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ORIGINAL RESEARCH

Combined aquaretic and diuretic therapy in acute heart failure

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Introduction: Acute heart failure (AHF) is a leading cause of hospitalization and readmission in the US. The present study evaluated maximum diuresis while minimizing electrolyte imbalances, hemodynamic instability, and kidney dysfunction, to achieve a euvolemic state safely in a shorter period of time.

Methods and results: A protocol of combined therapy with furosemide, metolazone, and spironolactone, with or without tolvaptan and acetazolamide, was used in 17 hospitalized patients with AHF. The mean number of days on combination diuretic protocol was 3.8 days. The mean daily fluid balance was 3.0 ± 2.1 L negative. The mean daily urine output (UOP) was 4.1 ± 2.0 L (range 1.8-10.5 L). There were minimal fluctuations in serum electrolyte levels and serum creatinine over the duration of diuretic therapy. There was no statistically significant change in patients' creatinine from immediately prior to therapy to the last day of therapy, with a mean increase in creatinine of 0.14 mg/dL (95% CI -0.03, +0.30, p=0.10).

Conclusion: Our strategy of treating AHF by achieving high UOP, while maintaining stable electrolytes and creatinine in a short period to euvolemic state, is safe.

Keywords: diuretics, aquaretic, acute heart failure, volume overload

Introduction

Acute heart failure (AHF) is a leading cause of hospitalization and readmission in the US.^{1,2} Worsening dyspnea is the most common reason for hospitalization, which is largely a result of sodium and water retention-related fluid overload.^{3,4} The standard management of AHF is removing the fluid primarily with loop diuretics or ultrafiltration.^{5,6} However, there are challenges associated with effective diuresis, not to mention possible adverse effects.^{7–9} As a result, many studies have shown that a significant proportion of patients with AHF are discharged with no to minimal weight loss and with persistently elevated natriuretic peptide levels and intra-cardiac pressures.¹⁰ These features are associated with subsequent adverse outcomes.¹¹

Simultaneous blockade of the nephron channels at different sites achieves higher diuresis effectively and safely.⁸ This is accomplished by the use of medications, including 1) acetazolamide, a reversible inhibitor of the enzyme carbonic anhydrase, which results in reduction in hydrogen ion secretion at the early proximal convoluted tubule with an increased excretion of sodium and water; 2) loop diuretics, which inhibit the Na+–K+–2Cl– pump in the thick ascending loop of Henle; 3) thiazides, which inhibit Na–Cl reabsorption in the distal convoluted tubule;¹² 4) potassium-sparing diuretics, which inhibit epithelial sodium channels (ENaC) at the collecting tubule;¹³

and 5) tolvaptan, which targets vasopressin V2 receptors in the collecting duct,¹⁴ leading to water loss (Figure 1). Various combinations of diuretic therapy in AHF have been reported.^{8,15,16}

The general theme of diuresis in AHF has been to relieve symptoms and achieve diuresis conservatively to a dischargeable state, which largely falls short of achieving a euvolemic state.¹⁰ We implemented a protocol of combined therapy with furosemide, metolazone, and spironolactone, with or without tolvaptan and acetazolamide, in an attempt to maximize diuresis while minimizing electrolyte imbalances, hemodynamic instability, and kidney dysfunction,¹⁷ with a goal to achieve a euvolemic state safely in a shorter period of time. Herein, we report our experience with this protocol in a retrospective series of patients with AHF and its impact on diuresis, kidney function, electrolyte balance, and hemodynamic stability.

