

# Satiety factor oleoylethanolamide recruits the brain histaminergic system to inhibit food intake

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**Key factors driving eating behavior are hunger and satiety, which are controlled by a complex interplay of central neurotransmitter systems and peripheral stimuli. The lipid-derived messenger oleoylethanolamide (OEA) is released by enterocytes in response to fat intake and indirectly signals satiety to hypothalamic nuclei. Brain histamine is released during the appetitive phase to provide a high level of arousal in anticipation of feeding, and mediates satiety. However, despite the possible functional overlap of satiety signals, it is not known whether histamine participates in OEA-induced hypophagia. Using different experimental settings and diets, we report that the anorexiant effect of OEA is significantly attenuated in mice deficient in the histamine-synthesizing enzyme histidine decarboxylase (HDC-KO) or acutely depleted of histamine via interocerebroventricular infusion of the HDC blocker  $\alpha$ -fluoromethylhistidine ( $\alpha$ -FMH).  $\alpha$ -FMH abolished OEA-induced early occurrence of satiety onset while increasing histamine release in the CNS with an H<sub>3</sub> receptor antagonist-increased hypophagia. OEA augmented histamine release in the cortex of fasted mice within a time window compatible to its anorexic effects. OEA also increased c-Fos expression in the oxytocin neurons of the paraventricular nuclei of WT but not HDC-KO mice. The density of c-Fos immunoreactive neurons in other brain regions that receive histaminergic innervation and participate in the expression of feeding behavior was comparable in OEA-treated WT and HDC-KO mice. Our results demonstrate that OEA requires the integrity of the brain histamine system to fully exert its hypophagic effect and that the oxytocin neuron-rich nuclei are the likely hypothalamic area where brain histamine influences the central effects of OEA.**

histamine receptors | behavioral satiety sequence | BSS | paraventricular hypothalamic nuclei | PVN

Eating behavior is regulated by central neurotransmitter systems and peripheral stimuli that interact to change the behavioral state and concur to alter homeostatic aspects of appetite and energy expenditure. The fatty acid amide oleoylethanolamide (OEA) is released by the small intestine in a stimulus-dependent manner and suppresses food intake by activating peroxisome proliferator-activated receptor- $\alpha$  (PPAR- $\alpha$ ) (1). Systemic administration of OEA induces c-Fos mRNA expression through the vagus nerve to the nucleus of the solitary tract (NST), supraoptic nuclei (SON), and paraventricular hypothalamic nuclei (PVN) and increases the expression of oxytocin (2, 3), which is involved in the central coordination of homeostatic signals and feeding behavior (4). However, it is not known whether OEA recruits other neurotransmitter systems in the brain to reduce food intake. Histaminergic neurons are clustered in the hypothalamic tuberomammillary nuclei (TMN). They send projections organized in functionally distinct circuits impinging on different brain regions (5), and their firing frequency changes according to the behavioral state (6). Brain histamine plays a fundamental role in eating

behavior as it induces loss of appetite and has long been considered a satiety signal that is released during food intake (7). Early studies showed that treatments increasing brain histamine availability or activating histamine H<sub>1</sub> receptor (H<sub>1</sub>R) in the CNS suppressed food intake (8–10), and enhanced c-Fos-like immunoreactivity in the PVN (10), whereas H<sub>1</sub>R blockade in the ventromedial hypothalamus (VMH) increased both meal size and duration, and suppressed the activity of glucose-responding neurons (11).

With these considerations in mind, we examined the hypothesis that peripherally administered OEA engages histamine signaling in the brain. We assessed the effect of OEA on cumulative food intake as well as on the expression of the behavioral satiety sequence (BSS) in mice defective in brain histamine, either because they lack the histamine-synthesizing enzyme histidine decarboxylase (HDC) or because acutely deprived of histamine by interocerebroventricular (i.c.v.) infusion of HDC inhibitor  $\alpha$ -fluoromethylhistidine ( $\alpha$ -FMH). We also assessed the effect of OEA following systemic administration of the H<sub>3</sub>R antagonist ABT-239 that transiently increases

## Significance

Several endogenous molecules contribute to the building of a complex network of neural and hormonal signals that align food intake and energy expenditure. Cerebral histamine works as a satiety factor by activating histamine H<sub>1</sub> receptor (H<sub>1</sub>R) in specific hypothalamic nuclei. Indeed, atypical antipsychotics presumably cause obesity by targeting brain H<sub>1</sub>R. The endogenous lipid messenger oleoylethanolamide (OEA) mediates fat-induced satiety by engaging sensory fibers of the vagus nerve that project centrally. We find that depletion of brain histamine blunts OEA-induced hypophagia in mice. Our study uncovers previously unidentified neural signaling mechanisms involved in the anorectic action of OEA. Our data offer new perspectives for developing more effective and safer pharmacotherapies to treat obesity and ameliorate the profile of centrally acting drugs.

