

MATURATION OF OOCYTES AND VITELLOGENESIS DURING THE DEVELOPMENT OF OVARY IN THE HOUSE CRICKET *GRYLLODES SIGILLATUS* (WALKER) ORTHOPTERA : GRYLLIDAE)

A DEVAPPAUL, A. RENI PRABHA, M.A. SUBRAMANIAN AND G. VARADARAJ
P.G. & Research Department of Zoology, C.N. College, Erode - 638 004, India.
(Received 01.04.98)

SUMMARY

In *G. sigillatus*, the ovaries are panoistic type in which the ovarioles exhibit a progressive increase in dimension as the development proceeds. The yolk deposition was found to commence on the 4th day of adult emergence in the basal oocyte which attained a fully mature stage on day 8. During this process, considerable changes were observed in both the oocytes and the follicular cells from the second day of female adult emergence. In the earlier adult instars, yolk spheres appeared in the cortical region of the ooplasm and large yolk spheres in the central region during vitellogenic phase. At the time of vitellogenesis, the follicular epithelial cells of the basal oocytes were tall and columnar with intercellular spaces. As the vitellogenesis progressed, the cells became cuboidal and the oocytes developed vitelline membrane and chorion. The present study also reveals that vitellogenesis in *Grylloides sigillatus* occurs both in the penultimate and basal oocytes simultaneously.

Key words : *Grylloides sigillatus*; Oocyte maturation; Vitellogenesis.

INTRODUCTION

The aspects of oogenesis, ovarian growth, site of yolk synthesis for the growing oocytes and mode of uptake of yolk precursors from the site of origin are available for cockroaches, grasshoppers and dragon flies (1-6) and not for house crickets. The ovarian development with reference to oogenesis and vitellogenesis has been described only in the field cricket *Plebeiogryllus guttiventris* (7). Therefore, an attempt has been made to study the maturation of oocytes and vitellogenesis in the ovary of female house cricket, *Grylloides sigillatus*.

MATERIALS AND METHODS

The adult crickets were collected from their habitats, brought to the laboratory and maintained in rectangular glass jars (20x20x18 cms), for the purpose of oviposition. The

hatched young ones were maintained in separate containers. The adults and nymphs were fed *ad libitum* with moist dog biscuit. The intermoult 8th and 9th instar nymphs and the adults of 1st - 8th days were chosen for the study. The females and instars were identified by the presence of ovipositor that appears on the emergence of 8th instar nymph.

A few (4 number of each nymph and adult) were dissected out, ovaries were removed and sections of 8 μ thickness were made (8), stained with Delafields haematoxylin and alcoholic eosin. Morphometric measurements were carried out using ocular micrometer fitted with stereoscopic compound microscope.

RESULTS AND DISCUSSION

The female reproductive system of *G. sigillatus* consists of a pair of panoistic type of ovaries each of which having 32-48 ovarioles. The oogonia are successively produced from the distal germarium and they increase in size gradually while passing down the vitellarium. The oocytes are linearly arranged with various stages of development. The first zone of vitellarium consists of immature oocytes usually 10 in number, with little quantity of cytoplasm and without follicular cells. The second zone also consists of immature oocytes but the number is less (3-4) without fully surrounded by follicular cells. The third zone includes the penultimate oocytes and the fourth zone basal oocytes (Fig. 1). The ovarioles exhibit a progressive increase

Table 1 : Morphometry of the ovarioles in various instars of female *G. sigillatus*.

