

Inhibition of Bicarbonate Absorption by Peptide Hormones and Cyclic Adenosine Monophosphate in Rat Medullary Thick Ascending Limb

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Abstract

In vitro microperfusion experiments were performed to examine the effects of peptide hormones on bicarbonate and ammonium transport by the medullary thick ascending limb (MTAL) of the rat. Arginine vasopressin (AVP; 2.8×10^{-10} M in the bath) reduced bicarbonate absorption by 50% (from 7.8 to 3.7 pmol/min per mm). AVP caused a similar reduction in bicarbonate absorption in tubules perfused with 10^{-4} M furosemide to inhibit net NaCl absorption. Glucagon (2×10^{-9} M in the bath) also reduced bicarbonate absorption (from 11.7 to 7.6 pmol/min per mm). The inhibition of bicarbonate absorption could be reproduced with either exogenous 8-bromo-cAMP or forskolin. With 8-bromo-cAMP (10^{-3} M) in the bath, addition of vasopressin to the bath did not significantly affect bicarbonate absorption. PTH significantly inhibited bicarbonate absorption, but the extent of inhibition was less than that observed with either AVP or glucagon. Vasopressin had no effect on net ammonium absorption in MTAL perfused and bathed with 4 mM NH_4Cl . These findings indicate that: (a) vasopressin, glucagon, and PTH directly inhibit bicarbonate absorption in the MTAL of the rat; (b) this inhibition occurs independent of effects on net NaCl absorption and appears to be mediated in part by cAMP; and (c) HCO_3^- and NH_4^+ absorption can be regulated independently in the MTAL. (*J. Clin. Invest.* 1990; 85:1006–1013.) vasopressin • cyclic AMP • thick ascending limb • renal bicarbonate transport • peptide hormones

Introduction

Medullary and cortical thick ascending limbs from rats absorb bicarbonate in vitro at rates that are comparable to rates measured in isolated, perfused cortical and medullary collecting ducts and that are sufficient to account for much of the 10–15% of filtered bicarbonate that is reabsorbed between the late proximal convoluted and early distal tubule of the rat in vivo (1, 2). The bicarbonate absorption occurs against both chemical and electrical gradients and is mediated predomi-

nantly by apical membrane sodium–hydrogen exchange (3). In contrast to segments of the proximal tubule and collecting duct, however, little direct information is available regarding factors that control bicarbonate absorption in the mammalian thick ascending limb.

Peptide hormones such as vasopressin, glucagon (Glu),¹ and PTH regulate a variety of cell functions in thick ascending limbs, including the transport of ions such as Na^+ , Cl^- , Ca^{2+} , and K^+ (4–8). Peptide hormones also have important effects on urinary acidification, both in physiologic and pathophysiologic conditions (9–13), and recent evidence suggests that the thick ascending limb may be a site of peptide hormone action on bicarbonate transport. Micropuncture studies in rats have shown that the acute infusion of vasopressin, Glu, or PTH can alter bicarbonate reabsorption by the superficial “loop segment,” the portion of the nephron between the late proximal convoluted and early distal tubule (10, 12, 14, 15). While these findings are consistent with an effect of peptide hormones on thick ascending limb bicarbonate transport, interpretation of the micropuncture experiments is complicated by the fact that the loop segment is composed of the proximal straight tubule, the thin descending limb, the thin ascending limb, and the early distal convoluted tubule, in addition to the medullary and cortical thick ascending limbs. Such heterogeneity precludes precise localization of the site(s) of hormone action. In addition, in experiments in which hormones are infused systemically, it is not possible to determine whether a change in bicarbonate absorption may be due to a direct action of the hormone on a particular nephron segment or to a change in some luminal or systemic factor that is related to the hormone and that indirectly influences bicarbonate transport. The possibility that different peptide hormones may affect different nephron segments or act via different cellular mechanisms is supported by the findings that vasopressin and Glu inhibited, whereas PTH stimulated, bicarbonate absorption in the loop segment (10, 14).

The purpose of the present study was to determine directly whether peptide hormones affect bicarbonate absorption in the medullary thick ascending limb of the rat. The isolated, perfused tubule technique was used so that the effects of different hormones on bicarbonate transport could be examined in the absence of changes in other systemic factors. In addition, experiments were performed to gain information regarding the signal transduction system that may be involved in the hormones' effects. Finally, because the medullary thick ascending limb also influences urinary net acid excretion by reabsorbing ammonium (1, 16–18), the effect of vasopressin on NH_4^+ transport also was examined. The results demonstrate that vasopressin, Glu, and PTH directly inhibit bicarbonate absorption in the rat medullary thick ascending limb, and that

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1. Abbreviations used in this paper: 8-Bromo-cAMP, 8-bromoadenosine 3':5'-cyclic monophosphate; Glu, glucagon.

the inhibition appears to be mediated in part by increased intracellular levels of cAMP. In contrast, vasopressin has no effect on net ammonium absorption, indicating that bicarbonate and ammonium transport rates can be regulated separately in this nephron segment.

