Augmented Natriuretic Response to Acute Sodium Infusion after Blood Pressure Elevation with Metaraminol in Normotensive Subjects *

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That patients with essential hypertension have an exaggerated natriuretic response to the rapid intravenous administration of sodium is well established (1-5). Other hypertensive states have also been reported to exhibit an exaggerated natriuretic response to acute salt loading (4, 6-11). Although the mechanisms responsible for this phenomenon remain unknown (5), one of the possibilities is that the elevation of the blood pressure per se may cause the enhanced natriuretic (10).

In the present investigation the natriuretic response to acutely administered sodium was studied under controlled conditions in normotensive subjects and after elevation of the blood pressure with the pressor amine, metaraminol. Subjects excreted two to four times as much sodium as when responding to a similar sodium load without metaraminol.

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Methods

Five normotensive male subjects (three Caucasians and two Negroes), ages 33 to 48, without evidence of cardiac, hepatic, or renal disease were studied. Three had minimal healing dermatological lesions, one had inactive duodenal ulcer, and the other had minimal inactive pulmonary tuberculosis.

Three subjects (group I, no. 1, 2, and 3) received a total daily intake of 150 mEq of sodium. They were provided a diet containing 10 mEq of sodium daily and in addition were given 35 mEq of sodium chloride in gelatin capsules with each meal and at bedtime. Two subjects (group II, no. 4 and 5) received a total daily sodium intake of 10 mEq. Daily urine excretion of sodium was determined, and the patients were in sodium balance at the time each of the following studies was performed. The particular chronological order in which these studies were done was varied among the subjects.

Group I. Daily sodium intake of 150 mEq

1) "Blank day." The subjects had breakfast and drank 20 ml of tap water per kg of body weight between 7 and 8 a.m. At 8 a.m. the subjects lay down and remained recumbent until 3 p.m., except for standing to void at half-hour intervals. After each voiding they were given by mouth 5 g of carbohydrate and sufficient water to maintain a stable urine flow. Blood pressure and pulse were recorded at 30-minute intervals. The sodium chloride capsules were omitted on the morning of each day of study. At 3 p.m., when the day's study was completed, the two omitted doses of salt were administered so that the daily total oral intake of sodium remained 150 mEq.

2) "Metaraminol control day." The protocol was the same as on "blank day," except that 10 to 16 mg of metaraminol (Aramine),¹ dissolved in 5% dextrose in water, was given intravenously from 9:30 a.m. to 1 p.m. at a rate (average of 53 μ g per minute) adjusted to raise the mean blood pressure an average of 27 mm Hg

¹ Merck Sharp & Dohme, Inc., Philadelphia, Pa.