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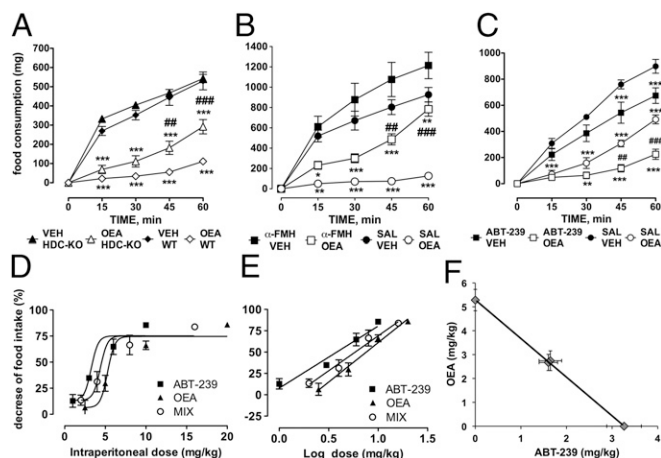
brain histamine release (12). Finally, we determined changes in neuronal activity by assessing the pattern of c-Fos expression induced by OEA in hypothalamic brain regions controlling feeding behavior and in brain regions regulating the motivational aspect of foraging in both HDC-KO and WT littermates. Our findings revealed that the anorexic effects of OEA are blunted in brain histamine-deficient mice, and establish new functional connections between peripherally acting hypophagic signals and brain histamine neurotransmission.

## Results

### Interaction Between Brain Histamine and OEA on Food Consumption.

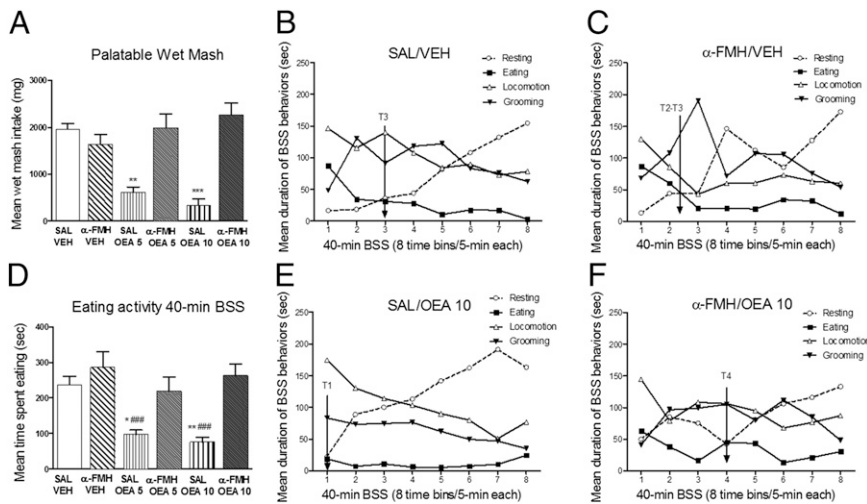
To test whether the integrity of the histaminergic system contributes to the anorexic effect of OEA, we used HDC-KO mice, which are unable to synthesize histamine. OEA was injected i.p. at 10 mg/kg to induce long-lasting appetite suppression (1); it does not readily enter the CNS (13) and does not cause taste aversion (14). Controls received equivalent volumes of the vehicle. OEA caused a profound reduction of the total amount of food consumed by WT mice compared with vehicle-treated littermates within the first 60 min after injection ( $P < 0.001$ ; Fig. 1A). Two-way ANOVA revealed an overall significant difference between groups [ $F_{(time)4,170} = 133.14$ ,  $P < 0.0001$ ;  $F_{(treatment)3,170} = 185.64$ ,  $P < 0.0001$ ;  $F_{(time \times treatment)12,170} = 12.93$ ,  $P > 0.0001$ ]. As expected, vehicle-treated HDC-KO mice consumed comparable amounts of food with respect to WT animals (histamine-deficient mice are not hyperphagic, nor

