AGE CHANGES OF THE LACRIMAL GLAND OF RABBITS (LIGHT AND ELECTRON MICROSCOPIC STUDIES)

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ABSTRACT
The dry eye in senile people is a major ophthalmologic problem; the aim of this work was to study the histological and histochemical changes with advancement of age in the lacrimal glands in rabbits, and correlating these changes with those, which might be detected at the ultra structural level.

Thirty healthy rabbits of both sexes were divided into three groups. Young age group (3-5 months), adult group (9-12 months), and senile group (24-36 months). The lacrimal glands, of each animal, were dissected out, isolated and processed; paraffin section stained with Hx&E., Mallory Trichrome and PAS stains. Fresh frozen cryocut section for localization of acid phosphatase enzyme activity were done. Epon embedded semithin sections stained with toluidine blue and ultra thin sections for Electone Microscopic study were prepared.

The rabbits has a dorsal lacrimal gland in the postero dorsal region of the eye ball, it is about 4mm in diameter, and Harderian gland lies in the antero ventral region of the eye ball, it is about 6mm in diameter. In young and adult age consists of tubulo acinar units separated by dense sheets

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of connective tissue septae into lobules. The acini lined by cubical cells, rounded basal nuclei and basophilic cytoplasm. In senile rabbits, the lacrimal gland showed thickening of the periductal and perivascular connective tissue infiltration of inflammatory cells and degeneration of some lacrimal acini.

The hardierian gland in young and adult rabbits consists of of tubulo acinar unit, divided into lobules by sheets of connective tissue. The acini composed of two types (type I and type II). type I had a narrow lumen, and their cells showed small vacuoles and rounded basal nuclei. The type II acini had a wider lumen, larger vacuoles, and rounded basal nuclei. In senile group the acini were closer, irregular in shape, showing outpocketing of the acinar cells. Perivascular and periductal cellular infiltration was noticed.

The lacrimal acinar cells in young and adult rabbits showed positive PAS reaction while they showed a weak positive reaction in senile ones. The acinar and ductal basement membrane of the acini and ducts were thickened and showed strong positive PAS reaction in senile age groups.

The lacrimal gland showed an increase in the amount of the collagenous fibers in adult animal and more condensation of these fibers were noticed around ducts and blood vessels in senile gland.

The basal portions of the acinar cells, in young age showed strong positive acid phosphatase reaction while the remaining cytoplasm showed moderate positive reaction. The senile gland showed weak reaction in the acinar cells. The epithelial lining of the ducts in all age group showed strong positive acid phosphatase reaction.

By E/M Some of the lacrimal acinar cells appeared darker than the others. The acinar cells showed supra nuclear Golgi complex, mitochondria and numerous secretory granules. The gland showed increase of the mitochondria in young age, while in senile rabbits signs of degeneration of the mitochondria in the form of pyknosis of the nuclei, and appearance vacuoles in the cytoplasm was noticed.
Review of Literature

The aqueous component of the tear film is responsible for keeping the cornea buffered, lubricated, nourished and protected. It is produced and secreted from the lacrimal gland (Dilly, 1994). Dry eye in senile people is a major problem present in some diseases such as kerato-conjunctivitis sicca and Sjogren's syndrome. (Farrell J, Patel S, Grierson DG, and Sturrock RD, 2003).

Sullivan and Krenzer, (1999) reported that Precorneal tear film is a sheet of tears, which cover the exposed inter palpebral portions of the globe and cornea.

Dry eye syndrome such as, keratoconjunctivitis sicca, Sjogren's syndrome and dacry adenitis are big problem in old age. Most dry eye symptoms result from an abnormal, non-lubricative ocular surface that increases shear forces under the eye lids, and diminishes the ability of the ocular surface to respond to environmental challenges. Michael, E. Sterm, ph, Roger, W and Beuerman, (1998).

Clao, (2000) included that the ocular surface of the eye and lacrimal gland function are tightly integrated unit. Dry eye conditions; damage the ocular surface, lead to further damage of lacrimal gland.

