Cytoplasmic kinases downstream of GPR30 suppress gonadotropin-releasing hormone (GnRH)-induced luteinizing hormone secretion from bovine anterior pituitary cells

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Abstract. GPR30 is known as a membrane receptor for picomolar concentrations of estradiol. The GPR30-specific agonist G1 causes a rapid, non-genomic suppression of gonadotropin-releasing hormone (GnRH)-induced luteinizing hormone (LH) secretion from bovine anterior pituitary (AP) cells. A few studies have recently clarified that protein kinase A (PKA) and phosphorylated extracellular signal-regulated kinase (pERK) might be involved in cytoplasmic signaling pathways of GPR30 in other cells. Therefore, we tested the hypothesis that PKA and ERK kinase (MEK) are important cytoplasmic mediators for GPR30-associated non-genomic suppression of GnRH-induced LH secretion from bovine AP cells. Bovine AP cells (n = 8) were cultured for 3 days under steroid-free conditions. The AP cells were previously treated for 30 min with one of the following: 5000 nM of PKA inhibitor (H89), 1000 nM of MEK inhibitor (U0126), or a combination of H89 and U0126. Next, the AP cells were treated with 0.01 nM estradiol for 5 min before GnRH stimulation. Estradiol treatment without inhibitor pretreatment significantly suppressed GnRH-induced LH secretion (P < 0.01). In contrast, estradiol treatment after pretreatment with H89, U0126 or their combination had no suppressive effect on GnRH-induced LH secretion. The inhibitors also inhibited the G1 suppression of GnRH-induced LH secretion. Therefore, these data supported the hypothesis that PKA and MEK (thus, also pERK) are the intracellular mediators downstream of GPR30 that induce the non-genomic suppression of GnRH-induced LH secretion from bovine AP cells by estradiol or G1.

Key words: Extra-cellular regulated kinases, Gonadotrope, G protein-coupled estrogen receptor-1, Protein kinase A, Ruminant

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Materials and Methods

All experiments were performed according to the Guiding Principles for the Care and Use of Experimental Animals in the Field of Physiological Sciences (Physiological Society of Japan) and approved by the Committee on Animal Experiments of the School of Veterinary Medicine, Yamaguchi University.

Effects of H89 and U0126 on LH secretion

This experiment was conducted to verify the effect of H89, U0126 and their combination (in the absence of estradiol) on the GnRH-induced LH secretion from bovine AP cells. Nett et al. [19] reported that LH and GnRH receptor levels in the AP were higher during the luteal phase than during the immediate post-estrus period in heifers. Therefore, APs were obtained from postpubertal Japanese Black heifers in the middle of the luteal phase (n = 8, 26 months of age) from a local slaughterhouse in Yamaguchi Prefecture. The protocol to obtain the AP from the head and the method of transporting the APs to the laboratory were as reported previously [11]. The experiment was repeated 8 times with each of the 8 different pituitary glands, using 4 wells per treatment. Each experiment began with enzymatic dispersal of AP cells by using a method previously described [20, 21], and cell viability of greater than 90% was confirmed by trypan blue exclusion. The total cell yield was $19.9 \times 10^6 \pm 0.9 \times 10^6$ cells per pituitary gland. The dispersed cells were then suspended in phenol red-free Dulbecco’s Modified Eagle’s Medium (DMEM; 21063-029, Gibco, Grand Island, NY, USA) containing 1% nonessential amino acids (100 ×; Gibco), 100 IU/ml penicillin, 50 µg/ml streptomycin, 10% horse serum (Gibco), and 2.5% fetal bovine serum (FBS; Gibco). The horse serum and FBS had previously been treated with dextran-coated charcoal to remove steroid hormones. After the cells (2.5 × 10^5 cells/ml, total 0.5 ml) were plated in 24-well culture plates (MS-80240; Sumitomo Bakelite, Tokyo, Japan), they were maintained at 37°C in a humidified atmosphere of 5% CO2 for 82 h. The wells were washed twice with PBS and then incubated with 490 µl of DMEM containing 0.1% BSA for 2 h.

