Research Article

Scanning Electron Microscopic Studies of the Pecten Oculi in the Quail (*Coturnix coturnix japonica*)

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The main purpose of this study is to extend the microscopic investigations of the pecten oculi in the quail in order to add some information on the unresolved functional anatomy of this unique avian organ. The pecten oculi of the quail was studied by scanning electron microscopy. Eighteen- to twenty-two highly vascularised accordion-like folds were joined apically by a heavily pigmented bridge of tissue, which holds the pecten in a fanlike shape, widest at the base. The structure of the double layered limiting membrane was recorded. The presence of hyalocytes with macrophage-like appearance was illustrated. It is assumed that the pecten oculi of the quail resembles that of the chicken. Illustrated morphological features of this species may add information on the active physiological role of the pecten. But still, the functional significance of this organ is a matter of controversies.

1. Introduction

The pecten oculi is a unique anatomical structure of the avian eye. Three types of pecten are recognized: the conical type (*pecten oculi conicus*) found only in kiwis, the vaned type (*pecten oculi vanellus*) found in the other extant Struthioniformes, and the tainamous and the pleated type (*pecten oculi plicatus*) found in all other birds [1]. The pecten has been frequently studied and extensive literature about pecten is available [2, 3]. Consequently, many investigations were carried out by light and transmission electron microscopy, regarding the structure of the pecten in various avian species (Table 1). In the great majority of the studies, light and transmission electron microscopy have been used.

The morphology of the pecten in the quail (*Coturnix coturnix japonica*) has been studied by scanning (SEM) and transmission (TEM) electron microscopy [4]. Moreover, branching of vessels in pecten oculi has also been investigated by light microscopy and stereomicroscopy [5]. However some morphological features such as the hyalocytes or the structure of the pectin-vitreal limiting membrane have not been recorded.

The present work aims to provide some unexplored morphological data of the pecten of the quail with the use of scanning electron microscopy. In addition, a concise review and discussion aim to summarize the up to date knowledge, in order to throw light on the functional structure of this organ.

2. Material and Methods

In the present study, six adult healthy quails of both sexes were used. Prior to decapitation, the animals were deeply anesthetized by inhalation of diethyl ether. The ocular bulbs were immediately removed and opened to facilitate access of the fixative used for immersion fixation. Ten of them were fixed in a sodium cacodylate buffered solution of 2% glutaraldehyde and 2% paraformaldehyde. Subsequently, pieces of the pectens were washed four times, for 15 min each time, with sodium cacodylate buffer (pH 7.2), then transferred for 1 h to 1% OsO$_4$, and dehydrated in graded acetone series. Tissues were critical point-dried in carbon dioxide (72–75 Barr), mounted onto stubs, and sputter coated with platinum and gold in a Bal-Tec sputter coater. Specimens were observed in a scanning electron microscope (JEOL, JSM 840).
Table 1: Research studies of pecten oculi in various avian species.

