Effect of the glucagon-like peptide-1 receptor agonist lixisenatide on postprandial hepatic glucose metabolism in the conscious dog

Mary Courtney Moore,1 Ulrich Werner,2 Marta S. Smith,1 Tiffany D. Farmer,1 and Alan D. Cherrington1

1Department of Molecular Physiology and Biophysics, Vanderbilt University School of Medicine, Nashville, Tennessee, and 2Sanofi Deutschland, Frankfurt, Germany

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Moore MC, Werner U, Smith MS, Farmer TD, Cherrington AD. Effect of the glucagon-like peptide-1 receptor agonist lixisenatide on postprandial hepatic glucose metabolism in the conscious dog. Am J Physiol Endocrinol Metab 305: E1473–E1482, 2013. First published October 22, 2013; doi:10.1152/ajpendo.00354.2013.—The impact of the GLP-1 receptor agonist lixisenatide on postprandial glucose disposition was examined in conscious dogs to identify mechanisms for its improvement of meal tolerance in humans and examine the tissue disposition of meal-derived carbohydrate. Catheterization for measurement of hepatic balance occurred $\approx 16$ days before study. After being fasted overnight, dogs received a subcutaneous injection of 1.5 $\mu$g/kg lixisenatide or vehicle (saline; control; $n = 6/group$). Thirty minutes later, they received an oral meal feeding (93.4 kJ; 19% protein, 71% glucose polymers, and 10% lipid). Acetaminophen was included in the meal in four control and five lixisenatide dogs for assessment of gastric emptying. Observations continued for 510 min; absorption was incomplete in lixisenatide at that point. The plasma acetaminophen area under the curve (AUC) in lixisenatide was 65% of that in control ($P < 0.05$). Absorption of the meal began within 15 min in control but was delayed until $\approx 30–45$ min in lixisenatide. Lixisenatide reduced ($P < 0.05$) the postprandial arterial glucose AUC $\approx 54\%$ and insulin AUC $\approx 44\%$. Net hepatic glucose uptake did not differ significantly between groups. Nonhepatic glucose uptake tended to be reduced by lixisenatide (6,151 ± 4,321 and 10,541 ± 1,854 $\mu$mol·kg$^{-1}$·min$^{-1}$ in lixisenatide and control, respectively; $P = 0.09$), but adjusted (for glucose and insulin concentrations) values did not differ (18.9 ± 3.8 and 19.6 ± 7.9 $\mu$mol·kg$^{-1}$·min$^{-1}$–1, lixisenatide and control, respectively; $P = 0.94$). Thus, lixisenatide delays gastric emptying, allowing more efficient disposal of the carbohydrate in the feeding without increasing liver glucose disposal. Lixisenatide could prove to be a valuable adjunct in treatment of postprandial hyperglycemia in impaired glucose tolerance or type 2 diabetes.

type 2 diabetes; hepatic glucose metabolism; mixed meal

THE INCRETIN HORMONE GLUCAGON-LIKE PEPTIDE-1 (GLP-1) has a number of beneficial direct and indirect effects on carbohydrate metabolism, including increasing insulin secretion in the presence of hyperglycemia, suppressing glucagon secretion, delaying gastric emptying, and stimulating glucose disposal by the liver and skeletal muscle (reviewed in Ref. 4). GLP-1 is released in response to meal ingestion, and therefore, it might be anticipated that it would have important therapeutic implications for patients with impaired glucose tolerance or type 2 diabetes. However, the biological half-life of GLP-1 is only $\sim 2$ min due to its rapid degradation by dipeptidyl peptidase IV (DPP-IV), and this limits the practicality of treatment with GLP-1 itself (26). Lixisenatide, a novel once daily prandial GLP-1 receptor agonist developed to provide the benefits of GLP-1 in a longer lasting formulation (12, 43, 60), was granted marketing authorization by the European Medicines Agency in February 2013 (16). It was also approved for use in Mexico, Australia, and Japan. It is considered a prandial agent because of its substantial effect in lowering prandial glucose concentrations and improving Hb A1C levels (43, 48, 50). Its very high affinity for the GLP-1 receptor (4–5 times that of GLP-1) has led to its being developed for once a day administration despite its relatively short half-life (2–4 h) (5). In dose-ranging studies, once daily treatment was observed to provide a better ratio of efficacy to tolerability than twice-daily dosing (43).

