Lymphatic diamine oxidase secretion stimulated by fat absorption is linked with histamine release

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Diamine oxidase (DAO) and histamine metabolism (8, 11). Histamine is abundant along the whole gastrointestinal (GI) tract, where it is mostly located in intestinal mucosal mast cells (MMC) (23). DAO mainly acts intracellularly (26). DAO is unevenly distributed, being absent in the small intestine, kidney, and placenta of humans and other mammals (18, 28). Intestinal DAO is synthesized continuously by mature enterocytes localized in the upper intestinal villi and is mainly associated with the basolateral aspect of enterocytes (3). DAO is stored in plasma membrane-associated vesicular structures in epithelial cells of intestine (27) and is secreted into the circulation in response to stimuli such as intravenous injection of heparin (25). As a rate-limiting enzyme in the terminal catabolism of histamine and polyamines in vivo, DAO acts as a physiological barrier against gut luminal histamine, putrescine, and cadaverine originating from food and intestinal microbiomes (26). DAO activity has been found to vary among individuals owing to polymorphisms and is potentially linked with food allergies and inflammatory disorders such as IBD (11).

Little information is available on the role of DAO in fat absorption. Our previous studies have shown that continuous duodenal infusion of triolein for 6 h gradually increases DAO activity in the intestinal lymph (34). However, why DAO activity is increased and whether it is linked to histamine release during fat absorption is far from clear. In the present study, we found that histamine is released into intestinal lymph during fat absorption and that it is probably involved in the regulation of the DAO secretion into lymph. This regulatory
effect of histamine is mainly mediated through H₄R and not through the other histamine receptors such as H₁R and H₂R. The results demonstrate for the first time the link between histamine and its degrading enzyme DAO released during fat absorption, and this may provide insight into our understanding of the mechanisms of food allergies and IBD.

MATERIALS AND METHODS

Materials

Liposyn II 20% was purchased from Hospira (Lake Forest, IL), and trilinolein from Nu-Chek-Prep (Elysian, MN). Histamine hydrochloride; the HR antagonists H₁R-pyrilamine maleate, H₂R-ranitidine, H₃R-thioperamide maleate, and H₄R-JNJ777120; and tricaprylin were purchased from Sigma-Aldrich Chemicals (St. Louis, MO). The radioactive [³H]putrescine was obtained from Amersham Biosciences (Piscataway, NJ). The histamine ELISA kit was purchased from Neogen (product no. 409010, Lexington, KY). The goat anti-mouse DAO polyclonal antibody (sc-67660) was from Santa Cruz Biotechnology (Santa Cruz, CA), and peroxidase-conjugated anti-goat secondary antibody was from DAKO (Glostrup, Denmark).

Experiments

Animals. Adult male Sprague-Dawley rats, weighing 240–350 g (Harlan, Indianapolis, IN), were used. Animals were allowed to acclimate to our animal facility for at least 2 wk prior to the experiment. During this period, the animals were fed rodent chow at libitum and housed in a room with a 12:12-h light-dark cycle. Both the temperature and the humidity of the room were maintained within the range 70–74°F and 40–60%, respectively.

