Fertilization *in vitro* of rat oocytes undergoing maturation in response to a GnRH analogue

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Summary. Oocytes were exposed to GnRHa to induce their maturation both *in vivo*, by administration of the hormone to hypophysectomized rats, and *in vitro*, in cultures of intact ovarian follicles. Mature oocytes obtained under both these conditions were then exposed *in vitro* to a sperm suspension for fertilization. Fertilization of control groups of oocytes, isolated from intact or hypophysectomized PMSG-primed hCG-induced ovulators, was $88.3 \pm 3.3\%$ ($n = 331$) and $90.0 \pm 2.8\%$ ($n = 427$), respectively, as compared to $82.8 \pm 3.2\%$ ($n = 413$) for oocytes isolated from hypophysectomized PMSG-primed GnRHa-induced ovulators. Fertilization rate in oocytes treated by GnRHa *in vitro* was $78.5 \pm 3.1\%$ ($n = 247$) as compared to $79.3 \pm 4.1\%$ ($n = 261$) in LH-treated oocytes. These results demonstrate that fertilizability of oocytes undergoing maturation in response to GnRHa is similar to that of oocytes induced to mature by LH. No differences could be detected in the proportions of abnormal oocytes (polyspermic, fragmented and dead) and the zygotes obtained after fertilization of GnRHa- or LH-treated oocytes showed similar ability to cleave.

Introduction

Until fairly recently it was generally accepted that the pituitary, which is stimulated by gonadotrophin-releasing hormone (GnRH) to synthesize and secrete gonadotrophins, is the only target for this hormone. However, later experiments have shown that GnRH and its agonist analogues can also elicit ovarian responses *in vitro*, in isolated ovarian cells (Clayton *et al.*, 1979; Hsueh & Erickson, 1979; Labrie *et al.*, 1979; Behrman *et al.*, 1980; Clark *et al.*, 1980; Hsueh & Jones, 1981; Knecht *et al.*, 1983) and *in vivo* in hypophysectomized rats (Ying & Guillemin, 1979; Hsueh *et al.*, 1980; Corbin & Bex, 1981; Ekholm *et al.*, 1981; Dekel *et al.*, 1983, 1985; Erickson *et al.*, 1983; Naor *et al.*, 1983). These findings, together with the demonstration of ovarian receptors for GnRH (Clayton *et al.*, 1979; Jones *et al.*, 1980; Clayton & Catt, 1981; Seguin *et al.*, 1982), suggested that GnRH can interact with the ovary in a direct manner.

It is known that the follicular oocyte in mammals is arrested at the diplotene stage of prophase of the first meiotic division (Austin, 1961) and that the physiological stimulus for resumption of meiosis is provided by the preovulatory surge of luteinizing hormone (LH) (Ayalon *et al.*, 1972). The observation that GnRH could mimic LH action, inducing meiosis resumption, was initially reported by Hillensjö & LeMaire (1980), who found that exposure of isolated ovarian follicles to GnRH or its agonist analogues *in vitro* results in maturation of the oocytes. The direct stimulatory action of GnRH on the ovary has also been demonstrated *in vivo*. Both oocyte maturation and ovulation were induced in hypophysectomized rats after administration of GnRH agonists (Hsueh & Erickson, 1979; Corbin & Bex, 1981; Ekholm *et al.*, 1981; Dekel *et al.*, 1983, 1985; Erickson *et al.*, 1983; Naor *et al.*, 1983).

The studies discussed above demonstrated that GnRH, like LH, can stimulate the oocyte to

mature, and can trigger the follicle to release the mature oocyte. LH action results in the release of a functional fertilizable oocyte; after sperm penetration this oocyte will develop into a normal embryo. Neither fertilization nor normal development, however, have yet been demonstrated following stimulation by GnRH. On the contrary, it has been reported that a significant fraction of degenerated oocytes is present in ovaries of GnRH-treated rats (Hsueh *et al.*, 1980; Erikson *et al.*, 1983; Naor *et al.*, 1983), indicating that the developmental ability of these oocytes may be impaired. The diagnosis of GnRH-induced oocyte maturation is based only on morphological markers such as germinal vesicle breakdown or polar body formation (Hillensjö & LeMaire, 1980; Ekholm *et al.*, 1981; Dekel *et al.*, 1983, 1985). Our study was designed to assess the ability of oocytes undergoing maturation in response to GnRH to be fertilized.

