Somatostatin restrains the secretion of glucagon-like peptide-1 and -2 from isolated perfused porcine ileum

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Abstract

Somatostatin restrains the secretion of glucagon-like peptide-1 and -2 from isolated perfused porcine ileum. Am J Physiol Endocrinol Metab 280: E1010–E1018, 2000.—Suspecting that paracrine inhibition might influence neuronal regulation of the entodocrine L cells, we studied the role of somatostatin (SS) in the regulation of the secretion of the proglucagon-derived hormones glucagon-like peptide-1 and -2 (GLP-1 and GLP-2). This was examined using the isolated perfused porcine ileum stimulated with acetylcholine (ACh, 10⁻⁶ M), neuretin C (NC, 10⁻⁸ M), and electrical nerve stimulation (NS) with or without α-adrenergic blockade (phentolamine 10⁻⁵ M), and perfusion with a high-affinity monoclonal antibody against SS. ACh and NC significantly increased GLP secretion, whereas NS had little effect. SS immunoneutralization increased GLP secretion eight- to ninefold but had little influence on the GLP responses to ACh, NC, and NS. Basal SS secretion (mainly SS28) was unaffected by NS alone. Phentolamine + NS and NC abstract strongly stimulated release mainly of SS14, whereas ACh had little effect. Infused intravascularly, SS14 weakly and SS28 strongly inhibited GLP secretion. We conclude that GLP secretion is tonically inhibited by a local release of SS28 from epithelial paracrine cells, whereas SS14, supposedly derived from enteric neurons, only weakly influences GLP secretion.

Gastrointestinal hormones; immunoneutralization

GLUCAGON-LIKE PEPTIDE (GLP)-1 is a peptide hormone secreted from the distal part of the small intestine and the colon in response to meal ingestion (18). It plays an important role as an incretin hormone, stimulating insulin secretion (40), and also functions as one of the gastrointestinal hormones; immunoneutralization

MATERIALS AND METHODS

This study conformed to the Danish legislation governing animal experimentation (1987) and was carried out after permission was granted from the National Superintendence for Experimental Animals.

Perfusion experiments. Young pigs of the YDL-strain weighing 15–20 kg were anesthetized with ketamine chloride (Ketalar, Parke-Davis, Morris Plains, NJ) for premedication, pentobarbital for induction, and chloralose (50 mg/kg) and N₂O/O₂ for maintenance. After laparotomy, an 80-cm segment of the central ileum, including its arterial and venous supply, was isolated, excised, and positioned floating in Ringer solu-
carbon dioxide in oxygen with a multibulb oxygenator. The perfusate was gassed continuously with a mixture of 5% fortid, Dumex, Copenhagen, 5 mg/l) was added to prevent abattoir), and a cyclooxygenase inhibitor (indomethacin, Conbovine erythrocytes (obtained 2–3 days earlier at a local and a mixture of amino acids (Vamin, 14 g/l, Pharmacia) in an amount yielding a total amino acid concentration of 5 mmol/l. The perfusion medium also contained 20% freshly washed bicarbonate solution containing, in addition, 0.1% human artery by use of a pulsatile finger pump (Ole Dich, Copenhagen, 5 ml/min), ensuring an oxygen delivery of 0.75 mol O2/min, which is about twofold in excess of the oxygen consumption of the preparation. Perfusion pressure was recorded constantly via a sidearm to the arterial catheter by a Statham transducer connected to an amplifier and a recorder. Motor activity of the gut was estimated visually, but contractions of the gut were also reliably reflected as short-lasting increments in perfusion pressure (spikes), validated in experiments involving intraluminal manometry (38). The frequency of such increments (which did not increase the basal perfusion pressure) was recorded. A bipolar platinum electrode, shaped like a hook, was positioned carefully around the supplying artery and the network of nerve fibers surrounding it, without damage to the fibers, and was kept in place by a loose ligature. Square wave impulses were delivered by a nerve stimulator. The intestinal lumen was perfused at a rate of 3 ml/min with preheated perfusate containing 15 mmol/l glucose. The venous effluent was collected for 1-min periods, chilled in ice, and centrifuged immediately after start of perfusion. The supernatants were frozen at 4°C within a few minutes. The supernatants were frozen was prepared from a total of 19 pigs. In six perfusion experiments, randomized order. After another 10-min rest period, an terminal decapeptide of gastrin-releasing polypeptide (GRP), addition was considered essential to eliminate the inhibitory effect of the sympathetic innervation on GLP and other hormone secretions of the mucosa. Electrical stimulation of the mixed intestinal nerve (as we use here) invariably activates the sympathetic fibers, and removal of their inhibitory effect may be necessary to observe the effect of stimulator neurons (25). After 10 min, the NS was repeated. After a further 10-min rest, 5-min infusions of acetycholine (ACh; Sigma, St. Louis, MO, via preweighed vials and at a final concentration of 10–8 mol/l) were given via the arterial line in randomized order. After another 10-min rest period, an infusion of high-affinity (affinity constant, or Kd, = 1011 l/mol) monoclonal somatostatin (SS) antibodies (20), which had been dialyzed overnight (against saline), was started, in an amount yielding an SS binding capacity of 6 mmol/l. After 25 min, 5-min periods of NS or ACh and NC infusions were carried out as before. In six additional perfusion experiments, a 5-min infusion of SS14 at 10–8 mol/l (final perfusate concentration) was carried out after the initial equilibration period, and in four perfusion experiments, stepwise increasing dose-response experiments were carried out for both SS14 and SS28 (at 10–10 to 10–8 mol/l). At the end of the experiments, NC was infused as before, without or with SS14 and SS28, both at 10–8 mol/l. In three perfusion experiments, monoclonal antibodies against 2,4,6-trinitrophenyl (3) were added to yield y-globulin concentrations similar to the anti-SS experiments. In an earlier study, when the same experimental model was used, it was demonstrated that vascular and secretory responses remained unchanged for up to 4 repeated instances of NS (25).