Methods

Pathogen-free male Sprague-Dawley rats (Taconic Farms, Inc., Germantown, NY) weighing 60–70 g were housed in filter-top isolation cages with autoclaved food (NIH 31 diet; Ziegler Brothers, Inc., Gardners, PA) and bedding. They had free access to food and distilled water up to the time of experiments. The rats were injected intraperitoneally with 2 mg furosemide 5–10 min before being anesthetized with sodium pentobarbital (50 mg/kg i.p.). The left kidney was exposed via an abdominal incision, bathed in situ for 1–2 min with ice-cold dissecting solution, and then removed and sliced for tubule dissection. As discussed previously (1, 18, 19), these procedures help preserve the viability of rat renal tubules in vitro. Medullary thick ascending limbs were dissected at 10°C from the inner stripe of the outer medulla in a solution that contained (in mM): 145 NaCl, 2 K₂HPO₄, 1.2 Na₂SO₄, 1 CaCl₂, 1 Ca lactate, 0.5 Mg citrate, and 5.5 glucose. The dissection fluid was gassed with 100% oxygen and titrated to pH 7.4.

The tubules were transferred to a temperature-controlled chamber, mounted on concentric pipettes, and perfused in vitro at 37°C as previously described (1, 3, 18). Tubule fluid emerging from the distal end of the tubules was collected for timed intervals into calibrated constriction pipettes for calculation of tubule fluid flow rate and for analysis of total CO₂ or total ammonia concentration. In bicarbonate transport experiments, perfusion rate averaged 1–2 nl/min per mm; in ammonium transport experiments, a higher perfusion rate (5–6 nl/min per mm) was used to prevent development of limiting transepithelial ammonium concentration gradients (18). In all experiments the luminal perfusion solution contained (in mM): 146 Na⁺, 4 K⁺, 122 Cl⁻, 25 HCO₃⁻, 2 Ca²⁺, 1.5 Mg²⁺, 2 phosphate, 1.2 SO₄²⁻, 1 citrate, 2 lactate, and 5.5 glucose. The bath solution was identical except for addition of 0.2% bovine albumin (fraction V, essentially fatty acid free; Sigma Chemical Co., St. Louis, MO). Experimental agents were added to the bath solution as described below. In ammonium transport experiments, the perfusate and bath also contained 4 mM NH₄Cl. All solutions were equilibrated with 95% O₂, 5% CO₂ and were pH 7.47–7.48 at 37°C. The oxygen and CO₂ tensions in the bath were maintained during the experiments by continuously flowing freshly gassed solution into the perfusion chamber as previously described (1, 18).

Arginine vasopressin (AVP), Glu, PTH (1–34 fragment, bovine), 8-bromoadenosine 3':5'-cyclic monophosphate (8-bromo-cAMP), and forskolin were purchased from Sigma Chemical Co. Glu was prepared as a concentrated stock solution in dilute HCl; forskolin was prepared and stored as a stock solution (1 mg/ml) in 95% ethanol. Final ethanol concentration in forskolin bath was 0.05%. An equal concentration of ethanol was added to the control bath in all forskolin experiments.

The general protocol was similar with each of the experimental agents. Tubules were equilibrated for 20–60 min at 37°C with the control bath solution² and the perfusion reservoir was adjusted to establish a steady luminal flow rate. After equilibration, two to four tubule fluid collections were obtained for determination of control bicarbonate or ammonium transport rate (control period). The bath solution was then changed to one containing one of the experimental agents and the tubule allowed to re-equilibrate for 10–30 min. Two to

2. After warming tubules to 37°C, the bicarbonate absorption rate increases to a stable value over a period of ~ 30 min. Therefore, a 20–60-min equilibration period was allowed. The increase in bicarbonate absorption presumably reflects a change in some regulatory factor, possibly a fall in cell cAMP (see Results).

Table I. Effect of AVP on Bicarbonate Transport

	V	[Total CO ₂]		JT _{CO₂}	V _{TE}
		Perfused	Collected		
	nl/min per mm	mM		pmol/min per mm	mV
Control	1.0±0.1	25.9±0.2	17.2±1.0*	7.8±0.4	12.5±1.1
Vasopressin	0.9±0.1	25.9±0.2	21.5±0.8*	3.7±0.5	15.1±0.7
P	NS		<0.001	<0.001	<0.05

Values are means±SE. Vasopressin was 2.8×10^{-10} M in the bath. No. of experiments = 8 (means do not include values from three tubules in Fig. 1 studied with NH₄Cl in perfusate and bath). Mean tubule length = 0.57 ± 0.02 mm. Total CO₂ concentration in the bath was 26.8 ± 0.4 mM in control and 26.7 ± 0.3 mM with vasopressin. V is fluid collection rate; JT_{CO₂} is absolute rate of total CO₂ absorption; V_{TE} is transepithelial voltage, oriented lumen positive with respect to bath. P values compare control and vasopressin (paired t test). * Collected total CO₂ concentration significantly different from perfused (paired t test).

seven additional tubule fluid collections were then obtained (experimental period). Finally, the agent was removed from the bath and control measurements were repeated (recovery period). Complete recovery generally was observed within 30–40 min. In some experiments, only two periods (either control and experimental or experimental and recovery) were studied. Also, longer time periods were used with some agents (see Results).

