Mechanisms of the antidiabetic effects of the \( \beta_3 \)-adrenergic agonist CL-316243 in obese Zucker-ZDF rats

XILIN LIU, FLEURETTE PÉRUSSE, AND LUDWIK J. BUKOWIECKI
Department of Physiology, Faculty of Medicine, Laval University, Quebec City, Canada G1K 7P4

Liu, Xilin, Fleurette Perusse, and Ludwik J. Bukowiecki. Mechanisms of the antidiabetic effects of the \( \beta_3 \)-adrenergic agonist CL-316243 in obese Zucker-ZDF rats. Am. J. Physiol. 274 (Regulatory Integrative Comp. Physiol. 43): R1212–R1219, 1998.—Previous studies have demonstrated that chronic cold exposure activates the sympathetic nervous system, increases energy expenditure, improves glucose tolerance, enhances insulin sensitivity, and stimulates glucose uptake in peripheral tissues (brown and white adipose tissues (BAT and WAT) and muscles) of normal rats. The goal of the present studies was to test whether the selective \( \beta_3 \)-adrenergic agonist CL-316243 (CL) would mimic the beneficial effects of cold exposure in lean and obese ZDF/Gmi-fa male (ZDF) rats, a new model of type II diabetes. In obese ZDF rats, chronic infusion of CL (1 mg·kg\textsuperscript{-1}·day\textsuperscript{-1} for 14 days) significantly decreased body weight gain, food intake, and WAT weight. It also increased total tissue cytochrome oxidase activity, not only in BAT (15 times), but also in WAT (2–4 times), suggesting that it progressively enhanced mitochondrial biogenesis in adipose tissues. CL treatment normalized hyperglycemia and reduced hyperinsulinemia and circulating free fatty acid (FFA) levels. It also improved glucose tolerance and reduced insulin response during an intravenous glucose tolerance test. In general, the beneficial effects of CL were more pronounced in obese than in lean rats. Hyperinsulinemic-euglycemic glucose clamp studies performed with \( [2-\text{H}] \text{deoxyglucose} \) method revealed that CL markedly improved insulin responsiveness in obese rats (3–4 times) and increased glucose uptake in BAT (21 times), WAT (3 times), skeletal muscles (2–3 times), and in the diaphragm (2.8 times), but not in the heart. It is concluded that chronic CL treatment improves glucose tolerance and insulin responsiveness in obese ZDF rats by a mechanism similar to that induced by chronic cold exposure, i.e., by stimulating facultative thermogenesis, mitochondrial biogenesis, and glucose utilization in BAT and WAT. In addition to this mechanism, the reduction in plasma FFA levels induced by chronic CL treatment may further contribute to enhance glucose uptake in skeletal muscles (a tissue that does not express typical \( \beta_3 \)-adrenergceptors) via the "glucose-fatty acid cycle". The anti-obesity and antidiabetic properties of CL suggest that selective \( \beta_3 \)-adrenergic agonists may represent useful agents for the treatment of type II diabetes.

CHRONIC COLD EXPOSURE activates the sympathetic nervous system, increases energy expenditure, improves glucose tolerance, enhances insulin sensitivity, and stimulates glucose uptake in peripheral tissues (brown adipose tissue (BAT), white adipose tissue (WAT), the heart, diaphragm, and skeletal muscles) (44, 52, 54, 55). Cold exposure exerts these antidiabetic effects despite the fact that it decreases plasma insulin levels and increases plasma norepinephrine concentration. The beneficial effects of cold exposure may be partly mimicked by chronic norepinephrine treatment in vivo (33) or by electrical stimulation of the ventromedial hypothalamus (51). In vitro studies have revealed that norepinephrine stimulates glucose uptake in isolated brown adipocytes (even in the absence of extracellular insulin) and potentiates the glucose-stimulatory effects of insulin (35). The stimulatory effects of norepinephrine can be blocked by the nonspecific \( \beta \)-agonist propranolol or by inhibiting mitochondrial fatty acid oxidation with methylpalmitoxor, a specific inhibitor of mitochondrial carnitine acyltransferase (35). This suggests that norepinephrine stimulates glucose uptake in BAT because it stimulates mitochondrial fatty acid oxidation and the glycolytic flux. Glycolysis presumably provides the necessary ATP for activating fatty acids when oxidative phosphorylation is uncoupled by fatty acids bound to the mitochondrial uncoupling protein (for reviews, see Refs. 24, 25).