Lymph and duodenal cannulation. The surgical procedure and the postoperative care have been described previously as described (16). All procedures were approved by the University of Cincinnati Internal Animal Care and Use Committee and complied with the NIH Guide for the Care and Use of Laboratory Animals. Briefly, animals were fasted overnight before surgery. Under isoflurane anesthesia, the superior mesenteric lymph duct was cannulated with soft vinyl tubing (0.8 mm OD) according to the method described (16). A drop of cyanoacrylate glue (Krazy Glue, New York, NY) was used to secure the lymph cannula. Intraduodenal (i.d.) cannulation was performed by inserting a silicone tube (1.6 mm OD) ~2 cm into the duodenum via a fundal incision of the stomach. The tubing was secured by a transmural suture in the duodenum, and the fundal incision was closed by a purse-string suture. Postoperatively, the animals were kept in Bolllman restraining cages. Although the animals were restrained, they had considerable freedom to move forward, backward, and sideways. The restraining was necessary to prevent the animals from chewing and damaging the cannula. The animals were infused id with 5% glucose in saline (145 mM NaCl, 4 mM KCl, and 0.28 M glucose). Beginning 16 h before the nutrient study (which occurred on the following morning), the 5% glucose in saline was switched to saline alone and was infused overnight at a rate of 3 ml/h until the following morning, when the saline solution was replaced with the nutrient infusate described below. Fasting lymph was collected for 1 h before the start of the nutrient infusion. The nutrient infusate was given as a single bolus through the duodenal infusion cannula. Lymph was collected continuously at 30-min intervals during the first hour after the nutrient infusion and hourly thereafter over the remaining 5-h time course.

Preparation of nutrient infusate. Two groups of animals were tested and infused id with a single bolus of 3 ml of normal saline (control group) or lipid, consisting of 2.215 ml of Liposyn II (20%) + 0.785 ml of saline with the caloric content of 4.43 kcal/3 ml. Following the nutrient bolus infusion, the continuous id infusion of saline was temporarily halted for 30 min to avoid overdistension of the small intestine and saline infusion continued afterward. Liposyn II 20% consists of a 50:50 blend of safflower and soybean oil with a caloric content of 2 kcal/ml. The caloric content of the full dose (4.4 kcal) of fat was equivalent to half of the total daily fat intake of the rat.

To determine the importance of the dose and chain length of fatty acids on lymphatic DAO secretion, the following groups of animals were studied. Four groups of animals were id infused different amounts of Liposyn II (0.55, 1.1, 2.2, and 4.4 kcal/3 ml) to detect the dose effect. In addition, the effect of long-chain triacylglycerol (TG) trilinolein (C18:2, n-6 TG), the major composite (65.8%) of Liposyn II, and medium-chain TG tricaprylin (C8:0) was also studied with id bolus (3 ml) infusions of phosphate-buffered, taurocholate-stabilized emulsions containing 120 µmol trilinolein or 120 µmol tricaprylin. The vehicle control emulsion contained 8.7 µmol egg phosphatidylcholine, 7.8 µmol cholesterol, and 57 µmol sodium taurocholate, sonicated in 3 ml phosphate-buffered saline (pH 6.4) (12).

To determine the role of histamine and the histamine antagonists, 10 mg/kg histamine was intraperitoneally (ip) administered into fasting rats and then lymph was collected every 10 min for 60 min. The dose of histamine (10 mg/kg) was calculated according to the previous reports (33). The four types of potent and specific HR antagonists pyrilamine maleate (H₁B, 10 mg/kg), ranitidine (H₂B, 8 mg/kg), thioperamide maleate (H₃B, 10 mg/kg), or JNJ777120 (H₄B 37 mg/kg) were ip administered into fasting rats 30–45 min before id bolus infusion of 3 ml (4.4 kcal) Liposyn II followed by saline infusion. Then lymph samples were collected at 30-min intervals for 180 min.

DAO activity measurement. DAO activity was measured by a radiometric assay described by Forget et al. (7) with slight modifications. Briefly, the reaction mixture consisted of a 50-µl enzyme sample and a 50-µl substrate mixture. The substrate mixture was a combination of cold and labeled putrescine prepared in 0.1 M sodium phosphate buffer at pH 7.2 with a ratio of 10 µl of [³H]putrescine per 1,000 µl of 0.9 mM unlabeled putrescine. The final concentration of substrate putrescine was 0.45 mM with the total count of ~0.8 million dpm. The reaction mixture was then incubated at 37°C for 30 min. The reaction was stopped by the addition of 10 µl of 0.1 M aminoguanidine, an inhibitor of DAO, followed by the addition of 50 µl sodium carbonate pH 12.2 (187.5 mM final). The reaction product Δ₁-pyridine was extracted twice with 500 µl of toluene. The radioactivity was determined by liquid scintillation counting. Activities were shown to be proportional to time and quantity of putrescine and enzyme. The DAO activities were calculated in milliliters per milliliter of lymph (mU/ml), where 1 unit oxidized 1 µmol of putrescine per hour at 37°C.