Cells were pretreated with DMEM (5 µl) alone or with DMEM containing 500 µM of a PKA inhibitor, H89 (Lkt Laboratories, St. Paul, MN, USA), and/or 100 µM of a MEK inhibitor, U0126 (Enzo Biochem, New York, USA). After 30 min of incubation, DMEM was added to each culture well. We added DMEM at the same time as adding estradiol or G1 in order to follow the same protocol as that used for estradiol and G1 in the following experiment. Cells were incubated with gentle shaking for 5 min, after which they were incubated for 2 h with 5 µl of 100 nM GnRH (Peptide Institute, Osaka, Japan) to stimulate LH secretion. Fig. 1 summarizes the final concentrations of H89, U0126 and GnRH for each treatment. As previously reported [22], LH secretion was stimulated by increasing the amounts of GnRH, with a peak at 1 nM GnRH, and reducing secretion at GnRH concentrations higher than 1 nM. Therefore, the final concentration of GnRH used in this study was 1 nM in all treatments, except in the “controls,” which were pretreated with 5 µl of DMEM for 30 min, treated with 5 µl of DMEM for 5 min and then treated with 5 µl of DMEM without GnRH for 2 h. The 30-min treatment time and the concentrations of H89 and U0126 used in the present study are commonly used in such studies [23] and are identical to those used in an investigation of signaling pathways in lactotroph cells [24]. After 2 h of incubation, the medium was collected for immunoassay of LH.

Fig. 1. Comparison of the effects H89 and U0126 on GnRH-induced LH secretion from cultured bovine AP cells. The final concentrations of H89, U0126 and GnRH were 5000 nM, 1000 nM and 1 nM, respectively. LH concentrations in control cells (cultured in medium alone) were averaged, and the mean LH concentrations of treated groups were expressed as percentages of the average control value. a vs. b: significant difference (P < 0.05)

Effects of H89 and U0126 on the suppression of GnRH-induced LH secretion by estradiol and G1, respectively

Three separate experiments were conducted to evaluate the effect of H89, U0126 and their combination on the suppression of GnRH-induced LH secretion from bovine AP cells by estradiol and G1, respectively. Anterior pituitaries were obtained from postpubertal Japanese Black heifers in the middle of the luteal phase (n = 8, 26 months of age).

Each experiment was repeated 8 times with each of the 8 different pituitary glands, using 4 wells per treatment. After enzymatic dispersal of AP cells, the cells were cultured in the medium described in the previous section for 82 h. The wells were washed twice with PBS and then incubated with 485 µl of DMEM containing 0.1% BSA for 2 h. Cells were pretreated with 5 µl of DMEM alone or with 5 µl of DMEM containing 500 µM of H89 (final concentration, 5000 nM) and/or 100 µM of U0126 (final concentration, 1000 nM). For the estradiol experiment, after 30 min of incubation, either 5 µl of DMEM alone or 5 µl of DMEM containing 1 nM estradiol (final concentration, 0.01 nM; Wako Pure Chemical Industries, Osaka, Japan) was added to each culture well. For the G1 experiment, G1 (final concentration, 0.01 nM; Azano Biotech, Albuquerque, NM, USA) was used instead of estradiol.

The cells were incubated with gentle shaking for 5 min, after which they were incubated for 2 h with 5 µl of 100 nM GnRH (Peptide Institute, Osaka, Japan) to stimulate LH secretion. The final concentrations of H89 and U0126 were 5000 nM and 1000 nM respectively, and the combination (in the absence of estradiol) on the GnRH-induced LH secretion from bovine AP cells.

The concentrations of GnRH used in this study were 1 nM in all treatments, except in the “controls,” which were pretreated with 5 µl of DMEM for 30 min, treated with 5 µl of DMEM for 5 min and then treated with 5 µl of DMEM without GnRH for 2 h. The 30-min treatment time and the concentrations of H89 and U0126 used in the present study are commonly used in such studies [23] and are identical to those used in an investigation of signaling pathways in lactotroph cells [24]. After 2 h of incubation, the medium was collected for immunoassay of LH.