Chronic (12 wk) treatment of obese prediabetic Zucker diabetic fatty (ZDF) rats with lixisenatide reduced basal glucose concentrations and improved oral glucose tolerance (61). In lean glucose-tolerant ZDF rats, on the other hand, lixisenatide treatment did not alter blood glucose (61). Acute treatment of conscious dogs with lixisenatide at doses of 0.15–5 $\mu$g/kg prior to an oral glucose tolerance test significantly reduced the maximum glucose response (49–73%) and the plasma insulin concentrations (60–75%) (60).

The effect of 12 wk of lixisenatide monotherapy on mixed-meal disposition has been reported in humans with type 2 diabetes (17). Lixisenatide treatment reduced the 2-h postprandial glucose concentrations seven- to eightfold compared with placebo and was associated with significant improvement in Hb A1C (doubling the number of subjects with Hb A1C $< 7\%$). Recent investigations have confirmed the benefit of lixisenatide in reducing postprandial glucose levels in individuals with type 2 diabetes following the consumption of a standard meal (1, 17, 39, 46, 47, 50, 56).

GLP-1 is an insulin secretagogue, and lixisenatide is fully active at the GLP-1 receptor. It has been shown to preserve $\beta$-cell mass and insulin mRNA expression in the db/db mouse and to preserve both the first- and second-phase insulin response in the ZDF rat (62). It also augmented the first-phase insulin response to an intravenous glucose challenge in nondiabetic humans (6) and improved $\beta$-cell function assessed by homeostasis model assessment-B in a 24-wk study of humans with type 2 diabetes (1). Nevertheless, the reduction in 2-h postprandial glucose concentrations with lixisenatide treatment in humans was associated with a reduction in 2-h postprandial insulin concentrations as well (13, 51). Thus, consistent with the results from humans treated with GLP-1 (28, 33), the effect of lixisenatide in improvement of glucose tolerance does not seem to be attributable simply to enhanced insulin secretion.

The liver plays an extremely important part in the disposal of carbohydrate from a glucose load or meal (11, 14), but it is difficult to quantify the role of the liver in glucose disposal in the human under physiological conditions because of the in-
vasiveness of the catheterization required. Recent data indicate
that lixisenatide delays gastric emptying (27), and Woerle et al. (64)
reported that the splanchic bed disposed of more of the
carbohydrate from a mixed meal when gastric emptying in
humans was delayed by pramlintide administration. Lixisen-
late’s impact on the relative roles of the liver and extrahepatic
tissues in glucose disposal have not been examined under
physiological conditions. For this reason, the current studies
were carried out to examine the effect of lixisenatide on the
disposition of a mixed meal in the conscious dog, a model in
which it is possible to quantify hepatic balance precisely.

METHODS

Animals and experimental preparation. The protocol was approved
by the Vanderbilt University Institutional Animal Care and Use
Committee, and the animals were housed and cared for according to
Association for Assessment and Accreditation of Laboratory Animal
Care guidelines. The studies were carried out in conscious overnight-
fasted male or female mongrel dogs (20.1–26.4 kg) that were fed once
daily a diet of meat and chow providing 31% protein, 52% carbohy-
brate, 11% fat, and 6% fiber based on dry weight. Approximately 16
days before study, each dog underwent a laparotomy for placement of
ultrasonic flow probes (Transonic Systems, Ithaca, NY) around the
portal vein and the hepatic artery as well as insertion of silicone
rubber catheters for sampling in a hepatic vein, the portal vein, and a
femoral artery, as described in detail elsewhere (31). The animals
were studied only if they met established criteria prior to study (31).
On the morning of the study, catheters and flow probe leads were
exteriorized from their subcutaneous pockets under local anesthesia
(31). An angiocath (Deseret Medical, Sandy, UT) was inserted into a
cephalic vein for infusion of indocyanine green dye.

