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Sympathetic activity controls fat-induced OEA signaling in small intestine

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Abstract

Ingestion of dietary fat stimulates production of the small-intestinal satiety factors oleoylethanolamide (OEA) and *N*-palmitoyl-phosphatidylethanolamine (NPPE), which reduce food intake through a combination of local (OEA) and systemic (NPPE) actions. Previous studies have shown that sympathetic innervation of the gut is necessary for duodenal infusions of fat to induce satiety, suggesting that sympathetic activity may engage small-intestinal satiety signals such as OEA and NPPE. In the present study, we show that surgical resection of the sympathetic celiac superior mesenteric ganglion, which sends projections to the upper gut, abolishes feeding-induced OEA production in rat small-intestinal cells. These effects are accounted for by suppression of OEA biosynthesis, and are mimicked by administration of the selective β_2 -adrenergic receptor antagonist, ICI-118,551. We further show that sympathetic ganglionectomy or pharmacological blockade of β_2 -adrenergic receptors prevents NPPE release into the circulation. In addition, sympathetic ganglionectomy increases meal frequency and lowers satiety ratio, and these effects are corrected by pharmacological administration of OEA. The results suggest that sympathetic activity controls fat-induced satiety by enabling the coordinated production of local (OEA) and systemic (NPPE) satiety signals in the small intestine.

Keywords

Oleoylethanolamide; *N*-palmitoyl-phosphatidylethanolamine; sympathetic; feeding

Introduction

The lipid-derived satiety factors, oleoylethanolamide (OEA) (Rodríguez de Fonseca et al., 2001; Fu et al., 2007) and *N*-palmitoyl-phosphatidylethanolamine (NPPE) (Gillum et al., 2008), are critically involved in the regulation of fat intake. These substances are produced in the proximal small intestine after ingestion of dietary fat and exert their anorectic effects through seemingly distinct physiological mechanisms (Rodríguez de Fonseca et al., 2001; Fu et al., 2007; Gillum et al., 2008; Schwartz et al., 2008). Evidence indicates that OEA

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operates as a local messenger within the gut and promotes satiety by recruiting afferent fibers of the vagus nerve (Rodriguez de Fonseca et al., 2001; Gaetani et al., 2003; Proulx et al., 2005; Fu et al., 2007; Fu et al., 2008), whereas NPPE is thought act as a blood-borne hormone and to reduce food intake by direct activation of neuronal circuits in the hypothalamus (Gillum et al., 2008).

The biochemical pathways underlying the production of OEA and NPPE also appear to be markedly different. OEA biosynthesis in response to dietary fat occurs primarily, if not exclusively, in absorptive epithelial cells of the duodenum and jejunum (Fu et al., 2007; Schwartz et al., 2008; Gillum et al., 2008). These cells utilize food-derived oleic acid to generate the phospholipid, *N*-oleoyl-phosphatidylethanolamine (NOPE), which is then cleaved by *N*-acyl-phosphatidylethanolamine-selective phospholipase D (NAPE-PLD) to release OEA (Fig. 1) (Fu et al., 2007; Schwartz et al., 2008). OEA-mediated signaling at peroxisome proliferator-activated receptors type- α (Fu et al., 2003) engages vagal afferents in the gut to promote satiety (Rodriguez de Fonseca et al., 2001) and is terminated by OEA hydrolysis (Fig. 1) (Fu et al., 2007). Small-intestinal NPPE biosynthesis is also stimulated by dietary fat (Gillum et al., 2008), but is not dependent on food-derived oleic acid and its precise regulation is still unknown (Fig. 1) (Fu et al., 2007; Schwartz et al., 2008).

An important question that remains to be addressed is whether the nervous system participates in the control of OEA- and NPPE-mediated signaling. The sympathetic division of the autonomic nervous system, in particular, is likely to play a role in this process. Sympathetic preganglionic neurons synapse at prevertebral ganglia, forming the celiac-superior mesenteric ganglion complex, which sends noradrenergic projections to multiple visceral organs, including the intestine (Lindh et al., 1988; Chevendra and Weaver, 1991; Parr et al., 1993; Furness et al., 2000; Nasser et al., 2006). Surgical removal of this complex impairs the ability of intra-duodenal infusions of fat to reduce food intake, which suggests that the outflow of sympathetic nerves to the gut may contribute to fat-induced satiety (Sclafani et al., 2003). Moreover, previous work from our laboratory has shown that sympathetic activation stimulates OEA biosynthesis in rat white adipose tissue (LoVerme et al., 2004). In the present study, we have used a combination of surgical and pharmacological approaches to examine whether sympathetic activity is implicated in the control of OEA and NPPE mobilization in the rat small intestine.