Sullivan, D.A, wickham LA, and Krenzer, (1997) and Azzarolo. et al, (1997), stated that the lacrimal gland function was significantly influenced by sex hormone. Androgens were showing to exert essential and specific effect on maintaining normal glandular function to suppress inflammation in normal and autoimmune animal models. Sullivan and Krenzer, (1999), included that androgen deficiency may promote the progression of Sjogren's syndrome and its associated lacrimal gland inflammation, Meibomian gland dysfunction, and severe dry eye.

Azzarolo, (2003), stated that lacrimal glandular atrophy was observed after ovariectomy, he suggested that in addition to androgen, estrogen also seem to play a role to maintain lacrimal gland structure and function.

Cornell-Bell, DA Sullivan, and MR Allansmith, (1985), stated that acinar
area in lacrimal glands of males was larger than that of females. These findings suggest that gender differences in lacrimal gland morphology may be a general phenomenon in a variety of species.

*Evaporation of tears:*

The evaporation rate is normally low because of protective oily layer; no more than 20-25% of the total tears secreted are lost by evaporation per day, remaining fluid is drained through the nasolacrimal duct into the nose. In the absence of the protective oily layer, the rate of evaporation is increased 10 to 20 times. The toxicity of human tears is subject to a dynamic change because of the evaporation process and rate of tear flow. The overall thickness of the tear film is about 4-9μm but the eye can manage up to 30mm without over flow. (Peiffer, R.L and Petersen Jones, 1997).

Most dry-eye symptoms result from an abnormal, non lubricative ocular surface that increases tear forces under eyelids and diminishes the ability of the ocular surface to respond to environmental changes. This ocular surface dysfunctions may result from immuno-compromise persons due to systemic autoimmune disease, locally from a decrease in systemic androgen support to the lacrimal gland as seen in aging and most frequently in the menopausal female. (Stem, 1998).

The function of the lacrimal gland is significantly influenced by sex hormone (androgen). Androgen is shown to exert essential and specific effects on maintaining normal glandular functions and to suppress inflammation in normal and autoimmune animal models. As the activity of androgen in lacrimal gland is proposed to be attributed to its ability to induce the accumulation of anti-inflammatory cytokines. (Azzarolo, AM, Olsen, E and Huang, ZM, 1997).

Androgen deficiency may promote the progression of Sjogren’s syndrome and its associated lacrimal gland inflammation, Meibomian gland dysfunction and severe dryness of the eye. (Sullivan and Krenzer, 1999).

Ovariectomy result in the decrease of the level of estrogen which
lead to necrotic change in the acinar cell and inflammatory response of the lacrimal gland by lymphocytic infiltration. This suggests that estrogen play a role to maintain lacrimal gland structure and function. (Elhusseni, H. M, 2003).

Keratoconjunctivitis sicca and dry eye is associated with reduced tear secretion and therefore a reduction in the effectiveness of lacrimal gland function. This have been seen to occur with increasing age. The difference between the young and aged rat glands include deposition of connective tissue, inflammatory cell infiltration, periductal fibrosis, acinar atrophy. Also morphological change of the acini which occur with age from serous to seromucous and then to mucous acini, resulting in a reduction in the presence of protein secretory granules. The mast cell increases with tissue damage. This explains the phenomenon of reduced tear secretion with ageing. (Draper, et al 1997).

Lacrimal gland is the primary tissue involved in the secretory immune system of the eye. This system serves as the first line of defense to protect the ocular surface against microbial agents and toxic compounds. So, it significantly limits the eye susceptibility to infection and allergic diseases and help to maintain both corneal and conjunctival integrity and visual function. (Sullivan and Sato, 1999).

Certain disease, such as Sjogren's syndrome (SS) results in secretory dysfunction of the lacrimal gland. There are progressive lymphatic proliferations which correlates with loss of secretory acini. (Yoshino; et.al 1996).

Decrease or even prevention of tear drainage by lacrimal occlusion is of great clinical benefit for patients suffering from dry eye syndrome. Itching, burning, foreign body sensation and light sensitivity are markedly improved following progressive lacrimal passage occlusion. (Navacastaneda; et al 2003).