Total carbon dioxide concentrations in perfusion and bathing solutions and in collected tubule fluid were measured by microcalorimetry as previously described (1, 3). Total ammonia concentrations were measured by microfluorometry (1). Transepithelial voltage was measured between calomel cells using a digital electrometer (1, 18). Because net fluid transport is absent in rat thick ascending limbs in both the presence and absence of vasopressin (1, 19–21), absolute rates of total CO₂ or total ammonia absorption (J_i, picomoles/minute per millimeter) were calculated as $J_i = V(C_o - C_i)/L$, where V is fluid collection rate (nanoliters/minute), C is total CO₂ or total ammonia concentration in perfused (o) and collected (i) fluid (millimolar), and L is perfused tubule length (millimeters).³ A mean fluid collection rate, transepithelial voltage, and total CO₂ or total ammonia absorption rate were calculated for each period studied in a given tubule. When control measurements were made at the beginning and end of an experiment, the control values were averaged. Single tubule values, presented in the figures, were averaged to obtain the group means presented in tables (n = No. of tubules). Differences between means were evaluated using the paired t test, with a P value < 0.05 considered statistically significant.

Results

Effect of vasopressin on bicarbonate absorption. The effects of AVP on bicarbonate transport by the medullary thick ascending limb are shown in Table I (mean values) and Fig. 1 (individual data). Vasopressin (2.8×10^{-10} M) added to the bath reduced net bicarbonate absorption from 7.8 to 3.7 pmol/min per mm (Table I). The inhibition of bicarbonate absorption

3. In this paper, as in previous papers (1, 3, 17), total CO₂ absorption is referred to generally as bicarbonate absorption and total ammonia absorption is referred to as ammonium absorption. When mechanisms of ammonium transport are discussed, the chemical formulae NH₄⁺ and NH₃ are used to indicate the particular species transported.

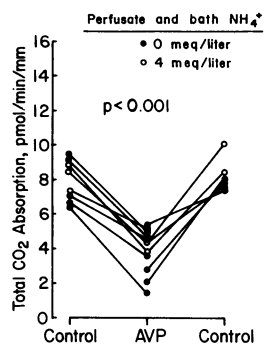


Figure 1. Effect of AVP (2.8×10^{-10} M in the bath) on total CO_2 absorption by the rat medullary thick ascending limb. Solid and open circles are mean values for single tubules; lines connect paired measurements made in the same tubule. P value is for paired t test.

was reversible (Fig. 1) and could be observed within 10–15 min after addition of vasopressin to the bath. Vasopressin caused a small but significant increase in the transepithelial voltage (Table I). In three additional tubules studied with 4 mM NH_4Cl in the perfusate and bath, vasopressin (2.8×10^{-10} M in the bath) reduced bicarbonate absorption rate from 8.3 ± 0.5 to 4.9 ± 0.6 pmol/min per mm (open circles, Fig. 1).

Effect of vasopressin on bicarbonate absorption in the presence of furosemide. To determine whether the inhibition of bicarbonate absorption may have been the indirect result of an effect of vasopressin on transcellular NaCl transport, the effect of vasopressin was studied in tubules perfused with furosemide to inhibit net NaCl absorption (Table II, Fig. 2). With 10^{-4} M furosemide in the luminal perfusate, vasopressin (2.8×10^{-10} M in the bath) reduced net HCO_3^- absorption from 11.5 to 8.1 pmol/min per mm (Table II). The inhibition was reversed when vasopressin was removed from the bathing solution (Fig. 2). Thus, the effect of vasopressin to inhibit HCO_3^- absorption was similar in the presence and absence of furosemide. With furosemide, the transepithelial voltage was near zero and was unaffected by vasopressin (Table II).

Effect of Glu on bicarbonate absorption. The effects of Glu are shown in Table III and Fig. 3. Glu (2×10^{-9} M) added to the bath reduced net bicarbonate absorption from 11.7 to 7.6 pmol/min per mm (Table III). This effect was reversible (Fig. 3). Glu increased the transepithelial voltage in four of five tubules; however, the net increase in voltage was small and was of borderline significance ($0.05 < P < 0.1$).

Table II. Effect of AVP on Bicarbonate Transport (10^{-4} M Luminal Furosemide)

	V nl/min per mm	[Total CO_2]		$J\text{TCO}_2$ pmol/min per mm	V_{TE} mV
		Perfused mM	Collected mM		
Control	1.1 ± 0.1	25.4 ± 0.2	$15.1 \pm 0.8^*$	11.5 ± 1.0	1.5 ± 0.2
Vasopressin	1.1 ± 0.1	25.4 ± 0.2	$18.0 \pm 1.1^*$	8.1 ± 1.2	1.3 ± 0.1
P	NS		<0.005	<0.005	NS

Values are means \pm SE. Vasopressin was 2.8×10^{-10} M in the bath. In both control and vasopressin conditions, the luminal perfusate contained 1×10^{-4} M furosemide. No. of experiments = 5. Mean tubule length = 0.49 ± 0.03 mm. Total CO_2 concentration in the bath was 25.7 ± 0.4 mM in control and 25.9 ± 0.5 mM with vasopressin. V , $J\text{TCO}_2$, V_{TE} , P values, and * as in Table I.