Analyses of the perfusion effluents. All effluent fractions were analyzed for SS and GLP–1 by use of previously described radioimmunoassays (17, 31). The SS assay recognizes equally well SS14 and SS28. The GLP-1 assay was carried out using the antibody 89390, which has an absolute requirement for the amidated COOH terminus of the GLP-1 molecule and therefore recognizes GLP-1(7–36)amide and the inactive metabolite, GLP-1(9–36)amide. The effluent fractions from three of the immunoneutralization experiments (the effluents from the other experiments were not available for analyses) and the effluent fractions from the four dose-response experiments with SS14 and SS28 were analyzed for GLP-2 content by use of the antibody 92160, which recognizes the NH2 terminus of GLP-2 and, therefore, only fully processed biologically active GLP-2. Details of the assay are described in Ref. 43. The detection limits of the assays were below 5 pmol/l.

Chromatography. Pools of effluent samples, collected during different experimental conditions (in the basal state, during NS + phentolamine perfusion, during NC + phentolamine perfusion, and during SS antibody perfusion), were applied to Sep-Pak C18 cartridges (Water-Millipore, Milford, MA) and eluted with 70% ethanol + 0.1% trifluoroacetic acid. The effluent was dried, redissolved in albumin-containing (0.1%) phosphate buffer (0.05 mM, pH 7.5), and applied to a Sephadex G50 fine K16/100 column (Pharmacia, Uppsala, Sweden) eluted with the same buffer at 4°C.

The eluted fractions resulting from gel filtration of the pools of venous effluent obtained in the basal state, during NS + phentolamine, and during NC + phentolamine perfusion, were analyzed for immunoreactive glicentin, GLP-1, and GLP-2 by use of the following antisera: 4304 (recognizes the glucagon part of glicentin, previously described in Ref. 21), 2135 (a so-called side-viewing antibody, recognizing a midregion and, therefore, all forms of GLP-1, previously described in Ref. 30), 89390 (described above and in Ref. 31), 92071 (a COOH-terminally directed antibody, recognizing specifically glycine-extended GLP-1–(7–37), previously described in Ref. 31), 92160 (described above), and another side-viewing GLP-2 antiserum that recognizes a midregion of GLP-2 (Peninsula Laboratories, cat. no. RAS 7176, Merseyside, St Helens, UK). For the side-viewing GLP-2 assay, we employed monoclonidated rat GLP-2, with an Asp33 Tyr33 substitution as tracer but otherwise similar assay conditions (43).