| Species                        | Method | Number of pleats | Order         | Family        | Visual activity | References |
|--------------------------------|--------|------------------|---------------|---------------|----------------|------------|
| Common loon (Gavia immer)      | LM, TEM| 14-15            | Gaviiformes   | Gaviidae      | Diurnal        | [15]       |
| Pigeon (Columba livia)         | LM, TEM| 15-17            | Columbiformes | Columbidae    | Diurnal        | [16]       |
| Mallard (Anas platyrhynchos)   | LM, TEM| 12-14            | Anseriformes  | Anatidae      | Diurnal        | [17]       |
| Black kite (Milvus migrans)    | SEM    | 12               | Falconiformes | Accipitridae  | Diurnal        | [18]       |
| Red-tailed hawk (Buteo jamaicensis) | LM, TEM| 17-18            | Accipitridae  | Accipitridae  | Diurnal        | [18]       |
| American crow (Corvus brachyrhynchos) | LM, TEM| 22-25            | Passeriformes | Corvidae      | Diurnal        | [19]       |
| Nighthawk (Chordeiles minor)   | LM, TEM| 4-5              | Caprimulgiformes | Caprimulgidae | Nocturnal     | [20]       |
| Emu (Dromaius novaehollandiae) | LM, TEM| 3-4              | Struthioniformes | Dromaiidae    | Diurnal        | [21]       |
| Australian galah (Eolophus roseicapillus) | LM, TEM| 20-25            | Psittaciformes | Psittacidae   | Diurnal        | [22]       |
| Great horned owl (Bubo virginianus) | LM, TEM| 7-8              | Strigiformes  | Strigidae     | Nocturnal      | [22]       |
| Chicken (Gallus domesticus)    | LM, SEM| 18–22            | Galliformes   | Phasianidae   | Diurnal        | [23]       |
| Great blue heron (Ardea herodias) | LM, TEM| 14-15            | Pelecaniformes | Ardeidae      | Diurnal        | [24]       |
| Barred owl (Strix varia)       | TEM    | 8–10             | Strigiformes  | Strigidae     | Nocturnal      | [25]       |
| Common buzzard (Buteo buteo)   | LM, SEM| 17               | Falconiformes | Accipitridae  | Diurnal        | [26]       |
| Budgerigar (Melopsittacus undulatus) | LM, SEM, TEM| 10–12            | Psittaciformes | Psittacidae   | Diurnal        | [27]       |
| Jungle Crow (Corvus macrorhynchos) | LM, TEM| 24-25            | Passeriformes | Corvidae      | Diurnal        | [28]       |
| Spotted eagle owl (Bubo bubo africanaus) | LM, TEM| 5-6              | Strigiformes  | Strigidae     | Nocturnal      | [29]       |
| White stork (Ciconia ciconia)  | LM, SEM| 15–17            | Ciconiiformes | Ciconiidae    | Diurnal        | [30]       |
| Eurasian sparrowhawk (Accipiter nisus) | LM | 11-12            | Falconiformes | Accipitridae  | Diurnal        | [31]       |
| Japanese quail (Coturnix coturnix japonica) | LM | 19              | Galliformes   | Phasianidae   | Diurnal        | [32]       |

3. Results

The pecten of the quail was a fan-like intraocular structure situated over the head of the optic nerve. It was trapezoidal shaped and black in color. Pleated type pecten oculi was comprising 18–22-folds which were joined api-cally on a fibrous bridge (Figures 1(a), 1(b), and 1(c)). The pleats were attached basally at intervals along the entire length of the linear portion of the optic disc (Figures 1(a) and 1(d)). The pectin was measured 4-5mm in length at its base, whereas the bridge measured 2.5–3mm. The mean height and width of pleats were 1mm and 100 𝜇m, respectively. A superficial membrane surrounding the pecten oculi was separating it from the vitreous body (Figure 2(a)). This structure was composed of two different laminae (Figure 2(b)). The basal lamina surrounded each blood vessel. It was attached on the outer aspect of the blood vessels and contained a large number of delicate reticular microfibrils (Figure 2(b)). The outermost had the appearance of amorphous substance (Figure 2(b)). The surface of the pleats showed a dense vascular network with branching and anastomosing vessels. Within this network extravascular spheroidal bodies were observed and identified as pigmented cells (Figure 3(a)). On the pectinal surface and especially on the pleats, amoeboid phagocyte-like cells, the hyalocytes were found (Figure 3(a)). These cells were not observed at the bridge of the pecten. They were seen isolated, double, or forming clusters both in the external and internal faces of the pleats. The hyalocytes had various shape, size, and number of cytoplasmatic processes. Their length varied from 6 𝜇m to 10 𝜇m whereas their width was 4-5 𝜇m. Their external morphology exhibited undulating membrane with filopodia (Figure 3(b)). The transverse section of the vascular network of the pecten revealed the profile of numerous capillaries with diameter ranging from 5 to 10 𝜇m. Among them, each fold contained two or more larger blood vessels, which were difficult to differentiate as either arterioles or venules (Figure 4). Their diameter frequently exceeded the 20 𝜇m.

4. Discussion

The pecten oculi of the quail (Coturnix coturnix japonica), in concert with other avian species, is a pigmented highly vascular, fanlike structure situated over the head of the optic nerve and projecting out into the vitreous. My observations
Figure 1: (a) Scanning electron micrograph of the pecten’s overview. B: the base of the pecten; b: the bridge of the pecten; arrow: the vitreopectineal membrane; the asterisks denote the pleats. (b) Scanning electron micrograph of three pleats ending at the bridge of the pecten. The asterisk shows the bridge. Thin arrows: pigmented cells. (c) Scanning electron micrograph of the fibrillar tissue of the bridge. (d) Scanning electron micrograph of a pleat arising from the base. Note the presence of hyalocytes (arrows).