On the other hand, pharmacological studies revealed that BAT and WAT contain at least three types of \( \beta \)-adrenoceptors (ARs) (for a review, see Ref. 30). Binding studies using hydrophilic radioligands performed on intact brown adipocytes showed that the low-affinity \( \beta_3 \)-ARs are 10 times more abundant than the high-affinity \( \beta_2 \)-ARs, whereas \( \beta_2 \)-ARs appear to be mainly localized in cells other than typical brown adipocytes, possibly in endothelial cells forming the numerous capillaries irrigating BAT (10). Other metabolic studies indicated that norepinephrine, at concentrations usually found in the circulation (1–25 nM), controls both lipolysis and respiration mainly via \( \beta_2 \)-ARs, whereas at much higher levels, presumably occurring in the synaptic cleft after sympathetic stimulation (by cold exposure, diet, stress, etc.), norepinephrine regulates these metabolic processes via both \( \beta_2 \)- and \( \beta_3 \)-adrenergic pathways (3). Until recently, it was generally considered that \( \beta_2 \)-ARs were absent in the heart (containing mainly \( \beta_1 \)-ARs) and in skeletal muscles (containing mainly \( \beta_2 \)-ARs), but it appears that the heart may contain functional \( \beta_2 \)-ARs (15) and/or "atypical" (\( \beta_4 \))-ARs (27). Although the role of these receptors still remains to be defined in different species, the presence of a variety of \( \beta \)-ARs in different tissues opens up the possibility of developing new drugs that specifically activate thermogenesis in adipose tissues and consequently increase glucose utilization, without stimulating the heart or muscles (2, 7).

On this basis, we tested whether the selective \( \beta_3 \)-agonist CL-316243 [disodium(R,R)-5-[2-[2-(3-chlorophenyl)-2-hydroxyethyl]-amino]propyl]-1,3-benzo[d]oxole-2,2-dicarboxylate (CL)] would mimic the beneficial effects of cold exposure (that were mainly studied in normal rats) in lean and obese ZDF/Gmi-fa male (ZDF) rats. The obese ZDF rat is a recently developed model of
obesity and type II diabetes that is characterized by elevated plasma levels of insulin, glucose, triglyceride, and cholesterol (41). In this new model, hyperglycemia can be detected at ~7 wk of age, and obese animals are clearly diabetic (blood glucose of ~25 mM) by 10 wk of age. Initially, the diabetic animals are markedly hyperinsulinemic, but hyperinsulinemia progressively decreases with age. Downregulation of glucose transporters in the pancreas (GLUT-2) and in muscles (GLUT-4) may contribute to the development of diabetic hyperglycemia (14, 47). Because of these characteristics and the consistency in the development of diabetes, the obese ZDF represents an ideal model for investigating the effects of antidiabetic drugs on type II diabetes (49).

In preliminary experiments, we observed that CL did not reduce hyperglycemia in diabetic ZDF rats when given under acute conditions (single intravenous injections or subcutaneous infusions for a few days). Therefore, we tested the long-term effects of CL that was chronically infused via an osmotic minipump during 2 wk. It was found that chronic CL treatment progressively normalizes glycemia, reduces insulinemia, and decreases the levels of circulating free fatty acids (FFA) in obese diabetic ZDF rats. This treatment also markedly improved their glucose and insulin responses during an intravenous glucose tolerance test (IVGTT). Hyperinsulinemic-euglycemic clamps combined with the [2-3H]deoxyglucose ([2-3H]DG) method revealed that chronic CL treatment markedly increases insulin responsiveness in obese rats and that it increases glucose uptake in BAT, WAT, the diaphragm, and skeletal muscles, but not in the heart.

