Stimulation of fibronectin production and deposition by chicken granulosa cells *in vitro* by epidermal growth factor and transforming growth factor α

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Experiments were conducted *in vitro* to examine the effect of epidermal growth factor (EGF) and transforming growth factor α (TGF- α) on the production of fibronectin (deposited, secreted into medium and cell-associated) by hen granulosa cells isolated from the largest (F_1 ; about 35 mm in diameter, mature) and third largest (F_3 ; 15–20 mm in diameter) preovulatory follicles, as well as from a pool of immature small yellow follicles (6–8 mm in diameter). The cells were incubated in culture wells coated with type IV collagen or in wells without collagen coating, and the amounts of fibronectin produced were measured using a specific ELISA. The total amount of fibronectin produced by unstimulated cells was greatest in wells containing F_1 cells and increased with time. The amount of fibronectin deposited by unstimulated cells was greatest in wells containing F_1 cells and was much higher in collagen-coated wells than in uncoated wells. Both EGF and TGF- α increased the quantity of fibronectin deposited by granulosa cells in collagen-coated and uncoated wells. Fibronectin secreted into the medium by unstimulated cells also increased with the stage of follicular maturation and was enhanced by EGF and TGF- α . The quantity of cell-associated fibronectin in granulosa cells in collagen-coated and uncoated wells was also increased by these growth factors. Type IV collagen did not have any appreciable effect on the amount of fibronectin present in the incubation medium or on cell-associated fibronectin. Because of its marked effect on deposited fibronectin, there were greater total quantities of fibronectin in culture wells coated with type IV collagen. These results demonstrate that EGF and TGF- α stimulate the production and deposition of fibronectin by chicken granulosa cells. The results also suggest that in combination with collagen IV, these growth factors can regulate the formation of the extracellular matrix (for example, basal lamina) of the hen ovarian follicle.

Introduction

The amount of the extracellular matrix glycoprotein, fibronectin, has been shown to increase in tissues that are undergoing modification (physiological or pathological), such as during embryogenesis (Linder *et al.*, 1975; Critchley *et al.*, 1979), inflammation (Fyrand, 1980; Clark *et al.*, 1981), tumour growth (Kimata *et al.*, 1982; Grimwood *et al.*, 1984), or wound repair (Kurkinen *et al.*, 1980; Grinnell *et al.*, 1981). Fibronectin binds to cell surfaces and biological materials (for example, collagen) simultaneously and influences the attachment, morphology and growth of cells (Klebe, 1974; Pearlstein, 1976). It is produced by bovine (Savion and Gospodarowicz, 1980; Bernath *et al.*, 1990), rat (Skinner *et al.*, 1985; Dorrington and Skinner, 1986; Carnegie, 1990), and chicken (Asem *et al.*, 1992, 1994; Asem and Novero, 1993a, b; Novero and Asem, 1993) ovarian granulosa cells *in vitro*. Fibronectin has been immunolocalized to the basal lamina (a basement membrane) of chicken

(Yoshimura *et al.*, 1985; Callebaut *et al.*, 1988), rat (Bagavandoss *et al.*, 1983) and bovine (Yoshimura and Okamoto, 1991) ovarian follicles. Basement membranes are extracellular matrices that perform a wide variety of functions such as molecular filtration, adhesion, cellular growth, and differentiation during important physiological processes.

The differentiation of rat and pig granulosa cells is influenced to a large extent by growth factors such as epidermal growth factor (EGF; Hsueh *et al.*, 1981; Schomberg *et al.*, 1983; May *et al.*, 1988; Bendell and Dorrington, 1990), transforming growth factor α (TGF- α ; Adashi *et al.*, 1987), transforming growth factor β (May *et al.*, 1988), and insulin-like growth factor I (May *et al.*, 1988; Roy and Greenwald, 1991), with pronounced effects on DNA synthesis (Hammond and English, 1987; May *et al.*, 1988). The differentiation of hen granulosa cells is accompanied by the production of large quantities of progesterone (Hammond *et al.*, 1981; Asem *et al.*, 1983; Bahr *et al.*, 1983; Etches and Duke, 1984; Marrone and Hertelendy, 1983).

