Modulation of the Plasma Ghrelin Level by Intraventromedial Hypothalamic Microinjection of D2 Receptors Agonist and Antagonist

Maedeh Ghasemi¹, Nasrin Mehranfard², Afsaneh Eliassi³, Hojjat Allah Alaei¹,†

Abstract
Dopamine is well recognized to regulate feeding via hypothalamic circuits in the brain. However, the mechanism by which dopamine controls food intake through hypothalamic pathways remains largely unknown. Recent findings propose that metabolic circuit and especially peripheral orexigenic/anorexigenic signals may play a critical role in the regulation of food intake. Therefore, in the present study, we hypothesized that ghrelin might modulate the orexigenic effect of ventromedial hypothalamic (VMH) dopamine receptors. To address this issue, we assessed the effects of intraventromedial administration of dopamine D2 receptors agonist, Quinpirole, and antagonist, Sulpiride, on plasma levels of ghrelin in 20-h fasted rats. Results showed that the pharmacological activation of D2 receptors increases the plasma ghrelin level persisted over 1h. This stimulatory effect of dopamine receptor activation on ghrelin was reduced by D2 receptors blockade in the VMH. In some experiments, atropine (cholinergic antagonist; 1 mg/kg s.c.) was administered 30 min before drug or saline injections which resulted in a marked suppression of fasted and stimulated ghrelin secretion. Together, our novel data implicates ghrelin in the orexigenic action of VMH D2 receptors signaling with a possible functional role of vagal pathway in the control of dopamine-evoked appetite (ghrelin) induction.

Keywords: Dopamine, D2 receptor, Ghrelin, Ventromedial nucleus, hypothalamus, Atropine

Introduction
Food intake is particularly controlled by the hypothalamus and involves homeostatic and hedonic pathways [1,2]. Impairment in nutrient sensing pathways in the hypothalamus has been found to induce obesity [3]. Several neuroendocrine hormones especially ghrelin regulate food intake through an effect on the hypothalamus [4]. Ghrelin is the primary gut orexigenic signal that is involved in short and long-term regulation of appetite principally via acting on the hypothalamus [5,6] and can enhance food intake in both animals and humans [7,8]. There is some evidence that showing dopamine (DA) neurotransmission also can exert a regulatory effect on food intake [9,10]. For example, DA-deficient mice exhibited behaviors associated with seeking and ingesting food, but they could not consume food enough to survive [11]. Additionally, the DA antagonists caused serious feeding deficits in animals [12,13]. The dopamine D2 receptors have been shown to
predominantly mediate the regulatory effect of the DA on food intake [14-16] within several specific sites of the hypothalamus including ventromedial nucleus [17,18]. However, whether modifying effect of hypothalamic dopamine receptor activity on energy homeostasis and appetite behaviours is mediated by regulation of ghrelin concentration is unknown. Interestingly, previous studies demonstrated that activation of the D2 receptor significantly reduces both basal and stimulated plasma leptin levels in human and rats, suggesting D2 receptor neurotransmission mediates energy balance via regulating leptin levels [19,20]. Although the precise mechanism(s) of hypothalamic dopaminergic regulation of energy homeostasis by peripheral homeostatic modulators is (are) unknown, it seems that normal functioning of dopamine receptors in the brain play an important role in the maintenance of normal eating behavior and in the regulation of energy balance through physiological state associated hormones (i.e., hunger and satiety hormones). Since, there is no adequate data in literature about a relationship between D2 receptor activation and ghrelin level, therefore in this study; we investigated the existence of a possible correlation between plasma levels of ghrelin and hypothalamic D2 receptors in ventromedial nucleus of the hypothalamus in rats. Together, our results showed that there is a potential interaction between the dopaminergic transmission in hypothalamus and appetite-regulating hormones that likely depend on vagal cholinergic pathway.

Materials and Methods

■ Animals

Male Wistar rats (220–250 g, Pasteur Institute of Iran, Tehran) were housed under controlled conditions (23 ± 2 °C, 30- 40% humidity and light on 07:00–19:00) with free access to standard rat chow and tap water in. They were deprived of food, but not water, for 20–24 h prior to experiments. All experimental protocols were approved by the Ethical Committee of Isfahan University of Medical Sciences (Isfahan, Iran) in accordance with the Guide for Care and Use of Laboratory Animals (National Institute of Health Publication No.80-23, revised 1996).