Calculations. Because of the constant perfusion flow, the hormone effluent concentrations parallel secretion rates. The data are presented as means ± SE. Changes in hormone secretion as a function of time were evaluated by ANOVA for repeated measures, or, alternatively, mean hormone outputs for the immediate prestimulatory 5-min periods were compared with mean 5-min stimulated plateau responses (calculated for the 3- to 7-min period after start of NS) by use of a
Table 1. Perfusion pressure and motor activity

<table>
<thead>
<tr>
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<th>Perfusion Pressure, % of basal</th>
<th>Motor Activity, spikes/5 min</th>
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<tr>
<td></td>
<td>Means ± SE</td>
<td>Changes vs. basal</td>
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<td>Basal condition</td>
<td>100</td>
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<tr>
<td>8 Hz</td>
<td>133 ± 10*</td>
<td>P &lt; 0.050</td>
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<tr>
<td>8 Hz + phentolamine</td>
<td>111 ± 3</td>
<td>P &lt; 0.050</td>
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<tr>
<td>NC + phentolamine</td>
<td>105 ± 1</td>
<td>P &lt; 0.050</td>
</tr>
<tr>
<td>NC + phentolamine + somatostatin antibodies</td>
<td>106 ± 1</td>
<td>P &lt; 0.001</td>
</tr>
<tr>
<td>ACh + phentolamine</td>
<td>103 ± 1</td>
<td>ns</td>
</tr>
<tr>
<td>ACh + phentolamine + somatostatin antibodies</td>
<td>117 ± 6</td>
<td>P &lt; 0.050</td>
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<tr>
<td>ACh + phentolamine + somatostatin antibodies</td>
<td>114 ± 4</td>
<td>P &lt; 0.050</td>
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8 Hz, nerve stimulation; NC, neuromedin C; ns, nonsignificant. *Significantly different from 8 Hz + phentolamine and 8 Hz + phentolamine + somatostatin antibodies, P < 0.050; †significantly different from 8 Hz + phentolamine and 8 Hz + phentolamine + somatostatin antibodies, P < 0.010; ‡significantly different from NC + phentolamine + somatostatin antibodies, P < 0.050.

RESULTS

The results concerning perfusion pressure and motor activity responses to infusions and nerve stimulations are summarized in Table 1. NS significantly increased perfusion pressure, and phentolamine strongly reduced this response, illustrating successful stimulation of sympathetic efferent fibers (eliciting vasoconstriction and therefore increasing perfusion pressure) and the expected effect of α-adrenergic blockade on this response. As expected, NS had little effect on motor activity alone but markedly enhanced motor activity after α-adrenergic blockade, illustrating effective stimulation of nonadrenergic motor neurons under these circumstances (38). Both ACh and NC significantly stimulated motor activity, demonstrating adequate activation of cholinergic and bombesin receptors.

The main finding of the present study is the dramatic increase in GLP-1 and GLP-2 secretion after SS immunoneutralization (Fig. 1). After 4–5 min, both GLP-1 and GLP-2 secretion increased, reaching plateau levels of 813 ± 141 and 995 ± 385% of basal secretion after 15–20 min. In preliminary experiments in which immunoneutralization was carried out without phentolamine, similar increases in GLP-1 secretion were observed (4- to 8-fold increases, n = 3, not shown). Thus the effect of immunoneutralization was independent of the presence of absence of noradrenergic blockade. For GLP-1, plateau values averaged ~400 pmol/l, whereas the GLP-2 plateau amounted to ~800 pmol/l. Because of this unexpected difference, we investigated the molecular nature of the secreted peptides by chromatographic analysis of pooled, concentrated perfusion effluents from four perfusion experiments. The results are shown in Fig. 2. Glucagon (glicentin) immunoreactivity eluted at a distribution coefficient (Kd) of 0.35, corresponding to the elution position of synthetic and natural glicentin. GLP-2 immunoreactivity eluted at Kd 0.55–0.60, regardless of the assay employed (sideviewing or NH2 terminal). The side-viewing GLP-1 assay (code no. 2135) identified a peak at Kd 0.65, and a peak with a similar Kd was obtained with the assay for glycine-extended GLP-1 (code no 92071). The assay for amidated GLP-1 revealed a peak around Kd 0.60.

