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Research Article

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Anion Transport Processes in the Mammalian Superficial Proximal Straight Tubule

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ABSTRACT The experiments reported in this paper were designed to evaluate some of the characteristics of anion transport processes during fluid absorption from superficial proximal straight tubules isolated from rabbit kidney. We measured net chemical Cl^- flux during fluid absorption from tubules perfused and bathed with Krebs-Ringer buffers containing 113.6 mM Cl^- , 10 mM acetate, and 25 mM HCO_3^- at pH 7.4; assessed the effects of carbonic anhydrase inhibitors on net fluid absorption in the presence and absence of CO_2 ; and evaluated the influx and efflux coefficients for [^{14}C]-acetate transport at 37°C, at 21°C, and in the presence of carbonic anhydrase inhibitors. The experimental data show that, for this nephron segment, net Cl^- flux accompanies approximately 27.5% of net Na^+ absorption; and net Cl^- absorption may be accounted for by a passive transport process, primarily diffusional in nature. Fluid absorption in this nephron segment is reduced 40–60% by carbonic anhydrase inhibitors, but only when the tubules are exposed to 95% O_2 –5% CO_2 rather than 100% O_2 . Thus, it seems probable that approximately half of Na^+ absorption in these tubules may be rationalized in terms of a carbonic anhydrase-dependent CO_2 hydration process. In addition, there may occur in these isolated proximal tubules an acetazolamide-insensitive moiety of HCO_3^- absorption comparable to that observed for proximal tubules in vivo. Finally, we provide evidence that net efflux of luminal acetate is due to metabolic energy-dependent processes

other than CO_2 hydration and may, under appropriate conditions, account for approximately one-fourth of net Na^+ absorption.

INTRODUCTION

The characteristics of salt and volume absorption by the mammalian proximal straight tubule, a nephron segment ordinarily inaccessible to in vivo micropuncture, are not fully understood. When superficial proximal straight tubule segments isolated from rabbit kidney are perfused and bathed at 37°C with symmetrical Krebs-Ringer (KR)¹ solutions containing 113.6 mM Cl^- , 25 mM HCO_3^- , and 10 mM acetate at pH 7.4 (HCO_3^- -KR buffer), isotonic fluid absorption may be accounted for quantitatively by net lumen-to-bath Na^+ flux (1, 2); the transepithelial voltage is -1.0 to -1.5 mV, lumen negative (1, 3); and, in contrast to results obtained in isolated proximal convoluted tubules (4–6), neither net fluid absorption nor the transepithelial voltage depends appreciably on the presence of glucose and/or alanine in perfusing solutions (1).

As in the in vivo mammalian proximal tubule (7, 8), the Cl^- content of tubular fluid in isolated superficial proximal straight tubules perfused with symmetrical HCO_3^- -KR solutions rises appreciably during volume absorption (1). And in the setting of lumen-to-bath Cl^- concentration gradients, the transepithelial voltage in Cl^- -permselective superficial proximal straight tubules becomes lumen positive: when these tubules are perfused with Cl^- -KR/ CO_2 buffers² containing 138.6 mM

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¹Abbreviations used in this paper: J^d , diffusional component of flux; J^e , entrained component of flux; J_v , net fluid transport; K^e , efflux; K^i , influx; KR, Krebs-Ringer solution; P , ionic permeability coefficient; σ , reflection coefficient.

²Two types of KR buffers containing high $\text{Cl}^-/\text{HCO}_3^-$ ratios are referred to in this paper. One of these, described above, contains 3.8 mM HCO_3^- , pH 6.6, and is bubbled with

Cl^- , 3.8 mM HCO_3^- , and 10 mM acetate (pH 6.6), and bathed with HCO_3^- -KR buffers, the transepithelial voltage is in the range +3.0 to +3.8 mV, lumen positive (1, 9). Similar considerations may apply to transepithelial voltages in isolated superficial proximal convoluted tubules (4–6), and the recent micropuncture observations of Barratt et al. (10) have provided evidence for lumen-positive transepithelial voltages in late portions of the *in vivo* superficial proximal tubule.

In contrast, Kawamura et al. (11) have observed that in juxtamedullary proximal straight tubules transepithelial lumen-to-bath NaCl concentration gradients result in lumen negative transepithelial voltages. These workers have also noted that proximal straight tubules originating from juxtamedullary glomeruli are Na^+ , rather than Cl^- permselective. Accordingly, we wish to emphasize that the experimental results and conclusions presented in this paper are restricted to transport events in proximal straight tubules originating from glomeruli in superficial regions of rabbit cortex.

Our earlier observations (9) showed that passive net NaCl flux from lumen to bath, driven by transepithelial Cl^- concentration gradients, attendant lumen-positive transepithelial voltages, and, to a lesser extent, entrainment with volume flow, might account for approximately 40% of the fluid absorbed from superficial proximal straight tubules perfused and bathed with Cl-KR/ CO_2 and HCO_3^- -KR solutions, respectively. Nevertheless, it is clear that fluid absorption in this nephron segment depends, in the final analysis, on transport processes requiring metabolic energy, since (a) when these tubules are perfused and bathed with symmetrical HCO_3^- -KR solutions, both net fluid transport (J_v) and transepithelial voltage (V_e) become zero either upon cooling from 37° to 21°C or at 37°C, by adding 0.1 mM ouabain to the bath (1, 2, 9); and (b) the rise in intraluminal Cl^- concentration during fluid absorption (1) must depend on preferential absorption of anions other than Cl^- with Na^+ . Clearly, additional information is required to describe salt flows in this nephron segment.

In the case of anion transport, certain issues merit particular consideration. In isolated superficial proximal straight tubules (1), we noted that net Na^+ absorption was approximately 15–20% of the unidirectional Cl^- flux, estimated by using $^{36}\text{Cl}^-$, either from lumen to bath or from bath to lumen. Since the magnitude of net Na^+ flux was within the range of experimental error for unidirectional tracer Cl^- fluxes, it was not possible to evaluate quantitatively the Cl^- component of

the anion flux accompanying net Na^+ absorption. Second, HCO_3^- and acetate are, after Cl^- , the predominant anions in the HCO_3^- -KR solutions we used to perfuse isolated superficial proximal straight tubules (1, 9). Accordingly, it is appropriate to inquire about the magnitude of Cl^- flux accompanying Na^+ absorption, the contributions of HCO_3^- and/or acetate as counterions for Na^+ absorption, and hence to the rise in tubular fluid Cl^- concentration during volume absorption, and to consider the possible mechanisms involved in HCO_3^- and acetate absorption.

The experiments reported in this paper were designed to evaluate aspects of these issues in superficial proximal straight tubules isolated from rabbit kidney. We measured net Cl^- flux during fluid absorption by chemical techniques, assessed the effects of carbonic anhydrase inhibitors on net fluid absorption in the presence and absence of CO_2 , and evaluated the influx and efflux coefficients for [^{14}C]acetate transport under a variety of experimental conditions. The experimental data show that, for this nephron segment, net Cl^- flux accompanies approximately 27.5% of net Na^+ absorption; and net Cl^- absorption may be accounted for by a passive transport process, primarily diffusional in nature. Fluid absorption in this nephron segment is reduced 40–60% by carbonic anhydrase inhibitors, but only when the tubules are exposed to 95% O_2 –5% CO_2 rather than 100% O_2 . Thus it seems probable that at least half of Na^+ absorption in these tubules may be rationalized in terms of a carbonic anhydrase-dependent CO_2 hydration process (12–20). In addition, in these isolated proximal tubules an acetazolamide-insensitive moiety of HCO_3^- absorption may occur, comparable to that observed for proximal tubules *in vivo* (12–20). Finally, we provide evidence that net efflux of luminal acetate occurs as a consequence of metabolic energy-dependent processes other than CO_2 hydration and may, under appropriate conditions, account for approximately one fourth of net Na^+ absorption.

METHODS

The techniques used for studying transport events in proximal straight tubules isolated from superficial regions of rabbit renal cortex are nearly the same as those developed originally by Burg et al. (2, 21). Details of the methodology, as utilized in this laboratory, have been presented previously (1, 9); unless otherwise indicated, these techniques were utilized without modification in the present experiments.