■ Drugs

Quinpirole hydrochloride (dopamine D2 receptors agonist), Sulpiride (dopamine D2 receptors antagonist) and atropine sulfate monohydrate (cholinergic antagonist) were purchased from Sigma (St Louis, MO, USA).

■ Stereotoxic surgery

After 1 week of habituation to the housing conditions, for cannulation and verification of placement, rats were deeply anesthetized with chlorate hydrate (400 mg/kg; i.p.) and fitted with a 23-gauge stainless steel cannula placed just above the right ventromedial hypothalamic (VMH).

Stereotoxic coordinates were determined from the rat brain atlas of Paxinos and Watson; VMH, lateral: +0.7 mm from midline; dorsoventral: -2.4 mm from the bregma [21]. The injector extended 1 mm beyond the end of the guide cannula. After surgery, the animals were allowed a 5-day recovery period before experimental trials. Placement of cannula in the VMH was confirmed after experiment by fixation of brains in formalin and then 100 μm thick sections were taken and cannula and injection tracks were examined with light microscopy.

Only data obtained in rats verifying histologically the placements of cannula were used (Figure 1). The total number of injected rats in the VMH was 36 (each group; n=6). Five rats were excluded due to cannula misplacement. Injection of compound drugs or saline was performed under chlorate hydrate anesthesia using a Hamilton 1 μl syringe through silastic tubing in a volume of 0.5 μl over 30 s. After the injection, the needle was left in place for an additional 30 s.

■ Treatments

In a first and second set of experiments, we considered the effects of VMH -microinjected Quinpirole (D2 receptor agonist) and Sulpiride (D2 receptor antagonist) on plasma concentration of ghrelin. In the first set of experiments, rats in control and treatment groups, received an effective dose of D2 receptor agonist, Quinpirole (0.5μg), and antagonist, Sulpiride (0.005 μg) or saline (0.5 μl) into the VMH, respectively, after a 20-h fasting period. The effective dose of D2 receptor agonist and antagonist was based on that used in other study in our lab (data not published). To investigate the time-course effect of the VMH injection of drugs; blood sampling was performed at 0, 10, 20, 40 and 60 min after drugs injection. Basal blood sampling (0 min) was immediately followed by the VMH injection and then at 10, 20, 40 and 60 min after the injection, blood
samples were collected. For protection of blood samples against ghrelin degradation, coated tubes with protease inhibitors were used.

Tail blood sampling technique was used to collect blood (at the amount of 500 μl) from the rats around 9:30 AM on the fasted day.

In the second set of experiments, we studied the role of the vagus nerve in the regulation of hypothalamic D2 receptors-mediated energy homeostasis. Rats were pre-treated with subcutaneous (s.c.) injection of atropine or saline, and 30 min later D2 receptor agonist, Quinpirole, and antagonist, Sulpiride, at doses of 0.5μg and 0.005 μg or saline (0.5 μl) was administered into VMH. Blood samples were collected at 0, 10, 20, 40 and 60 min after the drug injections for determination of hormone secretion output. After blood sampling, plasma ghrelin was measured using rat ghrelin EISA kit with intra-assay precision CV<10% (Zellbio Company ELISA kit-Germany). All experiments were performed at 9:30 AM.

**Statistical Analysis**

Results are shown as mean ± SEM. The differences between groups in the hormone study were checked by repeated measures analysis of variance (ANOVA) (time course analysis of ghrelin), one-way ANOVA followed by Turkey’s HSD test (hormone analysis at one time). Student’s paired t-test was used to evaluate differences between two points in one group. P<0.05 was considered statistically significant.

**Results**

In the present study, we evaluated whether modifying effect of hypothalamic D2 receptors on energy homeostasis can be mediated through the regulation of ghrelin secretion (probably mediated by vagal pathway). To do this, the effective doses of D2 receptor agonist and antagonist were selected and examined.

