

Basic Study

Vanadium-dependent activation of glucose transport in adipocytes by catecholamines is not mediated via adrenoceptor stimulation or monoamine oxidase activity

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Author contributions: Carpéné C designed the studies, performed the rat experiments, reviewed the literature, and wrote the manuscript; Tavernier G performed the mouse experiments, contributed to the literature review and revised the manuscript; Fontaine J isolated the cells for *in vitro* studies; Morin N was involved in the generation and analysis of the data.

Institutional review board

statement: The study was approved by the I2MC Institutional Review Board: Institut des maladies métaboliques et cardiovasculaires.

Institutional animal care and use

committee statement: Mice were housed and manipulated according to the INSERM guidelines and European Directive 2010/63/UE by competent and expert technicians or researchers in animal care facilities with agreement number A 31 555 04 and

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Abstract

BACKGROUND

Benzylamine and methylamine activate glucose uptake in adipocytes. For tyramine, this effect has even been extended to cardiomyocytes.

AIM

To investigate the effects of catecholamines and other amines on glucose uptake.

METHODS

A screening compared 25 biogenic amines on 2-deoxyglucose (2-DG) uptake activation in rat adipocytes. Pharmacological approaches and transgenic mouse models were then used to decipher the mode of action of several hits.

RESULTS

In rat adipocytes, insulin stimulation of 2-DG uptake was reproduced with catecholamines. 100 $\mu\text{mol/L}$ or 1 mmol/L adrenaline, noradrenaline, dopamine and deoxyepinephrine, maximally activated hexose transport only when sodium orthovanadate was added at 100 $\mu\text{mol/L}$. Such activation was similar to that already reported for benzylamine, methylamine and tyramine, well-recognized substrates of semicarbazide-sensitive amine oxidase (SSAO) and monoamine oxidase (MAO). Several, but not all, tested agonists of β -adrenoreceptors (β -ARs) also activated glucose transport while α -AR agonists were inactive. Lack of blockade by α - and β -AR antagonists indicated that catecholamine-induced 2-DG uptake was not mediated by AR stimulation. Adipocytes from mice lacking β_1 -, β_2 -

C 31 555 07. The experimental protocol was approved by the local ethical committee CEEA nb122.

Conflict-of-interest statement: The authors declare no competing financial interests.

Data sharing statement: No additional data are available.

ARRIVE guidelines statement: The authors have read the ARRIVE guidelines, and the manuscript was prepared and revised according to the ARRIVE guideline.

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Manuscript source: Invited manuscript

Specialty type: Endocrinology and metabolism

Country/Territory of origin: France

Peer-review report's scientific quality classification

Grade A (Excellent): 0
Grade B (Very good): 0
Grade C (Good): 0
Grade D (Fair): 0
Grade E (Poor): 0

Received: August 31, 2020

Peer-review started: August 31, 2020

First decision: October 5, 2020

Revised: October 12, 2020

Accepted: October 26, 2020

Article in press: October 26, 2020

Published online: December 15, 2020

P-Reviewer: Choi MR

and β_3 -ARs (triple KO) also responded to millimolar doses of adrenaline or noradrenaline by activating hexose transport in the presence of 100 $\mu\text{mol/L}$ vanadate. The MAO blocker pargyline, and SSAO inhibitors did not block the effects of adrenaline or noradrenaline plus vanadate, which were blunted by antioxidants.

CONCLUSION

Catecholamines exert unexpected insulin-like actions in adipocytes when combined with vanadium. For limiting insulin resistance by activating glucose consumption at least in fat stores, we propose that catecholamine derivatives combined with vanadium can generate novel complexes that may have low toxicity and promising anti-diabetic properties.

Key Words: Adipocyte; Amine oxidases; Insulin; Diabetes; Vanadium; Obesity

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Core Tip: In rat and mouse fat cells, the combination of catecholamines with vanadium reproduces the sugar entry activation already reported for benzylamine, methylamine or tyramine. Glucose transport stimulation is observed only with catecholamines at millimolar doses and in the presence of a vanadium dose that is ineffective on its own. The synergism between adrenaline or noradrenaline and vanadate is not mediated by adrenoceptors and resists amine oxidase inhibitors; while it is sensitive to antioxidants. Since vanadium exhibits antidiabetic properties, but with toxicological concerns, it is proposed that the combination of catecholamine derivatives plus vanadate salts might generate complexes with safer blood glucose-lowering properties.

Citation: Fontaine J, Tavernier G, Morin N, Carpené C. Vanadium-dependent activation of glucose transport in adipocytes by catecholamines is not mediated *via* adrenoceptor stimulation or monoamine oxidase activity. *World J Diabetes* 2020; 11(12): 622-643

URL: <https://www.wjnet.com/1948-9358/full/v11/i12/622.htm>

DOI: <https://dx.doi.org/10.4239/wjd.v11.i12.622>

INTRODUCTION

Type 2 diabetes is one of the morbid complications of obesity. The hypertrophied adipose tissues (AT) do not play only a quantitative but also a qualitative role in glucose handling. Due to the elevated inflammation state and the oxidative stress found in pathological adipose tissue, its constitutive cell types (adipocytes, precursor cells and immune cells) modulate the adipokines they release in the organism, leading to a state of insulin-resistance. With the exception of adiponectin, which facilitates the action of insulin, most of the adipokines are proinflammatory or impair insulin antihyperglycemic actions: interleukin (IL)-6, tumor necrosis factor α , and many others, which can be exemplified by one of them, the well-named resistin.

To treat such diabetic insulin resistant- and insulin deficient-states, the search for molecules that reproduce totally or partially the insulin activation of glucose transport has been included in the antidiabetic strategies developed so far. In the search for activators of glucose uptake that may replace insulin, the adipocytes themselves constitute a valuable cell model since their glucose transport capacity is highly responsive to insulin, at least in non-obese, non-diabetic rodents.

Glucose entry can be measured as an acute response after short-term exposure of adipocytes to the tested molecules followed by a transport assay consisting of the measurement of the intracellular uptake of a non-metabolizable analogue of glucose, such as 2-deoxyglucose (2-DG). Using this model, it was previously observed that the reactive oxygen species (ROS), especially hydrogen peroxide, partially mimic the insulin stimulation of glucose uptake^[1] while an excess of pro-oxidant agents, leading to oxidative stress hampers insulin action^[2]. Notably, the combination of hydrogen peroxide with vanadium, known to generate peroxovanadate, results in a more potent stimulation of glucose uptake as it acts downstream of the insulin receptor directly on

S-Editor: Gao CC**L-Editor:** Webster JR**P-Editor:** Ma YJ

the turnover of glucose transporters (GLUT4) between intracellular vesicles and extracellular surface^[3].

Similarly, it has been reported that hexose transport stimulation can be obtained by incubating fat cells with benzylamine and sodium orthovanadate^[4,5]. Indeed, the amine is oxidized by amine oxidases that generate benzaldehyde, ammonium and hydrogen peroxide during the oxidative deamination. Among these end-products, it is hydrogen peroxide that reacts with vanadate to generate peroxovanadate thereby inhibiting tyrosine phosphatases involved in the turn-off of glucose carrier translocation to the cell surface^[6]. Since benzylamine is a reference substrate for a class of amine oxidases that is abundant in adipocytes, the semicarbazide-sensitive amine oxidase (SSAO)^[7,8], we expected to extend such 'insulin like' property to other substrates. As SSAO has been renamed primary amine oxidase (PrAO) because it is blocked by many other inhibitors in addition to semicarbazide and due to its activity on a wide range of primary amines^[9], the number of potential insulin-mimicking agents was largely increased among the SSAO/PrAO substrate candidates. This was the case for methylamine, another SSAO/PrAO substrate of reference, which was demonstrated to activate glucose transport in rat adipocytes^[10] and in human fat cells^[11]. Methylamine is also active in mouse fat cells^[12]. Both benzylamine and methylamine have revealed anti-hyperglycemic effects in mice^[13,14], rats^[15], and rabbits^[10], and these effects are abolished in mice invalidated for SSAO/PrAO, also known as vascular adhesion protein-1 (VAP-1)^[16].

Alongside SSAO/PrAO, other amine oxidases are present in adipocytes, which also release hydrogen peroxide when oxidating their substrates. Among them, monoamine oxidases (MAO) have been reported to be involved in tyramine stimulated glucose transport in rodent adipocytes and cardiomyocytes^[17], which is also enhanced by vanadium^[18]. Similar to benzylamine, tyramine also exhibits long-term insulin-like actions such as the stimulation of adipose differentiation of preadipocytes^[19].

In this context, we aimed to verify in adipocytes whether the insulin-like properties of benzylamine plus vanadium could be reproduced or improved by exposing the fat cells to other amines, in order to retrieve novel insulin-mimickers on the basis of glucose transport activation, and candidates leading to potential antidiabetic strategies. We thus performed a screening of various amines, with a focus on testing them with and without vanadium in the incubation medium of freshly isolated rat adipocytes. The addition of sodium orthovanadate (Na₃VO₄) at 100 μmol/L in the incubation medium was chosen for this comparative approach as this has been successfully performed in rat adipocytes^[18]. This led us to use the diminutive term vanadate throughout this study although vanadyl and vanadate forms can coexist together with multiple oxidation states or oligovanadate forms in biological systems^[20]. Our screening was mainly limited to naturally occurring amines, since we previously reported insulin-like properties of non-natural compounds^[21], arylalkylamines^[22,23] and the hexakis (benzylammonium) decavanadate^[24], which have not been approved for further studies due to toxic side-effects.

The following results will provide evidence for the novel, insulin-like effect of catecholamines. Catecholamines are monoamine neurotransmitters composed of a catechol cycle (benzene with two hydroxyl side groups) and a side-chain amine: dopamine (with NH₂), noradrenaline (with OH and NH₂, also known as norepinephrine), and adrenaline (also named epinephrine, with OH and NH-CH₃). Catecholamines, under certain conditions, are able to activate glucose transport in fat cells in an unexpected manner, different to that of the reference SSAO/PrAO and MAO substrates: benzylamine, methylamine and tyramine.

MATERIALS AND METHODS

Chemicals

(+/-)-Adrenaline (also named epinephrine), (-)-noradrenaline (also named norepinephrine or arterenol), dopamine, tyramine, benzylamine, sodium orthovanadate, (-)-isoprenaline (also named isoproterenol), collagenase, dimethyl sulfoxide and most of the other reagents were from Sigma-Aldrich-Merck (Saint Quentin Fallavier, France). Brimonidine was a gift from the late Dr. Paris H. (Toulouse, France), while CL 316243 was kindly given by Dr. Lafontan M. (Toulouse, France). [³H]-2-deoxyglucose was from Perkin Elmer (Boston, MA, United States).