obese up to 12 wk of age) (15), whereas the anorexic effect of OEA was significantly diminished in HDC-KO mice ( $P < 0.01$  at 45 min;  $P < 0.001$  at 60 min). The effect of OEA was short lived as no difference in food consumption was observed among experimental groups 4 h after OEA injection. Histamine-deficient mice develop metabolic changes with age (15). Hence, to exclude the involvement of compensatory mechanisms in HDC-KO mice, we measured food intake also in CD1 mice that received an i.c.v. infusion of the HDC suicide inhibitor  $\alpha$ -FMH at a dose that abolishes basal and stimulated histamine release (see Fig. 3B). Mice were tested at lights off (1900 h), when they have a greater incentive to eat and the activity of histamine neurons is higher (16). Two-way ANOVA revealed an overall significant difference among groups [ $F_{(treatment)3,104} = 20.67$ ,  $P < 0.0001$ ;  $F_{(time)4,104} = 121.36$ ,  $P < 0.00001$ ;  $F_{(time \times treatment)12,104} = 13.30$ ,  $P < 0.0001$ ]. OEA caused a profound reduction in the total amount of food consumed by CD1 mice that received an i.c.v. infusion of saline (Fig. 1B). Mice in the  $\alpha$ -FMH/vehicle group tended to eat more than the saline/vehicle animals, although this effect did not reach statistical significance. In  $\alpha$ -FMH-treated mice, OEA-induced hypophagia was significantly less prominent than in the saline/OEA group. The effect reached statistical significance 45 min after OEA injection ( $P < 0.01$ ). The overall trend is very similar to the results obtained with HDC-KO mice during the lights-on phase, although during the lights-off phase all experimental groups ate consistently larger amounts of food. We then argued that, if OEA modulates feeding by recruiting the central histamine system, augmenting histamine release pharmacologically would potentiate the anorexic effect of exogenous OEA. We measured food-consumption in 12 h-fasted CD1 mice treated with a combination of OEA (10 mg/kg) and ABT-239 (3 mg/kg), an H<sub>3</sub>R antagonist that increases histamine release by blocking H<sub>3</sub> autoreceptors (12). Two-way ANOVA followed by Bonferroni's post hoc test revealed a significant difference between groups [ $F_{(time)4,104} = 392.5$ ,  $P < 0.0001$ ;  $F_{(treatment)3,104} = 31.81$ ,  $P < 0.0001$ ;  $F_{(time \times treatment)12,104} = 29.04$ ,  $P < 0.0001$ ]. Each compound significantly decreased food intake and coadministration of compounds determined a further decrease ( $P < 0.01$  at 45 min and  $P < 0.001$  at 60 min; Fig. 1C). We then determined the type of interaction between the two compounds by performing an isobolographic analysis as described in *SI Materials and Methods*. We observed a dose-dependent hypophagic effect for each compound and for OEA-ABT-239 mixtures administered in fixed ratios (Fig. S1). The ED<sub>50</sub> and 95% confidence interval, obtained from regressions of dose-response curves (Fig. 1D and E), were (in milligrams per kilogram) ED<sub>50</sub>ABT-239 = 3.262 (2.883–3.641), ED<sub>50</sub>OEA = 5.283 (4.836–5.731), and ED<sub>50</sub>mix = 4.389 (3.727–5.050). In the isoblogram, the dose of ABT-239 needed to reach 50% of the effect is plotted on the abscissa and the isoeffective dose of OEA on the ordinate (Fig. 1F). The straight line connecting these two points represents the theoretical additive effect. A Student *t* test revealed no significant differences between the experimental ED<sub>50</sub>mix and the theoretical ED<sub>50</sub>add = 4.275 (3.731–4.813). Furthermore, the interaction index  $\gamma = 1.03$  indicates an additive interaction between OEA and ABT-239 (17).



**Fig. 1.** Interactions between OEA and brain histamine on food consumption. (A) Time course of the effect of systemic administration of OEA (10 mg/kg, i.p.) or vehicle (VEH) on cumulative food intake in 12 h-fasted HDC-KO and WT mice during the lights-on period (0900 h). Each point represents mean  $\pm$  SEM of 11–8 mice. \*\*\* $P < 0.001$  vs. respective controls. #### $P < 0.001$ ; ## $P < 0.01$  vs. OEA-treated mice WT or HDC-KO mice by two-way ANOVA and Bonferroni's test. (B) Time course of the effect of  $\alpha$ -FMH administration (5  $\mu$ g; i.c.v.) or saline (SAL) on OEA-suppressed food intake in 12 h-fasted mice. Each point represents mean cumulative food consumption  $\pm$  SEM of seven to eight mice during the lights-off period (1900 h). \*\*\* $P < 0.001$ , \*\* $P < 0.01$ , \* $P < 0.05$  vs. respective controls; ## $P < 0.01$ , # $P < 0.05$  vs. OEA-treated WT or  $\alpha$ -FMH-injected mice; two-way ANOVA and Bonferroni's test. (C) Increased brain histamine boosts OEA-induced suppression of food intake. Time course of 10 mg/kg OEA and 3 mg/kg ABT-239 effects on cumulative food intake in 12 h-fasted CD1 mice during the lights-on period (0900 h). Each point represents mean  $\pm$  SEM of seven to eight mice. \*\*\* $P < 0.001$ , \*\* $P < 0.01$ , OEA and ABT-239 vs. respective controls; #### $P < 0.001$ ; ## $P < 0.01$ , ABT-239/OEA vs. saline/OEA. (D and E) Sigmoidal and log-derived dose-response curves for OEA, ABT-239, and the combined compounds. Each point represents mean cumulative food consumption  $\pm$ SEM of seven to nine mice expressed as percent food consumption of vehicles. (F) Isoblogram for OEA-ABT-239 combined effects on food consumption.