- **Time course effects of D2 receptor agonist (Quinpirole) and antagonist (Sulpiride) on the plasma concentration of ghrelin**

  The ghrelin secretion pattern over time (a period of 1 h) was evaluated following microinjection of Quinpirole (0.5 μg) into the VMH. As shown in Figure 2A, plasma ghrelin level was significantly changed in different groups

  \[ F (4, 24) = 9.273, P<0.00 \]. Further analysis revealed a significant increase (p<0.001) in plasma ghrelin from the basal levels of ~100% to 181% at 10 min and to 187% at 60 min, compared with saline injection (Figure 2B). Moreover, the application of Sulpiride significantly decreased (p<0.01) plasma ghrelin from the basal levels of ~100% to 65% at 40 min and to 51% at 60 min, compared with saline injection (basal level: 1191.61 ± 49.92; at 10 min: 1140.55 ± 97.50 and at 60 min: 1228.79 ± 73.38, Figure 2A). Sulpiride injection had no significant effect on ghrelin concentration at time of 10 (106%) and at 20 min (96%) (Figure 2B) compared with saline treated rats.

Atropine induced a vagal cholinergic dependent decrease in plasma ghrelin levels in drug- treated rats

We also examined the possibility that the vagal cholinergic pathway might be implicated in the stimulation of ghrelin secretion by VMH injection of Quinpirole.

As shown in Figure 3A, atropine injection in 20-h fasted rats, significantly changed plasma ghrelin levels in different groups \[ F (4, 24) = 4.53, P<0.00 \]. Further analysis revealed a significant decrease (p<0.00) in plasma ghrelin from the basal levels of 1204.01 ± 21 pg/dl to 357 ± 70.75 (Figure 3A) (31%; Figure 3B) at 10 min, and to 361.59 ± 11 pg/dl (Figure 3A) (32%; Figure 3B) at 60 min compared with control group. Increased ghrelin level by Quinpirole was prevented by atropine pretreatment (Fig. 3A).
Plasma ghrelin concentration fluctuates with feeding status (i.e., hunger vs. satiety). Existing literature demonstrates the ghrelin action on regulation of both normal and cue-driven feeding through the midbrain dopaminergic system. Ghrelin and hunger state augment dopamine release, thereby increasing food seeking behaviour [22-24]. Moreover, cue-associated dopamine release has been shown to trigger ghrelin release and increase its plasma level [6]. However, it is unknown whether the hypothalamic dopamine projections would physiologically affect homeostatic regulators of energy including ghrelin?

Normal function of dopaminergic neurotransmission has been shown to be essential for feeding and survival [25]. It is now clear that any abnormality involving the dopaminergic system will be reflected by changes in feeding behavior. Pervious pharmacological studies demonstrated that dopamine D2 agonists inhibit food intake by the action on hypothalamic neuropeptide Y. It has also been observed that D1 and D2 receptors synergize to inhibit feeding in healthy and diabetic rats and that a combination of D1/D2 receptor agonists are needed to alleviate obesity [26]. In contrast, adult Drd2-/- mice (D2R knockout mice), hypothalamic low D2 receptor expression and D2 receptor blockade by Sulpiride or antipsychotics were associated with a hyperphagic response that were found only in obese rats [15,27]. This supports the idea that abnormal dopaminergic transmission is involved in the pathogenesis of hyperphagia during obesity. Even though the hypophagic effects of dopamine receptor agonists have been characterized, it has been reported that Quinpirole (D2-like receptor agonist) administration produces a complex behavioral effect, decreasing palatable food consumption while it enhances the ingestion of standard food in rats [28]. Also Salimi, et al. observed that activation of D2 receptors of paraventricular nucleus (PVN) increases food intake and intra-
PVN microinjection of sulpiride decreases food intake by blocking D2 receptors in this area [29]. Taken together, increasing studies demonstrated that modulation of the hypothalamic dopamine receptor activity induced significant changes on food intake [17,9,10,12,28,30,31]. Nevertheless, the mechanism by which hypothalamic dopaminergic pathways control energy balance remains as an open question. It seems that metabolic circuit, especially variations in hyperphagic or hypophagic peripheral signals play a critical role in the regulation of eating behavior [32]. In this regard, in the present study, we investigated whether modifying effect of the hypothalamic D2 receptors on energy balance may be mediated through the regulation of ghrelin secretion. We provide in this work first evidence for an important role of the hypothalamic dopamine signaling in the regulation of ghrelin plasma levels via the connection in which activation of D2 receptors in the VMH increases ghrelin concentration. We also found that the stimulatory effect of dopamine D2 receptors on plasma ghrelin concentration is markedly blocked by the pretreatment with atropine.

