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Kenzo Sato, Richard L. Dobson

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2,4-dinitrophenol (10^{-4} M) reverses ouabain-induced inhibition of lactate and $^{14}\text{CO}_2$ production but only partially reverses inhibition induced by $\text{Na}^+ + \text{K}^+$ deprivation, indicating that metabolic inhibition by ouabain is secondary to the inhibition of sodium transport. There is no synergism between Mecholyl and epinephrine. The absence of any significant inhibitory effects by acetazolamide (Diamox) or HCO_3^- -free media suggests that H^+ transport may not be important in sweat gland function. In contrast to a report by Wolfe et al., human eccrine sweat glands show considerable oxidative activity ($^{14}\text{CO}_2$ production of 0.42-0.72 nmol/gland/h). These observations are discussed in terms of the linkage between sweat gland energy metabolism and sodium transport.

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Glucose Metabolism of the Isolated Eccrine Sweat Gland

II. THE RELATION BETWEEN GLUCOSE METABOLISM AND SODIUM TRANSPORT

KENZO SATO and RICHARD L. DOBSON

From the Department of Dermatology, University of Oregon Medical School, Portland, Oregon 97201, and the Department of Dermatology and Physiology, University of Nijmegen School of Medicine, Nijmegen, The Netherlands

ABSTRACT This paper attempts to further clarify the characteristics of Mecholyl- or epinephrine-stimulated glucose metabolism in the isolated monkey eccrine sweat gland with special emphasis on its relationship to increased sodium transport. The Mecholyl- or epinephrine-stimulated glucose metabolism (as estimated by either lactate or $^{14}\text{CO}_2$ production or both) is seen only in the secretory coil and not in the duct. It is markedly suppressed in the absence of glucose, Na^+ , or K^+ . It is inhibited by ouabain (10^{-3} M) and partially suppressed in a low-sodium (40 mM), high-potassium (100 mM) medium.

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Dr. Sato's present address is: Max-Planck-Institute für Biophysik, 6 Frankfurt a.M. 70, Germany.

Dr. Dobson's present address is: Department of Dermatology, the State University of New York at Buffalo, School of Medicine, Buffalo, N. Y. 14203.

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INTRODUCTION

When freshly isolated eccrine sweat glands are incubated aerobically in vitro, cholinergic and adrenergic drugs markedly stimulate both lactate and $^{14}\text{CO}_2$ production (1). Atropine and phentolamine, specific pharmacologic inhibitors of these respective drugs, and ouabain, an inhibitor of $(\text{Na}^+ + \text{K}^+)\text{-ATPase}$, abolish these stimulatory effects (1). Under physiological conditions the oxidation of glucose is coupled to the regeneration of ATP and orthophosphate (2). This led us to assume that the stimulation of glucose metabolism by cholinergic or adrenergic drugs resulted from an enhanced sodium transport, because the major function of the eccrine sweat gland is to secrete water and electrolytes. This assumption was supported by the stoichiometric relation between net sodium flux and ATP formation by the eccrine sweat gland after cholinergic stimulation (3). However, a similar stoichiometric relation does not apply after adrenergic stimulation, because the in vivo sweat rate is disproportionately lower than the amount of glucose metabolized in vitro.

In these studies we attempted to elucidate the mechanisms by which cholinergic and adrenergic drugs stimulate in vitro metabolism of the eccrine sweat gland, especially in terms of its relationship to sodium transport. The studies include an investigation of a differential metabolic response of the duct and secretory coil and of the effects of electrolytes, 2,4-dinitrophenol (DNP,¹ Merck Chemical Div., Merck & Co., Rahway, N. J.)

¹Abbreviations used in this paper: DNP, dinitrophenol; KRB, Krebs-Ringer bicarbonate solution; KT, Krebs-Tris buffer; POPOP, *p*-bis-2,5-phenyloxazolilbenzene; PPO, 2,5-diphenyloxazole.

and acetazolamide (Diamox, Lederle Laboratories, Pearl River, N. Y.) on glucose metabolism of the sweat gland coil (duct + secretory coil) in vitro. We also compared the glucose metabolism of human eccrine sweat glands to that of monkey sweat glands. Evidence was obtained to suggest that the metabolic stimulation in incubated eccrine sweat gland by cholinergic and adrenergic drugs reflects mainly sodium transport, an idea supported by the recent observation³ that both cholinergic and adrenergic drugs induce sweating from an isolated eccrine sweat gland in vitro.

METHODS

Glucose metabolism of the isolated monkey palm eccrine sweat gland

Skin biopsy specimens (1.5 cm × 0.5 cm) were obtained from the palm of 22 adult rhesus monkeys and 4 adult stump-tailed macaques tranquilized with phencyclidine HCl (Sernyl, Parke, Davis & Co., Detroit, Mich.). In most monkeys, several skin specimens were excised from the same palm. In such cases rebiopsy was performed after at least 4 wk and the area of skin adjacent to the previous excision was avoided. Either Krebs-Tris buffer (KT) 140 mM NaCl, 5 mM KCl, 1.4 mM CaCl₂, 2.5 mM MgCl₂, 20 mM Tris buffer, pH 7.4, and 11 mM glucose gassed with O₂ or Krebs-Ringer bicarbonate (KRB) (125 mM NaCl, 5 mM KCl, 1.2 mM MgCl₂, 1.0 mM CaCl₂, 25 mM NaHCO₃, 1.2 mM NaH₂PO₄, and 11 mM glucose, pH 7.4 gassed with 95% O₂ and 5% CO₂) was used as the incubation medium unless otherwise specified.