Effects of H89 and U0126 on the suppression of GnRH-induced LH secretion by estradiol and G1, respectively

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Each experiment was repeated 8 times with each of the 8 different pituitary glands, using 4 wells per treatment. After enzymatic dispersal of AP cells, the cells were cultured in the medium described in the previous section for 82 h. The wells were washed twice with PBS and then incubated with 485 µl of DMEM containing 0.1% BSA for 2 h. Cells were pretreated with 5 µl of DMEM alone or with 5 µl of DMEM containing 500 µM of H89 (final concentration, 5000 nM) and/or 100 µM of U0126 (final concentration, 1000 nM). For the estradiol experiment, after 30 min of incubation, either 5 µl of DMEM alone or 5 µl of DMEM containing 1 nM estradiol (final concentration, 0.01 nM; Wako Pure Chemical Industries, Osaka, Japan) was added to each culture well. For the G1 experiment, G1 (final concentration, 0.01 nM; Azano Biotech, Albuquerque, NM, USA) was used instead of estradiol.

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The concentrations of GnRH used in this study were 1 nM in all treatments, except in the “controls,” which were pretreated with 5 µl of DMEM for 30 min, treated with 5 µl of DMEM for 5 min and then treated with 5 µl of DMEM without GnRH for 2 h. The 30-min treatment time and the concentrations of H89 and U0126 used in the present study are commonly used in such studies [23] and are identical to those used in an investigation of signaling pathways in lactotroph cells [24]. After 2 h of incubation, the medium was collected for immunoassay of LH.
of the 2 agents used can significantly suppress GnRH-induced LH secretion from bovine AP cells. In all 2 experiments, after 2 h of incubation, the medium was collected for LH immunoassay.

Radioimmunoassay to measure LH concentration in culture media

LH concentrations in the culture media were assayed in duplicate by double-antibody radioimmunoassay using \(^{125}\)I-labeled bLH and anti-oLH-antiserum [AFP11743B and AFP192279, National Hormone and Pituitary Program of the National Institute of Diabetes and Digestive and Kidney Diseases (NIDDK), Bethesda, MD, USA]. This assay has been previously described in detail [12] and has been used to measure changes in blood LH concentration in Holstein heifers before and after GnRH treatment [25]. The detection limit was 0.40 ng/ml. At 2.04 ng/ml, the intra- and inter-assay coefficients of variation were 3.6% and 6.2%, respectively.

Statistical analysis

LH concentrations in the control samples for each pituitary were averaged, and the mean value was set as 100%. LH concentrations in the treated samples for each pituitary were averaged, and the mean LH values were expressed as a percentage of the control value. Data were analyzed using StatView version 5.0 for Windows (SAS Institute, Cary, NC, USA). The statistical significance of differences in LH concentration were analyzed by one-factor analysis of variance (ANOVA) followed by post hoc comparisons using Fisher’s protected least significant difference test. The level of significance was set at \(P < 0.05\). Data are expressed as the mean ± standard error of the mean (SEM).

Results

Effects of H89 and U0126 on GnRH-induced LH secretion in AP cells

Figure 1 depicts the effects of H89 alone, U0126 alone and the combination of both inhibitors on GnRH-induced LH secretion from cultured AP cells in the absence of estradiol. The LH concentration in the medium of GnRH wells was higher than in the control wells. None of the treatments had an effect on GnRH-induced LH secretion.

Effects of H89 and U0126 on estradiol-mediated suppression of GnRH-induced LH secretion in AP cells

Figure 2 depicts the effects of H89 and U0126 on estradiol-mediated suppression of GnRH-induced LH secretion from cultured AP cells. Treatment with 0.01 nM estradiol in the absence of inhibitors suppressed GnRH-induced LH secretion. In contrast, pretreatment with H89 alone, U0126 alone or their combination inhibited estradiol suppression of GnRH-induced LH secretion. The effects among treatments with H89 alone, U0126 alone or their combination did not differ significantly.