Animals and rodent adipocyte preparations

Male Wistar rats from Charles River (L'Arbresle, France) were euthanized after

overnight fasting when 2- to 3-mo old, according to Inserm guidelines. The generation of "β-less" mice was performed at the Centre Médical Universitaire (Genève, Switzerland). β-less ($\beta_1^{-/-}\beta_2^{-/-}\beta_3^{-/-}$) and wild-type (WT) ($\beta_1^{+/+}\beta_2^{+/+}\beta_3^{+/+}$) strains were obtained by intercrossing β_3 -AR knockout mice^[25] and β_1/β_2 -AR knockout mice^[26], kindly provided by Dr. Kobilka B.K. (Howard Hughes Medical Institute, Stanford, CA, USA). $\beta_1^{+/-}\beta_2^{+/-}\beta_3^{+/-}$ offspring were crossed to generate established WT and β-less colonies on the same, mixed genetic background (129 Sv/ev, 129 Sv/J, FVB/N, C57BL/6J, and DBA/2). After being transferred by shipment adapted for genetically modified organisms, the mice were bred under SOPF status at the animal unit of UMS006 (Toulouse, France); genotypes were determined by Southern blot as described elsewhere^[27]. Studies were performed on 4 β-less males aged 32 wk, while 13 WT mice with a mean age of 22 wk were studied in parallel. All the rodents were treated in accordance with the ARRIVE guidelines (Animal Research: Reporting of In Vivo Experiments)^[28].

Adipocyte preparations were obtained by collagenase digestion of perigonadic, retroperitoneal, perirenal and inguinal fat pads as described previously^[24]. The same procedure was applied for rat and mouse adipocyte preparations, as previously reported^[29]. Briefly, adipose tissues were minced with scissors in Krebs-Ringer salt solution pH 7.5 containing 15 mmol/L sodium bicarbonate, 10 mmol/L HEPES and 3.5% fat-depleted bovine serum albumin. As the only source of glucose for the cell preparations tested for glucose uptake assays was the non-metabolizable analogue described below, 2 mmol/L pyruvate was present in the medium for energy supply. After digestion with 1 mg/mL collagenase type II for approximately 45 min under agitation, buoyant adipocytes were separated by filtration through nylon mesh and washed twice with medium to eliminate collagenase and obtain functional adipocyte suspensions, ready for use in the initial screening or the complementary approaches. Suspensions of freshly isolated rat adipocytes were used at approximately 15 mg lipids/400 μL unless otherwise stated.

Glucose transport assay

An isotopic dilution of the non-metabolizable glucose analogue [³H]-2-DG^[30] was prepared to reach a final concentration of 0.1 mmol/L when added to 400 μL of fat cell suspension as described previously^[31]. This radioactive tracer was added after 45 min preincubation with the tested or reference agents. Then, the fat cells were incubated for additional 10 min in the presence of [³H]-2-DG (approximately 1300000 dpm/vial) and stopped with 100 μL of 100 μmol/L cytochalasin B. 200 μL of cell suspension were immediately transferred to plastic centrifugation microtubes prefilled with dinonyl phthalate (density 0.98 g/mL) before a 40 s spin, to separate the buoying adipocytes from the medium as described previously^[10]. The upper part of the tubes, containing radiolabelled hexose internalized in intact fat cells above the silicon layer was then counted in scintillation vials. The extracellular [³H]-2-DG present in this upper part of the tubes, and which was not internalized in cells was determined with adipocytes whose transport activity was previously blocked by cytochalasin B at time 0. It did not exceed 1% of the maximum 2-DG uptake in response to insulin and was subtracted from all assays. No radioactivity was found in the silicon layer separating the upper and lower part of the microtubes (not shown), in accordance with the entirely hydrophilic nature of 2-DG^[30].

Setting of negative and positive controls for glucose uptake-based screening of amines

Irrespective of the size of the molecular library used for a screening to test the properties of chemicals on hexose uptake it is mandatory to define the baseline and the maximal level of the transport of a non-metabolizable analogue of glucose, and to compare the unknowns to these negative and positive cut-offs. To perform a pilot approach comparing a total of 25 amines, we used a method previously validated, based on the above described uptake of [³H]-2-deoxyglucose, which has been revealed to be efficient in detecting insulin-like agents, such as vanadium derivatives^[24] or inhibitors such as β-AR agonists^[31]. The negative and positive controls were simply normalized, by setting baseline uptake at 0%, while maximal response to 100 nmol/L insulin was set at 100%. Each of five subsets of five amines was tested for 45 min with rat adipocytes, in parallel with the above-mentioned controls. Since each subset was tested on three independent adipocyte preparations, a total of fifteen adipocyte preparations isolated from fifteen rats was required for this first comparative approach. In the subsequent studies aiming at deciphering the mode of action of the best hits, the same protocol and data processing were used except that, alongside all

the required negative and positive controls, pharmacological agents were added during the 45-min preincubation period and remained present during 2-DG uptake assays. As each step of these mechanistic investigations was performed on groups of 4 to 6 rats, 38 rats were necessary for testing successively MAO and PrAO inhibitors, their interplay with vanadium, as well as the influence of different doses of dopaminergic and β - or α_1 -/ α_2 -adrenergic receptor antagonists or agonists. Thus, a total of 53 rats were used for both the pilot and mechanistic halves of the screening.

Lipolytic activity determination

Tested agents were added to 400 μ L of fat cell suspension as above for glucose uptake, with the following differences. The agents were incubated with the fat cells at 37°C under gentle shaking for 90 min in a buffer containing 5.5 mmol/L glucose instead of pyruvate and 2-DG. Incubations were stopped on ice. Lipolysis was determined by using glycerol release as an index as already documented, considering that free fatty acid release exhibits parallel variations in our experimental conditions^[52].

Statistical analysis

Results are presented as means \pm SE of (*n*) observations. All the statistical analyses for comparisons between parameters obtained in the presence of 100 μ mol/L vanadium and their respective control used the paired Student's *t* test. Other statistical comparisons used ANOVA followed by post-hoc Dunnett's multiple comparisons test, analyzed with Prism 6 for Mac OS X (from GraphPad software, Inc).

RESULTS

Screening of amines activating hexose transport in rat adipocytes under the influence of vanadium

Our preliminary screening approach aimed to detect naturally occurring amines that can mimic the previously reported effects of benzylamine, tyramine and methylamine on the uptake of the non-metabolizable glucose analogue 2-DG in fat cells. **Figure 1** displays data retrieved from 25 natural amines tested. The left panel of **Figure 1** shows that none of the amines increased hexose transport activity more than 20% of the maximal stimulation elicited by 100 nmol/L insulin (positive control, set at 100%). When tested alone, several amines even inhibited basal 2-DG uptake (baseline, set at 0%), leading to negative percentages. This was the case for n-decylamine, N-acetyl spermidine and trimethylamine. In contrast, methylamine, adrenaline, noradrenaline and tryptamine tended to slightly activate 2-DG uptake when tested at 0.1 and 1 mmol/L. Although such weak activations were detected, they were not considered positive hits since they never exceeded 20% of the maximal insulin effect, which was the threshold considered for the screening.

More importantly, when sodium orthovanadate was added at 100 μ mol/L at the beginning of the 45-min incubation period with amines, a totally different pattern was obtained. The chosen dose of vanadate, already reported to potentiate tyramine^[18] and benzylamine^[5] actions in rodent adipocytes was unable on its own to modify basal or insulin stimulated uptake, but facilitated the activation of glucose uptake by most of the tested amines. Several of them almost reached the maximal activation obtained with insulin itself. The powerful insulin-mimicking effect of methylamine on 2-DG uptake^[11] was confirmed since it was the best hit of the screening (**Figure 1**, right panel). Ranking the tested amines according to the response they elicited at 1 mmol/L in the presence of 100 μ mol/L vanadate showed that the millimolar dose of deoxyepinehrine, adrenaline, dopamine, and noradrenaline reproduced more than 75% of the insulin-induced maximal stimulation of glucose transport (**Figure 1**, right panel). A large proportion of the tested amines enhanced glucose uptake in a dose-dependent manner, except for benzylamine, n-decylamine and n-pentylamine, which were more efficient at 0.1 mmol/L than at 1 mmol/L. Since 100 nmol/L insulin increased the 2-DG uptake of isolated rat adipocytes by more than ten-fold (**Figure 2**), the stimulation reached after 45 min exposure to millimolar doses of these natural amines has to be definitely considered as substantial and highly significant, since the basal uptake was increased more than ten times. For tyramine and benzylamine (reference substrates of MAO and PrAO already demonstrated to activate 2-DG uptake in rodent adipocytes^[4,5,18]), the stimulatory property on glucose entry was also confirmed. As our aim was to detect amines that might reproduce or even overpass the previously described synergism between vanadium and benzylamine^[33],

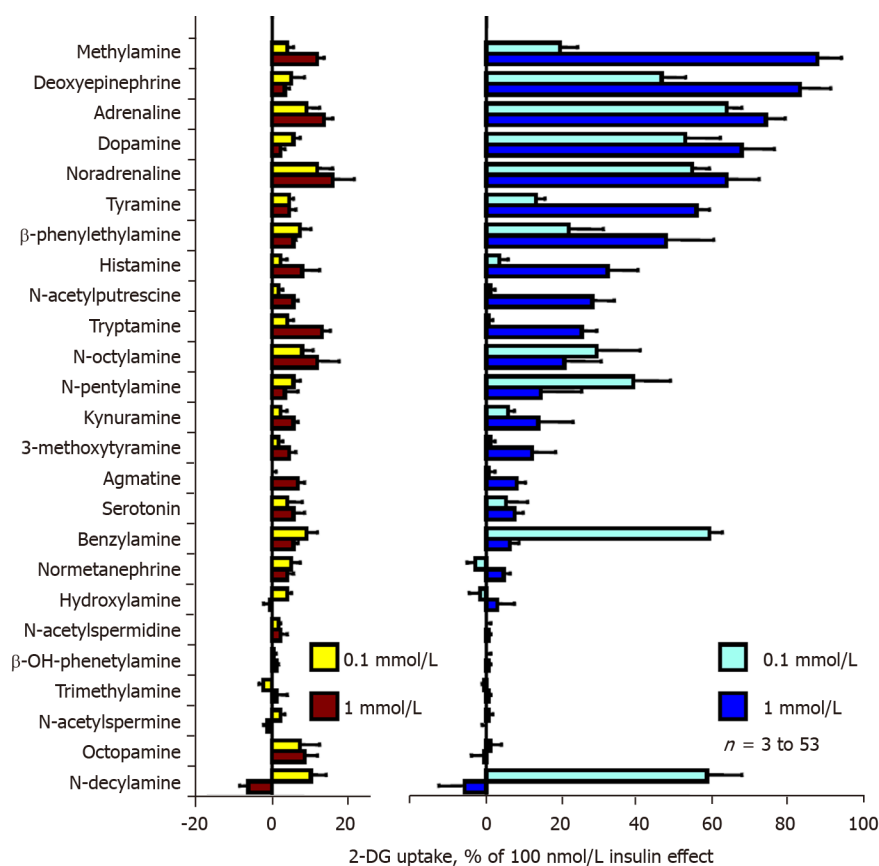


Figure 1 Screening of amines enhancing glucose transport in rat adipocytes under the influence of vanadate. Suspensions of rat fat cells were incubated without (left) or with 100 $\mu\text{mol/L}$ sodium orthovanadate (right) for 45 min in the presence of the indicated agents at 0.1 mmol/L (clearer columns) or 1 mmol/L (dark columns) just prior to the [^3H]-2-deoxyglucose (2-DG) transport assay for 10 min. 2-DG uptake was expressed as the percentage of maximal stimulation by 100 nmol/L insulin, with baseline set at 0%. Rank order of the amines was based according to their effect at 1 mmol/L + 0.1 mmol/L vanadate. The mean \pm SE from at least 3 determinations (for the agents almost inactive, at the bottom of the graph) up to 53 observations, for positive hits such as deoxyepinephrine, adrenaline (epinephrine), noradrenaline (norepinephrine), dopamine, methylamine, tyramine and benzylamine, with the seven most efficient amines being significant at $P < 0.001$. 2-DG: 2-Deoxyglucose.

