

Supplemental online content for:

## **Financial Implications of Early Hospital Discharge After AML-Like Induction Chemotherapy: A 4-Year Retrospective Analysis**

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**eTable 1. Characteristics of EHD and Inpatient Control Patients With Blood Count Recovery**

Parameter	EHD Cohort n (%)	Inpatient Cohort n (%)	P Value
Total, N	153	41	
Patient demographics			
Age, mean (range), y	56 (18–79)	58 (25–80)	.15
Sex			.48
Female	68 (44)	21 (51)	
Male	85 (56)	20 (49)	
Travel time to outpatient clinic, mean (range), min	21 (1–124)	63 (4–155)	<.001
Insurance type			.40
Medicare	56 (37)	20 (49)	
Medicaid	18 (12)	6 (15)	
Private	66 (43)	13 (32)	
Other	13 (8)	2 (5)	
Disease characteristics			
Disease			.24
AML	124 (81)	37 (90)	
MDS	29 (19)	4 (10)	
Secondary disease	46 (30)	7 (17)	.12
Cytogenetic risk			.016
Favorable	20 (14)	11 (30)	
Intermediate-1	47 (34)	16 (43)	
Intermediate-2	30 (21)	2 (5)	
Adverse	43 (31)	8 (22)	
Chemotherapy administered			.014
HiDAC-containing regimens	148 (97)	36 (88)	
7 + 3 (± additional drug)	5 (3)	3 (7)	
Cytarabine/Decitabine <sup>a</sup>	0 (0)	2 (5)	
Clinical/Laboratory findings at baseline			
Performance status			.0011
0–1	133 (87)	26 (63)	
2–4	20 (13)	15 (37)	
Creatinine, mg/dL	0.89 (0.39–4.36)	0.85 (0.47–2.51)	.23
Total bilirubin, mg/dL	0.81 (0.2–7.7)	0.78 (0.3–3)	.66
Albumin, g/dL	3.89 (2.3–5)	3.43 (2.3–4.2)	<.001
WBC count, thousand/microL	20.08 (0.39–261.48)	32.15 (0.47–144.96)	.018
% peripheral blood blasts	24 (0–97)	29.78 (0–94)	.078
TRM score	3.88 (0–24.22)	8.56 (0.3–42.02)	.0011

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eTable 1. Characteristics of EHD and Inpatient Control Patients With Blood Count Recovery (cont.)			
Parameter	EHD Cohort n (%)	Inpatient Cohort n (%)	P Value
Clinical/Laboratory findings postchemotherapy			
Performance status			<.001
0-1	137 (90)	18 (44)	
2-4	16 (10)	23 (56)	
Creatinine, mg/dL	0.71 (0.32-2.65)	0.72 (0.25-2.71)	.37
Total bilirubin, mg/dL	0.98 (0.2-5.1)	1.12 (0.3-2.6)	.049
Albumin, g/dL	3.56 (2.1-5)	3.04 (1.9-3.9)	<.001
WBC count, thousand/microL	0.89 (0-19.79)	0.68 (0-14.73)	.14
% peripheral blood blasts	1.35 (0-46)	0.73 (0-30)	.19
TRM score	5.77 (0.34-28.12)	14.21 (1.23-77.63)	<.001
Treatment response			.36
No CR/CRi	12 (8)	5 (12)	
CR or CRi	141 (92)	36 (88)	

Abbreviations: AML, acute myeloid leukemia; CR, complete remission; CRi, complete remission with incomplete hematologic recovery; EHD, early hospital discharge; HiDAC, high-dose cytarabine; MDS, myelodysplastic syndromes; TRM, treatment-related mortality; WBC, white blood cell.

<sup>a</sup>Cytarabine, 100 mg/m<sup>2</sup> × 7 days, and decitabine, 20 mg/m<sup>2</sup> × 10 days.