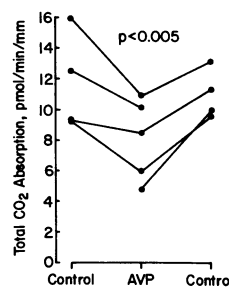


Figure 2. Effect of AVP (2.8×10^{-10} M in the bath) on total CO_2 absorption in the presence of furosemide. Furosemide (1×10^{-4} M) was present in the luminal perfusate throughout the experiments. Solid circles, lines, and P value as in Fig. 1.

Effect of PTH on bicarbonate absorption. PTH (2×10^{-9} or 3×10^{-8} M) added to the bath caused a small but significant decrease in net bicarbonate absorption (Table IV, Fig. 4). The inhibition of bicarbonate absorption was reversible (Fig. 4) but was smaller in magnitude (19%) than that observed with vasopressin or Glu (35–50%, Tables I and III). PTH had no significant effect on transepithelial voltage (Table IV).

Effects of 8-bromo-cAMP and forskolin on bicarbonate absorption. To determine whether vasopressin and Glu may have inhibited bicarbonate absorption via their action to stimulate intracellular production of cAMP, the effects of exogenous 8-bromo-cAMP and forskolin (an agent that increases cell cAMP production) were examined in the absence of added hormone. 8-Bromo-cAMP (1×10^{-3} M in the bath) reduced net bicarbonate absorption by 37%, from 13.1 to 8.2 pmol/min per mm (Table V, Fig. 5). Forskolin (1×10^{-6} M in the bath) reduced bicarbonate absorption by 30%, from 16.7 to 12 pmol/min per mm (Table V, Fig. 6). The effect of both agents to inhibit bicarbonate absorption was reversible (Figs. 5 and 6). Both 8-bromo-cAMP and forskolin significantly increased the transepithelial voltage (Table V).

Effect of vasopressin on bicarbonate absorption in the presence of 8-bromo-cAMP. To examine further whether cAMP may be responsible for the transport effects observed with vasopressin, the effects of vasopressin were studied in the presence of exogenous 8-bromo-cAMP. With 8-bromo-cAMP (1×10^{-3} M) in the bath, adding vasopressin (2.8×10^{-10} M) to the bath had no significant effect on bicarbonate absorption (Table VI, Fig. 7). In the presence of 8-bromo-cAMP, vasopressin also had no significant effect on transepithelial voltage (Table VI).

Effect of vasopressin on ammonium absorption. The effect of AVP on ammonium transport by the medullary thick as-

Table III. Effect of Glu on Bicarbonate Transport

	V nl/min per mm	[Total CO_2]		$J\text{TCO}_2$ pmol/min per mm	V_{TE} mV
		Perfused mM	Collected mM		
Control	1.1 ± 0.1	25.3 ± 0.3	$14.7 \pm 0.7^*$	11.7 ± 1.0	7.8 ± 0.8
Glu	1.1 ± 0.1	25.3 ± 0.3	$18.4 \pm 0.6^*$	7.6 ± 0.8	9.6 ± 1.2
P	NS		<0.005	<0.005	NS

Values are means \pm SE. Glu was 2×10^{-9} M in the bath. No. of experiments = 5. Mean tubule length = 0.56 ± 0.03 mm. Total CO_2 concentration in the bath was 25.4 ± 0.4 mM in control and 25.4 ± 0.4 mM with Glu. V , $J\text{TCO}_2$, V_{TE} , P values, and * as in Table I.

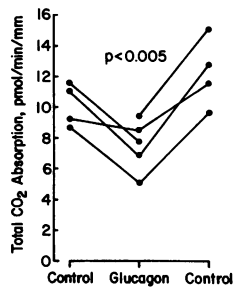


Figure 3. Effect of Glu (2×10^{-9} M in the bath) on total CO_2 absorption by the rat medullary thick ascending limb. Solid circles, lines, and P value as in Fig. 1.

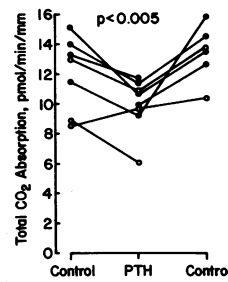


Figure 4. Effect of PTH in the bath on total CO_2 absorption by the rat medullary thick ascending limb. Solid circles, 2×10^{-9} M PTH; open circles, 3×10^{-8} M PTH. Lines and P value as in Fig. 1.

ending limb is shown in Table VII and Fig. 8. Vasopressin (2.8×10^{-10} M) added to the bath did not significantly affect net ammonium absorption. To determine whether exposure times longer than those used in the bicarbonate transport experiments were required to observe an effect of vasopressin on ammonium absorption, tubules in some of the experiments were exposed to bath vasopressin for up to 100 min after an initial 60–90-min control period. Even with these prolonged time periods, no effect of vasopressin on ammonium absorption was observed.

