Abstract. Furosemide inhibits 3-O-methyl-D-glucose equilibrium flux in isolated adipocytes. The inhibition is saturable with an increasing concentration of furosemide and shows a noncompetitive type of kinetics. Both basal and insulin-stimulated fluxes are equally affected by the inhibition. Hydrochlorothiazide and piretanide also inhibit the flux with a similar potency, whereas bumetanide, a more potent diuretic, is much less potent. To understand the molecular basis of this inhibition, effects of furosemide on the glucose-sensitive cytochalasin B binding activities of adipocytes were studied. Furosemide inhibits the glucose-sensitive cytochalasin B binding of both microsomal and plasma membrane preparations. For both preparations, the inhibition is time dependent and only slowly reversible, is saturable with an increasing concentration of furosemide, shows a noncompetitive type of kinetics with apparent $K_i$ (the inhibitor concentration that gives the half-maximum effect) of 3.5 and 0.7 mM after 2 and 18 h incubation, respectively, and is essentially identical between the basal and insulin-stimulated adipocytes. The inhibition develops with a first-order rate constant of approximately 0.12/h at 4°C. These results indicate that furosemide inhibits glucose transport in adipocytes by directly inactivating transport carriers of both plasma membranes and microsomal reserve pool. This inactivation of glucose carrier may play a part in the diuretic-induced glucose intolerance frequently observed during diuretic therapy.

Introduction

Significant glucose intolerance often develops in patients receiving prolonged treatment with diuretic agents, e.g., furosemide, thiazide, etc. (1–3). The exact mechanisms operative in this effect remain unclear and controversial. The postulated mechanisms fall into two broad classes: (a) due to alterations in pancreatic insulin secretion (4), and (b) due to extrapancreatic effects independent of insulin secretion, such as decreased glucose utilization or transport, and/or decreased insulin sensitivity at the peripheral tissue level (5, 6).

Alterations in carbohydrate economy conceivably may be due to effects of the diuretic molecule at any of the specific loci that govern such economy. Depletion of total body potassium stores is known to cause glucose intolerance by reducing pancreatic beta cell responsiveness to glucose (7–9). More recently, Helderman et al. (10) performed glucose clamp studies on normal volunteers and concluded that thiazide-induced glucose intolerance is a result of potassium depletion, which reduces insulin output by the above mechanism.

However, there is also a body of experimental and clinical evidence that suggests that mechanisms independent of pancreatic insulin output or potassium balance may contribute to glucose intolerance caused by diuretics, especially of the benzenothiadiazine class (6, 11). We have previously shown that furosemide inhibits glucose transport directly in human erythrocytes (12). We now extend our observations to an insulin-sensitive tissue, e.g., adipocytes, to show that furosemide inhibits basal and insulin-stimulated glucose transport by directly interacting with the glucose carrier molecule. We suggest that this direct inactivation of glucose carrier may contribute significantly in the glucose intolerance observed during diuretic therapy.

Methods

Materials. Collagenase was purchased from Worthington Biochemical Corp. (Freehold, NJ). 3-O-methyl-D-glucose (3-OMG), L-glucose, phloretin, furosemide, hydrochlorothiazide, and cytochalasin B were purchased from Sigma Chemical Co. (St. Louis, MO). [3H]Cytochalasin B, 3-O-methyl [3H]-D-glucose were obtained from New England Nuclear (Boston, MA). Bovine albumin powder (CRG-7) was obtained from Armour Pharmaceuticals (Kankakee, IL). Piretanide was a generous gift from Mr. Val Wagner, Hoechst Roussel Pharmaceutical (North Somerville, NJ), and Bumetanide, a generous gift of Dr. P. Sorter, 1984 and in revised form 12 July 1984.
Hoffman LaRoche, Inc. (Nutley, NJ). Insulin (porcine crystalline) was from Eli Lilly & Co. (Indianapolis, IN). All other chemicals were standard commercial products of reagent grade quality.