Stated briefly, 2.5–4.0 mm segments of proximal straight tubules were obtained by gentle teasing from cortical and outer medullary regions of rabbit renal cortex. The dissections were carried out at 0–5°C in the HCO_3^- -KR bathing media described below. The isolated tubule segments were transferred to a thermoregulated ($\pm 0.5^\circ\text{C}$) chamber, sucked into holding and collecting pipets, and perfused with a

95% O_2 –5% CO_2 ; for convenience, such a solution will be termed Cl-KR/ CO_2 buffer. The second type of Cl-KR solution contains no HCO_3^- , pH 7.4, and is bubbled with 100% O_2 ; this solution will be termed Cl-KR/ O_2 buffer.

microsyringe pump in series with a perfusion pipet advanced approximately 0.2 mm into the tubule lumen.

The HCO₃-KR perfusing solutions used in most experiments contained 105 mM NaCl, 25 mM NaHCO₃, 10 mM Na acetate, 4 mM Na₂HPO₄/NaH₂PO₄, 5 mM KCl, 1.8 mM CaCl₂, 1 mM MgSO₄, 8.3 mM glucose, and 5.0 mM L-alanine; the HCO₃-KR baths contained, in addition to these constituents, 6.0% bovine serum albumin. Both solutions were bubbled continuously with 95% O₂-5% CO₂. The Cl-KR/O₂ perfusing and bathing media were identical to HCO₃-KR solutions except that Cl⁻ replaced HCO₃⁻; the Cl-KR/O₂ solutions were bubbled with 100% O₂ rather than CO₂. Other changes in the composition of perfusing and/or bathing solutions are indicated in the text. All solutions were adjusted to 300 mosmol/liter, pH 7.4.

Net fluid absorption (J_v , nl min⁻¹ mm⁻¹ of tubule length) was measured as described previously (1) from the difference between perfusion and collection rates, with exhaustively dialyzed [*methoxy*-³H]inulin as the volume marker; a positive sign for J_v denotes volume efflux from lumen to bath. Since the mean internal diameter of these tubules is 22.3 μ m (1), J_v values may be converted from nanoliters min⁻¹ mm⁻¹ to cm³ s⁻¹ cm⁻² by the factor 2.38×10^{-5} cm min mm nl⁻¹ s⁻¹ (9). The J_v measurements in the present experiments were in the range 0.1–0.45 nl min⁻¹ mm⁻¹ for tubules 2.5–4.0 mm in length; for isolated proximal convoluted tubules, J_v values in the range of 1.0 nl min⁻¹ mm⁻¹ have been reported for segments 0.5–1.5 mm in length (2, 22, 23). In other words, although J_v is smaller for straight tubules than for proximal convolutions, absolute differences between perfusion and collection rates, and hence the precision of J_v measurements, are comparable in the two segments.

Since certain of the fluid absorption rates reported in this paper (e.g., Tables V, VI, VIII, and XI) were in the range of 0.1–0.2 nl min⁻¹ mm⁻¹, it is relevant to inquire about the extent to which random variations in radioactivity counting might account for such J_v values. In the experiments reported in this paper (66 tubules), the mean tubule length was 3.50 ± 0.07 (mean $\pm$ SEM), the mean perfusion rate was 11.59 ± 0.25 nl/min, and the mean [³H]inulin radioactivity in the perfusing solutions was 28.75 ± 1.05 cpm/nl. The collected fluid was sampled at 7- to 10-min intervals; and, in all cases, perfusing solution standards and collected fluid samples were counted for 10-min periods. With the same perfusing solutions, the coefficient of variation for the liquid scintillation counter used in our laboratory is, at a maximum, $\pm 0.3\%$ for a 10-min counting period.

Accordingly, in an average 3.5-mm tubule perfused for 8 min at 11.59 nl/min with a solution containing 28.75 cpm/nl [³H]inulin, the total radioactivity collected (in the absence of bath leaks) was 26,657 counts/10 min with an average random variation of ± 80 counts/10 min. Thus, the random error in estimating perfusion rate would be 11.59 ± 0.035 nl/min, which, for the tubule length of 3.5 mm, could result in errors in J_v measurements amounting to approximately 0.01 nl min⁻¹ mm⁻¹, or about 10% of the lowest observed J_v value significantly different from zero (Table XI, 0.13 nl min⁻¹ mm⁻¹). Such error in J_v determinations would be random, while the results in Table XI, for example, are specific and statistically reliable in paired observations on the same tubule.

Transepithelial electrical potential differences (V_e , in millivolts, lumen with respect to bath) were measured as described previously (1, 9) with glass 0.9% NaCl/4% agar electrodes placed in both perfusing and bathing solutions.

The electrodes were connected via Tygon tubing (Chamberlain Engineering Corp., Akron, Ohio) filled with 3 M KCl to calomel half-cells. Since the perfusing and bathing solutions were almost identical in ionic composition for all the experiments reported in this paper, no corrections for liquid junction potentials were necessary. The length constant for the proximal straight tubule is approximately 0.1 mm (3). Consequently, V_e measurements in the present paper provided direct information only about electrical events vicinal to the tip of inner perfusion pipet.

Chloride concentrations in perfusing and collected fluids were measured as described previously (9) with a microtitration method (24). In experiments in which the Cl⁻ concentration of collected fluid was determined, samples for the measurement of J_v were alternated with samples for Cl⁻ measurement; the latter were stored in glass micropipets between oil columns until the day after the experiment (a representative experiment according to this protocol, described in the Results section, is shown in Table I). All samples of collected fluid were compared to standard solutions with varying NaCl concentrations but the same concentrations of all other constituents as perfusing solutions. For the sake of uniformity, samples of perfusing solutions were treated in the same manner and also analyzed for Cl⁻ concentrations with the ultramicro method. Duplicate ultramicro and large sample determinations (with an Aminco-Cotlove chloride titrator: American Instrument Co., Silver Springs, Md.) of chloride concentrations agreed within 2%.

[¹⁴C]Acetate fluxes, measuring either the disappearance of tagged acetate from luminal solutions or tagged acetate flux from bath to lumen, were carried out as described previously (1), with 25 or 50 μ Ci/ml of isotope (New England Nuclear, Boston, Mass.) in perfusing or bathing solutions, respectively. In luminal efflux experiments, the bath tracer concentration was uniformly less than 1% of the tracer concentration in the perfusate. In bath-to-lumen tracer fluxes, the tracer concentration in collected tubular fluid was, at a maximum, 4% of the bath tracer concentration, which was constant. The acetate influx (K_{Ac}^i) and efflux (K_{Ac}^e) coefficients, both in centimeters per second, were computed from bath-to-lumen tracer fluxes and luminal tracer effluxes, respectively, as described previously (1). As stressed previously (1), K_{Ac}^e includes, in principle, terms for passive and active ion transport, exchange diffusion, and metabolic consumption of luminal acetate by tubular epithelial cells.

Acetazolamide and ethoxzolamide in chemically pure form were kindly provided by Dr. T. Maren (University of Florida, Gainesville, Fla.). Other reagents, isotope counting techniques, and chemical determinations were as described previously (1, 9). Measurements in a given tubule were used to compute a mean value for that tubule; generally, there were three to four measurements per tubule for a given set of experimental conditions. The mean values for individual tubules (e.g., the representative experiment in Table I) were then used to calculate a mean value $\pm$ SEM for the indicated number of tubules. The experimental results were expressed in this manner. When control and experimental observations were made within the same tubule, P values for mean paired differences were computed from Student's t test by comparing the differences to zero.

RESULTS

The contribution of Cl⁻ to net anion flux. We noted earlier (1) that in superficial proximal straight tubules perfused and bathed at 37°C with HCO₃-KR solutions,

the net lumen to bath Na^+ flux^{*} ($J_{\text{Na}}^{\text{net}}$, eq $\text{min}^{-1} \text{mm}^{-1}$), computed from the difference between unidirectional lumen to bath and bath to lumen $^{\text{ss}}\text{Na}^+$ fluxes, was in close agreement with the value of $J_{\text{Na}}^{\text{net}}$ predicted from simultaneous measurements of net fluid absorption for an isotonic process. Specifically, in tubules having a mean fluid absorption rate of $0.49 \text{ nl min}^{-1} \text{mm}^{-1}$, the predicted net Na^+ flux (computed as the product of J_v and the Na^+ concentration in the perfusate) was $65.5 \text{ peq min}^{-1} \text{mm}^{-1}$ while the observed net Na^+ flux was $63.8 \text{ peq min}^{-1} \text{mm}^{-1}$. Based on these results, values for the expected net lumen to bath Na^+ flux used in this paper were computed as the product of J_v and the Na^+ concentrations of perfusing solutions.