TABLE I
Sodium excretion in subjects provided a 150-mEq sodium diet

Subject Age Blood pressure	Time	Mean blood pressure	Na	Flow	Clearances		
					CCr	CIn	CPAH
		mm Hg	$\mu\text{Eq}/\text{min}$	ml/min	ml/min	ml/min	ml/min
Blank day							
1 38 110/68	7:41- 9:30 a.m.	75	163	11.2	145	156	707
	9:30-10:30 a.m.	81	160	9.6	147	151	685
	10:30-12:00 noon	83	155	4.9	136	148	626
	12:00- 1:00 p.m.	83	143	8.8	129	156	667
	1:00- 3:00 p.m.	89	144	8.3	126	150	588
Metaraminol control day*							
	7:33- 9:30 a.m.	81	227	12.4	139	137	652
	9:30-10:30 a.m.	105	191	11.4	114	146	568
	10:30-12:00 noon	110	219	11.6	132	153	574
	12:00- 1:00 p.m.	105	228	12.1	146	149	556
	1:00- 3:00 p.m.	91	208	9.1	143	148	604
Saline control day†							
	7:38- 9:30 a.m.	83	206	12.5	148	156	758
	9:30-10:30 a.m.	82	232	11.6	140	155	660
	10:30-12:00 noon	88	314	13.1	147	152	606
	12:00- 1:00 p.m.	87	386	14.4	144	169	609
	1:00- 3:00 p.m.	82	328	9.0	152	164	644
Metaraminol-saline day‡							
	7:37- 9:31 a.m.	78	224	14.2	178	186	862
	9:31-10:30 a.m.	94	207	14.0	186	197	699
	10:30-12:00 noon	113	413	18.6	176	199	667
	12:00- 1:00 p.m.	108	520	18.8	178	194	679
	1:00- 3:00 p.m.	88	378	10.7	175	181	709
Blank day							
2 48 125/82	8:00- 9:30 a.m.	103	162	10.4	118		
	9:30-10:30 a.m.	100	106	9.1	113		
	10:30-12:00 noon	96	100	6.8	101		
	12:00- 1:00 p.m.	95	93	6.1	105		
	1:00- 3:00 p.m.	96	83	5.4	100		
Metaraminol control day*							
	8:30- 9:30 a.m.	90	77	5.3	127		
	9:30-10:30 a.m.	107	75	5.9	133		
	10:30-12:00 noon	112	88	6.5	123		
	12:00- 1:00 p.m.	111	85	7.4	122		
	1:00- 3:00 p.m.	90	79	4.8	123		
Saline control day†							
	8:30- 9:30 a.m.	87	157	9.6	148		
	9:30-10:30 a.m.	90	130	8.7	148		
	10:30-12:00 noon	95	199	9.3	145		
	12:00- 1:00 p.m.	97	198	8.3	145		
	1:00- 3:00 p.m.	100	138	6.2	136		
Metaraminol-saline day‡							
	8:17- 9:30 a.m.	87	114	9.2	109		
	9:30-10:30 a.m.	110	134	10.3	116		
	10:30-12:00 noon	117	239	9.1	111		
	12:00- 1:00 p.m.	113	357	9.1	110		
	1:00- 3:00 p.m.	86	105	4.5	112		

* Metaraminol was infused intravenously from 9:30 a.m. to 1:00 p.m.

† Two thousand ml of sodium chloride-lactate solution was infused intravenously from 10:30 a.m. to 12:00 noon.

‡ Metaraminol was infused intravenously from 9:30 a.m. to 1:00 p.m., and 2,000 ml of sodium chloride-lactate solution was administered intravenously from 10:30 a.m. to 12:00 noon.

TABLE 1—(Continued)

Subject Age Blood pressure	Time	Mean blood pressure	Na	Flow	Clearances		
					C _{Cr}	C _{In}	C _{PAH}
		mm Hg	$\mu\text{Eq}/\text{min}$	ml/min	ml/min	ml/min	ml/min
Blank day							
3 36 118/78	8:30–9:30 a.m.	88	127	8.5	95		
	9:30–10:30 a.m.	88	120	7.5	84		
	10:30–12:00 noon	92	66	5.1	67		
	12:00–1:00 p.m.	95	78	4.1	81		
	1:00–3:00 p.m.	92	69	4.6	99		
Metaraminol control day*							
	8:01–9:30 a.m.	89	106	10.8	102		
	9:30–10:30 a.m.	112	87	8.5	101		
	10:30–12:00 noon	120	100	6.2	99		
	12:00–1:00 p.m.	119	102	6.9	92		
	1:00–3:00 p.m.	95	61	2.8	95		
Saline control day†							
	8:30–9:30 a.m.	90	138	9.3	124		
	9:30–10:30 a.m.	90	135	6.8	113		
	10:30–12:00 noon	89	144	8.4	103		
	12:00–1:00 p.m.	90	156	7.0	99		
	1:00–3:00 p.m.	90	139	4.5	94		
Metaraminol-saline day‡							
	8:04–9:30 a.m.	86	173	9.7	133		
	9:30–10:30 a.m.	111	211	9.5	109		
	10:30–12:00 noon	123	286	12.0	114		
	12:00–1:00 p.m.	124	197	10.8	111		
	1:00–3:00 p.m.	99	187	3.7	119		

above the premetaraminol level. The blood pressure and pulse rate were recorded every 2 to 5 minutes. The mean blood pressure was calculated as the diastolic pressure plus one-third of the pulse pressure.

