



Background

- Fluid overload is a common cause of hospitalization in patients with acute decompensated heart failure (ADHF).¹⁻⁵
- The Healthcare Cost and Utilization Project found that 25% of patients with ADHF are readmitted to the hospital within 30 days.⁶
- Practice guidelines recommend oral loop diuretics be initiated prior to discharge in order to confirm their effectiveness for patients admitted with heart failure (HF).^{1,2}
- Limited guidance is provided on how diuretics should be managed as patients approach euvolemia.
- A study of 433 U.S. hospitals found 24% of patients were administered intravenous (IV) diuretics on the day of discharge.⁷

Methods

Study Design

- Retrospective, single center, cohort study
- Two groups: observation on an oral loop diuretic < 24 hours versus observation on an oral loop diuretic \geq 24 hours

Primary Objective: Determine if the duration of observation on oral loop diuretics was associated with 30-day HF readmission.

Secondary Objectives:

- 60- and 90-day all-cause and HF readmission rates
- Hospital length of stay
- Predictors of observation periods of < 24 hours duration

Figure 1. Patient Selection

Inclusion

- Adults
- Diagnosis of ADHF
- IV loop diuretic
- Oral loop diuretic on discharge

Exclusion

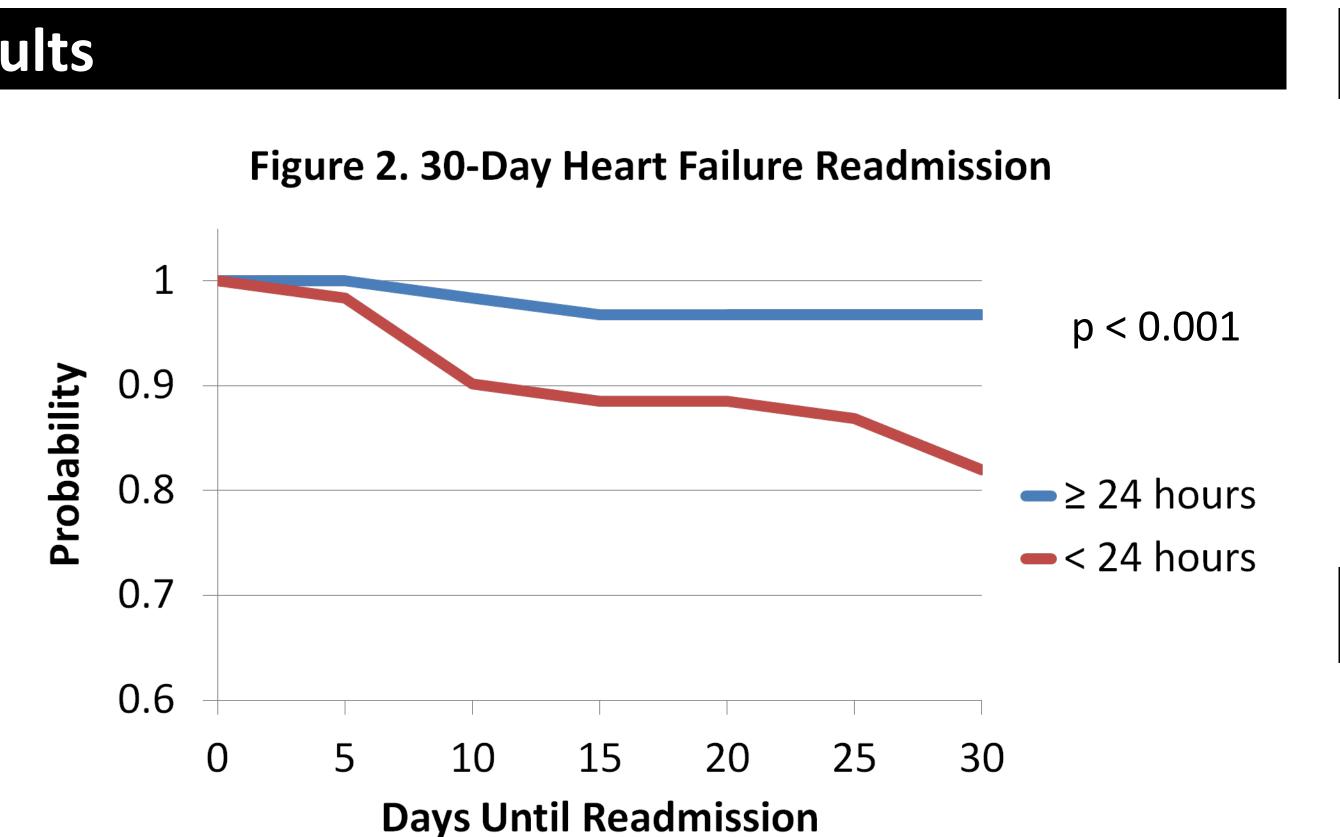
- Aldosterone antagonist > 50 mg
- Cirrhosis
- Dialysis
- Hospital stay < 48 hrs
- Palliative
- care