Material and methods: Thirty healthy rabbits of both sexes were divided into three groups. Young age group (3-5 months), adult group (9-12...
months), and senile group (24-36 months). The animal were killed by decapitation and lacrimal glands, of each animal, were dissected out, isolated and processed; paraffin sections stained with Hx&E., Mallory Trichrome stains and PAS reaction. Fresh frozen cryocut sections prepared for localization of acid phosphatase enzyme activity. Epon embedded semithin section stained with toluidine blue and ultra thin sections prepared for Electrone Microscopic study.

RESULTS

Light microscope study:

1) In the first group (young age, 3-5 months) of rabbits.

Dorsal lacrimal gland:

The dorsal lacrimal gland appeared to be consisted of small serous acini with narrow lumen, acinar cells were pyramidal in shape with rounded basal nuclei and granular basophilic cytoplasm (Fig.1). Interlobular connective tissue contained ducts and blood vessels. The ducts had wide lumen rounded basal nuclei. (Fig.2).

A minimal amount of positively stained collagenous fibers were found mainly in the interlobular connective tissue around ducts and blood vessels. (Fig 3).

Strong positive PAS reactions was observed in the basement membrane of acini, duct cells, and cytoplasm of acinar cells (Fig 4).

Some acinar cells showed strong positive acid phosphatase reaction. and negative reaction in some interlobular and periacinar connective tissue (Fig 5).

Semithin sections of the dorsal lacrimal gland, showed acinar secretory cells were packed with secretory granules. some of the acini was stained darker than the others (Fig, 6).

Harderian gland; The acini of the harderian gland were composed of two types of cells (type I and type II). In type I cells the acini had a narrow lumen, their cells showed small sized vacuoles, and rounded basal nuclei. Type II cells had a wider lumen, their cells had larger vacuoles, rounded basal nuclei and less amount of cytoplasm (Fig 7). The interlobular con-
nective tissue contained ducts and blood vessels. The duct cells had central rounded nuclei (Fig 8). It showed moderate amount of positively stained collagenous fibers mainly in the interlobular connective tissue around the duct and blood vessels in type I acini (Figs. 9). Type II acini showed mild amount of collagenous fibers around the duct and blood vessels and negative collagenous fibers in the interlobular connective tissue (Figs. 10).

The gland showed moderate positive PAS reaction in the basement membrane of acini, and cytoplasm of acinar cells (Fig 11).

Strong positive acid phosphatase reaction in some acinar cells and negative reaction in some interlobular and peri-acininar connective tissue were observed (Fig, 12).

2) In the second group (adult age, 9-12 months) of rabbits.

Dorsal lacrimal gland;
The lacrimal acini were apparently large in size they were lined by columnar cells with rounded basal nuclei and basophilic granular cytoplasm. Myoepithelial cells surrounding the acini (Fig 13).

Moderate amounts of positively stained collagenous fibers were laid around intra and interlobular connective tissue mainly around fibrous capsule, ducts and blood vessels (Fig. 14).

The positive PAS reaction were seen in the peri acinar, interlobular connective tissue and acinar basement membrane with negative reaction in the acinar cells. (Fig 15).

Harderian gland;
Acini were apparently large in size, they were lined by columnar cells, and rounded basal nuclei and some acini lined by cuboidal cells with flattened basal nuclei. Interlobular duct has large lumen, lined by columnar cells with rounded nuclei. Myoepithelial cells surrounding the acini (Fig 16).

The moderate amounts of positively stained collagenous fibers were seen lie around intra and interlobular connective tissue mainly around fi-
brous capsule, duct and blood vessels (Figs.17).

A strong positive PAS reaction is seen in the peri acinar, interlobular connective tissue and acinar basement membrane and moderate positive reaction in the acinar cell (Fig 18).

3) In the third group (senile age, 2-3 years) of rabbits.

Dorsal lacrimal gland;

Acini were large in size, and had wide irregular lumen. Some acini showed outpocketing of the glandular cells and some show cellular degeneration. Acinar cells were cuboidal with rounded basal nuclei (Fig19). The ducts were intralobular inside the lobules, and interlobular between the lobules in interlobular septae. The cells of the ducts were cuboidal, with central nuclei. Myoepithelial cells surrounding the acini, and cellular infiltration was noticed especially around the duct (Fig.20).