Experimental design. Each experiment consisted of a 60-min equil-
ibration period (−90 to −30 min), a 30-min basal period (−30 to 0
min), and a 510-min experimental period (0–510 min). At −90 min,
a continuous infusion of indocyanine green dye (0.08 mg/min; Sigma,
St. Louis, MO) was begun in all dogs. At −30 min, seven dogs
(lixisenatide group) received a subcutaneous injection of lixisenatide
1.5 μg/kg (Sanofi, Paris, France), and seven dogs (control group)
received a subcutaneous vehicle (0.9% saline) injection (Fig. 1). At 0
min the study timer was stopped, and a liquid mixed meal [93.4 kJ/kg;
energy supplied as 19% protein (BeneProtein, Novartis Medical Nu-
trition, Minneapolis, MN), 71% glucose polymers (Polycose; Abbott
Nutrition, Columbus, OH), and 10% lipid (Microlipid; Nestlé Health-
care, Minnetonka, MN)] was drunk by the dogs as it was gradually
injected into the mouth with a syringe. Five-hundred milligrams of
acetaminophen (paracetamol; Sigma) was added to the meal for the
last four of the control dogs and the last five of the lixisenatide dogs
to assess gastric emptying. Ingestion of the meal took no longer than
3 min, and the timer was restarted as soon as meal intake was
complete.

Samples were taken from the femoral artery, hepatic portal vein,
and hepatic vein during the 30 min before the meal and every 15–30
min during the 510-min postprandial period to assess the gastrointes-
tinal, hepatic, and nonhepatic disposition of the meal. One dog in each
group vomited within 1 h after the meal was delivered. These dogs
were deleted from the database, and thus the final groups contained six
dogs each. At the end of study, the dogs were euthanized with an
overdose of anesthetic, a laparotomy was performed, and the stomach
was opened and drained for measurement and visual examination of
the gastric fluid.

Hematocrit, blood lactate, alanine, and glyceral and plasma glu-
cose, insulin, glucagon, and cortisol were measured as described
previously (31). Plasma acetaminophen concentrations were measured
by HPLC following the addition of 2-acetaminophenol as the internal
standard and deproteination of the plasma with Ba(OH)2 and ZnSO4
by a modification of the methods of Ameer et al. (3) and O’Connell
and Zurzola (36). Plasma GLP-1 was measured with an ELISA
method that specifically quantifies the active forms of GLP-1 (Linco
Research, St. Charles, MO).

Calculations and data analysis. Hepatic blood flow was measured
using ultrasonic probes and indocyanine green extraction. The two
methods yielded similar results, but the data reported here were
analyzed with the ultrasonic-determined flows because their mea-
surement did not require an assumption regarding the relative contri-
bution of arterial and portal flow to total hepatic blood flow. The rate
of glucose delivery to the liver, or hepatic glucose load, was calcu-
lated as: loadin = (Fa × Sh) + (Fp × Sp), where Fa and Fp represent
blood or plasma flow (as appropriate for the substrate under exami-
nation) in the artery and hepatic portal vein, respectively, and Sh and
Sp represent the blood or plasma substrate concentrations in the two
vessels. Net hepatic substrate balance (NHB) was calculated as loadout
− loadin, where loadout = Fa × Sh, Fa is the total hepatic blood or
 plasma flow, and Sp is the substrate concentration in the hepatic vein.
Net hepatic fractional substrate extraction was calculated as net
hepatic substrate balance/loadin. Net gut glucose clearance was calcu-
lated as (Sa − Sp) × Fg. Nonhepatic glucose uptake was net gut
 glucose output minus net hepatic glucose uptake (NHBG), which was
adjusted for changes in the mass of the glucose pool. Glucose
 clearance was calculated as hepatic or nonhepatic glucose uptake,
as appropriate, divided by the arterial glucose concentration. Net
hepatic carbon retention, shown previously to be a close estimate of hepatic
glycogen synthesis (53), was calculated as NHGU minus net hepatic
lactate output (in glucose equivalents).