Western blot analysis. The lymph samples (10 µl each well) were loaded onto 4–20% polyacrylamide gradient gel and electrotransferred to polyvinylidene difluoride membranes. Following transfer, the membranes were treated with blocking reagent [5% nonfat milk and 0.1% Tween 20 in Tris-buffered saline (TBS)], after which the membranes were incubated with goat anti-mouse DAO antibody (diluted 1:500) at 4°C overnight. After rinsing with TBS with 0.1% Tween 20, the membranes were further incubated with peroxidase-conjugated anti-goat secondary antibody at 1:2,000 for 1 h and then developed with ECL Western blot detection kit (GE Healthcare) according to the manufacturer’s protocol and were quantified with the Total Lab Quant Analysis Software (TL100, FOTOLOGYNE, Harland, WI).

ELISA analysis of lymphatic histamine. Lymph histamine levels were measured with histamine ELISA kits. Briefly, 50 µl of lymph was mixed with 50 µl of enzyme conjugate at room temperature for 45 min. After three times rinsing with 300 µl washing buffer, 150 µl of substrate was added to each well of the plate, allowing the plate to incubate at room temperature for 30 min; then the plate was read in a microplate reader with a 650-nm filter. The histamine levels were calculated according to the manufacturer’s instruction.

Chemical assays of protein in lymph. Protein concentration was measured by the Bradford method with bovine serum albumin (BSA).
as a standard. Standard curves were constructed, and concentrations of the samples were calculated from the standard curves.

Statistical Analysis

The data shown are mean values ± SE. To compare groups through the 3-h or 6-h infusion, a two-way repeated-measures ANOVA was used. For comparison of data with two independent variables, a two-way ANOVA was used. A t-test was used for the rest of the analyses for comparing only two groups. Differences between treatments were considered significant when $P < 0.05$.

RESULTS

Effect of Duodenal Feeding of Nutrients on Lymphatic DAO Activity and DAO Output

DAO activities in the fasting lymph of lipid or saline-infused rats were comparable (25.7 ± 0.8 and 22.4 ± 0.4 mU/ml, respectively; $P > 0.05$). There was, however, a significant increase in DAO activity in lymph from 0.5 to 3 h after bolus infusion of 3 ml Liposyn II (4.43 kcal) with a peak of 115.4 ± 21.6 mU/ml at 1 h, which is a ~3.5-fold increase compared with control ($n = 6$, $P < 0.01$). Animals that received saline showed no significant alterations in lymph DAO activity throughout the entire 6 h (Fig. 1A).

When taking into account the lymph flow rate, which was not significantly altered relative to the saline control group (data not shown), lymphatic DAO output (the product of DAO activity and lymph flow rate) during 3-h period was significantly increased in the lipid group (435.8 ± 31.0 in lipid, vs. 178.8 ± 12.9 mU in saline group, $n = 6$, $P < 0.01$) (Fig. 1B). Our data therefore indicate that a bolus infusion of lipid increases intestinal DAO activity and DAO output in lymph.

To determine whether the increased DAO activity was due to increased DAO protein secretion into lymph, we conducted a Western blot analysis. As shown in Fig. 1C, the lymphatic DAO content increased in Liposyn II-infused rats, starting at 0.5 h, peaking at 1 h, and returning to the fasting level by 3 h postinfusion, whereas no increase in DAO content was observed in the saline group. The calculation of the density of the bands in the Western blot showed a 3.7 ± 0.3-fold increase at the peak DAO secretion at 1 h (Fig. 1D), which was consistent with the extent of the increase in the lymph DAO activity. This result indicates that fat absorption induces DAO secretion into lymph, thereby increasing lymphatic DAO activity.