![Fig. 1. Effect of infusion of monoclonal somatostatin antibodies (binding capacity 6 nmol/ml and affinity constant (Kd) of 10-11 [mol] on glucagon-like peptide-1 (GLP-1) secretion (A) and GLP-2 secretion (B). Values are means ± SE of 6 perfusion experiments (n = 3 for GLP-2).](http://ajpendo.physiology.org/)}
Fig. 2. Gel filtration profiles of immunoactive glicentin and GLP-1 and -2 in pools of effluent from isolated perfused ileum collected during somatostatin immunoneutralization. Eluted immunoactivity is expressed as a percentage of total eluted amount, plotted as a function of the coefficient of distribution ($K_d$). Values are means ± SE of results from gel filtration of 4 effluent pools. A: results of side-viewing assays for glucagon (x, comprising glicentin and oxyntomodulin) and GLP-1 (●) and the NH$_2$-terminal assay for GLP-2 (□). B: results of side-viewing assay for GLP-2 (□) and COOH-terminal assays for amidated (●) and glycine-extended GLP-1 (▲). In C, total amounts of immunoactivity eluted as measured in each assay are compared with amount measured with side-viewing GLP-1 assay (code 2135), set to 100%. Significant differences: **$P < 0.010$; ***$P < 0.001$. In separate experiments, synthetic glicentin was found to elute at $K_d$ 0.35, oxyntomodulin at 0.75, GLP-2 at 0.55–0.60, and amidated and glycine-extended GLP-1 at 0.60 and 0.65, respectively.

Fig. 3. Effect of nerve stimulation (8 Hz) on secretion of GLP-1 (A), GLP-2 (B), and somatostatin (C) without additions (left), during phentolamine administration ($10^{-5}$ M; middle), or during infusion of somatostatin antibodies and phentolamine ($10^{-5}$ M) infusion (right). Values are means ± SE of 6 perfusion experiments ($n = 3$ for GLP-2).
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Figure 3 shows the results of electrical mixed NS, either alone or in the presence of phentolamine (10^{-5} mol/l) and phentolamine + SS antibodies. GLP-1 secretion was unaffected by NS regardless of the perfusion conditions, whereas GLP-2 secretion decreased to a nadir of 52 ± 9% after 3 min of NS. After addition of phentolamine and phentolamine + SS antibody, GLP-2 secretion increased insignificantly during NS to maximally 297 ± 129 and 159 ± 30% of basal, respectively. SS secretion was uninfluenced by NS in the control experiments but increased greatly (to 470 ± 80% of prestimulatory levels, P < 0.010) during phentolamine perfusion. Chromatographic analysis revealed that the SS immunoreactivity released in the basal state mainly consisted of SS28, whereas during NS + phentolamine, mainly SS14 was released (Fig. 4). SS secretion could not be measured during antibody perfusion. Figure 5 shows the NC stimulation experiments during phentolamine alone and during phentolamine + SS antibody perfusion. GLP-1 secretion, expressed in percentage of basal, was significantly enhanced by NC during phentolamine as well as antibody perfusion (to 293 ± 43%, P < 0.010 and 149 ± 16%, P < 0.050, respectively; P < 0.050 for the difference). The absolute output of GLP-1 was larger during NC + antibody perfusion (3.4 ± 0.6 vs. 1.1 ± 0.1 pmol/min, P < 0.050). NC, in all three cases, stimulated GLP-2 secretion under both phentolamine and phentolamine + SS antibody (to 265 ± 42 and 175 ± 52% of prestimulatory levels), but the difference did not reach significance. SS secretion was greatly augmented by NC (to 437 ± 73%, P < 0.010), and chromatographic analysis revealed that SS14 was the predominant molecular form (Fig. 4).

Figure 6 shows the ACh stimulation experiments during phentolamine alone and during phentolamine + SS antibody perfusion. GLP-1 secretion showed a somewhat delayed increase in response to ACh, both during phentolamine and SS antibody perfusion (to 169 ± 12 and 116 ± 4%, in the 5- to 8-min period after start of ACh, P < 0.010 and 0.050, respectively). The absolute outputs of GLP-1 did not differ significantly (0.8 ± 0.2 vs. 1.3 ± 0.4 pmol/min, respectively). ACh also tended to increase GLP-2 secretion under both conditions (to 265 ± 50 and 111 ± 18% of basal secretion, but the changes did not reach statistical significance). ACh did not affect SS secretion.

