Luminal leptin inhibits L-glutamine transport in rat small intestine: involvement of ASCT2 and B₀AT1

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Ducroc R, Sakar Y, Fanjul C, Barber A, Bado A, Lostao MP. Luminal leptin inhibits l-glutamine transport in rat small intestine: involvement of ASCT2 and B₀AT1. Am J Physiol Gastrointest Liver Physiol 299: G179–G185, 2010. First published May 6, 2010; doi:10.1152/ajpgi.00048.2010.—L-glutamine is the primary metabolic fuel for enterocytes. Glutamine from the diet is transported into the absorptive cells by two sodium-dependent neutral amino acid transporters present at the apical membrane: ASCT2/SLC1A5 and B₀AT1/SLC6A19. We have demonstrated that leptin is secreted into the stomach lumen after a meal and modulates the transport of sugars after binding to its receptors located at the brush border of the enterocytes. The present study was designed to address the effect of luminal leptin on Na⁺-dependent glutamine (Gln) transport in rat intestine and identify the transporters involved. We found that 0.2 nM leptin inhibited uptake of Gln and phenylalanine (Phe) (substrate of B₀AT1) using everted intestinal rings. In Ussing chambers, 10 mM Gln absorption followed as Na⁺ induced short-circuit current was inhibited by leptin in a dose-dependent manner (maximum inhibition at 10 nM; Iₑ₅₀ = ~0.1 nM). The absorption was also decreased by leptin. Western blot analysis after 3-min incubation of the intestinal loops with 10 mM Gln, showed marked increase of ASCT2 and B₀AT1 protein in the brush-border membrane that was reduced by rapid preincubation of the intestinal lumen with 1 nM leptin. Similarly, the increase in ASCT2 and B₀AT1 gene expression induced by 60-min incubation of the intestine with 10 mM Gln was strongly reduced after a short preincubation period with leptin. Altogether these data demonstrate that, in rat, leptin controls the active Gln entry involved in the interorgan nitrogen flux. Gln is also the most important precursor of the nucleosides and glucose synthesis and is well known that leptin is a multifunctional hormone that is also involved in immune and neuroendocrine functions and nutrient absorption (22). This action is consistent with the production of leptin by many other tissues (1, 15, 23), as well as the expression of its receptors in peripheral tissues (14, 22). Indeed, we have demonstrated that leptin receptor is expressed in both the apical and basolateral membrane of intestinal absorptive cells (3). Other authors also showed that leptin is secreted into the gastric lumen after a meal by pepsinogen-containing secretory granules of chief cells (11), which also contain the leptin-soluble receptor, indicating that this leptin receptor isoform is also released into the gastric lumen (12). Bound to this receptor, leptin remains stable in the gastric juice, despite the severe conditions of pH and proteolytic activity in the gastric lumen, and is able to reach the intestinal lumen (17). Accordingly, we have demonstrated that leptin present in the intestinal lumen inhibits sugar absorption in vivo by regulating the Na⁺/glucose cotransporter SGLT1 (19). A similar effect was previously reported in vitro (2, 16, 18, 21) with implication of both PKC and PKA activation (4, 16). Interestingly, luminal leptin enhances the intestinal transport of dipeptides by the H⁺/peptide transporter PEPT1 and CD147/medium-chain triglyceride (MCT)-1 mediated uptake of butyrate in mice and Caco-2 cells (9, 10), as well as fructose transport by the facilitative glucose transporter GLUT5 in rat intestine in vivo (25). Given that leptin seems to regulate different nutrient transporters, one could anticipate that it may also modulate amino acid transporters in the intestine. However, there are not data in this respect yet. Considering the importance of Gln for the whole organism and in the intestine itself, the aim of the present study was to investigate the effect of luminal leptin on Gln transport and the target transporters of the hormone. We found that both ASCT2 and B₀AT1 are involved in Na⁺-dependent uptake of glutamine in rat intestine and regulated by apical leptin. The present results give new insights into the role of leptin as a major gastrointestinal hormone regulating intake of rich energy molecules.