Instars	Germarium		Vitellarium	
	Length(μ)	Width(μ)	Length(μ)	Width(μ)
8th intermoult nymph	210.50 \pm 9.90	14.84 \pm 1.89	592.12 \pm 31.56	67.52 \pm 1.84
9th intermoult nymph	312.12 \pm 18.82 (48.27)	37.00 \pm 2.14 (149.32)	991.43 \pm 28.00 (67.44)	9.105 \pm 4.24 (34.84)
0 day adult	496.20 \pm 13.30 (58.97)	40.60 \pm 4.70 (9.72)	1457.00 \pm 69.00 (46.95)	97.00 \pm 4.00 (6.53)
2 day adult	512.00 \pm 38.00 (3.18)	42.00 \pm 5.00 (3.45)	1793.00 \pm 65.00 (23.06)	101.00 \pm 9.40 (4.12)
4 day adult	640.00 \pm 31.56 (25.00)	42.00 \pm 2.80	1919.00 \pm 33.00 (7.03)	180.00 \pm 13.00 (78.22)
6 day adult	760.00 \pm 30.64 (34.37)	42.40 \pm 1.40 (0.95)	3826.00 \pm 62.00 (99.37)	390.00 \pm 18.90 (116.66)
8 day adult	866.00 \pm 85.23 (13.95)	45.30 \pm 3.50 (6.84)	4594.00 \pm 88.50 (20.07)	558.00 \pm 32.00 (43.07)

Each value is the mean \pm SD of 15 observations. All values are significant at $P < 0.05$. Values in parenthesis indicate % increase over previous instar.

Table 2 : Morphometry of penultimate oocyte and nucleo-cytoplasmic ratio in different instars of female *G. sigillatus*.

Instars	Length(μ)	Width(μ)	Nucleo-cytoplasmic ratio
8th intermoult nymph	68.43 \pm 711	48.84 \pm 6.89	0.452
9th intermoult nymph	140.46 \pm 19.28 (105.26)	66.81 \pm 6.46 (36.79)	0.209
0 day adult	263.00 \pm 64.10 (87.24)	71.14 \pm 4.45 (6.48)	0.118
2 day adult	313.70 \pm 15.39 (19.28)	71.43 \pm 4.10 (0.40)	0.104
4 day adult	376.63 \pm 27.10 (20.06)	75.25 \pm 7.33 (5.35)	0.094
6 day adult	451.60 \pm 19.26 (19.91)	95.43 \pm 5.24 (26.82)	0.059
8 day adult	548.80 \pm 24.00 (21.52)	116.00 \pm 8.34 (21.56)	0.022

Each value is the mean \pm SD of 10 observations.

All values are significant at $P < 0.05$. Values in parenthesis indicate % increase over previous instar.

Table 3 : Morphometry of basal oocyte and nucleo-cytoplasmic ratio in different instars of female *G. sigillatus*.

Instars	Length(μ)	Width(μ)	Nucleo-cytoplasmic ratio
8th intermoult nymph	83.68 \pm 7.25	42.86 \pm 3.65	0.204
9th intermoult nymph	241.52 \pm 14.92 (188.62)	61.65 \pm 5.41 (43.84)	0.102
0 day adult	393.75 \pm 30.24 (63.03)	78.00 \pm 4.24 (26.52)	0.070
2 day adult	430.80 \pm 49.66 (9.92)	111.00 \pm 15.10 (42.30)	0.034
4 day adult	576.63 \pm 63.80 (33.85)	157.50 \pm 9.50 (41.89)	0.020
6 day adult	1568.00 \pm 250.67 (171.92)	311.00 \pm 27.70 (97.46)	0.0004
8 day adult	1793.70 \pm 240.60 (14.39)	340.00 \pm 38.00 (9.32)	0.0003

Each value is the mean \pm SD of 10 observations.

All values are significant at $P < 0.05$. Values in parenthesis indicate % increase over previous instar.