MATERIALS AND METHODS

Animals and treatments. ZDF and their lean littermates were obtained from Genetic Models at the age of 7 wk and were housed in individual cages at 23°C with a 12:12-h light-dark cycle. The rats received Purina chow and water ad libitum and were used 3–5 wk after their arrival.

CL administration. CL was obtained from Lederle Laboratories, American Cyanamid, Pearl River, NY (7). The drug was dissolved under sterile conditions in distilled water containing sodium metabisulfite (0.2 mM) and was administered for 14 days at a dose of 1 mg·kg⁻¹·day⁻¹ via osmotic minipumps (Alza, Palo Alto, CA; model 2002) that were implanted in the back of the animals under isoflurane (Anaquest, Mississauga, ON, Canada) anesthesia. Control animals received the carrier solution (33).

IVGTTs. One week before the IVGTTs and the hyperinsulinemic-euglycemic clamps (see below), two polyethylene extension tubes were connected to the indwelling catheters of the jugular vein (PE-50) and carotid artery (PE-10). A four-way stopcock was used to infuse glucose, insulin, and radioabeled tracers into the jugular vein, whereas the carotid artery was used for blood withdrawal. A first blood sample was taken and analyzed on the glucose analyzer. Then, insulin (100 mU·kg⁻¹·min⁻¹) and glucose (2.78 M) were infused in parallel. The rate of glucose infusion was adjusted to maintain euglycemia (5.3–6.6 mM), and blood glucose concentration was tested at 5-min intervals. Sixty minutes after the initiation of the hyperinsulinemic-euglycemic clamp, [2-3H]DG and [14C]sucrose were intravenously injected for measurements of the rates of glucose uptake in peripheral tissues, as described above.

Statistics. The data were statistically analyzed using either the unpaired t-test or one-way ANOVA followed by the Fisher’s protected least-significant difference post hoc test. Results are expressed as means ± SE.

RESULTS

Effects of CL on body weights, food intake, and tissue weights. Total body weights, daily food intake values, and the weights of several adipose depots were markedly increased in the obese rats (Table 1). However, the individual weights of all six skeletal muscles studied were decreased, whereas the weights of the diaphragm, the heart, and the liver were not significantly different. CL treatment (1 mg·kg⁻¹·day⁻¹ for 14 days) significantly decreased body weight gain much more in obese than in lean rats. It also decreased the mean value of daily food intake in the obese, but not in the lean rats. However, this decrease mainly occurred during the first 2–3 days of the 2-wk treatment period (daily food intake values were not statistically significant between treated and untreated obese rats from day 3 to day 14, not shown). In obese rats, CL increased interscapular BAT weight, but decreased the weights of epididymal and retroperitoneal WAT depots as well as that of the liver. It also slightly increased the mass of several muscles and that of the heart. In lean rats, CL did not significantly affect food intake and body weight gain, but it decreased epididymal and retroperitoneal WAT weight, slightly decreased the weight of the gastrocnemius muscle, and did not affect the weights of other organs.

