Proliferation of granulosa and thecal cells in germinal disc and non-disc regions during follicular growth in Japanese quail (Coturnix coturnix japonica): bromodeoxyuridine incorporation in situ

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Proliferation of granulosa and thecal cells was analysed during ovarian follicular growth in laying Japanese quail. The birds were injected intraperitoneally with bromodeoxyuridine (BrdU) 10 or 4 h before ovulation, that is, before or after a preovulatory LH surge, respectively, and incorporation of BrdU by follicular tissues was detected immunocytochemically. Cells labelled with BrdU were seldom seen in the most immature follicles in the ovarian cortex, whereas many granulosa and thecal cells were labelled with BrdU in medium-sized white yolky follicles (approximately 13.3% and 14.4% in granulosa and theca layers, respectively). Ten and four hours before ovulation, the granulosa cells in the germinal disc and non-disc regions of the third largest yellow yolky follicle (F3) were labelled with BrdU (approximately 8.4% and 9.4% in germinal disc; 6.1% and 9.0% in the non-disc region), but only those in the germinal disc region were labelled (approximately 5.4% and 4.0%) in the largest yellow yolky follicle (F1). The percentage of thecal cells labelled with BrdU 4 h before ovulation was significantly higher than the percentage labelled 10 h before ovulation, and was higher in F3 (approximately 11.7%) than in F1 follicles (approximately 5.4%) 4 h before ovulation. These results show that proliferation of granulosa and thecal cells occurs in both germinal disc and non-disc regions in growing follicles, but when a follicle matures proliferation is reduced and in the case of granulosa cells it is restricted to the germinal disc region.

Introduction

The avian ovary contains numerous small non-yolky cortical follicles embedded in the ovarian stroma, and small white and large yellow preovulatory follicles. The follicular wall surrounding the oocyte consists of the granulosa and thecal layers. The germinal disc, which is believed to be a centre for oocyte growth (Yoshimura et al., 1994; Yoshimura and Bahr, 1995), is localized on the surface region of the oocyte (Yoshimura et al., 1993a). The number of granulosa cells increases approximately fivefold during the rapid growth phase of preovulatory follicles in domestic hens (Gilbert et al., 1980). The rate of proliferation of thecal cells increases as the cortical follicle develops into a white yolky follicle, and continues at a high rate during yellow-yolky follicular growth, but decreases at the final stage of follicular maturation (Callebaut et al., 1990).

The primary factors that control follicular growth are plasma gonadotrophins (Palmer and Bahr, 1989), although growth factors are also involved in the local regulation of cell proliferation (Onagbesan et al., 1994; Law et al., 1995). A study using electron microscopy shows that mitosis of granulosa cells is more frequent in the germinal disc region than in the non-disc region (Perry et al., 1978a). In support of this observation, flow cytometric analysis and [³H]thymidine incorporation studies show that granulosa cells isolated from the germinal disc region proliferate more than do those isolated from the non-disc region (Marrone et al., 1990; Tilly et al., 1992). A role for the germinal disc region in follicular growth has also been demonstrated by studies showing that destruction of the germinal disc region results in the induction of atresia (Jackson et al., 1993; Yoshimura et al., 1994; Yoshimura and Bahr, 1995). Therefore, in addition to gonadotrophins, communications between cells in germinal disc and non-germinal disc regions appear to play an important role in the control of follicular cell proliferation.

Although previous studies have investigated factors controlling the proliferation of follicular cells, most of them have used isolated granulosa cells. Callebaut et al. (1990) investigated cell mitosis in growing follicles using autoradiography, but they did not report on differences between germinal disc and non-disc regions. Therefore, more detailed studies on the in situ proliferation of granulosa and thecal cells during follicular growth are needed to clarify the mechanism of follicular growth. Our goal was to determine the mechanism by which cell proliferation is controlled during follicular growth. In this study the proliferation of granulosa and thecal cells of quails

Revised manuscript received 26 January 1996.

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were quantified in situ by use of bromodeoxyuridine (BrdU) to investigate the significance of the germinal disc region in ovarian follicular development.

**Materials and Methods**

Female Japanese quails (Coturnix coturnix japonica), regularly laying six or more eggs in a sequence, were kept in individual cages under a photoperiod of 14 h light:10 h dark and provided with food and water ad libitum.

The birds were injected i.p. with BrdU (Yamasa Co., Chiba; 40 mg kg⁻¹ body mass) dissolved in saline, 10 or 4 h before the expected time of ovulation, that is, before or after an LH surge, respectively (Doi et al., 1980). They were killed by decapitation, and the largest (F1) and third largest follicles (F3), white follicles (approximately 2–3 mm in diameter), and ovarian stroma containing cortical follicles were removed 1 h after BrdU injection. The surface of follicles was covered with OCT compound (Miles Inc., IN) and snap-frozen in isopentane–dry ice mixture. Cryostat sections (10 µm) were air dried on slides, and fixed by sequential exposure to ice-cold paraformaldehyde–picric acid (15 min), methanol (10 min) and acetone (10 min), followed by washing in PBS for 15 min (3 x 5 min).

