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Research Article

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Transmissible Gastroenteritis

MECHANISMS RESPONSIBLE FOR DIARRHEA IN AN ACUTE VIRAL ENTERITIS IN PIGLETS

D. G. BUTLER, D. G. GALL, M. H. KELLY, and J. R. HAMILTON

From the Department of Pediatrics, University of Toronto and the Research Institute, The Hospital for Sick Children, Toronto M5G 1X8 Ontario, Canada

ABSTRACT We studied 3-wk-old piglets 40 h after experimental infection with transmissible gastroenteritis (TGE) virus to identify the mechanisms of diarrhea in this disease and to better understand infectious diarrhea in humans. Using continuous segmental marker perfusion in four regions along the gut, we found significant increases in net intraluminal accumulation of water and electrolytes only in the proximal jejunum, the region infected by the virus. In this jejunal segment studied in vivo, unidirectional sodium flux, extracellular fluid (ECF) to lumen, significantly increased, lumen to ECF significantly decreased, compared with matched littermates. The standard perfusate rendered hypertonic by adding mannitol (450 mosmol/kg), in the same segment of normal pigs, caused only an increase in ECF to lumen flux of sodium. TGE did not alter gross villous structure or intraluminal bacteria, bile salts, lactate, pH, or osmolality. Epithelial cell migration was accelerated in the jejunum of infected pigs. Isolated in suspension, these cells from TGE pigs exhibited increased active and passive sodium efflux, cells from mannitol-perfused pigs exhibited only increased active sodium efflux. In this viral enteritis, altered sodium transport occurring in the jejunum, the region of the intestine infected appears to be associated with defective epithelial cell function. The precise nature of the abnormalities in sodium transport, their relationship to disturbances of transport of other solutes, and to virus epithelial cell interaction remain to be defined.

INTRODUCTION

To better understand acute infectious diarrhea in humans we investigated the mechanisms causing diarrhea

in transmissible gastroenteritis, a specific enteric viral infection of swine. This disease, caused by a Corona virus (1), is characterized by vomiting, diarrhea, and dehydration within 24 h of infection (2). Piglets infected at birth almost all die whereas those infected at 3 wk of age survive and their diarrhea subsides within 2 wk (3). The maximum effect of this disease is reported to be on the proximal small intestine where the virus infects mature mucosal epithelial cells (3-7). We had found that piglets infected at 3 wk of age had massive diarrhea with increased quantities and concentrations of sodium, potassium, and chloride in stools (8). We will describe controlled in vivo segmental perfusion studies (9) in infected piglets that identify the proximal jejunum as the region of major electrolyte and water loss and characterize sodium transport at that site. We will also describe in vitro experiments which support the hypothesis that disordered sodium transport plays an important role in the pathogenesis of this diarrhea.

METHODS

Litters of conventional piglets of the York breed were weaned at 2 wk of age, and at operation under general anaesthesia a triple lumen tube (TLT)¹ was securely fixed in the bowel lumen via a fistula. TLT were constructed from polyvinyl tubing (8 french) so that the mixing and test segments were 10 and 30 cm, respectively. After operation the piglets were placed in individual holding pens and fed reconstituted whole evaporated cows' milk. They convalesced for 1 wk during which time they were trained to lie quietly when suspended in cloth slings. Experiments were begun on the 7th postoperative day when the piglets were fully recovered. Piglets to be infected were given orally 2 ml of a virus suspension containing the Purdue

¹ *Abbreviations used in this paper:* ECF, extracellular fluid; PEG, polyethylene glycol; TGE, transmissible gastroenteritis; TLT, triple lumen tube.

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strain of transmissible gastroenteritis (TGE) virus prepared as a bacterial-free homogenate of whole intestinal mucosa (4). Infected piglets were isolated in a special facility.

