Skin pattern structure and function of juvenile ages of *Chameleo chameleon*

Yosra A. Fouda a,b, Ahmed A. El Mansi a,*

a Zoology Department, Faculty of Science, Mansoura University, Mansoura, Egypt  
b Biology Department, Faculty of Science and Arts, Qilwah, Baha University, Saudi Arabia

**Abstract**

Little is known about the skin structure of juvenile chameleon especially its sensory function of their integumentary structure. Fifteen juvenile *Chameleo chameleon* are collected from Abu Rawash, Northern area of Giza, Egypt during Summer of 2015. It is belong to the order Squamata, family, Chamaeleonidae. Three ages are used in the present study and categorized according to the morphological criteria of head, abdomen and limb lengths. Dorsal abdominal surfaces are covered with abdominal scales of varying sizes either conical or elliptical-structures, regularly arranged in rows and imbricated with each other. Each scale possessed one cylindrical lenticular epidermal sense organ containing heavy sensillia. Histologically, the scales are characterized by wider conical surfaces and intermingled with another one by hinge region. The epidermal layer of outer scale surface is composed of five-layered stratified squamous epithelium including the stratum germinativum, intermediate zone of stratum spinosum and granulosum, α-keratin layer, β-keratin layer and outer superficial Oberhäutchen. Melanosomes are abundant in the intermediate zone as well as in the peripheral dermal layer underneath stratum germinativum layer. The melanosomes possessed long cellular processes with their content of melanin granules underneath the epidermis. The dermis is composed of upper collagenous and inner compact layer. Semithin sections revealed the presence of fibroblast cells, collagenous fibrils, nerve axons, melanosomes and mast cells in the connective tissue core. Increased immunoreaction of cytokeatin is observed in the epidermal layers of G3; meanwhile, an increased proliferation of epidermal and dermal cells was detected in G1. Transmission electron microscopy exhibited striking formation of dermal sense organs containing neuronal cells of both oligodendrocytes and Schwann cells with myelinated and unmyelinated nerve axons ensheathed externally by thin collagenous fibers. Finally, the author concluded that the juvenile chameleon skin is keratinized with obvious external and internal sensation and abundant mast cells within dermis giving characteristic immunity. The melanosomes are dispersed within epidermis and dermis allowing the animal to maintain its color alterations according to the surrounding environment.

© 2017 Mansoura University. Production and hosting by Elsevier B.V. This is an open access article under the CC BY-NC-ND license ([http://creativecommons.org/licenses/by-nc-nd/4.0/](http://creativecommons.org/licenses/by-nc-nd/4.0/)).

**1. Introduction**

Squamata possessed varying skin structures. As a result of living in desert, the hardness of reptilian integument keeps the animal internal organs from damage and dehydration [1].

The scaly covering of epidermis represents the most adaptive function for water retention that maintain flourishing and accommodation of reptilian for a terrestrial life [2]. Characteristic variations of scale shape, size, and their overlapping are represented in reptilian species [3].

The epidermis mainly composed of five major layers, Oberhäutchen, β-keratin layer, α-keratin layer, intermediate zone and stratum germinativum [4–7]. The stratum germinativum or basale is made up of single layer of highly proliferative cuboidal cells and the source of keratinocytes progenitors [4]. After proliferation of the neo-keratinocytes, it migrates and differentiates along the way to the outer Oberhäutchen. The intermediate zone contains keratinocytes at different developmental stages where lipids and proteins are packaged in granular structures. Later, the lipid envelopes are created by releasing the lipid content to form the corneoocytes of the superficial layer [8]. The α-layer is soft and elastic and has specific connections to the preceding scales and the β-layer, which is relatively hard, forms the surface of the scales [4].

The dermis is mainly composed of connective tissue and collagen fibers with many important structures including chromatophores, lymphatic vessels, nerves and blood vessels [4].
The skin of reptilian species varied in their distribution and densities of epidermal melanocytes and dermal melanophores, lipophores, and iridophores [7,9,10]. In Chameleons and some anole lizards, the chromatophores represent the main components of a dermal chromatophoric unit in combination with melanophores [11].