Effects of CL on the total protein content and cytochrome oxidase activity in BAT and WAT. In previous studies, we found that total cytochrome oxidase activity...
(an index of total tissue mitochondrial content) was markedly decreased in interscapular BAT of obese SHR/N-cp rats, another model of type II diabetes that results from the cp mutation (4, 37). Morphometric stereologic analysis of brown adipocytes in lean and obese SHR/N-cp rats revealed that the total number of mitochondria per brown adipocyte decreased from >4,000 mitochondria per adipocyte in lean rats to ~250 in obese animals (21), resulting in a marked decrease of norepinephrine-stimulated respiration (4, 37). To assess whether BAT oxidative capacity was also reduced in obese ZDF rats, total protein content and cytochrome oxidase activity were measured in several adipose depots. In agreement with our previous observations, it was found that cytochrome oxidase activity was markedly reduced in interscapular BAT of obese ZDF rats (Table 2). Significantly, CL treatment increased cytochrome oxidase activity 5.4 times in interscapular BAT of obese rats, whereas it increased the same parameter by only 2.1 times in lean animals. It also increased total tissue protein content in obese rats. Furthermore, this treatment increased total protein content and cytochrome oxidase activity in the epididymal (2.0 and 4.5 times) and retroperitoneal (2.4 and 2.4 times) WAT depots of obese rats, whereas in lean animals, it only increased cytochrome oxidase activity in retroperitoneal WAT (1.8 times). Thus the defective BAT mitochondrial oxidative capacity in obese diabetic rats is normalized after CL treatment, whereas WAT oxidative capacity appears to be even enhanced compared with controls (Table 2). Recent observations revealed that CL treatment also increases uncoupling protein content or its expression in BAT and WAT of obese ZDF rats (F. D'Allaire and L. J. Bukowiecki, unpublished observations), obese Zucker rats (19), yellow KK obese mice (30), ob/ob mice (1), and diet-induced obese rats (18).

Effect of CL on plasma glucose, insulin, and FFA levels. Untreated obese ZDF rats were markedly diabetic (their plasma glucose levels were nearly 5 times more than those of lean controls). They were also significantly hyperinsulinemic, and their plasma FFA levels were approximately six times greater than those of lean rats (Fig. 1). Two weeks of treatment with CL normalized plasma glucose levels of obese animals to essentially normoglycemic concentrations (6.4 mM). CL also significantly decreased plasma insulin and FFA levels, although not entirely to control values. In contrast, CL treatment slightly decreased the levels of glucose and FFA in lean rats, without significantly affecting plasma insulin levels.

Table 1. Effects of CL treatment on body weight gain, food intake, and tissue weights in lean and obese ZDF rats

<table>
<thead>
<tr>
<th>Tissues</th>
<th>Lean (6)</th>
<th>Lean-CL (6)</th>
<th>Obese (5)</th>
<th>Obese-CL (5)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Body weights before treatments</td>
<td>372.7 ± 9.5</td>
<td>376.7 ± 6.5</td>
<td>454.6 ± 16.4</td>
<td>483.6 ± 5.8§</td>
</tr>
<tr>
<td>Body weight gain</td>
<td>13.3 ± 2.9</td>
<td>7.2 ± 3.3</td>
<td>-7.4 ± 8.0§</td>
<td>-53.4 ± 5.7§</td>
</tr>
<tr>
<td>Daily food intake during the treatments</td>
<td>18.4 ± 0.3</td>
<td>19.5 ± 0.3</td>
<td>33.9 ± 2.2§</td>
<td>27.1 ± 1.8§</td>
</tr>
</tbody>
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Table 2. Effects of CL on protein and cytochrome oxidase content of BAT and WAT depots in lean and obese ZDF rats