MATERIALS AND METHODS

Animals. Male Wistar rats weighing 220–260 g were obtained from Charles River Laboratories, L’Arbresle, France and the Applied Pharmacology Research Center (CIFA) of the University of Navarra,
Pamplona, Spain. They were caged under standard laboratory conditions with tap water and regular food provided ad libitum, in a 12-h:12-h light/dark cycle at a temperature of 21–23°C and fasted for 16–18 h, with free access to water, before the experiments. The animals were treated in accordance with the European Community Guidelines concerning the care and use of laboratory animals. The animal studies were performed under license from the veterinary department of Paris, France (to A. Bado and R. Ducroc, authorization no. 75-955 of September 2, 2004, and no. 75-174 of October 9, 2003, respectively; agreement no. B75-18-02; decision no. 05/12 established on July 12, 2005 by Prefecture of Police de Paris, France). The experimental protocol was approved by the Animal Research Ethics Committee of the University of Navarra, with the no. 064-06.

**Everted intestinal ring uptake assays.** Rats were anesthetized by intraperitoneal injection of a mixture (4:1) of ketamine chlorohydrate (Ketolar; Pfizer Orion, Espoo, Finland), at a dose of 0.25 ml per 100 g body wt. Uptake of Gln or Phe by everted jejunal rings was determined as previously described (21). Briefly, rats were anesthetized, and a segment (20–25 cm) of jejunum was quickly excised, rinsed with ice cold saline solution (NaCl 0.9%), everted, and cut into ~30-ng pieces. Groups of six rings were incubated for 15 min at 37°C under continuous shaking and gassed with O₂ in Krebs-Ringer-Tris containing 0.5 nM Gln or 50 μM Gln (Sigma-Aldrich, St. Louis, MO) and 0.064 μCi/ml L-[¹⁴C(U)]-glutamine (218 mCi/mmol; American Radiolabeled Chemicals, St. Louis, MO) or 0.5 nM Phe (Merck, Darmstadt, Germany) and 0.064 μCi/ml L-[¹⁴C(U)]-phenylalanine (370 mCi/mmol; American Radiolabeled Chemicals), in the absence (control) and in the presence of 0.2 nM recombinant rat leptin (Peprotech EC, London, UK). After the incubation period, rings were washed in ice-cold saline solution and radioactivity incorporated into the tissue was determined as previously described (21). Briefly, rats were anesthetized and laparotomized. Three small intestinal loops (~8 cm starting 15 cm from the cecum were prepared and filled with 1 nM leptin or saline. After 3-min in vivo incubation, loops were filled with 10 mM Gln for 60 min. After this time, rats were killed, loops were dissected and rinsed in saline, and mucosa was scrapped off on ice. Total RNA was extracted from the mucosa samples with the Trizol reagent (Qiagen, Valencia, CA). The first-strand cDNA was synthesized by reverse transcription from 5 μg of total RNA with SuperScript II reverse transcriptase (Invitrogen, Cergy-Pontoise, France). Quantification of cDNA was performed with a Light Cycler System (Roche Diagnostics, Meylan, France) according to the manufacturer’s instructions. Primers were as follows: rASCT2, 5′-CCATCGGAAAGGGAACTCT-3′ and 5′-CTTCAAGAGCCTCCTTCGAA-3′; rBoAT1, 5′-TTACCAAGTCGGGTTG-3′ and 5′-GATGGCCTCGTCATGCAAG-3′; r5′-GATGGCGGTCCCATGACGAT-3′; r18S, 5′-CCCTGCGCTT- TGGTACACACC-3′ and 5′-GATCCGGAGGCCGCTCATA-3′. They were designed with oligo 4 software and synthesized by Eurogentec (Southampton, UK). The comparative ΔΔCT-method was used for relative mRNA quantification of target genes, normalized to protein 18S and a relevant control equal to 2-ΔΔCT.

**Chemicals.** Recombinant murine leptin was purchased from PreProtec EC (London, UK). All other chemical reagents were purchased from Sigma. Leptin antagonist L39A/D40A (kind gift of Prof. Arie G. Gertler) was diluted in saline.