Methods

Patient populations and diuretic protocol

The project protocol was approved by the institutional review board of Stony Brook University. Data were de-identified and stored on the secured drive of Stony Brook University. We retrospectively evaluated 17 patients admitted with signs and symptoms of fluid overload related to AHF. This was clinically determined by the on-service attending physician using the Framingham criteria for heart failure (HF) and echocardiographic findings.^{18,19} These signs and symptoms included the documented presence of exertional dyspnea, orthopnea/paroxysmal nocturnal dyspnea, jugular venous distention, rales, lower extremity edema, nocturnal cough, tachycardia >120 bpm, and pleural effusion or pulmonary edema as seen on radiologic imaging. We collected data on consecutive patients who underwent a combination regimen of diuretics, including intravenous (IV) furosemide (bolus or infusion), metolazone, and spironolactone, with or without tolvaptan and acetazolamide. Each diuretic was started, while serum electrolytes were closely monitored throughout the day. IV furosemide, oral metolazone, and oral spironolactone were initiated first with a goal to achieve a target fluid loss of 4-5 L/day. Tolvaptan was added if serum Na was <135 mmol/L, and dose was titrated to maintain a serum sodium of 131-139 mmol/L (based on the initial value, maximum correction was 6-8 mmol/L, which did not exceed 139 mmol/L in 24 hours). Acetazolamide was given as a daily IV bolus if serum bicarbonate was >30mmol/L and to maintain serum bicarbonate 24-30 mmol/L (Table 1). Close monitoring of serum electrolytes was performed to ensure that safe levels of sodium and potassium were maintained. Serum electrolyte levels were measured every 6 hours over each day during combination diuretic treatment. Data on net fluid balance and mean arterial pressure (MAP), as well as serum N-terminal pro b-type natriuretic peptide (BNP), were trended. Data on cardiac structure and function were obtained from the most recent transthoracic echocardiogram. After initial diuresis to a clinically euvolemic state as determined



Figure I Schematic of sites of action and mechanism of ion channel blockade by selected diuretics in the nephron.

Table I Combination diuretics and aquaretic protocol

Diuretics	Laboratory monitoring	Electrolyte goal	Target fluid loss
*Furosemide	Serum electrolytes q6H	Na 131–139 mmol/L	4–5 L/day
IV infusion 10–40 mg/h or boluses			
^Spironolactone 25 mg QD–50 mg PO TID	Serum electrolytes q6H	Potassium 4–4.5 mmol/L	
Metolazone 5 mg BID–10 mg PO BID	Serum electrolytes q6H		
[#] Tolvaptan 15 mg QD to 30 mg PO QD	Serum electrolytes q6H	Na 131–139 mmol/L	
^{\$} Acetazolamide 500 mg IV QD	Serum electrolytes q6H	HCO ₃ 24–30 mmol/L	

Notes: *Doses of furosemide and metolazone were titrated based on UOP. ^Spironolactone dose was titrated to maintain serum potassium of 4–4.5 mmol/L (mEq/L); *Tolvaptan dose was titrated to maintain a serum sodium of 131–139 mmol/L (mEq/L); based on the initial value, the maximum correction was 6–8 mmol/L (mEq/L) and not to exceed 139 mmol/L (mEq/L) in 24 hours. \$Acetazolamide was used to maintain serum bicarbonate 24–30 mmol/L (mEq/L). Abbreviations: IV, intravenous; PO, by mouth; UOP, urine output.

by the cardiology and the nephrology attendings, the patients were maintained and discharged on an individualized oral diuretics regimen as determined by the primary team and cardiology and nephrology services.

Statistical analysis

Descriptive statistics were performed. Categorical variables were presented as numbers and proportions, and continuous variables were presented as mean \pm standard deviation (SD). Categorical values were assessed with chi-square test and continuous variables with Student's t-test. Differences in pre- and posttreatment creatinine and hemoglobin/hematocrit were assessed using data from immediately prior to initiation and last day of intensive combined diuretic therapy. The SD of daily electrolyte levels was averaged for each patient over the course of their therapy, and the mean was obtained of all patients. Given the tendency of extreme values to skew means, a range was also obtained of the highest and lowest electrolyte values over the course of the entire diuretic regimen. Patients with HF with reduced ejection fraction (HFrEF) were defined as those with EF \leq 50%, while patients were considered to have HF with a preserved EF (HFpEF) if they had EF \geq 50%.²⁰ All calculations were performed using SAS software, version 9.3 (SAS Institute Inc., Cary, NC, USA).