Immediately after excision, the skin specimens were sliced freehand with a razor blade and were rinsed in several changes of either ice-cold KT or KRB solution. Fresh, unstained sweat gland coils (secretory portion plus coiled duct) were isolated in either ice-cold KT or KRB under a stereoscopic microscope using two pairs of Dumont No. 5 stainless steel tweezers (Dumont Tweezer, Hammel, Riglander & Co., Hudson, N. Y.) (1, 4). The method for sweat gland incubation and determination of lactate and CO₂ production have been described in detail (1).

When only lactate production was studied, three to five sweat gland coils were incubated in a test tube containing 200 or 400 μ l of an appropriate Tris-buffered medium with or without the addition of drugs. All incubations were performed in duplicate for 1 h in a closed water bath at 38°C unless otherwise specified. Air was used as the gas phase. ¹⁴CO₂ production was measured by a method modified from that of Adachi and Uno (5). [6-¹⁴C]glucose (The Radiochemical Centre, Amersham, England, sp. act. = 3.0 mCi/mM) in appropriate dilution (usually 0.15 mCi/mM glucose in a final concentration) was used as a tracer. When KRB was used as a medium, an incubation test tube (8 mm ID × 50 mm) was filled with 5% CO₂ and 95% O₂, promptly capped with Parafilm (American Can Co., Neenah, Wis.), and connected to an upper test tube containing paper impregnated with hyamine hydroxide [*p*-(diisobutylcresoxy-ethoxyethyl) dimethyl benzyl ammonium hydroxide, Packard Instrument Co., Inc., Downers Grove, Ill.] by a rubber sleeve. The trapping of CO₂ by hyamine hydroxide was thus prevented

³ Sato, K. 1973. Sweat induction from an isolated eccrine sweat gland. *Am. J. Physiol.* In press.

during the incubation period and the pH of media was maintained at 7.4 ± 0.03. After a 1-h incubation at 38°C, the thin layer of Parafilm was torn with a needle inserted through the rubber sleeve and the gas phase of the incubation tube exposed to the hyamine hydroxide-impregnated paper. 30 μ l of cold 40% trichloroacetic acid was injected through the rubber sleeve with a needle and syringe to generate CO₂, which was completely trapped by the hyamine hydroxide-impregnated paper during the next 4 h. An aliquot of 100 μ l was taken from the incubation medium for the enzymatic determination of lactate (6) except that fluorescence was measured with a Farrand fluorometer (Farrand Optical Co., Inc., Valhalla, N. Y.). The ¹⁴CO₂-hyamine in the filter paper was eluted with scintillation fluid consisting of toluene (Merck), 0.5% 2,5-diphenyloxazole (PPO) (Packard) and 0.03% *p*-bis-2,5-phenyloxazoilbenzene (POPOP) (Packard) and counted with a Nuclear-Chicago liquid scintillation counter (Nuclear-Chicago Corp., Des Plaines, Ill.) fitted with external standards for correction of quenching and calculation of the radioactivity. The same procedure was used when KT was used as a medium except that an incubation test tube was filled with 100% O₂ and connected directly to an upper test tube without Parafilm. To exclude the possible contribution of bacterial metabolism during the incubation, all values of lactate and ¹⁴CO₂ production were compared with simultaneously incubated samples consisting of either incubation medium alone or medium containing boiled sweat glands. Unless otherwise described, both lactate and ¹⁴CO₂ production were expressed as nanomoles per gland per h, because in monkeys the size of the sweat gland was rather uniform between 25 and 30 μ g wet weight/gland for rhesus monkeys (3) and 30–35 μ g/gland for stump-tailed macaques with a mean protein content of 7.46% (4). In addition, despite the variation in the removal of attached collagen fiber, the difference in the dry weights of the glands in an individual rhesus monkey was rather small (mean 2.73 ± 0.14 (SE) μ g, range 2.2–3.5 μ g (7)).

The effect of glucose-free medium on lactate production. Sweat glands were isolated in ice-cold KT medium containing 5.5 mM glucose within 30 min of excision to minimize a decrease in endogenous glycogen or glucose or both. Those sweat glands to be incubated in the glucose-free medium were washed briefly in ice-cold glucose-free KT medium and gently blotted before being transferred to the incubation test tubes.

The differential response of the duct and the secretory coil to Mecholyl and epinephrine. Approximately 60 freshly isolated, unstained sweat gland coils were separated into the secretory coil and coiled duct in ice-cold KT as described previously (4). Approximately 10 segments of either secretory coil or duct were incubated in each test tube containing KT with or without Mecholyl (acetyl- β -methylcholine, Sigma Chemical Co., St. Louis, Mo.) or epinephrine (Sigma). 4–5 h was required from the time of biopsy to the beginning of incubation. Since the tissue size of the separated segments varied, the protein content of the tissue was determined in each test tube by the method of Lowry, Rosebrough, Farr, and Randall (8). Both ¹⁴CO₂ and lactate production were expressed as nanomoles per μ g protein per hour (a single sweat gland coil contains approximately 2 μ g protein).