**Palatable Wet Mesh Intake and Time Spent Eating in Histamine-Depleted Mice.** Palatable wet mesh (PWM) consumption was measured in  $\alpha$ -FMH i.c.v.-infused mice treated with OEA during lights off. This protocol allowed us to use the BSS paradigm to disclose the specificity of potential anorexiants agents on satiety development (18). One-way ANOVA revealed a significant treatment effect on PWM intake ( $F_{5,39} = 16.95$ ,  $P < 0.0001$ ; Fig. 24) during the 40-min BSS testing.  $\alpha$ -FMH had no effect on consumed PWM. Post hoc comparisons revealed a significant decrease of PWM intake in saline/OEA (5 mg/kg,  $P < 0.01$ ; 10 mg/kg,  $P < 0.001$ ) groups compared with controls.  $\alpha$ -FMH-infused mice treated with OEA did not significantly differ from



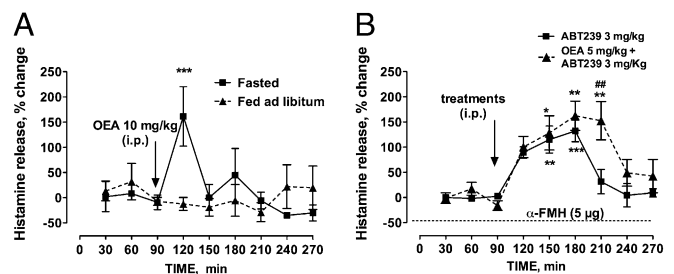
**Fig. 2.** Effects of OEA on the BSS in normal and histamine-depleted mice. *A* and *D* show mean cumulative PWM intake ( $\pm$ SEM) and mean time spent in eating activity ( $\pm$ SEM) over the 40-min BSS, respectively ( $n = 8$  for each experimental group). *A* and *D* also illustrate the temporal development of eating, locomotion, grooming, and resting duration (seconds) in saline/vehicle (*B*),  $\alpha$ -FMH/vehicle (*C*), saline/10 mg/kg OEA (*E*), and  $\alpha$ -FMH/10 mg/kg OEA (*F*). Represented on the x axes are eight time bins (T) of 5 min each for a total of 40 min of BSS analysis. \*\*\* $P < 0.001$ , \*\* $P < 0.01$ , \* $P < 0.05$  vs. respective controls; ### $P < 0.001$  vs.  $\alpha$ -FMH/vehicle treated mice by one-way ANOVA and Bonferroni's test. Experiments were carried out during the lights-off period (1900 h).

controls in the amount of PWM consumed (Fig. 2*A*). Moreover, one-way ANOVA disclosed a significant treatment effect on time spent eating over the 40-min BSS test ( $F_{5,39} = 8.60$ ,  $P < 0.0001$ ; Fig. 2*D*). Post hoc analysis showed a significant decrease in total time spent eating only in saline/OEA (5 and 10 mg/kg) groups, compared with saline/vehicle and  $\alpha$ -FMH/vehicle groups (Fig. 2*D*). No significant effects of OEA were detected in histamine-depleted mice compared with controls.

**Effect of OEA on BSS Development in Histamine-Depleted Mice.** We tested the effect of histamine depletion on the BSS in mice infused i.c.v. with  $\alpha$ -FMH. Two-way ANOVA showed the significant main effect of time for all of the behavioral categories examined (eating,  $F_{7,273} = 8.06$ ,  $P < 0.0001$ ; locomotion,  $F_{7,273} = 13.72$ ,  $P < 0.0001$ ; resting,  $F_{7,273} = 18.15$ ,  $P < 0.0001$ ; and grooming  $F_{7,273} = 4.13$ ,  $P < 0.001$ ). The time-dependent changes in the evolution of these behavioral patterns reflect the gradual decline of active behaviors (i.e., eating, grooming, and locomotion) and the parallel increase in resting that typically describe the expression of the BSS (Fig. 2*B, C, E, and F* and Fig. S2*A and B*). One-way ANOVA for each behavior revealed a significant treatment effect only for the duration of eating during 40-min BSS testing (Fig. 2). Post hoc comparisons performed on the duration of eating showed that only saline/OEA groups (Fig. 2*E* and Fig. S2*A*) significantly reduced such activity compared with saline/vehicle and  $\alpha$ -FMH/vehicle groups (Fig. 2*B and C*). No significant effects of treatment on time spent resting [ $F_{5,39} = 0.79$ , not significant (n.s.)], motor ( $F_{5,39} = 0.59$ , n.s.), or grooming activities ( $F_{5,39} = 1.81$ , n.s.) were found among all experimental groups. As shown in BSS temporal patterns, both control groups displayed an initial peak of feeding response followed by its decline over time and subsequent transition to resting that gradually increased around time bin T3 (Fig. 2*B and C*). This temporal pattern was different in mice with intact brain histamine treated with OEA (Fig. 2*E* and Fig. S2*A*). OEA administration produced a shifting of about 10-min of satiety and resting cooccurrence, thus emphasizing the premature onset of satiety in these animals with resting rapidly replacing eating activity (from a T3 to T1 interval) in a basically well-preserved BSS structure. In contrast, infusion of  $\alpha$ -FMH prevented the effects of OEA on satiety onset, as shown in Fig. 2*F* and Fig. S2*B*.