Results

Baseline patient characteristics

Of the 17 patients, nine were male and eight were female, with a mean age of 69.0 ± 15.4 years (range 38-93 years). Patients had a mix of comorbid conditions, including hypertension (n=15), coronary artery disease (n=10), and diabetes mellitus (n=8). The mean weight was 98.9 ± 21.7 kg (range 70-141 kg). All patients had known chronic kidney disease, with a mean estimated glomerular filtration rate (eGFR) of 30.6 ± 13.6 mL/ min/1.73 m² (range 8-59 mL/min/1.73 m²).

Nine patients had HFrEF; the overall mean EF was $44.1\pm18.0\%$ (range 20%–72%). There were no statistically

significant differences in age, GFR, or duration of diuretic therapy between patients with HFpEF or HFrEF. Seven patients were managed in an intensive care unit, nine in a step-down unit, and one on the general medicine ward. One patient had a functioning renal allograft. No patients needed ultrafiltration. Two of the patients were on IV inotropes. The average pro-BNP levels on admission were 10457.2 \pm 11084.6 pg/mL (range 422–32494 pg/mL), n=12. Three of the patients (18%) were readmitted back to the hospital within 30 days of discharge.

Euvolemic state

Patients had presented in a volume-overloaded state as defined previously, with the most common presenting symptom of exertional dyspnea (n=9) and the most common sign reported as pedal edema (n=12). The vast majority of patients showed a significant improvement in signs and symptoms of fluid overload at the completion of diuretic therapy (Table 2).

Diuretic use and fluid balance

The mean number of days on combination diuretic protocol was 3.8 (range 1–8 days). The mean daily fluid balance was 3.0 ± 2.1 L negative. The mean daily urine output (UOP) was 4.1 ± 2.0 L (range 1.8-10.5 L). The MAP did not fluctuate significantly, with an average MAP of 79.8 ± 5.4 mmHg in each patient. When serial weights were recorded, the mean weight loss was 7.0 kg over the duration of diuretic therapy (n=9, range 1.0-14.2 kg).

Average daily doses were furosemide 235 ± 149 mg IV (mean dose of 2.7 mg IV/kg body weight), spironolactone 82 ± 37 mg orally, and metolazone 8 ± 4 mg orally. When used, acetazolamide (n=1) dose was 500 mg IV and tolvaptan (n=3) dose was 21 ±8 mg orally.

Electrolyte changes

There were minimal fluctuations in serum electrolyte levels and serum creatinine over the duration of diuretic therapy (Table 3). The changes in serum sodium, potassium, and eGFR in 24-hour intervals as well as the daily mean UOP are shown in Table 4. There was no statistically significant change in patients' creatinine from immediately prior to therapy to the last day of therapy, with a mean increase in creatinine of 0.14 mg/dL (95% CI -0.03, +0.30, p=0.10). Eight out of 17 (47%) patients required potassium supplementation over the course of diuresis, with an average daily dose of 38.6±25.7 mEq (range 25–100 mEq). Individual patients' data are shown in Table 5.

Discussion

We report a series of 17 patients with AHF due to severe volume overload who achieved clinical euvolemic status

Table 2 Number of patients with signs and symptoms	of	fluid
overload prior to and after diuretic therapy		

Sign/symptom	Present	Present
	before	after
	therapy	therapy
Pedal edema	12	0
Pleural effusion	10	4
Rales	9	5
Exertional dyspnea	9	0
Pulmonary edema	8	0
Orthopnea/paroxysmal nocturnal dyspnea	5	0
Jugular venous distention	4	0
Tachycardia >I20 bpm	3	0
Nocturnal cough	0	0

 Table 3 Serum electrolyte and creatinine fluctuations over the course of treatment