A representative experiment illustrating the method used for measuring net lumen to bath Cl^- flux is presented in Table I. Alternating samples of collected fluid were taken for J_v and Cl^- determinations. These data were used to compute mean values for J_v and collected fluid Cl^- concentration in that tubule; the mean perfusion and collection rates for the tubule were computed from the values of these variables measured during the J_v periods.

Table II presents the results of five such experiments in as many individual tubules. The mean value for J_v , $0.38 \text{ nl min}^{-1} \text{mm}^{-1}$, was approximately the same as reported previously (1, 9), and the Cl^- concentration of the collected fluid was $10.9 \pm 1.4 \text{ meq/liter}$ higher than that in the perfusate ($P < 0.002$). The rate of tubular chloride perfusion was computed from the product of the perfusion rate and the perfusate chloride

^{*} Net ion absorption ($J_{\text{ion}}^{\text{net}}$) in this paper is expressed as $\text{peq min}^{-1} \text{mm}^{-1}$ of tubule length. Since the mean internal diameter of these tubules is $22.3 \mu\text{m}$ (1), $J_{\text{ion}}^{\text{net}}$ may be converted from $\text{peq min}^{-1} \text{mm}^{-1}$ to $\text{neq s}^{-1} \text{cm}^{-2}$ of apparent luminal surface area by the factor 0.02381.

TABLE I
A Representative Experiment Measuring the Cl^- Component of Net Anion Flux

Time	Perfusion rate	Collection rate	J_v	$[\text{Cl}^-]_{\text{collected fluid}}$
min	nl/min	nl/min	nl min ⁻¹ mm ⁻¹	mM
39-49	14.28	13.24	0.30	—
49-57	—	—	—	127.0
57-66	11.91	10.29	0.46	—
66-74	—	—	—	121.0
75-82	13.56	12.33	0.35	—
82-91	—	—	—	124.0
91-100	12.31	10.63	0.48	—
Mean values	13.01	11.62	0.40	124.0

Proximal straight tube segments were perfused and bathed with $\text{HCO}_3\text{-KR}$ buffer solutions at 37°C . Time represents the time elapsed after sacrifice of the animal. As described previously (1), the perfusion and collection rates were measured from the rates of delivery into the collected fluid of [*methoxy-³H*]inulin and volume, respectively. The chloride concentration of the perfusate, measured as described previously (9) with an ultramicro method (24), was 115 meq/liter .

concentration, and the rate of tubular chloride collection was computed as the product of the collection rate and the chloride concentration in the collected fluid. $J_{\text{Cl}}^{\text{net}}$, the net chloride flux from lumen to bath, was expressed as the difference between the rates of Cl^- perfusion and collection per millimeter of tubule length. The last column of Table II shows that $J_{\text{Cl}}^{\text{net}}$ was $27.5 \pm 2.3\%$ of the expected $J_{\text{Na}}^{\text{net}}$.

We measured the Cl^- concentrations of perfusing and collected fluids for tubules perfused and bathed with $\text{HCO}_3\text{-KR}$ solutions at 21°C , since, under these conditions, J_v and V_v are both zero (1, 9; Tables V and VIII). In five tubules perfused and bathed at 21°C , the perfusate Cl^- concentration was $116.4 \pm 1.6 \text{ mM}$, the collected fluid Cl^- concentration was $116.2 \pm 1.8 \text{ mM}$, and the mean paired difference between these two con-

TABLE II
The Cl^- Component of Net Anion Flux

Tubule	Perfusate		Collected fluid		$\Delta[\text{Cl}^-]$	J_v	$J_{\text{Cl}}^{\text{net}}$	$J_{\text{Na}}^{\text{net}}$	$J_{\text{Cl}}^{\text{net}}/J_{\text{Na}}^{\text{net}}$
	$[\text{Cl}^-]$	Rate	$[\text{Cl}^-]$	Rate					
mm	mM	nl/min	mM	nl/min	mM	nl min ⁻¹ mm ⁻¹	peq min ⁻¹ mm ⁻¹		%
3.45	117.0	10.26	131.2	8.65	14.2	0.46	19.0	67.7	28.0
3.25	116.0	8.62	126.3	7.67	10.3	0.29	9.6	42.7	22.4
2.95	115.0	11.92	122.0	10.73	7.0	0.40	21.0	58.9	35.6
3.5	115.0	13.01	124.0	11.62	9.0	0.39	16.1	57.4	28.0
3.6	120.0	8.81	134.5	7.54	14.0	0.35	13.0	51.5	23.3
Mean	116.6		127.6		10.9	0.38	15.7	55.6	27.5
±SEM	±0.9		±2.3		±1.4	±0.03	±2.0	±2.0	±2.3

$P < 0.002$

The data presented are from five experiments at 37°C , each conducted as outlined in Table I. $\Delta[\text{Cl}^-]$ represents the difference in Cl^- concentration between perfused and collected fluids. Net $J_{\text{Cl}}^{\text{net}}$ was computed from differences between the rates of Cl^- perfusion and collection. $J_{\text{Na}}^{\text{net}}$, the expected net Na^+ flux, was calculated as described in Results.

TABLE III
Effect of Acetazolamide on J_v and V_o : HCO_3^- -KR
Perfusate and Bath

Acetazolamide	J_v	V_o
mM	nl min ⁻¹ mm ⁻¹	mV
0	0.26±0.03	-1.3±0.20
0.22	0.11±0.02	-1.6±0.21
Mean paired difference	0.16±0.02 $P < 0.001$	0.32±0.08 $P < 0.01$
	(n = 10)	
0	0.34±0.03	-0.62±0.14
2.25	0.18±0.02	-0.62±0.11
Mean paired difference	0.16±0.03 $P < 0.001$	0.00±0.10 $P > 0.10$
	(n = 7)	

Proximal straight tubule segments were perfused and bathed with HCO_3^- -KR buffer solutions at 37°C. When present, acetazolamide was added to both perfusate and bath at the indicated concentrations. Paired observations on each tubule were carried out in all instances. All values of J_v and V_o in the presence or absence of acetazolamide differed from zero at $P < 0.001$. The mean perfusion rates for the sets of experiments involving 0.22 mM and 2.25 mM acetazolamide were 12.29 ± 0.28 and 11.15 ± 0.62 , respectively.

centrations was 0.2 ± 0.1 mM ($P > 0.1$). Thus, we are confident that the rise in tubular fluid concentration at 37°C (Table II) was not due to an experimental artifact arising during tubular perfusion.

The effect of carbonic anhydrase inhibitors. It is clear from the results shown in Table II that anions other than Cl^- were preferentially absorbed with Na^+ . Accordingly, we assessed the effects of carbonic anhydrase inhibitors on J_v and V_o (Tables III and IV) in paired control and experimental (i.e., in the presence of acetazolamide or ethoxzolamide) measurements in each of the tubules tested. The order in which observations were made was not varied; rather, one or the other inhibitor was always added after control observations. However, the J_v and V_o measurements in the presence of acetazolamide or ethoxzolamide were completed within 100 min after initiating tubule perfusion, and our earlier observations (1) indicate that, within this time span, values for J_v and V_o in a given tubule perfused and bathed with HCO_3^- -KR solutions are stable.

Table III illustrates the effects of adding the relatively hydrophilic agent (15, 16) acetazolamide to both perfusing and bathing solutions. At 0.22 mM acetazolamide, J_v fell approximately 61%, while V_o rose slightly. The former seems to be the maximal degree to which the carbonic anhydrase inhibitor reduced fluid absorption from tubules exposed to symmetrical HCO_3^- -KR solutions, since at approximately 10-fold higher con-

centrations of acetazolamide, 2.25 mM, J_v was inhibited 47% and V_o was unchanged. In like manner, Table VIII (cf. below) shows that addition of 0.16 mM ethoxzolamide, a relatively lipophilic agent (15, 16), to bathing media inhibited J_v by approximately 40%. These observations are comparable to the effects of acetazolamide on fluid absorption from proximal tubules during in vivo micropuncture (14, 17-19).