There were more deposition of collagenous fibers showed in periductal, perivascular and peri acinar connective tissue (Figs. 21).

A strong positive PAS reaction were seen in the acinar basement membrane, weak reaction in the cytoplasm of acinar cells (Fig22).

A strong positive Acid phosphatase reaction showed in some cells lining the lacrimal duct. Moderate reaction in some interlobular and peri-acinar connective tissue and weak reaction in the acinar cells (Fig.23).

Semithin sections of dorsal lacrimal gland, stained with toluidine blue showed acini separated by connective tissue septae containing blood vessels. Acinar cells were cuboidal surrounded by basement membrane, and myoepithelial cells. Secretory vacuoles were present in the apical part of the acinar cells, where the rounded nuclei of the cells were basal in position (Fig24). Interlobular ducts were seen, and also showing some secretory vacuoles (Fig 25).

Harderian gland:

The acini were closer, irregular in shape, some showing outpocketing of the acinar cells. Acinar cells were cuboidal with basal nuclei, cellular degeneration of the acini also noticed.
(Figs. 26, 27). The duct had cuboidal cells with basal nuclei, perivascular and periductal cellular infiltrations were noticed (Fig 27). Myoepithelial cells were noticed surrounding the acini (Fig 26).

More deposition of collagenous fibers was noticed around ducts and blood vessels and minimal amount of collagen fibers in the inter and intralobular connective tissue in cells with large vacuoles, more deposition of collagen fiber in the inter and intralobular connective tissue cells with small vacuoles (Fig. 28).

A strong positive PAS reaction was observed in the ductal cells, ductal basement membrane, moderately positive PAS reaction in acinar basement membrane, perivascular and interlobular connective tissue and negative reaction in the acinar cell (Fig 29).

Mild acid phosphatase reaction was seen in some acinus cell, and weak reaction in some interlobular and periacinar connective tissue (Fig 30).

Electron microscopic result:
In the lacrimal gland of the young age rabbits, the lacrimal acini contained acinar cells filled with secretory granules, rounded or oval basal nuclei. Most of those granules had granular contents, while few of them had homogenous electron dense contents. Supranuclear Golgi complex and mitochondria were apparent. (Figs. 31, 32). The lacrimal gland of senile rabbits show some intact acinar cells, while others had ill defined cell boundaries and the nuclei of some of them were pyknotic, nuclei have high content of condensed chromatin. The cytoplasm contained coalescent secretory granules, the secretory granules might be fused together which had dark or light varieties. It also contained cytoplasmic vacuoles, Supranuclear Golgi complex, mitochondria and vacuoles. (Figs 33, 34)
Figure 1: A photomicrograph of a section in the dorsal lacrimal gland of three month rabbit showing glandular acini (A), with narrow lumen (L), the nuclei are rounded and lies in the basal portion of the cell (N). Acini separated by interlobular connective tissue (T). (Hx&E X 200)

Figure 2: A photomicrograph of a paraffin section in the dorsal lacrimal gland of three month rabbit showing interlobular duct (D), with wide lumen (L), with rounded basal nuclei (N). (Hx&E X 400)

Figure 3: A photomicrograph of a paraffin section in the dorsal lacrimal gland of three month rabbit showing collagen fiber (F) around interlobular connective tissue (T) mainly around interlobular duct (D) and blood vessels (V). (Mallory Trichrome X 100)

Figure 4: A photomicrograph of a paraffin section in the dorsal lacrimal gland of three month rabbit showing strong positive PAS reaction in ductal cells (s), acinar basement membrane (s), and cytoplasm of acinar cells (S). (PAS X400)

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Figure 5: A photomicrograph of frozen cryocut section in the dorsal lacrimal gland of three month rabbit showing positive acid phosphatase reaction in the acinar cell (S). And negative reaction in the interlobular and periacinar connective tissue (N). (Acid phosphatase X 200)