Experiments. In the first series of experiments we measured net water and electrolyte flux by marker perfusion in four specific regions of the bowel. Piglets were then infected and 40 h later subjected to a second perfusion study. Simultaneously, identical studies were carried out on uninfected match-fed littermates. We studied 15 piglets intubated in the upper jejunum 30 cm beyond the ligament of Treitz; 5 intubated in the mid-jejunum 250 cm above the cecum; 5 intubated in the ileum 30 cm above the cecum, and 3 intubated in the colon. The perfusion solution contained 120 mM Na⁺, 40 mM K⁺, 115 mM Cl⁻, 45 mM HCO₃⁻, 20 mM glucose, and as a nonabsorbable marker, polyethylene glycol 4,000 (PEG 4,000) 5 g/liter, giving an osmolality of 320 mosmol/kg water and pH 7.80 at 37°C. Preliminary studies had established that PEG 4,000 was a satisfactory marker in this model, that steady-state conditions existed in test segments after 60 min of continuous perfusion at 1.8 ml/min in small bowel and 7.5 ml/min in colon, and that transit times through test segments were 15 and 20 min, respectively. Preceded by an 8-h fast and then a 1-h equilibration period, each perfusion study lasted 1½ h. Consecutive 15-min samples were collected by a staggered technique (9) from proximal and distal sampling sites of the TLT using controlled gravity flow into containers chilled at 4°C. The collected juice was filtered at 10°C for 18 h. Samples for glucose and lactate measurements were deproteinized with 0.6% perchloric acid and all juice was frozen at -30°C until analyzed. The pigs were killed at the end of the second perfusion study. The location of the perfusion tube was identified and in each case it had not shifted from its inserted position. Tissue from the perfused segment was obtained by scraping the mucosa from a 15-cm segment with a glass slide. This tissue was homogenized in buffered saline (pH 8.0) containing 20 mM Mg⁺⁺ to yield a protein concentration of 8–12 µg/ml and analyzed for adenyl cyclase activity. These experiments identified the proximal jejunum as the region in which the major losses of water and electrolytes occurred.

A second series of experiments was designed to study sodium and water transport in the proximal jejunum in TGE in more detail. The piglets were prepared identically to those described above with TLT implanted in the proximal jejunum. However, for these experiments the litters were divided into three groups, infected and control as above, and an added group of normal uninfected piglets which were perfused with hypertonic mannitol (the perfusate described above to which we added mannitol, 23.5 g/liter giving an osmolality of 450 mosmol/kg water). In preliminary studies we had established that this degree of hypertonicity induced a net sodium flux across normal jejunum similar to that observed in TGE. As before, we measured net flux of water and electrolyte before and after infection (15 piglets), comparing our data with those from uninfected controls (6 piglets) and those receiving the hypertonic perfusion (13 piglets). In all animals we measured unidirectional sodium flux by adding a tracer dose of ²²Na (50 µCi/liter) to the perfusion solution. We had determined that in this model reflux of ²²Na from ECF to the lumen of the test segment was insignificant. During these studies we also collected proximal jejunal juice for bacteriological studies and bile salt measurements, before and after infection. All piglets were killed after the post-

infection perfusion study and the intestinal cells were isolated immediately from the perfused segments for measurement of sodium efflux rate constants. 2-cm pieces of the perfused segments were immediately stapled on paraffin blocks, fixed in Suza's fixative for 24 h, then transferred to 95% ethanol and processed routinely for staining with hematoxylin and eosin. For electron microscopy, jejunal mucosa was fixed in 3% glutaraldehyde buffer, rinsed in buffer, postfixes in 1% osmium tetroxide in veronal acetate buffer, and embedded in epon. Sections were cut with a diamond knife, stained with uranyl acetate and lead citrate, and examined in a Philips EM 200 electron microscope (Philips Electronic Instruments, Mount Vernon, N. Y.)

Suspensions of isolated epithelial cells were prepared by everting 15-cm lengths of the perfused segments of proximal jejunum over a spiral rod, immersing it in ice-cold buffered saline (2 mM Tris pH 7.40 at 4°C), and subjecting it to high frequency, low amplitude vibration for 30 min as described by Levine and Weintraub (10). The cell pellet obtained by centrifuging this suspension at 50 g was washed twice in ice-cold buffered saline (pH 7.40 at 4°C) before measurements of Na⁺ efflux were made. This technique isolates, almost exclusively, epithelial cells from the villi rather than the crypt cells.