It seems reasonable to surmise from these data that both acetazolamide (Table III) and ethoxzolamide (Table VIII) inhibited carbonic anhydrase-dependent HCO_3^- absorption (12-20). But it is possible that the effects of these agents on J_v were nonspecific. For example, Grantham et al. observed (25) that the diuretics acetazolamide, chlorothiazide, and furosemide each reduced fluid absorption in isolated rabbit proximal tubules by 30-80%. The recent observations of Burg et al. indicate clearly that diuretics such as mersalyl (26), furosemide (27), and ethacrynic acid (28) inhibit primarily active Cl^- absorption in isolated segments of the thick ascending limb of Henle's loop. Since the rabbit proximal straight tubule secretes organic acids (29, 30), it might therefore be argued (25) that the effects of acetazolamide and ethoxzolamide on net fluid absorption in these tubules (Tables III, VIII) were the consequence of fluid secretion coupled to net bath to lumen flux of these agents. This possibility was not evaluated explicitly, since we did not measure fluxes of these diuretics. However, Table IV shows that when the tubules were

TABLE IV
Effect of Carbonic Anhydrase Inhibitors on J_v and V_o :
 Cl-KR/O_2 Perfusate and Bath

Inhibitor	J_v	V_o
	nl min ⁻¹ mm ⁻¹	mV
None	0.37±0.13	-0.88±0.05
0.22 mM acetazolamide	0.36±0.05	-0.70±0.11
Mean paired difference	0.01±0.02 $P > 0.5$	0.18±0.15 $P > 0.5$
	(n = 5)	
None	0.37±0.02	-0.93±0.18
0.16 mM ethoxzolamide	0.35±0.02	-1.00±0.1
Mean paired difference	0.03±0.01 $P > 0.05$	0.07±0.08 $P > 0.05$
	(n = 4)	

Proximal straight tubules were perfused and bathed with Cl-KR/O_2 buffer solutions at pH 7.4 bubbled with 100% O_2 . When present, acetazolamide was added to both perfusing and bathing fluids, while ethoxzolamide was added only to bathing solutions. In all instances, paired observations on each tubule were carried out. In the experiments with acetazolamide and ethoxzolamide, the mean perfusion rates were 11.85 ± 0.91 and 11.45 ± 0.45 , respectively.

perfused and bathed with Cl-KR/O₂ solutions exposed to 100% O₂ rather than 95% O₂-5% CO₂ (conditions which, for these tubules, give comparable values for J_v ; ref. 1), neither 0.22 mM acetazolamide nor 0.16 mM ethoxzolamide had a detectable effect on J_v . Consequently it is reasonable to infer that the inhibition of fluid absorption produced by these agents (Tables III, VIII) was due to decrements in the rate of CO₂ hydration. According to this view, the results in Tables III, and VIII indicate that 40–60% of the expected net Na⁺ flux in these tubules was coupled to CO₂ and carbonic anhydrase-sensitive HCO₃⁻ absorption.

The acetazolamide-insensitive moiety of J_v . It is well known that 40–50% of proximal tubular HCO₃⁻ absorption continues after virtually complete (i.e., greater than 99.99%) inhibition of renal carbonic anhydrase activity by acetazolamide (12–20). Thus we examined in part the acetazolamide-insensitive fraction of fluid absorption in these tubules. Table V indicates, in accord with earlier observations (1), that cooling from 37° to 21°C in random paired experiments reduced the acetazolamide-insensitive moiety of net fluid absorption in tubules exposed to symmetrical HCO₃-KR solutions from 0.21 to 0.02 nl min⁻¹ mm⁻¹; the former differed significantly from zero ($P < 0.001$), while the latter was indistinguishable from zero ($P > 0.5$).

We also assessed the magnitude of net Cl⁻ flux with respect to the expected net Na⁺ flux in the presence of 2.25 mM acetazolamide (Table VI). To exclude possible contributions of glucose (31), alanine, or acetate absorption (Tables VII–IX) to acetazolamide-insensitive fluid absorption, the HCO₃-KR perfusates, but

TABLE V
Effect of Cooling on the Acetazolamide-Insensitive Moiety of J_v

Temperature	Acetazolamide	J_v
°C	mM	nl min ⁻¹ mm ⁻¹
37	2.25	0.20 ± 0.03
21	2.25	0.02 ± 0.01
Mean paired difference		0.18 ± 0.03 $P < 0.02$ ($n = 4$)

Proximal straight tubules were perfused and bathed with HCO₃-KR solutions in the presence of 2.25 mM acetazolamide in both solutions. J_v and V_e were measured at 37° and 21°C. In all instances, paired observations on each tubule were carried out in random order. The values of J_v at 37°C differed from zero at $P < 0.001$, while the values of J_v at 21°C were indistinguishable from zero ($P > 0.5$). The mean perfusing rate was 9.64 ± 0.63.

not the baths, used in these experiments were modified: glucose and alanine were replaced isosmotically with urea; and acetate was replaced isosmotically with isethionate, a highly impermeant anion ($P_{\text{isethionate}}$ in these tubules is 0.023 μm/s; [9]). As is shown in Table XI, luminal isethionate, sulfate, and phosphate, in the absence of luminal Cl⁻, acetate, glucose, alanine, and HCO₃⁻ concentrations greater than 3.8 mM, are unable to support rates of fluid absorption distinguishable from zero. And our prior observations have shown that in these tubules, oppositely directed transepithelial concentration gradients for urea and alanine plus glucose (specifically, urea in luminal solutions and glucose plus

TABLE VI
The Cl⁻ Component of Net Anion Flux for the Acetazolamide-Insensitive Moiety of J_v

Tubule	Perfusate		Collected fluid		$\Delta[\text{Cl}^-]$	J_v	$J_{\text{Cl}^{\text{net}}}$	$J_{\text{Na}^{\text{net}}}$	$J_{\text{Cl}^{\text{net}}}/J_{\text{Na}^{\text{net}}}$
	[Cl ⁻]	Rate	[Cl ⁻]	Rate					
mm	mM	nl/min	mM	nl/min	mM	nl min ⁻¹ mm ⁻¹	peq min ⁻¹ mm ⁻¹		%
2.34	117.5	8.08	120	7.54	2.5	0.26	19.1	38.2	50.0
3.7	118.0	8.12	117.7	7.60	-0.3	0.14	18.2	20.6	88.3
3.4	118.8	9.45	122.6	8.80	3.8	0.18	12.9	26.4	48.9
3.7	116.0	12.50	120.5	11.20	4.5	0.35	27.8	51.8	53.6
3.9	116.0	15.14	116.7	14.72	0.7	0.11	10.1	16.2	62.3
3.8	120.5	10.35	120.6	9.96	0.1	0.10	12.1	14.7	82.3
Mean	117.8	10.60	119.7	9.97	1.88	0.19	16.7	28.0	64.2
±SEM	±0.7		±0.9		±0.82	±0.04	±2.6	±5.9	±7.0
$P > 0.05$									

Cl⁻ concentrations of perfused and collected fluids were measured for tubules perfused and bathed at 37°C with solutions containing 2.25 mM acetazolamide. $\Delta[\text{Cl}^-]$, $J_{\text{Cl}^{\text{net}}}$, and the expected $J_{\text{Na}^{\text{net}}}$ were computed as described for Table II. The HCO₃-KR perfusates used in these experiments were modified in the following manner: glucose and alanine were replaced isosmotically with urea, and acetate was replaced by isethionate. The bath was unmodified HCO₃-KR. The values for J_v were uniformly different from zero ($P > 0.001$).

alanine in bathing solutions) do not result in J_v values different from zero (9). Thus, in Table VI, HCO_3^- and Cl^- were the only potentially suitable counterions in the perfusing solutions for expected net Na^+ flux resulting in rates of fluid absorption significantly different from zero.