Materials and Methods

Animals

A total of 205 adult male Sprague-Dawley rats (250-300 g) were purchased from Charles River (Wilmington, MA) and housed in standard cages (4/cage) at room temperature (22°C). They were maintained on a 12-hour light/dark cycle (on at 6:30 am, off at 6:30 pm) and had free access to water and standard chow pellets (Research Diets, New Brunswick, NJ), unless otherwise noted. When experimentation began, the rats were housed individually in suspension cages consisting of wire bottoms to prevent coprophagia, and habituated to the experimental settings for 4 days. All procedures met the National Institutes of Health guidelines for the care and use of laboratory animals, and were approved by the Institutional Animal Care and Use Committee of the University of California, Irvine.

Chemicals

Prazosin, CGP20712A, SR59230A, ICI-118,551, isoproterenol, and salbutamol were purchased from Sigma-Aldrich (St. Louis, MO). Drugs were prepared at concentrations that allowed administration in 2 ml/kg of an appropriate vehicle (described below). 1,2-dipalmitoyl-*sn*-glycero-3-phosphoethanolamine was from Sigma and [³H]-anandamide

(arachidonoyl-1-[³H]ethanolamide) from American Radiolabeled Chemicals (Saint Louis, MO). [²H₄]-OEA, 1,2-dipalmitoyl-*sn*-glycero-3-phosphoethanolamine-*N*-heptadecanoyl, and 1,2-dipalmitoyl-*sn*-glycero phosphoethanolamine-*N*-heptadecanoyl were synthesized in the lab (Fu et al., 2007) using appropriate fatty acid chlorides purchased from Nu-Chek Prep (Elysian, MN).

Surgical procedures

Resection of the celiac superior mesenteric ganglion complex was performed as previously described (Sclafani et al., 2003). In brief, animals were anesthetized with an intraperitoneal (IP) injection of ketamine (100 mg/kg) and xylazine (10 mg/kg). A midline incision was made through the skin and abdominal muscle, and the stomach and spleen were exposed and gently retracted. The descending aorta was visualized, and the ganglion complex was isolated with fine forceps and severed for removal using microsurgical scissors. Care was taken not to damage the adjacent celiac and superior mesenteric arteries. The abdominal muscle wall was closed using plain gut 4-0 suture (Fischer Scientific; Houston, Texas) and the skin sutured using stainless steel wound clips. Sham-operated control animals underwent the same procedure, except that the sympathetic ganglion complex was left intact. For pain management, animals received buprenorphine (0.02 mg/kg, IP) just prior to and following surgery. Testing started 7 days after surgery. Following experimentation, verification of correct ganglionectomy was made by determining the absence of neural tissue at the level of the celiac and mesenteric artery branching at the descending aorta (see Figure 2 E-F).

Experimental design

1. Effects of celiac superior mesenteric ganglionectomy—Forty-eight rats were randomized into four groups of ganglionectomized and sham-operated ($n=12$ for each surgery condition for rats re-fed standard rodent chow, and $n=12$ for each surgery condition for rats re-fed a high-fat diet; experimentation on groups receiving different diets occurred on separate days). Seven days after surgery, each group was further divided into 2 subgroups: food-deprived rats ($n = 6$), which had no access to food for 24 h, and re-fed rats ($n=6$), which were allowed to eat standard rodent chow or the high-fat diet for 30 min following a 24-h fast. All animals within a diet group were killed at the same time.

2. Effects of adrenergic receptor antagonists—Each receptor antagonist was tested on a separate group of 24 rats. The animals were randomized into two groups: food-deprived ($n=12$) and re-fed ($n=12$) (see above). Each group was further divided into a vehicle subgroup ($n=6$), which received 2 ml/kg of dimethylsulphoxide (30% in saline), and a drug subgroup ($n=6$), which received one of the following antagonists: CGP2071 (β_1 receptor; 1 mg/kg), ICI-118,551 (β_2 receptor; 1 mg/kg), SR59230A (β_3 receptor; 1 mg/kg) or prazosin (α receptor; 5 mg/kg). Vehicle or drugs were administered by IP injection 30 min before tissue collection (jejunum, blood).

3. Effects of β -adrenergic receptor agonists—Thirty-six food-deprived animals were divided into 3 groups: vehicle ($n=6$), which received an IP injection of saline (2 ml/kg); isoproterenol (β_1/β_2 receptors; 2.5, 5, 10 mg/kg, $n = 6$ per dose) and salbutamol (β_2 receptors; 1 or 5 mg/kg, $n = 6$ per dose). Six free-feeding animals were also included. Drugs were administered by IP injection 30 min before tissue collection (jejunum).