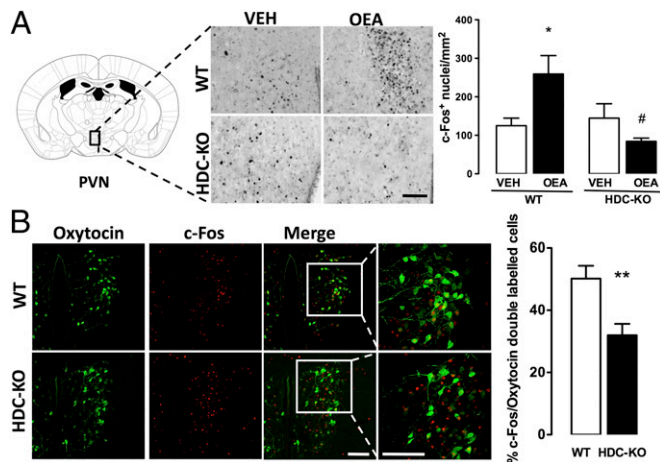
**Functional Interaction Between OEA and Brain Histamine.** In line with the idea that OEA may exert its anorectic effects by modulating brain histamine signaling, we found that systemic administration of 10 mg/kg OEA increased histamine release from the prefrontal cortex by  $\sim 160\%$ . Furthermore, OEA increased histamine release

only in 12 h-fasted but not in satiated mice (Fig. 3*A*). The increase in histamine was transient and reached a maximum within 30 min of OEA administration. Mean spontaneous histamine release was not significantly different between experimental groups ( $53.7 \pm 18.8$  fmol per 30 min-fasted mice;  $61.9 \pm 33$  fmol per 30 min-fed ad libitum mice). We then measured histamine release from the prefrontal cortex of freely moving mice that received i.p. administration of 3 mg/kg ABT-239. As shown in Fig. 3*B*, ABT-239 increased significantly histamine release by  $\sim 130\%$ . Histamine mean spontaneous release was  $49 \pm 12$  fmol/30 min. Co-administration of OEA and ABT-239 further increased cortical histamine release (Fig. 3*B*). Two-way ANOVA followed by Bonferroni's post hoc test showed a significant main effect between groups [ $F_{(treatment)1,61} = 6.12$ ,  $P < 0.01$ ;  $F_{(time)8,61} = 11.6$ ,  $P < 0.0001$ ; and  $F_{(treatment \times time)8,61} = 1.37$ , n.s.]. Mean spontaneous histamine release was  $34.3 \pm 14.1$  fmol/30 min. To ascertain whether our  $\alpha$ -FMH protocol caused a prolonged depletion of releasable histamine stores from the mouse brain, we measured histamine release from the mouse cortex 24 h after  $\alpha$ -FMH i.c.v. infusion. Cortical basal histamine contents of  $\alpha$ -FMH-treated mice were below detection level and no further histamine release was elicited when mice received ABT-239 (Fig. 3*B*).



**Fig. 3.** Functional interaction between brain histamine and OEA. (*A*) Effect of OEA administration on histamine release in the prefrontal cortex of freely moving mice. Represented are means  $\pm$  SEM of three fasted and four fed ad libitum mice. (*B*) Effect of ABT-239 and OEA-ABT-239 on cortical histamine release in freely moving mice. Represented are means  $\pm$  SEM of four and five mice, respectively. ABT-239 and OEA were administered i.p. separately and  $\alpha$ -FMH (5  $\mu$ g) was administered i.c.v. Baseline histamine release was not detectable in  $\alpha$ -FMH-treated mice or during ABT-239 administration (dashed line;  $n = 4$ ). Experiments were carried out between 0900 and 1600 h. \*\*\* $P < 0.001$ , \*\* $P < 0.01$ , \* $P < 0.05$ , vs. baseline values; ### $P < 0.01$  vs. ABT-239, two-way ANOVA and Bonferroni's test. For clarity, we reported only the significant differences vs. the last baseline sample before drug treatment, although differences were significant vs. all baseline samples.





**Fig. 4.** OEA-induced c-Fos expression in the hypothalamic PVN nucleus is blunted in HDC-KO mice. (A) Brain coronal sections showing the effect of vehicle or 10 mg/kg OEA on c-Fos protein expression in WT and HDC-KO mice in PVN. (Scale bar: 100  $\mu$ m.) Quantitative data are expressed in the bar graphs as means  $\pm$  SEM; \* $P$  < 0.05 vs. respective controls; # $P$  < 0.05 vs. WT/OEA by one-way ANOVA and Newman-Keuls test. (B) Immunohistochemical detection of oxytocin and c-Fos in the PVN neurons of WT and HDC-KO mice treated with OEA. (Scale bar: 50  $\mu$ m.) The bar graph shows the percentage of oxytocin-immunopositive neurons expressing c-Fos in OEA-treated WT and HDC-KO mice; \*\* $P$  < 0.01, unpaired  $t$  test.  $n$  = 3–5 mice per experimental group.