Figure 2: (a) Scanning electron micrograph of the vitreopectineal limiting membrane (arrow). (b) Scanning electron micrograph of the vitreopectineal limiting membrane. Note the outer layer (left side) and the inner layer (right side).

are in agreement with previous studies in the quail [4, 5]. The morphometrical data of this study are in the same range with that of Orhan et al. [5]. The general morphology of the quail’s pecten resembles that of chicken [6–8].

The limiting membrane which covers the pecten has been illustrated for the first time in terms of SEM. Besides the numerous morphological investigations of the avian pecten, some of them have illustrated the limiting membrane by means of TEM [7, 9]. Our observations are similar to reports of chicken [7] and pigeons [9]. The authors demonstrated the double layered limiting membrane by TEM. The double layer constitutes a barrier between the pecten and the vitreous body.

The presence of hyalocytes on the pectineal surface of the quail has been demonstrated for the first time by means of SEM, in the present study. Previous investigations in the developing quail revealed the phagocytic activity of the hyalocytes [10]. These specialized cells have also been observed
in the chicken [8], the mallard [11], and the budgerigar [12]. In contrary, they have not been observed in many other avian species [2, 9, 13]. Fischlschweiger and O’Rahilly [7] described with means of TEM the presence of cells on the external aspect of the covering membrane. These cells, termed “pericellular cells,” were thought by the authors to be a special type of astroglia. As these cells display phagocytosis and from their histochemistry, it has been concluded that the hyalocytes are a subtype of macrophage [8]. My observations agree with the positive relationship between the distribution of hyalocytes and blood vessels in the chicken [8]. In fact, the hyalocytes were numerous on the free surface of the pleats excluding the bridge in which the relatively fewer vessels were located in the deep portion. Llombart et al. [10] proposed that hyalocytes might prevent the movement of macromolecules into the retina by their macrophagic activity.

The presence of pigmented cells is a constant feature of all pectens recorded to date. In all species described, there is a marked increase in melanocytes in the apex or bridge region of the pecten [2]. Overlying and within the vascular network, a close association between blood vessels and pigmented cells is evident. This association may serve as a protective mechanism of the vessels against the effects of ultraviolet light [12, 14].

Fine structural studies to date indicate that while the pleated pecten was widespread and essentially similar in most species, there were some variations in such parameters as shape, size, number of folds, and the thickness of capillary basal laminae [15]. The number of pleats varies among the species which possess a pleated type pecten (Table 1). SEM studies showed the approximate number of pleats was 18–22 in the chicken [6, 23]. In the present study the number ranged from 18 to 22 similar to chicken and to quail [4, 5]. Many attempts have been made to correlate the number of pleats with the function of the pecten. From a survey of the existing literature (Table 1) no correlation could be found regarding the number and the zoological classification of the birds. The number of pleats in the quail is similar to the fowl [6, 23]. Both of them have a zoological closeness since they belong to the same family of Phasianidae. On the other hand, the budgerigar [12] and the Australian galah [21] which belong to the family of Psittacidae have 10–12 and 20–25 pleats, respectively. The size of the pecten and the number of pleats do not appear to relate directly to the size of the eye itself. On the other hand, the size of the pecten and the number of pleats seem to pertain to the behavior of the bird in relation to its general activity and its environmental lighting. Furthermore, these variations seem to correlate with the diurnal activity and/or visual requirements of the species (Table 1). The same deduction has been recorded by Pettigrew [31] who compared the number of pectineal pleats of 8 diurnal and 8 nocturnal birds. However this correlation may not be evidential; for example, the emu has a pecten with 3–4 pleats [20].