Feeding behavior

Animals (9 sham-operated rats and 10 ganglionectomized rats) were habituated to the test cages for 3 days prior to trials. Feeding behavior was analyzed in rats starting at the onset of the dark phase (6:45 pm), and lasted for 24 h. Free feeding was analyzed for 5 consecutive

days, and the average of these 5 trials was calculated for each parameter. Food intake was recorded using the automated system TSE Drinking & Feeding Monitor (TSE Technical & Scientific Equipment GmbH, D-61350 Bad Homburg, Germany). This system consists of cages equipped with baskets filled with standard chow pellets and connected to weight sensors that record the duration of each meal (each time food is taken from the basket), the quantity of food consumed, and the time at which the event occurred. Weight variations were monitored every second, and the threshold for an eating episode was set at 0.6 g and > 1 min. Feeding parameters (Gaetani et al., 2003) were divided into the following categories: average meal frequency (meals/h), average meal size (MS; g/kg), average post-meal interval (PMI; min), average satiety ratio [SR; ratio between average PMI and meal size, min/(g/kg)]. Feeding data were analyzed as total food ingested per kg of body weight and as cumulative food intake (g/kg) over 24 h. The significance of differences between sham-operated and ganglionectomized rats were analyzed by Student's *t*-test.

Lipid extractions—Animals were anesthetized with isoflurane, the jejunum was rapidly removed, snap-frozen in liquid N₂ and stored at -80°C until analyses. Blood was collected via cardioventricular puncture in heparin-containing vials and centrifuged at 2000×g for 30 min to obtain plasma. Samples of jejunal tissue were homogenized in 1 ml of methanol containing [2H₄]-OEA as internal standard. Lipids were extracted with chloroform (2 vol) and the extracts were washed with water (1 vol). Organic phases were collected and dried under N₂. OEA, NOPE and NPPE were fractionated by open-bed silica gel column chromatography, as previously described (Astarita et al., 2009). Eluted fractions were dried under N₂ and reconstituted in 60 ml of methanol for analyses.

Liquid Chromatography/Mass Spectrometry (LC/MS) analyses—We used an 1100-LC system coupled to a 1946D-MS detector (Agilent Technologies, Inc., Palo Alto, CA) equipped with an electrospray ionization (ESI) interface. Lipids were separated on a XDB Eclipse C₁₈ column (50 × 4.6 mm i.d., 1.8 μm, Zorbax), eluted by a gradient of methanol in water (from 85% to 90% methanol in 2.5 min) at a flow rate of 1.5 ml/min. Column temperature was kept at 40°C. MS detection was in the positive ionization mode, capillary voltage was set at 3 kV, and fragmentor voltage at 120V. Lipids were quantified with an isotope-dilution method (Giuffrida et al., 2000), monitoring sodium adducts of the molecular ions [M+Na]⁺ in the selected ion-monitoring (SIM) mode (Fu et al., 2007). For NOPE and NPPE analysis, we used a Poroshell 300SB C₁₈ column (2.1 × 75 mm i.d., 5 μm, Agilent) eluted with a linear gradient of methanol in water containing 5 mM ammonium acetate and 0.25% acetic acid (from 85% to 100% in 4 min), at a flow rate of 1 ml/min. NOPE and NPPE were identified using an Ion Trap XCT (Agilent) system in the negative ionization mode; the capillary voltage was set at 4.5kV, the skim1 at -40V, and the capillary exit at -151 V. NOPE and NPPE were identified by their retention times and LC/MSⁿ properties as described elsewhere (Fu et al., 2007). Extracted ion chromatograms were used to quantify each precursor ion by selecting two or more characteristic product ions: 1-stearoyl-2-arachidonoyl-*sn*-glycero-phosphoethanolamine-*N*-oleoyl (for NOPE), 1-stearoyl-2-arachidonoyl-*sn*-glycero-phosphoethanolamine-*N*-palmitoyl (for NPPE) and 1,2-dipalmitoyl-*sn*-glycero-3-phosphoethanolamine-*N*-heptadecanoyl, which was used as an internal standard (Fu et al., 2007).

NAPE-PLD assay—Jejunal tissue was homogenized in ice-cold Tris-HCl (50 mM, pH 7.5, 10 vol) containing 0.32 M sucrose. Homogenates were centrifuged at 1,000×g for 10 min and the supernatants (0.1 mg protein) were incubated at 37°C for 30 min in Tris-HCl (50 mM, pH 7.4, 0.2 ml) containing 0.1% Triton X-100, phenylmethylsulfonyl fluoride (1 mM) and 1,2-dipalmitoyl-*sn*-glycero-3-phosphoethanolamine-*N*-heptadecanoyl (0.1 mM). Reactions were stopped by adding 0.6 ml of chloroform-methanol (2:1, vol/vol) containing

N-heptadecenoyl-[²H₄]-ethanolamide as internal standard. After centrifugation at 1,000×g at 4°C for 10 min, the organic layers were collected and dried under N₂. The residues were suspended in 50 μl of chloroform/methanol (1:3, vol/vol) and analyzed by LC/MS. We quantified *N*-heptadecenoylethanolamide with an isotope-dilution method, monitoring sodium adducts of molecular ions in the SIM mode (Fu et al., 2007).