The area under the curve (AUC) was calculated for the postprandial
period (0–510 min) using the trapezoidal rule, with the AUCs expre-
ssed as change from the basal concentrations or rates (ΔAUC),
unless otherwise indicated.

SigmaStat (SPSS, Chicago, IL) was used for statistical analysis.
Time course data were evaluated with repeated-measures ANOVA.
Where significant differences between groups in their responses over
time were found, Tukey’s test was utilized for post hoc analysis to
identify the time points where the groups differed. Unpaired Student’s
t-test was used for comparison of AUCs.

RESULTS

Acetaminophen concentrations. In the four control and five
lixisenatide dogs that received acetaminophen to assess gastric
emptying (Fig. 2), the peak plasma concentration in the lix-
isenatide group was only 53% of that in the control dogs (P <
0.05), whereas the overall ΔAUC in the lixisenatide dogs was
65% of that in the control group (924 ± 228 vs. 1,412 ± 240
ng·ml⁻¹·510 min⁻¹, P < 0.05).

Glucose metabolism. The basal arterial and portal blood
glucose concentrations were ~4.6 mmol/l in both groups
(Fig. 3). In the control group, the arterial glucose concentra-

Fig. 1. Timeline of procedures. A continuous peripheral venous infusion of
indocyanine green (ICG) began at −90 min, followed by drug or vehicle (0.9%
saline) injection at −30 min. The meal was given as soon as the 0-min baseline
blood sampling was complete.
The peak rates of net gut glucose output were 59.9 mmol/kg observed in the lixisenatide group was 7.2 mmol/l above basal at the end of study. The mean arterial blood glucose concentration remained elevated above basal for 4 h after the meal. Because of this, the ΔAUC for hepatic glucose load in the lixisenatide group was only 28% smaller than in the control group (P = 0.1). The control group switched from net hepatic glucose output to NHGU within 30 min after the meal, but the shift to NHGU was delayed until 90 min in the lixisenatide group, and the rate remained significantly lower in lixisenatide through 300 min (Fig. 4). NHGU ceased by ~160 min in the control group but was ongoing at the end of study in the lixisenatide group, so the ΔAUC of NHGU was not different in lixisenatide or control (P = 0.24). Net hepatic fractional extraction of glucose and hepatic glucose clearance followed a pattern similar to that of NHGU. The AUC of net hepatic carbon retention did not differ between groups (P = 0.24; Fig. 4).

The assimilation of the meal was incomplete at the end of study in the lixisenatide group, as can be seen from the acetaminophen concentrations and net gut glucose balance. The acetaminophen concentrations provide an independent tool for estimating the time remaining for meal assimilation. If the rate of change of acetaminophen concentrations over the last hour of study is projected forward until the concentrations fall to zero in the lixisenatide group, it can be estimated that gastric release of the acetaminophen (and presumably the meal) would have continued for ~4 h past the end of the study. If the mean rate of NHGU observed between groups over the last hour of study had continued for an additional 4 h, then the AUC of NHGU in the lixisenatide group would total 10.3 ± 3.3 mmol/kg, remarkably similar to the AUC of 10.7 ± 1.1 mmol/kg observed in the control group.

Nonhepatic glucose uptake was significantly greater in the control group than in the lixisenatide group for >3 h after the meal (Fig. 5). The ΔAUC of nonhepatic glucose uptake for the postprandial period was reduced >40% in lixisenatide (10,541 ± 1,854 and 6,151 ± 4,321 mmol/kg·h, P = 0.09). Since the glycemic concentrations differed, nonhepatic glucose clearance was calculated (Fig. 5). Although there was a significant group × time effect of lixisenatide on nonhepatic glucose clearance, the ΔAUC did not differ between groups (P = 0.3). When nonhepatic glucose clearance was divided by the insulin concentrations, neither the individual time points nor the AUCs differed (P = 0.9 for comparison of AUCs; Fig. 5).