3) "Saline control day." The protocol was the same as on "blank day," except that 2,000 ml of a solution containing 125 mEq per L of sodium, 100 mEq per L of chloride, and 25 mEq per L of lactate was administered intravenously from 10:30 a.m. to 12 noon.

4) "Metaraminol-saline day." The protocol was the same as on "blank day," except that the metaraminol solution was infused between 9:30 a.m. and 1 p.m. as described for the "metaraminol control day" and that the hypotonic sodium solution was given from 10:30 a.m. to 12 noon as described for the "saline control day."

Group II. Daily sodium intake of 10 mEq

These two subjects were studied as described for "saline control day" and "metaraminol-saline day." The protocols were the same, except that the oral water load was 500 ml.

Creatinine, sodium, potassium, chloride, and total solutes were measured in serum and urine. Endogenous creatinine (C_{Cr}), osmolal (C_{osm}), and free water (C_{H_2O}) clearances were calculated in all subjects. Inulin and para-aminohippurate clearances (C_{In} , C_{PAH}) were estimated in Subject 1 throughout the studies by a standard technique (12) and without catheterization of the bladder. Sodium and potassium were measured with a Baird flame photometer with an internal lithium standard. The Aminco-Cotlove automatic chloride titrator was

FIG. 1. SODIUM EXCRETION IN ONE SUBJECT (No. 1) ON "BLANK DAY," "METARAMINOL CONTROL DAY," "SALINE CONTROL DAY," AND "METARAMINOL-SALINE DAY." Daily sodium intake, 150 mEq.

used to measure chloride. Creatinine was determined in urine by the method of Peters (13) and in serum by the technique of Hare (14). Inulin was measured by the anthrone procedure of Young and Raisz (15). Para-aminohippurate was determined by the method of Goldring and Chasis (16). The Fiske osmometer was employed to measure concentration of total solutes.

Results

1) "*Blank day.*" The average mean blood pressure for group I was 90 mm Hg. C_{Cr} , C_{In} , and C_{PAH} were within the normal range, except for Subject 3, who on this day had a moderately low C_{Cr} (Table I).

2) "*Metaraminol control day.*" During metaraminol infusion the average mean blood pressure for group I was 111 mm Hg. The rate of excretion of sodium was unaltered (Table I, Figure 1). There was no difference in the maximal increase of sodium excretion from the mean preinfusion rate between the "metaraminol control day" and the "blank day" (Table III, Figure 2). Neither

C_{Cr} nor C_{In} changed significantly (more than 15%) (Table I).

3) "*Saline control day.*" After the sodium infusion there was no change in mean blood pressure. The rate of excretion of sodium increased in the five subjects studied (Tables I, II, and III; Figures 1 and 2).

In four of the five subjects, during the saline administration neither C_{Cr} nor C_{In} was significantly different from the preinfusion periods (Tables I and II). Subject 3 had a slight decrease in C_{Cr} . Subject 1 had a 16% decrease in C_{PAH} . After the salt load there were no significant changes in the serum concentration of sodium (-0.4 to $+3$ mEq per L), chloride (-3.2 to $+3.9$ mEq per L), or potassium (-0.3 to $+0.2$ mEq per L).

4) "*Metaraminol-saline day.*" The average mean blood pressure during the metaraminol and saline infusions was 112 mm Hg. The excretion of sodium increased in all the subjects (Tables I,

FIG. 2. SODIUM EXCRETION IN THREE SUBJECTS (Nos. 1, 2, 3) ON "BLANK DAY," "METARAMINOL CONTROL DAY," "SALINE CONTROL DAY," AND "METARAMINOL-SALINE DAY." Daily sodium intake, 150 mEq. The maximal increase of sodium excretion (from 9:30 a.m. to 3 p.m.) from the mean preinfusion rate (8:30 to 9:30 a.m.) in microequivalents per minute (peak rate minus preinfusion rate) is on the vertical scale. Each column represents an individual subject studied on the 4 experimental days. (On "blank day," Subject 2 had the highest rate of sodium excretion before 9:30 a.m.)