Despite the numerous publications, the functional significance of the pecten remains controversial. It is more than a century ago when the function of the pecten has challenged the imagination of investigators. The pecten according to numerous authors is mainly concerned with the nourishment of the avascular avian retina [17, 20]. The intriguingly dense meshwork of capillaries points to erectile tissue of the pecten which potentially could serve as sun shade [27]. The double
limiting membrane provides the mechanical support of the pecten and a barrier between the pecten capillaries and the hyaloid body. In the same capillaries, blood which is circulated, could serve as carrier of various sensory molecules. The hyalocytes probably serve as sweepers of unwanted material within the hyaloid body [8]. Frost and Mourtisen [32] claimed that the pecten could possibly function as an ‘ocular sextant.’ Maybe the pecten could potentially be engaged in avian navigation.

5. Conclusions

The purpose of the present study was to give some additional information on the morphology of the avian pecten and to explore possible relations between the structure and function of this organ. The appearance of the hyalocytes, as well as the illustration of the bilaminar structure of the covering membrane, was demonstrated for the first time. These morphological characteristics add information on the morphology of the avian pecten and as well as the illustration of the bilaminar structure of the covering membrane. However, there is need for a further approach to throw light in this enigmatic organ.

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References

[1] H. E. Evans and G. R. Martin, “Organa sensuum,” in Handboook of Avian Anatomy-Nomina Anatomica Avium, J. J. Baumel, A. S. King, J. E. Breazile, H. E. Evans, and J. C. Vanden Berge, Eds., pp. 585–611, Publications of Nuttal Ornithological Club, 1993.
[2] V. Brach, “The functional significance of the avian pecten: a review,” Condor, vol. 79, no. 3, pp. 321–327, 1977.
[3] H. Wolburg, S. Liebner, A. Reichenbach, and H. Gerhardt, “The pecten oculi of the chicken: a model system for vascular differentiation and barrier maturation,” International Review of Cytology, vol. 187, pp. 111–159, 1999.
[4] A. S. Ramos and M. D. Tingari, “A correlative scanning and transmission electron microscope study of the pecten oculi of Japanese quail,” Canadian Journal of Zoology, vol. 62, no. 12, pp. 2457–2464, 1984.
[5] I. ¨O. Orhan, O. Ekim, and A. G. Bayraktaroğlu, “Morphological investigation of the pecten oculi in quail (Coturnix coturnix japonica),” Veterinary Journal of Ankara University, vol. 58, no. 1, pp. 5–10, 2011.
[6] M. Uehara, S. Oomori, H. Kitagawa, and T. Ueshima, “The development of the pecten oculi in the chick,” Japanese Journal of Veterinary Science, vol. 52, no. 3, pp. 503–512, 1990.
[7] W. Fischlschweiger and R. O’Rahilly, “The ultrastructure of the pecten oculi in the chick,” Acta Anatomica, vol. 65, no. 4, pp. 561–578, 1966.
[8] M. Uehara, T. Imagawa, and H. Kitagawa, “Morphological studies of the hyalocytes in the chicken eye: scanning electron microscopy and inflammatory response after the intravitreous injection of carbon particles,” Journal of Anatomy, vol. 188, no. 3, pp. 661–669, 1996.
[9] E. Raviola and G. Raviola, “A light and electron microscopic study of the pecten of the pigeon eye,” American Journal of Anatomy, vol. 120, no. 3, pp. 427–462, 1967.
[10] C. Llombart, V. Nacher, D. Ramos et al., “Morphological characterization of pecteneal hyalocytes in the developing quail retina,” Journal of Anatomy, vol. 215, no. 3, pp. 280–291, 2009.
[11] C. R. Braekevelt, “Fine structure of the pecten oculi of the mallard (Anas platyrhynchos),” Canadian Journal of Zoology, vol. 68, no. 3, pp. 427–432, 1990.
[12] A. Micali, A. Pisani, C. Ventrici et al., “Morphological and morphometric study of the pecten oculi in the budgerigar (Melopsittacus undulatus),” Anatomical Record, vol. 295, no. 3, pp. 540–550, 2012.
[13] C. R. Braekevelt, “Electron microscopic observations on the pecten of the Nighthawk (Chordeiles minor),” Ophthalmologica, vol. 189, no. 4, pp. 211–220, 1984.
[14] S. G. Kiama, J. Bhattacharjee, J. N. Maina, and K. D. Weyrauch, “A scanning electron microscope study of the pecten oculi of the black kite (Milvus migrans): possible involvement of melanosomes in protecting the pecten against damage by ultraviolet light,” Journal of Anatomy, vol. 185, no. 3, pp. 637–642, 1994.
[15] C. R. Braekevelt, “Fine structure of the pecten oculi of the common loon (Gavia immer),” Canadian Journal of Zoology, vol. 64, no. 10, pp. 2181–2186, 1986.
[16] C. R. Braekevelt, “Fine structure of the pecten of the pigeon (Columba livia),” Ophthalmologica, vol. 196, no. 3, pp. 151–159, 1988.
[17] L. de Schaepdrijver, P. Simoens, H. Lauwers, and J. P. de Geest, “Retinal vascular patterns in domestic animals,” Research in Veterinary Science, vol. 47, no. 1, pp. 34–42, 1989.
[18] C. R. Braekevelt, “Fine structure of the pecten oculi of the red-tailed hawk (Buteo jamaicensis),” Anatomia, Histologia, Embryologia, vol. 20, no. 4, pp. 354–362, 1991.
[19] C. R. Braekevelt, “Fine structure of the pecten oculi in the American crow (Corvus brachyrhynchos),” Anatomia, Histologia, Embryologia, vol. 23, no. 4, pp. 357–366, 1994.
[20] C. R. Braekevelt, “Fine structure of the pecten oculi of the emu (Dromaius novachollandiae),” Tissue and Cell, vol. 30, no. 2, pp. 157–165, 1998.
[21] C. R. Braekevelt and K. C. Richardson, “Fine structure of the pecten oculi in the Australian galah (Eolophus roseicapillus) (Aves),” Histology and Histopathology, vol. 11, no. 3, pp. 565–571, 1996.
[22] C. R. Braekevelt, “Fine structure of the pecten oculi in the great horned owl (Bubo virginianus),” Histology and Histopathology, vol. 8, no. 1, pp. 9–15, 1993.
[23] T. Amemiya, “A transmission and scanning electron microscopic study of the pecten of the chick eye,” Japanese Journal of Ophthalmology, vol. 84, no. 4, pp. 295–303, 1980.
[24] S. G. Kiama, J. N. Maina, J. Bhattacharjee, and K. D. Weyrauch, “Functional morphology of the pecten oculi in the nocturnal spotted eagle owl (Bubo bubo africanus),” and the diurnal black kite (Milvus migrans) and domestic fowl (Gallus gallus var. domesticus): a comparative study,” Journal of Zoology, vol. 254, no. 4, pp. 521–528, 2001.
[25] C. R. Braekevelt, “Electron microscopic observations on the pecten of the great blue heron (Ardea Herodias),” Histology and Histopathology, vol. 6, no. 3, pp. 345–351, 1991.
[26] B. J. Smith, S. A. Smith, and C. R. Braekevelt, “Fine structure of the pecten oculi of the barred owl (Strix varia),” Histology and Histopathology, vol. 11, no. 1, pp. 89–96, 1996.