Materials and Methods

Hormones. The GnRH agonist analogue [D-Ser(t-Bu)⁶]des-Gly¹⁰-GnRH-*N*-ethylamide (GnRHa), used in this study was purchased from Hoechst AG (Frankfurt, West Germany). To study the effect of LH *in vitro* NIH-LH-S20 was added to the culture medium. For hormonal preparation of the rats, PMSG (Gestyl: Organon, Oss, The Netherlands) and hCG (Pregnyl: Organon) were used.

Oocyte maturation and ovulation in vivo. We have characterized the induction of oocyte maturation and ovulation by GnRHa in hypophysectomized rats of our departmental colony (Dekel *et al.*, 1985). This rat model was used in the present study as the source for oocytes undergoing maturation in response to GnRHa *in vivo*. Briefly, 25-day-old rats were injected with PMSG (15 i.u./rat) immediately after hypophysectomy and with GnRHa (2 µg/rat) or hCG (4 i.u./rat) 48 h later. The rats were killed by 20 h after hormone administration. The ovulated oocytes were recovered from the oviducts and transferred to a sperm cell suspension for in-vitro fertilization.

Maturation of oocytes in vitro. Oocytes matured *in vitro* in response to GnRHa were obtained by the use of a follicle-enclosed oocyte culture system. The effect of both GnRHa and LH on oocytes in this system has previously been studied by us (Dekel *et al.*, 1983). Briefly, follicles were isolated from 27-day-old rats, 48 h after PMSG (15 i.u./rat) administration. The isolated follicles were transferred to L-15 Leibovitz tissue culture medium (Gibco, NY, U.S.A.) supplemented with 10% fetal bovine serum (Sera-Lab, Crawley Down, U.K.) and antibiotics (100 U penicillin/ml, 50 µg streptomycin/ml; Sigma, St Louis, MO, U.S.A.) and incubated in a shaking water bath at 37°C in an atmosphere of O₂:N₂ (1:1 v/v) in the absence or the presence of GnRHa (10⁻⁷ M) or LH (5 µg/ml). Oocytes recovered from the follicles at 18 h after exposure to the hormones were transferred to a sperm suspension for in-vitro fertilization.

Fertilization in vitro. Spermatozoa were collected from the uteri of mature cyclic rats soon after mating (Shalgi *et al.*, 1981) and the sperm concentration was adjusted to 4–7 × 10⁵/ml by dilution with Rat Fertilization Medium (Kaplan & Kraicer, 1978). Samples of sperm suspension (200 µl) were incubated under heavy paraffin oil at 37°C in an atmosphere of 5% CO₂ in air. The oocytes were introduced to the sperm suspension (10–15 per sample) after 3 h of sperm incubation. The ova were examined by Nomarski interference-contrast microscopy for fertilization and first cleavage after 24 h. Ova were considered fertilized when a sperm flagellum was seen in the vitellus (Shalgi & Phillips, 1982). The numbers of fragmented, dead and polyspermic oocytes were also recorded. The fraction of oocytes under each category was calculated. All data were tested using χ^2 distribution tests. Significance was defined as $P < 0.05$.

A control group of oocytes (isolated from the oviducts of 28-day-old, intact PMSG/hCG-treated rats) was included for each experiment. This control group was used to standardize the results according to daily variations in fertilization rates which may result from the use of different donors of spermatozoa (Shalgi *et al.*, 1981).

Results

Fertilization rate of the control groups of oocytes isolated from intact or hypophysectomized PMSG-primed, hCG-induced ovulators was 88.3 ± 3.3% ($n = 331$) and 90.0 ± 2.8% ($n = 427$) respectively. The success rate of fertilization of oocytes undergoing maturation and ovulation *in vivo* in response to GnRHa was not significantly different (82.8 ± 3.2, $n = 413$). Fertilization rate of oocytes exposed to GnRHa *in vitro* was lower than that of control oocytes, isolated from the oviducts of intact PMSG/hCG-treated rats (78.5 ± 3.1, $n = 247$, compared to 91.7 ± 2.4, $n = 379$), but did not differ significantly from that of oocytes exposed *in vitro* to LH (79.3 ± 4.1, $n = 261$ $P > 0.1$). The extent of fragmentation and death was also significantly higher in the oocytes exposed to hormones *in vitro* ($P < 0.01$), with no differences between the two hormonal treatments ($P > 0.1$) (Table 1).