TABLE II
Sodium excretion in subjects provided a
10-mEq sodium diet

Subject	Age	Blood pressure	Time	Mean blood pressure	Na	Flow	C _{Cr}
				mm Hg	μEq/min	ml/min	ml/min
Saline control day*							
4 38 118/70			8:00- 9:30 a.m.	84	16	2.8	163
			9:30-10:30 a.m.	84	19	1.5	148
			10:30-12:00 noon	87	122	9.8	144
			12:00- 1:00 p.m.	86	174	4.3	143
			1:00- 3:00 p.m.	80	190	2.6	153
Metaraminol-saline day†							
			8:00- 9:30 a.m.	84	44	1.1	152
			9:30-10:30 a.m.	102	46	3.7	144
			10:30-12:01 p.m.	110	745	13.5	141
			12:01- 1:00 p.m.	114	1,429	15.9	147
			1:00- 3:00 p.m.	82	326	2.6	131
			Saline control day*				
5 33 110/64			8:00- 9:30 a.m.	74	25	7.9	130
			9:30-10:30 a.m.	73	34	12.2	120
			10:30-12:00 noon	73	56	8.7	117
			12:00- 1:00 p.m.	78	130	5.6	118
			1:00- 3:00 p.m.	74	176	8.1	123
Metaraminol-saline day†							
			8:00- 9:30 a.m.	80	15	8.3	103
			9:30-10:30 a.m.	96	13	10.3	107
			10:30-12:00 noon	119	95	8.7	112
			12:00- 1:00 p.m.	117	263	6.5	114
			1:00- 2:41 p.m.	88	159	5.1	114

* Two thousand ml of sodium chloride-lactate solution was infused intravenously from 10:30 a.m. to 12:00 noon.

† Metaraminol was infused intravenously from 9:30 a.m. to 1:00 p.m., and 2,000 ml of sodium chloride-lactate solution was administered intravenously from 10:30 a.m. to 12:00 noon.

II, and III). The maximal increase of sodium excretion from the mean preinfusion rate on this day was an average of 261% greater than on the "saline control day" in the three subjects of group I (Figure 2).²

Similar results were obtained in the two subjects studied while taking the 10-mEq sodium diet. Figure 3 illustrates the results in one of these subjects.

There were no significant changes in C_{Cr} or C_{In} after the metaraminol and saline infusions in four of the five subjects (Tables I and II). Subject 3 had a slight decrease in C_{Cr}. In Subject 1 the C_{PAH} decreased 21% after the vasopressor infusion.

² Angiotensin II was employed in Subject 1 with different results. The excretion of sodium after the administration of angiotensin II was markedly reduced, as described by others (17-20). This effect of angiotensin II on sodium excretion was not markedly modified by the rapid iv administration of sodium.

FIG. 3. SODIUM EXCRETION IN ONE SUBJECT (No. 4) ON "SALINE CONTROL DAY" AND "METARAMINOL-SALINE DAY." Daily sodium intake, 10 mEq.

TABLE III
Sodium excretion in normotensive subjects after blood pressure elevation with metaraminol

Subject	Experimental day	Rate of sodium excretion		
		Preinfusion*	Peak†	Peak - preinfusion‡
		μEq/min	μEq/min	μEq/min
1	Blank	154	166	12
	Metaraminol control	232	236	4
	Saline control	213	410	197
	Metaraminol-saline	216	620	404
2	Blank	153	121	-32§
	Metaraminol control	77	104	27
	Saline control	157	239	82
	Metaraminol-saline	90	374	284
3	Blank	127	139	12
	Metaraminol control	103	113	10
	Saline control	138	162	24
	Metaraminol-saline	187	315	128
4	Saline control	21	212	191
	Metaraminol-saline	58	1,842	1,784
5	Saline control	34	194	160
	Metaraminol-saline	23	323	300

* Mean rate of sodium excretion from 8:30 to 9:30 a.m.