Integumentary sense organs represent the mechanosensory receptors in reptilian species such as Amphibolurus barbatus [12] and Iguanian lizards [13]. These sense organs serve as mechano and thermoreceptors and possibly sensitivity to humidity [12,13]. It is also served as complex touch corpuscles on the skin surface of the head of snakes [14,15]. Small tactile mechanosensory epidermal sense organs are also reported in 13 fully aquatic and two semi-aquatic species of elapids [16].

The present study aimed to illustrate the integumentary neuroepithelial interaction of juvenile chameleon and its capacity of proliferation through light, immuno and ultrastructural studies.

2. Materials & methods

Fifteen juvenile individuals of Chameleo chameleon, Linnaeus (1758) (Class: Reptilia; Order, Squamata; Suborder Sauria; Family, Chamaeleonida; Genus, Chamaeleo; Species, Chameleo chameleon) were collected from Abou–Rawash desert, Giza Governorate, Egypt during summer 2015. The specimens were categorized into three stages according to the variations of total body length, head length, tail length, body width and limbs length as represented in Table 1 and Fig. 1.

2.1. Histological investigation

2.1.1. Hematoxylin eosin staining

Dorsal skin of trunk region of different developing stages is fixed in 10% phosphate buffered formalin (pH 7.4), dehydrated in ascending percentages of ethyl alcohol, cleared in xylene and mounted in molten paraplast at 58–62 °C. Five μm sections were cut, stained with hematoxylin and eosin and investigated under a bright field Olympus light microscope.

2.1.2. Immunohistochemical staining of CK and PCNA

Immunostaining was performed on formalin-fixed and paraffin-embedded tissues. Hydrated tissue sections were firstly incubated in 2% hydrogen peroxide endogenous peroxidase activity for 5 min to block peroxidases. The slides were then incubated overnight at 4 °C in a humidified chamber with the primary antibodies of cytokeratin (Cat. sc-25280, 1:50, mouse, Santa Cruz) and PCNA (Cat. sc-56, 1:400, mouse, Santa Cruz). After rinsing with a phosphate-buffered solution, the specimens were incubated in biotinylated secondary antibody for 50 min at room temperature followed by treatment with Avidin–Biotin–horseradish peroxidase and staining with 0.04% 3,3′-diamino-benzidine tetrahydrochloride and Hematoxylin. Negative control was carried out by using the primary antibody.

2.2. Scanning electron microscope

Skin samples were fixed in 2.5% glutaraldehyde in cacodylate buffer. This was followed by dehydration in ascending percentages of ethyl alcohol and critically point drying, and coating with gold in platinum-palladium ion-sputtering and investigating under Jeol scanning electron microscope, JSM-5400LV.

2.3. Transmission electron microscope

Fresh samples were fixed in 2.5% glutaraldehyde in cacodylate buffer (pH 7.4, post-fixed in 1% osmium tetroxide, dehydrated in propylene oxide and embedded in araldite resin and allow for complete polymerisation at 60 °C. Semithin sections (1 μm) were cut at Ultracut Reichert-Jung ultramicrotome with the aid of glass knives, stained with toluidine blue and examined under light microscope. For electron microscopy ultrathin sections were carried out, stained with uranyl acetate and lead citrate and examined with a Joel CX 100.
transmission electron microscope operated at an accelerating voltage of 60 kV.

2.4. Statistical analysis

Data were presented as mean ± standard error. The statistical analysis was performed using SPSS (version 18) software package for windows.

3. Results

3.1. Scanning electron microscopy observations

The selected ages of studied young Chameleo chameleon are determined according to the length of fore and hind limb and head length (from tip of snout to the corner of the neck), Snout vent length and whole body length and tail length (Table 1 and Fig. 1). The dorsal abdominal scales are of varying sizes and appeared either conical or elliptical-shaped structures, regularly arranged in rows and overlapped with each other. The scale width is larger than its vertical axis. Micro-ridges are observed throughout its whole surface. A deep furrow separate the scale from each other. Large lenticular epidermal sense organs are emerged from the sensory lenticular dome-shaped structure (Fig. 2A–D).