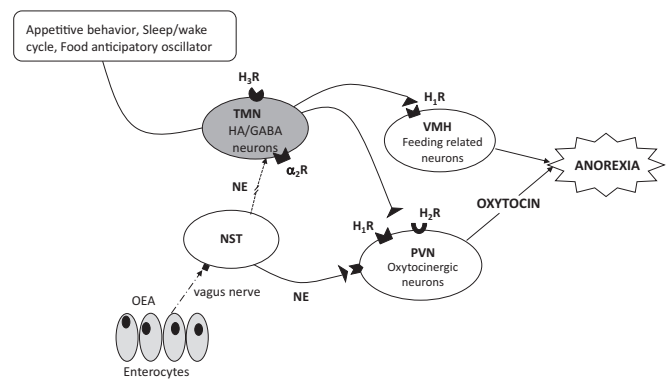
**OEA-Induced c-Fos Expression in HDC-KO Mice.** In an attempt to clarify how brain histamine deficit may prevent the satiety effect of OEA, we measured c-Fos protein expression in the brain of HDC-KO and WT mice following OEA administration. OEA did not change c-Fos expression in the VMH ( $F_{3,18} = 0.081$ , n.s.) of either WT or HDC-KO mice (Fig. S3C). As expected from previous results obtained in rats (2), systemic OEA increased significantly c-Fos expression in the PVN of WT mice (Fig. 4A;  $F_{3,18} = 4.817$ ,  $P$  < 0.05), whereas no differences were found in the PVN of HDC-KO mice treated with vehicle or OEA. OEA activates preferentially oxytocin neurons in the PVN (2). We found that the percentage of oxytocin neurons expressing c-Fos immunofluorescence was significantly lower in the PVN of OEA-treated HDC-KO mice compared with OEA-treated WT mice (Fig. 4B;  $P$  < 0.01), despite HDC-KO mice expressing higher levels of oxytocin-positive neurons (WT =  $166 \pm 17$  per square millimeter vs. HDC-KO =  $235 \pm 14$  per square millimeter;  $P$  < 0.01, unpaired  $t$  test). We also evaluated c-Fos expression in limbic brain structures related to feeding behavior that receive histaminergic innervation. Interestingly, in the nucleus accumbens that receives TMN input regulating exploratory behavior (19) indispensable during food provisioning, OEA administration decreased c-Fos expression in both HDC-KO and WT mice ( $F_{3,16} = 5.550$ ,  $P$  < 0.05; Fig. S3B). In the infralimbic cortex, which presumably implements arousal during appetitive behavior (20) and provides the TMN with an essential input for the appetitive function of histaminergic neurons (21), no differences were found in c-Fos expression in either WT or HDC-KO mice receiving OEA (Fig. S3A). We also observed a significant increase in the percentage of c-Fos positive TMN neurons in OEA- compared with vehicle-treated WT mice (OEA =  $12.1 \pm 1.35\%$ ; vehicle =  $5.4 \pm 1.4\%$ ;  $P$  < 0.01, unpaired  $t$  test; Fig. S4).

## Discussion

Satiety-related signals are integrated at the cellular and system level to give reliable and appropriate behavioral responses by recruiting specific neuronal populations. OEA is synthesized in several peripheral tissues and in the CNS (22) and in mammals it has been described as a mediator of numerous metabolic

processes (4). OEA secreted by enterocytes serves as a fast-sensing molecule that signals to satiety centers in the brain by engaging vagus nerve sensory fibers (2, 4, 23) and suppresses feeding by indirectly activating central oxytocin transmission in the PVN and SON (3). However, it is not known if other neurotransmitter systems integrate the peripheral signaling of OEA with effector hypothalamic nuclei. Brain histamine affects feeding behavior in a complex and not fully understood fashion (Fig. 5). It is fundamental for appetitive and aversive responses during motivated behavior (24), and blockade of histamine  $H_1$ R in the hypothalamus is believed to be responsible for the weight gain and metabolic dysregulation associated with the clinical use of atypical antipsychotics (25). Several peptides and hormones—such as leptin, corticotropin-, TSH-releasing hormones, and nefastin-1—are satiety modulators acting, at least in part, through histamine neurons activity (26–28). So far, there has been no information regarding brain histamine taking part in the anorexic effects of a modulator that, at the dose used in this study, does not easily pass the blood–brain barrier. Here, by using different experimental settings, we invariably show that chronic or acute histamine deficiency significantly attenuates OEA-induced hypophagia. OEA's effects in normal mice were not completely reversed in HDC-KO or  $\alpha$ -FMH-treated mice; indeed, it has been recently proposed that an increase in peripheral fatty acid oxidation and ketogenesis (29), as well as direct activation of the area postrema (30), may contribute to OEA's satiety effects. Nonetheless, our use of different experimental settings gave comparable results, supporting our view that OEA requires the integrity of the histamine system to fully exert its hypophagic effect.