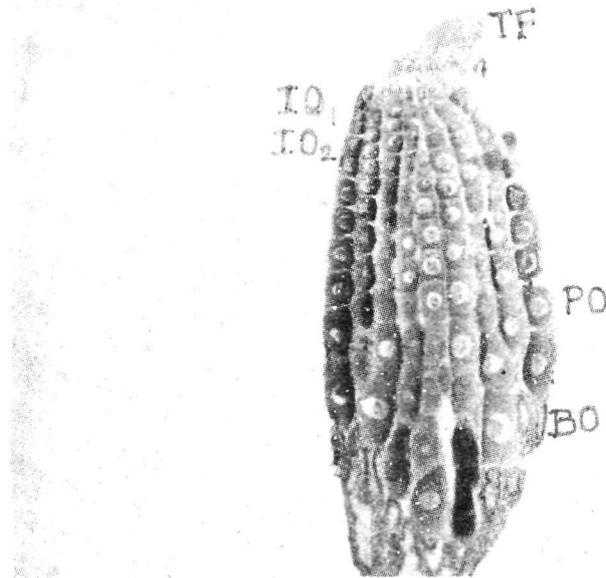


Fig. 1 : L. S. of ovary showing terminal filament (TF), immature oocytes of zone I (IO_1), immature oocytes of zone II (IO_2), penultimate oocytes (PO) and basal oocytes (BO). X100.

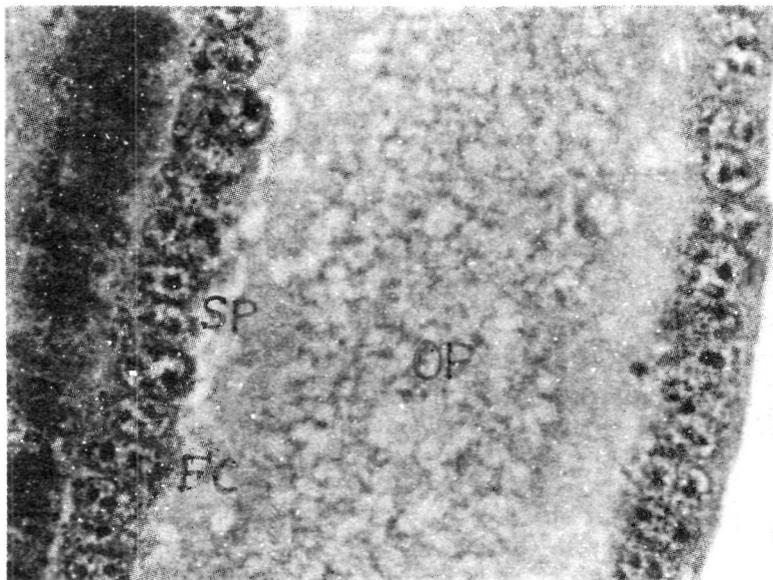


Fig. 2 : L. S. of 2-day basal oocyte (previtellogenic) showing clear space (SP) between ooplasm (OP) and follicular epithelium (FC). X400

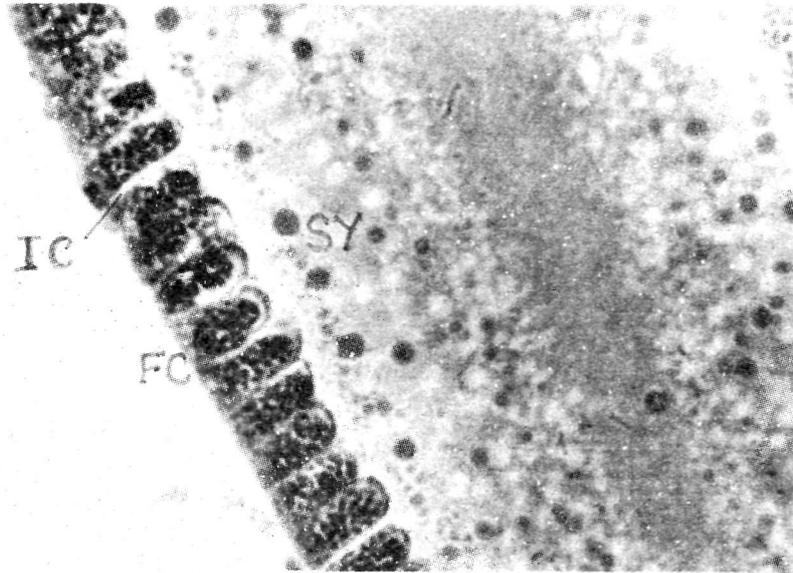


Fig. 3 : L. S. of early vitellogenic oocyte showing follicular epithelium (FC), intercellular space (IC) and small yolk spheres (SY). X400.