In the third experiment the rate of jejunal epithelial cell migration on the villus was measured in normal and TGE infected 3-wk-old piglets. Four piglets from a litter of six were randomly chosen and each given a standard oral dose of TGE virus. The other two piglets were left as match-fed controls. 16 h after infection, each piglet was given a subcutaneous injection of tritiated thymidine at the rate of 0.3 µCi/g body weight (sp act 18.4 Ci/mM). 24 h postinfection, one control and two infected piglets were killed and at 40 h after infection, one control and two infected piglets were killed. From each animal, a 3-cm length of intestine, starting 30 cm distal to the ligament of Treitz was opened along the mesenteric border, stapled flat on a paraffin block, serosal side down, placed in Bouin's fixative for 12 h, and transferred to 70% alcohol. Tissue sections 4 µm thick were cut and prepared for autoradiography, exposed for 1 mo at 10°C, and stained with hematoxylin and eosin. To determine migration rates of the epithelial cells, the distance from the base of the crypt to the foremost labeled cells was measured using a previously calibrated eyepiece. From the difference of the means of measurements made on piglets killed at 24 and 40 h a rate of migration over the 16 h interval was calculated.

Analytical techniques. Sodium and potassium were measured by flame spectrophotometry and chloride by titration using standard laboratory techniques. Osmolality was determined by freezing point depression, glucose by the glucose oxidase method (11), lactate using lactic dehydrogenase (Boehringer Mannheim Corp., New York), and PEG spectrophotometrically by the method of Hyden (12). We measured ²²Na activity in a Gamma Counter (Nuclear Chicago, Corp., Des Plaines, Ill.). Qualitative and semi-quantitative measurements of bile salts in intestinal juice were made by thin-layer chromatography. We separated conjugated bile salts (glycocholate, glycodeoxycholate, glycochenodeoxycholate, taurocholate, taurodeoxycholate, taurochenodeoxycholate) using Kottke's solvent system (13) and free bile salts (cholate, deoxycholate, chenodeoxycholate) with Sundaram's solvent (14). All chromatographs were subjected to the same spray for color detection (15). Bacterial content of juice was estimated by direct examination of gram-stained specimens and quantitative counts

TABLE I
Analyses of Variance Comparing Net Electrolyte and Water Fluxes and Mean Electrolyte Concentrations in Proximal Jejunum of Normal and Infected Piglets and Normal Piglets Perfused with Hypertonic Solution (NH Piglets)*

	Na ⁺ , mM/h/cm				[Na ⁺]	H ₂ O, ml/h/cm					Cl ⁻ , mM/h/cm				[Cl ⁻]	K ⁺ , mM/h/cm				[K ⁺]
	Mean	SD	n	P <		Mean	SD	n	P <		Mean	SD	n	P <		Mean	SD	n	P <	
Control	+0.009	±0.049	28	—	0.125	+0.153	±0.430	28	—		+0.020	±0.058	28	—	0.121	+0.029	±0.015	28	—	0.026
Infected	-0.119	±0.092	28	0.005	0.124	-0.920	±0.752	28	0.005		-0.104	±0.073	28	0.005	0.118	+0.003	±0.013	27	0.01	0.027
NH	-0.105	±0.081	36	0.005	0.111	-1.218	±0.980	36	0.005		-0.106	±0.099	36	0.005	0.110	+0.017	±0.019	36	0.05	0.025

Positive values indicate net absorption out of intestinal lumen. Negative values indicate net secretion into lumen.

* Millimoles per milliliter.

made after 48 h and serial dilutions (10^{-1} – 10^{-5}) plated on blood agar, aerobically and anaerobically; MacConkey's agar, aerobically and Rogosa agar, anaerobically. Each light microscopic slide of jejunal mucosa was examined by one observer (Dr. Butler) unaware of whether the tissue was from an infected or uninfected piglet. A standard calibrated micrometer eyepiece was used to measure villous height and crypt depth. Epithelial cell morphology and round cell infiltration were also rated. Adenyl cyclase activity (basal and fluoride stimulated) was measured on whole mucosal homogenates (16). Optimal conditions for the assay in pig tissue had been determined in preliminary studies.