tyramine^[18], or methylamine^[34], such confirmatory observations totally supported our chosen screening approach and amines of reference. Unfortunately, the screening did not allow the detection of a molecule that behaves more efficiently than the reference amines in stimulating glucose consumption. Nevertheless, such an approach identified various catecholaminergic neurotransmitters as potent activators of glucose uptake when present at 0.1 mmol/L together with vanadium. In fact, it was not so amazing to find adrenaline, noradrenaline and dopamine among the best hits of this preliminary screening since they share with the reference amines the capacity to be substrates of amine oxidases. However, a different readout was obtained with histamine, a well-known substrate of diamine oxidase, which was hardly more active at 1 than at 0.1 mmol/L, and with serotonin (a MAO-A substrate also known as 5-hydroxytryptamine), which was almost ineffective.

Noradrenaline and adrenaline were consequently considered as the major novel hits of the screening. On the large number of observations obtained during the primary screening and its further exploration steps, these two catecholamines clearly reproduced approximately 70%-80% of insulin stimulation on glucose entry when present at 0.1-1 mmol/L in combination with 0.1 mmol/L vanadate (cumulated number of preparations was $n = 53$ for each catecholamine). Although we were aware of the apparently extra-physiological nature of such an *in vitro* effect, we further explored its putative mechanisms of action, keeping in mind that these neurotransmitters are well-known substrates of central MAOs, which deaminate them into the corresponding aldehydes and hydrogen peroxide^[35].

Effects of pargyline and semicarbazide on amine-dependent glucose transport in rat adipocytes

100 $\mu\text{mol/L}$ vanadate did not alter basal 2-DG uptake in rat adipocytes in these

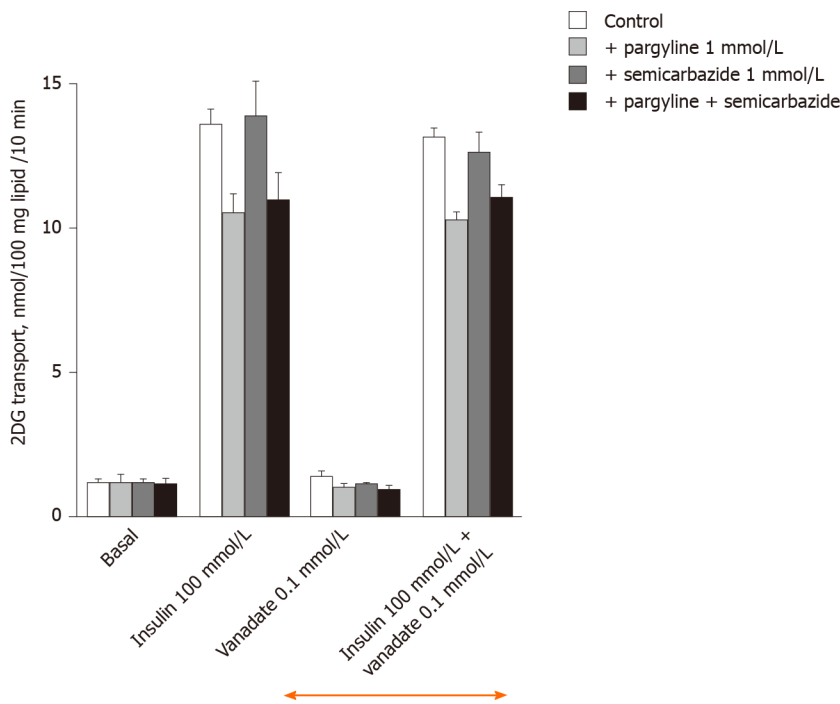


Figure 2 Lack of influence of amine oxidase inhibitors and of 100 $\mu\text{mol/L}$ vanadate on basal and insulin-stimulated hexose uptake in rat adipocytes. Rat fat cells (approximately 18 mg lipid/400 μL) were incubated for 45 min with the indicated concentrations of agents without (control, open columns), or with 1 mmol/L pargyline and 1 mmol/L semicarbazide, respectively (grey columns) or in combination (dark columns); then [^3H]-2-deoxyglucose uptake (2-DG) was assayed over a 10-min period. The mean \pm SE of 4 separate experiments was determined. There was no significant difference between inhibitors and each respective control. Similarly, no difference was found between corresponding conditions (basal or 100 nmol/L insulin-stimulated 2-DG transport) without (left) and with 100 $\mu\text{mol/L}$ vanadate (right, above orange arrow) by the paired *t* test. 2-DG: 2-Deoxyglucose.

mechanistic investigations, as during the preliminary screening experiments. This was also the case for 1 mmol/L pargyline (MAO inhibitor) or 1 mmol/L semicarbazide (SSAO/PrAO inhibitor) (Figure 2). The maximal effect of the high insulin dose (100 nmol/L) was also unaffected by 100 $\mu\text{mol/L}$ vanadate or by the inhibitors, either when tested separately or in combination (Figure 2). These observations fully support why the dose of 0.1 mmol/L vanadate is described as 'ineffective on its own' since it did not alter basal or insulin-stimulated uptake whereas higher doses of vanadium clearly activate glucose transport^[36].

The stimulation of 2-DG uptake induced by the combination of 1 mmol/L hydrogen peroxide and 100 $\mu\text{mol/L}$ vanadium corresponds to a well-recognized potentiation of the separate effects of ROS and metal ions^[20,37], and results from the non-enzymatic generation of peroxovanadate, a potent insulin mimicker^[38], the efficacy of which was confirmed here (Figure 3). Notably, the effect of hydrogen peroxide + vanadium was not impaired by the MAO and PrAO inhibitors. In contrast, the blockade of MAO and/or PrAO by pargyline and/or semicarbazide prevented tyramine, benzylamine, N-acetylputrescine, tryptamine and histamine from activating glucose transport (Figure 3). A complete inhibition of the latter amines was observed with the combination of pargyline and semicarbazide. However, for benzylamine and histamine, maximal inhibition was attained with semicarbazide alone, without any further inhibition obtained by its combination with pargyline (Figure 3). Surprisingly, the amine oxidase inhibitors, used either separately or in combination, were totally unable to alter the activation of glucose uptake by deoxyepinephrine, adrenaline, dopamine or noradrenaline (Figure 3). This finding was unexpected as it was initially supposed that catecholamines activated 2-DG uptake in rat adipocytes in a manner similar to that of other widely recognized substrates of MAO or PrAO^[7,17].

In fact, noradrenaline, adrenaline, dopamine and deoxyepinephrine can be considered as "false hits" in our screening. Elevated concentrations of these catecholamines stimulated hexose uptake in a manner that resembled that of hydrogen peroxide, but which was basically distinct from that of other MAO or PrAO substrates; including those used as internal references (benzylamine, methylamine, tyramine). In other words, their mechanism of action appeared independent from MAO or PrAO activity.

At this point, it is worth mentioning that the "insulin-like" effect of several

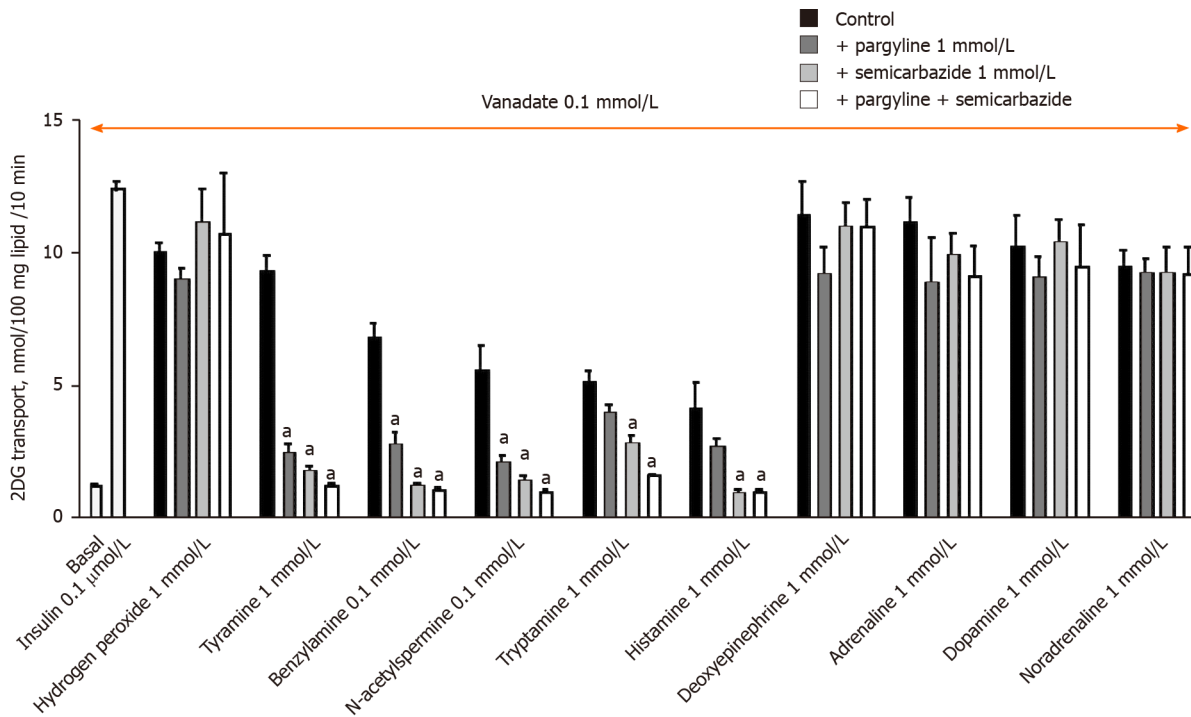


Figure 3 Inhibition by pargyline, semicarbazide, and their combination on amine-induced hexose uptake in rat adipocytes in the presence of 100 $\mu\text{mol/L}$ sodium orthovanadate. Immediately prior to the 10-min 2-deoxyglucose (2-DG) uptake assay, rat fat cells were incubated for 45 min as in Figure 2 in the presence of 100 $\mu\text{mol/L}$ vanadate (orange arrow) without (basal) and with 100 nmol/L insulin (light grey columns), or with the indicated agents (control, black columns). 1 mmol/L pargyline or 1 mmol/L semicarbazide (grey columns) and their combination (open columns) were added to each agent (hydrogen peroxide or amine) tested at 1 mmol/L, except benzylamine, which was tested at 0.1 mmol/L. A significant reduction of 2-DG uptake when compared to the respective control was observed at: $^*P < 0.02$. The mean \pm SE of 4 adipocyte preparations was determined. 2-DG: 2-Deoxyglucose.

catecholamines could not be qualified as non-specific since not observed with most of the other amines tested at 1 mmol/L. This was probably not a toxic effect, as noxious chemical species are rather inhibiting than facilitating or mimicking insulin action^[39].