Discussion

Peptide hormones such as vasopressin and PTH influence urinary net acid excretion and have been shown to alter luminal acidification in the proximal tubule, distal tubule, and cortical and inner medullary collecting ducts (9–15, 22–26). The present experiments were designed to determine directly whether peptide hormones affect bicarbonate absorption in the medullary thick ascending limb of the rat. The results show that vasopressin, Glu, and PTH inhibit bicarbonate absorption, and that this inhibition can be reproduced by procedures that increase intracellular levels of cAMP. Thus, peptide hormones and factors that regulate intracellular cAMP are likely to be important determinants of luminal acidification in the medullary thick ascending limb.

Effects of vasopressin on bicarbonate transport. Bicarbonate absorption was markedly inhibited by 2.8×10^{-10} M vasopressin (Fig. 1). This hormone concentration was used because it represents the upper end of the physiologic range of vasopressin concentrations, and because it had been shown pre-

viously to stimulate net NaCl absorption in the isolated, perfused rat medullary thick ascending limb (21). The inhibition of bicarbonate absorption was observed with or without NH_4Cl in the perfusate and bath, and was sustained for as long as vasopressin was present in the bathing solution.

The direct inhibition of bicarbonate absorption in the medullary thick ascending limb *in vitro* can explain, in part, the recent observation that an acute infusion of 1-deamino-8-D- AVP decreased fractional bicarbonate reabsorption in the loop segment of the Sprague-Dawley rat *in vivo* (12). The inhibition *in vivo* was observed in hormone-deprived rats in which the potential effects of endogenous peptide hormones were minimized by hypotonic volume expansion, thyroparathyroidectomy, and somatostatin infusion, but could not be detected in intact rats that received only hypotonic volume expansion (12). These findings suggest that factors other than vasopressin influence loop segment bicarbonate transport; however, they do not preclude an important regulatory role for the action of vasopressin on the medullary thick ascending limb for several reasons: (a) the medullary thick ascending limb may be relatively short in superficial nephrons, minimizing the impact of vasopressin on overall loop bicarbonate transport; (b) changes in bicarbonate absorption in the cortical thick ascending limb may compensate for changes in the medullary segment (see below); and (c) direct effects of vasopressin on the medullary thick ascending limb may be opposed by other luminal or systemic factors that change with vasopressin and that indirectly influence loop bicarbonate transport.

In contrast to its action to inhibit bicarbonate absorption in the medullary thick ascending limb, vasopressin stimulates net bicarbonate absorption in the distal tubule and collecting ducts (12, 25, 26) and increases urinary net acid excretion (11, 12). The physiologic importance of increased net acid excretion with vasopressin may be to maintain plasma pH during states of antidiuresis when water retention would tend to dilute the plasma bicarbonate concentration (12). A possible means by which the thick ascending limb could contribute to the vasopressin-related stimulation of urine acidification would be as follows. If vasopressin inhibited bicarbonate absorption in the medullary thick ascending limb but not in the cortical thick ascending limb (as is the case for vasopressin stimulation of NaCl absorption [6]), then an increase in bicarbonate absorption by the cortical segment, in response to the increased bicarbonate load, could compensate for the decrease in absorption in the medullary segment. The net result would be a shift in bicarbonate absorption from the medullary to the cortical thick ascending limb with little or no change in overall loop bicarbonate transport. This would have the important effect of reducing net bicarbonate delivery to the medullary interstitium, which could limit the bicarbonate concentration

Table IV. Effect of PTH on Bicarbonate Transport

	V	[Total CO_2]		$J\text{TCO}_2$	V_{TE}
		Perfused	Collected		
	nl/min per mm	mM	mM	pmol/min per mm	mV
Control	1.1 ± 0.1	25.2 ± 0.4	$14.4 \pm 0.8^*$	12.5 ± 0.8	9.4 ± 1.1
PTH	1.2 ± 0.1	25.2 ± 0.4	$16.6 \pm 0.7^*$	10.1 ± 0.7	10.0 ± 0.5
P	NS		<0.001	<0.005	NS

Values are means \pm SE and are combined results of experiments with 2×10^{-9} and 3×10^{-8} M PTH in the bath. Total no. of experiments = 8 ($n = 5$ with 2×10^{-9} M PTH; $n = 3$ with 3×10^{-8} M PTH). Mean tubule length = 0.50 ± 0.03 mm. Total CO_2 concentration in bath was 25.5 ± 0.2 mM in control and 25.6 ± 0.4 mM with PTH. V , $J\text{TCO}_2$, V_{TE} , P values, and * as in Table I.