To measure sodium efflux rate constants in intestinal epithelial cells, approximately 0.5 ml of packed cells was added to 3 ml of either a medium containing NaCl 120 mM, Tris HCl 20 mM, K₂HPO₄ 3 mM, MgCl₂ 1 mM, CaCl₂ 1 mM, glucose 10 mM, bovine serum albumin 1 mg/ml, ²²Na 2.5 μg/ml (osmolality 290 mosmol/kg water, pH 7.40 at 37°C) or to the same medium to which ouabain 10^{-8} M had been added. The mixtures were incubated at 37°C for 30 min in a Dubnoff metabolic shaker and immediately centrifuged for 1 min at 4°C, and washed twice in ice-cold buffered isotonic MgCl₂ (110 mM MgCl₂, 2 mM Tris, pH 7.40). While maintaining dilution at 1:30 the cells were resuspended in tracer-free flux medium with or without ouabain. The sodium efflux was ascertained by measuring the rate of appearance of ²²Na in the medium as described for Na⁺ efflux from the red blood cells (17) and expressed as the total sodium efflux constant/h (*K⁺Na⁺) and the passive sodium efflux constant/h (*K⁺Na⁺) (18). The passive efflux rate constant is defined as the ouabain-insensitive portion of the total efflux rate constant remaining after maximal inhibition. Optimal con-

ditions for both cell isolation (temperature, vibration time, and centrifugal force) and Na⁺ efflux (duration and temperature of loading, composition of solution to remove excess isotope, and ouabain concentration required for maximum inhibition) had been determined in preliminary studies. The integrity of the isolated cells was evaluated by trypan blue exclusion (19) and we demonstrated that 80% excluded trypan blue up to 3 h after isolation. Sodium efflux constants determined on cell suspensions from control and experimental groups were stable up to 3 h after isolation and were independent of the number of cells incubated.

Calculations. Standard formulas were used to calculate the net fluxes of water and electrolyte (9) and the unidirectional Na⁺ fluxes (20) across the test segment of bowel in vivo. We compared means of the changes observed in these flux data between the two perfusion studies in control and test groups by an analysis of variance. When significant "F" values occurred the means were ranked using Scheffe's method for multiple comparisons (21). Data from the in vitro experiments were compared using standard Student's *t* tests.

RESULTS

In vivo net flux studies. The proximal jejunum was the only site of significant net electrolyte and water loss after infection with the virus. Table I shows the analyses of variance between net fluxes in the proximal jejunum of the three study groups, control, infected, and those perfused with a hypertonic solution (NH piglets). In infected piglets and NH piglets the reversal in the net fluxes of Na⁺, Cl⁻, and water was significant (*P* <

TABLE II
Analyses of Variance Comparing Flow Rate, Osmolality, and pH Entering and Leaving Proximal Jejunal Test Segment during Continuous Segmental Perfusion in Control and Infected Piglets and Normal Piglets Perfused with Hypertonic Solution (NH Piglets)

	Flow, ml/min								Osmolality, mosmol/kg water								pH							
	Entering				Leaving				Entering				Leaving				Entering				Leaving			
	Mean	SD	n	P <	Mean	SD	n	P <	Mean	SD	n	P <	Mean	SD	n	P <	Mean	SD	n	P <	Mean	SD	n	P <
Control	1.70	±0.50	25	—	1.67	±0.63	25	—	312	±7	25	—	303	±13	25	—	8.57	±0.30	25	—	8.58	±0.35	25	—
Infected	1.51	±0.29	23	0.05	1.95	±0.49	23	0.01	312	±9	23	—	309	±17	20	—	8.69	±0.19	23	—	8.53	±0.44	22	—
Hypertonic	2.18	±0.50	31	0.01	2.79	±0.63	31	0.005	371	±18	31	0.005	329	±15	31	0.005	8.35	±0.42	31	0.01	8.21	±0.47	31	0.01

TABLE III
Analyses of Variance Comparing Net Fluxes of Electrolytes, Water in Mid-Small Bowel, Distal Ileum, and Colon in Piglets before and 40 h after Infection (TGE)