**eTable 2. Safety and Resource Utilization of EHD and Inpatient Control Patients With Blood Count Recovery**

Parameter	EHD Cohort Mean (Range)	Inpatient Cohort Mean (Range)	P Value
Total, N	153	41	
Days on study	22.81 (14–41)	18.56 (5–34)	<.001
Days as inpatient	6.10 (0–25)	18.56 (5–34)	<.001
Days as outpatient	16.71 (3–38)	—	—
Time spent as outpatient, %	72.45 (11–100)	—	—
Number of readmissions	1.08 (0–3)	—	—
Days in ICU	0.14 (0–8)	0.59 (0–11)	.067
Study days spent in ICU, %	0.70 (0–38.10)	3.09 (0–57.89)	.067
Early death, n (%)			
No early death	151 (99)	40 (98)	.51
Died ≤30 days after start of study	2 (1)	1 (2)	—
Physician visits per outpatient study day	0.06 (0–0.33)	—	—
RN/APP visits per outpatient study day	0.10 (0–0.50)	—	—
Outpatient laboratory visits per outpatient study day	0.44 (0–1.00)	—	—
ED visits per study day	0.04 (0–0.16)	—	—
Study days on IV antimicrobials, %	36.84 (0–100)	52.22 (0–100)	.01
RBC transfusions per study day			
Total	0.27 (0.04–0.80)	0.40 (0.13–0.92)	<.001
Inpatient	0.61 (0–2.33)	0.40 (0.13–0.92)	<.001
Outpatient	0.15 (0–0.65)	—	—
Platelet transfusions per study day			
Total	0.24 (0.04–1.06)	0.41 (0.10–1.20)	<.001
Inpatient	0.38 (0–2.33)	0.41 (0.10–1.20)	.19
Outpatient	0.19 (0–1.00)	—	—

Abbreviations: APP, advanced practice provider; ED, emergency department; EHD, early hospital discharge; ICU, intensive care unit; IV, intravenous; RBC, red blood cell; RN, registered nurse

<b>eTable 3. Charges per Day for EHD and Inpatient Control Patients With Blood Count Recovery</b>			
<b>Charge Category</b>	<b>EHD Cohort Charges per Day (Range), USD</b>	<b>Inpatient Cohort Charges per Day (Range), USD</b>	<b>P Value</b>
Total, N	153	139	
<b>All categories</b>			
Combined	3,826 (905–9,954)	8,404 (4,596–26,320)	<.001
Inpatient only	8,119 (4,828–16,068)	8,404 (4,596–26,320)	.59
<b>Facility/Provider</b>			
Combined	1,214 (104–4,060)	3,592 (2,806–9,907)	<.001
Inpatient only	3,419 (1,693–6,410)	3,592 (2,806–9,907)	.14
<b>Emergency department</b>			
Combined	103 (0–480)	0 (0–0)	<.001
<b>Transfusion</b>			
Combined	1,111 (225–3,780)	1,875 (628–7,933)	<.001
Inpatient only	1,667 (306–6,299)	1,875 (628–7,933)	.39
<b>Laboratory/Pathology</b>			
Combined	686 (87–2,398)	916 (338–3,031)	.0037
Inpatient only	993 (307–2,897)	916 (338–3,031)	.15
<b>Imaging</b>			
Combined	172 (0–852)	333 (0–1,708)	.038
Inpatient only	502 (0–2,904)	333 (0–1,708)	.011
<b>Pharmacy</b>			
Combined	497 (10–1,797)	1,538 (130–3,330)	<.001
Inpatient only	1,531 (265–3,500)	1,538 (130–3,330)	.42
<b>Miscellaneous</b>			
Combined	72 (0–598)	150 (0–991)	.43
Inpatient only	86 (0–1,217)	150 (0–991)	.0012

Abbreviation: EHD, early hospital discharge.