Regarding the effect of dopamine, its resistance to the blockade by the dopaminergic antagonist metoclopramide (Figure 4) did not suggest any mediation *via* dopaminergic receptor activation. Similarly, dopaminergic receptors were apparently not involved in the stimulation of 2-DG uptake by adrenaline or noradrenaline (Figure 4). Thus, further exploration consisted of testing whether adrenergic receptors, which bind catecholamines, were involved in their stimulatory action on hexose uptake. Hence, diverse β -adrenoreceptor (β -AR) agonists have already been reported to modulate glucose transport activity in adipocytes, but with contradictory effects, from stimulatory^[40-42] to inhibitory^[31,43] depending on the nature (white, beige, brown) of the adipocytes, the animal species^[44], and the experimental conditions tested.

Short-term effects of adrenergic agents on glucose transport in rat adipocytes

Further pharmacological tests required to study the activation of hexose transport by high doses of catecholamines were performed with increasing doses of α -adrenoreceptor (α -AR) agonists in rat adipocytes, still in the presence of 100 $\mu\text{mol/L}$ vanadate: amidephrine, a α_1 -AR agonist, and brimonidine, an α_2 -AR agonist. Figure 5 indicates that stimulation with millimolar doses of catecholamines was not reproduced with these selective activators of α_1 - and α_2 -ARs in rat adipocytes, at least after short-term exposure to doses from 10 nmol/L up to 1 mmol/L.

Studies performed with β -AR agonists also indicated that several of them were totally unable to immediately activate glucose uptake, from micromolar to millimolar doses: procaterol (β_2 -AR agonist), CGP 12177 and CL 316243 (β_3 -AR agonists) (Figure 5, right panel). Only the β_1 -receptor agonist dobutamine and the pan-agonist (active on β_1 -, β_2 -, β_3 -ARs) isoprenaline reproduced at submillimolar and millimolar concentrations the hexose uptake stimulation observed with catecholamines. As these β -AR agonists are secondary amines structurally close to catecholamines, they can undergo similar amine oxidation, and thereby stimulate uptake independently from β -AR activation, either *via* a MAO-, or PrAO-dependent mechanism (as depicted for

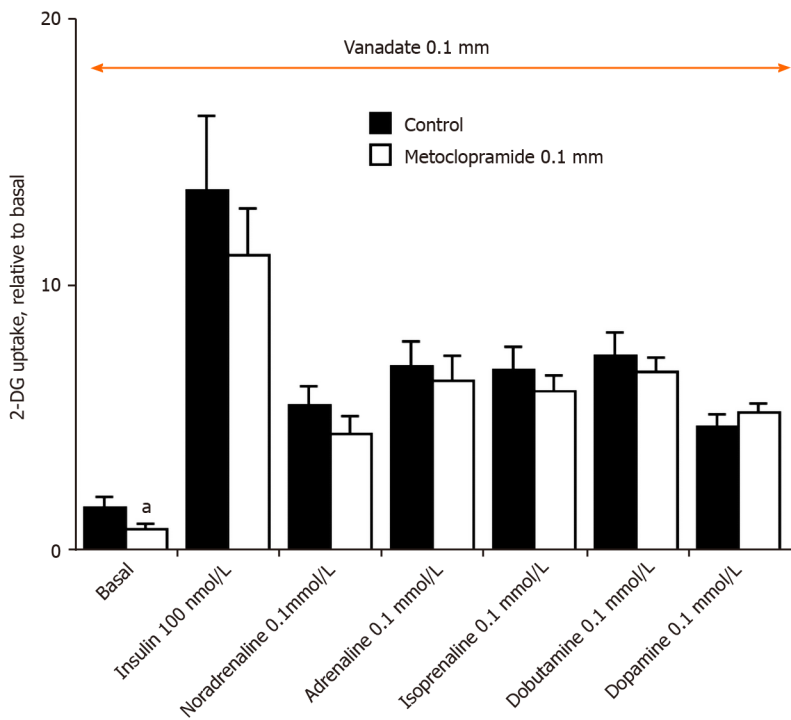


Figure 4 Lack of influence of the dopaminergic antagonist metoclopramide on insulin- or amine-stimulated hexose uptake in rat adipocytes. Rat fat cells were incubated for 45 min with the indicated concentrations of agents without (control, closed columns), or with 0.1 mmol/L metoclopramide (open columns); then uptake of the glucose analogue 2-deoxyglucose (2-DG) was assayed over 10 min. 2-DG uptake was expressed as the fold increase relative to basal uptake set at 1.0. The mean \pm SE of 4 separate experiments performed in the presence of 100 μ mol/L vanadate (orange arrow) was determined. A significant difference between metoclopramide and the respective control was observed at: ^aP < 0.05. 2-DG: 2-Deoxyglucose.

tyramine and tryptamine in Figure 3) or *via* an auto-oxidation process, as proposed for adrenaline.

At this step, the use of several AR-antagonists was necessary to study whether the effects of adrenaline and noradrenaline, but also those of isoprenaline, dobutamine and dopamine were resistant to adrenergic blockade. The selective α_2 -AR antagonist, methoxy-idazoxan (also known as RX 821002), did not influence at 10 μ mol/L the effects of adrenaline or noradrenaline on glucose transport in rat adipocytes in the presence of vanadate (Figure 6). However, this dose-dependent experiment definitely showed that a substantial activation of hexose uptake is detected only with 0.1 and 1 mmol/L of noradrenaline and adrenaline, while lower doses (*e.g.*, 10 μ mol/L) only tend to moderately enhance the transport activity (Figure 6). No clear-cut effect of α_2 -AR blockade could be detected on any of these two halves of the dose-response curves (Figure 6). Similar observations were made with the α_1 -AR antagonist prazosin at 10 μ mol/L (not shown). Increasing the doses of these two α -AR antagonists did not provide more conclusive findings (Figure 7). In addition, these high doses of α -AR antagonists did not hamper clearly the uptake stimulation by noradrenaline and adrenaline. As 0.1 mmol/L RX 821002 also altered the activation by the β -AR agonist isoprenaline, no definitive interpretation could be established from these tests, save that a lack of selectivity is often an expected pitfall of the use of large concentrations of recognized selective agonists and antagonists. Such loss of selectivity is often associated with cytotoxic effects and impairment of integrated biological responses. This was likely the case for RX 821002, which remained ineffective at 10 μ mol/L in most of the studied conditions (Figure 6), while it was inhibitory at 0.1 mmol/L irrespective of the stimulation studied (Figure 7). This also applied to the millimolar dose of prazosin, but was not the case for noradrenaline and adrenaline since their complete dose-response curves exhibited an almost sigmoid shape in the range of studied concentrations without any inhibition at the higher doses tested (Figure 6).

In this context, the β -AR antagonists were tested in order to delineate whether they could block the effects of high doses of catecholamines plus vanadate on 2-DG uptake at a maximal dose limited to 0.1 mmol/L for managing selectivity. At this dose, bupranolol or propranolol did not hamper basal or insulin-stimulated hexose uptake. They also did not affect the tested amines (Figure 8). They were therefore discarded in the predominant participation of β -adrenergic receptors in the catecholamine-

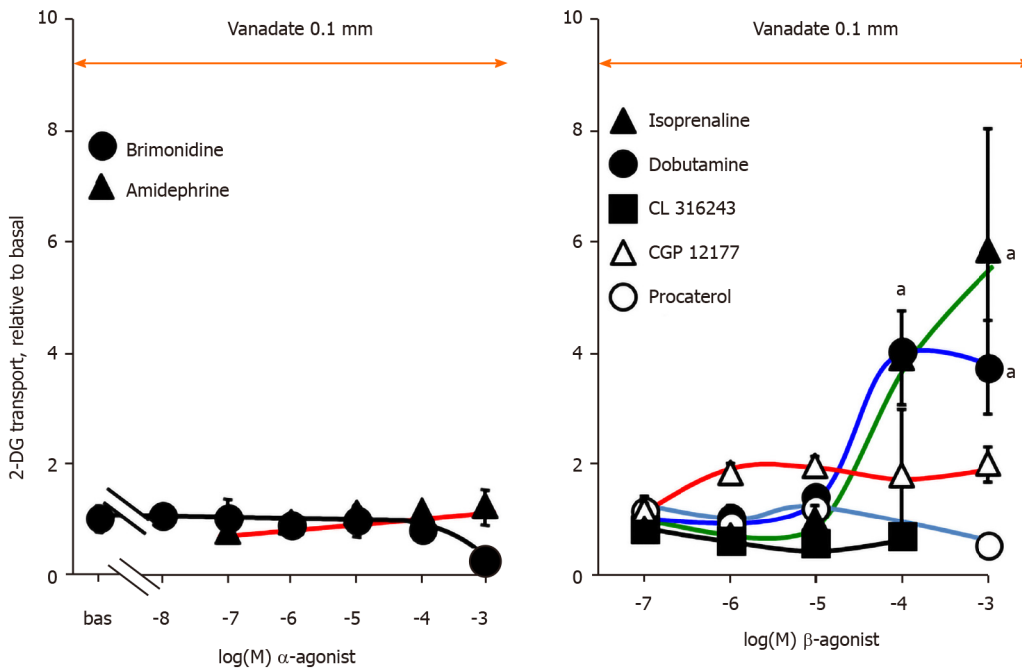


Figure 5 Effects of α - and β -adrenergic agonists on glucose transport in rat adipocytes. 2-Deoxyglucose uptake assay was performed as in Figure 3 with 100 $\mu\text{mol/L}$ vanadate (orange arrows) and increasing doses of the indicated α - (left panel) or β -adrenergic agonists (right panel). Glucose transport was expressed as the fold increase relative to basal uptake (bas) set at 1.0. The mean \pm SE of 4-6 adipocyte preparations was determined. Only high doses of isoprenaline and dobutamine induced responses were significantly different from baseline at: $^aP < 0.05$. 2-DG: 2-Deoxyglucose.