The effect of Na⁺ and K⁺ deprivation in the medium on lactate production. Five incubation media were prepared by changing the Na⁺ or K⁺ concentrations in the KT medium; i.e., a complete KT medium, a Na⁺- and K⁺-free medium, in which Na⁺ and K⁺ were replaced by 140 mM

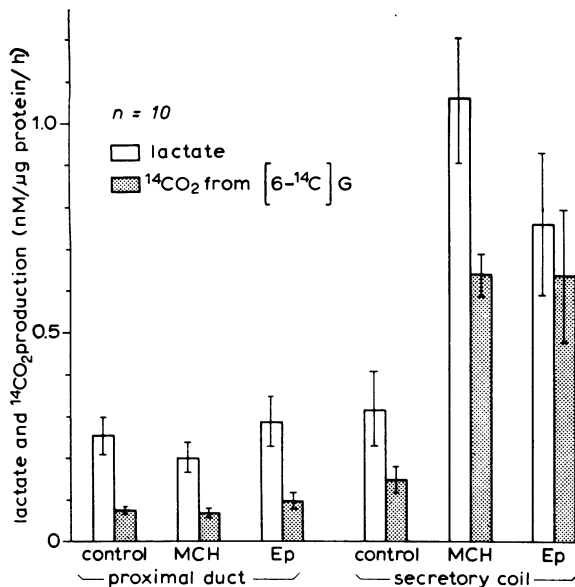


FIGURE 1 Lactate and ¹⁴CO₂ production in the coiled (proximal) duct and the secretory coil. The columns and vertical bars represent mean \pm SEM. Note that both lactate and ¹⁴CO₂ production are expressed as nanomoles per μ g protein per hour. Unless otherwise stated, Mecholyl (MCH) was added to a final concentration of 4.2×10^{-4} M and epinephrine (EP) to a final concentration of 5.5×10^{-5} M throughout the study. *n*, number of experiments. [6-¹⁴C]G, [6-¹⁴C]glucose.

choline chloride, a Na⁺-free, 140 mM K⁺ medium, a 140 mM Na⁺ and K⁺-free medium and a 40 mM Na⁺ and 100 mM K⁺ medium. The isolated sweat gland coils were quickly dipped in 150 mM Tris buffer (pH 7.5) and blotted on ashless filter paper before transfer to the incubation medium.

The effect of 40 mM Na⁺ + 100 mM K⁺ medium on lactate production. In each experiment, two systems were run simultaneously. The first system, used as a control, consisted of duplicate incubation test tubes containing 400 μ l KT with or without Mecholyl. 10 sweat gland coils were incubated in each test tube for 10 min at 38°C and then placed in an ice bath. The sweat glands were collected as a batch, blotted with filter paper, quickly transferred to new test tubes containing the same medium as in the first incu-

TABLE I
Effect of Glucose in the Incubation Medium
on Lactate Production

Drug addition	Medium	
	Glucose-containing	Glucose-free
No drug addition	0.64 \pm 0.06	0.10 \pm 0.03
Mecholyl (4.2×10^{-6} M)	2.87 \pm 0.27	0.08 \pm 0.03
Epinephrine (5.5×10^{-6} M)	1.16 \pm 0.29	0.10 \pm 0.03

The results are the mean \pm SEM of four experiments. All the values are expressed as nmol lactate/gland/h.

bation period, and reincubated at 38°C for 10 min. The same procedures were repeated for a third 10-min incubation. The time required for transferring the sweat glands from one test tube to another was less than 1 min. The second system was processed in the same manner except that 100 mM K⁺ + 40 mM Na⁺ medium was used for the second 10-min incubation instead of KT medium. After the incubation, aliquots were taken from the media for lactate determination.

The effect of DNP on the metabolic inhibition induced by ouabain or Na⁺ and K⁺ deprivation. The effects of varying concentrations of DNP were first tested on lactate production in KT media. Based on this dose-response study, the effect of 10⁻⁴ M DNP, which gave the maximal stimulation of lactate production, on the metabolic stimulation induced by Mecholyl or epinephrine (in KRB) and on the metabolic inhibition induced by 10⁻³ M ouabain (in KRB) or Na⁺ + K⁺-deprivation (in KT), was studied. Na⁺ + K⁺-free medium was prepared as previously described.

The effects of buffer and Diamox on lactate and ¹⁴CO₂ production. Lactate and ¹⁴CO₂ production were first compared between KT and KRB. The effects of varying concentrations of Diamox (10⁻² M–10⁻⁵ M) were then studied on lactate and ¹⁴CO₂ production with KRB as a medium.

The synergistic effect of Mecholyl and epinephrine. Mecholyl (4.2×10^{-6} M) and epinephrine (5.5×10^{-6} M) were added to the KRB either singly or in combination and their effects on lactate and ¹⁴CO₂ production were compared.