	.		
Electrolyte	Mean*	SD^	Range*
	(mmol/L)	(mmol/L)	(mmol/L)
Sodium	138.0	1.3	5.1
Potassium	4.0	0.3	1.1
Chloride	93.9	3.0	8.9
Bicarbonate	30.0	2.2	8.0
Magnesium	2.1	0.1	0.5
Phosphate	4.3	0.5	1.1
Creatinine	2.33	0.16	0.55

Notes: *Mean of individual daily electrolyte concentrations averaged across all patients. ^SD of individual daily electrolyte fluctuations averaged across all patients. *The range of individual daily electrolyte fluctuations averaged across all patients.

with a diuretic and aquaretic regimen including furosemide, metolazone, and spironolactone within a mean of 4 days. Acetazolamide and tolvaptan were used as needed. Renal function and serum electrolytes remained stable. Given the frequent readmissions and lengthy hospitalizations for AHF, this strategy accelerated restoration to a clinically euvolemic state.

The current pattern of diuretic use in AHF suggests that patients are inadequately diuresed by the time of discharge. The 30-day all-cause readmission rate after AHF remains ~25% for Medicare patients.² Although the factors contributing to this trend are multifactorial, one potential reason is that patients are not effectively treated before discharge.²¹ A combined analysis of the Diuretic Optimization Strategies Evaluation (DOSE) and the Cardiorenal Rescue Study in Acute Decompensated Heart Failure (CARRESS-HF) trials has demonstrated that more than one-third of the patients still had congestion when they were discharged from the index hospitalization.⁶ Our goal in this case series is to show that patients have effectively and safely achieved a euvolemic state in a short time interval.

The mean duration of diuretic therapy was <4 days, and the mean daily UOP was >4 L, and this was achieved with relatively stable kidney function and hemodynamic and electrolyte parameters. There was only a non-significant trend toward increase in creatinine. Likewise, the changes in electrolytes occurred within generally accepted parameters. A high dose of spironolactone also helped maintain potassium levels within the target range in the setting of significant diuresis. We emphasize that while the combination diuretic therapy was successfully administered in settings ranging from intensive care units to general medicine wards, this was accomplished with the use of frequent serum blood testing and a close working relationship between the nephrology service and consulting teams.

Our data are limited to in-hospital outcomes. The effect of our strategy on post-discharge outcomes requires a comparator group and longer term follow-up. Although based on

Table 4 Dail	y changes in	serum electroly	ytes, creatinine,	eGFR, and mean	daily urine	over 96 hours
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	24 hours	48 hours	72 hours	96 hours
Change in serum potassium, mmol/L (±SD)	-0.20 (0.43)	+0.17 (0.42)	-0.11 (0.39)	-0.06 (0.38)
Change in serum sodium, mmol/L (±SD)	-0.45 (1.95)	-0.38 (1.82)	-0.27 (1.53)	+0.58 (0.51)
Change in serum bicarbonate, mmol/L (±SD)	+1.89 (2.55)*	+1.46 (2.83)	-0.22 (1.79)	+0.86 (0.13)
Change in serum creatinine, mg/dL (±SD)	+0.05 (0.20)	+0.64 (0.18)	+0.02 (0.20)	+0.01 (0.15)
Change in eGFR, mL/min/1.73 m ² (±SD)	-0.29 (4.82)	-1.31 (3.95)	+0.39 (3.07)	+2.76 (6.63)
Mean total UOP, mL (±SD)	4123 (2272)	4679 (2122)	3971 (2201)	2623 (1419)

Note: *Statistically significant change from prior at p < 0.05.

Abbreviations: eGFR, estimated glomerular filtration rate; SD, standard deviation; UOP, urine output.