Duration of Oral Loop Diuretics and 30-Day Readmission in Patients With Acute Decompensated Heart Failure

Results								
Table 1. Patient Characteristics	Observed < 24 hr (n = 61)	Observed > 24 hr (n = 62)	P-value	Figure 2. 30-1	Day Heart Fa	ailure Readn	nission	
Age (years)	60.8 (14.1)	56.7 (13.8)	NS					
Male	41 (67.2)	45 (72.6)	NS	1				< 0.001
Black	38 (62.3)	40 (64.5)	NS				р	< 0.001
Body weight (kg)	97.5 (25.2)	98.3 (25.5)	NS	0.9 0.9				
Ejection fraction	25.1 (14.7)	23.4 (15.2)	NS	bab				
Prior 30-day readmission	15 (24.6)	6 (9.7)	NS	<u>ä</u> 0.8				24 hours
Non-ischemic etiology	29 (47.5)	37 (59.7)	NS	A 0.7			— < 2	24 hours
NYHA Class				0.7				
H	10 (16.4)	10 (16.1)	NS	0.6				
	29 (47.5)	25 (40.3)	IND	0 5 10) 15	20 25	30	
Past medical history					_		50	
Atrial fibrillation	25 (41.0)	16 (25.8)	NS	Days Until Readmission				
Chronic kidney disease	26 (42.6)	27 (43.5)	NS					
Coronary artery disease	31 (50.8)	28 (45.2)	NS	Table 2. Outsource	Observed	Observed	Duralua	P-value
Diabetes mellitus	29 (47.5)	24 (38.7)	NS	Table 2. Outcomes	< 24 hr	<u>> 24 hr</u>	P-value	(Adj)
Admission medications					(n = 61)	(n = 62)		
ACEI or ARB	33 (58.9)	26 (43.3)	NS	Primary Outcome				
Beta-blocker	48 (85.7)	35 (60.3)	0.005	30-day HF readmission	11 (18.0)	2 (3.2)	0.023	<0.001
Aldosterone antagonist	20 (35.7)	19 (32.2)	NS	Secondary Outcomes				
Loop diuretic	49 (78.9)	44 (70.0)	NS NC	60-day HF readmission	18 (29.5)	6 (9.7)	0.014	< 0.001
IV inotrope	2 (3.4)	6 (9.8)	NS	90-day HF readmission	23 (37.7)	12 (19.4)	0.049	< 0.001
Admission laboratories	138.9 (5.1)	139.8 (5.6)	NS	HF readmission	34 (55.7)	23 (37.1)	0.049	< 0.001
Sodium (mEq/L) Potassium (mEq/L)	4.1 (0.7)	4.2 (0.7)	NS	All-cause readmission	39 (64.0)	38 (61.3)	NS	0.009
Serum creatinine (mg/dL)	1.5 (0.8)	1.7 (0.9)	NS	Length of stay (days)	7.8 (6.2)	10.6 (7.4)	0.023	NS
BUN (mg/dL)	35.6 (23.1)	37.6 (27.5)	NS		, , , ,			
NT-proBNP (pg/mL)	11257 (9790)	10588 (9256)	NS	Cardiology follow-up	43 (70.5)	47 (75.8)	NS	NS
IV inotrope during admission	19 (31.1)	28 (45.2)	NS	Time to follow-up (days)	15.0 (19.2)	13.7 (11.6)	NS	NS
\geq 1 dose of oral loop diuretic	42 (68.9)	62 (100)	< 0.001	*Data are number (%) of patients				
IV loop diuretic on day of	· · ·	.		⁺ Adj = adjusted, HF = heart failure				
discharge	26 (42.6)	0 (0.0)	<0.001					
Weight change (kg)	-4.8 (4.4)	-6.8 (6.6)	NS	Table 3. Variables	Estimat	e (SD) 9	5% CI	P-value
Time observed on oral loop			.0.001	Observation < 24 hr	6.0 (0).8) 1.4	-42.4	0.033
diuretic (hr)	7.6 (9.1)	86 (122.2)	<0.001	Age	1.0 (0.3) 1.0 -) – 1.1	NS
Discharge medications				Readmission within 30 day	•	•	- 16.7	0.032
ACEI or ARB	37 (60.7)	34 (54.8)	NS		•	-		
Beta-blocker	48 (78.7)	46 (74.2)	NS	ACEI or ARB at discharge	1.1 (0	J./) 0.3	8 – 4.1	NS
Aldosterone antagonist	24 (39.3)	29 (46.8)	NS	Beta-blocker at discharge	2.1 (0	0.9) 0.4	- 17.6	NS
Loop diuretic dose (FE)	116.4 (74.6)	126.1 (81.3)	NS	IV dobutamine at discharg	e 3.0 (1	L.3) 01	- 33.5	NS
IV inotrope	15 (24.2)	6 (9.8)	NS		•	•		
[*] Data presented as mean (SD) or n (%)				IV milrinone at discharge	4.4 (1	L.1) U.5	- 35.0	NS