Isolation of adipocytes. Adipocytes were isolated from the epididymal fat pads of Sprague Dawley (150–200 g) male rats given free access to lab chow. Fat cells were prepared by the method described by Rodebell (13) and modified as follows. Minced fat pads were put in several 25-ml polyethylene scintillation vials with 15 ml of Krebs-Ringer bicarbonate buffer containing collagenase (1.5 mg/ml), bovine serum albumin (40 mg/ml), and d-glucose (3 mM) as described (14). The vials were then gassed briefly (10–20 s) with a 95% O₂/5% CO₂ mixture, sealed, and incubated with gentle shaking at 37°C for 1 h, at the end of which time the suspension was filtered through a nylon mesh to remove all extraneous tissue. The fat cells were then washed four times in a Krebs-Ringer buffer containing bovine serum albumin (40 mg/ml) and d-glucose (3 mM) (this buffer is referred to as incubation buffer) by centrifugation (900 g for 1 min, which gives 80–90% hemocrit).

Preparation of 150,000 g total particulate fraction. Isolated fat cells were hypotonically lysed by vortexing the cells for 60 s in a hypotonic buffer containing MgCl₂ (2.5 mM), CaCl₂ (0.1 mM), and KHCO₃ (1 mM) adjusted to pH 7.2. The suspension was then centrifuged (1,000 g for 1 min), the resulting supernatant (containing fat cake and unlysed cells) was separated from lysate and the lysates were pooled and stored on ice. This procedure was repeated with subsequent supernatants four times until only predominantly the fat cake remained, which was discarded. The lysates were spun down in a Beckman ultracentrifuge (Beckman Instruments, Inc., Fullerton, CA) at 150,000 g for 60 min at 5°C. The 150,000-g pellet was then resuspended in the hypotonic buffer referred to above and stored in liquid nitrogen until used.

Preparation of plasma membrane and microsome fractions. Plasma and microsomal membrane fractions were prepared by the differential centrifugation method described by McKeel and Jarett (15) and modified by Cushman and Wardzala (16). Using a medium (medium 1) composed of 10 mM Tris HCl, pH 7.4, EDTA (1 mM), and sucrose (0.25 M). Where insulin and/or furosemide-treated cells were used furosemide and/or insulin was also added to medium 1. The suspension was then transferred in 30-ml aliquots to a glass homogenizer (Arthur H. Thomas Co., Philadelphia, PA) with the pestle rotating at 2,000 rpm. During the homogenization the homogenate was kept in ice. The homogenate was then centrifuged at 16,000 g for 15 min. The fat cake was discarded. To isolate the microsomal membrane, the 16,000 g supernatant was spun for 90 min at 200,000 g, and its pellet was resuspended in the hypotonic buffer and stored in liquid nitrogen.

To isolate plasma membranes, the 16,000 g pellet of homogenized cells was resuspended in medium 1 and centrifuged for 10 min at 1,000 g. The subsequent pellet was discarded and the supernatant was centrifuged at 17,000 g for 20 min. The pellet was resuspended in 2 ml of medium 1 and layered on a 15-ml of medium (medium 11) composed of sucrose (1.12 M), Tris (20 mM), and EDTA (1 mM), pH 7.4, and centrifuged at 110,000 g for 60 min following the method of Cushman (17). This procedure gave a single band of plasma membrane that was visible on the upper portion of medium 11. This portion was removed and suspended in 20 times the volume of medium 1 and centrifuged for 20 min at 17,000 g. The pellet was resuspended in the hypotonic buffer and stored in liquid nitrogen.

Insulin treatment. Immediately following isolation (for cytochalasin B binding studies) or following incubation with 3-OMG (for flux measurements, see below) 12-ml aliquots of adipose cells (40–50% cytocr) were distributed in 25-ml polyethylene scintillation vials containing 10 ml of incubation media and enough insulin to achieve final concentrations at either 0 or 7.0 nM. When specified, furosemide was also added to this stage. The cells were then incubated for 45 min at 37°C with gentle shaking.

Furosemide treatment. Furosemide stock solutions were prepared in 75% methanol in water adjusted to pH 10.0 with NaOH. Furosemide treatments were started by adding an appropriate stock solution to the cell or membrane suspension to give a specified final concentration without exceeding methanol content of 1%, and the mixture was incubated for a given time. Controls contained the same amount of methanol at the same pH. For cytochalasin B binding to membrane preparations, the furosemide treatments were done either with intact cells prior to their membrane isolation (pretreatment) or with isolated membranes (posttreatment). For the binding studies, the temperature during furosemide treatment was 37°C (pretreatment) or 4°C (posttreatment).