Hormonal response. The arterial plasma insulin response in the lixisenatide group was 44% smaller than in the control group (ΔAUC 51,813 ± 9,735 vs. 87,088 ± 18,875 pmol·l⁻¹·min⁻¹, P < 0.05), and the hepatic sinusoidal insulin response also tended to be blunted (37% less in lixisenatide than control), as shown in Fig. 6. Arterial plasma glucose concentrations did not differ in the two groups and remained nearly basal throughout (Fig. 6). Arterial and portal vein GLP-1 concentrations demonstrated a robust increase in the lixisenatide group, but it continued until the end of the study in the lixisenatide group. As a result, the ΔAUC of gut glucose output did not differ significantly between groups (P = 0.2).

Because of the overall reduction in glycemia, the hepatic glucose load was lower in the lixisenatide group than in the control group for ~5 h after the meal (P < 0.05; Fig. 4). The hepatic load had returned to basal in the control group by the end of study, whereas it remained significantly elevated in the lixisenatide group. Because of this, the ΔAUC for hepatic glucose load in the lixisenatide group was only 28% smaller than in the control group (P = 0.1). The control group switched from net hepatic glucose output to NHGU within 30 min after the meal, but the shift to NHGU was delayed until 90 min in the lixisenatide group, and the rate remained significantly lower in lixisenatide through 300 min (Fig. 4). NHGU ceased by ~160 min in the control group but was ongoing at the end of study in the lixisenatide group, so the ΔAUC of NHGU was not different in lixisenatide or control (P = 0.24). Net hepatic fractional extraction of glucose and hepatic glucose clearance followed a pattern similar to that of NHGU. The AUC of net hepatic carbon retention did not differ between groups (P = 0.24; Fig. 4).

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Metabolite concentrations and balance. During the first half-hour of the postprandial period, the arterial concentrations of nonesterified fatty acids (NEFA) were significantly higher in the lixisenatide vs. the control group (Table 1). Although the NEFA concentrations in the lixisenatide group remained nu-

![Graph](image-url)
Arterial and portal vein blood glucose, the arterial-portal (A-P) glucose gradient, and net gut glucose balance in the control and lixisenatide groups; n = 6/group. Data are means ± SE. There was a significant (P < 0.05) group × time interaction for all time course data; *data points that differed on post hoc analysis. The histograms show the areas under the curve (AUCs) for the postprandial period; †P < 0.05 between groups using an unpaired t-test.

Arterial blood glycerol concentrations in the basal state were lower in the lixisenatide group than in the control group, but the early postprandial concentrations did not differ between groups (Table 1). As a result, the decline in arterial glycerol concentrations between the basal period and the 30-min postprandial sample was three times as large in the control group as in the lixisenatide group (P < 0.05). During the last 150 min of study, glycerol concentrations in both groups began to return toward basal. During this time, the concentrations in the control group again were significantly greater than those in the lixisenatide group. Net hepatic glycerol uptake did not differ between groups (P = 0.3).

