Introduction

Rodent pests, distributed worldwide, are a huge menace in that they affect human health by spreading diseases and human welfare by damaging food crops and stored food grains. Soft-furred field rat, *Millardia meltada*, is one such rodent pest widely distributed in India, Pakistan, Nepal, Bangladesh and Sri Lanka, mostly found in gravelly areas, bunds of fields, cultivated areas of tropical and sub-tropical dry deciduous forests, tropical grasslands, irrigated croplands and grasslands with gravel, wherein the preferred habitats are agriculture lands, water courses, embankments, and dry rocky hills. This rat’s infestation causes pre-harvest losses of up to 1-10% in rice and wheat crops (Greaves 1989). The very high reproductive fitness of rats in general and field rat in particular overrides the conventional pest control strategies whereupon control of this pest has been gruesome. Effective olfactory (chemical) communication between male and female rats plays a vital role in this high reproductive fitness (Archunan 2009). In most mammals, the olfactory communication is mediated by pheromones which are emitted into the environment via urine, feces, saliva, and the secreted fluids of diverse integumentary scent glands. Latero-caudal, supraorbital, orbital, circumcaudal, infracaudal, preputial, clitoral, armpit, metatarsal, tarsal, preorbital, interdigital, flank, cheek, chin, anal, perineal, ventral, midventral, supracaudal, etc. are some such integumentary scent glands. They are located under the dermal layer of the skin (Balakrishnan and Alexander 1985; Archunan and Ponmanickam 2010; Rajagopal and Archunan 2011; Alexandre-Pires et al. 2014; Yilmaz et al. 2017).

Among the integumentary scent glands of rats prepu-
tial glands in the male and clitoral glands in the female are very prominent (Kannan et al. 1998; Kannan and Archunan 2001; Archunan and Ponmanickam 2010). These two glands are the prime sources of pheromones in rats and have several critical functions including conspecific attraction through releasing specific volatiles as well as non-volatile substances, mother-young bond-age, etc. (Archunan 2009; Archunan and Ponmanickam 2010). According to Zhang et al. (2008) the male rats are attracted towards the females by odors emanating from the clitoral gland and this attraction is more at the time of estrus than the other phases of the reproductive cycle. Achiraman et al. (2011) found the concentration of squalene to be significantly higher in clitoral gland secretion at the time of estrus which, therefore, could be an ovulation-indicating chemosignal in female rat. Zhang et al. (2008) identified farnesol in female rats and squalene in male rats as the major chemosignals. The predominant pheromone compound farnesol and its carrier protein α₂u-globulin have been reported in preputial gland secretion of the laboratory rat (Ponmanickam and Archunan 2006; Ponmanickam et al. 2009; Ponmanickam et al. 2010) and house rat (Kamalakkannan et al. 2006; Rajkumar et al. 2010). The preputial gland becomes atrophied after castration, and testosterone replacement restores it to its original size, which suggest that the preputial gland and the glandular proteins, *vis a vi* pheromone carrier protein, are testosterone dependent (Kamalakkannan et al. 2006; Ponmanickam et al. 2010). It is interesting to note that rat pub preputial gland releases a pheromone, dodecyl propionate, which regulates the maternal anogenital licking behavior that forms an aspect of the mother-young bond (Brouette-Lahlou et al. 1999; Ponmanickam et al. 2009). The preputial gland also enhances the poison bait efficiency by inhibiting the bait shyness when tested in house rat (Selvaraj and Archunan 2006).