Table 1. Numbers of fertilized oocytes with polyspermia, fragmented or dead oocytes, and 2-cell embryos after incubation for 24 h with spermatozoa

	Fragmented/dead oocytes		Polyspermic oocytes		2-Cell embryos	
	Matured <i>in vivo</i> (%)*	Matured <i>in vitro</i> (%)*	Matured <i>in vivo</i> (%)†	Matured <i>in vitro</i> (%)†	Matured <i>in vivo</i> (%)†	Matured <i>in vitro</i> (%)†
Control	0/331 (0)	32/379 (8.4)	48/289 (16.6)	65/351 (18.5)	115/289 (39.8)	135/351 (38.5)
GnRHa	39/413 (9.4)	44/247 (17.8)	38/330 (11.5)	58/199 (29.1)	138/330 (41.8)	16/199 (0.8)
LH/hCG	29/427 (6.7)	45/261 (17.3)	68/374 (18.2)	65/208 (31.2)	201/374 (53.7)	5/208 (0.02)
No hormone	—	3/187 (1.6)	—	1/ 15 (6.0)	—	0/ 15 (0)

*Of total oocytes incubated with spermatozoa.

†Of fertilized oocytes.

Some of the oocytes diagnosed as fertilized were penetrated by more than one spermatozoon cell. No significant differences in the rate of polyspermic oocytes was observed between the two hormonal treatments ($P > 0.1$) (Table 1). However, polyspermia was significantly more prominent in the oocytes exposed to both hormones *in vitro* than in oocytes exposed to the hormones *in vivo* ($P < 0.01$).

The reduced ability of the fertilized oocytes undergoing maturation *in vitro* to develop was clearly manifested by the stage of the first cleavage. While about 50% of the oocytes undergoing maturation *in vivo* in response to hCG or GnRHa had the potential to cleave after fertilization, almost none of the oocytes matured *in vitro* proceeded to the 2-cell stage embryo (Table 1).

Discussion

Our present study demonstrates that the potential of oocytes undergoing maturation in response to GnRHa stimulation to be fertilized is similar to that of oocytes stimulated by LH/hCG. Moreover, their ability to develop further into a 2-cell embryo is also similar. The exploration of the mode of the direct action of GnRHa on the oocyte is of immediate clinical importance. GnRH was initially used to treat infertile women with gonadotrophin deficiency. Since hypothalamic GnRH is known to control the secretion of gonadotrophins by the pituitary, it is believed that GnRH treatment activates the pituitary–gonadal axis to induce follicular development and ovulation. However, since it is clear that, at least in the rat, GnRH acts on extrapituitary targets as well (for review see Hsueh & Jones, 1981), the direct interaction of GnRH with the ovary should be taken into consideration. Moreover, treatment with GnRH agonists is not limited to infertile patients. Massive doses of this hormone are administered daily to young girls with idiopathic precocious puberty (Yen, 1983). GnRH exerts its action on large as well as small antral follicles (N. Dekel, unpublished). As a significant fraction of oocytes at terminal stages of degeneration can be observed in the ovaries of GnRH-treated rats (Hsueh *et al.*, 1980; Erikson *et al.*, 1983; Ranta *et al.*, 1982) and as chronic treatment with GnRH stimulates excessive recruitment of oocytes followed by their massive degeneration (Naor *et al.*, 1983), there could be a possible clinical hazard using this therapy.

The studies on GnRH, discussed above, have all been performed in the rat, and an important question is whether such findings can be extended to other species. Attempts to answer this question by investigating whether ovarian GnRH receptors are present has yielded negative results for woman (Clayton & Huhtaniemi, 1982), sheep, cattle and pigs (Brown & Reeves, 1983). However, direct ovarian effects of GnRH agonists have been shown in the pig (Massicotte *et al.*, 1980), cow (Milvae & Hansel, 1980) and man (Tureck *et al.*, 1982), and specific but “low affinity”

GnRH-agonist receptors have been demonstrated in human corpus luteum (Popkin *et al.*, 1983). It is possible that inability to demonstrate receptors for GnRH in ovaries of some species results from technical problems associated with such studies because, even in rat ovaries, GnRH receptors are few in number.

Our findings cannot point towards any impaired fertilizability of oocytes exposed to GnRH. Furthermore, they do indicate that oocytes undergoing maturation in response to GnRH have a potential to cleave similar to that of LH-stimulated oocytes. They cannot, however, give any indication as to the potential of these oocytes to develop beyond the 2-cell stage. The development of the embryos derived from oocytes undergoing maturation in response to GnRH is now being investigated in our laboratory. The results obtained in the present study using oocytes exposed to hormones (GnRHa as well as LH) *in vitro* clearly indicate that this system cannot provide oocytes which have the ability to develop beyond the 2-cell stage after fertilization. For our further studies, therefore, only oocytes that have been exposed to hormones *in vivo* will be used.

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