<table>
<thead>
<tr>
<th>Tissues</th>
<th>Lean (6)</th>
<th>Lean-CL (6)</th>
<th>Obese (5)</th>
<th>Obese-CL (5)</th>
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</thead>
<tbody>
<tr>
<td>Interscapular BAT</td>
<td></td>
<td></td>
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</tr>
<tr>
<td>Total proteins, mg</td>
<td>16.8 ± 0.7</td>
<td>22.0 ± 3.7</td>
<td>10.3 ± 2.4</td>
<td>27.3 ± 2.2†</td>
</tr>
<tr>
<td>Total cytochrome oxidase content, µmol O₂/min</td>
<td>24.5 ± 0.9</td>
<td>52.0 ± 7.9†</td>
<td>8.7 ± 1.1‡</td>
<td>46.8 ± 1.1‡</td>
</tr>
<tr>
<td>Epididymal WAT</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Total proteins, mg</td>
<td>52.1 ± 8.5</td>
<td>51.2 ± 7.6</td>
<td>25.5 ± 7.4</td>
<td>51.2 ± 1.6*</td>
</tr>
<tr>
<td>Total cytochrome oxidase content, µmol O₂/min</td>
<td>0.32 ± 0.07</td>
<td>0.43 ± 0.08</td>
<td>0.24 ± 0.05</td>
<td>1.07 ± 0.27‡</td>
</tr>
<tr>
<td>Retroperitoneal WAT</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Total proteins, mg</td>
<td>22.0 ± 2.2</td>
<td>15.2 ± 1.5</td>
<td>40.5 ± 12.9</td>
<td>98.5 ± 15.1§</td>
</tr>
<tr>
<td>Total cytochrome oxidase content, µmol O₂/min</td>
<td>0.13 ± 0.02</td>
<td>0.23 ± 0.03*</td>
<td>0.30 ± 0.06§</td>
<td>0.71 ± 0.05§</td>
</tr>
</tbody>
</table>

All values represent means ± SE. Numbers in parentheses indicate number of rats in each group. * and †Significant differences from rats of same phenotype; ‡ and §significant differences from lean rats at P < 0.05 or P < 0.01 levels, respectively.
Effects of CL on the glucose and insulin responses to an IVGTT in lean and obese ZDF rats. CL treatment markedly decreased the glucose response in obese ZDF rats during an IVGTT (Fig. 2). This effect was evident at all time points after the intravenous injection of glucose, as well as when the data were expressed as total surfaces under the glucose curve (Fig. 2, inset). In lean rats, the β3-agonist slightly decreased plasma glucose levels at several, but not all, time points after glucose administration, resulting in a small, statistically nonsignificant, decrease in the total glucose area. Similarly, CL decreased the insulin response, mainly in obese animals (Fig. 3). Thus both the glucose and insulin responses were significantly decreased in treated obese ZDF rats, providing a first indication that CL increases insulin responsiveness in these animals.

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DISCUSSION

The main goal of the present studies was to determine whether treatment of diabetic rats with CL exerts a beneficial effect in a new rat model of type II diabetes.
to investigate, in a second step, the mechanisms of action of this selective β3-agonist. In the first series of experiments, we found that CL, when given for 2 wk to ZDF diabetic rats, normalizes their plasma glucose levels and significantly decreases their plasma insulin and FFA concentrations (Fig. 1). These observations agree with previous findings, showing that β3-agonists display antidiabetic effects in various animal models of type II diabetes (1, 48). In the present experiments, CL treatment improved not only basal glycemia, but also glucose tolerance (Fig. 2). During the entire IVGTT, both plasma glucose and insulin concentrations of treated rats remained decreased in comparison with untreated animals, providing a first indication that CL improves insulin responsiveness in peripheral tissues.

This conclusion was supported by the hyperinsulinemic-euglycemic clamp experiments (Figs. 4–6). They revealed that CL markedly increased insulin responsiveness: nearly four times more glucose had to be infused in CL-treated than in untreated obese animals to achieve euglycemia. It was very difficult in the clamp experiments to significantly reduce the elevated glycemia of untreated obese rats without increasing plasma insulin concentrations to levels close to the insulin V_max for net glucose utilization (28). This situation is rather different from what happens with other rat models of obesity that generally develop milder forms of insulin resistance not associated with frank diabetes. Nevertheless, the fact that chronic CL treatment normalized plasma glucose levels in ~2 wk (Figs. 1–3) is remarkable in view of the marked insulin resistance of obese ZDF rats.