**Statistical analysis.** All results were expressed as means ± SE. Student’s t-test or one-way ANOVA with Tukey-Kramer multiple comparison post test when appropriate were performed using GraphPad Prism version 4.0 for Windows (GraphPad Software, San Diego, CA) or SPSS v15. The level of significance was set at P < 0.05.

**RESULTS**

**Leptin inhibits glutamine uptake in everted rings.** The effect of leptin on Gln uptake was first studied on preparations of everted intestinal rings after 15-min incubation. As shown in Fig. 1A, uptake of 0.5 mM Gln was significantly reduced (~35%) by 0.2 mM leptin. The hormone also significantly reduced, by ~25%, uptake of 50 μM Gln, a concentration close to the Kₜ₀ for ASCT2 (6) (Fig. 1B). To evaluate the contribution of B⁰AT1 on Gln absorption at both concentrations, we examined the effect of 25 mM Phe (specific substrate of B⁰AT1) on Gln uptake. As depicted in Fig. 1A, Phe reduced 0.5 mM Gln uptake by ~65%, indicating a higher contribution of B⁰AT1 on Gln uptake compared with ASCT2. However, uptake of 50 μM Gln was not inhibited by Phe (Fig. 1B), demonstrating that at this
uptake but was also inhibited (\(\text{ration as Fig. 1}\)). The uptake of 0.5 mM Phe, assayed in the same tissue preparation only ASCT2 was contributing to Gln absorption.

Luminal leptin reduces glutamine-induced \(I_{sc}\). Leptin effect was further investigated in Ussing chambers, a polarized system that permits the access to either side of the tissue preparation. After the intestinal mucosa was isolated in the chamber and allowed to reach a steady state (usually 40 min), tissues were challenged with 10 mM Gln, which induced a rapid (less than 2 min) and significant rise in \(I_{sc}\) (\(\sim 23 \mu \text{A/cm}^2\)). The increase in \(I_{sc}\) was the result of the \(\text{Na}^+\) mucosal-to-serosal movement that sustained the amino acid entry through sodium-dependent amino acid transporter(s). In accordance with the results obtained with intestinal rings (Fig. 1, A and C), 10 mM Phe induced a small \(I_{sc}\) which was \(\sim 12\%\) of that for Gln (Fig. 2A). Luminal addition of 10 mM Ala also significantly raised \(I_{sc}\) (\(\Delta I_{sc} \sim 15 \mu \text{A/cm}^2\)), whereas no change in \(I_{sc}\) was observed after 10 mM methyl aminoisobutyric acid (data not shown), indicating the absence of system A participation in the electrogenic uptake of Gln.

Again, in line with the data obtained with intestinal rings (Fig. 1A), 20 mM Phe reduced the Gln-induced \(I_{sc}\) by \(\sim 70\%\) when it was added to the mucosal bath 3 min before 10 mM Gln challenge (Fig. 2B) and leptin (10 nM) inhibited by \(\sim 60\%\) Gln and Ala-induced \(I_{sc}\) (Fig. 2C).

Inhibition of 10 mM Gln transport across intestinal mucosa, following rapid incubation with leptin in the mucosal reservoir, was found to be concentration dependent (Fig. 2D). Inhibition was significant with 0.01 nM (\(\sim 30\%\)), maximal for 10 nM (\(\sim 80\%\)), and decreased with 100 nM leptin up to \(\sim 50\%\). The concentration producing a half-maximal inhibition of Gln transport (IC50) was 0.1 nM (Fig. 2D). Leptin (10 nM) reduced the \(I_{sc}\) induced by 30 mM Gln (26% inhibition, \(P < 0.05\)) and also the small \(I_{sc}\) induced by 1 mM Gln although, in this case, the inhibition did not reach statistical significance (data not shown).