The histomorphology of male preputial and female clitoral glands has been studied in a very few rat and mouse species. For instance, Knoblaugh et al. (2018) found that the preputial glands of rats and mice are lobulated and consist of a connective tissue capsule that surrounds large, cavernous ducts lined by stratified squamous epithelium and acini. The acini are composed of eosinophilic, pale, foamy, secretory sebaceous cells with dark nuclei and peripheral, flat, elongated basal cells (Ponmanickam et al. 2016). Gourbal and Gabrion (2006) investigated the histomorphological alterations induced by the parasite (*Taenia crassiceps*) in male preputial and female clitoral glands of mice, and found disorganization of the acinar cells of male preputial gland, but no impact was reflected in the histomorphology of the female clitoral glands. It was suggested that this difference between infected male and female mice might be related to the different sex hormones (Gourbal and Gabrion 2006). According to Ramachandran et al. (2018) the histological preparations of the preputial gland of *M. meltada* show acinar cells with sebum. Immunohistochemical analysis revealed the presence of α₂u-globulin, the carrier protein, in the sebum. The sebum is secreted into the central duct of the preputial gland and excreted through the urethra when it contains volatile compounds (e.g., farnesol and 6-methyl-1-heptanol) for chemical communication. Barring this, there has been no focused study of comparative histomorphology of male preputial glands and female clitoral glands. Hence, the present investigation was taken up to explore the sex-specific differences in the histomorphology of preputial and clitoral glands of soft-furred field rat, *M. meltada*.

**Materials and Methods**

**Animal**

Adult male and female rats *Millardia meltada* were collected from paddy fields in and around Madurai and Tiruchirappalli districts and housed separately in polypropylene cages (40 x 25 x 15 cm) with rice husk to 2 cm height as bedding material, at 12 h light: 12 h dark cycle, and temperature 24 ± 1 °C, when they were fed with formulated rat pellet food (Sai Durga Feeds and Foods, Bangalore) and water *ad libitum*. The bedding material was changed twice a week. The study was conducted under approval from the Institutional Animal Ethics Committee (IAEC) of Bharathidasan University, Tiruchirappalli, India (Approval No. BDU/IAEC/2012/71).

**Dissection of preputial and clitoral glands**

Six adult intact male and female rats were dissected under sodium pentobarbital anesthesia (Fig. 1). The preputial and clitoral glands were removed carefully, and the rats were sacrificed under excess ketamine (2 mg/kg, iv). The length and width of the glands were measured using a graph sheet, and then weighed in a monopan balance (Rajagopal and Archunan 2011).

**Histological study**

Immediately thereafter the glands were fixed separately in Bouin’s fluid and subjected to routine histological analysis (Humason 1979). After several changes of 70% alcohol, until the yellow color of Bouin’s fluid disappeared, the tissues were dehydrated by passing through ascending grades of alcohol, cleared in xylene, infiltrated with molten paraffin, and finally embedded in paraffin wax. Transverse and longitudinal sections at 3–5 μm...
thickness were obtained using a rotary microtome (Leica, Germany). The sections, thus obtained, were stained in Harris hematoxylin and eosin, dehydrated using alcohol, cleared in xylene and mounted in DPX adhesive resin.

Data processing

The data with respect to length, width and weight were used to calculate the respective means and standard deviations. Paired sample $t$-test was conducted and $p$ value less than 0.05 was taken to indicate significant difference.

Results

Morphological observation

The scent glands are located underneath the dermis and embedded in the subcutaneous fat. The preputial glands are bilateral and located in the subcutaneous adipose tissue, laterocranial to the penis (Fig. 1A), while the clitoral glands are situated on either side of the clitoris and immediately adjacent to the inguinal portion of the mammary gland (Fig. 1B). The paired glands (both preputial and clitoral) are yellowish-brown, pear-shaped, and dorsoventrally compressed. The preputial gland of adult male rat measured $14.16 \pm 3.81$ mm long, and $4.41 \pm 1.05$ mm wide and weighed $132.6 \pm 12.10$ mg. The clitoral gland of adult female rat measured $11.25 \pm 2.13$ mm long, and $2.95 \pm 0.75$ mm wide and weighed $97.5 \pm 10.85$ mg. Thus, the morphometric parameters [length: $t = 10.03$, df = 11; width: $t = 5.99$, df = 11 and weight: $t = 13.47$, df = 11] of preputial and clitoral glands differ significantly ($p < 0.05$) (Fig. 2).