[27] M. E. Gültiken, D. Yildiz, B. Onuk et al., “The morphology of the pecten oculi in the common buzzard (Buteo buteo),” Veterinary Ophthalmology, vol. 15, no. supplement 2, pp. 72–76, 2012.

[28] M. L. Rahman, E. Lee, M. Aoyama, and S. Sugita, “Light and electron microscopy study of the pecten oculi of the Jungle Crow (Corvus macrorhynchos),” Okajimas Folia Anatomica Japonica, vol. 87, no. 3, pp. 75–83, 2010.

[29] B. Onuk, S. Tutuncu, A. Alam, M. Kabak, and N. G. Ince, “Macroanatomic, light and scanning electron microscopic studies of the pecten oculi in the stork (Ciconia ciconia),” Microscopy Research and Technique, vol. 76, no. 9, pp. 963–967, 2013.

[30] J. M. Rajab, “Morphological and histological description of the pecten oculi in the Sparrow Hawk (Accipiter nisus),” Diyala Journal of Pure Sciences, vol. 8, no. 1, pp. 8–19, 2012.

[31] J. D. Pettigrew, “A note on the eyes of the Letter winged Kite (Elanus scriptus),” Emu, vol. 82, pp. S305–S308, 1983.

[32] B. J. Frost and H. Mouritsen, “The neural mechanisms of long distance animal navigation,” Current Opinion in Neurobiology, vol. 16, no. 4, pp. 481–488, 2006.