By monitoring the natural progression of feeding behavior and its transition from eating to resting (i.e., satiety occurrence), the BSS can provide a detailed analysis of the structure of feeding behavior (18, 31). With respect to controls and  $\alpha$ -FMH-injected animals, OEA administration produced a temporal shifting toward earlier intervals of the cooccurrence of eating and resting behaviors. As illustrated by the preservation of the mutual relationship between feeding and nonfeeding behaviors, OEA induced a robust acceleration of satiety development without disrupting the basic structure of the feeding cycle, whereas histamine depletion abolished OEA-induced premature onset of satiety. Also, the BSS allows the continuous and complete



**Fig. 5.** Schematic drawing illustrating the putative interactions between OEA and the central histaminergic system. Histamine neurons are localized exclusively in the TMN of the posterior hypothalamus. Putative driver and modulatory inputs to the TMN are designated according to (24). The broken line designates presumed noradrenergic excitatory input from the NST to the TMN. According to our hypothesis, OEA induces anorexia indirectly stimulating histamine neurons in the TMN that project to the PVN. Activation of  $H_1$  and  $H_2$  receptors on feeding-related neurons in the PVN stimulates oxytocin release. Histamine mediates suppression of food intake also independently of OEA activating neurons in the VMH (47).

examination of food intake and provides the most accurate and detailed profile of feeding behavior, which are largely superior to interval sampling techniques (18). To understand which brain regions require histamine signaling for OEA-induced hypophagia, we used c-Fos expression as marker of functional activity. We found that lack of central histamine dampens OEA-induced increase of c-Fos expression in oxytocin PVN neurons of HDC-KO mice. The PVN integrates central and peripheral satiety signals, and H<sub>1</sub>R activation within the PVN has been implicated in the neuronal regulation of appetite (Fig. 5), as reported by decreased food intake following brain infusion of an H<sub>1</sub>R agonist and increased c-Fos-like immunoreactivity within the PVN (10). As pharmacological blockade of oxytocin receptors in the brain prevents OEA anorexic effects (2), we believe that histamine signaling on oxytocin PVN neurons is necessary for OEA to fully exert its anorexic effect (Fig. 5). In this regard, intranuclear and systemic release of oxytocin in response to suckling is controlled by H<sub>1</sub>R and H<sub>2</sub>R within the PVN (32). It was recently shown that noradrenergic NST–PVN projections are involved in the activation of the hypothalamic oxytocin system (33). In the TMN,  $\alpha$ 2A adrenoreceptors inhibit GABAergic transmission to TMN neurons (34). It is conceivable that NST adrenergic fibers projecting to the TMN disinhibit TMN neurons that in turn facilitate oxytocin release from the PVN to mediate OEA's prosatiety effect. Indeed, OEA increased c-Fos expression in a subgroup of TMN neurons, corresponding to the E2/E3 region in the mouse (35), that presumably are organized in a functionally distinct circuit, impinging on selected brain regions (36). c-Fos/HDC double-immunoreactive neurons in food-deprived mice were few, which is consistent with results obtained in rats under scheduled feeding (37). Interestingly, we also found that OEA decreased neuronal activity within the nucleus accumbens, as suggested by the low expression of c-Fos in both HDC-KO and WT mice. Although in this context we did not investigate this aspect further, it is interesting that OEA restores gut-stimulated dopamine release in the striatum of high-fat-fed rats increasing the reward value of unpalatable, yet healthier food (38).

As brain histamine signaling in the PVN seems to be involved in the acute effects of OEA on food consumption, we expected OEA to increase brain histamine release. Although technical limitations do not allow performing *in vivo* microdialysis experiments in the mouse PVN, we found in support of our hypothesis that OEA increased cortical histamine release. We further tested our hypothesis of brain histamine and OEA signaling functional interactions by measuring food consumption after coadministration of OEA and ABT-239, an H<sub>3</sub>R antagonist that blocks both auto- and heteroreceptors and increases transiently central release of histamine (12). Previous reports indicate that H<sub>3</sub>R antagonists decrease food intake in several mammalian species (reviewed in ref. 39), and here we show for the first time, to our knowledge, such an effect for ABT-239 as well. Furthermore, ABT-239 increased cortical histamine, and in agreement with feeding behavior, we observed a further increase in histamine release following ABT-239 and OEA coadministration. In our model we hypothesize that the hypophagic effects of OEA and ABT-239 converge onto a common pathway, as also strongly suggested by the isobolographic analysis of feeding behavior. Histaminergic neurons may also induce hypophagia by targeting other brain regions not affected by OEA, as is the case of leptin that recruits histamine neurotransmission in the VMH (40). Also, H<sub>3</sub>R antagonists and OEA regulate the release of several neurotransmitters other than histamine (41, 42), which may contribute to the hypophagic effects independently of each other.

Recently, Torrealba and coworkers, using a behavioral protocol that separates the appetitive from the consummatory phase of feeding, showed that histamine is differentially involved in these two aspects of feeding behavior and that histamine is