The metabolic role of the dopaminergic system and its interaction with ghrelin has been reported in previous studies [24,33-35]. The importance of this interaction in the energy control center is highlighted by a co-localization of growth hormone secretagogue receptor-1A and D2 receptors in the hypothalamus [36]. Actually, a significant point of our study is the fact that dopamine has a role outside of the classical pathway in which it regulates ghrelin level possibly via a two-way link between dopamine signaling and ghrelin. Hypothalamic regions implicated in ghrelin signaling, lateral and ventromedial hypothalamic nuclei as well as arcuate nucleus, form neurocircuits with the reward pathways that is likely to be important in integrating feeding control [33]. Tracing studies show that neurons in the ventral tegmental area (VTA) receive projections from hypothalamus and infusion of ghrelin into the lateral hypothalamus or directly into regions related to pleasure, reward and motivation stimulates VTA neurons, causing release of dopamine [34]. Also, DA projections to hypothalamus are implicated in homeostatic regulation of food intake. These data establish a physiological role of hypothalamic dopamine signaling in stimulating ghrelin secretion. Nevertheless, future studies are needed to elucidate which hypothalamic neuronal populations and neurotransmitters are involved in ghrelin secretion.

The vagovagal circuitry within the dorsal vagal complex (DVC) via descending inputs from the lateral hypothalamus, PVN, VMH, and arcuate nuclei constitutes the primary means whereby higher centers exert a direct influence over gastrointestinal function [37-40]. Furthermore, the ventromedial hypothalamus elicits anorexigenic and orexigenic signals to paraventricular nucleus and lateral hypothalamic nucleus, and then lateral hypothalamic nucleus exerts a direct pathway to dorsal motor nucleus. These lead to efferent output via the dorsal motor nucleus of the vagus, which activates the vagus nerve to store energy [32,39].

In this context, we found the possibility that dopamine-containing projections to the DVC may also play a role in the central regulation of ghrelin release. This is based on the fact that administration of D2 receptor agonist induced a vagal cholinergic-dependent increase in plasma ghrelin levels and peripheral pretreatment of atropine markedly suppressed this effect. A relationship between vagus nerve activity and ghrelin release has also been reported in previous studies. For example, it has been demonstrated that the rise of ghrelin levels in fasted condition is mediated via the vagal nerve connections to the stomach mucosa [41]. Hosoda, et al. and Williams, et al. reported that vagal-induced elevation of plasma ghrelin in fasting completely is suppressed by subdiaphragmatic vagotomy or atropine pretreatment, and this result was mimicked by treatment with the atropine [42,43]. More directly, cholinergic agonists or antagonists increase or reduce circulating ghrelin levels in humans, respectively [44,45]. Additionally, vagal-cholinergic (muscarinic) activation can stimulate gastric ghrelin gene expression and ghrelin secretion [46]. Since in the present study, atropine was peripherally injected, acetylcholine receptors in other regions modulating appetite and perhaps in reward centers of brain might also be involved in the regulation of ghrelin secretion. For example, there are few data that reveal muscarinic acetylcholine receptors (M3 receptors) in the hypothalamus are mainly targets for obesity and food intake. The lack of M3 receptors have been reported to protect against some forms of obesity as well as to ameliorate impairments in glucose and energy homeostasis [47,48]. Furthermore, the blockade of muscarinic acetylcholine receptors in the
ventral tegmental area could disrupt food-related learning in rats [49].

Together, our results suggest that the hypothalamic dopamine signaling might follow a mechanism through which vagal outflow to the gastric system is increased and results in enhanced ghrelin secretion, although further studies are needed to assess the mechanism(s) involved in the hypothalamic dopamine regulation of food intake.

**Conclusion**

Collectively, these findings suggest that dopamine is a physiological regulator of ghrelin release and support the role of hypothalamic dopamine-vagal efferent activation in stimulating ghrelin secretion. It also seems that dopamine signaling, cholinergic pathway and ghrelin secretion make a straight pathway and any point blockade of the pathway may interrupt orexigenic action of the hypothalamic dopamine.

**Conflicts of interest**

The authors declare that no competing interests exist.

**Acknowledgments**

This work was supported by the Isfahan University of Medical Sciences, Isfahan, Iran [Grant No. 194071, 2015]. The funders had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript.

**References**

26. Rusk IN, Cooper SJ. Parametric studies of selective D1 or D2 antagonists: effects on appetitive and feeding behaviour. Behav. Pharmacol 5(6), 615-622 (1994).