The experimental results, shown in Table VI, illustrate several noteworthy characteristics. First, in the presence of acetazolamide (and in the absence of luminal acetate, glucose, and alanine), increases in tubular fluid Cl^- concentrations (e.g., see Table II) were blunted. In Table VI, the Cl^- concentration of collected fluid was 119.7 ± 0.9 mM, i.e., negligibly greater (1.88 ± 0.82 mM; $P > 0.05$) than that in the perfusate. Second, the observed net Cl^- flux accounted for approximately two-thirds of the expected net Na^+ flux. By exclusion, we argue that approximately one third of the expected net Na^+ flux was accompanied by HCO_3^- , although the acetazolamide concentration used in these experiments, 2.25 mM, was ten times that required for maximal inhibition of fluid absorption from symmetrical HCO_3^- -KR solutions (Table III). Finally, it is interesting to note that the mean value $\pm$ SEM for the acetazolamide-insensitive component of fluid absorption listed in Table VI did not differ significantly ($P > 0.5$) from the comparable value of J_v at 37°C shown in Table V. One cannot infer explicitly from this that the temperature-dependent, acetazolamide-insensitive moiety of fluid absorption in Table V occurred independently of luminal glucose, alanine, or acetate, since the experiments in Tables V and VI were carried out on different tubules. However, Table VI indicates per se that acetazolamide-insensitive ion and fluid transport occurred in these tubules in the presence of luminal Cl^- and HCO_3^- and in the absence of either luminal glucose,

TABLE VII
Acetate Influx and Efflux Coefficients

K_{Ae}^o	K_{Ae}^i	J_v
$\mu\text{m/s}$		$\text{nl min}^{-1} \text{mm}^{-1}$
0.46 ± 0.03	0.14 ± 0.02	0.39 ± 0.02
Mean paired difference:		
0.32 ± 0.05		
$P < 0.01$		
$(n = 4)$		

Efflux (K_{Ae}^o) and influx (K_{Ae}^i) coefficients for acetate were measured as described in Methods from luminal disappearance rates and bath to lumen fluxes, respectively, of [^{14}C]acetate at 37°C . The perfusing and bathing solutions were standard HCO_3^- -KR buffers containing 10 mM acetate. K_{Ae}^o and K_{Ae}^i were measured in each tubule; the order in which K_{Ae}^o and K_{Ae}^i were measured was varied at random among the individual tubules. The mean perfusing rate was 8.89 ± 2.24 .

TABLE VIII
Effect of Cooling or Carbonic Anhydrase Inhibition on the Acetate Efflux Coefficient

Temperature	Ethoxzolamide	K_{Ae}^o	J_v
$^\circ\text{C}$	mM	$\mu\text{m/s}$	$\text{nl min}^{-1} \text{mm}^{-1}$
37	0	0.40 ± 0.02	0.32 ± 0.03
21	0	0.18 ± 0.02	0.01 ± 0.01
Mean paired difference		0.22 ± 0.03	0.31 ± 0.03
		$P < 0.001$	$P < 0.001$
$(n = 6)$			
37	0	0.44 ± 0.02	0.41 ± 0.03
37	0.16	0.50 ± 0.04	0.25 ± 0.04
Mean paired difference		0.06 ± 0.03	0.16 ± 0.02
		$P > 0.5$	$P < 0.002$
$(n = 5)$			

K_{Ae}^o and J_v were measured as described in Table VIII for tubules perfused and bathed with HCO_3^- -KR solutions. In the upper part of the table, the effect of reducing temperature from 37° to 21°C was examined; the order in which experiments was carried out was varied at random. In the lower part of the table, the effect of adding 0.16 mM ethoxzolamide to the bathing solutions was examined; in all cases, control observations preceded those in the presence of ethoxzolamide. Paired observations on the same tubule were carried out in all instances. In the experiments involving cooling, the mean perfusion rate was 11.56 ± 0.82 ; in the experiments with ethoxzolamide, the mean perfusion rate was 11.78 ± 0.73 .

alanine, and acetate, or oppositely directed transepithelial gradients for relatively permeant and impermeant anions, such as Cl^- and HCO_3^- , respectively (9).

[^{14}C]Acetate flux coefficients. Table VII shows the results of experiments designed to measure the flux coefficients for [^{14}C]acetate. The influx coefficient was computed, from expressions presented previously (1), from the appearance rate of [^{14}C]acetate in luminal solutions. In other words, K_{Ae}^i represents the bath-to-lumen acetate flux coefficient. On the other hand, K_{Ae}^o was calculated, as described previously (1), from the disappearance rates of [^{14}C]acetate from luminal solutions. Thus, K_{Ae}^o is a luminal efflux coefficient rather than a unidirectional lumen-to-bath flux coefficient, and may include not only terms for passive and/or active [^{14}C]acetate flux from lumen to bath, and exchange diffusion of acetate, but also cellular consumption of luminal acetate (1), a phenomenon recently proposed on the basis of proximal tubular micropuncture studies in the rat to account for net luminal efflux of the organic anion lactate (32).

Table VII shows that, in random paired observations in four tubules, K_{Ae}^o was approximately three times greater than K_{Ae}^i , the mean paired difference between

these values being 0.32 ± 0.05 ($P < 0.01$). Table VIII shows that, in paired observations, cooling from 37° to 21°C reduced K_{Ac}^* more than twofold, while 0.16 mM ethoxzolamide in the bath reduced fluid absorption by approximately 40% (cf. also Table III) but had no effect on K_{Ac}^* . We also assessed directly the effects of CO₂ omission on K_{Ac}^* . In five tubules, random paired measurements of K_{Ac}^* were made under the following conditions: either when the perfusate and bath each contained HCO₃-KR bubbled at pH 7.4 with 95% O₂ 5% CO₂ at 37°C; or when the perfusate and bath each contained Cl-KR/O₂ bubbled at pH 7.4 with O₂ at 37°C. The mean paired difference for K_{Ac}^* under these two conditions was -0.11 ± 0.05 ($P > 0.2$). These data, taken together with the results in Tables VII and VIII, indicate clearly that K_{Ac}^* exceeded K_{Ac}^i , and that the difference between K_{Ac}^* and K_{Ac}^i was temperature-dependent but unaffected either by concentrations of ethoxzolamide sufficient to inhibit fluid absorption or by exclusion of CO₂ and HCO₃⁻ from the system.

The contribution of luminal acetate efflux to fluid absorption. Since the pK of acetic acid is 4.7, the ratio of acetate anion to acetic acid, in the pH range 6.6–7.4, varies between 80 and 500. Thus, regardless of the molecular events responsible for the fact that K_{Ac}^* exceeded K_{Ac}^i (Tables VII and VIII), it is clear that net disappearance of acetate from luminal fluids was accompanied by net cation flux from lumen to bath and by net fluid absorption. We were not able to measure chemically the content of acetate in collected fluids. Accordingly, net luminal acetate effluxes could not be computed from the experiments shown in Tables VII and VIII for the following reasons. First, the results in Table II indicate that tubular fluid Cl⁻ concentrations rose during net fluid absorption; consequently, the acetate concentration in tubular fluid may not have been constant. Second, during the luminal [¹⁴C]acetate efflux experiments shown in Tables VII and VIII, 10 mM unlabeled acetate was present in the bathing solutions. Since bath to lumen flux of unlabeled acetate could have reduced the [¹⁴C]acetate specific activity (counts per minute [¹⁴C]acetate per mole unlabeled acetate) in luminal solutions, the [¹⁴C]acetate concentration of collected fluid could not be used to compute the unlabeled acetate concentration of collected fluid.

To circumvent these difficulties, we measured luminal effluxes of [¹⁴C]acetate in experiments in which isethionate replaced acetate in HCO₃-KR bathing solutions. Table IX presents the results of a series of experiments in which random paired measurements of K_{Ac}^* and J_v were carried out either in the presence or absence of acetate in the bathing solutions. With the latter exception, the experimental conditions were identical to those used for assessing simultaneously net fluid absorption and net Na⁺ flux (1), net fluid absorption and net Cl⁻

TABLE IX
Effect of Omitting Acetate from the Bath on K_{Ac}^* and J_v

Bath acetate	K_{Ac}^*	J_v
mM	$\mu\text{m/s}$	$\text{nl min}^{-1} \text{mm}^{-1}$
10	0.34 ± 0.01	0.32 ± 0.04
0	0.37 ± 0.02	0.31 ± 0.04
Mean paired difference	0.03 ± 0.01 $P > 0.5$	-0.01 ± 0.01 $P > 0.5$

($n = 5$)

K_{Ac}^* and J_v were measured as described in Table VIII from the disappearance rates of [¹⁴C]acetate from luminal fluids in tubules perfused with standard HCO₃-KR buffers containing 10 mM acetate. The bathing solutions were HCO₃-KR buffers containing either 10 mM or zero acetate. In the latter case, acetate was replaced by 10 mM isethionate. Paired observations at zero and 10 mM acetate in the bath were carried out in each of the five tubules; the order in which acetate was omitted from the bath was varied at random among the five tubules. The mean perfusing rate was 11.72 ± 0.98 .

flux (Table II), net fluid absorption and the acetate influx and efflux coefficients (Table VII), and the effects of carbonic anhydrase inhibitors on net fluid absorption and transepithelial voltages (Table IV). The results in Table IX show clearly that, within the limits of experimental error, neither K_{Ac}^* nor J_v was affected by replacing acetate in the bath with isethionate. Specifically, the mean paired differences for K_{Ac}^* and J_v in the presence and absence of 10 mM bath acetate were $+0.03 \pm 0.02$ ($P > 0.5$) and -0.01 ± 0.01 ($P > 0.5$), respectively.