For a prolonged incubation with furosemide, an adipose tissue culture method was used. Under sterile conditions isolated fat pads were gently rinsed, cut in small pieces, and placed in 100 × 20-mm tissue culture dishes (3 g/dish) containing Parker's medium 199 following the method of Maloff and Lockwood (18). When used enough furosemide to give a final concentration of 0.1 mM was added to the incubation medium. The dishes were then incubated at 37°C for 48 h under an atmosphere of 95% O₂/5% CO₂ with the media changed after 24 h.

Measurement of 3-OMG flux. Glucose transport was assessed by measuring equilibrium exchange rates of 3-OMG using tritium-labeled 3-OMG as a tracer and a modification of the oil flotation method of Whitesell and Gliemann (19). Isolated cells were incubated with a given concentration of 3-OMG in a Krebs-Ringer buffer (10% cytocr) for 30 min at room temperature. The cell suspension was then briefly centrifuged (30 s at 500 g) to reduce its volume to a 40–50% cytocr. Insulin or furosemide when used, and 3-OMG solution (for controls) were added this stage to given final concentrations and hemocrits of 22–27% (see above). 200-μl aliquots of this cell suspension were transferred into centrifuge tubes (12 × 75-mm polypropylene), and then, incubated without shaking for 45 min at 37°C. The flux measurements were initiated by injecting 10 μl of 0.9 μCi of labeled 3-OMG directly into the cell suspension with an automatic pipette and vortexed gently for 3 s. The flux was terminated by the addition of 3.5 ml of ice cold Krebs-Ringer bicarbonate buffer containing 0.3 mM of phloretin and 0.15% vol/vol of ethanol. 1 ml of ice-cold silicone oil (relative density, 0.99; viscosity, 100 centistokes) was layered on top of the mixture and the tubes were centrifuged immediately for 1 min at 2,500 g. The packed cells on top were then removed with a pipe cleaner and placed in counting vials with 10 ml of scintillation fluid (Hydrofluor, National Diagnostics, Somerville, NJ) vortexed vigorously and their radioactivities were counted. Extracellular trapped radioactivity was measured by adding phloretin solution to cells before injection of isotopic 3-OMG.

Cytochalasin B binding assays. Cytochalasin B binding assay was carried out by a centrifugation method that was described from our laboratory (20) with some modification. Binding assay mixture contained membrane preparation (40–80 μg protein, as measured by Pierce reagent [21] using the Pierce standard) in the hypotonic buffer described previously, 0.01 μCi [3H]cytochalasin B with a given concentration of unlabeled cytochalasin B, and 500 mM L- or D-glucose, made up to a final volume of 200 μl. When so specified (posttreatment) furosemide was also added at this stage. The mixtures were incubated in 230 μl
cellulose propionate centrifuge tubes for 30 min at 4°C, then centrifuged at 200,000 g for 60 min at 4°C. After centrifugation, 50 μl of supernatant was removed for measurement of the free [3H]cytochalasin B. The remaining supernatant was aspirated completely and inner wall of tubes were scrubbed with cotton swabs to remove any residual medium. Pellets in centrifuge tubes were transferred directly to a 5-ml scintillation vial containing 4 ml of scintillation fluid (Hydroluor) for measurement of bound [3H]cytochalasin B. Radioactivities of both pellets (previously solubilized with 2% sodium dodecyl sulfate for 12 h) and supernatants were counted in a liquid scintillation spectrophotometer (Nuclear Chicago, Mark II, Chicago, IL). Drug treatments did not affect protein recovery in pellets under the condition used.

**Results**

Rate of equilibrium exchange of 3-OMG by isolated adipocytes of rat epididymal pads measured at 37°C as a function of the sugar concentration showed a simple, saturation kinetics with a Michaelis constant (Kₘ) of 5-7 mM and ϕ_max (the maximal rate) of ~0.27 μmol/s per ml cell water at 37°C (Fig. 1). The same batch of isolated adipocytes that were treated with insulin (7 nM) for 30 min at 37°C prior to the flux measurement (insulin-stimulated state) showed approximately a threefold high ϕ_max value compared with basal state without significant change in Kₘ value (Fig. 1).