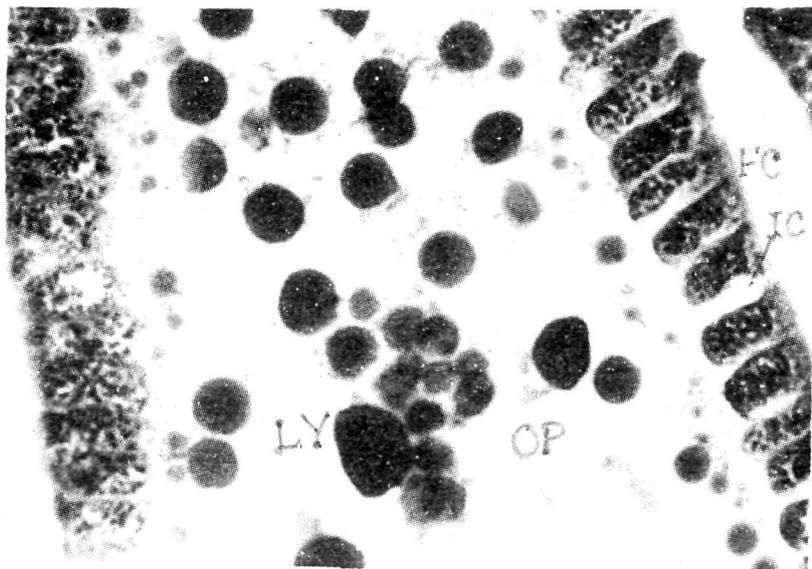


Fig. 4 : L. S. of midvitellogenic oocyte showing ooplasm (OP) with yolk spheres (LY) and follicular epithelium (FC) with intercellular spaces (IC). X400.

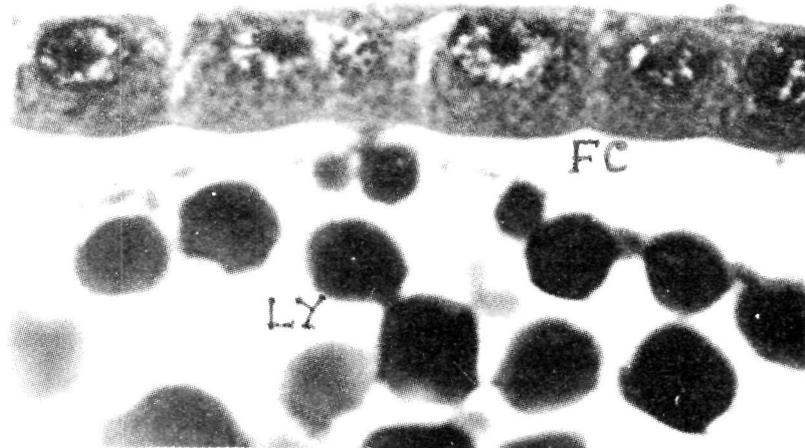


Fig. 5 : L. S. vitellogenic oocyte showing large yolk spheres (LY) and flat follicular epithelial cells (FC). X100.