FAAH assay—Jejunal tissue was homogenized and homogenates were centrifuged at 1,000×g for 10 min. Supernatants (50 μg protein) were incubated with [³H]-anandamide (10,000 dpm, specific activity, 20 Ci/mmol) for 30 min at 37°C in 0.5 ml of 50 mM Tris-HCl buffer (pH 8.0) containing fatty acid-free bovine serum albumin (0.05% weight/vol). Reactions were terminated adding 1.5 ml of chloroform/methanol (2:1, vol/vol) and phases were separated by centrifugation. The aqueous layers were collected and [³H]-ethanolamine was quantified by liquid scintillation counting.

Statistical analyses—Results are expressed as the mean ± S.E.M. The significance of differences between groups was evaluated by two-way analysis of variance (ANOVA), followed by a Dunnett's test for multiple comparisons. Analyses were conducted using GraphPad Prism software (GraphPad Software; San Diego, CA) and differences were considered significant if *p* < 0.05.

Results

Sympathetic denervation abolishes food-induced OEA mobilization

As previously shown for intact rats (Fu et al., 2007; Schwartz et al., 2008), re-feeding with standard rodent chow after a 24-h fast increased the levels of OEA [Fig. 2A, Two-way ANOVA, *F*(1,20)=14.47, *p*=0.001] and its precursor NOPE [Fig. 2B, Two-way ANOVA, *F*(1,20)=6.71, *p*=0.017] in jejunal tissue of sham-operated rats. *Ex vivo* measurements (Fu et al., 2007; Schwartz et al., 2008) showed that these effects are accompanied by an increase in NAPE-PLD activity [Fig. 2C, Two-way ANOVA, *F*(1,20)=6.03, *p*=0.018] and a decrease in FAAH activity [Fig. 2D, Two-way ANOVA, *F*(1,20)=11.11, *p*=0.002]. Surgical resection of the sympathetic celiac superior mesenteric ganglion, which extends projections to the upper gut (Lindh et al., 1988; Chevendra and Weaver, 1991; Parr et al., 1993; Furness et al., 2000), disabled food-dependent OEA mobilization in the jejunum (Fig. 2A). The surgical procedure markedly reduced OEA biosynthesis, preventing food-dependent changes in both NOPE production and NAPE-PLD activity (Fig. 2B,C), and lowered FAAH activity in food-deprived animals (Fig. 2D). Re-feeding with a high-fat diet (60% total kcal from fat) increased jejunal OEA levels in sham-operated rats [Table 1, Two-way ANOVA, *F*(1,20)=5.8, *p*=0.025], as seen in animals re-fed with standard chow (Figure 2A). Removal of the sympathetic ganglion attenuated, but did not completely block this response (Table 1). The surgical procedure removed the ganglion complex in its entirety (Fig. 2E-F). In addition, we found no marked difference in standard chow or high-fat consumption and body weight between ganglionectomized and sham-operated animals, except for a minor, but significant, increase in free-feeding 24-h standard chow intakes (Table 2). The results suggest that the sympathetic nervous system serves an obligatory function in food-induced OEA mobilization.

Role of β₂-adrenergic receptors

β₂-Adrenergic receptors are highly expressed in the rodent small intestine (Nasser et al., 2006). Pharmacological blockade of these receptors with the selective antagonist ICI-118,551 prevented food-induced OEA mobilization in rat jejunum [Fig. 3A, Two-way ANOVA, *F*(1,20)=26.3, *p*=0.001]. Moreover, β₂-receptor blockade impaired the ability of re-feeding to alter NOPE levels [Fig. 3B, Two-way ANOVA, *F*(1,20)=5.2, *p*=0.048] and

NAPE-PLD activity [Fig. 3C, Two-way ANOVA, $F(1,20)=6.84$, $p=0.014$] and reduced FAAH activity in food-deprived animals [Fig. 3D, Two-way ANOVA, $F(1,20)=10.78$, $p=0.003$]. Underscoring the pharmacological selectivity of these effects, systemic administration of drugs that antagonize other adrenergic receptors – including α (prazosin), β_1 (CGP2071) and β_3 receptors (SR59230A) – did not alter jejunal OEA mobilization (Table 3). Furthermore, systemic administration of adrenergic receptor agonists – including isoproterenol (2.5, 5 and 10 mg/kg), which targets β_1/β_2 receptors, or salbutamol (1 and 5 mg/kg), which preferentially targets β_2 receptors – exerted no detectable effect on jejunal OEA content under food-deprived conditions (Table 4). The same agonist dosages were found to markedly stimulate OEA production in rat white adipose tissue (LoVerme et al., 2006). We interpret these results to indicate that sympathetic activity facilitates small-intestinal OEA production, but is not sufficient to elicit this effect in the absence of food intake.