Arterial blood lactate and alanine concentrations (Table 1) were significantly greater in the control vs. lixisenatide group during the postprandial period (P < 0.05), but net hepatic lactate output and alanine uptake, as well as net gut alanine output, did not differ significantly between groups (P = 0.2, 0.4, and 0.2, respectively). Neither the concentrations nor the net hepatic balances of glutamine and glutamate differed significantly between groups (data not shown).
Fig. 4. Hepatic glucose load (A), net hepatic glucose uptake (B), net hepatic fractional glucose extraction (C), hepatic glucose clearance (D), and net hepatic carbon retention (E) in the control and lixisenatide groups. There was a significant \( P < 0.05 \) group × time interaction for all parameters. *Data points that differed on post hoc analysis. There were no significant differences in postprandial AUCs between groups (histograms); \( n = 6 \)/group. Data are means ± SE.
Meal histograms show the AUC0–510 min for nonhepatic glucose uptake. *Data points that differed on post hoc analysis. The formula volume was with what appeared to be curdled formula. The formula volume exceeded that in the lixisenatide group, all stomachs contained bile-stained fluid. In the control dogs, only a small amount (5–10 ml) of clear fluid was visible in the stomach at the end of study, whereas a placebo-treated group exhibited no decrease in the stomach at the end of study, are also consistent with a delay in nutrient assimilation in the lixisenatide group. Indeed, although measurement of the stomach contents can provide only a rough estimate of the amount of the diet in the stomach at the end of study, the post-study volume of intragastric formula in the lixisenatide group was approximately equivalent to the amount that remained unabsorbed based on the net gut glucose output data. The reduction in the rate of gastric emptying with lixisenatide treatment is consistent with results from [13C]octanoic acid breath tests in humans with type 2 diabetes. The tendency toward blunting of the net gut output of alanine and the significant reduction in the arterial concentrations of alanine, as well as the presence of some of the meal in the stomach at the end of study, are also consistent with a delay in nutrient assimilation in the lixisenatide group. Indeed, although measurement of the stomach contents can provide only a rough estimate of the amount of the diet in the stomach at the end of study, the post-study volume of intragastric formula in the lixisenatide-treated dogs was approximately equivalent to the amount that remained unabsorbed based on the net gut glucose output data. The reduction in the rate of gastric emptying with lixisenatide treatment is consistent with results from [13C]octanoic acid breath tests in humans with type 2 diabetes. After 4 wk of daily treatment, the gastric emptying rate for a labeled breakfast meal decreased 85% compared with the pretreatment measurement in the lixisenatide-treated group, whereas a placebo-treated group exhibited no decrease in
gastric emptying (27). GLP-1 may alter gastric emptying by both reducing antral activity and stimulating pyloric contraction (54, 63). The delay of gastric emptying by GLP-1, GLP-1 receptor agonists, or DPP-IV inhibitors apparently involves central neural mediation (15, 24, 40, 52), although whether or not the vagal afferent innervation is required for the response is unclear (24, 32).

Similar to NHGU, nonhepatic glucose uptake was significantly greater in the control group than in the lixisenatide group for more than 3 h after meal ingestion, but nonhepatic glucose uptake continued at a substantial rate in the lixisenatide group at the end of the experiments. Although the AUC of nonhepatic glucose uptake tended to be reduced in the lixisenatide group, when the rate was adjusted for the differences in glycemia (i.e., nonhepatic glucose clearance), the result was more similar between groups. When nonhepatic glucose clearance was further adjusted for the prevailing insulin concentrations, no differences were evident between groups. Thus, there was no evidence for either a specific hepatic or a nonhepatic effect of lixisenatide.

Glycemic levels during the first half of the postmeal period in the current studies were significantly blunted in the dogs receiving lixisenatide. This is particularly relevant, given that the 2-h postload glucose concentration during an OGTT has been found to be predictive of both the progression of prediabetic individuals to diabetes and the risk of cardiovascular disease in type 2 diabetes (8, 10, 20, 35). Hb A1c is heavily influenced by postprandial glycemic levels (21).

In obese prediabetic ZDF rats, 12 wk of treatment with lixisenatide vs placebo reduced Hb A1c significantly (1.7%) (61). Consistent with this, 12 wk of lixisenatide monotherapy in adults with type 2 diabetes significantly decreased 2-h postprandial plasma glucose concentrations and excursions (change from basal concentrations) after a standardized breakfast meal, reduced fasting plasma glucose, and doubled the number of subjects with Hb A1c concentrations <7% compared with placebo treatment (17). Subsequent randomized clinical investigations in adults with type 2 diabetes inadequately controlled with metformin (1), basal insulin with or without metformin (46, 47), basal insulin with or without a sulfonylurea (56), or pioglitazone (39) demonstrated that the addition of lixisenatide once daily for periods ranging from 13 to 24 wk was effective in reducing postprandial plasma glucose after a standardized test meal and lowering Hb A1c relative to placebo (43). Given the relationship between elevations in Hb A1c and cardiovascular disease and other complications of diabetes (7, 57),
higher insulin levels might have been observed in dogs if lixisenatide had been administered. Relative to placebo, this finding is consistent with the idea that individual responses to GLP-1 cannot be predicted. This is particularly true in subjects with type 2 diabetes, where there was no effect of 10 days of treatment with lixisenatide. However, in subjects with type 2 diabetes, the GLP-1 receptor agonist, exhibits a similar postprandial insulinemic response as the 0- to 4-h postprandial insulin AUC in a dose-dependent manner. Using the specific GLP-1 receptor antagonist exendin (9–39), Shirra et al. (55) determined that GLP-1’s reduction of fasting and postprandial glycemia. However, in the current acute study, there was only a small effect, if any, of lixisenatide on insulin concentrations following a mixed meal (59).

