CONCANAVALIN A INHIBITS THE BASAL AND STIMULATED HORMONE RELEASE FROM PITUITARY CELLS IN MONOLAYER CULTURES

Gabriel PONSIN+, Ashok KHAIR, Jolanta KUNERT-RADEK, Thérèse BENNARDO and Marian JUTISZ
Laboratoire des Hormones Polypeptidiques, CNRS, 91190 Gif sur Yvette, France

Received 8 February 1980

1. Introduction

Many of the properties of the plasma membrane glycoproteins have been investigated by using several lectins, including concanavalin A (con A). Although the precise mechanism of lectin action on the cell surface is not known, interaction of con A with plasma membrane involves the recognition of sugar residues [1]. Moreover, the binding of con A to cell surface macromolecules can be specifically reversed by the addition of α-methyl-D-mannoside or α-methyl-D-glucoside [2]. The interactions of con A with plasma membranes have been shown to cause a number of biological effects including cell agglutination [3], transformation of lymphocytes [4] and mitotic induction [5] and to affect membrane enzyme activities such as 5'-nucleotidase [6], ATPase [7] and sialyltransferase [8]. Con A has also been found to modify the binding [9-11] or the activity [12-14] of certain hormones, and to mimic several hormonal effects [15,16].

Here we demonstrate that con A is able to inhibit the basal and stimulated hormone release by rat pituitary cells in culture.

2. Materials and methods

Culture medium and fetal calf serum were obtained from GibcoBiocult (Paisley). [3H]Leucine, [3H]-glucosamine and [3H]thymidine were from CEA (Saclay). Plastic Petri dishes were purchased from Falcon Plastics Co. (Oxnard, CA). Thyrotropin releasing hormone (TRH) was obtained from UCB (Brussels) and GnRH was a generous gift of Dr R. Studer from Hoffman La Roche (Basel). Con A was from Pharmacia (Uppsala) and α-methyl-D-mannoside from Sigma (St Louis, MO). All other chemicals were of reagent grade.

2.1. Cell culture and incubations

Dispersed rat anterior pituitary cells in monolayer cultures were prepared according to [17]. After dispersion, the cells were suspended in medium F-10 with the addition of fetal calf serum (2.5%), rat serum (15%), N2-hydroxyethylpiperazine-N2-ethane sulfonic acid (Hepes, 25 mM) and a mixture of antibiotics (50 IU penicillin/ml, 50 μg streptomycin/ml and 2.5 μg fungizone/ml). Each Petri dish was seeded with 4-8 X 10⁵ cells in 1 ml/culture medium. Usually 5 dishes/group were used. The cells were used on day 5 of the culture when adrenocorticotrophic hormone (ACTH), thyroid stimulating hormone (TSH), prolactin (PRL) and growth hormone (GH) release were studied. For follicle stimulating hormone (FSH) and luteinizing hormone (LH) studies, the cultures were used on day 7 with a change of medium on day 4. After washing the cells, the incubations were carried out in the synthetic medium F-10 for 4 h. For the studies with high K⁺, the cells were incubated in Krebs-Ringer bicarbonate buffer containing 59 mM K⁺ (instead of Na⁺), 16 mM glucose and 15 mM Hepes.

2.2. Radioimmunoassays

Radioimmunoassays for rat TSH, PRL and GH were performed using kits provided by the NIAMDD rat pituitary hormone distribution program. ACTH

+ Present address: Laboratoire de Médecine Expérimentale, FRA INSERM no. 30, Faculté de Médecine Alexis Carrel, 69372 Lyon Codex 2, France
was radioimmunoassayed according to [18], using antiseraum kindly provided by Dr Vaudry. LH and FSH were estimated by radioimmunoassay as in [19], using anti-ovine LHβ and anti-ovine FSH obtained from Dr Kerdelhue (this laboratory). Our own preparations of rat LH (1.26 × NIH-LH-S₁) and rat FSH (37.60 × NIH-FSH-S₁) were used as standard preparations.

2.3. Incorporation of labelled precursors

After incubation in the presence of labelled precursors (2.5 μCi/dish), the cells were washed and 0.5 ml water were added. The dishes were frozen and thawed. The broken cells were detached with a rubber policeman, transferred into test tubes and centrifuged at 2000 × g for 20 min. The clear supernatants were used for measuring the incorporation of [3H]leucine and [3H]glucosamine into trichloroacetic acid-precipitable proteins and for measuring the incorporation of [3H]thymidine into deoxyribonucleic acid (DNA) [20].

2.4. Hypothalamic extracts

Lyophilised hypothalamic extracts (HE) used for ACTH studies [21] were dissolved in phosphate-buffered saline so that the equivalent of 0.2 hypothalamus/dish was added.