Amount of Fat Affects DAO Activity

To determine the effect of the amount of fat fed on DAO activity, increasing caloric doses (0.55, 1.1, 2.2, and 4.4 kcal, all in 3 ml) of Liposyn II were infused into the duodenum. As shown in Fig. 2A, the 2.2- and 4.4-kcal doses raised the levels of lymphatic DAO activity above that of the saline control at the 1 h following lipid infusion (1.9 ± 0.2 and 2.8 ± 0.8-fold over saline, respectively). The differences in DAO activity among 4.4-, 2.2-, 1.1-, and 0.55-kcal doses as well as 2.2 vs. 1.1- and 0.55-kcal dose groups were all significant ($P < 0.05$, $n = 6$), showing a dose-dependent pattern (Fig. 2A). In addition, cumulative DAO secretion, calculated as the area under the curve (AUC) over the 3-h lymph collection period (Fig. 2B), increased in response to larger amounts of dietary lipid: from 114.3 ± 2.4 for the 1.1-kcal lipid dose to 177.9 ± 9.9 for the 4.4-kcal lipid dose ($P < 0.05$), exhibiting a dose-dependent relationship.

Type of Fat Affects DAO Activity

Our previous study has shown that the 18-carbon chain (C18) triolein more effectively stimulated DAO release into lymph than tricaprylin (C8), when continuously given id in
equimolar amounts (34). In the present study we further examined the effect of the long-chain TG trilinolein on DAO activity because linoleic acid (C18:2, n-6) is the major component (65.8%) of Liposyn II. Figure 3 shows that after the id bolus infusion of 120 mol trilinolein emulsion plus vehicle, lymphatic DAO activity significantly increased, with the peak value of 130.6 (41.8 mU/ml vs. the vehicle control (lipid emulsion containing phosphatidylcholine and sodium taurocholate) value of 15.5 ± 4.3 mU/ml (n = 6, P < 0.01). The DAO activity returned to fasting level by 3 h. In contrast, the bolus infusion of 120 μmol tricaprylin (C8) emulsion produced only a marginal increase in lymphatic DAO activity with a value of 35.5 ± 11.0 mU/ml (P < 0.05, vs. control, n = 6) at 1 h postinfusion (Fig. 3). The DAO output in lymph was significantly increased in trilinolein-infused rats compared with the tricaprylin group (data not shown). These results confirmed that long-chain TG (both triolein studied previous and trilinolein in this study) is more effective than medium-chain TG in inducing an increase in DAO secretion.

Release of Histamine into Lymph After Nutrient Infusion

We hypothesized that the infusion of a lipid meal causes the secretion of histamine, which in turn stimulates the secretion of DAO to control the unwanted effects of histamine. To test this hypothesis, we measured lymphatic histamine concentration in Liposyn II and trilinolein-infused rats. As shown in Fig. 4, A and B, a significant 1.1-fold and 1.3-fold increase of the peak histamine level was observed after Liposyn II or trilinolein infusion, respectively. The output peaked at 1 h postinfusion in both groups (38.5 ± 1.9 ng/ml in the Liposyn II group vs. 18.5 ± 3.2 ng/ml in saline group, n = 6, P < 0.01, and 35.0 ± 3.6 in the trilinolein group vs. 15.0 ± 1.6 ng/ml in vehicle control rats, P < 0.05). Our data demonstrate that id infusion

![Fig. 2. Dose-dependent effect of Liposyn II on lymphatic DAO activity. A: responses to 4 infused lipid doses (0.55, 1.1, 2.2, and 4.4 kcal). B: area under the curves (AUC). Values are means ± SE. *P < 0.05, **P < 0.01, 4.4-kcal dose vs. the rest doses of Liposyn II, #P < 0.05, ##P < 0.01 vs. 1.1 and 0.55 kcal.](image)