Fibronectin is produced by chicken granulosa cells *in vitro* such that the amount produced is increased with advancing

stages of follicular maturation (Asem *et al.*, 1992; Novero and Asem, 1993). Recent studies have demonstrated that EGF and TGF- α control the transcription of genes encoding enzymes involved in steroid-hormone biosynthesis in hen granulosa cells (Li and Johnson, 1993). Because both fibronectin and steroid-hormone biosynthesis and secretion by hen granulosa cells are influenced by the stage of follicular development, and because EGF and TGF- α regulate the expression of steroidogenic enzymes in immature hen granulosa cells, it was hypothesized that these growth factors regulate fibronectin production by hen granulosa cells. The aim of the present study was to examine the effect of EGF and TGF- α on fibronectin production by hen granulosa cells, especially those obtained from immature, small yellow follicles (6–8 mm in diameter). Because fibronectin interacts with other macromolecules that participate in the structural organization of basement membranes (Ruoslahti *et al.*, 1981; Hynes and Yamada, 1982), and because type IV collagen is a major (and unique) component of basement membranes (Timpl, 1989; Yurchenco and Schittny, 1990), experiments were also designed to assess the effect of this type of collagen on fibronectin production by hen granulosa cells.

Materials and Methods

Hormones and chemicals

Mouse submaxillary gland EGF and synthetic TGF- α were obtained from ICN Biochemicals (Irvine, CA). Hepes, collagenase Type 1A, soybean trypsin inhibitor, BSA (Fraction V), leupeptin, phenylmethylsulfonyl fluoride (PMSF), Triton X-100, pepstatin, EDTA-Na, penicillin G, streptomycin, and fungizone were purchased from Sigma Chemical Co. (St Louis, MO). Medium 199 containing Hank's salts was from Gibco-BRL (Grand Island, NY). Fibronectin standard (purified human fibronectin) and rabbit anti-chicken fibronectin serum were purchased from Chemicon International (Temecula, CA). Immulon 2 microtitre plates were obtained from Dynatech Laboratories (Chantilly, VA). Peroxidase-conjugated goat anti-rabbit immunoglobulin G (IgG) and 2,2'-azino-di-3-ethylbenzthiazoline sulfonate were purchased from Boehringer-Mannheim (Indianapolis, IN). Falcon 96-well plates (uncoated) and coated with mouse type IV collagen were purchased from Collaborative Biomedical Products, Becton Dickinson Labware (Bedford, MA).

Animals, collection of follicles and isolation of granulosa cells

Single-comb White Leghorn hens in their first year of reproductive activity (obtained from Purdue Baker Farms, West Lafayette, IN) were caged individually in a windowless, ventilated room under a photoperiod of 14 h light:10 h dark. The birds had free access to a layer ration and tap water. The time of daily egg lay of each bird was noted to the nearest 30 min. They were killed by cervical dislocation 10–14 h before the expected time of ovulation of the largest preovulatory follicle. The granulosa cell layer from the largest preovulatory follicle (F_1 , about 35 mm in diameter, mature) and the third largest preovulatory follicle (F_3 , 15–20 mm in diameter, developing)

were separated from the theca layer as described by Gilbert *et al.* (1977). The immature small yellow follicles were placed in a 60 mm Petri dish containing 1–2 ml ice-cold saline. A 3–5 mm cut was made in the wall of the follicle and the yolk allowed to fall out. While holding one edge of the thecal layer with fine forceps, the follicular wall was shaken gently in the saline to release the granulosa cell layer. The granulosa cells from all types of follicle were dissociated in Medium 199 (with Hank's salts) containing 10 mmol Hepes l^{-1} , pH 7.4, 500 U collagenase ml^{-1} , 0.2 mg trypsin inhibitor ml^{-1} , 100 000 U penicillin G l^{-1} , 100 mg streptomycin l^{-1} , and 250 μg fungizone l^{-1} (Novero and Asem, 1993). Cell viability, determined by the trypan blue exclusion method, was routinely greater than 95% for cells obtained from the F_1 follicles and greater than 90% for cells obtained from the F_3 follicles or from small yellow follicles. Follicles were classified according to the criteria of Robinson and Etches (1986).