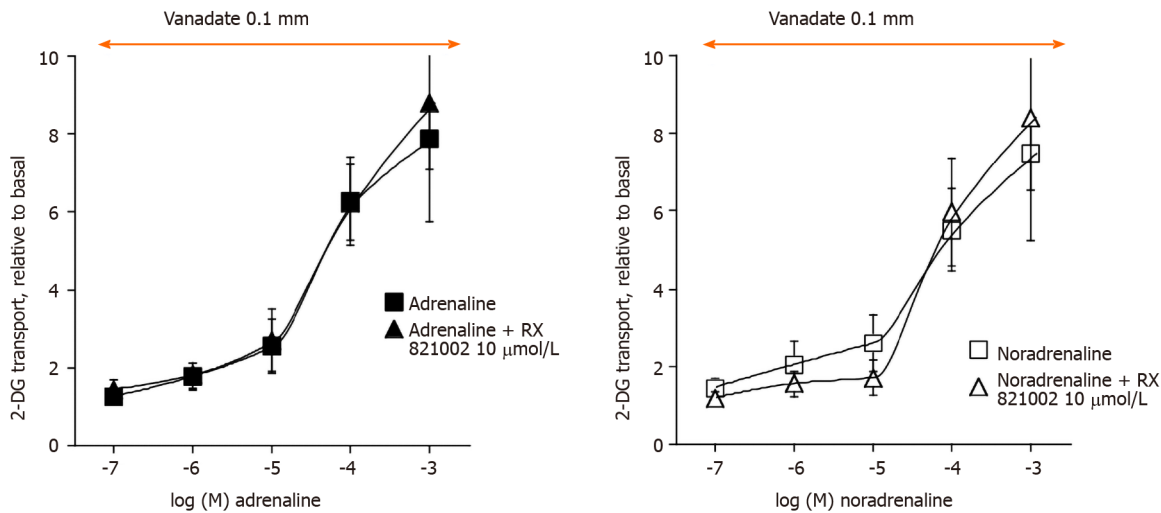


Figure 6 Dose-dependent activation by adrenaline and noradrenaline of hexose uptake in rat adipocytes: influence of the α_2 -adrenoreceptors antagonist methoxy-idazoxan. Rat fat cells were incubated for 45 min with the indicated concentrations of adrenaline (left panel) or noradrenaline (right panel) without (control, squares), or with 0.01 mmol/L RX 821002 (triangles) prior to the 2-deoxyglucose uptake assay. Hexose transport was expressed as the fold increase relative to basal set at 1.0. The mean \pm SE of 4 separate experiments performed in the presence of 100 $\mu\text{mol/L}$ vanadate (orange arrow) was determined. No significant difference was found between the inhibitor and the respective control. 2-DG: 2-Deoxyglucose.

induced activation of hexose uptake. Again, a high dose of another β -AR antagonist, CGP 20712, did not bring further clear information since this selective β_1 -AR blocker inhibited basal and insulin-stimulated uptake (Figure 8). Taken together, these findings did not support a major contribution of α -ARs or β -ARs in the effects of high doses of adrenaline. The complex influence of several β -AR agonists and β -AR-blockers on hexose transport did not permit us to demonstrate whether totality or a portion of the insulin-like effects of catecholamines plus vanadate were independent from β -AR activation.

Instead of performing more in-depth investigations in the same animal model, we took advantage of the availability of mice from the " β -less" lineage, *i.e.*, mice devoid of

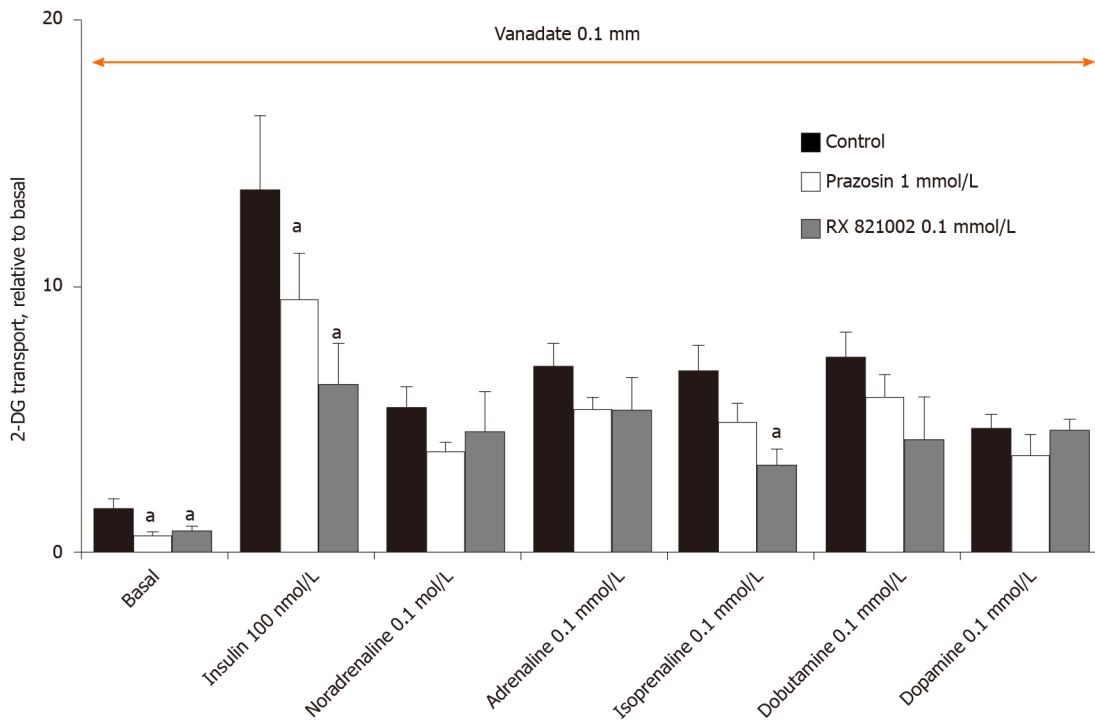


Figure 7 Inhibition of basal and insulin-stimulated hexose uptake in rat adipocytes by high doses of the α_1 -adrenoreceptors antagonist prazosin and the α_2 -adrenoreceptors antagonist methoxy-idazoxan. Rat fat cells were incubated for 45 min with the indicated concentrations of agents without (control, closed columns), and with 1 mmol/L prazosin (open columns) or 0.1 mmol/L RX 821002 (shaded columns) prior to 2-deoxyglucose uptake, expressed as the fold increase relative to basal. The mean \pm SE of 4 separate experiments performed in the presence of 100 μ mol/L vanadate was determined. Significant difference between the inhibitor and the respective control was observed at: ^a $P < 0.05$. 2-DG: 2-Deoxyglucose.

β -adrenergic receptors with triple knock-out of β_1 , β_2 , β_3 -adrenoreceptors^[27]. Our hypothesis-driven research was: if adipocytes from "β-less" mice, which are genetically invalidated for the three β -AR subtypes expressed in adipocytes, still respond to high doses of catecholamines by increasing glucose uptake, this should signify that β -ARs are not involved in the action of high doses of neurotransmitters.

Insulin-dependent and -independent stimulation of glucose transport in mouse adipocytes

Glucose transport in adipocytes from "β-less" mice was sensitive to insulin as was the case for control WT mice. While basal uptake was slightly higher in the triple KO strain, the dose-dependent response of fat cells to insulin stimulation was not dramatically blunted (Figure 9). As for rat adipocytes, maximal activation of 2-DG uptake was obtained with 100 nmol/L insulin, although its magnitude was slightly reduced in "β-less" mice. When tested in the presence of 100 μ mol/L vanadate, increasing doses of adrenaline and noradrenaline activated hexose uptake, leading to a similar degree of activation at 1 mmol/L in both genotypes (Figure 9). The activation by lower doses of catecholamines was even higher in "β-less" mice than in controls. These findings indicated that β -AR stimulation was not necessary for the activation of glucose transport by high doses of catecholamines. However, since 100 μ mol/L vanadate abolished the small difference between basal uptake in control and triple KO mice (Figure 9), further verifications were performed. In fact, the dose of 100 μ mol/L vanadium chosen from previous dose-response studies performed in rats^[18] appeared suitable for studies on mouse adipocytes since it did not alter the response to insulin while it allowed the detection of an "insulin-like" response to tyramine or benzylamine in both genotypes (Figure 10). The use of mouse adipocytes also validated the relevance of a synergism between vanadate and millimolar doses of catecholamines, when considering short-term activation of glucose uptake. Again, the comparison of all our tested conditions indicated that adrenaline and noradrenaline, when tested alone at high doses without vanadium, did not notably stimulate glucose transport in fat cells. These results also indicated that the response to insulin could be slightly altered by the lack of β -ARs, although the exact mechanism involved and the occurrence of compensatory mechanisms could not be established.

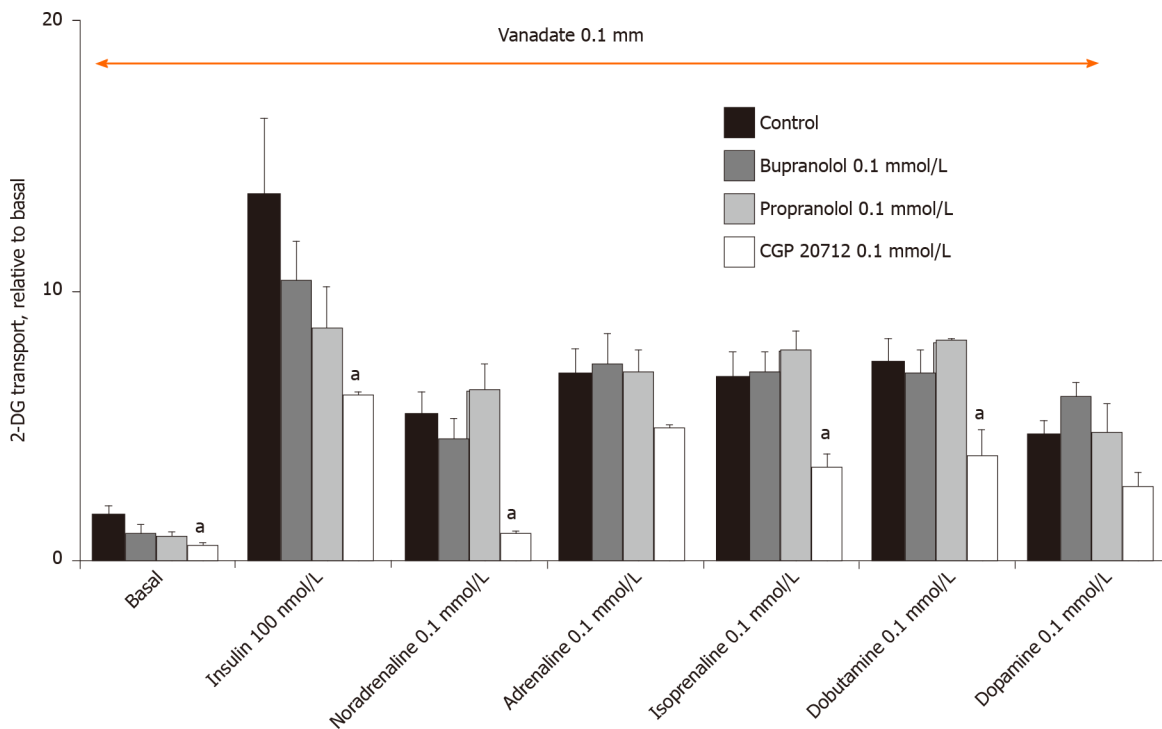


Figure 8 Influence of β -adrenoreceptors antagonists on hexose uptake in rat adipocytes. Rat fat cells were incubated for 45 min with the indicated concentrations of agents without (control, closed columns), and with 0.1 mmol/L bupranolol (shaded columns), 0.1 mmol/L propranolol (grey columns) or 0.1 mmol/L CGP 20712 (open columns) prior to 2-deoxyglucose uptake, expressed as the fold increase relative to basal. The mean \pm SE of 4 separate experiments performed in the presence of 100 μ mol/L vanadate was determined. Significant difference between the inhibitor and the respective control was observed at: $^aP < 0.05$. 2-DG: 2-Deoxyglucose.