		Na ⁺ , mM/h/cm			H ₂ O, ml/h/cm			Cl ⁻ , mM/h/cm			K ⁺ , mM/h/cm		
		Mean	SD	n	Mean	SD	n	Mean	SD	n	Mean	SD	n
Mid-small bowel	Control	+0.012	±0.026	8	+0.085	±0.077	8	+0.019	±0.014	8	+0.015	±0.008	8
	Infected	-0.033	±0.017	8	-0.312	±0.175	8	-0.016	±0.013	8	+0.008	±0.007	8
Distal ileum	Control	+0.009	±0.060	12	+0.043	±0.530	12	+0.030	±0.045	12	+0.019	±0.019	12
	Infected	-0.043	±0.054	12	-0.361	±0.453	12	+0.003	±0.037	12	+0.010	±0.006	12
Colon	Control	+3.05	±4.24	6	+2.18	±4.95	6	+4.59	±1.09	6	+1.31	±1.25	6
	Infected	+6.68	±2.16	6	+6.82	±9.17	6	+4.79	±1.70	6	+1.20	±0.68	6

Positive values indicate net absorption out of intestinal lumen. Negative values indicate net secretion into lumen.

0.005) as was the decrease in the net flux of potassium. Whether caused by infection or hypertonic perfusion, the induced net losses of Na⁺ and Cl⁻ into the lumen were similar. However, compared with infected pigs, hypertonic perfusion produced a significantly greater net loss of water into the intestinal lumen ($P < 0.01$) and a significantly greater net K⁺ flux out of the lumen ($P < 0.05$). Table II shows that the flow rate entering the test segment was lowest in infected piglets, intermediate in the controls, and highest in the NH piglets. The flow rate leaving the test segment in infected piglets was significantly greater than in control piglets and significantly less than in the normal piglets given the hypertonic solution. Osmolality at either end of the test segment was similar for infected and control piglets, but as expected, it was significantly greater in NH piglets (Table II). The pH of intestinal contents did not differ between groups (Table II). Net flux data from studies on the more distal segments are summarized in Table III. Infection produced no significant effect.

In vivo unidirectional sodium flux studies. The combined results of the unidirectional sodium flux studies between lumen and ECF of proximal jejunum in normal and infected piglets and NH piglets are listed in Table IV. Unidirectional sodium flux from lumen to ECF ($\phi^{Na^+L \rightarrow ECF}$) in infected piglets was significantly decreased ($P < 0.005$); the opposite unidirectional so-

dium flux ($\phi^{Na^+ECF \rightarrow L}$) in infected pigs was significantly greater than that observed in controls ($P < 0.01$). The $\phi^{Na^+L \rightarrow ECF}$ in the NH piglets was similar to control values but the $\phi^{Na^+ECF \rightarrow L}$ was significantly greater ($P < 0.005$). In NH piglets $\phi^{Na^+ECF \rightarrow L}$ was significantly greater than in infected piglets ($P < 0.01$). Thus we observed changes in both unidirectional sodium fluxes after infection whereas only the $\phi^{Na^+ECF \rightarrow L}$ was altered during hypertonic perfusion.

The numbers and types of bacterial organisms present in the proximal intestinal lumen were not altered by the viral infection (Table V). Lactate concentrations ($\times 10^{-4}$ mM/ml) entering, 1.77 ± 0.82 (mean $\pm$ SD) and leaving, 1.94 ± 0.99 the proximal jejunal segment measured in 8 piglets 40 h after infection did not differ significantly from concentrations measured before infection, 1.65 ± 0.68 and 1.96 ± 0.40 , respectively. Bile salt concentrations (mM/liter) in juice from intubated proximal jejunum after infection, both conjugated 0.68 ± 0.61 and free, 1.21 ± 0.25 also did not differ significantly from control data 0.78 ± 0.44 and 1.46 ± 0.38 . There was a preponderance of free bile salts in the juice of both groups.