Lactate and ¹⁴CO₂ production of isolated human eccrine sweat glands

After intradermal anesthesia with 1% procaine, which has no inhibitory effect on monkey sweat gland metabolism,³ 1 \times 0.5-cm elliptical skin specimens were excised from the flexor forearm of three healthy men, aged 22–24, and the uninvolved flexor forearm of a 30-yr-old woman with mild atopic dermatitis. The sweat gland isolation, incubation, and lactate and ¹⁴CO₂ determination were the same as previously described. KRB was used throughout. Since sweat gland size differed markedly from subject to subject in humans, sweat glands were recollected after ¹⁴CO₂ trapping and homogenized, and the mean protein content per gland was measured for each individual by the method of Lowry et al. (8).

RESULTS

Glucose metabolism of the isolated monkey palm eccrine sweat gland

The effect of glucose-free medium on lactate production. The eccrine sweat gland contains glycogen, which shows a variable decrease after sweat secretion (9). CO₂ production as determined in the present study would be underestimated if endogenous (cellular) glycogen or glucose or both provide a considerable portion of the total energy. Since lactate and ¹⁴CO₂ production are quantitatively parallel under aerobic conditions (1), only the former was compared in glucose-free and glucose-containing media to estimate the role of endogenous glycogen or glucose or both as energy sources. As

³ Sato, K. Unpublished observations.

shown in Table I, lactate production in the glucose-free medium was considerably lower than in the glucose-containing medium after pharmacologic stimulation. In the glucose-free medium, Mecholyl or epinephrine failed to stimulate lactate production. After Mecholyl stimulation, lactate production in glucose-free medium was only 3% of that in glucose-containing medium.

The differential response of the duct and the secretory coil to Mecholyl and epinephrine. Fig. 1 shows lactate and $^{14}\text{CO}_2$ production by the coiled duct and the secretory coil. Stimulation of lactate and $^{14}\text{CO}_2$ production by Mecholyl or epinephrine occurs only in the secretory coil. Glucose metabolism by the duct was low, unresponsive to Mecholyl or epinephrine, and equal to or slightly lower than in the unstimulated secretory coil.

The effect of Na^+ and K^+ deprivation on lactate production. Both sweat secretion and sodium reabsorption are dependent on the energy-requiring, membrane ($\text{Na}^+ + \text{K}^+$)-ATPase system (3, 4, 10, 11). Both Na^+ and K^+ must be present for the activation of ($\text{Na}^+ + \text{K}^+$)-ATPase (3, 10). As shown in Fig. 2, the omission of either Na^+ or K^+ or both produced a lack of activation by Mecholyl but considerable activation by epinephrine. In the 40 mM $\text{Na}^+ + 100$ mM K^+ medium, lactate production was slightly activated by Mecholyl.

The effect of a 40 mM $\text{Na}^+ + 100$ mM K^+ medium on lactate production. Since the assumed intracellular Na^+ and K^+ concentration of the salivary gland is 40 and

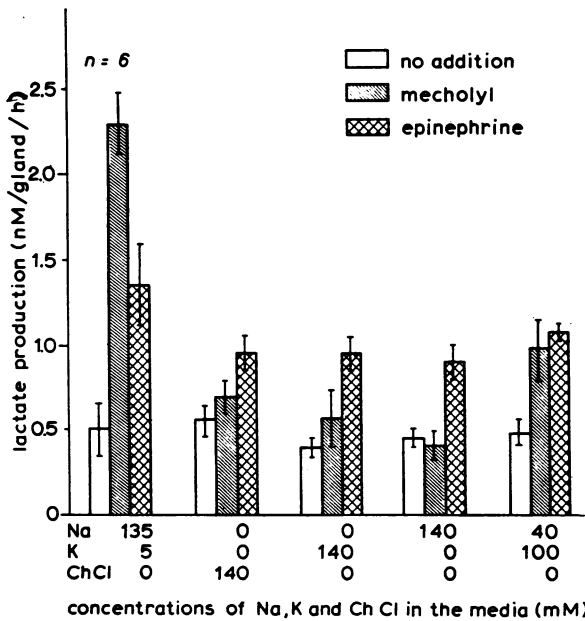


FIGURE 2 The effect of cations on Mecholyl- or epinephrine-stimulated lactate production. Lactate and $^{14}\text{CO}_2$ production are expressed as nanomoles per gland per hour in this and following figures. ChCl, choline chloride.