Effects of H89 and U0126 on G1-mediated suppression of GnRH-induced LH secretion in AP cells

Figure 3 depicts the effects of H89 and U0126 on the G1-mediated suppression of GnRH-induced LH secretion from cultured AP cells. Treatment with 0.01 nM G1 in the absence of inhibitors suppressed GnRH-induced LH secretion. In contrast, pretreatments with H89 alone, U0126 alone or the combination of H89 and U0126 inhibited the G1-mediated suppression of GnRH-induced LH secretion. The
effect of U0126 alone was weaker (P < 0.01) than that of H89 alone and the combination treatment.

Discussion

In the present study, pretreatment with PKA or MEK inhibitors inhibited the estradiol- or G1-induced suppression of GnRH-induced LH secretion. The inhibitors had no significant effect on nonactivated GPR30. Therefore, we addressed each inhibitor separately and the possible association between PKA and MEK signaling as the cytoplasmic pathways for GPR30 activated by estradiol and G1.

A few studies have recently indicated that PKA is a potential intracellular downstream mediator of the GPR30 pathway in non-gonadotroph cells, mouse trigeminal ganglia [13] and the rat liver [14], although the role of PKA is not linked to controlling LH secretion. G1, as well as estradiol, increases the current amplitude of voltage-gated Na+ channels in human breast cancer cells, and a PKA inhibitor can abolish such an effect [26]. Therefore, the present data suggested that PKA might be an intracellular mediator downstream of GPR30 that induces estradiol-mediated suppression of LH secretion from AP cells in a non-genomic manner. Further studies are required to clarify the contribution of Na+ channels to estradiol’s non-genomic suppression of LH secretion.

Estradiol increases pERK in the ovine AP both in vivo and in vitro within 15 min of treatment [7, 15]. The GPR30 antagonist G36 inhibits ERK phosphorylation by estrogen in SKBr3 cells [16]. Estrogen activates ERK even in ER-negative SKBr3 cells [27]. Therefore, the present findings suggest that MEK and pERK could be other intracellular mediators downstream of GPR30 that induce estradiol-mediated suppression of LH secretion from AP cells via a non-genomic mechanism.

Considering the inhibitory effect of estradiol mediated by GPR30 on pain perception, PKA might be an upstream mediator of MEK [28]. In this study, there was no synergistic effect between U0126 and H89 on the estradiol-mediated suppression of GnRH-induced LH secretion. Therefore, PKA could be a downstream mediator of GPR30 and an upstream mediator of MEK. However, the effect of pretreatment with U0126 alone was weaker against G1-mediated suppression than pretreatment with H89 alone or with the combination of H89 and U0126. Therefore, there could be another mediator downstream of PKA that mediates GPR30 activation. Terasawa and Kenealy [29] reported that estradiol affects various pathways in GnRH neurons and induces cross-talk between cell surface receptors, GPR30 and the nuclear receptors ERα and ERβ. Treatment with ERβ-specific ligand (diarylpropionitrile) as well as estradiol and G1 was reported to rapidly increase pERK in inflammatory breast cancer cell lines [30], suggesting that a combination of ERβ and GPR30 is involved in promoting invasion through the activation of MEK in the non-genomic signaling pathway. When viewed in the context of this recent study, data from the present study suggest that GPR30 plays an important role in the suppression of LH secretion but that ERα and ERβ could also be involved in inducing the rapid suppression of LH secretion.

LH is secreted from the AP into circulating blood in a pulsatile manner during most of the estrous cycle [31, 32]. The pulsatile secretion of GnRH from the hypothalamus into hypophyseal portal blood is the most important factor that controls the parameters of pulsatile LH secretion, particularly the LH pulse frequency [33]. However, changes in the PKA and MEK pathways in AP cells might contribute to other parameters of pulsatile LH secretion, namely, the LH amplitude and mean LH concentration [34, 35].

In conclusion, our study supported the hypothesis that PKA and MEK are intracellular mediators downstream of GPR30 that induce non-genomic suppression of GnRH-induced LH secretion from bovine AP cells by estradiol and G1.

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