![Fig. 3. Comparison of the effect of long-chain triacylglycerol (TG) trilinolein with medium-chain TG tricaprylin on DAO activity. Values are means ± SE, n = 6. **P < 0.01 vs. vehicle emulsion; ##P < 0.01 vs. tricaprylin.](image)

![Fig. 4. Lymphatic histamine concentrations after id infusion of Liposyn II (4.4 kcal) (A) or 120 μmol trilinolein (B). *P < 0.05, **P < 0.01 vs. saline or vehicle, n = 6.](image)
of Liposyn II containing mainly n-6 long-chain fatty acids (LCFA), or pure trilinolein induces histamine release into lymph. This data is suggestive of the induction of DAO release by histamine, which in turn is tied to active intestinal fat absorption.

**Effect of Histamine on Lymphatic DAO Activity**

To address the question of whether histamine is a mediator of DAO secretion, we examined the effect of histamine on lymphatic DAO activity by ip injection of histamine (10 mg/kg) into fasting rats. As shown in Fig. 5A, 10 mg/kg ip histamine resulted in a quick increase in the lymphatic histamine level up to ~200 ng/ml at 10 min and a subsequent decline to baseline by 20 min. At the same time, ip histamine substantially increased the lymphatic DAO activity by 1.1-fold. The increase lasted until 30 min postinjection with a peak at 20 min (Fig. 5B). Our data suggest a close relationship between the increase of lymphatic histamine and DAO activity. It is noteworthy that the effect of ip histamine on DAO activity is transient; after 30 min when the lymph histamine concentration returned to the basal level, lymphatic DAO activity was also restored to the baseline (Fig. 5B).

The effectiveness of ip histamine was confirmed by dramatic elevations in mesenteric lymph flow rate and lymphatic protein transport. As shown in Fig. 5, C and D, the maximal lymph flow rate and the peak lymphatic protein concentration increased by 1.36-fold and 6.43-fold at 20 and 10 min after ip histamine, respectively. Our result confirms the increase in vascular permeability by histamine and suggests that it mediates the secretion of DAO.

**Effect of Histamine Receptor Antagonists on Lymphatic DAO Activity**

To further examine whether endogenous histamine mediates DAO secretion during fat absorption, four types of specific HR blockers [pyrilamine (H1B), ranitidine (H2B), thioperamide (H3B) or JNJ 7777120 (H4B)] were ip injected at 30–45 min before id infusion of Liposyn II (4.4 kcal/3 ml) into rats, based on the reports that all four types of HRs are distributed along GI tract in rats (5, 11). As shown in Fig. 6A, neither pyrilamine nor ranitidine nor thioperamide, nor pyrilamine plus ranitidine, inhibited the lipid-induced increase of DAO activity. The H4B JNJ 7777120, however, significantly reduced the DAO activity, the AUC over 3 h following lipid infusion was reduced by 65.9% from 827.9 ± 15.6 to 282.1 ± 7.9 mU·ml⁻¹·h⁻¹ (P < 0.01, n = 6). Western blot analysis further confirmed that the ip H4B decreased the lymph DAO secretion by ~64.3 and 33.3% at 1 and 1.5 h after lipid infusion, respectively (Fig. 6, B and C). These results indicate that histamine, through H4R, is involved in mediating fat-induced DAO secretion into lymph.