**Mucosal vs. serosal leptin.** The possible effect of leptin on Gln-induced \(I_{sc}\) acting from the serosal side was also studied in Ussing chambers. As shown in Fig. 2E, 10 nM leptin after 2 min in the serosal bath also decreased the 10 mM Gln-induced \(I_{sc}\) by \(\sim 40\%\). This decrease, however, was smaller than that produced by the hormone acting from the mucosal side, which was around 80%. After 10 min, leptin from the serosal side had no more inhibitory effect while it was still active from the mucosal side (50% inhibition).

**Leptin inhibition of glutamine transport is blocked by leptin receptor antagonist.** To examine whether the inhibitory effect of leptin was dependent on leptin receptor, we studied the effect of L39A/D40A, a mutated leptin-based peptide acting as a leptin antagonist (26). As shown in Fig. 2F, addition of 50 nM L39A/D40A to the mucosal bath immediately before 10 nM leptin reversed leptin inhibition of Gln-induced \(I_{sc}\), whereas 5 nM L39A/D40A had no effect.

**Leptin reduces glutamine-induced ASCT2 and B0AT1 protein expression.** Western blot assays were performed to investigate whether leptin could modify the expression in the plasma membrane of ASCT2 and B0AT1 transporters. As shown in Fig. 3, incubation of the intestinal loops with 10 mM Gln during 3 min increased the presence of ASCT2 and, in a lesser extent, of B0AT1 in the BBMV (Fig. 3, A and B). This increase was inhibited when the loops had been previously incubated with 1 nM leptin for 3 min, showing that leptin reduced Gln-induced ASCT2 and B0AT1 expression in the plasma membrane.

**Fig. 1.** Effect of leptin on glutamine (Gln) and phenylalanine (Phe) uptake by everted intestinal rings. Uptake of 0.5 mM Gln (A), 20 mM Phe (B), or 0.5 mM Phe (C) was measured in the absence or presence of 0.2 nM leptin after 15-min incubation. Uptake of Gln was also measured in the presence of 25 mM Phe (A and B). Experiments in A and C were performed in the same experimental group. The results are expressed as means \(\pm \text{SE; } ***P < 0.001\) (\(n = 6\)–18 from 3 animals).

concentration only ASCT2 was contributing to Gln absorption. The uptake of 0.5 mM Phe, assayed in the same tissue preparation as Fig. 1A, was found less than 10% of 0.5 mM Gln uptake but was also inhibited (\(\sim 45\%\)) by leptin (Fig. 1C).
Leptin reduces glutamine-induced ASCT2 and B0AT1 mRNA expression. Finally, the effect of luminal leptin on Gln-induced ASCT2 and B0AT1 gene expression was studied. Luminal infusion of 10 mM Gln for 60 min markedly increased the level of ASCT2 and B0AT1 mRNA (6.7-fold and 2.8-fold, respectively compared with their relative controls; Fig. 4, A and B). When the intestinal loops were first incubated with 1 nM leptin for 3 min before the Gln infusion, the increase in ASCT2 and B0AT1 mRNA was significantly reduced (~55%). These results indicate that leptin can also regulate Gln-induced ASCT2 and B0AT1 mRNA expression.

DISCUSSION

The present results demonstrate that, in rat intestine, leptin inhibits the absorption of Gln and the protein and mRNA expression of ASCT2 and B0AT1, the two major Gln trans-

**Fig. 2.** Leptin inhibition of glutamine induced short-circuit current ($I_{sc}$) in Ussing chamber. A: $I_{sc}$ induced by 10 mM Gln, alanine (Ala), and Phe ($n = 8–29$). B: effect of 20 mM Phe on 10 mM glutamine-induced $I_{sc}$ ($n = 4$). C: effect of 10 nM mucosal leptin on $I_{sc}$ induced by 10 mM Gln or Ala. D: dose response for mucosal leptin inhibition of Gln-induced $I_{sc}$. Leptin was added in the mucosal bath 2 min before tissues were challenged with 10 mM Gln. Values for $I_{sc}$ were standardized to control value. Each point represents the mean ± SE of 4–6 noncumulative values from 4 separate experiments. E: effect of serosal leptin (sx) vs. mucosal leptin (mq) at 10 nM concentration on 10 mM Gln-induced $I_{sc}$ (control) ($n = 6$). *P < 0.05 vs. control; #P < 0.05 vs. 2 min mq. F: action of leptin is receptor specific; leptin antagonist L39A/D40A was added to the mucosal bath 5 min before the addition of leptin, and 3 min later tissues were challenged in the mucosal side with 10 mM Gln ($n = 6$).
by hormones and peptides is not yet completely understood. Growth hormone (GH) and epidermal-growth factor (EGF) can enhance intestinal Gln uptake (27), and a possible hormonal stimulation of ASCT2 in vivo has been suggested (24).