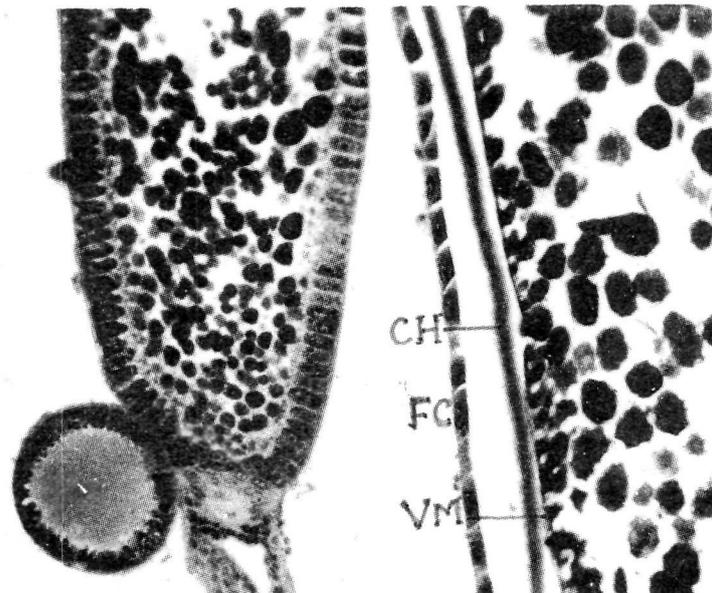


Fig. 6 : L. S. of late vitellogenic oocytes showing vitelline membrane (VM), chorion (CH) and flat follicular epithelial cells (FC). X100.

in dimension as the development proceeds (Table 1). These observations corroborate with the findings of many other orthopteran insects. (9,12)

It is evident that the penultimate and basal oocytes increase in their dimension whereas, the nucleocytoplasmic ratio decreases during the progress of development of *G.sigillatus*. (Tables 2 and 3). These could be due to vitellogenesis and maturation of oocytes in the vitellarium. In the present study, the yolk deposition in the basal oocyte is found to commence on the 4th day of adult emergence and a fully mature stage is attained on the 8th day.

Elliott and Gillott(6) have shown that a clear space appears shortly before vitellogenesis between the apical regions of the follicular cells and ooplasm in *Melanoplus sanguinipes*. The same is also true in *G.sigillatus* in which the basal oocyte contains ooplasm with clear spaces all along the epithelial lining and central vacuolated region without yolk granules. The follicular cells are columnar and compactly arranged without intercellular space. (Fig. 2)

In the present experimental insect, considerable changes have been observed both in the oocytes and in the follicular epithelial cells from the 2nd day of adult female emergence onwards. The differentiation of follicular epithelium is to be an essential pre-requisite for yolk deposition (13,14). Follicular cells of the basal oocytes of *G.sigillatus* are tall columnar type with intercellular spaces at the time of vitellogenesis (Fig. 3). These intercellular spaces could provide a passage for the uptake of extraovarian protein into the oocytes as observed in many orthopteran insects (15). The presence of yolk granules in the apex of the follicular cells observed in the present study (Fig. 3) supports the concept of intracellular pathway for protein accumulation inside the oocyte. Several investigators have reported that during the formation of yolk, proteins from haemolymph are transported through the intercellular spaces of follicular cells to the developing oocytes in insects (16-18). The same could also hold good in the present study where the follicle cells could play a decisive role in vitellogenesis by transporting substances from haemolymph into oocytes.

It has been recorded that yolk vesicles that appear in the periphery of the oocyte coalesce to produce large yolk spheres in insects (19,20). In *G.sigillatus* also yolk spheres appear in the cortical region of the ooplasm at initial adult stages (Fig. 4) and large yolk spheres in the central region of the ooplasm in the vitellogenic phase (Fig. 5). As the vitellogenesis progresses, the epithelial cells become cuboidal or squamous in the basal oocyte (Fig. 6.). This change in the configuration of the follicular cells is associated with the change in the nucleo - cytoplasmic ratio. This falls in line with the findings of Elliott and Gillott (6) and Sareen and Thukral (14) in orthopteran insects. At this stage, the follicular cells become secretory and the oocyte develops vitelline membrane (1 μ in thickness) and chorion (3-4 μ in thickness) (Fig. 5) in between follicular cells and ooplasm.

In locusts, deposition of yolk occurs only in the basal oocytes and in the penultimate oocytes after the release of basal oocytes from the ovary (5,6,20). However, in *G.sigillatus* the penultimate oocyte also contains small yolk spherules near the apices of the follicular

epithelium even when the basal oocytes with yolk spheres remain in the ovary. This suggests that vitellogenesis occurs both in penultimate and basal oocytes simultaneously in the selected species of house cricket; probably as an adaptation for higher fecundity.