**Effect of Lipid Infusion and Histamine Receptor Antagonists on Lymph Flow Rate and Protein Transport**

The lymph flow rates were not significantly altered in Liposyn II-infused rats, nor in trilinolein and tricaprylin-in-
fused rats and in the HRs antagonist-administered groups (data not shown), suggesting a complex regulation of lymph flow rate, in which histamine may not be the key factor. On the contrary, id Liposyn II caused a significant increase in lymph protein concentrations as well as protein flux (a product of lymph protein concentration and lymph flow rate), starting at 1 h, peaking at 2 h, and then gradually declining by 4 h following lipid infusion (Fig. 7A). In addition, the lymphatic protein flux over 3 h following lipid infusion showed a significant dose-dependent increase from 27.5 ± 8.6 mg·h in 0.55 kcal to 93.0 ± 13.9 mg·h (P < 0.01, n = 5–6) in the 4.4-kcal

**Fig. 6.** Effect of ip histamine receptor blockers (HB) on Liposyn II-induced lymph DAO activity. A: lymph DAO activity after ip H1B–H3B; n = 6. B: representative blot shows the DAO content in lymph at fasting and after ip H4B followed by the infusion of Liposyn II. C: quantitative data of the DAO secretion in lymph. Means ± SE, *P < 0.05, **P < 0.01 vs. control.

Fig. 7. Lymphatic protein flux after id bolus infusion of Liposyn II (4.4 kcal) (A); AUC of different doses of Liposyn II (0.55, 1.1, 2.2, 4.4 kcal) (B). Lymphatic protein concentrations after infusion of long-chain TG trilinolein or medium-chain TG tricaprylin (C) or after ip H1B, H2B, H1B+H2B, and H3B followed by Liposyn II infusion (D). Means ± SE, n = 6, *P < 0.05, **P < 0.01 vs. controls, #P < 0.05, ##P < 0.01 vs. 1.1 and 0.55 kcal.
lipid-infused group (Fig. 7B). Furthermore, infusion of tricaproin
lein induced a significant increase in lymphatic protein trans-
port, whereas tricaprylin did not (Fig. 7C). These data indicate
that the absorption of Liposyn II, mainly containing linoleic
acid, increases lymphatic protein transport in a dose-dependent
manner.

Histamine, via increasing the intestinal vascular permeabil-
ity (9, 21), increases lymph protein transport, as shown in Fig.
5D. To determine the participation of histamine in the in-
creased lymph protein transport during fat absorption, HR
antagonists were ip administered before id infusion of Liposyn
II. As shown in Fig. 7D, ip pyrilamine (H1B) or ranitidine
(H2B), or thioperamide (H3B) or JNJ 7777120 (H4B) (data not
shown) did not significantly alter the lymph protein concen-
trations nor the protein flux in response to lipid infusion, except
for a 35% (P < 0.05, n = 6) and a 39% (P < 0.05, n = 6)
reduction at 90 min after lipid infusion in ranitidine or thio-
peramide pretreated rats. However, a significant 67.8% (P <
0.01, n = 6) reduction of lymph protein concentrations was
observed in pyrilamine (H1B, 10 mg/kg ip) plus ranitidine
(H2B, 8 mg/kg) administered animals (Fig. 7D). Our data
demonstrate that lymphatic protein transport increases in re-
response to fat absorption; histamine, via increasing the intestinal
vascular permeability, may play a role in the regulation of
fat-induced lymph protein transport through additive effects on
H1R and H2R. These data are consistent with the previous
finding showing that histamine H1R antagonists pyrilamine
plus H2R antagonists burimamide prevented intestinal lymph
protein transport during id olive oil feeding (32). Thus the
histamine receptor (H4R) involved in DAO secretion stimu-
lated by fat absorption is different from the histamine receptors
(H1R + H2R) involved in the increase in lymph protein flux.

DISCUSSION

In the present study, using the conscious lymph fistula rat
model, we demonstrated that fat absorption induces histamine
release as well as DAO secretion in mesenteric lymph. The
basal level of plasma DAO is very low (25). The mesenteric
lymph collected in our experiment had not entered the circu-
lation. Therefore, the DAO in the mesenteric lymph, which had
not been metabolized by the liver, directly reflects the secretion
of intestinal DAO. In addition, histamine appears to be rapidly
metabolized in the circulation because its half-life is estimated
at <1 min (24). The measurement of both molecules using
mesenteric lymph provides a more direct way to study the in
vivo secretory function of intestinal villi. The released histo-
amine induced by fat absorption in turn stimulates the release of
DAO through the H4R. This is a first demonstration of the
release of histamine being linked to the release of DAO by fat
absorption, but the physiological function of this link is far
from clear.