Leptin is now well documented as an important regulator of nutrient transporters from the apical membrane of the enterocytes. Indeed, leptin was shown to control butyrate MCT-1-mediated absorption (10) and di-/tripeptide transporter PepT-1 (9) and modulate the activity of glucose and fructose transporters through its action on SGLT1 (16, 19, 21), GLUT2, and GLUT5 (25). A significant role for leptin in controlling active transport of Gln was previously established in human placental villous fragments where leptin stimulates the activity of system

![Fig. 3](image-url) Effect of luminal leptin on Gln-induced expression of ASCT2 and B0AT1 proteins in brush-border membrane (BBM). BBM vesicles were obtained from mucosa of NaCl-treated (control), 1 nM leptin-treated, and 10 mM Gln-treated (3 min) intestinal loops, and ASCT2 (A) and B0AT1 (B) protein expression were analyzed by Western blot. Representative Western blot images are shown. Intensity of the immunoreactive bands was quantified and expressed in relation to control. Results are expressed as means ± SE for 3–4 rats per experimental group. **P < 0.01 and ***P < 0.001 vs. control. #P < 0.05 vs. Gln.

![Fig. 4](image-url) Effect of leptin on Gln-induced ASCT2 and B0AT1 gene expression. Total RNA was extracted from mucosa of NaCl-treated (control), 1 nM leptin-treated, and 10 mM Gln-treated (60 min) intestinal loops, and ASCT2 (A) and B0AT1 (B) mRNA expression were analyzed by RT-PCR. Results are expressed as ratio of transporter/18S mRNA expression in relation to control and presented as means ± SE of 3–4 rats per experimental group. *P < 0.05; **P < 0.01, and ***P < 0.001 vs. control; #P < 0.05 vs. Gln.
A, thus controlling the availability of this important fuel to placenta cells (20).

Here, the functional studies were first performed in vitro using everted intestinal rings. The same technique was used to demonstrate for the first time that leptin inhibits sugar intestinal transport (21). Even though with this technique the mucosa is exposed to the medium and the access of the hormone to the enterocyte is mostly through the apical membrane, a possible basolateral action of leptin could not be discarded. Therefore, we further measured Gln transport in Ussing chambers, where contribution of mucosal vs. serosal leptin could be distinctly examined and mucosal-to-serosal Gln entry could be followed as sodium-induced $I_{sc}$. As observed with intestinal rings, leptin pretreatment from the mucosal side induced a rapid and marked reduction of the Gln-induced $I_{sc}$. This effect was concentration dependent with characteristics of inhibition found in the nanomolar range (16, 21). The effect of leptin from the serosal side was found significantly less pronounced and disappeared at the longest incubation time (10 min). These results may reflect the fact that leptin exposed to the serosal side needs to diffuse a few layers of muscle and connective tissues before reaching its receptors on the basolateral membrane of the enterocytes, with possible reduction of the effective concentration. In addition, the membrane surface area for leptin action is smaller when it acts from the serosal side compared with the apical side, and the distance between the receptor and the Gln transporters is higher. After 10 min, the lack of leptin action from the serosal side can be also related to the decrease of its effect from the apical side.