Sympathetic activity controls blood levels of NPPE and NOPE

Re-feeding after a 24-h fast increased plasma NPPE levels in sham-operated rats, as previously shown for intact rats (Gillum et al., 2008) [Fig. 4A, Two-way ANOVA, $F(1,20)=12.56$, $p=0.004$]. This response was greatly reduced in rats in which the celiac superior mesenteric ganglion had been removed (Fig. 4A) or which had received an injection of the β_2 -adrenergic antagonist ICI-118,551 (1 mg/kg) 30 min prior to food exposure [Fig. 4B, Two-way ANOVA, $F(1,20)=12.00$, $p=0.0032$]. Similar results were obtained with the OEA precursor, NOPE, in those receiving surgery [Fig. 4C, Two-way ANOVA, $F(1,20)=12.36$, $p=0.0015$] or pharmacological treatment [Fig. 4D, Two-way ANOVA, $F(1,20)=7.83$, $p=0.01$]. Notably, in contrast with a previous report (Gillum et al., 2008), we found that both baseline and food-stimulated NOPE levels in plasma were significantly higher than corresponding NPPE levels (Fig. 4). We conclude that sympathetic neural activity controls, through activation of β_2 -adrenergic receptors, the release of both NPPE and NOPE into the general circulation.

Sympathetic denervation modifies meal patterns

OEA modifies food intake by reducing daily meal frequency, and increasing satiety ratio [ratio between average post-meal interval (PMI) and meal size (Gaetani et al., 2003)]. Free-feeding ganglionectomized animals displayed increases in average 24-h meal frequency (Fig. 5A1), and decreases in average meal size (Fig. 5A2), average PMI (Fig. 5A3), and average satiety ratio (Fig. 5A4), versus sham-operated controls. Treatment with an effective dose of OEA [10mg/kg, IP, at time of dark onset (Gaetani et al., 2003)] completely normalized all daily free-feeding parameters affected by ganglionectomy (Fig. 5B1-4). Ganglionectomized animals re-fed after a 24-h fast displayed greater average meal frequencies (Fig. 5C2), and decreased average PMI (Fig. 5C3) and average satiety ratio (Fig. 5C4), versus sham-operated controls. We found no statistically significant changes in average meal size (Fig. 5C1). The results indicate that, under conditions where OEA signaling is disrupted by ganglionectomy, normal meal patterns controlled by OEA become aberrant.

Discussion

Dietary fat stimulates enterocytes in the proximal small intestine to produce the endogenous lipid amide, OEA, which may regulate satiety through a local autocrine or paracrine mechanism (Rodríguez de Fonseca et al., 2001; Fu et al., 2003; Schwartz et al., 2008). Consistent with this view, systemic administration of OEA inhibits food intake in rodents, and this effect is (i) prevented by treatment with the sensory-fiber neurotoxin, capsaicin (Rodríguez de Fonseca et al., 2001); and (ii) mimicked by virally induced over-expression of

NAPE-PLD in the small intestine, which locally elevates tissue OEA levels (Fu et al., 2008), but not by infusion of OEA into the brain ventricles (Rodriguez de Fonseca et al., 2001). These findings suggest that OEA regulates feeding by recruiting sensory vagal afferents in the gut. In further support of this possibility, OEA increases expression of c-Fos (a marker of neuronal activation) in the nucleus of the solitary tract of the brainstem and in the paraventricular and supraoptic nuclei of the hypothalamus, which receive direct and indirect neural inputs from the sensory vagus (Rodriguez de Fonseca et al., 2001). Furthermore, systemic administration of OEA stimulates expression of the hypothalamic neuropeptide, oxytocin, and intracerebroventricular infusion of an oxytocin receptor antagonist prevents OEA-induced hypophagia (Gaetani et al., 2010). These studies collectively suggest that small-intestinal fluctuations in OEA mobilization regulate satiety by triggering the central release of oxytocin through the sensory vagus (Gaetani et al., 2010). Similarly to OEA, NPPE inhibits food intake and stimulates c-Fos expression in the paraventricular and supraoptic nuclei of the hypothalamus (Gillum et al., 2008). However, the effects of systemic NPPE administration are not altered by subdiaphragmatic vagotomy and are reproduced by intracerebroventricular infusion of NPPE, suggesting that this compound may reach its target sites in the brain through the general circulation (Gillum et al., 2008). Thus, a plausible interpretation of the available data is that dietary fat stimulates small intestinal cells to generate mediators that act both locally (OEA) and systemically (NPPE) to modulate satiety responses.

In the present study, we have shown that (i) surgical removal of the sympathetic innervation of the gut prevents the food-induced mobilization of OEA and NPPE; (ii) this effect is mimicked by pharmacological blockade of β_2 -adrenergic receptors, but not by administration of other adrenergic receptor antagonists; (iii) β_2 -adrenergic agonists fail to stimulate OEA mobilization when administered alone to food-deprived animals; and (iv) sympathetic ganglionectomy alters meal patterns that are under the control of OEA. These results indicate that fat-induced signaling through OEA and NPPE, though primarily regulated by dietary fat ingestion, is subjected to the integrative control of the sympathetic nervous system.