3. Results

As shown in fig.1, both basal and GnRH-stimulated FSH release were inhibited by con A in a dose-related manner. At 100 μg/ml, con A was able to decrease the basal FSH release by ~90% and to completely inhibit the releasing action of GnRH. The inhibition patterns of the basal and stimulated release of LH, TSH, ACTH, GH and PRL, observed in the presence of con A, were dose-related in a similar way (not shown).

Whatever the secretagogue used, physiological (TRH, GnRH, HE) or pharmacological (8 Br-cAMP, high K'), con A was able to decrease the release of each pituitary hormone tested (fig.2). The effect of con A was partially reversed by α-methyl-D-mannoside (1 mg/ml). In contrast, con A was unable to modify the incorporation of [3H]leucine and [3H]glucosamine into trichloroacetic acid-precipitable proteins and the incorporation of [3H]thymidine into DNA (table 1).

---

Fig.1. Effect of increasing concentrations of con A on the basal (○—○) and the GnRH (9 ng/ml) (●—●) induced release of FSH by pituitary cells in culture, during a 4 h incubation.

Fig.2. Effect of con A (50 μg/ml) on the basal and stimulated hormone release by pituitary cells in culture during 4 h incubations. LH and FSH release was stimulated by GnRH (9 ng/ml), that of TSH by TRH (10 ng/ml), 59 mM K⁺ and 8 Br-cAMP (1 mM), that of ACTH by HE (0.2 hypothalamus/dish), 59 mM K⁺ and 8 Br-cAMP (3 mM), that of PRL by TRH (10 ng/ml) and 59 mM K⁺ and that of GH by 8 Br-cAMP (1 mM). The concentration of α-methyl-D-mannoside was 1 mg/ml.
Table 1

Lack of effect of con A (100 ng/ml) on the incorporation of \[^{3}H\]leucine and \[^{3}H\]glucosamine into trichloroacetic acid-precipitable proteins and on the incorporation of \[^{3}H\]thymidine into DNA

<table>
<thead>
<tr>
<th>Group</th>
<th>Incorporation of [^{3}H]Leucine into proteins (cpm)</th>
<th>Incorporation of [^{3}H]Glucosamine into proteins (cpm)</th>
<th>Incorporation of [^{3}H]Thymidine into DNA (cpm)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>135 502 ± 9363</td>
<td>13 855 ± 1182</td>
<td>88 186 ± 3749</td>
</tr>
<tr>
<td>Con A (100 µg/ml)</td>
<td>148 832 ± 13 157</td>
<td>17 351 ± 372</td>
<td>86 895 ± 6815</td>
</tr>
</tbody>
</table>

The incubations were carried out for 4 h in the presence of the labelled precursors (2.5 µCi/ml)

4. Discussion

This paper shows that con A is able to decrease the hormone release from rat pituitary cell cultures. This was true for every hormone tested. The effect of con A was dose-related and partially reversed by α-methyl-D-mannoside.

Con A has been demonstrated to inhibit the binding of prolactin to mammary tumors [9] and to particulate fractions of rat liver [10]. The binding of insulin to both intact cells and solubilized receptors was also inhibited by con A [11]. In contrast, con A had a protective effect on the stimulation of the adenylate cyclase activity of adrenocortical plasma membranes, by ACTH [12], whereas it facilitated [13,14] or decreased [14] the in vitro thyroid activation by TSH, depending on lectin concentration. Taking these results into account, one could postulate that the inhibition of the pituitary hormone release observed here was due to an alteration by con A of the binding of the hypothalamic releasing hormones tested (TRH, GnRH and CRF from HE) to their specific receptors. However this explanation cannot account for the inhibition of the hormone release stimulated by either 59 mM K⁺ or 8 Br-cAMP. It could be thought that the permeability of the pituitary cells to both K⁺ and 8 Br-cAMP was decreased because of a major alteration of the plasma membrane transport processes by con A. This is unlikely since our results showed that the incorporation of \[^{3}H\]leucine and \[^{3}H\]glucosamine into trichloroacetic acid-precipitable proteins, and the incorporation of \[^{3}H\]thymidine into DNA were not altered by con A, indicating that the membrane transport of amino acids, carbohydrates and nucleotides were not modified. Since the release of each pituitary hormone tested was decreased by con A, whatever the secretagogues used, an alternative explanation for our results would be that con A acted at the hormone release level. This would be in good agreement with the fact that the basal hormone release was also inhibited by the lectin. Therefore, our results suggest that in addition to its effect on the interactions of certain hormones with their receptors [9–14], con A may alter the function of other carbohydrate containing membrane components involved in the transport of hormones across the plasma membrane, during their release.

Acknowledgements

We are indebted to Dr A. Parlow and to the NIAMDD rat pituitary hormone distribution program for providing us the radioimmunoassay kits. This work was supported by grants from Délégation Générale à la Recherche Scientifique et Technique (contracts no. 78.7.0324 and 78.7.2002).

References