As a detailed examination of the dose-response to catecholamines in triple KO mice showed that 2-DG uptake was even more readily activated by 10 and 100 μ mol/L noradrenaline in β -less than in WT mice (Figure 9), it was suggested that β -AR activation could even limit the effect observed in WT mice or rats. In keeping with this was the tendency of the β_3 -AR agonist CL 3162343 to inhibit basal 2-DG uptake in WT mice, which occurred at relatively low doses: 1-10 μ mol/L (Figure 11). Moreover, CL 316423 did not exhibit any potentiation with vanadate and definitely behaved differently from the catecholamines regarding glucose transport regulation (Figure 11). Nonetheless, it was verified that this β_3 -AR agonist was devoid of noticeable lipolytic activity in adipocytes from " β -less" mice: at 10 μ mol/L it hardly reproduced a fifth of the glycerol release found in WT mice (not shown), as previously reported^[27]. Such a defect in lipid mobilization was in entire agreement with the increased fatness we observed in the triple KO group, since the inguinal fat stores weighed 0.67 ± 0.12 g vs 0.20 ± 0.05 g in four " β -less" and control mice ($P < 0.02$).

At this stage, rather than deciphering the exact role of β -ARs in the regulation of glucose handling by mouse adipocytes, it was the possible influence of hydrogen peroxide in the observed synergism between vanadate and amines that was further analyzed. As with rat adipocytes, 1 mmol/L hydrogen peroxide activated 2-DG uptake to only $14.7\% \pm 3.4\%$ of the maximal insulin effect when tested alone, while when combined with 100 μ M vanadate, this ROS reproduced $85.0\% \pm 6.0\%$ of the insulin effect in WT mouse adipocytes ($n = 8$, $P < 0.001$). To determine whether amine oxidase inhibitors and/or antioxidants were able to prevent the synergism between adrenaline or noradrenaline and vanadate in mouse adipocytes, we tested pargyline, semicarbazide, N-acetylcysteine, glutathione and ascorbic acid. Again, insulin 100 nmol/L was used as a reference for glucose transport maximal activation. The insulin-induced stimulation almost remained unaltered by the tested agents (Figure 11). In contrast, the effects of adrenaline and noradrenaline were inhibited by antioxidants (notably abolished by 1 mmol/L ascorbate) while both resisted pargyline plus semicarbazide.

Together, these findings indicated that, in mouse and in rat, the catecholamines were active at high dose in the presence of vanadium in a manner that was: (1) Not predominantly reproduced by α -AR or β -AR activation; (2) Not mediated by MAO or PrAO activity; and (3) Dependent on redox mechanisms, probably *via* the generation

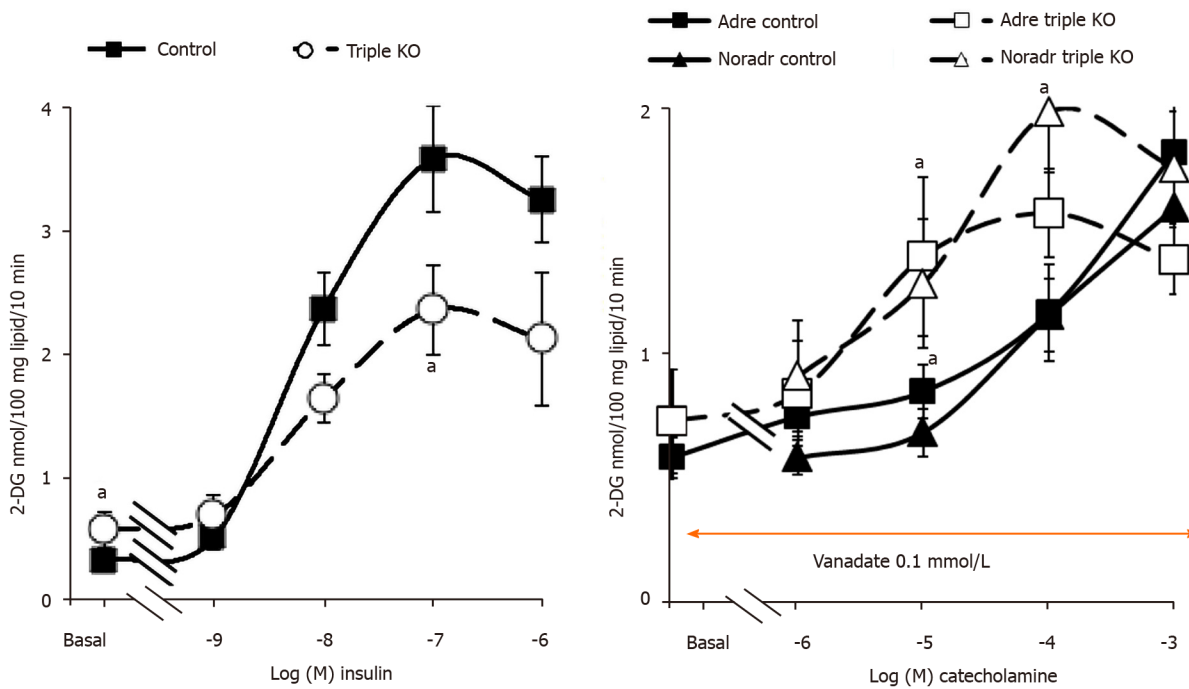


Figure 9 Glucose transport activation by insulin, adrenaline and noradrenaline in adipocytes from mice expressing or lacking β -adrenoceptors. Left panel: Insulin dose-dependent stimulation of 2-deoxyglucose uptake in adipocytes from wild-type (control, black symbols) and "β-less" mice lacking β_1 -, β_2 -, β_3 -adrenoceptors (triple KO, open symbols). Right panel: Stimulation of hexose uptake by adrenaline (adre, squares) and noradrenaline (noradr, triangles) in the presence of 100 μ mol/L vanadate (orange arrow). The mean \pm SE of 4 (triple KO, dotted lines) to 13 (control, solid lines) adipocyte preparations was determined. A significant difference from the corresponding control was observed at: ^a*P* < 0.05. 2-DG: 2-Deoxyglucose.

of ROS that mimicked the synergism found between hydrogen peroxide and vanadium.

Although the *in vitro* effects of the combination of catecholamines and vanadium were observed at relatively high concentrations, they were of considerable magnitude and activated glucose uptake in a rapid and reproducible manner in rodent adipocytes, and therefore could be qualified as insulin-like. All these observations probably do not have a direct physiological relevance for a better knowledge of glucose homeostasis, but they might add a novel insight on the development of vanadium-based antidiabetic approaches, as discussed below.

DISCUSSION

A stimulatory effect of high doses of the catecholamines dopamine, noradrenaline and adrenaline was constantly observed on glucose uptake when rodent adipocyte preparations were incubated in the presence of vanadium. The effect of these catecholamines cannot be classified as artifactual since not mandatorily obtained with other biogenic amines or sympathomimetic agents. Among the other most active biogenic amines was deoxyepinephrine, also known as epinine or N-methyldopamine, which is also a catecholamine. The catecholamine effect on glucose entry is relatively rapid and of high magnitude and can be termed an "insulin-like effect". Although not totally deciphered, its mechanism of action is different to other recognized substrates of MAO or SSAO/PrAO since the synergism between catecholamines and vanadium on glucose transport activation resists blockade by amine oxidase inhibitors, by contrast to previous observations made in similar conditions with tyramine^[18], benzylamine^[33] or methylamine^[10,11]. This catecholamine effect is also distinct from the noxious effects of the highest doses of other tested agents, which hamper rather than activate glucose transport in insulin-sensitive cells. In fact, inhibitory effects have been observed under identical conditions with supra-therapeutic doses of pharmacological agents (prazosin, CGP 20712, RX 821002) or with a millimolar dose of the polyamine n-decylamine. These inhibitions were much less interesting than the clear-cut activation of glucose transport provoked by the synergism between 0.1 or 1 mmol/L catecholamines and 0.1 mmol/L sodium orthovanadate.