It is well known that fat absorption in the small intestine
induces a series of physiological reactions including increases in
mesenteric blood flow (1, 6) and vascular permeability (9, 21)
as well as alterations in intestinal motility (15). These changes may
involve the participation of the whole neural-hormonal-immune
network including vagal nerves, enteric neurons, GI hormones
such as cholecystokinin, neurotensin, and GLP-1 (9, 22), as well
as the intestinal immune system (10, 19). Our present study
demonstrated that histamine is released during fat absorption,
suggesting that histamine may play a role in this process. Since
gut histamine is mostly located in intestinal mast cells (23), the
histamine released during fat absorption may originate from the
intestinal MMC. This hypothesis is supported by our recent
studies in which we demonstrated that the intestinal MMC are
activated after id infusion of Liposyn II, causing a peak of
~20-fold increase in rat MMC protease II (RMCPII), a specific
marker of MMC degranulation, in intestinal lymph at 1 h after
lipid infusion (10).

The role of histamine during fat absorption is probably
related to its vasoactive properties. Histamine increases both
mesenteric blood flow and vascular permeability (9, 21, 32).
This vasoactive effect of histamine is further confirmed in the
present study, in which ip histamine increased both lymph flow
rate and lymph protein influx (Fig. 5, C and D). However, in
response to the bolus lipid infusion, no significant increase in
lymph flow rate was observed, nor was it decreased after
pretreatment with ip HR antagonists. This apparent lack of
response may be related to our experimental design. Consid-
ering that bolus infusion of 3 ml of the emulsion may cause
overdistension of the small intestine, we temporarily stopped
the saline infusion (3 ml/h) after id bolus for 30 min before
resuming. Despite the lack of the effect on lymph flow, the
significant increase in lymph protein transport in response to
Liposyn II infusion clearly hints the involvement of histamine
in regulating vascular permeability. The significant inhibition
of the lymph protein flux in response to the administration of
H1R and H2R blockers (Fig. 7D) supports this view. The
physiological significance of the elevation of the lymphatic
protein transport during fat absorption is unclear. It has been
reported that the integrity of the intestinal epithelium is com-
promised during fat absorption (13, 30). In the present study,
we showed that it is tricaproin, but not tricaprylin, that in-
creased protein transport in lymph, further indicating that
LCFA rather than medium-chain fatty acids (MCFs) induce
protein influx into lymph. LCFA are mostly transported as
chylomicrons (CM) in lymph, whereas the majority of MCFs is
transported via the portal circulation (29). Dietary fats (mainly
LCFA) are digested and absorbed by the enterocytes and
packaged into CM. The LCFA containing CMs are then trans-
ported into the expanded intercellular space. The basement
membrane of the intestinal villi is obviously a barrier to the
passage of the CM from the intercellular space to the lamina
propria. We have previously shown that CM travel into the
lamina propria by breakages of the basement membrane (30).
It is therefore plausible to assume that histamine induces an
increase in the hydration (expansion) of the interstitial matrix
in the lamina propria by increasing vascular permeability, thus
facilitating the diffusion the CM particles into the central
lacteals (31). Histamine may also participate in the gut mucosal
repair during fat absorption as it does during intestinal isch-
emia-reperfusion-induced injury (8). Our previous study re-
vealed that the jejunal epithelium is temporarily injured and the
“injury” is presumably repaired rapidly during fat absorption
(13). It is conceivable that the role of histamine in healing the
ischemia-reperfusion-induced mucosal damage may also apply
to the temporary mucosal damage induced by fat absorption.