Experiments

The granulosa cells were plated at a density of $1.0\text{--}2.0 \times 10^5$ live cells ml^{-1} in 96-well Falcon plates, and incubated at 37°C in serum-free Medium 199 (with Hank's salts) buffered with 10 mmol Hepes l^{-1} (pH 7.4) and supplemented with 0.1% (w/v) BSA (Novero and Asem, 1993) in the presence or absence of different concentrations of EGF or TGF- α for 12 or 24 h. The incubation medium was not changed during the incubation period. The cells were not exposed to fetal calf serum or any other media supplement at any time during these experiments. At the end of the incubation period (12 or 24 h), the medium, designated as the soluble fibronectin, was removed and stored at $-20^\circ C$ until assayed for fibronectin (cumulative).

The wells were rinsed twice (37°C) with PBS containing 0.85% NaCl (w/v) and 10 mmol NaH_2PO_4 l^{-1} , pH 7.4, and the rinse was discarded. The cells that were attached to the wells were lysed in PBS (PBS–Triton solution) containing 0.5% Triton X-100 (v/v), 0.5 mg leupeptin l^{-1} , 1 mmol EDTA-Na l^{-1} , 0.7 mg pepstatin l^{-1} , and 0.2 mmol PMSF l^{-1} as described by Novero and Asem (1993). The PBS–Triton solution (granulosa cell lysate) was removed and stored at $-20^\circ C$ until assayed for cell-associated fibronectin (Novero and Asem, 1993). The wells were rinsed again with PBS (the rinse was discarded), and the residual fibronectin content (deposited fibronectin) was measured in the wells (Novero and Asem, 1993).

Total fibronectin produced in each incubation well was calculated as the sum of fibronectin secreted into the medium plus deposited fibronectin plus cell-associated fibronectin. Deposited fibronectin and cell-associated fibronectin are collectively referred to as insoluble fibronectin. The method used for the lysis of cultured granulosa cells did not result in the loss of fibronectin deposited (Novero and Asem, 1993). Medium (soluble) and cell-associated fibronectin were quantified by a competitive ELISA, whereas fibronectin deposited in the wells was quantified with a noncompetitive ELISA (Novero and Asem, 1993). Experiments with EGF or TGF- α were conducted with the same population of cells.

At the beginning of distribution of cells for incubation, aliquots of cell suspension equal to the amount incubated were

Fig. 1. Effect of epidermal growth factor (EGF) on total fibronectin production by chicken granulosa cells. Granulosa cells obtained from (a) F_1 (large, mature), (b) F_3 (third largest, developing), and (c) small yellow follicles were cultured for ($\blacktriangle$) 12 h or ($\square$) 24 h in the absence or presence of EGF. Each point is the mean $\pm$ SE of three separate experiments each conducted in triplicate. Where an error bar is absent, it is smaller than the size of the symbol. * $P < 0.05$ compared with the corresponding control value (0 EGF nmol l^{-1}).

placed in microfuge tubes (three tubes per follicle type). The suspension was centrifuged at the beginning of each incubation period, and the fibronectin content of the supernatant was quantified and defined as the medium fibronectin at the start. The cell pellet was lysed and the amount of fibronectin in the lysate was quantified and designated as the cell-associated (cellular) fibronectin at the start.

Measurement of medium (soluble) fibronectin

Fibronectin secreted into the incubation medium (soluble fibronectin) was measured by ELISA as described by Novero and Asem (1993). The primary antibody used was raised in rabbits against fibronectin purified from chicken plasma. The interassay and intra-assay coefficients of variation were 10% and 8%, respectively.

Measurement of cell-associated or deposited (insoluble) fibronectin

Cell-associated fibronectin was measured after cell lysis using the competitive ELISA as described by Novero and Asem (1993). Fibronectin deposited in incubation wells was measured by noncompetitive ELISA, also described by Novero and Asem (1993). The interassay and intra-assay coefficients of variation were 9% and 7%, respectively. The type IV collagen used to coat the wells did not crossreact with the anti-chicken fibronectin serum.