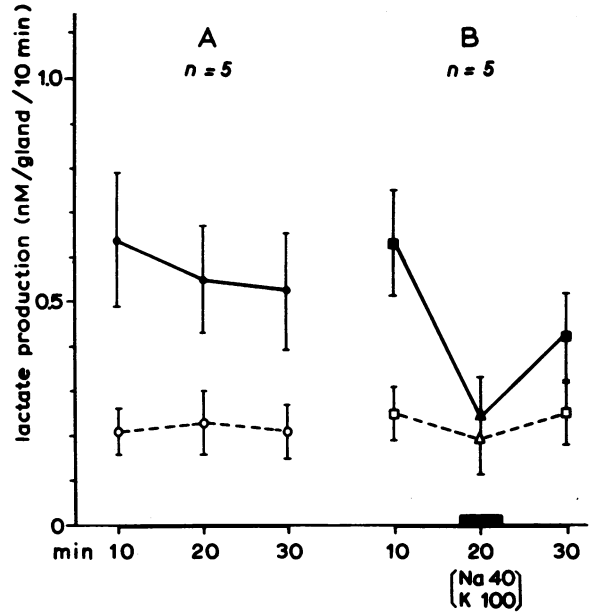


FIGURE 3 The effect of 40 mM $\text{Na}^+ + 100$ mM K^+ medium on lactate production. Open symbols (\circ , \square , and \triangle) signify unstimulated sweat glands, and closed symbols (\bullet , \blacksquare , and \blacktriangle) Mecholyl-stimulated sweat glands. The horizontal bar on the abscissa (B, 20 min) signifies the 40 mM $\text{Na}^+ + 100$ mM K^+ medium. A is the control system, where KT medium was used throughout the three 10-min incubation periods.

100 mM, respectively (12, 13), the effects of these electrolyte concentrations on lactate production were further studied. In this medium, the stimulating effect of Mecholyl on lactate production was markedly suppressed (Fig. 3B, 20 min) and was partially restored when the tissue was subsequently transferred to the complete KT medium.

The effect of DNP on the metabolic inhibition induced by ouabain or Na^+ and K^+ deprivation. Both ouabain-containing and Na^+ - and K^+ -free media inhibit ($\text{Na}^+ + \text{K}^+$)-ATPase activity (3) and Mecholyl- or epinephrine-stimulated glucose metabolism in isolated eccrine sweat glands (1). If the inhibition of energy metabolism by ouabain or Na^+ and K^+ deprivation is due solely to their effects on active sodium transport and not to damage to the metabolic machinery secondary to an altered cellular homeostasis, then DNP, an uncoupler of oxidative phosphorylation, should reverse the metabolic inhibition thus incurred. Fig. 4 shows the effect of 10^{-4} M DNP on Mecholyl- or epinephrine-stimulated lactate and $^{14}\text{CO}_2$ production. DNP stimulated both lactate and $^{14}\text{CO}_2$ production of unstimulated sweat glands. The addition of Mecholyl to DNP further stimulated both $^{14}\text{CO}_2$ and lactate production. As illustrated in Fig. 5, 10^{-8} M ouabain alone strongly inhibited both unstimu-

TABLE II
Effects of Incubation Media on Lactate and CO₂ Production

	No drug addition		Mecholyl (4.2 × 10 ⁻⁶ M)		Epinephrine (5.5 × 10 ⁻⁶ M)	
	KRB*	KT†	KRB	KT	KRB	KT
Lactate nmol/gland/h	0.45±0.10	0.63±0.12	1.77±0.09	1.71±0.20	1.06±0.13	1.33±0.14
	<i>P</i> > 0.2		<i>P</i> > 0.7		0.1 < <i>P</i> < 0.2	
CO ₂ nmol/gland/h	0.36±0.07	0.49±0.09	1.85±0.14	1.53±0.16	0.94±0.16	1.33±0.08
	<i>P</i> > 0.3		0.10 < <i>P</i> < 0.15		0.05 < <i>P</i> < 0.10	

The results on lactate and ¹⁴CO₂ production are the mean ±SEM of five experiments.

* Krebs-Ringer bicarbonate.

† Krebs-Tris buffer.

lated and Mecholyl stimulated lactate and ¹⁴CO₂ production below the baseline level (control "none" in Fig. 5). Epinephrine-stimulated lactate production was slightly higher than the baseline level but ¹⁴CO₂ production was below the control level. The addition of DNP reversed the inhibitory effect of ouabain to approximately 80% of the Mecholyl-stimulated lactate and ¹⁴CO₂ production ("control-MCH" in Fig. 5). The removal of Na⁺ and K⁺ from the media resembled the effect of ouabain (Fig. 6). However, the addition of DNP to Na⁺- and K⁺-free media enhanced both lactate and ¹⁴CO₂ production to a much smaller extent than in the ouabain-containing complete media.

The effects of buffer and Diamox on lactate and ¹⁴CO₂ production. The KT buffer used in this and previous studies (1, 14) lacks HCO₃⁻ and H₂PO₄⁻. In some tis-

ues, the lack of HCO₃⁻ and the addition of Diamox lead to a marked decrease in cellular metabolism and function (15). Table II lists lactate and ¹⁴CO₂ production in KRB and KT media. ¹⁴CO₂ production in the presence of Mecholyl tended to be lower in KT than in KRB, whereas epinephrine-stimulated lactate and ¹⁴CO₂ production tended to be higher in KT than in KRB, although these were statistically not significant. As shown in Table III a slight inhibitory effect of Diamox on Mecholyl-stimulated ¹⁴CO₂ production was seen only at the highest concentration of Diamox (10⁻² M).