3.2. Histological observations

At light microscopic level, the epidermal scales are of varying shapes taking conical, polygonal, and tubercular shaped structure. The scales are overlapped and conjugated with each other in hinge region. The scale is characterized by its wider conical surfaces and intermingled with the successive one by the hinge region. The epidermal layer of outer scale surface is composed of five-layered stratified squamous epithelium comprising the stratum germinativum, intermediate zone with different keratinocytes, α-keratin, β-keratin and the outermost Oberhautchen. The inner scale surface is made of less thickened epidermis comparing with the hinge regions which attained more thickness. Melanosomes formed a network underneath the germinativum layer and possessed long cellular processes rich with their content of melanin granules underneath the epidermis. The inner core of the scale is composed of dermal collagenous tissue arranged regularly parallel with the epidermis and rich in fibroblasts, collagenous fibrils and blood capillaries (Fig. 3A-C1). Semithin sections revealed that the dermis rich in fibroblast cells, collagenous fibrils, nerve axons, melanosomes and mast cells with secretory granules dispersed in the connective tissue core (Fig. 5A&B).

3.3. Immunohistochemical observations

Following immunohistochemical staining with cytokeratin, a dark-brown reaction is detected in the epidermal cell layers,
mostly increased in stage 2 and 3 (Table 2, Fig. 4A–C). Also, Immunohistochemical staining of PCNA revealed highest proliferation and differentiation rate of epidermal cells, fibroblast cells especially in the young age (G1) (Table 2, Fig. 4A1-C1).

3.4. Transmission electron microscopic observations

At ultrastructural level, the epidermis is composed of five layers. Basally, germinativum cells having centrally located nuclei with dispersed euchromatin and characteristic nucleoli. Fine keratohyalin granules are dispersed in between stratum germinativum cells. The intermediate contains abundant keratinocytes with cytoplasm rich in keratohyalin granules. Electron-dense smooth granules of melanosomes are detected in between the keratinocytes in the stratum granulosum layer. The epidermal surface is covered by thickened keratinized layer, the Oberhautchen (Fig. 5C&D). Underneath the epidermis, numerous mast cells with cytoplasm rich in secretory granules were observed. Bundles of sensory organs are distributed in the dermis and ensheathed with collagenous fibrils. It is composed of myelinated and unmyelinated nerve axons. Two types of neurons are detected. First, oligodendrocyte appears with peculiar electron-dense nuclear chromatin. The other type, Schwann cell with myelinated axon is observed (Fig. 6A–D).

4. Discussion

The observed dorsum body scales are overlapping with each other and take conical, polygonal and tubercle-shaped structure.
Developed Oberhautchen with characteristic α & β keratin make the hardness of epidermal surface. The epidermis is composed of stratum germinativum, intermediate zone of stratum spinosum and granulosum cells, α keratin, β keratin and the superficial Oberhautchen.

Similar findings were reported in skin of Indian Chameleon (Chamaeleo zeylanicus) [17]. Fine keratinohyaline granules are dispersed throughout the epidermal cells and characteristic differentiation of Oberhautchen into α and β keratins. Cornification is highly detected in old ages meanwhile highly proliferated cells are observed in young ages. Similar findings were reported by Alibardi [18,19] during embryonic development in the lizard Anolis lineatus. Development of the Oberhautchen with characteristic keratinization especially of the outer scale surfaces during early stages reflected the protective barrier and adaptation to the terrestrial life [20].

Abundant distribution of melanosomes is detected in peripheral epidermal layer as well as in collagenous dermal layer underneath the epidermis. The dermal melanophores are characterized by their dendritic arborization. Similar distribution of melanosomes was reported in Plestiodon laticinctus [10].

These melanophores are characteristic pattern of reptilia especially chameleon species facilitating thermoregulation via increasing body temperature in cold climate by increasing light

Fig. 4. (A-C2). Photomicrograph of histological sections of skin of Chameleo chameleon immunohistochemically stained with anti-CK and anti-PCNA. A&A1. Stage G1. B&B1. Stage G2. C&C1. Stage G3. A-C. CK immunostaining showing increased cytokeratin staining of Oberhautchen layer in G2 and G3. A1-C1. PCNA immunostaining showing increased proliferation of epidermal cells and fibroblasts in connective tissue core in G1. Abbreviations: α-K, Alpha-keratin; β-K, Beta keratin; BV, Blood vessel; CF, Collagen fibers; D, Dermis; Ep, Epidermis; HR, Hing region; me, Melanosomes; Ob, Oberhautchen.
penetration across the skin [21] as well as production of various coloring of reptiles skin [22].