Intestinal morphology. By light microscopy there were no significant abnormalities in the dimensions of villi or crypts in infected piglets. The cells at the tips of villi appeared cuboidal rather than columnar as one

TABLE IV
Analyses of Variance Comparing Unidirectional Sodium Fluxes between Intestinal Lumen and Extracellular Fluid in Proximal Jejunum of Control and Infected Piglets and Normal Piglets Perfused with Hypertonic Solution (NH Piglets)

Comparison	Direction							
	L $\rightarrow$ ECF, mM/h/cm				ECF $\rightarrow$ L, mM/h/cm			
	Mean	SD	n	P <	Mean	SD	n	P <
Control	0.133	±0.090	13	—	0.127	±0.079	13	—
Infected	0.071	±0.049	11	0.005	0.192	±0.101	11	0.01
NH	0.147	±0.075	13	—	0.260	±0.104	13	0.005

TABLE V
Colony Counts* and Bacterial Species Isolated per Milliliters Intestinal Juice Collected from Intubated Proximal Jejunum of Piglets before and 40 h after Infection with Transmissible Gastroenteritis Virus

Medium	Control (preinfection)				Infected				
	Mean	SD	n	Species	Mean	SD	n	P <	Species
Blood agar (Aerobic)	125	±136	5	<i>E. coli</i> Proteus Hemolytic	111	±52	5	—	<i>E. coli</i> Proteus
Blood agar (Anaerobic)	134	±132	5	<i>E. coli</i>	129	±60	5	—	<i>E. coli</i>
MacConkey's agar (Aerobic)	157	±145	5	<i>E. coli</i> Proteus	96	±35	5	—	<i>E. coli</i> Proteus
Rogassa agar (Anaerobic)	Very large numbers of lactobacilli too fine to count. However, no growth at 10 ⁻⁶ dilution.				Very large numbers of lactobacilli too fine to count. However, no growth at 10 ⁻⁶ dilution.				

* ×10⁵.

sees in the normal. By electron microscopy, various changes were observed in some villous absorptive cells whereas others retained normal morphology (Figs. 1 and 2). The microvilli were shorter and less densely packed than in the normal cell. The filaments of the core of most microvilli and of the terminal web were less defined. In the apical cytoplasm, organelles were fewer and less well defined; some dilation and vesiculation was observed in the endoplasmic reticulum. Separation of the plications of lateral cell membrane was observed between adjacent cells on many villi but junctional complexes appeared intact.

Neither the length of jejunal villi in control piglets (258±28 μm) and infected piglets (237±69 μm) or crypt depth in control piglets (226±15 μm) and infected piglets (216±34 μm) were significantly different. However, jejunal epithelial cells migrated along villi of control piglets at a rate of 13 μm/h compared to a rate of 20 μm/h in infected piglets. At these rates the intestinal mucosa in control and infected piglets would be renewed in approximately 38 and 23 h, respectively.

In vitro studies. Adenyl cyclase activity (basal or fluoride stimulated) in small or large intestine was unchanged after infection (Table VI).

The results of our *in vitro* studies on sodium efflux from isolated epithelial cells are listed in Table VII. Cells from infected piglets showed a significantly greater passive (ouabain-insensitive) Na⁺ efflux constant ($P < 0.01$) and total Na⁺ efflux constant ($P < 0.005$) than cells from controls. The passive Na⁺ efflux constant was unchanged in cells obtained from NH piglets whereas the total Na⁺ efflux constant was significantly increased ($P < 0.005$). The passive Na⁺ efflux constant for cells

obtained from infected piglets was significantly greater than in cells from NH piglets ($P < 0.005$) whereas the total Na⁺ efflux constants were similar.

DISCUSSION

Our studies have identified the proximal jejunum as the major site of electrolyte and water secretion in TGE. Since no compensatory mechanisms were detected in more distal intestinal segments, secretion in the upper intestine must have contributed substantially to the diarrhea observed. The experiments reported have been concerned primarily with the relationship between disordered sodium transport in the upper intestine and TGE diarrhea. We recognize that additional intestinal

TABLE VI
Adenyl Cyclase Activity,* Basal and Fluoride Stimulated in Whole Mucosal Homogenates from Control Piglets and Piglets 40 h after Infection

Intestinal segment		Control			Infected (TGE)		
		Mean	SD	n	Mean	SD	n
Proximal jejunum	Basal	153	8	6	120	12	10
	Fluoride	711	97	6	605	71	10
Mid-small intestine	Basal	113	17	6	115	11	10
	Fluoride	669	40	6	810	87	10
Distal ileum	Basal	118	24	6	112	14	10
	Fluoride	600	63	6	744	57	10
Colon	Basal	187	25	6	149	20	10
	Fluoride	869	40	6	845	61	10

* Micromoles cAMP per milligrams protein.