In the six experiments in which SS14 was infused to a final concentration of 10^{-9} mol/l, GLP-1 secretion decreased to 79 ± 7%, P < 0.050. Because of this finding, a dose-response study was carried out. Figure 7 shows the effect of SS28 and SS14 infused to final concentrations of 10^{-10}, 10^{-9}, and 10^{-8} mol/l on GLP-1 and GLP-2 secretion. SS28 dose dependently inhibited both GLP-1 and GLP-2 secretion, reaching 38 ± 10% of basal secretion for GLP-1 and 27 ± 7% for GLP-2 at 10^{-8} mol/l (P < 0.01). SS14 had weak and inconsistent inhibitory effects on GLP secretion. During co-infusion of SS28 and NC, GLP-1 secretion was inhibited to 66 ± 5% and GLP-2 secretion to 50 ± 9% (P < 0.05), both significantly different from the stimulated secretion observed with NC alone (P < 0.01). In contrast, during

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co-infusion of SS14 and NC, both GLP-1 and GLP-2 secretion increased (P = 0.08 and 0.09, respectively) to levels not significantly different from those obtained with NC alone. Addition of the 2,4,6-trinitrophenyl antibodies had no influence on any registered parameter (data not shown).

DISCUSSION
The most important finding of the present study is the dramatic increase in the secretion of GLP-1 and -2 in response to somatostatin immunoneutralization. The antibodies were administered as a highly purified solution of γ-globulins, and there was no effect of...
infusion of control antibodies, speaking against unspecific effects. In addition, the high concentrations of the glucagon-like peptide measured during immunoneutralization were verified by chromatographic analysis of the effluent (see further discussion). Furthermore, the inhibitory effect of exogenous somatostatin was confirmed. It must be concluded, therefore, that under the prevailing experimental conditions, local somatostatin secretion exerts a marked restraint of the secretory activity of the L cells, the endocrine cells known to be the source of the glucagon-like peptides. This raises the question of the cellular source of somatostatin involved.

In previous studies, we found that the density of endocrine epithelial somatostatin-secreting cells is lower in the ileum compared with the more proximal segments of the gut (41). On the other hand, the ileal mucosa harbors a dense network of somatostatin-containing nerve fibers, which seem to originate in the nerve cell bodies of the submucous plexus (41). The predominant secretory product of the endocrine somatostatin-producing cells is thought to be SS28 (corresponding to the COOH-terminal 28 amino acids of prosomatostatin) (2, 12), whereas the main product of the intestinal neurons is SS14 (the COOH-terminal 14 amino acids of prosomatostatin) (37). In the present study, the chromatographic analysis of ileal effluent revealed that almost all of the immunoreactivity released in the basal state corresponded to SS28. Together, these findings indicate that the source of somatostatin interacting with the L cells in the basal state is the endocrine cells, whereas somatostatin released during nerve stimulation or neuropeptide C infusion corresponds to SS14, and therefore presumably is released from the enteric neurons. In addition, our experiments showed that SS28 was much more efficacious and potent as an inhibitor of GLP secretion than SS14. We conclude, therefore, that the local somatostatinergic control of GLP secretion is exerted by paracrine cells of the mucosa releasing mainly SS28, which is likely to interact with sstr-5 receptors on the L cells, because sstr-5 is the only somatostatin receptor with a preference for SS28 (33). Brubaker et al. (5) reported that GLP-1 was capable of releasing both SS14 and SS28 from rat intestinal cultures. In preliminary experiments using the experimental model of the present study, in which the morphological integrity of the tissue and the paracrine relationships are preserved, we found a dose-related release of somatostatin upon intravascular infusions with GLP-1, with $10^{-9}$ mol/l being the most effective concentration (unpublished studies). This suggests that GLP-1 and somatostatin secretion may be mutually interdependent in a paracrine relationship, whereby GLP-1 limits its own secretion by activating a SS28-mediated paracrine inhibition.

The possible influence of SS14 released from enteric neurons on GLP secretion was investigated in a series of nerve stimulation experiments. Electrical stimulation of the mixed extrinsic nerves innervating the ileal segment elicited vasoconstriction in the segment, attesting to the efficiency of stimulation. However, this had little effect on the secretion of the GLPs or somatostatin, although a tendency to inhibition was noted. This is in agreement with the results of yet unpublished studies, in which intra-arterial norepinephrine inhibited GLP release (Mineo and Holst, unpublished studies). In the present experiments, addition of the $\alpha$-adrenergic blocker phentolamine reduced the hypertensive response as expected (36), but now nerve stimulation strongly stimulated somatostatin release. This suggests that the intestinal somatostatinergic neurons are innervated by inhibitory, noradrenergic fibers as well.
as strongly stimulatory fibers. Acetylcholine alone had no effect on somatostatin release and is therefore unlikely to act as the stimulatory transmitter; vasoactive intestinal polypeptide, which is released from intrinsic ileal neurons in the pig under similar conditions (25), might be responsible. GLP secretion was slightly augmented by nerve stimulation during α-adrenergic blockade. In addition, acetylcholine weakly but significantly stimulated secretion, in agreement with previous investigations performed in vivo and in other species, suggesting that acetylcholine may stimulate L cell secretion (16, 34, 44). Thus it appears that cholinergic mechanisms may lead to a weak stimulation of GLP-1 secretion.