The sympathetic nervous system plays a key function in the control of energy balance (Berthoud, 2008; Lambert et al., 2010). Current research is primarily focused on the role played by this system in peripheral glucose and lipid metabolism (Haque et al., 1999; Bartness et al., 2005; Nogueiras et al., 2007; Nogueiras et al., 2009; Shiuchi et al., 2009), yet the existence of links between feeding state and sympathetic activity has long been recognized. Early work by Young and Landsberg has shown that fasting is associated with a decrease in sympathetic outflow (Young and Landsberg, 1977a), whereas sucrose feeding and chronic feeding of lard are accompanied by an increase in sympathetic outflow (Young and Landsberg, 1977b; Young et al., 1994). These changes parallel those undergone by small-intestinal OEA, the levels of which also decrease during fasting and increase during re-feeding (Fu et al., 2003; Fu et al., 2007; present study). More recently, Sclafani and collaborators found that surgical ablation of the celiac superior mesenteric ganglion complex – the same intervention used here to remove sympathetic innervation of the gut – prevents the satiating actions of duodenal fat infusion (Sclafani et al., 2003). These observations are consistent with the findings in this report, and raise the intriguing possibility that sympathetic activity modulates satiety in response to fat by facilitating fat-dependent OEA and NPPE signaling in the small intestine.

An important question that remains to be addressed pertains to the mechanism through which sympathetic activity facilitates small-intestinal OEA biosynthesis. Postganglionic-sympathetic efferents targeting the intestine terminate in the submucosal and myenteric plexus (Nasser et al., 2006; Lomax et al., 2010), but do not reach the enterocytes where

OEA is synthesized (Fu et al., 2007; Schwartz et al., 2008). In addition, β_2 receptors are localized to enteric neurons in the submucosal and myenteric plexus of the rodent intestine, which implies that adrenergic activity controls enterocyte OEA biosynthesis through an indirect mechanism (e.g., a diffusible factor or reflexes mediated by the enteric nervous system). Despite these uncertainties, our results, showing that both blockade of β_2 receptors and ganglionectomy abolish feeding-induced OEA biosynthesis, clearly identify a postganglionic sympathetic involvement in this response.

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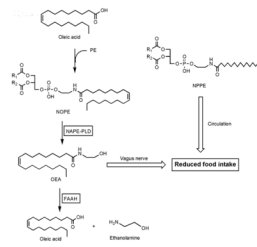


Figure 1.

Diagram illustrating OEA and NPPE metabolism in rodent small intestine. Enterocytes in duodenum and jejunum incorporate food-derived oleic acid into the phospholipid, *N*-oleoyl-phosphatidylethanolamine (NOPE) (Schwartz et al., 2008). Newly formed NOPE is hydrolyzed by *N*-acyl phosphatidylethanolamine-specific phospholipase D (NAPE-PLD) to yield OEA (Schwartz et al., 2008). OEA signaling is terminated by enzyme-mediated hydrolysis, which is primarily catalyzed by fatty acid amide hydrolase (FAAH) (Fu et al., 2007). Dietary fat stimulates the small-intestinal mobilization of NPPE through a mechanism that remains to be elucidated. Newly formed NPPE is released into the lymphatic system and reaches the brain *via* the general circulation (Gillum et al., 2008).

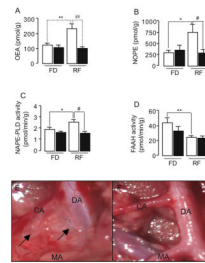


Figure 2.

Sympathetic denervation abolishes feeding-induced OEA mobilization in rat proximal small intestine. Effects of 30-min re-feeding after 24-h food deprivation (RF) or food deprivation alone (FD) on jejunal levels of (A) OEA, (B) NOPE, (C) NAPE-PLD activity, and (D) FAAH activity in sham-operated (open bars) or ganglionectomized rats (closed bars). Photomicrographs represent an intact celiac-superior mesenteric ganglion complex in a sham-operated rat (E), and the complete absence of the ganglion complex in a ganglionectomized rat (F). DA=descending aorta, CA=celiac artery, MA=mesenteric artery. Closed arrows indicate the ganglion complex. * $p < 0.05$ or ** $p < 0.01$ versus same surgical group; # $p < 0.05$ or ## $p < 0.01$ versus sham-operated controls. Results are expressed as mean \pm SEM; $n = 6$ /condition.