These isolated adipocytes were incubated with varying concentrations of furosemide for 45 min at 37°C and the effects of this incubation on the equilibrium exchange flux of 3-OMG were studied (Figs. 1, 2, and 3). For each 3-OMG concentration used, furosemide inhibited the flux significantly (Fig. 2). Furosemide treatments also reduced the 3-OMG space of adipocytes significantly and this effect is apparently dose-dependent (Fig. 2). This is most likely due to cell shrinkage (23) or cell lysis. Nevertheless, analyses of the flux data using the sugar space observed (Sₛ) for each furosemide concentration shows that the inhibition increases in a simple, saturable manner as the concentration of furosemide in the incubation was increased (Fig. 3 A). The furosemide concentration that reduced the flux rate by 50% (apparent inhibition constant, Kᵢ) is ~4-5 mM irrespective of the three 3-OMG concentrations used (Fig. 3 A). When the same data were plotted as a function of 3-OMG for each furosemide concentration (Fig. 1), it is evident that the inhibition is due to a reduction in ϕ_max without any significant change in the Kₘ value, indicating a noncompetitive mode of inhibition.

The 3-OMG fluxes of insulin-stimulated adipocytes were also inhibited by furosemide (Figs. 1 and 3 B). The pattern of

![Figure 1](image1.png)

**Figure 1.** Rates of equilibrium exchange of 3-OMG by isolated adipocytes as a function of 3-OMG concentration. The half-time of equilibration (t₁/₂) was estimated from the time course of the equilibrium analyzed as a first order process. See Figs. 2 and 3 for details. Data are plotted according to a linear transformation of Michaelis-Menten equation:

\[ C/\phi = [K_m + C]/\phi_{max}, \]

and the relationship (21):

\[ t_{1/2} = 0.693 \cdot \frac{V_{ir} C}{(V_i + V_o) \cdot \phi}, \]

where C, ϕ, Kₘ, ϕ_max, Vᵢ, and Vₒ denote 3-OMG concentration, flux rate, Michaelis-Menten constant, the maximal flux rate, intracellular water volume, and extracellular water volume, respectively. Furosemide concentrations used are none (●), 1 (○), 3 (■), and 6 (▲) mM. Straight lines are drawn by eye to best fit to the data points, obtained from noninsulin-treated cells (broken lines) and insulin-treated cells (solid lines).

![Figure 2](image2.png)

**Figure 2.** Time course of the equilibrium influx of 3-OMG by isolated adipocytes without (○) and with 1 (●), 3 (■), and 6 (▲) mM furosemide treatments, respectively. Cells were incubated with 20 mM 3-OMG before the isotopic uptake measurement as described in Methods. Data were analyzed based on a simple model of tracer exchange in a closed system of two compartments in inset according to the equation (22):

\[ \log \frac{S_a - S_b}{S_a} = \frac{-1}{2.3} \cdot \frac{V_i C C}{V_o \cdot \phi}, \]

where Sₐ, Sₒ, and Sₐ denote cellular radioactivities at a given time, time of complete equilibration (15 min), time of start of experiment, respectively, and t denotes time. The time course of the tracer equilibrium exchange in this plot fits a reasonably good straight line, from which t₁/₂ (the time required for one half of the complete equilibration, in seconds) can be estimated for each 3-OMG concentration.
transport glucose between plasma and its membrane vs. microsomes)

Figure 3. Inhibition of equilibrium exchange of 3-OMG by isolated adipocytes as a function of furosemide concentration. Replot of the data of Fig. 1 according to the equation (24):

\[
\frac{C}{\phi} = \frac{1}{\phi_{\text{max}}} \left[ K_n + C \right] \left[ 1 + \frac{I}{K_i} \right],
\]

where I and K_i are inhibitor concentrations and inhibition constants, respectively, and the relationship of t_{1/2} to C/\phi (21) as shown in the legend of Fig. 1. The three different concentrations of 3-OMG used were 1 (○), 5 (△), and 20 mM (●) for the basal (left) and insulin-stimulated (right) adipocytes.

the inhibition was essentially identical to that of the basal state showing noncompetitive inhibition and an apparent K_i value of \( \sim 4.5 \text{ mM} \).

Piretanide and hydrochlorothiazide also inhibited 3-OMG equilibrium flux in a similar manner as furosemide, with a K_i value of \( \sim 4 \text{ mM} \) (data not shown). Bumetanide, however, did not show any significant inhibition at the concentrations up to 3 mM (solubility limit).

Cytochalasin B, a well-established specific inhibitor of glucose transport of adipocytes, binds to purified microsomal and plasma membranes of adipocytes (16, 17). Scatchard analyses of these bindings indicated that there are at least two saturable binding components at the ligand concentration range of up to \( 10^{-5} \text{ M} \). The ligand binding to one of these components is displaced specifically by D-glucose, but not by L-glucose (16). The difference in the ligand bindings observed between these two isomers are taken as the glucose-sensitive binding (16, 20).