The incorporated BrdU was detected as described by Soriano and Del Rio (1991). First, the sections were incubated with 2.5 µg proteinase K ml⁻¹ dissolved in 0.02 mol Tris–HCl 1⁻¹ buffer (pH 7.4) at 37°C for 10 min. After a brief rinsing with PBS, sections were treated with 0.1 mol HCl 1⁻¹ at room temperature for 10 min. Then incubated with 2 mol HCl 1⁻¹ at 37°C for 20 min. After washing with PBS for 15 min (3 x 5 min), the sections were incubated with 1% (w/v) BSA in PBS for 15 min. They were then incubated with mouse anti-BrdU monoclonal antibody (Sanbio, Uden) diluted to 1:10 in PBS for 2 h, followed by washing with PBS for 15 min (3 x 5 min). The primary antibody was detected using a Vectastain ABC-PO kit (Vector Lab. Inc., Burlingame, CA) following the supplier’s instructions, with 3',3-diaminobenzidine–4 HCl as substrate. After washing with water, sections were counterstained with haematoxylin. Specificity for BrdU was demonstrated using control sections prepared in the same manner except that the primary antibody was replaced by normal mouse IgG (20 µg ml⁻¹ in PBS). No nuclear staining was seen in control sections, showing that immunolabelling for BrdU was specific.

Follicles were taken from groups of five birds either 4 h or 10 h before ovulation. The number of cells labelled with BrdU and the total number of cells were counted on colour photographs. The number of cells was counted in 0.4 mm wide sections of granulosa and thecal layers and parallel to the surface of a follicle. The total numbers of cells counted in a photograph were approximately 60–100 and 160–220 in granulosa and thecal layers, respectively. The ratio of the number of labelled cells to the total number of cells in each follicle was calculated from the average counts made in two different photographs from each follicle. Values were expressed as the mean ± SEM for five follicles. Statistical comparisons among groups were made by Duncan’s multiple range test, and statistical significance was taken as P < 0.05.

**Results**

The labelling of cells with BrdU in either cortical or white-yellow follicles was similar 4 h and 10 h before ovulation. Only a few fibroblasts surrounding the granulosa layer and a few granulosa cells were labelled with BrdU in cortical follicles (Fig. 1a). In the white follicles protruding from the ovarian surface, 13.3 ± 3.3% of granulosa and 14.4 ± 2.9% of thecal cells were labelled with BrdU (Fig. 1b).

In F3 follicles 10 h before ovulation, 8.4% and 6.1% of granulosa cells were labelled with BrdU in the germinal disc and non-disc regions, respectively (Figs 1c, d and 2a). In F1 follicles 10 h before ovulation, 5.4% of the granulosa cells in the germinal disc region were labelled with BrdU, whereas no labelled granulosa cells were observed in the non-disc region (Figs 1e, f and 2a). In F3 follicles 4 h before ovulation, approximately 9% of granulosa cells of both germinal disc and non-disc regions were labelled with BrdU. In F1 follicles 4 h before ovulation, 4.0% of granulosa cells incorporated BrdU into the germinal disc region but no labelled cells were seen in the non-disc region (Fig. 2b). The differences in the populations of labelled granulosa cells between 10 h and 4 h before ovulation were not significant in either F1 or F3 follicles.

There was no significant difference between the percentage of BrdU-labelled thecal cells in the germinal disc and non-disc regions (Figs 1c–f and 2c). However, the percentage of labelled thecal cells in F3 follicles was higher than in F1 follicles 4 h before ovulation (P < 0.05). The percentage of labelled thecal cells in F1 and F3 follicles was greater at 4 h than at 10 h before ovulation (P < 0.05).

**Discussion**

The observation that cells labelled with BrdU were seldom seen in cortical follicles, but were more abundant in granulosa and thecal cells of white follicles supports the view (Callebaut et al., 1990) that granulosa and thecal cells proliferate when cortical follicles develop into white follicles. This proliferation of follicular cells may occur in association with cell differentiation, since during the transition of cortical to white follicles, androgen and oestrogen receptors appear in granulosa and thecal interstitial cells (Yoshimura et al., 1993b, 1995) and in association with the development of steroidogenic cells containing P450 aromatase (Nijita et al., 1991).