**eTable 4. Characteristics of Medically Matched EHD and Inpatient Control Patients**

Parameter	EHD Cohort n (%)	Inpatient Cohort n (%)	P Value
Total, N	152	49	
Patient demographics			
Age, mean (range), y	56.31 (18–83)	54.92 (24–90)	.65
Sex			.19
Female	60 (40)	25 (51)	
Male	90 (60)	24 (49)	
Travel time to outpatient clinic, mean (range), min	17.27 (1–58)	53.32 (2–155)	<.001
Insurance type			.11
Medicare	57 (38)	22 (45)	
Medicaid	18 (12)	11 (22)	
Private	62 (41)	12 (24)	
Other	13 (9)	4 (8)	
Disease characteristics			
Disease	122 (80)	46 (94)	.026
AML	30 (20)	3 (6)	
MDS	49 (33)	17 (35)	
Secondary disease			.86
Cytogenetic risk			.43
Favorable	16 (12)	9 (20)	
Intermediate-1	44 (32)	16 (36)	
Intermediate-2	31 (23)	7 (16)	
Adverse	45 (33)	13 (29)	
Chemotherapy administered <sup>a</sup>			
HiDAC-containing regimens	142 (95)	45 (92)	.40
7 + 3 (± additional drug)	6 (4)	2 (4)	
Cytarabine/Decitabine <sup>a</sup>	2 (1)	2 (4)	
Clinical/Laboratory findings at baseline			
Performance status			1.00
0–1	144 (96)	47 (96)	
2–4	6 (4)	2 (4)	
Creatinine, mg/dL	0.91 (0.48–4.36)	0.9 (0.47–1.48)	.51
Total bilirubin, mg/dL	0.79 (0.2–2.9)	0.67 (0.3–1.6)	.23
Albumin, g/dL	3.89 (2.5–5.0)	3.7 (2.5–4.7)	.0062
WBC count, thousand/microL	17.47 (0.39–261.48)	34.33 (0.62–247.93)	.0052
% peripheral blood blasts	21.57 (0–97)	35.2 (0–97)	.0022
TRM score	3.32 (0–16.80)	4.58 (0–32.15)	.41
Clinical/Laboratory findings postchemotherapy			
Creatinine, mg/dL	0.73 (0.35–2.65)	0.72 (0.40–1.41)	.54
Total bilirubin, mg/dL	0.95 (0.2–3.0)	0.8 (0.3–2.6)	.20
Albumin, g/dL	3.55 (2.1–5.0)	3.37 (2.4–4.6)	.023
WBC count, thousand/microL	1.14 (0–37.20)	0.75 (0–9.38)	.41
% peripheral blood blasts	1.43 (0–46)	0.84 (0–38)	.30
TRM score	5.25 (0.34–13.56)	5.55 (0.11–16.26)	.75

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eTable 4. Characteristics of Medically Matched EHD and Inpatient Control Patients (cont.)			
Parameter	EHD Cohort n (%)	Inpatient Cohort n (%)	P Value
Treatment response			.004
No CR/CRi	24 (16)	18 (37)	
CR/CRi	128 (84)	31 (63)	
Off-study reason			<.001
Blood count recovery	130 (87)	15 (31)	
Day 42	8 (5)	1 (2)	
Death	4 (3)	0 (0)	
Hospital discharge	0 (0)	31 (63)	
New AML therapy	6 (4)	2 (4)	
Transfer of care to outside facility	2 (1)	0 (0)	

Abbreviations: AML, acute myeloid leukemia; EHD, early hospital discharge; CR, complete remission; CRi, complete remission with incomplete hematologic recovery; HiDAC, high-dose cytarabine; MDS, myelodysplastic syndromes; TRM, treatment-related mortality; WBC, white blood cell.

<sup>a</sup>Cytarabine, 100 mg/m<sup>2</sup> x 7 days, and decitabine, 20 mg/m<sup>2</sup> x 10 days.