Table 1. NEFA, glycerol, lactate, and alanine concentrations and balances

<table>
<thead>
<tr>
<th>Group and Parameter</th>
<th>Basal Period</th>
<th>Postprandial Period, min</th>
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<tbody>
<tr>
<td></td>
<td>30</td>
<td>60</td>
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<tr>
<td>Arterial plasma NEFA, (\mu\text{mol}l^{-1})</td>
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<tr>
<td>Control</td>
<td>283 ± 44</td>
<td>150 ± 35</td>
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<tr>
<td>Lixisenatide</td>
<td>558 ± 153†</td>
<td>443 ± 156</td>
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<td>Arterial blood glycerol, (\mu\text{mol}l^{-1})</td>
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<tr>
<td>Control</td>
<td>26 ± 6</td>
<td>21 ± 5</td>
</tr>
<tr>
<td>Lixisenatide</td>
<td>40 ± 10</td>
<td>30 ± 10</td>
</tr>
<tr>
<td>Net hepatic glycerol uptake, (\mu\text{mol}kg^{-1}\text{min}^{-1})</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Control</td>
<td>0.5 ± 0.2</td>
<td>0.3 ± 0.1</td>
</tr>
<tr>
<td>Lixisenatide</td>
<td>1.2 ± 0.3</td>
<td>0.6 ± 0.2</td>
</tr>
<tr>
<td>Arterial blood lactate, (\mu\text{mol}l^{-1})</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Control</td>
<td>784 ± 178</td>
<td>1,122 ± 124</td>
</tr>
<tr>
<td>Lixisenatide</td>
<td>663 ± 158†</td>
<td>612 ± 121</td>
</tr>
<tr>
<td>Net hepatic lactate output, (\mu\text{mol}kg^{-1}\text{min}^{-1})</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Control</td>
<td>16.5 ± 3.8</td>
<td>18.6 ± 5.0</td>
</tr>
<tr>
<td>Lixisenatide</td>
<td>9.9 ± 5.8</td>
<td>10.6 ± 5.2</td>
</tr>
<tr>
<td>Arterial blood alanine, (\mu\text{mol}l^{-1})</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Control</td>
<td>417 ± 45</td>
<td>428 ± 46</td>
</tr>
<tr>
<td>Lixisenatide</td>
<td>346 ± 46†</td>
<td>311 ± 28†</td>
</tr>
<tr>
<td>Net gut alanine output, (\mu\text{mol}kg^{-1}\text{min}^{-1})</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Control</td>
<td>1.7 ± 0.5</td>
<td>2.1 ± 0.7</td>
</tr>
<tr>
<td>Lixisenatide</td>
<td>0.7 ± 0.2</td>
<td>1.2 ± 0.3</td>
</tr>
<tr>
<td>Net hepatic alanine uptake, (\mu\text{mol}kg^{-1}\text{min}^{-1})</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Control</td>
<td>3.5 ± 0.3</td>
<td>3.9 ± 0.8</td>
</tr>
<tr>
<td>Lixisenatide</td>
<td>2.7 ± 0.5</td>
<td>3.0 ± 0.6</td>
</tr>
</tbody>
</table>

Data are means ± SE. NEFA, nonesterified fatty acid. *Significant differences between groups in their responses over time identified by ANOVA (P < 0.05); †time points where post hoc testing indicated differences between groups.