We then hypothesized that the dramatic release of somatostatin that was observed during α-blockade might dampen the secretory response of the L cells. To test this hypothesis, nerve stimulation was repeated during somatostatin immunoneutralization, with a technique previously demonstrated to bring about a lasting and virtually complete elimination of the actions of exogenous as well as endogenous somatostatin (20). However, under these conditions (which included α-adrenergic blockade), nerve stimulation had no significant effects. Our immediate conclusion would be that somatostatin, released during nerve stimulation, had no influence on the GLP response to nerve stimulation. A similar conclusion was reached from the experiments involving the neurotransmitter neuromedin C. Neuromedin C, the COOH-terminal active fragment of mammalian bombesin or GRP, has been shown to provide a strong stimulus for GLP secretion (29), as also confirmed here. Neuromedin C, however, also stimulated somatostatin secretion, almost as much as nerve stimulation under phentolamine infusion. The GLP response to neuromedin C was preserved during somatostatin immunoneutralization. The fractional increase was much smaller than before antibody infusion, but the amount above that elicited by the immunoneutralization, was larger. Similar results were obtained with acetylcholine. These results would, again, be interpreted to indicate that somatostatin released in response to nerve stimulation had only a minor influence on nerve- or neurotransmitter-stimulated GLP-1 secretion. This notion is supported by the finding of a weak effect of SS14, the molecular form released during nerve stimulation, and by the finding that only SS28, but not SS14, could inhibit neuromedin C-stimulated GLP secretion.

From our experiments, we conclude that paracrine SS28 is a main regulator of GLP secretion and that the extrinsic innervation of the gut exerts a weak, presumably cholinergic stimulatory effect on GLP secretion that is independent of somatostatinergic intestinal neurons. GRP (or neuromedin C) is much more efficacious than acetylcholine, raising the possibility that GRP-producing neurons could be involved in a stimulatory secretory control of GLP secretion. However, in nerve stimulation experiments (Orskov C, Knuthsen S, and Holst JJ, unpublished studies), we have never been able to measure a co-release of GRP and GLPs, which seems to preclude an association between the two. With respect to somatostatin, we were unable to identify mechanisms that regulate the paracrine somatostatinergic cells restraining GLP secretion (except for the potential paracrine actions of GLP-1 itself). The epithelial somatostatin cells seem to function independently of the extrinsic innervation. Our results show that adrenergic mechanisms strongly inhibit somatostatin secretion but also seem to inhibit GLP secretion, so that diffuse activity in the extrinsic noradrenergic neurons does not cause a stimulated release of GLPs, despite lifting of the somatostatin restraint.

We were concerned about the discrepancy between the amounts of GLP-1 and GLP-2 secreted during somatostatin immunoneutralization; therefore, we performed a chromatographic analysis, which was facilitated by the large amount of immunoreactive material released. Of the glucagon-containing peptides, glicentin predominated; amidated GLP-1 [GLP-1-(7—36)amide] and glycine-extended GLP-1 [GLP-1-(7—37)] contributed about equally to the total amount of GLP-1 secreted, and GLP-2 occurred predominantly as a single component, identified equally well by side viewing and NH₂-terminal assays. Together the two forms of GLP-1 approximately equaled the amount of GLP-2, in agreement with the concept that the two are secreted in equimolar amounts. However, the large contribution of GLP-1-(7—37) was unexpected and distinguishes pigs from humans, in whom the amidated form predominates (31). Pigs seem to resemble rats in this respect (26). This observation illustrates the difficulties involved in determining secretion of GLP-1 in vivo, where side-viewing assays, measuring “total” GLP-1, cannot be used because of pancreatic secretion of GLP-1-containing molecules that are picked up by such assays (30). In addition, both NH₂-terminally modified and COOH-terminally modified molecular forms are circulating (8, 9), precluding the use of single antibody or sandwich approaches for detection of the total secretory rate of GLP-1. The rate of GLP-2 secretion, estimated using an NH₂-terminal assay, may thus provide the best estimate of L cell activity.

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