Three issues relating to the results in Table IX are particularly interesting. First, since K_{Ac}^* was unaffected by removal of bath acetate, it is improbable that acetate exchange diffusion occurred to a detectable degree. Second, K_{Ac}^i isethionate, the bath-to-lumen flux constant for isethionate in these tubules, is $0.023 \mu\text{m/s}$ (9), i.e., 7 and 23 times smaller than K_{Ac}^i or K_{Ac}^* , respectively (Table VII); thus, it seems unlikely that any appreciable electrically silent exchange of luminal acetate for bath isethionate occurred in the experiments shown in Table IX. Finally, there is compelling experimental evidence that bulk-phase unstirred layer effects do not complicate flux measurements with isolated perfused renal tubules (9, 33, 34). The latter implies that, in the experiments recorded in Table IX, acetate accumulation in the bath could not have significantly affected the unidirectional efflux of [¹⁴C]acetate from luminal solutions.

Given these assumptions, it seems reasonable to infer that the K_{Ac}^* measurements in the absence of bathing solution acetate (Table IX) may be used to calculate the net efflux—or disappearance—of acetate from luminal solutions, at least for these particular conditions.

The specific activity X_{Ac}^p (in counts per minute per mole) of acetate in the perfusate is:

$$X_{Ac}^p = \frac{[Ac]_p^*}{[Ac]_p} \quad (1)$$

where $[Ac]_p^*$ and $[Ac]_p$ are, respectively, the concentrations of labeled ($[^{14}C]$ acetate counts per minute per milliliter) and unlabeled (moles acetate per milliliter) species in the perfusate. In the absence of back flux of unlabeled acetate from bath to lumen, X_{Ac}^p equals X_{Ac}^{ef} (counts per minute per mole), the specific activity of acetate in collected fluid, and $[Ac]_{ef}$ (moles per milliliter), the unlabeled acetate concentration in collected fluid, is given by

$$[Ac]_{ef} = \frac{[Ac]_{ef}^*}{[Ac]_p^*} [Ac]_p \quad (2)$$

where $[Ac]_{ef}^*$ and $[Ac]_{ef}$ are the concentrations of the labeled (counts per minute $[^{14}C]$ acetate per milliliter) and unlabeled (moles acetate per milliliter) species in the collected fluid, respectively.

Table X shows that $[Ac]_{ef}$, computed according to Eq. 2 from the $[^{14}C]$ acetate efflux data in Table IX with no acetate in the bathing media, was 5.91 ± 0.77 mM. $[\bar{Ac}]$, the arithmetic mean of $[Ac]_{ef}$ and $[Ac]_p$, was 8.15 mM, and J_{Ac}^{net} , computed as (1, 9)

$$J_{Ac}^{net} = K_{Ac}^e [\bar{Ac}] \quad (3)$$

(with the mean value for K_{Ac}^e in Table IX in the absence of acetate in bathing media) was $0.301 \text{ neq s}^{-1} \text{ cm}^{-2}$ or $12.6 \text{ peq min}^{-1} \text{ mm}^{-1}$. In other words, for these conditions, net luminal efflux of acetate accounted for approximately 27.5% of the expected net Na^+ absorption.

Finally, we evaluated directly whether luminal acetate alone could support net ion and fluid absorption

TABLE X
An Acetate Contribution to Net Cation Flux

$[Ac]_{ef}$	$[\bar{Ac}]$	J_{Ac}^{net}	J_{Na}^{net}	$J_{Ac}^{net}/J_{Na}^{net}$
mM			peq/min/mm	
5.91 ± 0.77	8.15	12.6	45.8	2.75
	(n = 5)			

$[Ac]_{ef}$, the mean $\pm$ SEM for the acetate concentration in collected fluid, was computed according to eq. 2 with the luminal $[^{14}C]$ acetate efflux data listed in Table IX for the case of zero acetate in the bathing media. $[\bar{Ac}]$, the mean luminal acetate for these conditions, was taken to be the arithmetic mean of 10 mM Ac, the acetate concentration in perfusing solutions, and $[Ac]_{ef}$. J_{Ac}^{net} , the net rate of luminal acetate efflux for these conditions, was computed as described previously (1) from the product $K_{Ac}^e [\bar{Ac}]$, with the K_{Ac}^e value listed in Table IX for the case of zero bath acetate.

TABLE XI
Support of Fluid Absorption by Luminal Acetate

$[Ac]$ perfusate	Tempera- ture	J_s	ΔJ_s
mM	°C	nl min ⁻¹ mm ⁻¹	
0	37	-0.04 ± 0.02 $P > 0.5$	0.17 ± 0.05 $P < 0.002$
10	37	$+0.13 \pm 0.04$ $P < 0.002$	0.16 ± 0.04 $P < 0.002$
10	21	-0.04 ± 0.03 $P > 0.5$	
(n = 5)			

NaCl and $NaHCO_3$ in the HCO_3^- -KR perfusing solutions were uniformly replaced by Na isethionate, and alanine and glucose were isosmotically replaced with urea. Na isethionate also replaced Na acetate when the perfusing acetate was zero. Na isethionate replaced NaCl in the HCO_3^- -KR bathing solutions. In all solutions, K^+ and Ca^{++} were present as sulfate rather than Cl^- salts, and all solutions were bubbled with 95% O_2 -5% CO_2 . The final pH of the bathing solutions was 7.4. The final pH of the perfusing solutions was 6.6, which corresponds to an HCO_3^- concentration of 3.8 mM. In each of the five tubules studied, J_s was measured under each of the indicated experimental conditions. The order in which the various measurements were carried out was made random among individual tubules. The P values under J_s values indicate the levels at which these values differed from zero. Mean paired differences and P values for these differences are listed in the ΔJ_s column. The mean perfusing rate was 11.76 ± 1.04 .

(Table XI). In these experiments, the HCO_3^- -KR perfusing solutions were modified by isosmotic replacement of HCO_3^- and Cl^- with isethionate, isosmotic replacement of glucose and alanine with urea, and, when acetate was excluded, isosmotic replacement of the latter with isethionate. In the HCO_3^- -KR bathing solutions, isethionate replaced acetate. And, in both perfusing and bathing solutions, K^+ and Ca^{++} were present as sulfate rather than chloride salts. The final pHs of the bathing and perfusing solutions, all bubbled with 95% O_2 -5% CO_2 were 7.4 and 6.6, respectively.

Table XI shows the results of random paired J_s measurements under three sets of conditions: at 37°C, when the perfusate contained no acetate; at 37°C, when the perfusate contained 10 mM acetate; and at 21°C, when the perfusate contained 10 mM acetate. The rates of fluid absorption at 37°C in the absence of luminal acetate were indistinguishable from zero. Stated in another way, luminal isethionate, sulfate, phosphate, and 3.8 mM HCO_3^- (pH 6.6) were unable to support detectable rates of fluid absorption. However, when 10

mM acetate was added to perfusing solutions at 37°C, J_v rose to 0.13 ± 0.04 , which differed from zero at $P < 0.002$, and when the system containing 10 mM acetate in the perfusing solution was cooled to 21°C, J_v was again indistinguishable from zero, in accord with the fact that K_{Ac}^* was reduced more than twofold on cooling from 37° to 21°C (Table VIII).