Statistical analyses

Each experiment contained three replicate wells per treatment, and was repeated three times unless stated otherwise. Granulosa cells were obtained from follicles pooled from at

least two hens per experiment with each animal donating follicles of the same maturational stage. The amount of fibronectin in the medium at the start was subtracted from the fibronectin in the medium at the end of each incubation. The amount of cell-associated fibronectin at the start was subtracted from cell-associated fibronectin values obtained at the end of each incubation. Data from replicate experiments were analysed by analysis of variance followed by post-hoc Tukey test to determine significant differences among treatment means. The data were also subjected to linear regression analysis to obtain correlation coefficients (r) between EGF or TGF- α concentrations and the amount of fibronectin produced. Differences at a probability level of $P \leq 0.05$ were considered significant.

Results

The total amount of fibronectin produced (deposited plus medium plus cellular) by F_1 , F_3 and small yellow follicle granulosa cells in the absence and presence of EGF or TGF- α increased with advancing stages of follicular maturation (Figs 1 and 2). Addition of EGF for 12 or 24 h stimulated total fibronectin production by granulosa cells from F_1 , F_3 and small yellow follicles (Fig. 1). The correlation coefficients between the concentration of EGF and the total amount of fibronectin produced ranged from 0.63 to 0.84 for all follicle types and periods of incubation, and they were all significant ($P < 0.005$). Similarly, addition of TGF- α for 12 or 24 h stimulated total fibronectin production by granulosa cells from F_1 , F_3 and small yellow follicles (Fig. 2). The correlation coefficients between the concentration of TGF- α and the total amount of fibronectin produced ranged from 0.78 to 0.95 for all follicle types and periods of incubation, and they were all significant ($P < 0.005$).

Fig. 2. Effect of transforming growth factor α (TGF- α) on total fibronectin production by chicken granulosa cells. Granulosa cells obtained from (a) F₁ (large, mature), (b) F₃ (third largest, developing), and (c) small yellow follicles were cultured for 12 h (■) or 24 h (△) in the absence or presence of TGF- α . Each point is the mean $\pm$ SE of three separate experiments each conducted in triplicate. Where an error bar is absent, it is smaller than the size of the symbol. * $P < 0.05$ compared with the corresponding control value (0 TGF- α nmol l⁻¹).

Influence of collagen IV on fibronectin distribution

The influence of type IV collagen (a unique component of basement membranes) on the distribution of fibronectin produced (deposited, medium and cellular) by hen granulosa cells was examined. Granulosa cells were incubated in culture wells coated with type IV collagen. Uncoated plates served as controls.

Deposited fibronectin. The quantity of fibronectin deposited in wells coated with type IV collagen or in uncoated wells by unstimulated (no EGF or TGF- α added) F₁, F₃ or small yellow follicle granulosa cells increased concomitantly with advancing follicular development (Figs 3 and 4). The amount of fibronectin deposited by each cell type (unstimulated) was greater ($P < 0.05$) in collagen-coated wells than in uncoated wells (Figs 3 and 4). Addition of EGF to cells in uncoated wells increased the quantity of deposited fibronectin by F₁, F₃ and small yellow follicle granulosa cells (the value of r was between 0.88 and 0.99; $P < 0.01$). EGF also increased the amount of fibronectin deposited in collagen-coated wells by F₁, F₃ and small yellow follicle granulosa cells (r was between 0.86 and 0.97; $P < 0.05$).

Addition of TGF- α to granulosa cells in uncoated wells increased the quantity of deposited fibronectin by F₁, F₃ and small yellow follicle granulosa cells (r was between 0.87 and 0.95; $P < 0.05$). Similarly, TGF- α increased the quantity of fibronectin deposited in collagen-coated wells by F₁, F₃ and small yellow follicle granulosa cells (r was between 0.75 and 0.97; $P < 0.05$). The amount of fibronectin deposited after addition of EGF or TGF- α was greater ($P < 0.05$) in each cell type in collagen-coated wells than in uncoated wells (Figs 3 and 4). The quantity of fibronectin deposited by granulosa cells following exposure to EGF and TGF- α was less ($P < 0.05$) for cells obtained from small yellow follicles than from F₁ and F₃ follicles (Figs 3 and 4).