The density and diameter of sebaceous glands were found to vary between preputial gland of male and clitoral gland female rats under microscopic investigation. The paired $t$-test clearly showed that the density of sebaceous gland is significantly ($t = 7.97$, df = 11, $p < 0.05$) higher in male preputial gland $(14.65 \pm 1.14$ units/mm$^2$) than the female clitoral gland $(8.35 \pm 0.52$ units/mm$^2$). Further, the diameter of sebaceous gland varied significantly ($t = 13.58$, df = 11, $p < 0.05$) in the male preputial gland $(29.90 \pm 1.56$ µm) compared to female clitoral gland $(18.71 \pm$ µm). The density $(6.44 \pm 0.48$ units/mm$^2$) and diameter $(24.69 \pm 0.88$ µm) of apocrine gland was also noted in the preputial glands of male rat.

Histological observation

The secretory units of preputial gland are formed of se-
Figure 3. Hematoxylin-eosin-stained sections of preputial gland of male rat. A, Sebaceous glandular lobular portion; B, Apocrine glandular lobular portion; C, Ordinary sebaceous glandular lobules; D, Modified sebaceous glandular lobules; E, Sebaceous glandular portion with a secretory acinus with the lumen distended with secretory material released from the cells by necrosis. Abbreviations: ED: Excretory duct; E: Epidermis; SG: Sebaceous gland; AG: Apocrine gland; NHGL: Normal alveolar holocrine glandular lobules; SA: Serous acini; MA: Mucus acini; LD: Lipid droplets.
baceous secretory lobules and apocrine secretory lobules (Fig. 3A, B). On the other hand, the secretory units of clitoral glands are only sebaceous glandular lobules (Fig. 4A). The sebaceous secretory gland lobules of preputial as well as clitoral glands are formed of both ordinary and modified sebaceous glandular lobules (Fig. 3C, D; Fig. 4B, C). The ordinary sebaceous glandular lobules are small and superficially located. The secretory acini formed of the modified sebaceous glandular lobules are teardrop-shaped and very large as compared to acini formed of the ordinary type of glandular lobules. The modified sebaceous glandular lobules consist of four types of cells: peripheral cells, differentiating cells, mature cells and necrotic cells. Peripheral cells are flat with oval nucleus and form a thin layer. Differentiating cells are large, polyhedral in shape, and contain centrally located nucleus in the eosinophilic cytoplasm. The differentiating cells progressively transform into mature cells which have a foamy cytoplasm. Necrotic cells possess pyknotic nuclei and on lysis release the content into the lumina.

Figure 4. Hematoxylin-eosin stained sections of clitoral gland of female rat. A, Sebaceous glandular lobules; B, Ordinary sebaceous glandular lobules; C, Modified sebaceous glandular lobules; D, Sebaceous glandular portion with a secretory acinus distended with secretory material released from cells by necrosis. Abbreviations: ED: Excretory duct; E: Epidermis; SG: Sebaceous gland; NHGL: Normal alveolar holocrine glandular lobules; SA: Serous acini; MA: Mucus acini; LD: Lipid droplets.
In both preputial and clitoral glands the secretory material of sebaceous cells is released by holocrine mechanism into the branched tubuloalveolar system wherein the secretory acini open into numerous lateral ducts that fuse to form a wide central duct through which the secretion is conveyed to the lumen of the pouch (Fig. 3E; Fig. 4D). At the distal end of the gland the central duct, containing membrane-bound secretory granules, opens onto the skin at the transition of the parietal layer of the prepuce and the end of the urethra in the tip of the penis in the case of preputial gland and vagina in the case of clitoral gland. The sebaceous type acinar cells look foamy, with small granules in the cytoplasm. The glandular epithelioid acini are composed of flat basal cells adjacent to the basement membrane and lateral to the secretory cells in the central parts of the acini. The cells are closer to the center of the alveoli and become progressively large, and the cytoplasm is distended with fat droplets due to which the cytoplasm takes honeycomb-like appearance (Fig. 3C, D; Fig. 4B, C). The apocrine secretory units of preputial glands consist of an inner layer of cuboidal cells that rest directly on the basement membrane and an outer layer of myoepithelial cells. The cuboidal cells that line the secretory acini possess acidophilic vacuolated cytoplasm and spherical, centrally located nucleus. The cells often exhibit apical protrusions. The secretory material is released in the form of droplets by apocrine mechanism and get stored in the lumina (Fig. 3B).