Earlier studies suggested that the germinal disc region is the growth centre for the granulosa layer (Perry et al., 1978a; Marrone et al., 1990; Tilly et al., 1992). However, in the present study there was no difference in the incorporation of BrdU by granulosa cells in germinal disc and non-disc regions in the F3 follicles. The view that the germinal disc region is a centre for the growth of the granulosa layer therefore requires re-evaluation. It seems that a factor(s) present in the germinal disc region which specifically stimulates the granulosa cell proliferation is not exclusively involved in the growth of the granulosa layer. We assume that gonadotrophins and growth factors stimulate growth of the granulosa layer in both germinal disc and non-disc regions in growing follicles, because proliferation of cultured chicken granulosa cells is stimulated by FSH and LH (Yoshimura and Tamura, 1988), cAMP...
Fig. 1. Bromodeoxyuridine labelling in laying quail 10 h before ovulation in (a) cortical follicles in the ovarian stroma showing that only a few cells are labelled. (b) A white follicle showing numerous labelled cells in the granulosa and thecal layers. (c) and (d) The third largest follicle (F3) showing labelling in granulosa and thecal layers in both germinal disc (c) and non-disc regions (d). (e) and (f) The largest follicle (F1) showing that (e) granulosa and thecal cells are labelled in germinal disc, whereas (f) only thecal cells are labelled in the non-disc. Arrows show examples of labelled cells. C: loose connective tissue coat; D: germinal disc; G: granulosa layer; O: oocyte; S: ovarian stroma; T: thecal layer; Y: yolk. Sections were counterstained with haematoxylin. Scale bar represents 40 μm.

(Yoshimura and Tamura, 1991) and epidermal growth factor (Yoshimura and Tamura, 1988; Peddie et al., 1994). However, the precise mechanism by which gonadotrophins regulate the proliferation of granulosa cells remains unknown because we did not find any significant difference in BrdU incorporation by granulosa cells from follicles removed before and after the preovulatory LH surge (10 h and 4 h before ovulation, respectively).

Differentiation of granulosa cells during follicular maturation is marked by an increase in sensitivity to LH, and the capacity to synthesize progesterone (Calvo et al., 1981; Bahr et al., 1983), loss of oestrogen receptors and an increase in progesterone receptors (Yoshimura and Bahr, 1991; Yoshimura et al., 1995). There is therefore a correlation between differentiation and cessation of proliferation of granulosa cells in the non-disc region of the F1 follicle. However, this is not the case for granulosa cells in the germinal disc region in the F1 follicle since they were labelled with BrdU. We assume that the germinal disc region in the F1 follicle contains some factor(s) which suppresses granulosa cell differentiation or stimulates granulosa cell proliferation. The source of such factor(s) may be the germinal disc of the oocyte, which exerts paracrine effects on the granulosa cells. In mammals, it has been reported that the oocyte provides granulosa cells with some growth factor-like signals (Buccione et al., 1990; Vanderhyden et al., 1990, 1992).
Although granulosa cells in the non-disc region of the F1 follicle were not labelled with BrdU, in culture they proliferate in a similar manner to the corresponding cells from the F3 follicle (Yoshimura and Tamura, 1988, 1991). In addition, some granulosa cells from the non-disc region showed mitotic features when studied using flow cytometry or \(^{3}H\)thymidine incorporation in vitro (Marrone et al., 1990; Tilly et al., 1992). The reason why granulosa cells in the non-disc region proliferate in vitro but not in situ is uncertain. We assume that cell to cell communication is important in the regulation of cell proliferation in situ. The presence of gap junctions between granulosa cells and between granulosa cells and the oocyte suggests the presence of a mechanism for such cell to cell communication (Perry et al., 1978b; Yoshimura et al., 1993a).

The percentage of thecal cells in the F3 and F1 follicles labelled with BrdU 4 h before ovulation (after the preovulatory LH surge) was higher than 10 h before ovulation (before the LH surge). Overall, thecal cells were more extensively labelled with BrdU in F3 than in F1 follicles. It therefore seems likely that the LH surge increases the proliferation of thecal cells. Unlike granulosa cells, there were no significant differences in the numbers of labelled cells between the germinal disc and non-disc regions of either F1 or F3 follicles. This observation suggests that the rate of proliferation of thecal cells changes during the development of follicles and during the ovulatory cycle. Factors originating from the oocyte and the germinal disc region may not play an important role in the proliferation of thecal cells. Our results also support the observation that mitosis of thecal cells is maintained during follicular growth but decreases during the final stage of maturation (Callegari et al., 1990). Thecal tissue is a rich source of growth factors including epidermal growth factor and transforming growth factors \(\alpha\) and \(\beta\), and their production is greater in smaller follicles (Onagbesan et al., 1994; Law et al., 1995). It is possible that these growth factors are involved in the regulation of the proliferation of thecal cells.

In conclusion, we have confirmed that granulosa cells proliferate in both the germinal disc and non-disc regions in growing follicles, but granulosa cells in the germinal disc region continue to proliferate in the mature, F1 follicle. The proliferation of thecal cells is greater in smaller follicles than in mature follicles, especially after the preovulatory LH surge.

This work was supported by grants from Kieikai Foundation (Tokyo, Japan 150) and Grants-in-Aid from Ministry of Education, Science and Culture of Japan (06603556) to Y. Yoshimura.

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