Light and ultrastructural observations revealed abundant distribution of mast cells. Mast cells is highly important for the chameleon species due to its secretion of peptidases such as tryptases, chymases, carboxypeptidase A3, and dipeptidylpeptidase I (cathepsin C) which is responsible for host defense and homeostasis [23].

Immunohistochemical staining with cytokeratin show increased reaction in different epidermal layers in stage 2 and 3. Meanwhile, the immunoreaction of PCNA revealed highest proliferation and differentiation rate of epidermal cells in the young age (G1).

Alibardi [24] observed an increase in the three different anti-cytokeratins (α-keratin) in the epidermis of lizard epidermis of the lizard Podarcis sicula. Also, our results were consistent with

---

**Table 2**  
Immunoreactivity of CK and PCNA of skin of *Chameleo chameleon*.

|          | G1 | G2 | G3 |
|----------|----|----|----|
|          | CK | PCNA | CK | PCNA | CK | PCNA |
| Epidermis|    |      |    |      |    |      |
| SB       | –  | –   | +  | –    | +  | –   |
| SSP      | –  | –   | +  | –    | ++ | +   |
| SG       | +  | –   | ++ | –    | ++ | +   |
| Ob       | +  | +++ | ++ | +    | +++| +   |
| Epidermal-dermal junction | – | – | – | – | – | – |
| Dermis   | –  | –   | –  | –    | –  | –   |
Di-Poï and Milinkovitch [25] who reported an increase in cell proliferation rate in the dermo-epidermal elevation that generates the scales in lizards and snakes.

A striking finding is the presence of a single epidermal sense organ per each dorsal body scale. The sense organs take the lenticular-shape possess dense sensory hairlet-structures bearing bristles. Several studies reported similar sense organs although the emerging sensory hairlet structures are very few such as, *Oplurus fierinensis* [26], *Amphibolurus barbatus* [12], *Ceratophora, Draco*, *Phrynocephalus*, *Stellio*, and *Trapelus* (agamids) and in *Anolis, Chalarodon* and *Oplurus iguanids* [13] and *Stenodactylus Petrii* and *Ptyodactylus Guttatus* [27].

de Haan [28] studied several kinds of snakes such as *D. lineatus*, *M. monspessulanus* and *R. rubropunctatus* and reported the presence of cephalic epidermal sense organs on their upper head and serve for enabling correct skin during molting.

According to Ananjeva et al. [13] epidermal sense organs are important for serving mechanical and thermoreceptors and possibly sensitive to the humidity.

Furthermore, for the first to remarks, the epidermal sense organs have deep connections with integumentary sense organs embedded in the dermal layer. Ultrastructurally, it is appeared in the form oval-shaped bundles covered by a thin coat of collagenous fibrils. It is composed of two types of neurons including Schwann and oligodendrocyte cells. Myelinated and unmyelinated nerve axons are detected.

Similar findings of light microscopic observations were reported during embryogenesis of crocodile which serve as chemo- and thermosensory transduction channels [29].

Finally, the author concluded that the juvenile chameleon skin is keratinized with obvious external and internal sensation and abundant mast cells within dermis giving characteristic immunity.
The melanosomes are dispersed within epidermis and dermis allowing the animal to maintain its color alterations according to the surrounding environment.