Discussion

The significant difference of the morphometric parameters (i.e., length, width, and weight) between preputial and clitoral glands is to be perceived as manifestation of the sexual difference of the pheromone secreting glands between male and female soft-furred field rat which is in agreement with the condition in Iranian sheep (Abbasi et al. 2009), Egyptian sheep (Awaad et al. 2015), Awassi sheep (Yilmaz et al. 2017), and Bandicoot rat (Ponmanickam et al. 2016). In another context, the preorbital gland length, width and weight were higher in dominant male (i.e., adult) than subordinate male (i.e., sub-adult and adolescent) blackbucks (Rajagopal and Archunan 2011). According to Zhang et al. (2008) the scent glands of rats do not exhibit sexual differences, but gonadectomy resulted in significant decrease of size of the scent glands of both sexes. The dominant male mice are characterized by higher testosterone levels and heavier preputial glands than subordinate male mice (Mckinney and Desjardins 1973). Further, the diameter of sebaceous gland has a higher in size in male preputial gland compared to female clitoral gland. It is reported that the higher development of the sebaceous gland in the male fallow deer could depend on the higher social responsibility (dominance hierarchy) / production of testosterone (Schaal 1982; Moawad 2016). Therefore, the larger size and higher weight of male scent gland are to be taken to reflect manifestation of the androgen-support. These findings indicate that the size of scent glands is influenced by sex hormones and function to express their respective reproductive statuses to the opposite sex.

Microscopic examination of preputial and clitoral glands revealed a clear fibrous capsule around each. The histological structure of preputial gland is heterogeneous in that it has holocrine sebaceous and apocrine secretory cells, whereas sebaceous secretory cells alone are present in the entire clitoral gland. This is a clear histomorphological manifestation of sexual dimorphism. Similar findings have been made on intermandibular glands of lesser mouse deer (Agungpriyono et al. 2006), infraorbital gland of the barking deer (Adnyane et al. 2011) and preorbital gland of the blackbuck (Rajagopal and Archunan 2011), but the finding contradicts the observation made on infraorbital glands of the Japanese serow (Atoji et al. 1987), and Formosan serow (Atoji et al. 1996) and interdigital glands of sheep (Misk and Misk 2013). According to Zhang et al. (2008) rat male scent gland secretion is rich in squalene compared to female gland, whereas farnesol content was lower in male scent gland than female gland. Squalene is a female-attractant and farnesol a male-attractant. Thus, these compounds are sex-specific (Zhang et al. 2008). One of our studies showed that the amount of protein was higher in preputial glandular secretion than clitoral gland (Archunan et al. 2004). In addition, though the 20 kDa protein was found in both male and female scent glands of rat, the intensity was higher in male than female and it was suggested that this protein is indeed a sex-associated protein (Archunan et al. 2004). Thus, this study substantiates histological difference between the preputial and clitoral glands of soft-furred field rat indicating sexual dimorphism. We suggest that the sex-specific nature of pheromone and protein may be due to the variations in the histoarchitecture of the glands and sex hormone-dependence of their morphology and secretion.