important for motivation to eat (20, 21). Brain histamine then seems to have different roles during food anticipatory responses and food consumption (43). Such a complex orchestration may be served by different histamine neuronal subpopulations that are recruited at different times during the unfolding of a specific behavior (36). Histamine neurons send broad projections within the CNS that are organized in functionally distinct circuits impinging on different brain regions (5). This implies independent functions of subsets of histamine neurons according to their terminal projections and their selective participation in different aspects of behavioral responses. It is interesting that OEA elicits histamine release from the cortex of hungry mice, but it is ineffective in satiated animals. To our knowledge, this is the first study to report that endogenous molecules can affect histamine release differentially depending on the homeostatic state of the animal. Whether histamine modulation in the cortex is relevant for satiety to curb the incentive to eat is not known and certainly deserves further study. Our study uncovers the role of brain histamine in the signal transduction of OEA-mediated anorexic effects. We believe that understanding the role of the histaminergic system in driving or modulating feeding behavior is of therapeutic relevance; for instance, atypical antipsychotics are thought to cause obesity by targeting histamine H<sub>1</sub>R (25), and the orexigenic liability of these drugs parallels their binding potency for histamine H<sub>1</sub>R (44, 45). Our results may contribute to the development of more effective pharmacotherapy for the management of obesity and ameliorate the safety profile of centrally acting drugs. In this regard, the first H<sub>3</sub>R antagonist for narcolepsy treatment was recently filed with the European Medicine Agency (46), and its effectiveness in obese patients may be assessed.

## Materials and Methods

**Animals.** Male CD1 mice (8–9 wk, 25–35 g; Harlan), HDC-KO mice and WT littermates (129/Sv background; see *SI Materials and Methods* for genotyping procedures and Fig. S5) grown in the animal facility of Dipartimento di Neuroscienze, Psicologia, Area del Farmaco, e Salute del Bambino, were housed in a temperature-controlled room (22 ± 1 °C) with a 12:12-h light–dark cycle (lights on from 0700 to 1900 h), at constant temperature and humidity with a standard diet (4RF21; Mucedola s.r.l.) and freely available water. HDC-KO and WT were used at 2–3 mo of age (25–35 g). Mice were handled for 1 wk before experiments. Housing, animal maintenance, and all experiments were conducted in accordance with the Council Directive of the European Community (86/609/EEC) of the Italian Decreto Legislativo 116 (1992) and National Institutes of Health guidelines on animal care and were approved and supervised by a veterinarian.

**Evaluation of Cumulative Food Consumption.** Mice were tested during the lights-on (0900 h) or lights-off (1900 h) cycle after 12 h food deprivation while water remained available. A weighed amount of standard chow pellets was placed in the food rack, and food consumption evaluated as the difference in weight between that of initially provided food and that left in the rack, including spillage in the cage. Mice were randomly assigned to the experimental groups. The protocol for cerebral histamine depletion with  $\alpha$ -FMH *i.c.v.* injection is described in detail in *SI Materials and Methods*. Food consumption was measured 15, 30, 45, and 60 min at 2, 4, 6, 8, 10, and 24 h after food presentation. OEA was injected *i.p.* 15 min before food presentation in 12 h-fasted mice.

**Isobolographic Analysis.** The interaction between OEA and ABT-239 was evaluated by coadministration of fixed ratios of each drug and by performing an isobolographic analysis as detailed in *SI Materials and Methods*.

**BSS Analysis and PWM Intake Assessment.** Before  $\alpha$ -FMH infusions, mice were habituated for 7 d to a PWM diet. PWM was prepared daily as a mixture of ground standard dry powdered food pellets in sweetened condensed milk (Nestlé) solubilized in distilled water (1:1.5) and offered for 1 h/d (light cycle). To minimize diet spillage, PWM was provided in special feeder beakers. The BSS paradigm was performed as described previously (31) and in *SI Materials and Methods*.

**Microdialysis Experiments.** The effects of OEA and ABT-239 on brain histamine release were evaluated in freely moving CD1 mice implanted with microdialysis probes. For details on surgery, experimental protocols, and HPLC-fluorimetric assay to quantify histamine, see *SI Materials and Methods*. Experiments are also described in detail in *SI Materials and Methods*.

**Immunohistochemistry.** HDC-KO and WT littermates were maintained on the standard chow diet and food deprived for 12 h (between 2000 h and 800 h; water remained available) before i.p. administration of OEA (10 mg/kg) or saline. Two hours after injections, mice were deeply anesthetized with chloral hydrate and perfused transcardially with cold physiological saline followed by 4% (vol/vol) paraformaldehyde in 0.1 M phosphate buffer, pH 7.4. Brains were processed for standard immunostaining as detailed in *SI Materials and Methods*.

**Data and statistical analysis.** Statistical analysis was performed using Prism Software (GraphPad). Statistical significance of cumulative food consumption and histamine release in microdialysis experiments were analyzed by two-way ANOVA (time  $\times$  treatment) with Bonferroni's post hoc test. Differences in c-Fos expression were determined by one-way ANOVA with

Newman-Keuls post hoc test. An unpaired *t* test was used to determine statistical significance between theoretical and experimental ED<sub>50</sub>, as well as differences between WT and HDC-KO mice in single- and double-labeled PVN and TMN neurons. OEA and  $\alpha$ -FMH effects on the duration of each BSS-related behavior were assessed by two-way repeated-measures ANOVAs with treatment (six levels) as the between-subject variable and time (eight levels consisting in eight time bins of 5-min each, from T1 to T8) as the within-subject variable, with Bonferroni's post hoc test. The criterion value for all statistical tests was  $P \leq 0.05$ .

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