The synergistic effect of Mecholyl and epinephrine on lactate and ¹⁴CO₂ production. Kuno (16) claimed that in man epinephrine potentiates the sudorific action of acetylcholine, whereas Foster Ginsburg, and Weiner (17) could not confirm this. If the sites or mechanisms of action of these drugs differ, some synergistic effect might be expected when both drugs are used simultaneously. As tabulated in Table IV, the addition of epinephrine to Mecholyl failed to enhance Mecholyl-stimulated lactate and ¹⁴CO₂ production.

Lactate and ¹⁴CO₂ production of isolated human eccrine sweat glands

In this and in a previous communication (1), the monkey palm eccrine sweat gland has been used as a model of the human eccrine sweat gland. Wolfe et al. (14) have claimed that aerobic metabolism accounts for less than 1% of the total energy production in human eccrine sweat glands. Human as well as monkey sweat glands are well endowed with mitochondria (18), and arterial occlusion causes a sharp decrease in sweat rate and an increase in sweat Na⁺, K⁺, and lactate concentration in human (19). These observations suggest that aerobic metabolism must also play an important role in human eccrine sweat glands. Table V lists lactate and ¹⁴CO₂ production of isolated human sweat glands in four

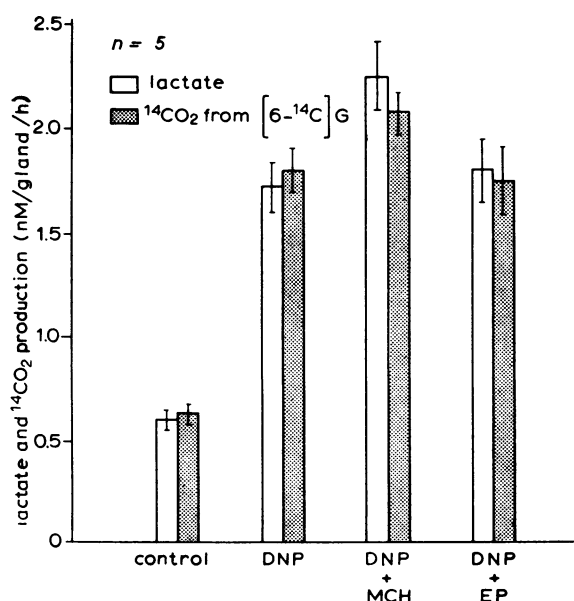


FIGURE 4 The effect of 10⁻⁴ M DNP on lactate and ¹⁴CO₂ production. Control means no addition of drugs.

⁴ Sato, K. Unpublished observations.

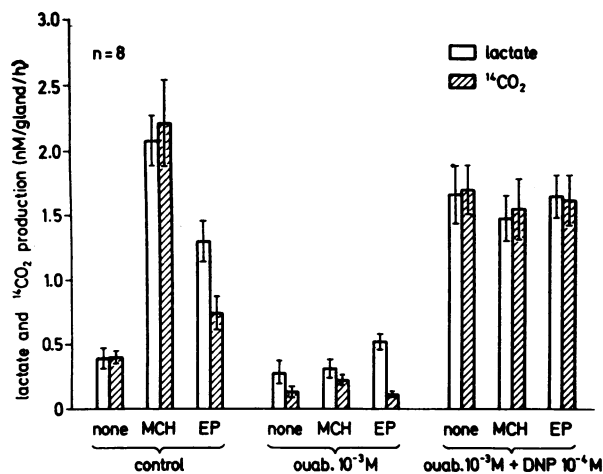


FIGURE 5 The effect of 2,4-DNP on the ouabain-induced inhibition of lactate and $^{14}\text{CO}_2$ production. The concentrations used: ouabain (ouab.) 10^{-3} M; DNP, 10^{-4} M.

subjects. In contrast to monkey sweat glands, in which molar production rates of lactate and $^{14}\text{CO}_2$ are approximately equal, human sweat glands yielded more lactate (from 1.9 to 3.4 nmol/gland/h) and less $^{14}\text{CO}_2$ (from 0.42 to 0.72 nmol/gland/h). The mean protein content per gland, and thus their sizes, varied widely (from 1.1 to 2.8 $\mu\text{g/gland}$) but the mean of four subjects, 2.3 $\mu\text{g/gland}$, was comparable to that of the monkey sweat gland, 2 μg (3).

DISCUSSION

The whole sweat gland coil (secretory coil plus proximal duct) used to a great extent in the present study consists of five different cell types: clear or secretory, dark, myoepithelial and ductal cells. The metabolic response measured in the whole sweat gland is thus the sum of the metabolic activities of all these cell types. We have assumed, however, that the major portion of the total energy production is related to the secretory and ductal cells since these cells are primarily involved in sodium transport. Furthermore, Mecholyl-induced metabolic stimulation can account for the net sodium flux across the secretory and ductal cells if we assume a Na^+/ATP ratio of three (3). Eccrine sweat glands are predominantly cholinergic organs, and sweating can be induced by either endogenous or exogenous cholinergic drugs. Although epinephrine and norepinephrine induce sweating (20–23), their physiological and functional significance are as yet enigmatic. The present study has shown that only the secretory coils, innervated by acetylcholinesterase-positive nerves (18, 24), respond to Mecholyl and epinephrine *in vitro* by increases in lactate and $^{14}\text{CO}_2$ production and that the ducts, which are not innervated, do not. The effect of