The Ussing chamber technique mimics the relevant action of leptin, which is produced and secreted by gastric cells (1, 11) together with its soluble receptor (12) and can flow along the digestive lumen to reach the small intestine and act as a physiological modulator. In fact, using leptin mутеїіїз, a leptin receptor antagonist (26), we demonstrate that this modulation requires an effective interaction of leptin with its specific receptor located at the BBM of the small intestine (3, 9).

Interestingly, a recent report shows that, in leptin receptor-deficient obese Zucker rats, Gln transport is highly increased (29). This is in line with the inhibitory action of leptin as a regulator of the Gln uptake here reported. It is tempting to speculate, therefore, that leptin may act as a brake on the entry of Gln, possibly in balance with another stimulating hormone (i.e., GH or EGF). Interestingly, this leptin-induced action is apparently different from the one recently proposed for angiotensin II that inhibits intestinal glucose uptake but not amino acid transport under normal and diabetic conditions (30, 31).

In the intestine, Gln transport is assumed to be predominantly achieved by the sodium-dependent transporter B0AT1 (28). In rat intestinal rings, uptake of Phe, a specific substrate of B0AT1, was also inhibited by leptin, indicating that B0AT1 was a target for the hormone. Interestingly, Phe uptake was very low compared with Gln uptake at the same concentration, which can be explained by the fact that B0AT1 shows lower affinity for its substrates ($K_{0.5}$ ranging from 1.4 to 4 mM) compared with ASCT2 ($K_{0.5} = \sim 20 \mu M$) (6). Nevertheless, 25 mM Phe was able to inhibit by $\sim 65\%$ Gln uptake, suggesting a higher contribution of B0AT1 on Gln absorption in rat intestine under this experimental condition. By employing Ussing chamber, we also found that Phe-induced $I_{sc}$ was lower compared with the Gln-induced $I_{sc}$. Aside from B0AT1, ASCT2 is the other sodium-dependent Gln transporter present in the apical membrane of small intestine (6). From the functional studies using 50 $\mu M$ Gln (concentration at which only ASCT2 is responsible for Gln absorption), we also demonstrate that ASCT2 is regulated by leptin.

To assess directly the presence of the two Gln transporters in rat BBM and responsiveness to luminal substrate and/or leptin, we further used biochemical and molecular approaches. Our results clearly indicate that, in the basal condition, both B0AT1 and ASCT2 are present in rat BBM. Their expression in the membrane can be rapidly increased by luminal Gln, and this increase can be reduced by leptin at nanomolar range. Interestingly, the ratio of ASCT2/B0AT1 protein expression in the basal state was low (Fig. 3) and confirmed the reported predominance of B0AT1 in intestinal BBM (28). However, after luminal challenge with Gln, more ASCT2 than B0AT1 protein was observed to swing into the BBM. These biochemical data would support a major importance of ASCT2 on Gln uptake in rat intestine. However, because of the differences in the experimental conditions, it is not possible to compare the functional and biochemical results to draw conclusions about which transporter is the principal in Gln entrance in the enterocytes. Nevertheless, both functional and biochemical data hallmark the contribution of ASCT2 and B0AT1 transporters in Gln uptake.

We further demonstrated that leptin, in addition to its effect on posttranslational regulation, can rapidly trigger transcriptional control of the two Gln transporters. Thus leptin inhibits the Gln-induced increase of ASCT2 and B0AT1 mRNA levels. Regulation of ASCT2 gene expression by Gln was previously reported in hepatic epithelial cells (7). The stimulation of expression in response to Gln was shown to involve, in part, binding of FXR/RXR to the ASCT2 promoter (8).

In summary, we demonstrate that luminal leptin can modulate the intestinal activity and expression of the Gln transporters ASCT2 and B0AT1 in the apical membrane of the enterocytes. Because Gln is known to have both acute and chronic effects on cell metabolism and function (13), the present findings of leptin involvement in the modulation of Gln uptake in both short- and long-term mode of action are believed to be of major importance for the biology of intestinal cells.

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DISCLOSURES

No conflicts of interest, financial or otherwise, are declared by the authors.

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LEPTIN MODULATES INTESTINAL GLUTAMINE ABSORPTION