Table	5 Indiv	vidual pa	tient	data												
Age	Sex	D/H/C	Ш	Days	Average	Average	Average	Mean daily	Mean daily	Mean	Mean daily	Mean	Daily	Mean	Initial	Final
(years)				therapy	furosemide	spironolactone	Metolazone	UOP (mL)	K (mmol)	daily Na	нсо	daily Cr	K-supp#	daily MAP	Hgb	Hgb
					(mg)	(mg)	(mg)			(Iomm)	(Iomm)	mg/dL	(mEq)	(mmHg)	(g/dL)	(Jp/g)
64	щ	H/C	30	2	300	25	4	3600	3.5	138.0	29.2	1.76	001	72	11.3	10.1
67	Σ	D/H/C	25	8^	285	130	17	4569	4.1	138.9	36.9	2.06	26.7	60	10.8	10.3
79	Σ	D/H/C	30	7^	231	121	6	2799	4.2	133.4	29.9	2.32	0	68	10.0	10.5
71	щ	D/H	72	2	365	150	15	3850	4.0	139.8	23.4	0.89	0	65	9.6	0
47	щ		30	٣	325	75	8	5688	4.1	134.5	30.4	10.1	0	71	7.6	8.2
59	щ	т	55	4	540	94	6	1764	4.3	142.6	28.4	2.09	0	84	7.6	9.2
89	щ	т	99	٣	483	50	01	3680	3.7	132.5	29.9	1.78	37.5	77	11.9	Π.Π
38	Σ		20	ß	144	90	7	3468	3.5	135.1	35.8	2.07	0	69	8.6	9.5
78	Σ	D/H/C	56	4	60	34	8	2481	4.2	135.3	36.8	2.21	0	65	11.2	12.9
49	Σ	D/H/C	56	4	113	94	6	4342	3.9	137.6	20.7	6.81	0	116	9.7	10.3
86	щ	т	7	4	203	125	4	2984	3.9	141.3	39.3	1.26	0	77	8.II	12.6
84	Σ	D/H/C	51	4	011	125	8	2426	4.1	142.0	30.0	2.05	0	88	9.1	9.1
68	щ	D/H/C	35	4	241	25	5	3745	4.2	136.5	25.8	3.43	25	60	II.3	13.9
71	Σ	D/H/C	6	2*	180	50	5	5765	4.0	136.5	31.3	2.49	25	103	12.8	14.4
72	щ	т	25	_	460	25	5	10500	3.8	137.7	24.0	2.85	27.5	76	10.3	II.3
58	Σ	H/C	25	4	100	75	5	5781	4.0	139.2	29.8	1.73	25	108	15.4	14.3
93	Σ	H/C	63	5	175	75	15	4671	3.8	144.5	20.2	2.82	42.5	67	7.7	9.0
Notes: * Abbrevi Hgb, hem	'Mean da I ations: ≀oglobin;	aily dose of D, diabete: F, female;	potass s; H, h, M, mal	ium supplem /pertension; e.	entation over cour C, coronary artery	se of diuresis in mEq. ^A f disease; EF, ejection fr	Patient also receive action; UOP, urine	ed tolvaptan to m output; K, serun	aintain target sod 1 potassium; Na,	um levels. *P	atient also receiv 1; HCO ₃ serum b	ed acetazolan vicarbonate; (nide to maint Cr, serum cre	ain target bicarb atinine; MAP, m	onate levels ean arterial	pressure;
,)															

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the association of persistent congestion and outcomes, these data are encouraging for potentially improving outcomes in AHF patents. This protocol was used with close monitoring of electrolytes under the supervision of an experienced team of cardiologists and nephrologists. This case series demonstrates the potential of combination diuretic therapy for allowing diuresis of a large amount of fluid safely and within a short period of time. Ultimately, this may prevent prolonged hospital stay and frequent readmissions, and such a strategy needs to be evaluated in a randomized controlled trial.

Conclusion

We extend our strategy in treating volume-overloaded AHF patients and achieving high UOP while maintaining stable electrolytes and creatinine in a short period to a euvolemic state. Further data on longer term outcomes are needed.

Disclosure

The authors report no conflicts of interest in this work.

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