Effects of furosemide on this glucose-sensitive, cytochalasin B binding to microsomal and plasma membrane preparations were studied by assaying the ligand binding in the presence of 500 mM L- and D-glucose. In a series of experiments (pretreatment series), cells were first incubated with a given concentration of furosemide prior to their membrane subfractionation. In other experiments (posttreatment series), the membranes were prepared first, then incubated in the presence of furosemide and its effect on the ligand binding was assessed. Both of the furosemide treatments inhibited the cytochalasin B binding of both the microsomal and plasma membranes equally, regardless of whether in basal or in insulin-stimulated states. This is illustrated in Fig. 4 with microsomal membrane preparation of insulin-stimulated adipocytes.

The effects of furosemide on the glucose-sensitive cytochalasin B binding activities of these membrane preparations at both basal and insulin-stimulated states were further analyzed by Scatchard plot. In each case, when the incubation time with furosemide is fixed, B_T (total binding capacity) of the glucose-sensitive binding was increasingly reduced as the furosemide concentration was raised in the incubation without affecting affinity (or K_D) of the binding significantly. This is illustrated in Fig. 5 using plasma membranes prepared from basal state of adipocytes as an example, and the results with all preparations are summarized in Table I. Pretreatment appears slightly more effective than posttreatment throughout all preparations. This may be simply due to the additional exposure to the drug during membrane preparation involved in the pretreatment experiments. Data presented in Table I do not support the contention that furosemide affects insulin-induced redistribution of cytochalasin B binding activities between plasma membrane vs. microsomes significantly.

This reduction of the B_T analyzed as a function of furosemide concentration for a fixed incubation time indicates that...
Microsomal and Prior to Preparations

Microsomes, Plasma membranes, 37°C. and effect on pads Adipose 1.50 glucose-sensitive cytochalasin not 3.70 1.16 furosemide concentration 2-h between in the states are findings that indicate the basal and insulin-stimulated states are equally sensitive to the inhibition.

The furosemide effect is saturable with a simple stoichiometry (Fig. 6). For a 2-h posttreatment (30 min at room temperature followed by 90 min at 4°C) the 50% inhibition occurred at the furosemide concentration of ~3.5 mM. This apparent Ki value was not significantly different between microsomes and plasma membranes. For an 18-h posttreatment, the concentration that effects 50% inhibition was reduced to ~0.9 mM for basal microsomes and 0.7 mM for insulin-stimulated microsomes. These findings indicate that the basal and insulin-stimulated states are equally sensitive to the inhibition.

The time dependence of the furosemide effect was already evident in the observed reductions in the apparent Ki value between 2-h and 18-h incubations (Fig. 6). Using a total particulate fraction of adipocytes (150,000 g fraction), the time course of the development of the inhibitory effect on the glucose-sensitive cytochalasin B binding was followed for 32 h (Fig. 7). With the furosemide concentration of 1 mM, the inhibition developed as an exponential function of time with a first order rate constant of ~0.12/h at 4°C.

In an attempt to examine if the furosemide effect is reversible, a 150,000 g preparation preincubated with 6 mM furosemide for 3 h at 4°C was, after separation by centrifugation, resuspended into a furosemide-free medium (a final furosemide concentration <0.06 mM) for 3 h at 4°C. This procedure reduced the inactivation on the glucose-sensitive cytochalasin B binding activity from 71% down to 25%, indicating that the inactivation is only slowly reversible (data not illustrated).

The effects of longer term incubation with a lower concentration of furosemide on 3-OMG flux was also examined. Adipose pads were incubated for 48 h at 37°C with or without 0.1 mM furosemide. Adipocytes were isolated from these pads, then treated with insulin (7 nM), and 3-OMG equilibrium flux was measured as described in Methods. The half time (t1/2) observed at the 3-OMG concentration of 5 mM were 22 and 43 s for control and furosemide-treated samples, respectively. This indicates significant (50%) inhibition of transport occurs at therapeutically attainable plasma concentration of furosemide with prolonged incubation of cells at 37°C. This should be compared with the effect on cytochalasin B binding of an 18-h incubation at 4°C with 0.1 mM furosemide of ~25% (calculated from data of open triangles in Fig. 6).