DISCUSSION

The experiments reported in this paper were intended to assess the nature and contributions of anion fluxes—specifically, net lumen to bath Cl^- flux, net HCO_3^- absorption, and net disappearance of acetate from luminal solutions—to expected net Na^+ flux and observed rates of fluid absorption in isolated rabbit superficial proximal straight tubules exposed to symmetrical HCO_3^- -KR solutions containing 95% O_2 -5% CO_2 at pH 7.4. We emphasize again that the experimental data, and the analysis which follows, are applicable only to these Cl^- permselective superficial proximal straight tubules, and not necessarily to juxtamedullary proximal straight tubules, since the latter are Na^+ rather than Cl^- permselective (11).

To analyze the present results, we made three general assumptions. First, as in previous studies (1, 9), we consider that temperature-sensitive events, e.g., fluid absorption from tubules perfused with symmetrical HCO_3^- -KR solutions (1, 9; Tables V and VIII) or the greater values of K_{Ac}^* with respect to K_{Ac}^1 (Tables VII and VIII), depend on conservative processes, such as net active ion transport from lumen to bath and/or cellular consumption of a particular ionic species. The assumption appears warranted since, for these conditions, both J_v and V_o become zero either by cooling from 37° to 21°C (1, 9; Tables V and VIII) or by exposure to ouabain at 37°C (1, 9); at 21°C, K_{Ac}^* falls to values nearly the same as those for K_{Ac}^1 at 37°C (Tables VII and VIII); and at 21°C, there is no rise in tubular fluid Cl^- concentration. Second, we consider that the primary effect of acetazolamide or ethoxzolamide in these tubules is to inhibit carbonic anhydrase-catalyzed CO_2 hydration (15, 16), since these agents reduce fluid absorption in tubules exposed to symmetrical HCO_3^- -KR solutions containing 95% O_2 -5% CO_2 but have no effect on fluid transport in tubules exposed to symmetrical Cl^- -KR/ O_2 solutions containing neither HCO_3^- nor CO_2 (Tables III and IV). Third, based on the facts that (a) NaCl dilution potentials in these tubules appear to be expressed across symmetrical extracellular interfaces containing neutral polar sites (1), and (b) these tubules have remarkably low transepithelial electrical resistances (i.e., approximately $5 \text{ ohm} \cdot \text{cm}^2$ [3]), we assume, as discussed in detail previously (1, 9), that passive ion permeation in these tubules, as in other electrically leaky epithelia (35, 36), involves a paracellular route.

In this frame of reference, five general conclusions emerge from the present studies. First, net Cl^- absorption accounted for approximately 27.5% of the expected net Na^+ flux during fluid absorption from tubules exposed to symmetrical HCO_3^- -KR solutions containing 95% O_2 -5% CO_2 (Table II); the rise in tubular fluid Cl^- concentrations during fluid absorption depended on net conservative luminal efflux of other anions, since tubular fluid Cl^- concentrations remained constant when fluid absorption was reduced to zero by cooling to 21°C. Second, CO_2 and carbonic anhydrase-sensitive net HCO_3^- absorption accounted for 40–60% of the expected net Na^+ flux in tubules exposed to symmetrical HCO_3^- -KR solutions containing 95% O_2 -5% CO_2 (Tables III, IV, VIII), and was responsible, at least in part, for the rise in tubular fluid Cl^- concentrations. Third, it seems probable that an acetazolamide-insensitive component of HCO_3^- absorption occurred, since Cl^- accounted for slightly less than two-thirds of the expected net Na^+ flux under conditions where HCO_3^- was the only other probable counterion for both expected net Na^+ flux and observed net fluid absorption rates significantly different from zero (Tables VI, XI). Fourth, the difference between K_{Ac}^* and K_{Ac}^1 depended on conservative processes unaffected by carbonic anhydrase inhibitors (Tables VII and VIII) or CO_2 exclusion from perfusing media (cf. Results). Under appropriate conditions, net disappearance of luminal acetate accounted for approximately one fourth of the expected net Na^+ flux during net fluid absorption (Table X). Finally, fluid absorption from tubules exposed to symmetrical HCO_3^- -KR solutions containing 95% O_2 -5% CO_2 was unaffected, within experimental error, by omitting acetate from bathing media (Table IX). We now consider these processes in further detail.

The mode of Cl^- transport. If passive Cl^- transport involves an extracellular route (1, 9), the contribution of passive Cl^- transport to net Cl^- absorption requires analysis of two events: Cl^- flux across junctional complexes; and Cl^- flux from intercellular spaces to bathing media. The observations of Welling and Grantham (37) indicate that the peritubular basement membranes of these tubules are rather permeable to albumin; consequently, we consider, as in an earlier analysis (9), that the Cl^- reflection coefficient for these peritubular basement membranes is zero. Given the hydraulic conductivity of these peritubular basement membranes, which exceeds $0.4 \text{ mm s}^{-1} \text{ atm}^{-1}$ (37), it may be shown (9) that, for J_v values in the range of $0.4 \text{ nl min}^{-1} \text{ mm}^{-1}$ (i.e., those reported in Table II), one requires negligible hydrostatic gradients, i.e., 0.1–0.2 cm H_2O , between intercellular spaces and bath. Finally, it seems probable that, in these tubules, there is little voltage attenuation (1) or Cl^- accumulation (9) in intercellular spaces.

Thus, to a crude first approximation, one may express the diffusional (J_{Cl}^d) and entrained (J_{Cl}^e) components of Cl^- flux in terms of bulk-phase driving forces and P_{Cl} , the observed transepithelial Cl^- permeability coefficient.

Given these considerations, the diffusional component of Cl^- flux may be approximated (9, 38–40) by

$$J_{Cl}^d \simeq -P_{Cl}([Cl]^\circ - [\bar{Cl}]^1) + \frac{P_{Cl}F}{RT} \bar{V}_e \frac{([\bar{Cl}]^1 + [Cl]^\circ)}{2} \quad (4)$$

where J_{Cl}^d (eq $cm^{-2} s^{-1}$) is the net transepithelial diffusional flux, P_{Cl} is the transepithelial Cl^- permeation coefficient, $[\bar{Cl}]^1$ is the arithmetic mean of the Cl^- concentrations in perfusing and collected solutions, $[Cl]^\circ$ is the bathing solution Cl^- concentration, and $\bar{V}_e$ is the arithmetic mean of the spontaneous transepithelial voltage at the perfusing and collecting ends of the tubules during fluid absorption. Previous studies have shown that for experimental conditions identical to those listed in Table II, P_{Cl} and $\bar{V}_e$ are $0.73 \mu m s^{-1}$ and -0.16 mV, respectively. From Table II, $[\bar{Cl}]^1$ is 122.1 mM. From these values together with the experimental conditions for Table II, J_{Cl}^d computed from eq. 4 is 0.453 neq $s^{-1} cm^{-2}$ or 19.0 peq $min^{-1} mm^{-1}$, in comparison to 15.7 peq $min^{-1} mm^{-1}$, the observed J_{Cl}^{net} in Table II.

The estimates of J_{Cl}^d according to eq. 4, in comparison to the observed values of J_{Cl}^{net} in Table II, imply that entrainment of solute and solvent flows contributed negligibly to net Cl^- transport. σ_{Cl} , the Cl^- reflection coefficient in these tubules, is in the range 0.78–0.95, and, for fluid absorption linked solely to passive ion flows, entrainment of Cl^- flux to solvent flow makes a relatively small contribution, with respect to Cl^- diffusion, to net Cl^- transport (9). In the present instance, the entrained component of net Cl^- flux may be approximated (1, 9, 38, 40) as

$$J_{Cl}^e \simeq \beta J_v (1 - \sigma_{Cl}) \frac{([\bar{Cl}]^1 + [Cl]^\circ)}{2}, \quad (5)$$

where β is the fraction of net fluid absorption traversing junctional complexes (9). The latter is not known for the present experimental conditions, and, as indicated above, only the range of values for σ_{Cl} have been evaluated (9). However, choosing the maximum value of unity for β and 0.87 for σ_{Cl} (i.e., the mean of 0.78 and 0.95, the range of σ_{Cl} values in these tubules; ref. 9), J_{Cl}^e computed according to eq. 5 is 0.138 neq $s^{-1} cm^{-2}$ or 5.8 peq $min^{-1} mm^{-1}$, or 30% of the J_{Cl}^d values computed from eq. 4. In short, these approximations are consistent with the view that net Cl^- absorption in tubules perfused

and bathed with symmetrical HCO_3^- -KR solution (i.e., Table III) was passive, mainly diffusional in nature.