Table V. Effects of 8-Bromo-cAMP and Forskolin on Bicarbonate Transport

	V nl/min per mm	[Total CO ₂]		JTCO ₂ pmol/min per mm	V _{TE} mV
		Perfused mM	Collected mM		
Control	1.0±0.1	25.0±0.2	11.6±1.4*	13.1±1.5	11.9±0.8
8-Bromo-cAMP	1.0±0.1	25.0±0.2	16.3±1.4*	8.2±1.0	13.8±1.1
P	NS		<0.001	<0.001	≈0.05
Control	1.7±0.2	25.7±0.2	15.8±0.7*	16.7±1.4	6.6±0.6
Forskolin	1.7±0.2	25.7±0.2	18.5±0.2*	12.0±0.9	9.0±0.7
P	NS		<0.01	<0.01	<0.025

Values are means±SE. 8-Bromo-cAMP was 1×10^{-3} M and forskolin was 1×10^{-6} M in the bath. No. of experiments = 8 with 8-bromo-cAMP and 5 with forskolin. Mean tubule length was 0.57 ± 0.03 mm with 8-bromo-cAMP and 0.46 ± 0.05 mm with forskolin. Total CO₂ concentration in bath was 25.4 ± 0.2 mM in control, 25.5 ± 0.1 mM with 8-bromo-cAMP, and 25.9 ± 0.3 mM with forskolin. V, JTCO₂, V_{TE}, P values, and * as in Table I.

surrounding the medullary collecting ducts and enhance collecting duct proton secretion. Such an effect may be particularly important with vasopressin, since concentrations of medullary solutes are greatest during antidiuresis. Further studies of the effects of vasopressin on the cortical thick ascending limb and on medullary acid-base handling will be required to evaluate this hypothesis.

Mechanisms of vasopressin action. The effect of vasopressin to inhibit bicarbonate absorption can be accounted for by its action to stimulate adenylyl cyclase and increase the intracellular production of cAMP. This view is supported by several observations: (a) vasopressin in physiologic concentrations (10^{-11} – 10^{-10} M) stimulates adenylyl cyclase and increases cAMP production in the rat medullary thick ascending limb (4, 8, 27); (b) an analogue of cAMP (8-bromo-cAMP) closely mimicked vasopressin action (Fig. 5); (c) the effect of vasopressin to inhibit bicarbonate absorption was blocked by exogenous 8-bromo-cAMP (Fig. 7); (d) forskolin, an agent that stimulates intracellular cAMP production (28, 29), inhibited bicarbonate absorption (Fig. 6); and (e) Glu, another hormone that stimulates adenylyl cyclase activity in medullary thick ascending limbs (4, 7, 8), inhibited bicarbonate absorption by an amount similar to that observed with vasopressin (Fig. 3).

One possible explanation for the inhibition of bicarbonate absorption would be that it occurs as an indirect result of the effect of vasopressin to stimulate net NaCl absorption. Vaso-

pressin (via cAMP) increases net NaCl absorption in mouse (6, 20) and rat (21) medullary thick ascending limbs and this effect is associated with increased apical sodium uptake via the Na⁺-K⁺-2 Cl⁻ cotransport system (6, 30). The vasopressin-dependent increase in Na⁺ uptake could increase cell Na⁺ activity, reduce the Na⁺ gradient driving apical membrane Na⁺-H⁺ exchange, and inhibit net bicarbonate absorption. To test this possibility, the effect of vasopressin was examined in tubules perfused with furosemide to inhibit Na⁺-K⁺-2 Cl⁻ cotransport and transcellular NaCl absorption (5). The results show that vasopressin inhibited bicarbonate absorption to a similar extent either in the presence or absence of luminal furosemide. Thus, the inhibition of bicarbonate absorption occurs independent of effects on net NaCl absorption. The experiments with furosemide also demonstrate that the inhibition of bicarbonate absorption can occur in the absence of a change in transepithelial voltage (Table II).

Another means by which vasopressin and cAMP could inhibit bicarbonate absorption would be by inhibiting apical membrane Na⁺-H⁺ exchange, which mediates most of transcellular bicarbonate absorption in the rat thick ascending limb (2, 3). cAMP and hormones such as PTH that stimulate production of cAMP, have been shown to inhibit Na⁺-H⁺ exchange and net bicarbonate absorption in proximal tubules (9, 22, 23, 31) and in renal brush border membranes (32). cAMP also inhibits Na⁺-H⁺ exchange in the apical membrane of the

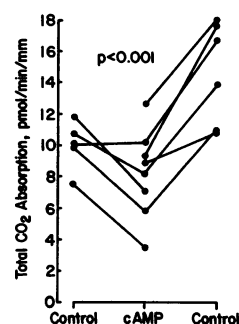


Figure 5. Effect of 8-bromo-cAMP (cAMP; 1×10^{-3} M in the bath) on total CO₂ absorption by the rat medullary thick ascending limb. Solid circles, lines, and P value as in Fig. 1.