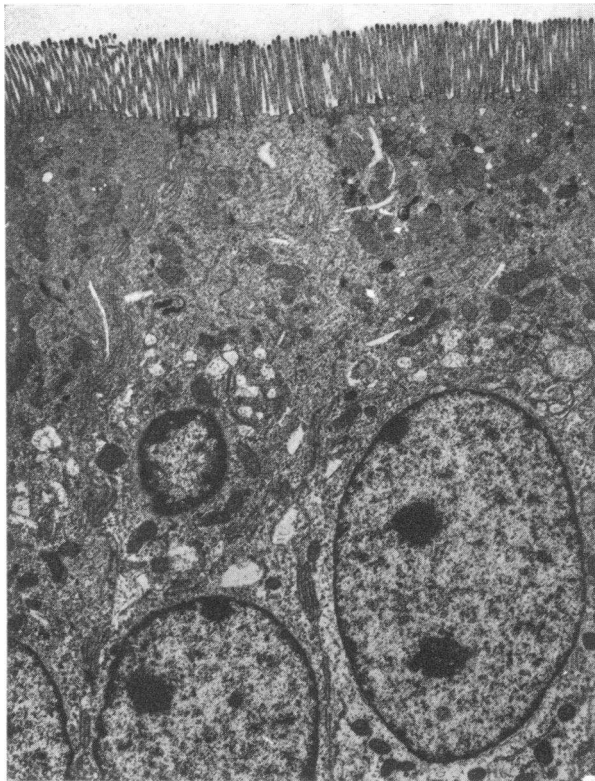


FIGURE 1 Electron micrograph, epithelial cells from lateral wall of jejunal villus of control piglet. Note long regular microvilli. Magnification $\times 5,400$.

transport processes may be affected by the disease, some of which may influence water and sodium transport.

The large net sodium flux into the jejunal lumen in TGE was mediated by a decreased $\phi^{Na^+L \rightarrow ECF}$ and an increased $\phi^{Na^+ECF \rightarrow L}$. Whatever the mechanisms causing these abnormalities, they differ significantly from those occurring in osmotically induced diarrhea. Perfusion of the upper jejunum with hypertonic mannitol caused a net sodium movement into the lumen similar to

that seen in TGE, but did not affect the unidirectional lumen to ECF flux. Several factors may have affected $\phi^{Na^+L \rightarrow ECF}$ in TGE and our data provide little insight into the location and precise nature of the defect. Apparently, because of the relatively low glucose concentration in our perfusate, the defect was identified with virtually no glucose perfusing the test segment. One clue to a possible factor of importance comes from our electron microscopic studies. They showed spotty shortening and sparsity of microvilli at the luminal surface, a finding that would reduce absorptive surface area and one that was predicted by our earlier observations of decreased activity of some brush border enzymes (8). Our data showed that the increased $\phi^{Na^+ECF \rightarrow L}$ in TGE similar to that seen during hypertonic perfusion could not be attributed to increased concentration of intraluminal solutes such as sugars or organic acid anions. Our earlier study also suggested the TGE diarrhea is not an osmotic diarrhea (8). The route for the excessive flow of sodium into the lumen could have been intercellular, transcellular, or both (22). The marked abnormalities in sodium efflux in isolated