† Maximal 30-minute period of sodium excretion from 9:30 a.m. to 3:00 p.m.

‡ Maximal increase in sodium excretion from the mean preinfusion rate.

§ On "blank day," Subject 2 had the highest rate of sodium excretion before 9:30 a.m.

During the saline infusion serum sodium concentration did not change significantly in four of the subjects (-0.1 to $+4.4$ mEq per L) and decreased 9.6 mEq per L in Subject 3. There were no significant changes in serum potassium concentration in four subjects (-0.6 to $+0.2$ mEq per L) and a decrease of 0.9 mEq per L in Subject 1. There were no significant changes in serum chloride concentration (-1.4 to $+3.8$ mEq per L).

The excretion of chloride and total solutes and urine flow generally followed the changes in sodium excretion in all the experimental days. There were no remarkable changes in the rate of excretion of potassium, except for a transient increase with the saline infusion in group II. The C_{osm} increased on the "metaraminol-saline day." There were no consistent changes in C_{osm} or C_{H_2O} in the other experimental days. The pulse rate decreased in all the subjects during the metaraminol infusion; Subject 2 had few isolated premature beats.

Discussion

The administration of metaraminol in amounts sufficient to elevate systemic blood pressure in normotensive subjects is evidently accompanied by an enhanced natriuretic response to rapidly administered sodium. The augmented natriuresis observed is modest in comparison to the markedly enhanced natriuresis observed in patients with essential hypertension (2, 5). Although this difference might be related to differences in the degree of hypertension or the duration of increased blood pressure, the present data do not permit further interpretation of this point.

Although the possibility of a direct effect of the vasopressor agent on sodium excretion cannot be excluded, it seems unlikely, considering that the administration of metaraminol alone did not result in increased sodium excretion. No significant changes in C_{Cr} or C_{In} were observed in this study.

This augmented sodium excretion was present at two different levels of salt intake, 10 and 150 mEq of sodium daily. Papper, Belsky, and Bleifer (5) demonstrated that the exaggerated natriuresis of patients with essential hypertension is apparent at all levels of salt intake ranging from 10 to 300 mEq daily.

There are observations implicating elevated blood pressure in the exaggerated natriuresis of hypertension. This phenomenon has been repeatedly reported in patients with essential hypertension (1-5). Patients with hypertension caused by chronic glomerulonephritis (9, 10), Cushing's syndrome (4, 6, 7), pheochromocytoma (4), primary aldosteronism (8, 11), and unilateral renal disease (10) have also been reported to have an exaggerated natriuretic response to acute salt loading, suggesting that elevated blood pressure could be a common denominator in this phenomenon.

Selkurt, Hall, and Spencer (21) and more recently Tobian, Coffee, Ferreira, and Meuli (22) reported studies in animals indicating that an increase in renal arterial perfusion pressure results in augmented sodium excretion. The mechanisms whereby this occurred were not clear. Tobian and his associates suggested the possibility of some intrarenal humoral mediation, and Selkurt and co-workers considered some intrarenal hemodynamic alterations.

The present data do not permit further delineation of the precise mechanisms involved in the enhanced natriuresis observed when a salt load is administered after systemic blood pressure has been elevated with metaraminol in normotensive subjects.

Summary

The natriuretic response to an acutely administered intravenous sodium solution (125 mEq per L) was studied under controlled conditions in five normotensive subjects and after elevation of the blood pressure with the pressor amine, metaraminol.

Under the described conditions, there was an augmented natriuretic response to the rapid administration of sodium in the five normotensive subjects. This augmented response was present at two different levels of dietary sodium intake. The vasopressor agent alone did not increase sodium excretion.

The mechanisms whereby this enhanced natriuresis occurs in normotensive subjects under the described conditions are not known.

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