Figure 6: A photomicrograph of a semitin section in the lacrimal gland of three month rabbit showing secretory acini, they have narrow lumen (L) and their cytoplasm are filled with secretory granules (G) some of the acinar cells appear darker (D) than the others (P). (Toluidine blue, X1000)

Figure 7: A photomicrograph of section in the Harderian gland of three month rabbit showing different size and shape of acini (A). type I (I) with narrow lumen (L), rounded basal nuclei (n) and type II (II) with wide lumen (L) rounded basal nuclei (n). (Hx&E X 200)

Figure 8: A photomicrograph of section in the Harderian gland of three month rabbit showing acinar cells with large vacuole (V) interlobular duct (D), duct cells have central rounded nuclei (n) and blood vessels (V). (Hx&E X 400)

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Figure 9: A photomicrograph of section in the Harderian gland of three month rabbit showing collagen fiber (F) around interlobular connective tissue (T) mainly around interlobular duct (D) in acini with small vacuole. (Mallory Trichrome X 200)

Figure 10: A photomicrograph of section in the Harderian gland of three month rabbit showing mild collagen fiber (f) around ducts (d), in acini (A) with wide lumen (L) and negative collagen fiber around interlobular connective tissue. (Mallory Trichrome X 200)

Figure 11: A photomicrograph of a paraffin section in the Harderian gland of three months rabbit showing moderate positive PAS reaction in the acinar basement membrane (M), and acinar cells (m). (PAS X400)

Figure 12: A photomicrograph of frozen cryocut section in the Harderian gland of three month rabbit showing strong positive acid phosphatase reaction in some acinar cell (S), and negative reaction in some interlobular and periacinar connective tissue (N). (Acid phosphatase X 200)
Figure 13: A photomicrograph of section in the dorsal lacrimal gland of rabbit aged one year showing lacrimal acini, with different lumen (L), they are lined by columnar cells (cl) with rounded nuclei positioned in the basal portion of the cell (N), interlobular Blood vessels (V). Myoepithelial cells surrounding the acini (my), as flattened nuclei (n). (Hx&E X 400)

Figure 14: A photomicrograph of section in the dorsal lacrimal gland of rabbit aged one year showing moderate amount of collagen fiber (F) in intra and interlobular connective tissue. And around blood vessels (V). (Mallory Trichrome X 400)

Figure 15: A photomicrograph of section in the dorsal lacrimal gland of rabbit aged one year showing positive PAS reaction in the peri acinar, interlobular connective tissue, acinar basement membrane (P), and negative reaction in the acinar cells (N). (PAS X400)

Figure 16: A photomicrograph of section in the Harderian gland of of rabbit aged one year showing large acini variable in size (A), interlobular duct (D) with central rounded nuclei (n), and large lumen (L). Myoepithelial cells surrounding the acini (my), as flattened nucleus (n). (Hx&E X 400)

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Figure 17: A photomicrograph of section in the Harderian gland of of rabbit aged one year showing moderate amount of collagen fiber (F) condensed around interlobular duct (D). (Mallory Trichrome X 200)

Figure 18: A photomicrograph of section in the Harderian gland of of rabbit aged one year showing strong positive PAS reaction in the peri acinar, interlobular connective tissue and acinar basement membrane (S).and moderate positive reaction in the acinar cell (M). (PAS X 400)

Figure 19: A photomicrograph of section in the dorsal lacrimal gland of rabbit aged three years showing glandular lobules containing large acini(A),irregular lumen(L),outpocketing of acinar cells(p), some of them appear degenerated(d).acinar cell cuboidal in shape(c)with basal nuclei(N). (Hx&E X 400)

Figure 20: A photomicrograph of section in the dorsal lacrimal gland of rabbit aged three years showing interlobular duct (D), myoepithelial cell with flattened nuclei(N). (Hx&E X 400)

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Figure 21: A photomicrograph of section in the dorsal lacrimal gland of rabbit aged three years showing condensed amount of collagen fiber (F) mainly around intralobular duct and periacinar connective tissue. (Mallory Trichrome X 400)