The maximal capacity of various tissues for glucose uptake in CL-treated animals varied with the following order: BAT > heart > diaphragm > skeletal muscles > WAT. This sequence of potencies agrees with previous observations made with normal rats treated with insulin (54) or norepinephrine (33) as well as with cold-exposed animals (55). It shows that BAT possesses a remarkable capacity for glucose utilization, either for storing it in the form of triglycerides or for oxidizing it for thermogenesis (in cold-exposed animals or in warm-exposed animals treated with norepinephrine or β3-agonists) (5, 24, 33, 55). In vitro studies confirmed that both insulin and norepinephrine stimulate glucose uptake and revealed that a very low (nanomolar) norepinephrine concentration potentiates the glucose stimulatory effects of insulin (35). This suggests that norepinephrine facilitates glucose entry into brown adipocytes by stimulating thermogenesis, fatty acid oxidation, and glycolysis. It is likely that CL acts in a fashion similar to that of norepinephrine, with the exception that this selective β3-agonist is not expected to activate β1-adrenergic pathways. However, maximal thermogenesis can be achieved in isolated brown adipocytes by stimulating them either with β1- or β3-agonists (3).

The mechanism by which CL improves insulin responsiveness and glucose uptake by peripheral tissues appears to be similar to that occurring during cold acclimation of warm-acclimated rats. Similar to long-term CL treatment, chronic cold exposure (4–5°C) improves glucose tolerance, increases insulin sensitiv-
ity, stimulates glucose uptake in peripheral tissues (BAT, WAT, skeletal muscles, and heart), and reverses the diabetogenic effects of high-fat feeding (44, 52–55). It exerts these beneficial effects despite the fact that it decreases plasma insulin concentration and increases norepinephrine levels. Chronic cold exposure also stimulates mitochondrial proliferation and mitochondrial uncoupling protein content (4–5°C) (16, 17, 22, 25). Furthermore, in CL-treated animals (Figs. 5 and 6), as in cold-exposed rats (54, 55), glucose uptake is increased much more in BAT (by 1–2 orders of magnitude) than in WAT or muscles. This order of potency agrees with the fact that BAT capacity for nonshivering thermogenesis is much higher than that of WAT or muscles. Cold exposure stimulates the release of norepinephrine from sympathetic nerves and consequently increases nonshivering thermogenesis in BAT and WAT, not only via β3-ARs, but also via β1-adrenergic pathways (β2-ARs are undetectable in brown adipocytes) (10, 24, 30, 31). Although the affinity of β1-ARs for norepinephrine is much higher than that of β2-ARs, the latter are ~10 times more numerous and appear to be resistant to catecholamine-induced desensitization or downregulation (9, 10, 23, 30, 39, 40). In muscles, norepinephrine may stimulate nonshivering thermogenesis and glucose uptake via β2-ARs and/or atypical (β2j)-ARs (12, 27, 34, 45, 54). Thus it is likely that CL mimics the β3-effects of norepinephrine in BAT and WAT, but its stimulatory effects in muscles are probably indirect because this tissue lacks typical β3-ARs. Most probably, the decrease in FFA levels induced by CL treatment (Fig. 1) enhances glucose uptake in muscles via the so-called Randle’s effect or glucose-fatty acid cycle (42). However, this remains to be directly demonstrated.