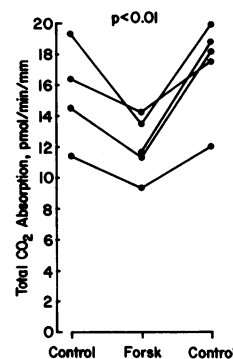


Figure 6. Effect of forskolin (Forsk; 1×10^{-6} M in the bath) on total CO₂ absorption by the rat medullary thick ascending limb. Solid circles, lines, and P value as in Fig. 1.

Table VI. Effect of AVP on Bicarbonate Transport in the Presence of 8-Bromo-cAMP

	V	[Total CO ₂]		JT _{CO₂}	V _{TE}
		Perfused	Collected		
		nl/min per mm	mM		
8-Bromo-cAMP	1.2±0.1	25.4±0.3	19.1±0.7*	7.3±0.4	9.5±1.0
Vasopressin + 8-bromo-cAMP	1.2±0.1	25.4±0.3	19.3±0.5*	6.9±0.5	8.8±1.1
P	NS		NS	NS	NS

Values are means±SE. Vasopressin was 2.8×10^{-10} M in the bath. The bath contained 1×10^{-3} M 8-bromo-cAMP throughout the experiments. No. of experiments = 7. Mean tubule length = 0.58 ± 0.02 mm. Bath total CO₂ concentration was 26.2 ± 0.5 mM in both 8-Bromo-cAMP and vasopressin + 8-bromo-cAMP conditions. V, JT_{CO₂}, V_{TE}, P values, and * as in Table I.

Necturus gallbladder (33). However, whether cAMP inhibits apical Na⁺-H⁺ exchange in the medullary thick ascending limb has not been determined.

Effects of Glu on bicarbonate transport. Glu directly inhibited bicarbonate absorption (Fig. 3). As discussed above, this effect can best be accounted for by the action of Glu to stimulate adenylyl cyclase and increase intracellular production of cAMP (4). The similarity of the effects of Glu and vasopressin on bicarbonate absorption in the medullary thick ascending limb supports the concept of multiple hormonal control in which different hormones that activate adenylyl cyclase in a particular nephron segment are predicted to exert a similar physiologic effect (4, 7).

In recent studies in hormone-deprived rats, the acute infusion of Glu to physiologic levels caused an increase in urinary bicarbonate excretion and a decrease in net acid excretion (13). The increase in bicarbonate excretion was associated with a decrease in fractional bicarbonate absorption by the loop segment (10, 15). Results of the present study indicate that this decrease can be explained, at least in part, by direct inhibition of bicarbonate absorption in the medullary thick ascending limb. At the present time the physiological importance of the effects of Glu on urinary acidification is unknown.

Effects of PTH on bicarbonate transport. Bichara et al. reported recently that infusion of PTH into thyroparathyroidectomized rats increased bicarbonate absorption along the superficial loop segment (14). Results of the present study show, however, that PTH directly inhibited bicarbonate absorption in the medullary thick ascending limb (Fig. 4). Furthermore, preliminary studies in this laboratory have shown that PTH also inhibits bicarbonate absorption in the rat cortical thick ascending limb (Good, D., unpublished observations).⁴ Thus, the increase in loop bicarbonate absorption seen with PTH administration in vivo does not appear to be due to direct hormonal stimulation of bicarbonate absorption in the thick ascending limb. Because the extent of inhibition of bicarbonate absorption by PTH in vitro was relatively small (< 20%), and because the effect was observed using PTH concentrations above the physiological range, it is uncertain whether inhibition of bicarbonate absorption in the medullary thick ascending limb represents an important physiologic effect of the hor-

4. In two experiments with cortical thick ascending limbs, bicarbonate absorption rate with 3×10^{-8} M PTH in the bath was 1–3 pmol/min per mm compared with 7–9 pmol/min per mm measured in controls.

none. If PTH does not significantly inhibit thick ascending limb bicarbonate absorption in vivo, then the stimulation of loop bicarbonate absorption observed after PTH infusion (14) may simply reflect the ability of the loop segment to increase its bicarbonate absorptive capacity in response to the increase in bicarbonate load (increase in luminal bicarbonate concentration) delivered to it from the proximal tubule.

The mechanisms by which PTH affects bicarbonate transport in the medullary thick ascending limb are uncertain. PTH inhibits bicarbonate absorption and Na⁺-H⁺ exchange in cells of the proximal tubule, and these actions are mimicked by analogues of cAMP (9, 22, 23, 31, 32). Several studies, however, have reported no effect of PTH on adenylyl cyclase activity in the medullary thick ascending limb (34–36). It is possible, however, that the sensitivity of the adenylyl cyclase assay may be insufficient, or that performance of the assay in lysed cells under nonphysiological conditions may be inadequate to detect modest (~ 50%) changes in cyclase activity that may occur in intact cells and be physiologically significant. Alternatively, PTH could influence bicarbonate transport in thick ascending limbs by other mechanisms such as activation of Ca²⁺-dependent messenger systems (37, 38), effects on prostaglandins, or inhibition of carbonic anhydrase (39).