Lixisenatide appears to hold promise for improving the care of selected individuals with type 2 diabetes. The data from the current study and a previous study in dogs undergoing an OGTT (60) are generally consistent with those in humans given exogenous GLP-1 infusions or short-term treatment with a DPP-IV inhibitor in that insulin concentrations were not increased in the lixisenatide group during most of the postprandial period. Continuous intravenous GLP-1 slowed gastric emptying and brought about a dose-dependent reduction in insulin concentrations after a meal in both normal volunteers and those with type 2 diabetes (28, 33). Only with the use of erythromycin as a prokinetic agent was it possible to observe an insulinotropic effect of GLP-1 following meal feeding in normal subjects (30). Moreover, despite the increase in insulin secretion during combined treatment with erythromycin and GLP-1, the glucose-lowering effect of GLP-1 was markedly reduced by inhibition of its ability to delay gastric emptying (30). On the other hand, in subjects with type 2 diabetes, there was no effect of 10 days of treatment with the DPP-IV inhibitor vildagliptin on either the rate of gastric emptying or insulin concentrations following a mixed meal (59). However, after longer periods (6–52 wk) of vildagliptin therapy, β-cell effectiveness (insulin secretion in relation to glycaemia) in individuals with prediabetes or type 2 diabetes was improved significantly relative to placebo (2, 49, 58). Thus it is possible that higher insulin levels might have been observed in dogs if lixisenatide treatment had continued chronically. Nevertheless, lixisenatide reduced both 2-h postprandial insulin concentrations as well as the 0- to 4-h postprandial insulin AUC in a dose-dependent manner in adults with type 2 diabetes inadequately controlled with metformin (44) while also reducing postprandial glucagon concentrations dose dependently (43). Exenatide, another short-acting GLP-1 receptor agonist, exhibits a similar postprandial insulin-lowering effect (25). The insulin concentrations in the control group showed substantial fluctuations. This was likely due to the pulsatile nature of insulin secretion in the dog, as in the human (19, 41, 42). With 15- to 30-min sampling intervals, the samples may capture concentrations anywhere on the curve from peak to trough. Additionally, the absorption of a meal is not a smooth process but instead occurs in a rhythmic pattern induced by the migrating myoelectric complex (9, 45). This has been observed to entrain insulin secretion (45). Overall, our data from the lixisenatide group agree with those of Lorenz et al. (27) obtained in the human in that postprandial insulin secretion appeared to be appropriate to the glucose stimulation. Lixisenatide appears to improve glucose disposal (glucose clearance) even in the absence of obvious enhancement of insulin secretion.

Using the specific GLP-1 receptor antagonist exendin (9–39), Shirra et al. (55) determined that GLP-1’s reduction of glucagon secretion can play a significant role in reducing fasting and postprandial glycemia. However, in the current acute study, there was only a small effect, if any, of lixisenatide to lower postprandial glucagon concentrations (Fig. 5).
Lixisenatide treatment slowed the suppression of NEFA concentrations and reduced the fall in glycerol concentrations in the postprandial period. These effects were likely related to the reduced insulin concentrations in the lixisenatide group. The lixisenatide group also had lower blood lactate concentrations in the early postprandial period, apparently due to a reduction in hepatic glycolysis related to the lower rates of NHGU in the lixisenatide group.

In conclusion, the reduction in the postprandial glycemic response in the current report, combined with the promising data from clinical trials (1, 17, 39, 46, 47, 50, 56), indicates that lixisenatide could be a useful adjunct in treatment of postprandial hyperglycemia in individuals with impaired glucose tolerance or type 2 diabetes. As suggested in humans (23, 34), our data demonstrate a delay in gastric emptying following administration of lixisenatide that allows more efficient disposal of the carbohydrate in the meal. Data from mice and humans suggest that this may be accompanied by improved postprandial triglyceride and lipoprotein metabolism (29, 37, 38, 65). There appears to be no specific effect of lixisenatide to increase liver glucose uptake.

GRANTS

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DISCLOSURES

U. Werner is an employee of Sanofi.

AUTHOR CONTRIBUTIONS

M.C.M., M.S.S., and T.D.F. performed the experiments; M.C.M., M.S.S., and T.D.F. analyzed the data; M.C.M. and A.D.C. interpreted the results of the experiments; M.C.M. prepared the figures; M.C.M. drafted the manuscript; M.C.M., U.W., and A.D.C. edited and revised the manuscript; M.C.M., U.W., and A.D.C. contributed to the conception and design of the research.

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