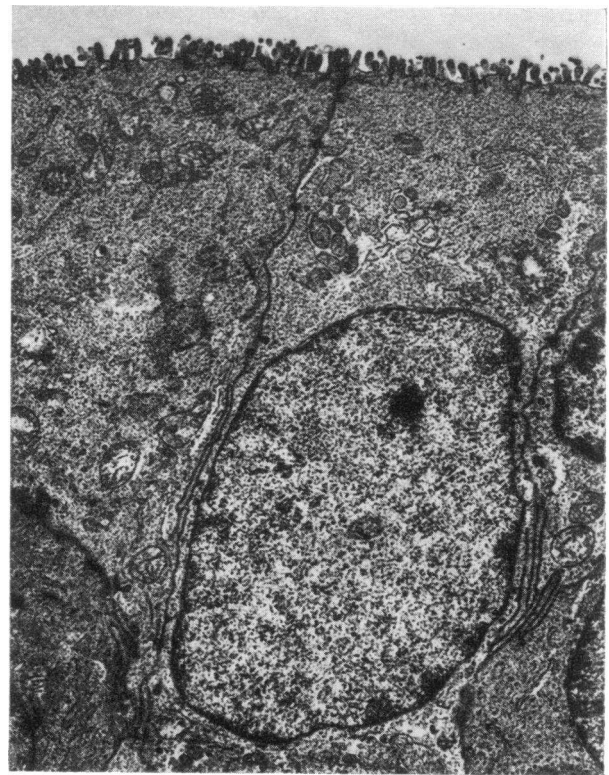


FIGURE 2 Electron micrograph, jejunal epithelial cells from TGE-infected piglet, demonstrating the most extensive ultrastructural abnormalities found. Sample site comparable with that seen in Fig. 1. Note shortened sparse microvilli. Magnification $\times 8,000$.

TABLE VII
Comparison of Passive and Total Sodium Efflux Constants Determined on Epithelial Cells Isolated from Proximal Jejunum of Control and Infected Piglets and Normal Piglets Perfused with the Hypertonic Solution (NH Piglets)

Com- parison	Sodium efflux constant, h^{-1}							
	Passive				Total			
	Mean	SD	n	P <	Mean	SD	n	P <
Control	3.11	± 1.27	35	—	7.16	± 1.78	27	—
Infected	4.36	± 1.21	30	0.01	13.80	± 1.69	10	0.005
NH	3.01	± 1.32	28	—	12.29	± 4.59	23	0.005

epithelial cells from infected pigs, studied under conditions identical to those used for controls, strongly suggest that cellular factors were contributory. These cellular factors are apparently not specific for TGE since hypertonic perfusion, to our surprise, induced a similar abnormality of total efflux. The factors causing this abnormality in response to mannitol may differ from TGE particularly since after mannitol passive efflux was unaffected. The abnormality of passive efflux observed in cells from infected pigs suggests increased cell membrane permeability. In TGE, unlike cholera (23) adenylyl cyclase activity stimulation was not a factor in inducing a secretory state in the intestinal epithelium.

We attributed the observed abnormalities of intestinal function to a direct response to the viral infection. In the lumen of the proximal test segment, bacterial flora did not differ from controls. Bile salts might have been altered by abnormal flora (9) and when altered might have disturbed sodium transport (24). Although there was a preponderance of free bile salts in both infected and control pigs, the study groups did not differ significantly. The upper jejunum, the region known to be infected in TGE (5) is also the region most severely affected functionally. In planning future exploration of this specific viral enteritis, it is useful to speculate on the possible relationship between virus, epithelial cell interaction, and intestinal transport. The TGE virus infects primarily mature cells on tips of jejunal villi (6). We and others have shown that after infected cells are shed, they are replaced by cells migrating from the crypts at an accelerated rate (25). In our pigs, studied 40 h after infection, we were assessing an epithelial surface clothed by cells rapidly conscripted from the crypts and probably not infected by virus. In their haste to repair, do these cells retain the relatively undifferentiated state they possess in the crypts? Our microscopic studies and those in the literature suggest they do (6, 26). If in fact they do, our finding of marked sodium secretion in TGE is in keeping with the current hypothesis that crypt cells, unlike mature cells on villi are normally secretory (27). Obviously, many questions are raised by our data: many remain unanswered. Future studies of this specific viral enteritis should greatly improve our understanding of a major health problem in infants and children, acute viral gastroenteritis.

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