Effect of vasopressin on ammonium transport. In addition to reabsorbing bicarbonate, a second means by which the thick ascending limb influences urinary net acid excretion is by reabsorbing ammonium (1, 18). Absorption of NH₄⁺ by the medullary thick ascending limb enhances urinary ammonium excretion by promoting countercurrent multiplication of ammonium in the renal medulla, which facilitates secretion of ammonium into the collecting ducts (16–18). Absorption of ammonium by the medullary thick ascending limb occurs predominantly by secondary active transport, mediated by sub-

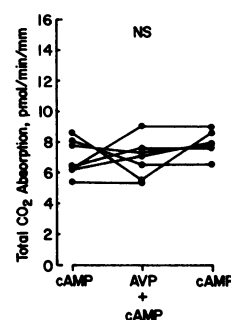


Figure 7. Effect of AVP (2.8×10^{-10} M in the bath) on total CO₂ absorption in the presence of 8-bromo-cAMP. 8-Bromo-cAMP (1×10^{-3} M) was present in the bath throughout the experiments. Solid circles and lines as in Fig. 1. NS, No significant difference (paired *t* test).

Table VII. Effect of AVP on Ammonium Transport

	<i>V</i>	[Am]		<i>J</i> Am	<i>V</i> _{TE}
		Perfused	Collected		
	nl/min per mm	mM	mM	pmol/min per mm	mV
Control	5.8±0.4	4.0±0.03	2.5±0.2*	9.2±1.6	3.2±0.3
Vasopressin	5.9±0.5	4.0±0.03	2.5±0.2*	9.4±1.9	3.6±0.5
<i>P</i>	NS		NS	NS	NS

Values are means±SE. Vasopressin was 2.8×10^{-10} M in the bath. No. of experiments = 6. Mean tubule length = 0.56 ± 0.02 mm. [Am] is total ammonia concentration; *J*Am is absolute rate of total ammonia absorption. Total ammonia concentration in bath was 4.0 ± 0.1 mM in both control and vasopressin conditions. *V* and *V*_{TE} are as in Table I. *P* values compare control and vasopressin (paired *t* test). * Collected total ammonia concentration significantly different from perfused (paired *t* test).

stitution of NH_4^+ for K^+ on the apical membrane $\text{Na}^+\text{-K}^+\text{-2 Cl}^-$ cotransport system (1, 40–42). The previous observation that vasopressin stimulates apical $\text{Na}^+\text{-K}^+\text{-2 Cl}^-$ uptake in mouse and rat medullary thick ascending limbs (6, 21, 30) suggested that NH_4^+ uptake via the cotransporter also might be accelerated. The results show, however, that vasopressin had no significant effect on net ammonium absorption (Table VII, Fig. 8). Therefore, vasopressin either did not stimulate apical NH_4^+ uptake, or an increase in NH_4^+ uptake was offset by another effect tending to reduce net NH_4^+ absorption.

The observation that vasopressin markedly inhibits net bicarbonate absorption with no effect on net ammonium absorption illustrates that ammonium and bicarbonate transport rates can be regulated independently in the thick ascending limb. In the medullary thick ascending limb, net ammonium absorption is due to direct NH_4^+ transport that is predominantly transcellular and active (40, 41). Furthermore, back-diffusion of NH_3 from bath to lumen, which could be influenced by luminal pH, comprises only a small fraction of the net ammonium flux due to the low NH_3 concentrations at physiological pH and to a relatively low transepithelial NH_3 permeability (17, 41).⁵ Thus, it is not surprising that NH_4^+ absorption in the medullary thick ascending limb might be relatively insensitive to changes in luminal pH. The observations that changes in potassium concentration inhibited net NH_4^+ absorption with no effect on bicarbonate absorption (18, 40) and that furosemide inhibited NH_4^+ absorption while stimulating net bicarbonate absorption (1, 3) illustrate further that NH_4^+ and HCO_3^- transport rates can vary independently in thick ascending limbs.

5. Using the collected fluid bicarbonate and ammonium concentrations in Tables I and VII, and assuming a transepithelial NH_3 permeability of 3×10^{-3} cm/s (41), the calculated change in NH_3 backflux with vasopressin would be ~ 0.9 pmol/min per mm. This represents an upper estimate since high flow rates used in the ammonium transport experiments would tend to minimize luminal pH changes. Thus, the change in NH_3 backflux would be $< 10\%$ of the net ammonium absorptive flux.

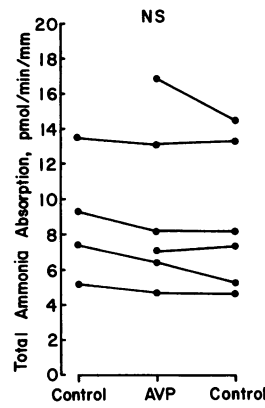


Figure 8. Effect of AVP (2.8×10^{-10} M in the bath) on ammonium absorption by rat medullary thick ascending limbs. Solid circles are mean values for single tubules. Lines as in Fig. 1. NS, No significant difference (paired *t* test).

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