Although histamine certainly has important physiological
function during fat absorption, it is equally important to have
an efficient system to deactivate it to avoid unwanted side
effects. Studies have shown that unwanted or excessive hista-
mine is detrimental and can lead to a number of pathophysiological conditions such as anaphylaxis, food allergy, and IBD (17, 26, 35). Enzymatic inactivation of circulating histamine is exclusively mediated by the degrading enzyme DAO. Our previous and present studies demonstrate that in the rat, id infusion of lipid, either continuously (34) or as a bolus, induces a significant increase of DAO secretion into intestinal lymph. Given the fact that DAO is synthesized and stored in enterocytes (3), it is conceivable that the intestinal mucosa contributes to the large increase in enzyme activity in the intestinal lymph. The rapid DAO release within 1 h in our present study probably reflects the release of the stored DAO rather than an increase in DAO synthesis. In addition, we have found that the lymphatic DAO activity (a measure of the secretion) depends on the amount fed and also types of fat infused: long-chain TG (trilinolein) promotes a significant release of histamine than medium-chain TG (tricaprylin). We propose that CM transport plays a role in DAO release by the intestinal mucosa and the purpose is to maintain optimal histamine concentration to mediate its many roles in intestinal fat absorption.

Lastly, in this study, we found that lymphatic histamine and DAO both peaked at the same time during fat absorption, suggesting a potential close relationship between the two. The present observation (Fig. 5B) that ip histamine (10 mg/kg), similar to id histamine (0.15–1.2 mmol/kg) administration in a previous study (33), induced the release of DAO into intestinal lymph further confirmed this close relationship, suggesting that histamine could be an endogenous mediator in the secretion of DAO to safeguard against deleterious effects of the excessive histamine secretion during fat absorption. Another interesting observation of this study is the particular histamine receptor involved in DAO release. All four classes of HRs (H1R, H2R, H3R, and H4R) are present in the rat small intestine (5, 14), distributing in intestinal musculature, mucosal epithelium, enteric nervous system, and immune/inflammatory cells (5). In this study, we found that only H4R antagonist, but not the other three HR blockers, inhibited fat-induced DAO secretion. The underlying mechanism of how activation of H4R regulates the fat-induced secretion of DAO is unknown. Recent studies revealed that H4R is primarily expressed on inflammatory/immune cells including eosinophils, mast cells, basophils, dendritic cells, and T cells (5). H4R expression is also present in the myenteric neurons of the rodent GI tract (20), but there is no report on rat enterocytes. The direct connection between H4R and DAO secretion by the enterocytes in response to lipid warrants further studies.

In conclusion, we have reported a number of interesting observations in this study. First, we demonstrated that histamine release is stimulated by fat absorption and this in turn stimulates the release of DAO. Second, this action of histamine is mediated through the H4R, which is different from the role of increasing protein flux mediated by the H1R and H2R. Histamine, thus, may play an important role in the CM trafficking from the intercellular space to the lamina propria and then subsequently into the lacteals during fat absorption. The DAO secreted during fat absorption may act as deamination mechanism to counterbalance an excess of histamine to exert unwanted side effects. It is tempting to speculate that an imbalance between the histamine-DAO system may be involved in intestinal disorders such as inflammatory bowel disorder.

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DISCLOSURES
No conflicts of interest, financial or otherwise, are declared by the author(s).

AUTHOR CONTRIBUTIONS
Y.J. conception and design of research; Y.J., Y.S., X.L., C.Z., Q.Y., and M.X. performed experiments; Y.J., Y.S., X.L., and C.Z. analyzed data; Y.J. and Y.S. interpreted results of experiments; Y.J. and Y.S. prepared figures; Y.J. drafted manuscript; Y.J., A.W., W.L., and P.T. edited and revised manuscript; Y.J., W.L., and P.T. approved final version of manuscript.

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