References

[1] Matoltsy G, Bereiter-Hahn J. Biology of the integument. Introduction. In: Bereiter-Hahn J, editor. Matoltsy of the integument. Berlin: Springer-Verlag; 1986. p. 1–7.
[2] Pough FH, Janis CM, Heiser JB. Vertebrate life. 9th ed. Boston: Pearson; 2013.
[3] Pough F, Andrews R, Cadle J, Crump M, Savitzky A, Wells K. Herpetology. Upper Saddler River, NJ, USA: Prentice-Hall; 2001.
[4] Jensen-Jarolim E. Comparative medicine: anatomy and physiology. Springer Science & Business Media; 2013.
[5] Alam AA, Daza JD, Abo Eleneen RE. Histology of the skin of three limbless snakes. J Anat 2016;299(7):979–89.
[6] Alibardi L. Adaptation to the land: The skin of reptiles in comparison to that of amphilians and endotherm amniotes. J Exp Zool B Mol Dev Evol 2003;298(1):12–41.
[7] Alibardi L, DeNardo DF. Ultrastructural and immunocytochemical features of the epidermis of the lizard Heloderma suspectum indicate richness in lipids and lack of a specialized shedding complex. Acta Zool (Stockholm) 2011;94(1):35–43.
[8] Hogan MB, Peele K, Wilson NW. Skin barrier function and its importance at the start of the atopic march. J Allergy (Cairo) 2012;2012(1):35–43.
[9] Bagnara JT. Developmental aspects of vertebrate chromatophores. Am Zool 1983;23:665–78.
[10] Maclean S. Ultrastructure of epidermal sensory receptors in Amphibolurus barbatus (Lacertilia, Agamidae). Cell Tissue Res 1980;210:435–45.
[11] Ananjeva WB, Dilmuchamedov ME, Matevysa TN. The skin sense organs of some Iguanian lizards. J Herpetol 1991;25(2):186–99.
[12] Jackson MK, Doetsch GS. Functional properties of nerve fibers innervating cutaneous corpuscles within cephalic skin of the Taxas rat snake. J Exp Neurol 1976;56:63–77.
[13] Von Düring M, Miller MR. Sensory nerve ending of the skin and deeper structures. In: Gans IC, editor. Biology of the reptilian, vol. 19. New York: Academic press; 1979. p. 407–41.
[14] Crowe-Riddell JM, Snelling EP, Watson AP, Suh AK, Partridge JC, Sanders KL. The evolution of scale sensilla in the transition from land to sea in elapid snakes. Open Biol 2016;6(6).
[15] Nissar S, Basha SH, Ramesh G, Venkatesan S, Allwin B. Gross and histomorphology of the skin of the Indian Chameleon (Chameleo zeylanicus). Ind J Vet Anat 2016;28(1):47–8.
[16] Alibardi L. Scale morphogenesis during embryonic development in the lizard Anolis lineatus. J Anat 1996;188:713–25.
[17] Alibardi L. Differentiation of the epidermis during scale formation in embryos of lizard. J Anat 1998;192:173–86.
[18] Chang C, Wu P, Baker RE, Maini PK, Alibardi L, Chuong CM. Reptile scale paradigm: Evo-Devo, pattern formation and regeneration. Int J Dev Biol 2009;53(5–6):813–26.
[19] Francois BL, Richard B. The chamelon handbook. 3rd ed. New York: Barron’s Educational Series; 2009.
[20] Teysier J, Saenko VS, Marel DV, Milinkovitch MC. Photonic crystals cause active colour change in chameleons. Nat Commun 2015;6:63–8.
[21] Trivedi NN, Caughey GH. Mast cell peptidases chameleons of innate immunity and host defense. Am J Respir Cell Mol Biol 2010;42:257–67.
[22] Alibardi L, Maurizi M, Taddei C. Immunocytochemical and electrophoretic distribution of cytokeratins in the resting stage epidermis of the lizard Podarcis sicula. J Exp Zool 2001;289(7):409–18.
[23] Di-Pui N, Milinkovitch MC. The anatomical placode in reptile scale morphogenesis indicates shared ancestry among skin appendages in amniotes. Sci Adv 2016;2:e1600708.
[24] Williams EE. A new look at the Iguania. In: Vanzolini PE, Heyer WR, editors. Proc. workshop on neotropical distribution patterns. Academy Brasil Clien Rio de Janeiro; 1988. p. 429–88.
[25] Darwish ST. Comparative light and ultrastructural studies of skin in Stenodactylus Petraii and Pteryodactylus Guttatus (Reptilia: Gekonidae). Egy J Exp Biol (Zool.) 2012;8(1):9–14.
[26] de Haan CC. Sense-organ-like parietal pits found in Psammophiini (Serpentes, Colubridae). CR Biol 2003;2003(326):287–93.
[27] Di-Pui N, Milinkovitch MC. Crocodilians evolved scattered multi-sensory micro-organs. Evo Devo 2013;4(1):19.