**eTable 5. Safety and Resource Utilization of Medically Matched EHD and Inpatient Control Patients**

Parameter	EHD Cohort Mean (Range)	Inpatient Cohort Mean (Range)	P Value
Total, N	152	49	
Days on study	24.00 (6–42)	15.98 (3–42)	<.001
Days as inpatient	6.24 (0–25)	15.98 (3–42)	<.001
Days as outpatient	17.76 (1–42)	—	—
Time spent as outpatient, %	72.38 (11–100)	—	—
Number of readmissions	1.10 (0–3)	—	—
Days in ICU	0.25 (0–16)	0 (0–0)	.08
Study days spent in ICU, %	1.01 (0–38.1)	0 (0–0)	.08
Early death, n (%)			.34
No early death	147 (97)	49 (100)	
Died ≤30 days after start of study	5 (3)	0 (0)	
Physician visits per outpatient study day	0.06 (0–0.33)	—	—
RN/APP visits per outpatient study day	0.10 (0–0.50)	—	—
Outpatient laboratory visits per outpatient study day	0.45 (0–1.00)	—	—
ED visits per study day	0.04 (0–0.17)	—	—
Study days on IV antimicrobials, %	37.05 (0–100)	40.90 (0–100)	.71
RBC transfusions per study day			
Total	0.27 (0.04–0.80)	0.40 (0–1.00)	<.001
Inpatient	0.60 (0–1.50)	0.40 (0–1.00)	<.001
Outpatient	0.16 (0–0.75)	—	—
Platelet transfusions per study day			
Total	0.24 (0.04–1.06)	0.4 (0.09–1.38)	<.001
Inpatient	0.38 (0–1.27)	0.4 (0.09–1.38)	1.00
Outpatient	0.19 (0–1.00)	—	—

Abbreviations: APP, advanced practice provider; ED, emergency department; EHD, early hospital discharge; ICU, intensive care unit; IV, intravenous; RBC, red blood cell; RN, registered nurse.



eTable 6. Characteristics of Medically Matched EHD and Inpatient Control Patients With Blood Count Recovery			
Parameter	EHD Cohort n (%)	Inpatient Cohort n (%)	P Value
Total, N	130	15	
Patient demographics			
Age, mean (range), y	55 (18–79)	51 (25–73)	.70
Sex			.18
Female	54 (42)	9 (60)	
Male	76 (58)	6 (40)	
Travel time to outpatient clinic, mean (range), min	17 (1–58)	89 (28–155)	<.001
Insurance type			.37
Medicare	47 (36)	7 (47)	
Medicaid	14 (11)	3 (20)	
Private	56 (43)	5 (33)	
Other	13 (10)	0 (0)	
Disease characteristics			
Disease			.30
AML	104 (80)	14 (93)	
MDS	26 (20)	1 (7)	
Secondary disease	40 (31)	3 (20)	.55
Cytogenetic risk			.021
Favorable	16 (14)	5 (38)	
Intermediate-1	39 (33)	6 (46)	
Intermediate-2	26 (22)	0 (0)	
Adverse	37 (31)	2 (15)	
Chemotherapy administered			.12
HiDAC-containing regimens	126 (97)	13 (87)	
7 + 3 (± additional drug)	4 (3)	2 (13)	
Clinical/Laboratory findings at baseline			
Performance status			.54
0–1	124 (95)	14 (93)	
2–4	6 (5)	1 (7)	.18
Creatinine, mg/dL	0.90 (0.48–4.36)	0.78 (0.47–1.25)	.86
Total bilirubin, mg/dL	0.78 (0.2–2.9)	0.77 (0.3–1.6)	.063
Albumin, g/dL	3.9 (2.5–5.0)	3.7 (2.9–4.2)	.022
WBC count, thousand/microL	16.58 (0.39–261.48)	28.01 (1.83–138.44)	.022
% peripheral blood blasts	21.19 (0–97)	34.27 (0–93)	.99
TRM score	3 (0–16.80)	3.14 (0.3–10.96)	.54

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**eTable 6. Characteristics of Medically Matched EHD and Inpatient Control Patients With Blood Count Recovery (cont.)**

Parameter	EHD Cohort n (%)	Inpatient Cohort n (%)	P Value
Clinical/Laboratory findings postchemotherapy			
Creatinine, mg/dL	0.72 (0.32–2.65)	0.63 (0.40–1.00)	.16
Total bilirubin, mg/dL	0.94 (0.2–3.0)	1.00 (0.3–2.6)	.64
Albumin, g/dL	3.56 (2.1–5.0)	3.21 (2.7–3.8)	.0035
WBC count, thousand/microL	0.83 (0–19.79)	0.39 (0–2.19)	.56
% peripheral blood blasts	1.41 (0–46)	0 (0–0)	.27
TRM score	4.85 (0.34–13.56)	4.82 (1.23–9.26)	1.00
Treatment response			
No CR/CRi	8 (6)	2 (13)	
CR or CRi	122 (94)	13 (87)	

Abbreviations: AML, acute myeloid leukemia; CR, complete remission; CRi, complete remission with incomplete hematologic recovery; EHD, early hospital discharge; HiDAC, high-dose cytarabine; MDS, myelodysplastic syndromes; TRM, treatment-related mortality; WBC, white blood cell.