As far as sebaceous secretory portion of both preputial and clitoral glands are concerned, we found ordinary as well as modified glandular lobules. The modified sebaceous glandular lobules have been found in the scent glands of several mammalian species such as antelope Madoqua sp. (Richter 1971), antelope/red duiker Cephalophus natalensis (Mainoya 1978), royal antelope Neotragus pygmaeus (Kuhn 1976), steenbok Raphicerus campestris (Cohen and Gerneke 1976), and oribi Ourebia ourebi (Sokolov et al. 1994). Nevertheless, both modified and ordinary sebaceous glandular lobules produce sebum which is an oily/waxy
substance documented to be concerned with olfactory communication in several mammals (Rajagopal and Archunan 2011; Yilmaz et al. 2017; Ramachandran et al. 2018; SankarGanesh et al. 2018). The modified sebaceous glandular lobules of the preputial glands of male mouse are larger than in the female clitoral glands, and this feature is greatly influenced by a variety of hormones (Bronson and Caroom 1971). An immunohistochemical study revealed the presence of α2u-globulin in the sebium produced in modified sebaceous glandular lobules of preputial gland. The sebum is discharged by holocrine mechanism into the central duct of the preputial gland and excreted through the urethra. The putative pheromones (e.g., farnesol) are bound to the α2u-globulin (Ramachandran et al. 2018). Ilayaraja et al. (2014) conducted research to confirm that farnesol binds with the α2u-globulin efficiently. The present study suggests that sebaceous secretory cells of scent glands of male and female rats secrete different odoriferous volatile substances. These substances provide for olfactory communication between conspecifics (Kannan et al. 1998; Zhang et al. 2008; Achiraman et al. 2011; Ramachandran et al. 2018).

In the present study, the apocrine glandular lobules were found only in the preputial gland, i.e., male. The myoepithelial cells observed in the apocrine glandular lobules revealed evidence of apocrine secretion of the secretory cells. The myoepithelial cells surround the secretory acini and facilitate discharge of the secretory material by way of their contraction (Abbasi et al. 2009; Awaad et al. 2015). Apocrine glands have been reported in sheep, horse, antelope, cow and a few marsupials, and these glands function as scent glands to produce pheromones (Robertshaw 1987; Rajagopal and Archunan 2011). According to Satoh et al. (1994) the myoepithelial cells of apocrine glands are endowed with capability to provide compression pressure for glandular expulsion of the secretory material. It is remarkable that the pheromonal compounds derived mainly from the sebaceous gland on discharge to the outside can adhere to objects in view of the carrier proteins (Atoji et al. 1987; Agungpriyono et al. 2006; Rajagopal and Archunan 2011). Previous studies have shown that the lumen of the apocrine gland is filled with dense secretory materials which are discharged directly onto the excretory duct through prepuce (Karahan et al. 2007; Abbasi et al. 2009; Rajagopal and Archunan 2011; Awaad et al. 2015). The present study suggests that the apocrine secretory gland in the male rats would produce sex-specific volatile substances as an aspect of the preputial glands that might facilitate better adhesion capacity and high persistence in the scented site during the expression of territorial/scent marking in male rats.

To the best of our knowledge this is the first comprehensive comparative histomorphological analysis of preputial and clitoral glands, the male and female scent glands respectively, of soft-furred field rat. Preputial glands of male rat are relatively larger in size than the clitoral glands of female rat. Preputial gland is composed of sebaceous and apocrine secretory glandular lobules whereas clitoral gland is made up only of sebaceous glandular lobules. These findings indicate that the scent glands of the soft-furred male and female field rats show sexual dimorphism. Based on the morphometric and histomorphological observation, we conclude that the soft-furred field rat scent glands (preputial and clitoral) would play role in the production of sex-specific volatile substances through sebaceous and apocrine secretory glands for olfactory communication. Further ultrastructural and immunohistochemical studies of different rodent species and correlation between pheromone production and behaviors would throw light on importance of these glands in pheromonal communication which can be made use of in the rodent pest management programs.

Acknowledgements

The authors thank the Principal and the Management of Thiagarajar College (Autonomous), Madurai, for providing facilities and the constant encouragement. GA thanks the University Grants Commission (UGC), New Delhi, for the award of UGC-BSR Faculty Fellowship. The authors declare that there is no conflict of interest.

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