TABLE III
The Effect of Varying Concentrations of Diamox on Lactate and CO_2 Production

Addition		n	Production	
Mecholyl	Diamox		Lactate	CO_2
			nmol/gland/h	nmol/gland/h
None	None	5	0.31 ± 0.02	0.62 ± 0.09
10^{-6} M	None	5	1.50 ± 0.16	$1.94 \pm 0.10^*$
10^{-6} M	10^{-2} M	4	1.60 ± 0.12	$1.55 \pm 0.06^*$
10^{-6} M	10^{-3} M	4	1.39 ± 0.18	1.71 ± 0.21
10^{-6} M	10^{-5} M	4	1.57 ± 0.16	2.05 ± 0.14

n, number of experiments.

* P value: $0.05 < P < 0.10$.

these drugs on cellular metabolism is unlikely to be nonspecific, i.e., unrelated to the cholinergic or adrenergic mechanisms of sweat secretion or due to the direct stimulation of some steps in a metabolic pathway in the duct since it fails to respond to these drugs. A more likely explanation would be that, like the acinar cell of the salivary gland (25), the contraluminal membrane of the secretory cell responds to Mecholyl and perhaps also to epinephrine by an increase in its permeability to external sodium (step 1 in Fig. 7), which in turn stimulates the sodium pump at the luminal (including intercellular canaliculi) side of the membrane (step 2). The stimulation of energy metabolism would then follow (step 3) (26). This assumption is supported by the observations that $(\text{Na}^+ + \text{K}^+)\text{-ATPase}$ activity is stimulated exponentially by Na^+ (3), and that lactate production is markedly suppressed when the sweat glands are transferred to the 40 mM $\text{Na}^+ + 100$ mM K^+ medium containing Mecholyl (Fig. 3). Under these circumstances, the concentrations of sodium and potassium

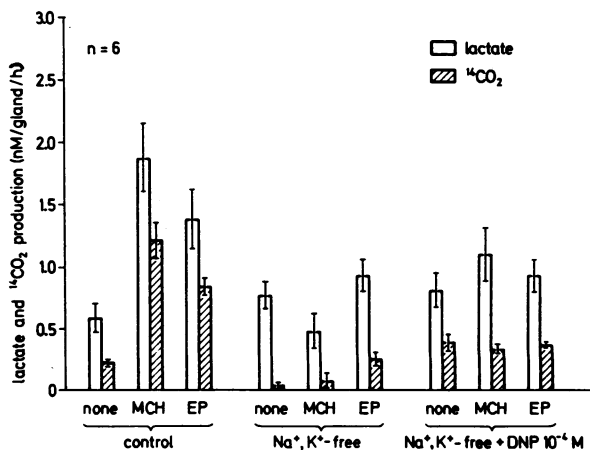


FIGURE 6 The effect of DNP on the metabolic inhibition induced by Na^+ and K^+ deprivation.

TABLE IV
The Effect of Epinephrine on Mecholyt-Stimulated Lactate and CO₂ Production

	No drug addition	Mecholyt (4.2 × 10 ⁻⁶ M)	Epinephrine (5.5 × 10 ⁻⁶ M)	Mecholyt (4.2 × 10 ⁻⁶ M) + Epinephrine (5.5 × 10 ⁻⁶ M)
Lactate nmol/gland/h	0.68 ± 0.07	2.24 ± 0.26*	1.31 ± 0.18	2.20 ± 0.15*
¹⁴ CO ₂ nmol/gland/h	0.31 ± 0.08	2.39 ± 0.39‡	0.79 ± 0.11	2.25 ± 0.22‡

The mean ± SEM of eight experiments.

* *P* > 0.8.

‡ *P* > 0.7.

are approximately equal inside and outside of the cell, so that Mecholyt-induced alterations in membrane permeability produce little or no net sodium influx (step 1), leading to much lower activation of steps 2 and 3. The absence of a synergistic effect between Mecholyt and epinephrine on lactate and ¹⁴CO₂ production suggests that both drugs act at the same site, probably at step 1, since neither Mecholyt nor epinephrine are stimulants of (Na⁺ + K⁺)-ATPase.⁵ However, the failure to completely abolish the epinephrine-stimulated lactate production by the highest concentration of ouabain (10⁻³ M) (1), by nitrogenated media (1), or by Na⁺- and K⁺-free media suggests that at least part of the epinephrine-induced metabolic stimulation is independent of the linkage between steps 2 and 3. This action of epinephrine cannot be explained by its effect on the adenylcyclase-cyclic AMP system since neither dibutyryl cyclic AMP nor theophylline nor a combination of both enhances lactate or ¹⁴CO₂ production (from [6-¹⁴C]glucose) (7). As shown in Figs. 4-6, DNP, an uncoupler of oxidative phosphorylation (27-29), stimulated lactate and ¹⁴CO₂ production almost as effectively as Mecholyt, and also