<b>eTable 7. Safety and Resource Utilization of Medically Matched EHD and Inpatient Control Patients With Blood Count Recovery</b>			
<b>Parameter</b>	<b>EHD Cohort Mean (Range)</b>	<b>Inpatient Cohort Mean (Range)</b>	<b>P Value</b>
Total, N	130	15	
Days on study	22.62 (14–38)	18.2 (5–28)	.0021
Days as inpatient	5.91 (0–25)	18.2 (5–28)	<.001
Days as outpatient	16.72 (3–38)	—	—
Time spent as outpatient, %	72.97 (11–100)	—	—
Number of readmissions	1.08 (0–3)	—	—
Days in ICU	0.15 (0–8)	0 (0–0)	.44
Study days spent in ICU, %	0.75 (0–38.1)	0 (0–0)	.44
Early death, n (%)			
No early death	128 (98)	15 (100)	1.00
Died ≤30 days after start of study	2 (2)	0 (0)	
Physician visits per outpatient study day	0.06 (0–0.33)	—	—
RN/APP visits per outpatient study day	0.10 (0–0.50)	—	—
Outpatient laboratory visits per outpatient study day	0.44 (0–1.00)	—	—
ED visits per study day	0.04 (0–0.16)	—	—
Study days on IV antimicrobials, %	36.76 (0–100)	40.13 (0–100)	.69
RBC transfusions per study day			
Total	0.27 (0.04–0.80)	0.37 (0.13–0.83)	.026
Inpatient	0.60 (0–1.50)	0.37 (0.13–0.83)	<.001
Outpatient	0.15 (0–0.65)	—	—
Platelet transfusions per study day			
Total	0.23 (0.04–1.06)	0.23 (0.10–0.56)	.94
Inpatient	0.35 (0–1.27)	0.23 (0.10–0.56)	.044
Outpatient	0.18 (0–1.00)	—	—

Abbreviations: APP, advanced practice provider; ED, emergency department; EHD, early hospital discharge; ICU, intensive care unit; IV, intravenous; RBC, red blood cell; RN, registered nurse

**eTable 8. Charges per Day for Medically Matched EHD Versus Inpatient Control Patients With Blood Count Recovery**

Charge Category	EHD Cohort (n=130) Charges per Day (Range), USD	Inpatient Cohort (n=15) Charges per Day (Range), USD	P Value
Total, N	130	15	
All categories			
Combined	3,792 (905–9,954)	6,429 (4,596–8,342)	<.001
Inpatient only	8,076 (4,828–16,068)	6,429 (4,596–8,342)	<.001
Facility/Provider			
Combined	1,200 (104–4,060)	3,331 (3,034–3,758)	<.001
Inpatient only	3,417 (1,693–6,410)	3,331 (3,034–3,758)	.81
Emergency department			
Combined	103 (0–480)	0 (0–0)	<.001
Transfusion			
Combined	1,091 (225–3,780)	1,216 (628–2,181)	.23
Inpatient only	1,604 (306–3,768)	1,216 (628–2,181)	.031
Laboratory/Pathology			
Combined	694 (87–2,398)	721 (353–1,195)	.52
Inpatient only	1,001 (307–2,897)	721 (353–1,195)	.018
Imaging			
Combined	167 (0–852)	79 (0–335)	.11
Inpatient only	526 (0–2,904)	79 (0–335)	<.001
Pharmacy			
Combined	494 (10–1,797)	1,018 (130–1,611)	<.001
Inpatient only	1,538 (265–3,500)	1,018 (130–1,611)	<.001
Miscellaneous			
Combined	71 (0–598)	65 (0–623)	.0023
Inpatient only	75 (0–1,217)	65 (0–623)	.58

Abbreviation: EHD, early hospital discharge.