Fibronectin in the medium. The amount of fibronectin secreted into the incubation medium by unstimulated (no EGF or TGF- α added) F₁, F₃ and small yellow follicle cells in wells coated with type IV collagen or in uncoated wells increased with advanced stages of follicular maturation (Figs 5 and 6). Addition of EGF increased the quantity of fibronectin secreted into the incubation medium in uncoated wells by granulosa cells obtained from F₁, F₃ and small yellow follicles (r was between 0.81 and 0.97; $P < 0.05$). Similarly, EGF increased the amount of fibronectin secreted into the incubation medium in collagen-coated wells by F₁, F₃ and small yellow follicle granulosa cells (r was between 0.86 and 0.95; $P < 0.05$) (Fig. 5). Like EGF, TGF- α increased the quantity of fibronectin secreted into the medium in uncoated wells by F₁, F₃ and small yellow follicle granulosa cells (r was between 0.76 and 0.83; $P < 0.05$) (Fig. 6). TGF- α also increased the amount of fibronectin secreted into the medium in collagen-coated wells by F₁, F₃ and small yellow follicle granulosa cells (r was between 0.88 and 0.95; $P < 0.01$) (Fig. 6). Unlike deposited fibronectin, the quantities of fibronectin secreted into the medium in collagen-coated wells (in the presence and absence of the growth factors) were not different ($P > 0.05$) from those in the uncoated wells (Figs 5 and 6). Fibronectin secreted into the medium following exposure to EGF and TGF- α was less ($P < 0.05$) from cells obtained from small yellow follicles than from cells obtained from F₁ and F₃ follicles (Figs 5 and 6).

Cellular (cell-associated) fibronectin. The quantity of cellular fibronectin produced by unstimulated F₁ granulosa cells (no EGF and TGF- α) was less ($P < 0.05$) than that produced by F₃ or small yellow follicle granulosa cells (Figs 7 and 8). EGF increased the quantity of cellular fibronectin (Fig. 7) in F₁, F₃ and small yellow follicle granulosa cells in uncoated wells (r was between 0.90 and 0.99; $P < 0.005$). Similarly, EGF

Fig. 3. Influence of type IV collagen on stimulation of fibronectin deposition by epidermal growth factor (EGF) in chicken granulosa cells. Granulosa cells obtained from (a) F_1 (large, mature), (b) F_3 (third largest, developing), and (c) small yellow follicles were cultured for 24 h in wells coated with type IV collagen (□) or in uncoated wells (▲) in the absence or presence of EGF. Inset: data for small yellow follicle granulosa cells are presented on an expanded scale. Each point is the mean $\pm$ SE of two separate experiments each conducted in triplicate. Where an error bar is absent, it is smaller than the size of the symbol. * $P < 0.05$ compared with the corresponding control value (0 EGF nmol l^{-1}).

Fig. 4. Influence of type IV collagen on stimulation of fibronectin deposition by transforming growth factor α (TGF- α) in chicken granulosa cells. Granulosa cells obtained from (a) F_1 (large, mature), (b) F_3 (third largest, developing), and (c) small yellow follicles were cultured for 24 h in wells coated with type IV collagen (Δ) or in uncoated wells ($\blacksquare$) in the absence or presence of TGF- α . Inset: data for small yellow follicle granulosa cells are presented on an expanded scale. Each point is the mean $\pm$ SE of two separate experiments each conducted in triplicate. Where an error bar is absent, it is smaller than the size of the symbol. * $P < 0.05$ compared with the corresponding control value (0 TGF- α nmol l^{-1}).

increased the quantity of cellular fibronectin (Fig. 7) in F_1 , F_3 and small yellow follicle granulosa cells in collagen-coated wells (r was between 0.89 and 0.98; $P < 0.01$). TGF- α caused an increase in the quantity of cell-associated fibronectin (Fig. 8) in F_1 , F_3 and small yellow follicle granulosa cells in uncoated wells

(r was between 0.92 and 0.98; $P < 0.005$). Likewise, TGF- α increased the quantity of fibronectin associated (Fig. 8) with F_1 , F_3 and small yellow follicle granulosa cells in collagen-coated wells (r was between 0.91 and 0.96; $P < 0.005$). The amount of fibronectin that was associated with granulosa cells following

Fig. 5. Influence of type IV collagen on stimulation of soluble (medium) fibronectin production by epidermal growth factor (EGF) in chicken granulosa cells. Granulosa cells obtained from (a) F_1 (large, mature), (b) F_3 (third largest, developing) and (c) small yellow follicles were cultured for 24 h in wells coated with type IV collagen ($\square$) or in uncoated wells ($\blacktriangle$) in the absence or presence of EGF. Each point is the mean $\pm$ SE of two separate experiments each conducted in triplicate. Where an error bar is absent, it is smaller than the size of the symbol. * $P < 0.05$ compared with the corresponding control value (0 EGF nmol l^{-1}).

