

CTN 1101 vs. The Community

Research hypothesis:

 Outcomes from a contemporary registry study will approximate outcomes from a phase III randomized clinical trial

Endpoints:

- Primary endpoint: Progression-free survival at 2-yr post-transplant
- Secondary endpoints: hematopoietic recovery, graft failure, acute and chronic GvHD, relapse, non-relapse mortality and overall survival

CTN 1101 vs. The Community

	CTN1101		Non-CTN1101		
Donor Type	BM	dUCB	BM	PB	dUCB
No. of patients	157	185	319	409	147
No. of centers	29	31	40	73	38

Effect of Center Expertise

Progression-Free Survival	Hazard Ratio (95% confidence interval)	P-value
Adjusted for transplant center		
Haplo-BM	1.00	
dUCB	1.27 (0.92-1.75)	0.162

Effect of Center Expertise

	Center experience with Haplo and dUCB										
	>10 UCB transplants	<=10 UCE	3 transpla	ants	<=10 UC	0 UCB transplants					
		>10 haplo transplants		<=10 haplo transplants							
	(n=117, 10 centers)	(n=110, 2 centers)			(n=140, 21 centers)						
Site Experience in prior year n Overall, Unadj. for Ctr. 367 Overall, Adj. for Ctr. 367 UCB>10 pts 117 UCB<=10 pts, Haplo>10 pts 110 UCB<=10 pts, Haplo<=10 pts 140 UCB<=10 pts, Haplo<=10 pts 140 HR for dUCB vs Haplo							0				

CTN 1101 Quality of Life

FACT-BMT: The Functional Assessment of Cancer Therapy – Bone Marrow Transplant subscale version 4.0 instrument: Physical Well-being, Social/Family Well-being, Emotional Wellbeing, and Functional Well-being.

MOS SF-36: The Medical Outcomes Study Short Form 36. Physical Functioning, Role Physical, Pain Index, General Health Perceptions, Vitality, Social Functioning, Role Emotional, and Mental Health Index.

Global HQL: Four standard questions to assessed patient self-assessed Karnofsky performance status, overall health and overall quality of life, (excellent, very good, good, fair, poor).

Occupational Functioning: Occupational functioning assessed current job status, type of work, number of hours of paid and unpaid work, school, importance of work and change in work goals.

EQ-5D: The EQ-5D collected data used to calculate patient-reported utilities for cost-utility analyses and contains a five item survey measuring mobility, self-care, usual activities, pain/discomfort and anxiety/depression.

Cost Effectiveness Analysis PI Scott Ramsey

- Single center studies suggested that early post-HCT cost after dUCB to be higher than haplo-BM
- Hypothesis: haplo-BM more cost effective than dUCB
- Plan: obtain 2-year cost data in CTN 1101 patients
- "Glitch": insurance companies declined to provide data
- Solution: to cross reference Vizient and CIBMTR data

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