HCO_3^- absorption. Acetazolamide and ethoxzolamide reduced fluid absorption by 40–60% with no consistent change in transepithelial voltage when the tubules were perfused and bathed with HCO_3^- -KR buffers containing 95% O_2 -5% CO_2 (Tables III and VIII), but had no effect on fluid absorption from CO_2 and HCO_3^- free media (Table IV). We infer from these data that approximately half of the expected Na^+ flux from tubules exposed to symmetrical HCO_3^- -KR solutions containing 95% O_2 -5% CO_2 was accompanied by HCO_3^- generated by carbonic anhydrase-catalyzed CO_2 hydration; and that HCO_3^- absorption was responsible for a major component of the rise in tubular fluid Cl^- concentrations (Table II). These data shed no light on the molecular events involved in HCO_3^- absorption from these tubule segments. But they are consistent with a number of partially divergent views (12–20) concerning the mode of HCO_3^- salvage by the proximal nephron, all of which involve, at a minimum, carbonic anhydrase-catalyzed hydration of CO_2 by tubular cells.

It is worth noting in this regard that appreciable HCO_3^- absorption by the in vivo proximal nephron continues in the presence of virtually complete inhibition of renal carbonic anhydrase activity (12–20). At least two hypotheses have been set forth to account for this phenomenon. One of these, developed by Rector (14), argues that H_2CO_3 recycling from tubular fluid in an acid disequilibrium state, to proximal tubular cells in an alkaline disequilibrium state, provides an additional source for cellular protons. A second argument, set forth independently by Maren (15, 20) and by Brodsky and Schilb (41), proposes direct transepithelial HCO_3^- transport which may be independent of cellular carbonic anhydrase activity.

A similar issue arises in connection with the present experiments. Table VI shows that in the presence of 2.25 mM acetazolamide, approximately one third of expected Na^+ absorption, i.e., 11.3 peq $min^{-1} mm^{-1}$, was not accounted for by the measured net Cl^- flux. And the other agents and/or anions in the perfusing solutions used in Table VI, i.e., urea, SO_4^{2-} , HPO_4^{2-}/HPO_4^{1-} , and isethionate, were not capable of supporting net solute or fluid absorption (Table XI). But the present experimental data provide no information about whether this anion moiety represents HCO_3^- absorption due to H_2CO_3 recycling from lumen to cell (14), a direct HCO_3^- absorption process (13, 20, 41), or transport of other, unidentified anions by a different process.

Acetate transport. It is evident from Tables VIII and IX that the greater values of K_{Ae}^* with respect to K_{Ae}^1 depend on carbonic anhydrase-insensitive, temperature-dependent events. At least two classes of explanations,

individually or in unison, might have accounted for the disparity between K_{ac}^o and K_{ac}^i . There may have been net lumen-to-bath transport, presumably active, of the acetate species. Alternatively, luminal acetate might have been consumed by tubular cells. Acetate is a renal metabolic substrate (42, 43), and it has been suggested that acetate, lactate, and pyruvate may be converted to acylglycines within renal tubular cells (44). Moreover, proximal tubular lactate absorption may depend on diffusion of lactate across luminal surfaces driven by lumen to cell lactate concentration gradients produced by cellular consumption of lactate (32). Consequently, it may be that in the present experiments (Table VII) K_{ac}^o exceeded K_{ac}^i because of preferential metabolic consumption of luminal rather than contraluminal acetate.

In either instance, i.e., net lumen-to-bath acetate flux or cellular consumption of luminal acetate, it is clear that net luminal acetate efflux may contribute, under appropriate conditions, to fluid absorption. Thus, in the experiments listed in Tables IX and X, where net luminal [^{14}C]acetate disappearance could be evaluated, the latter accounted for approximately one fourth of fluid absorption and expected net Na^+ flux. Likewise, in Table XI, the addition of acetate to luminal solutions increased temperature-sensitive fluid absorption from values indistinguishable from zero to $0.13 \text{ nl min}^{-1} \text{ mm}^{-1}$.

In the case of the acetate influx measurements (Table VII), certain factors are noteworthy. The bath to lumen influx coefficients of inorganic ions such as Na^+ and Cl^- are considered to be the passive ionic permeability coefficients for these ions in isolated proximal tubules (1, 9, 11, 22). But the K_{ac}^i values in Table VII might have depended not only on passive permeation of acetate via the paracellular route traversed by Na^+ and Cl^- in these tubules (1, 9), but also on either transcellular acetate transport via an organic acid secretory system (29, 30), or, since the ^{14}C collected in luminal fluids during K_{ac}^i measurements (Table VIII) was not identified uniquely as [^{14}C]acetate, or as cellular secretion of a ^{14}C -labeled metabolic product of acetate. When considered in this context, the values of K_{ac}^i in Table VII represent an upper limit for P_{ac} , the passive permeability coefficient for acetate in these tubules.

However, two considerations suggest that the values of K_{ac}^i may have depended primarily on a passive transport process. As shown in Table IX, exclusion of acetate from bathing media did not significantly reduce fluid absorption. Stated alternatively, metabolic consumption and/or transport of contraluminal acetate, for the conditions in Table VIII, had a negligible effect on ion and fluid transport in these tubules. Second, the values of K_{ac}^o at 21°C and zero net fluid absorption, $0.18 \pm 0.02 \text{ } \mu\text{m/s}$ (Table VIII), were in close accord with those for K_{ac}^i during fluid absorption at 37°C , 0.14 ± 0.02

$\mu\text{m/s}$ (Table VII). To the extent that these data indicate the absence of significant transcellular transport and/or metabolic usage of bathing fluid acetate, we may conclude tentatively that the passive permeability coefficient (P_{ac}) for acetate in these tubules was in the range of $0.14\text{--}0.18 \text{ } \mu\text{m/s}$ (Tables VII and VIII).

Uncertainties concerning NaCl absorption. Based on the results in Tables III–V, VIII, it seems reasonable to assume that approximately 40–60% of Na^+ absorption from tubules exposed to symmetrical HCO_3^- -KR solutions was accompanied by HCO_3^- . Likewise, if the results in Table X provide an index to net luminal acetate efflux, one might deduce that an appreciable fraction, but less than that in Table X, of Na^+ absorption was accompanied by acetate when the latter was present in bathing media (e.g., Table II). Such a stoichiometric relationship between Na^+ , HCO_3^- , and acetate seems at least plausible, since the results in Table II indicate directly that net NaCl absorption accounted for approximately one-fourth of the expected net Na^+ flux. In this connection, major questions arise concerning a detailed rather than approximating analysis of Cl^- flows in these tubules, and the mode of NaCl transport.

With regard to Cl^- transport, the sum of J_{Cl}^d and J_{Cl}^o with eq. 4 and 5 respectively, yields a value of $24.8 \text{ peq min}^{-1} \text{ mm}^{-1}$ for the approximate net passive Cl^- flux, substantially greater than the observed rate of $15.7 \text{ peq min}^{-1} \text{ mm}^{-1}$ (Table II) for net Cl^- absorption. These enabling calculations clearly require more rigorous evaluation. In regard to the mode of transport for the fraction of Na^+ absorption accompanying Cl^- (Table II), at least two possibilities might be considered. First, the spontaneous transepithelial voltage in tubules exposed to symmetrical HCO_3^- -KR solutions is -1.0 to -1.5 mV lumen negative at the start of perfusion (1, 3, 9), i.e., before the rise in tubular fluid concentration (Table II). Consequently it might be argued (1) that active, rheogenic Na^+ transport occurs in regions where the transepithelial voltage is lumen negative. Second, it seems probable that the transepithelial voltage becomes lumen positive coincident with the rise in tubular fluid Cl^- concentrations in these superficial proximal straight tubules (1, 9) and in isolated (4–6) and in vivo (10) superficial proximal convolutions (although not in juxtamodullary proximal straight tubules; ref. 11); and prior observations have provided direct experimental evidence for passive Na^+ absorption driven by lumen-positive transepithelial voltages in isolated proximal straight tubules (9). It is evident from these considerations that in the case of fluid absorption from symmetrical HCO_3^- -KR solutions, a rigorous quantitative analysis of Cl^- absorption and an assessment of the magnitudes of the passive and active moieties of Na^+ flux accompanying Cl^- will require, at a minimum, solution of the flow-

diffusion equations for solute and solvent flows in intercellular spaces for tubules under the conditions listed in Table II.

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