Fig. 6. Influence of type IV collagen on stimulation of soluble (medium) fibronectin production by transforming growth factor α (TGF- α) in chicken granulosa cells. Granulosa cells obtained from (a) F_1 (large, mature), (b) F_3 (third largest, developing), and (c) small yellow follicles were cultured for 24 h in wells coated with type IV collagen (Δ) or in uncoated wells ($\blacksquare$) in the absence or presence of TGF- α . Each point is the mean $\pm$ SE of two separate experiments each conducted in triplicate. Where an error bar is absent, it is smaller than the size of the symbol. * $P < 0.05$ compared with the corresponding control value (no TGF added).

exposure to EGF and TGF- α was greater ($P < 0.05$) in cells obtained from F_3 and small yellow follicles than in cells from F_1 follicles (Figs 7 and 8). Like fibronectin secreted into the medium, the quantities of cellular fibronectin in collagen-coated wells (in the presence and absence of the growth factors) were

not different ($P > 0.05$) from the amount of cellular fibronectin in uncoated wells (Figs 7 and 8).

The effect of collagen IV on the total amount of fibronectin produced (deposited, medium and cellular) was calculated. Because of its large effect on deposited fibronectin, there were

Fig. 7. Influence of type IV collagen on stimulation of cellular fibronectin production by epidermal growth factor (EGF) in chicken granulosa cells. Granulosa cells obtained from (a) F_1 (large, mature), (b) F_3 (third largest, developing) and (c) small yellow follicles were cultured for 24 h in wells coated with type IV collagen ($\square$) or in uncoated wells ($\blacktriangle$) in the absence or presence of EGF. Each point is the mean $\pm$ SE of two separate experiments each conducted in triplicate. Where an error bar is absent, it is smaller than the size of the symbol. * $P < 0.05$ compared with the corresponding control value (no EGF added).

Fig. 8. Influence of type IV collagen on stimulation of cellular fibronectin production by transforming growth factor α (TGF- α) in chicken granulosa cells. Granulosa cells obtained from (a) F_1 (large, mature), (b) F_3 (third largest, developing) and (c) small yellow follicles were cultured for 24 h in wells coated with type IV collagen ($\triangle$) or in uncoated wells ($\blacksquare$) in the absence or presence of TGF- α . Each point is the mean $\pm$ SE of two separate experiments each conducted in triplicate. Where an error bar is absent, it is smaller than the size of the symbol. * $P < 0.05$ compared with the corresponding control value (0 TGF- α nmol l^{-1}).

greater total quantities of fibronectin in culture wells coated with type IV collagen than in uncoated wells (data not shown).

Discussion

The present results demonstrate that EGF and TGF- α stimulate fibronectin production by chicken granulosa cells. The stimu-

lated production of fibronectin was reflected in each of the three compartments (medium, cell-associated and deposited) that make up total fibronectin. Stimulation of fibronectin deposition and secretion into medium by EGF and TGF- α in the present study increased with advancing stage of follicular development; however, cell-associated fibronectin stimulated by these factors was lower in granulosa cells isolated from the most mature (F_1) follicles. Similar observations were made for

the effects of FSH and 8-bromo-cyclic AMP on fibronectin production by chicken granulosa cells (Asem and Novero, 1993a; Novero and Asem, 1993).

Fibronectin produced by a wide variety of cells is incorporated into matrix material *in vitro* (Ruoslahti *et al.*, 1981; Hynes and Yamada, 1982), and the results of the present study confirm the recent reports that chicken granulosa cells produce and deposit fibronectin *in vitro* (Asem and Novero, 1993a; Novero and Asem, 1993). In view of the fact that type IV collagen is a major component of basement membranes with which other components of this structure interact (Timpl, 1989; Yurchenco and Schittny, 1990), the present data suggest that fibronectin produced by chicken granulosa cells can interact with collagen IV in the formation of basal lamina. Fibronectin has been immunolocalized to the basal lamina of ovarian follicles of domestic hens (Yoshimura *et al.*, 1985; Callebaut *et al.*, 1988). Although it remains to be proved, the quantity of fibronectin deposited by F₁, F₃ and small yellow follicles of granulosa cells in collagen-coated wells may be a reflection of the relative contribution of fibronectin by chicken granulosa cells at a particular stage of development to the formation of basal lamina *in vivo*. It is noteworthy that the maturation of the chicken ovarian follicle is associated with a marked increase in the size of the oocyte and an increase in the number of granulosa cells (Gilbert *et al.*, 1980). This growth must be accompanied by an increase in the production and deposition of various extracellular matrix proteins to maintain the continuity of the basal lamina.