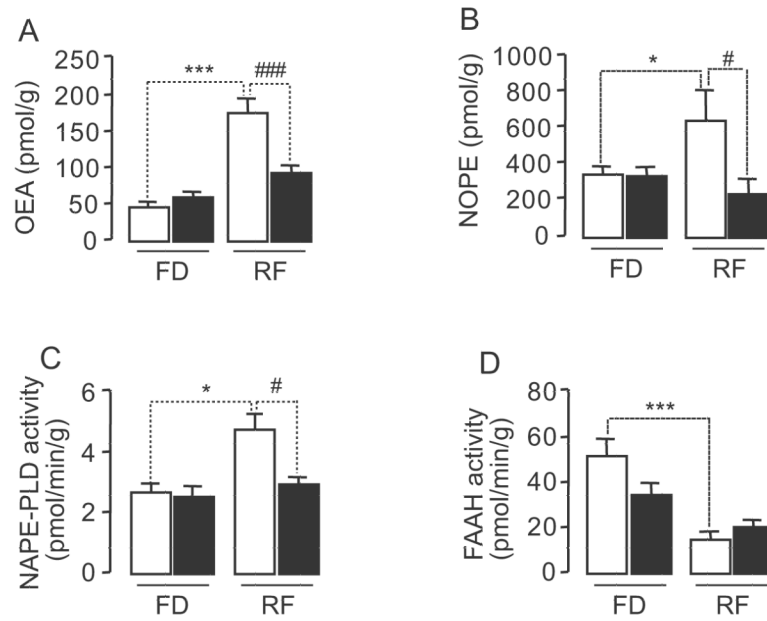


Figure 3. β_2 -Adrenergic receptor blockade abolishes feeding-induced OEA mobilization in rat proximal small intestine. Effects of 30-min re-feeding after 24-h food deprivation (RF) or food deprivation alone (FD) on jejunal levels of (A) OEA, (B) NOPE, (C) NAPE-PLD activity, and (D) FAAH activity in vehicle-treated rats (open bars) or rats treated with the β_2 receptor antagonist, ICI-118,551 (1 mg/kg, closed bars). * $p < 0.05$ or *** $p < 0.01$ versus same treatment group; # $p < 0.05$ or ### $p < 0.01$ versus sham-operated controls. Results are expressed as mean \pm SEM; $n = 6$ /condition.

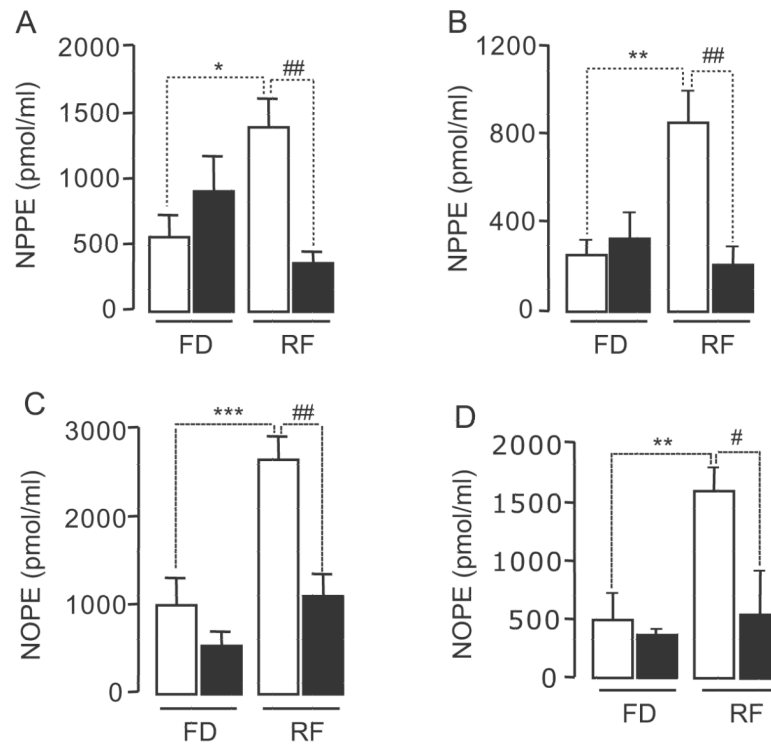


Figure 4. Sympathetic denervation lowers feeding-induced NPPE and NOPE levels in rat plasma. Effects of 30-min re-feeding after 24-h food deprivation (RF) or food deprivation alone (FD) on plasma levels of (A) NPPE, and (C) NOPE in sham-operated (open bars) or ganglionectomized rats (closed bars). Effects of 30-min re-feeding after 24-h food deprivation (RF) or food deprivation alone (FD) on plasma levels of (B) NPPE, and (D) NOPE in vehicle-treated rats (open bars) or rats treated with the β_2 receptor antagonist, ICI-118,551 (1 mg/kg, closed bars). * $p < 0.05$, ** $p < 0.01$, or *** $p < 0.001$ versus the same surgical or treatment group; # $p < 0.05$ or ## $p < 0.01$ versus sham-operated controls or those treated with vehicle. Results are expressed as mean \pm SEM; $n = 6$ /condition.