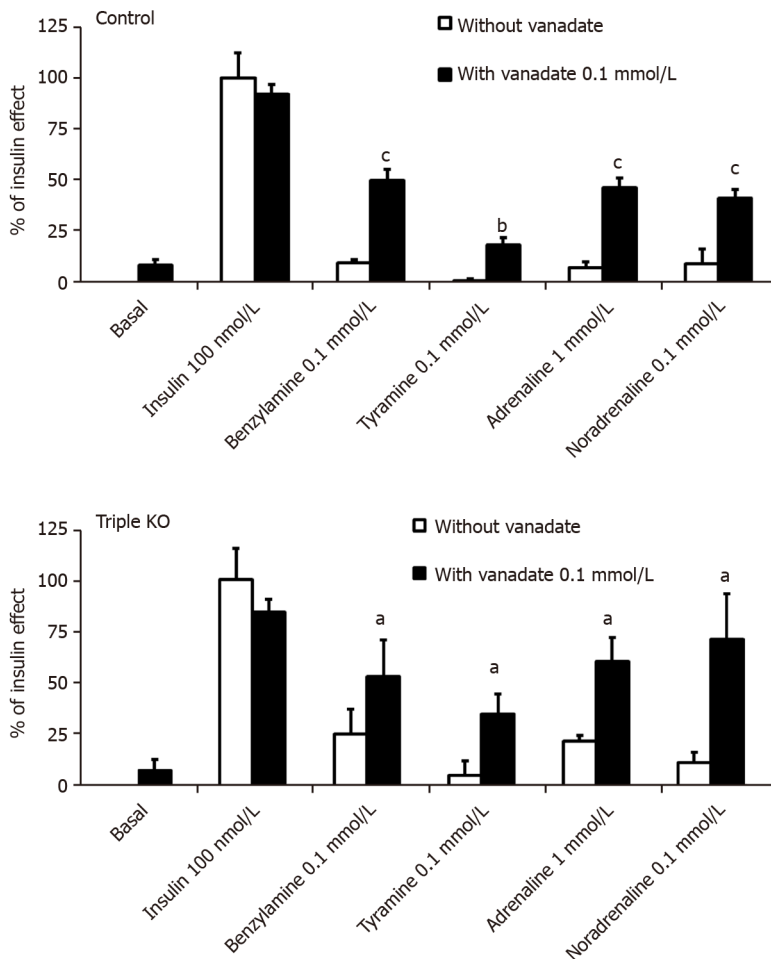


Figure 10 Influence of vanadate on hexose uptake into adipocytes from control and β -less mice. Suspensions of mouse fat cells from control (upper panel) or mice with triple KO for the β_1 -, β_2 -, β_3 -adrenoceptors (lower panel) were incubated for 45 min without (basal, set at 0%), with 100 nmol/L insulin (set at 100%), or with the indicated doses of amines alone (open columns), or in combination with 100 μ mol/L vanadate (dark columns), then [3 H]-2-deoxyglucose uptake was assayed over 10 min. The mean \pm SE of 4 (triple KO, lower panel) to 13 (control, upper panel) adipocyte preparations containing 17.2 ± 1.7 and 12.0 ± 1.6 mg lipids/400 μ L, respectively, was determined. A significant difference from the corresponding condition without vanadate was observed at: ^a $P < 0.05$, ^b $P < 0.01$, ^c $P < 0.001$.

Curiously, the catecholamine-dependent stimulation of glucose transport is not mimicked by β_1 -, β_2 -, β_3 -, α_1 - or α_2 -AR agonists (except those having a chemical structure close to that of catecholamines; *e.g.*, isoprenaline), even in the presence of vanadium, and is not clearly altered by adrenergic or dopaminergic antagonists. In addition, noradrenaline and adrenaline activate hexose transport in adipocytes from " β -less" mice, lacking β -ARs, demonstrating that such activation is adrenoceptor-independent. Sensitive to antioxidants, it likely results from a synergism between a still undefined oxidation or auto-oxidation product(s) of catecholamines reacting with vanadium, a metal known for decades to improve insulin sensitivity and to facilitate glucose handling^[23,45]. Indeed, potentiation between other amines and vanadium has already been reported to depend on peroxovanadate generation^[38]. It is widely recognized that metal ions favour the auto-oxidation of catecholamines and ROS generation^[46]. However, in the case of the interaction between catecholamine and vanadium studied here, the transition metal appears in turn to be changed to a novel state of oxidation by a redox-sensitive interplay. As vanadium can adapt multiple oxidation states, the forms of oxyanions generated in our preparations when sodium orthovanadate was mixed together with catecholamines and adipocytes remain undefined. Nevertheless, our experiments suggest that, after being oxidized, either spontaneously or enzymatically, catecholamines trigger a better signalling role for vanadium forms, thereby activating redox-sensitive processes such as glucose carrier recruitment and glucose transport activation. In this view, complex forms such as metavanadate and decavanate have been reported to be more active than other vanadium forms^[23,47,48].

The fact that ROS mediate some of dopamine signalling has been well characterized

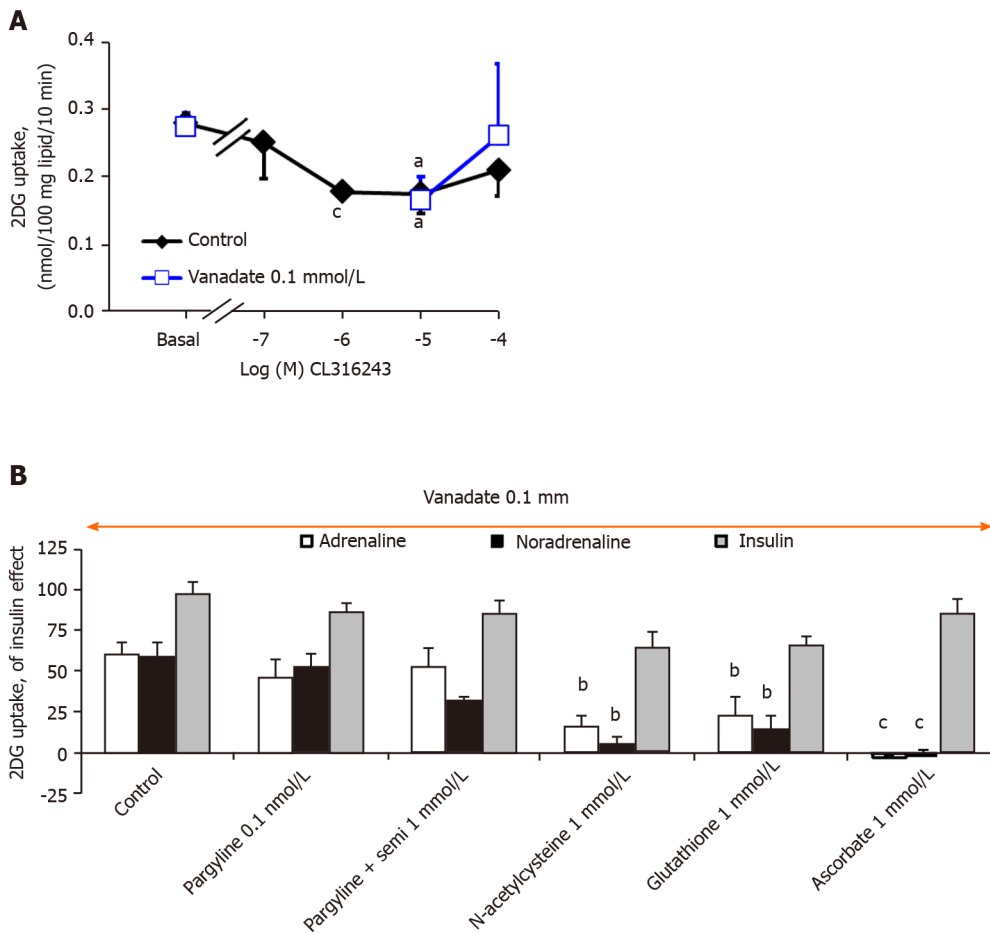


Figure 11 Influence of CL 316243, vanadate, amine oxidase inhibitors and antioxidants on glucose uptake in mouse adipocytes. Fat cells from wild-type mice were subjected to 2-deoxyglucose (2-DG) uptake in the presence of CL 316243, or catecholamines, insulin and vanadate. A: The β_3 -adrenoceptors agonist was incubated alone (control, closed diamonds) or with vanadate (open squares) at the indicated concentrations; B: 1 mmol/L adrenaline (open columns) and noradrenaline (black columns) or 100 nmol/L insulin (shaded columns) were incubated with 100 μ mol/L vanadate, without (control) or in the presence of pargyline 0.1 mmol/L, or pargyline + semicarbazide 1 mmol/L, or with 1 mmol/L of the antioxidants: N-acetylcysteine, glutathione, ascorbic acid. 2-DG uptake was expressed as the percentage of the insulin effect (which reached 1.73 ± 0.33 nmol 2-DG/100 mg lipids/0 min). The mean \pm SE of 4 determination was determined. A significant difference from the respective control was observed at: ^a $P < 0.05$, ^b $P < 0.01$, ^c $P < 0.001$. 2-DG: 2-Deoxyglucose.

in various cell types, such as in the renal proximal tubules^[49], but the exact cascade of events triggered by this and other catecholamines is still a matter of debate. It was out of the scope of the present study to totally decipher the mechanism underlying the synergism between catecholamines and vanadium, but it can be proposed that an auto-oxidation, or an oxidation independent of MAO and SSAO/PrAO activity, readily occurs at high doses of these neurotransmitters, and modifies the bioavailability of vanadium, improving its efficiency as an insulin mimicker, likely by turning off the activity of protein phosphatases^[45,50], especially those involved in the termination of insulin signalling^[23]. Unfortunately, we cannot attest whether the glucose uptake stimulated by catecholamines plus vanadate is a response to an increased energy demand of the fat cells, or is a direct consequence of insulin-sensitive glucose carrier recruitment to the membrane surface.

The lack of a clear α -adrenergic component in this catecholamine effect was evidenced by the use of several selective α_1 - and α_2 -AR agonists and antagonists, and confirms previous observations made in cultured mouse adipocytes^[51]. However, our *in vitro* observations are in apparent contrast with the decrease of interstitial glucose concentration induced by the α_1 -AR agonist norfenefrine and by noradrenaline in human abdominal subcutaneous adipose tissue during microdialysis experiments^[52]. In this clinical investigation, increased glucose uptake was accompanied by decreased blood flow. Both parameters were reversed upon pretreatment with the α_1 -AR antagonist urapidil^[52], and it has been proposed that adrenergic agents were mimicking the stimulatory effect of sympathetic nerves. Although a large contribution of α_1 -AR is not confirmed here, this clinical study argued as our findings for a noradrenaline activation of glucose entry into adipocytes that was independent of

insulin action.

The inhibitory effect of the β_3 -AR agonist CL 316243 observed in adipocytes from WT mice and reproduced to a lesser extent in rat adipocytes is in perfect agreement with previous reports on glucose transport inhibition in adipocytes by β -AR stimulation^[51,53,54]. This also explains the slightly enhanced basal uptake found in the " β -less" mice, but contrasts with other previous studies claiming that CL 316243 stimulates glucose uptake in mouse brown adipocytes in culture^[55] as well as β -AR activation under *in-vivo* conditions^[42]. Indeed, such controversy does not appear to have great relevance for the regulation of glucose uptake in human adipocytes, the β_3 -adrenergic component of which is remarkably weak^[56], and in which short-term CL 316243 administration neither induces alterations in the early insulin signalling cascade nor changes the basal level of glucose uptake^[57].

Another previous study that is in agreement with our unexpected observation of a catecholamine stimulation of glucose transport is the elegant demonstration performed by the Canadian group of Bukowiecki, showing that noradrenaline infusion in awake rodents stimulates glucose uptake in adipose tissue, especially brown fat depots^[58]. Obviously, the doses were lower in these *in vivo* experiments (noradrenaline was administered at 25 nmol/kg body weight/min) than in our *in vitro* studies. However, the activator effects of noradrenaline found in this study converge with those obtained with the microdialysis technique in human subcutaneous adipose tissue^[52] to support the functional relevance of our current observations.