A question that is often raised is how a potent lipolytic agent, such as CL, decreases the levels of circulating fatty acids instead of increasing them. We believe that this paradox is merely apparent because, in addition to stimulating lipolysis, β3-agonists markedly stimulate thermogenesis, particularly in BAT (3, 18, 19, 26, 46). Fatty acids represent the principal substrates used for thermogenesis by BAT (50), and activation of nonshivering thermogenesis by norepinephrine or β3-agonists stimulates fatty acid oxidation, decreases the circulating levels of fatty acids, and diminishes triglyceride stores in adipose tissues. Using an oxygen consumption system for continuously monitoring daily oxygen consumption in rats (43), we recently found that CL does in fact stimulate 23-h oxygen consumption in obese ZDF rats when infused during 14 days under the same experimental conditions as in the present experiments (11). Remarkably, the enhancement of oxygen consumption progressively increased from ~10% above basal values during the first 4 days of the infusion to 35% during the last 4 days (days 10–14). This progressive increase was accompanied by a parallel increase in total tissue cytochrome oxidase activity and uncoupling protein content, not only in BAT, but also in several WAT depots, confirming previous observations (26). We therefore hypothesize that CL exerts its beneficial action by 1) acutely enhancing thermogenesis in BAT, WAT, and possibly also other tissues and 2) by restoring to normal the defective BAT thermogenic capacity, as evidenced by its remarkable effect on total tissue cytochrome oxidase activity (Table 2). The fact that CL did not reduce hyperglycemia in diabetic ZDF rats at short term (single injection or infusion for <1 wk) strongly suggests that the β3-agonist acts by progressively stimulating mitochondrial biogenesis in BAT and restoring to normal the defective thermogenic capacity of obese rats (4, 36, 37). Another explanation for the decreased FFA levels in obese rats chronically treated with CL could be based on the increased insulin responsiveness of adipose tissues. Insulin is a potent antilipolytic hormone, and the increased insulin responsiveness induced by CL treatment may contribute to decreased lipolysis and plasma FFA levels.

The observation that CL increased glucose uptake ~10 times more in BAT than in WAT (per gram of tissue) (Fig. 5) agrees with the observations that BAT possesses a much higher capacity for heat production than WAT (Table 2, cytochrome oxidase data). Quantitatively, WAT may still represent a significant site of glucose and fat oxidation, because it is much more abundant than BAT, although it is difficult to precisely estimate the relative proportion of WAT versus BAT, particularly in obese rats. Nevertheless, the muscles remain the main anatomic site of glucose uptake in CL-treated rats under the clamp conditions described in Figs. 3–5. On the assumption that the muscles, WAT, and BAT, respectively, represent 30, 40, and 1% of the body weight of obese rats and by averaging the individual glucose uptake values for the different types of muscles and fat depots investigated, it can roughly be estimated that total BAT and WAT combined represent ~20% of glucose uptake in muscles (BAT 10% and WAT 10%).

In summary, it is concluded that chronic, but not short-term, CL treatment normalizes glycemia and increases insulin responsiveness and glucose uptake in adipose tissues and muscles, but not in the heart, of obese ZDF rats. It is suggested that the β3-agonist progressively increases the defective mitochondrial oxidative capacity in BAT and WAT of diabetic animals, thereby increasing energy expenditure and fat oxidation, and, consequently, reducing plasma FFA levels. This may lead to an enhancement of glucose utilization by skeletal muscles via the glucose fatty acid cycle. Although human β3-ARs appear to be different from rat β3-ARs, selective β3-agonists such as CL-316243 may represent useful agents for the treatment of obesity and type II diabetes (6, 32).

Perspectives

When we started to investigate the effects of cold exposure on glucose metabolism a few years ago, we had no idea that one day catecholamines or β-adrenergic agonists would represent potential agents for normalizing plasma glucose levels in type II diabetes. At that time, catecholamines, glucagon, and other lipolytic hormones were considered counterregulatory hormones, because they inhibited, generally at high phar-
macological doses, the beneficial effects of insulin. It took several years to demonstrate that the main effec-
tor of the stimulation of glucose uptake by peripheral
tissues in cold-exposed animals was norepinephrine
secreted from sympathetic nerves. The idea that physi-
ological conditions (cold exposure, exercise) or drugs
(adrenergic agonists) activating mitochondrial ATP
synthesis or heat production also stimulate glucose utili-
za tion in muscles and adipose tissues has opened and still
opens new avenues for developing more efficient and
more selective antidiabetic drugs. In recent years,
adipose tissues (brown or white) have been the focus of
search opportunities for developing new drugs facilitat-
ing glucose uptake directly in the skeletal muscles that
represent the main anatomic sites of glucose utilization
in humans as well as in laboratory animals.

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Address for reprint requests: L. J. Bukowiecki, Dept. of Physiology, Faculty of Medicine, Laval Univ., Québec City, Canada G1K 7P4.

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