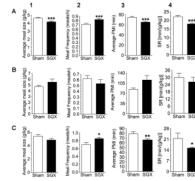


Figure 5.

Sympathetic denervation alters feeding behavior. Effects of sham operation (open bars) and ganglionectomy (closed bars) in free-feeding animals, on five-day averages of meal frequency (A1), meal size (A2), post-meal interval (PMI; A3), and satiety ratio (SR) (A4). Effects of OEA treatment in free-feeding ganglionectomized animals (closed bars) and vehicle treatment in sham-operated controls (open bars) on average 24-h meal frequency (B1), meal size (B2), PMI (B3), and SR (B4). Effects of sham-operation (open bars) and ganglionectomy (closed bars), in animals re-fed after 24-h food deprivation, on average 24-h meal frequency (C1), meal size (C2), PMI (C3), and SR (C4). Vh=vehicle, OEA=10mg/kg IP, Sham=sham operation, CGX=celiac-superior mesenteric ganglionectomy. * $p < 0.05$, ** $p < 0.01$, or *** $p < 0.001$ versus sham-operated controls. Results are expressed as mean \pm SEM. Sham operation, $n=9$; ganglionectomy, $n=10$.

Table 1

Sympathetic denervation attenuates high-fat induced OEA mobilization in rat proximal small intestine. Thirty-min of re-feeding (RF) after 24 h of food deprivation in sham-operated animals increased jejunal OEA levels compared to food deprivation alone (FD; *** $p < 0.001$). Jejunal OEA levels of ganglionectomized animals after re-feeding are attenuated when compared to those in re-feeding sham-operated animals (* $p < 0.05$); however, ganglionectomy fails to completely block increases in OEA levels of re-feeding animals versus those food-deprived (# $p < 0.05$). Results are expressed as mean \pm SEM; n = 6/condition.

	Sham	Ganglionectomy
FD	52.9 \pm 2.8	52.2 \pm 2.6
RF	85.0 \pm 6.0***	69.8 \pm 4.6*#

Table 2

Sympathetic denervation fails to affect body weights, or re-feeding of standard or high-fat chow after a fast, but modestly increases 24-h intakes. Ganglionectomy failed to affect body weights, and re-feeding of standard chow or a high-fat diet, when compared to sham-operated animals. A minor, but significant, increase in average 24-h intakes of standard chow was found in ganglionectomized animals versus sham-operated controls (* $p < 0.05$). Results are expressed as mean \pm SEM; $n = 6$ /condition.

	Body weight	24-h chow	Re-feed standard	Re-feed high-fat
Sham	294.2 \pm 4.0	78.8 \pm 1.0	5.0 \pm 0.5	5.6 \pm 0.9
Ganglionectomy	287.6 \pm 3.1	82.8 \pm 1.2*	4.5 \pm 0.4	5.1 \pm 0.2

Table 3

Administration of α -, β_1 -, or β_3 -adrenergic receptor antagonists does not affect jejunal OEA levels in re-fed rats. Thirty-min of re-feeding (RF) after 24 h of food deprivation in vehicle-treated animals increased jejunal OEA levels compared to 24-h food deprivation alone (FD). Systemic treatment with α - (prazosin), β_1 - (CGP2071), or β_3 - (SR59230A) adrenergic receptor antagonists did not alter OEA mobilization. Results are expressed as mean \pm SEM; n = 6/condition.

	α -adrenergic antagonist		β_1 -adrenergic antagonist		β_3 -adrenergic antagonist	
	V	Prazosin	V	CGP2071	V	SR59230
FD	68.1 \pm 7.4	55.4 \pm 5.4	58.2 \pm 8.3	67.1 \pm 3.5	90.3 \pm 9.5	114.2 \pm 8.9
RF	172.1 \pm 25	123.5 \pm 21.7	155.5 \pm 21.7	108 \pm 9.4	186.5 \pm 23.6	172.3 \pm 23.

Administration of β_2 -adrenergic receptor agonists does not affect jejunal OEA levels in food-deprived rats. The mixed β_1/β_2 -adrenergic receptor agonist, isoproterenol, or the selective β_2 -adrenergic agonist, salbutamol, failed to alter jejunal levels of OEA in food-deprived animals when compared to saline-treated controls. Results are expressed as mean \pm SEM; n = 6/condition.

Table 4

	Free-feed	Food deprivation					
		saline	Isoproterenol (mg/kg)			Salbutamol (μ g/kg)	
			2.5	5	10	1	5
OEA	213.8 \pm 25.2	93.4 \pm 1.6	113.2 \pm 26.5	127.4 \pm 43.4	105.7 \pm 22.9	83.4 \pm 14.9	77.8 \pm 5.6