REFERENCES

- 1 Nath V and Mohan P (1929). Studies on the origin of yolk IV. Oogenesis of *Periplaneta americana*. *J Morph* **48** : 253-279.
- 2 Hignam KC, Lusia O and Hill L (1963). The role of corpora allata during oocyte growth in the desert locust, *Schistocera gregaria* Forsk. *J Insect Physiol* **9** : 587-596.
- 3 Seshachar RR and Bagga S (1963). A cytochemical study of oogenesis in the dragonfly, *Pantala flavescens* (Fabricus). *Growth* **27** : 225-246.
- 4 Anderson E (1964). Oocyte differentiation and vitellogenesis in the roach *Periplaneta americana*. *J Cell Biol* **20** : 131-155.
- 5 Nath V, Mittal PK and Chandrasheikhar (1974). Morphological and cytochemical studies on the vitellogenesis of *Locusta migratoria* (L). *Res Bull Punjab Uni* **25** : 41-53.
- 6 Elliott RH and Gallott C (1976). Histological changes in the ovary in relation to yolk deposition, allatectomy and destruction of median neurosecretory cells in *Melanoplus sanguinipes*. *Can J Zool* **54** : 185-192.
- 7 Ramasamy K (1983). Neuroendocrine control of ovarian development in *Plebeiogryllus guttiventris* (Walk) (Orthoptera : Gryllidae), *Ph.D. Thesis*, Annamalai University, T.N., India.
- 8 Smith S G (1940). A new embedding schedule for insect cytology. *ST* **15** : 175-176.
- 9 Gupta PD (1948). Structure of the female reproductive organs in the orthopteroid insects. *Indian J Ent* **10** : 91-98.
- 10 Wuest J (1979). Histological and cytological studies on the ovary of *Nauphoeta cinerea* (Blattaria : Oxyhaloidea) during the first reproductive cycle. *J Inv Rep* **1** : 153-166.
- 11 Subramaniam M (1984). Studies on the male accessory reproductive glands and mating physiology in the house cricket, *Grylodes sigillatus* (Walker) Orthoptera : Gryllidae), *Ph.D. Thesis*, Univ. of Madras, Chennai, India.
- 12 Karuppanan U (1986). Studies on the development and growth rate of ovary in nymphal and adult stages of a mantid, *Euantissa pulchra* (Fabricus) (Dictyoptera : Mantidae), *The Indian Zoologist* **10** : 105-110.
- 13 Thukral D and Sareen ML (1979). Studies on the follicular epithelium in relation to vitellogenesis in Bruchids (Coleoptera). *Res Bull (Sci) of Punjab Univ* **30** : 21-26.

- 14 Sareen ML and Thukaral DA (1982). On the ovarian morphology in *Physiphora arena* (Diptera), *Res Bull (Sci) of Punjab Univ* **33** : 209-211.
- 15 Pratt JE and Davery KG (1972). The corpus allatum and oogenesis in *Rhodnius prolixus* (Stal) 1. The effect of allatextomy. *J Exp Biol* **56** : 201-214.
- 16 Ramamurthy PS (1964). On the contribution of the follicle epithelium to the deposition of yolk in the oocyte of *Panorpa communis* L. (Mecoptera). *Exp Cell Res* **33** : 601-605.
- 17 Stay B (1965). Protein uptake in the oocytes of the Cecropia moth, *J Cell Biol* **28** : 49-62.
- 18 Patchin S and Davery KG (1968). The histology of vitellogenesis in *Rhodnius prolixus*. *J insect Physiol* **14** : 1815-1820.
19. Tefler WH (1961). The route of entry and localization of blood proteins in the oocytes of saturnid moths, *J Biophys Biochem Cytol* **9** : 747-759.
20. Lusi O (1963). The histology and histochemistry of development and resorption in the terminal oocytes of the desert locust, *Schistocera gregaria*. *Q J Microsc Sci* **104** : 57-68.