The reason for increased deposition of fibronectin in wells coated with type IV collagen is not known. However, since it is a major (naturally occurring) component of basement membranes, collagen IV may increase the deposition of fibronectin by interacting with and sequestering fibronectin. Clearly, additional studies are needed to elucidate the exact mechanism by which collagen IV influences fibronectin deposition by chicken granulosa cells. The possibility remains that the effect of collagen IV on fibronectin deposition is not specific but is a general feature of extracellular matrix proteins (collagenous and non-collagenous) *in vitro*. Future studies with different types of collagen (for example type I or type III collagen) should provide information concerning the specificity of collagen IV on fibronectin deposition by chicken granulosa cells *in vitro*.

Fibronectin production in a number of cell types is stimulated by EGF (Chen *et al.*, 1977; Thorne *et al.*, 1987; Bade and Feindler, 1988). It induces fibronectin gene transcription in AKR-2B mouse embryo fibroblasts (Blatti *et al.*, 1988) and in rat liver epithelial cells (Seebacher *et al.*, 1988). The source of EGF that may influence chicken granulosa cell function *in vivo* is unknown. EGF or EGF-like substances are produced by the mammalian ovary (Rall *et al.*, 1985; Skinner *et al.*, 1987; Roy and Greenwald, 1990; Maruo *et al.*, 1993). Furthermore, TGF- α is also produced by the mammalian ovary (Kudlow *et al.*, 1987; Lobb *et al.*, 1989). Whether these growth factors are produced by the chicken ovary remains to be determined. However, the observations that EGF (mammalian) regulates a number of chicken granulosa cell functions (Pulley and Marrone, 1986; Yoshimura and Tamura, 1988; Tilly and Johnson, 1990; Li *et al.*, 1991) support the notion that this growth factor is a regulator of hen ovarian function.

It is noteworthy that the primary structure of the chicken EGF receptor has been shown to show considerable similarity to that of the human EGF receptor (Lax *et al.*, 1988). Human TGF- α binds the chicken EGF receptor with slightly higher affinity than the human EGF receptor (Lax *et al.*, 1988). The effects of TGF- α may be mediated by the mechanisms that transduce the action of EGF because of the structural similarity between TGF- α and EGF (Marquardt *et al.*, 1984). In addition, TGF- α binds to the EGF receptor in the plasma membrane (Massague, 1983; Marquardt *et al.*, 1984) and induces biological responses similar to those caused by EGF (Derynck, 1986). The growth factors (EGF and TGF- α) had no effect on number of granulosa cells in the present experiments (data not shown). It has been demonstrated that EGF, LH and FSH have no effect on the number of chicken granulosa cells *in vitro* until after 7 days in culture (Yoshimura and Tamura, 1988). The effects of EGF and TGF- α on fibronectin production (in the present studies) by chicken granulosa cells that were incubated for 12 or 24 h (in serum-free medium) may not therefore be ascribed to changes in the number of granulosa cells.

In summary, hen granulosa cells produce and deposit fibronectin. The deposition of fibronectin is greater in wells coated with type IV collagen (a unique component of basal lamina). Both EGF and TGF- α stimulated soluble (medium) and insoluble (deposited and cell-associated) fibronectin production by the cells. These data indicate that during follicular development, EGF and TGF- α in combination with collagen can increase fibronectin deposition in the extracellular matrix (for example basal lamina) by granulosa cells. The deposited fibronectin could be one of the factors that cause the reported changes in the morphology of granulosa cells during follicular development. In the chicken ovarian follicle, granulosa cells become flattened with advancing follicular development (Gilbert *et al.*, 1980).

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