[†]ACEI = angiotensin-converting enzyme inhibitor, ARB = angiotensin II receptor blocker, BUN= blood urea nitrogen, FE = furosemide equivalent, IV = intravenous, NYHA= New York Heart Association, NT-proBNP = N-terminal pro-B-type natriuretic peptide

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*ACEI = angiotensin-converting enzyme inhibitor, ARB = angiotensin II receptor blocker, CI = confidence interval, IV = intravenous

Limitations

- 24-hour threshold selected based on expert consensus recommendations rather than evidence from clinical trials
- Select number of confounders included in adjusted analysis
- Single center, retrospective design; demographics may not be representative of all centers
- Cannot completely rule out the potential for selection bias

Conclusion

- Observation on an oral loop diuretic \geq 24 hours was associated with significantly lower 30-, 60-, and 90-day HF readmission.
- Observation < 24 hours and prior 30-day readmission were independent prognostic factors for 30-day HF readmission.
- This simple and feasible practice should be strongly considered prior to discharging patients who initially present with ADHF.
- Prospective trials are warranted to confirm these results.

References

- ¹Yancy CW, Jessup M, Bozkurt B, et al. J Am Coll Cardiol. 2013;62(16):e147-239.
- ²Heart Failure Society of America. *J Card Fail*. 2010;16(6):e1-194. ³Ponikowski P, Voors AA, Anker SD, et al. *Eur Heart J*. 2016.
- doi:10.1093/eurheartj/ehw128.
- ⁴Gheorghiade M, Follath F, Ponikowski P, et al. *Eur J Heart Fail*. 2010;12(5):423-433.
- ⁵Collins S, Storrow AB, Albert NM, et al. J Card Fail. 2015;21(1):27-43.
- ⁶Agency for Healthcare Research and Quality. Healthcare Cost and Utilization Project.
- http://www.ahrq.gov/research/data/hcup/index.html.
- ⁷Fazel R, Strait KM, Bikdeli B, Dharmarajan K, Krumholz HM. J Card *Fail*. 2014;20(9):706-707.

Disclosures

Benjamin Laliberte, Pharm.D. - Nothing to Disclose