reversed the ouabain-induced inhibition of glucose metabolism. These effects of DNP are to be expected since uncoupling of oxidation from phosphorylation produces high cellular ADP levels (since ATP is not formed or is quickly hydrolyzed to ADP even if formed), which in turn enhance glycolysis (30) as well as oxidative phosphorylation (31). Furthermore, the reversal of ouabain-induced inhibition of glucose metabolism by DNP points to the specificity of the effect of ouabain on sodium transport and the tightness of the coupling between glucose oxidation (Fig. 7, step 3) and ATP utilization for sodium transport (step 2). If the metabolic inhibition by ouabain simply represents damage to the metabolic machinery caused either directly by ouabain or by alteration of cellular electrolyte balance (increase of Na⁺ and decrease of K⁺), then the stimulation of glucose metabolism in the presence of both ouabain and DNP is unexplained. The in vivo effects of ouabain on both sweat secretion and Na⁺ reabsorption have been previously studied (11). The action of ouabain on sweat gland sodium transport (11) and glucose metabolism supports our earlier postulates that both sweat secretion by the secretory cells and Na⁺ reabsorption by the ductal cells are mediated by a (Na⁺ + K⁺)-ATPase-depend-

⁵ Sato, K. Unpublished observations.

TABLE V
Lactate and CO₂ Production of Isolated Human Forearm Eccrine Sweat Glands

Sub.	Age	Sex	Protein*	Lactate			CO ₂		
				C	MCH	EP	C	MCH	EP
				nmol/gland/h			nmol/gland/h		
			μg/gland						
1	30	F	2.4	0.72	2.86	1.30	0	0.47	0.14
2	22	M	1.1	0.11	1.90	1.30	0	0.42	0.31
3	22	M	2.8	0.92	3.40	3.10	0	0.72	0.46
4	24	M	2.7	0.80	3.40	2.60	0	0.60	0.43
Mean			2.3	0.66	2.90	2.08	0	0.55	0.34

C, control (unstimulated); MCH, Mecholyt (4.2 × 10⁻⁴ M); EP, epinephrine (5.5 × 10⁻⁵ M).

* Mean protein content per gland. Experiments were performed in March and May 1971 in the Netherlands.

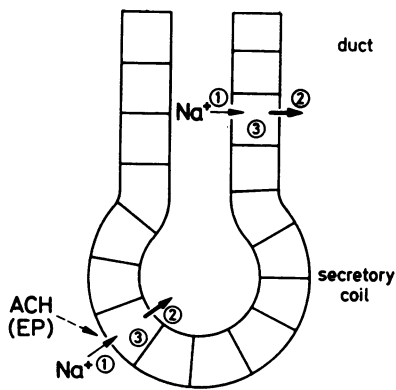


FIGURE 7 A schematic drawing of the three major steps in sodium secretion by the secretory coil and sodium reabsorption by the duct. Step 1, passive entry of Na^+ ; step 2, sodium pump ($[\text{Na} + \text{K}]\text{-ATPase}$); step 3, energy metabolism; ACH, acetylcholine; EP, epinephrine.

dent Na^+ pump (3, 4). The reversal by DNP of the metabolic inhibition caused by Na^+ and K^+ deprivation was only partial. This may indicate that Na^+ and K^+ deprivation caused irreversible damage to the cellular metabolic machinery or that uncoupling by DNP was incomplete, since DNP-sensitive, mitochondrial ATPase requires K^+ for its maximal activity (32). A third possibility is that the glucose entry into the cell is coupled to Na^+ influx and is thus inhibited in a Na^+ - and K^+ -free medium. However, this seems unlikely to be the cause of the low reversal by DNP of metabolic inhibition in the Na^+ - + K^+ -free media, since in the absence of Na^+ transport, as in the case after the addition of DNP to the unstimulated sweat gland (Fig. 6), sufficient glucose was available in the cell to stimulate glucose metabolism maximally. The failure of Diamox or a bicarbonate-free medium to significantly reduce sweat gland glucose metabolism suggests that hydrogen-ion transport does not play a major role at all stages of sodium transport.

In disagreement with the results of Wolfe et al (14), the present study showed considerable $^{14}\text{CO}_2$ production by human forearm sweat glands although the mean $^{14}\text{CO}_2$ -to-lactate ratio after Mecholyl stimulation of 0.21 is only one-fifth that of monkey palm sweat glands (1). If we assuming 6 mol CO_2 is equivalent to 38 mol ATP, and 1 mol lactate to 1 mol ATP, human forearm sweat glands produce a mean ATP of 3.5 nmol/gland/h by oxidation and 2.9 nmol/gland/h by glycolysis. The higher activity of the oxidative pathway in monkey palm sweat glands may be due partly to a species difference and partly to the fact that sweating on the palm in the monkey occurs almost continuously (33) whereas the thermally-responsive sweat glands of man are relatively inactive in a temperate climate during the months of March to May when these studies were performed.

The disagreement between our results and those of Wolfe et al. (14) can be partly explained since Wolfe et al. incubated sweat glands in a glucose-free medium to measure O_2 consumption whereas we used a glucose-containing medium.

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