Catecholamines are widely described to increase blood glucose and are termed catabolic hormones in Physiology, as they increase heart rate, blood pressure, and blood glucose levels; therefore, providing the organism with the capacity to better "fight or flight" under stress conditions; whilst the opposite is observed for the anabolic hormone insulin. Therefore, both the recognized hyperglycemic actions and the circulating levels of catecholamines appear to contrast strikingly with the "insulin-like" effects we report here. Before discussing the putative physiological relevance of our findings, it is necessary to reassess the range of the so-called physiological concentrations of catecholamines in biological systems, or at least in humans. It must be reminded briefly that the highest concentrations of catecholamines found in the granules of chromaffin cells of the adrenal glands can reach 600 mmol/L, while the circulating levels in blood are highly variable, but average the nanomolar range in man. The millimolar concentration of adrenaline required to observe synergism with vanadium on glucose uptake is therefore within these limits. If one considers only the catecholamine plasma levels, then the effects we report can be qualified as extra-physiological since high levels of catecholamines (also known as catecholamine toxicity) occur in pathological conditions only: Extreme stress, brainstem trauma, pheochromocytoma, and neuroblastoma. Nevertheless, adipocytes are not simply surrounded by plasma, and to the best of our knowledge, the range of catecholamine physiological levels found in the intercellular matrix neighbouring adipocytes is still not well established, not only due to the catecholamine short half-life and the high efficiency of their reuptake systems, but also when considering that the post-ganglionic orthosympathetic nerves can liberate some of their neurotransmitter vesicles in synaptic clefts very close to those adipocytes that are innervated. It can therefore be proposed that the mechanism described at high doses of catecholamines, may occur in discrete areas of adipose depots, in which local catecholamines are released. This aspect also relies on the influence of noradrenaline and innervation on adipose tissue development^[59], a research topic that has led to controversial findings regarding the origin of noradrenaline in this tissue during cold exposure. As the proposed local production of noradrenaline by activated macrophages of adipose tissue^[60] does not seem to have been confirmed in recent studies^[61], the debate regarding the relevance or the extraphysiological nature of the catecholamine doses used in our studies will only lead to speculative issues. For the sake of brevity, such debate needs to be closed and will give way to another discussion related to the multiplicity of catecholaminergic effects in adipocytes.

Adrenaline exerts multiple metabolic effects: it increases blood glucose at doses higher than those required for tachycardia induction; its stimulation of α_1 -ARs is glycogenolytic in liver and also contributes to hepatic neoglucogenesis, while β_2 -AR activation stimulates muscle glycogenolysis. Moreover, another widely known adrenergic effect contributes to increased blood glucose: The α_2 -AR mediated inhibition of insulin secretion in the endocrine pancreas. Therefore, how can a receptor-independent mechanism be required to increase glucose entry in several cell types when surrounding catecholamine levels are high?

One of our proposed answers is based on a functional point of view. Faced with overstimulation by catecholamines, adipocytes will strongly activate their lipolytic

activity, thereby releasing huge amounts of fatty acids (FA) that can be used by the body if energy expenditure is required or that can re-accumulate ectopically, a phenomenon known as lipotoxicity, involved in the onset of obesity complications. Indeed, it is conceivable that catecholamines stimulate both the mobilization of stored lipids and the utilization of glucose in adipocytes. This might facilitate a finely tuned management of triacylglycerol synthesis and breakdown, that varies rapidly between bouts of energy intake (post-prandial periods) or energy dissipation (exercise, fasting, cold exposure). Besides providing FA for other tissues *via* their lipolytic effect, catecholamines have recently been described to also increase FA oxidation rate in the adipocytes themselves, *via* the phosphorylation of signal transducer and activator of transcription 3 (STAT3), which, in turn regulates FA metabolism in a non-genomic manner^[62]. Alongside these two fates of released FA, mobilization and oxidation, there is also a third way, consisting in FA reesterification, which is predominant in adipocytes, since it limits excessive FA release and subsequent lipotoxicity. We therefore propose that, while increasing FA release and oxidation at low doses, catecholamines also increase FA reesterification when overstimulating adipocytes and when (dietary) vanadium is available. To esterify FA, adipocytes require a glycerol-P template, which is mainly provided from glucose since most of the fat cells are devoid of glycerokinase. Thus, we suggest that the effect we report here belongs to a possible feedback preventing adipocytes from excessive lipid mobilization when locally subjected to high doses of catecholamines, and this feedback requires enhanced glucose metabolism.

Another explanation of this unexpected effect of catecholamines relies on oxidative stress management. The autoxidation of catecholamines has been proposed to be a source of oxygen radicals that are noxious for cardiomyocytes, especially during ischemia-reperfusion injury^[63]. However, spontaneous catecholamine autoxidation is slow, but is considerably increased by enzymatic or metal catalysis, which results in the generation of the superoxide-free radicals, quinones and aminochromes. This process has been implicated in neurotoxicity. Catechol-O-methyltransferase (COMT) and MAO are the major enzymes involved in catecholamine metabolism and are supposed to prevent such cytotoxic species. However, when MAO, abundant in adipocytes^[64], oxidates amine substrates, it generates reactive aldehydes and hydrogen peroxide, *i.e.*, products which are as toxic as the substrates. The concomitant presence of vanadium together with catecholamines probably helps to circumvent the toxicological effects of each species. Their interaction, either spontaneous or enzymatically driven, seems to generate intermediates leading to a substantial increase of glucose uptake in a ROS-dependent manner. Peroxovanadate is known to be more efficient than vanadate in the stimulation of hexose uptake^[23]. Changes in the vanadium oxidation state generated in the presence of catecholamines might reproduce such a gain in insulin-like potency, resulting in uptake activations equivalent to those previously observed with benzylamine, being SSAO/PrAO-dependent or with tyramine, being mainly MAO-dependent.

By paying attention to the fact that overeating often accompanies the onset of insulin resistance, it is conceivable that the dietary intake of vanadium is sufficient to contribute to an alternative pathway for glucose uptake in the developing fat stores of obese subjects with insulin resistance, providing that stress conditions led to a rise in catecholamines to sufficient amounts. Indeed, regarding the stimulation of glucose uptake, it has already been described that vanadate is more efficient in insulin-resistant cells^[20].

Although the activation of glucose transport in rodent adipocytes up to maximal effects obtained with benzylamine and vanadate^[22,23] has not been overtly overpassed, our results open novel research aimed at improving the antidiabetic action of vanadium derivatives by limiting their cytotoxicity^[48,65,66]. One of the limitations of our study lies with the limited number of screened amines, since many aromatic or aliphatic amines—either naturally occurring or not—were not tested (*e.g.*, amphetamines). It is difficult to predict the behaviour of such amines not tested so far. Of note, trimethylamine, which is essentially a product of intestinal microbiota, and a precursor of trimethylamine N-oxide, currently associated with deleterious complications of obesity and diabetes^[67,68], was without any notable effect in the present screening, even at 1 mmol/L. However, evidencing several catecholamines as hits regarding glucose entry can be extended to many other amines susceptible to interplay with vanadium. In this view, the drug metformin, which resembles a polyamine, has been successfully coupled with vanadium complexes^[69] and one of the resulting compounds reported to exhibit hypoglycemic and lipid lowering properties after low dose administration in insulin-deficient or insulin-resistant diabetic rats^[70,71].

CONCLUSION

This study demonstrates, for the first time, that the combination of vanadium plus catecholamines strongly stimulates hexose uptake in rodent adipocytes and appears more promising than simply extraphysiological, in view of the efforts made recently to improve the benefits/risk ratio of vanadium-based antidiabetic approaches. This might open an additional, unexpected possibility to develop vanadium-containing antidiabetic agents that could become more active, less toxic and "drugable".

ARTICLE HIGHLIGHTS

Research background

Catecholamines have been demonstrated to induce oxidative stress by generating excessive reactive oxygen species (ROS) *via* amine oxidase-catalyzed oxidative deamination and *via* autoxidation. Like other metals, vanadium has been speculated to contribute to catecholamine oxidation. While catecholamines are recognized to mobilize lipids, vanadium is known to facilitate glucose consumption in fat cells. However, the effects of the combination of catecholamines plus vanadium on glucose utilization by adipose cells require clarification.

Research motivation

Vanadium is a potential antidiabetic agent, the use of which is limited by toxicological issues, since several vanadate salts tend to unselectively replace phosphate in many biological reactions. The search for other forms of biological vanadium complexes with either natural compounds or with designed molecules aims at developing novel less toxic and more potent antihyperglycemic agents.

Research objectives

To evaluate the impact of various biogenic amines, including the catecholamines, dopamine, adrenaline and noradrenaline, without and with vanadium, on glucose transport in adipose cells. To decipher the mode of action of the synergism between amines and vanadate regarding glucose transport activation.

Research methods

Research methods included animal husbandry, especially rats and mice deficient in all the β -adrenoceptor subtypes, preparations of freshly isolated adipocytes, quantitative exploration of glucose transport using uptake assays of its non-metabolizable analogue 2-deoxyglucose, together with appropriate use of pharmacological agents.

Research results

In adipose cells, we confirmed the strong stimulatory action of insulin on glucose transport, leading to a tenfold increase over baseline. This was not altered by the addition of 100 $\mu\text{mol/L}$ sodium orthovanadate or clearly reproduced by any of the 25 biogenic amines tested. However, when some amines were added at millimolar doses together with vanadium, they strongly increased glucose uptake up to 70% of the maximal response to insulin. This was the case for methylamine, benzylamine and tyramine, already demonstrated elsewhere to produce hydrogen peroxide when oxidized by monoamine oxidase (MAO) or semicarbazide-sensitive amine oxidase (SSAO) highly expressed in adipocytes. In addition to these amines of reference, known to exert insulin-like effects when combined with vanadate, the catecholamines dopamine, adrenaline and noradrenaline also stimulated glucose transport in a vanadium-dependent manner. Contrarily to reference amines, the stimulation by catecholamines was resistant to MAO and SSAO inhibition. Not all the tested α - and β -adrenergic agonists displayed a clear-cut stimulation of glucose uptake, and the effects of catecholamines were not inhibited by dopaminergic or adrenergic antagonists. These latter effects were even detected in mice genetically invalidated for β -adrenergic receptors. Only antioxidants, such as ascorbate, impaired the stimulation of glucose uptake by the combination of catecholamines plus vanadate.

Research conclusions

It is likely an interaction between vanadium and catecholamine autoxidation that generates intermediates activating in a ROS-dependent manner glucose transport in adipose cells.

Research perspectives

The observed synergism provides the basis for possible future research of novel vanadium/amine complexes with antidiabetic properties.

ACKNOWLEDGEMENTS

We thank the staff of animal unit CREFRE, currently headed by Xavier Collet, and especially its Rangueil satellite for housing wild type and transgenic rodents, and Anne Bouloumié for helpful discussions.

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