Discussion

We have shown in this study that furosemide inhibits the 3-OMG equilibrium exchange flux in isolated rat adipocytes. The inhibition is independent of insulin since flux in basal as well as insulin-stimulated adipocytes are affected equally. We
have also shown that furosemide inhibits the glucose-sensitive cytochalasin B binding activities of both plasma membranes and microsomal membranes of adipocytes. There is ample evidence that glucose-sensitive cytochalasin B binding proteins are the glucose transport carrier molecules in adipocytes and other cells (16, 20, 25). In adipocytes, these binding proteins exist in two pools, namely in plasma membranes and microsomal membranes, representing active and reserve pools of carrier, respectively (16, 17). A widely accepted current hypothesis proposes that insulin induces recruitment of carrier molecules from the microsomal reserve pool into the plasma membrane (16, 17), thereby stimulating glucose transport. In the present study we demonstrate that furosemide inhibits the D-glucose-displaceable cytochalasin B binding in both pools with affinities practically indistinguishable from each other. Furthermore, no significant differences were found between the inhibitions observed with samples pretreated with insulin versus noninsulin-treated controls. These data indicate that furosemide inactivates the glucose carrier molecules by direct interaction, and this is independent of their location (plasma membrane vs. microsomes) or of their state (basal- or insulin-stimulated). The posttreatment is almost as effective as the pretreatment, indicating that the intact cell structure is not prerequisite to the drug effect. There is no indication in our data that furosemide inhibits insulin-induced recruitment of cytochalasin B binding activity or glucose carriers from microsome to plasma membrane. This is consistent with the observation referred to above that furosemide affects the 3-OMG flux equally independent of insulin treatment.

The carrier inactivation by furosemide appears to be time dependent. The time course of the loss of D-glucose-displaceable cytochalasin B binding activity of total particulate fractions revealed that the inactivation develops slowly. It is a first order process with a rate constant of ~0.12/h for 1 mM of furosemide at 4°C. The inactivation is only slowly reversible, being sustained after several washings. When the incubation time with furosemide is fixed, the inhibition observed is a saturable function of inhibitor concentration, which follows noncompetitive mode of inhibition kinetics with one-to-one stoichiometry. The apparent inhibition constant ($K_i$) estimated by our analysis was between 3.5-4.5 mM for 2 h incubation in furosemide. This was the same for both the glucose-flux assay and the cytochalasin B binding assay. When incubation time in furosemide is prolonged to 18 h, the $K_i$ value was reduced to between 0.7 to 0.9 mM.

The in vitro inactivation of the glucose carrier by furosemide revealed in this study may have a clinical significance. It should be noted that $K_i$ values for the in vitro effects with a short-term incubation are considerably higher than observed or calculated serum furosemide concentrations in patients receiving therapeutic doses (<0.1 mM) (12, 26, 27). Nevertheless, in view of the time-dependent, progressive, and poorly reversible nature of the observed effects, even these low blood levels could significantly inhibit glucose transport in peripheral tissues in vivo if drug therapy is continued for extended periods of time. In fact, our preliminary results show that after 48 h incubation at 37°C furosemide at a concentration as low as 0.1 mM inhibits 50% of transport function. The comparison of effective in vivo vs. in vitro concentrations is complicated by possible enrichment of the drug at specific cellular sites and lack of detailed information on volume of distribution and systemic fate of diuretic agents. Evidence of progressive accumulation of furosemide in renal tissue has been observed (28, 29). The loop diuretic ethacrynic acid is also known to be accumulated by kidney slices to concentrations 10-fold higher than those in the incubation media (30). In other in vitro systems the effective concentrations of furosemide are not markedly different from those causing effects reported in this paper. The concentration of furosemide in perfusion medium that inhibits sodium reabsorption maximally in isolated perfused renal tubule is 1 mM (31). The effective concentration for inhibition of sodium transport in the toad bladder is 0.8 mM (32). Taken together, these considerations suggest that the drug-induced inhibition of glucose transport in peripheral tissues could contribute to the glucose intolerance observed in patients receiving diuretic therapy. Furthermore, our results demonstrate that hydrochlorothiazide and piretine were as effective as furosemide, while bumetanide was significantly less potent. This is in accordance with reports that bumetanide in equivalent diuretic dosage may be less prone to cause glucose intolerance (33, 34).

In conclusion, furosemide and other diuretics inactivate in vitro the glucose transport carrier of adipocyte independent of insulin. We propose that this may contribute to the glucose intolerance in patients receiving diuretic therapy.
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