Figure 22: A photomicrograph of section in the dorsal lacrimal gland of rabbit aged three years showing strong positive PAS reaction in (S) acinar basement membrane (b), and myoepithelial cells (my). And weak reaction in the acinar cell (W). (PAS X400)

Figure 23: A photomicrograph of section in the dorsal lacrimal gland of rabbit aged three years showing strong positive acid phosphatase reaction in the cell lining lacrimal main duct (S). (Acid phosphatase X 400)

Figure 24: A photomicrograph of a semi thin section in the lacrimal gland showing secretory acini, they have rounded basal nuclei (N) their cytoplasm is filled with secretory vacuoles (s), acini surrounded by basement membrane, nucleus of myoepithelial cell were noticed (m), connective tissue septae contain blood vessels (v). (To-luidine blue, X1000)

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Figure 25: A photomicrograph of a semitint section in the lacrimal gland showing secretory acini, they have wide lumens (L) with rounded basal nuclei (N) their cytoplasm is filled with secretory vacuoles (s). Connective tissue septae contain interlobular duct (D) with basal nuclei (N) and blood vessels (v). (Toluidine blue, X1000)

Figure 26: A photomicrograph of a paraffin section in the Harderian gland of three years in rabbit showing large acini (A), acinar cells cuboidal in shape (c) with apical vacuole (a) and basal nuclei (N). Myoepithelial cell (m) with flattened nuclei (N). (Hx&E X 400)

Figure 27: A photomicrograph of a paraffin section in the Harderian gland of rabbit aged three years showing, Blood vessels (V), interlobular duct (D) had cuboidal cell (c), basal nuclei (N), acini (A) show cellular degeneration (d). Cellular infiltration (i) was noticed around duct and blood vessels (Hx&E X 400).

Figure 28: A photomicrograph of a paraffin section in the Harderian gland of rabbit aged three years showing condensed amount of collagen fiber (F) mainly around interlobular connective tissue. (Mallory Trichrome X 400)
Figure 29: A photomicrograph of a paraffin section in the Harderian gland of rabbit aged two years showing positive PAS reaction in acinar, basement membrane, perivascularular and interlobular connective tissue (P). Negative reaction in the acinar cell (N). (PAS X400)

Figure 30: A photomicrograph of frozen cryocut section in the Harderian gland of rabbit aged three years showing mild positive acid phosphatase reaction in some acinus cell (m) and weak reaction in some interlobular and periacinar connective tissue (W). (Acid phosphatase X 400)

Figure 31: An electron micrograph of an acinar cell of young age rabbit lacrimal acinus. showing oval nuclei (N), mitochondria (M). And secretory vacuole also present (S). (Uranyl acetate and Lead citrate X6610)

Figure 32: An electron micrograph of an acinar cell of young age rabbit lacrimal acinus. showing rounded nuclei (N), Supranuclear Golgi complex (G), mitochondria (M), their cytoplasm is filled with a homogenous electron dens material (H). (Uranyl acetate and Lead citrate X5200)

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Figure 33: An electron micrograph of an acinar cell of senile age rabbit lacrimal acinus. It part of acinus is intact, its cell boundaries are defined, with rounded nuclei (N). The other part of acinus show ill defined cell boundaries and pyknotic nuclei (P). (Uranyl acetate and Lead citrate X 6610)

Figure 34: An electron micrograph of an acinar cell of senile rabbit aged lacrimal acinus showing nucleus N). Some mitochondria appear intact (M), and vacuoles are noticed (V). (Uranyl acetate and Lead citrate X 8900)

DISCUSSION

In the present study, the acini were enlarged and dilated with the progress of age. In younger age the acini were small serous acini with narrow lumen and acinar cell pyramidal in shape with rounded basal nuclei. In senile group the acini enlarge in size, with wide irregular lumen without pocketing of glandular cells, some acini show cellular degeneration, and cellular infiltration. The acinar cells were cuboidal with rounded basal nuclei. Some of these acinar cells, on using toluidine blue stain, appeared darker than the others. These dark cells may contain in their secretory granules, more acidic and neutral glycoprotein than the pale cells. These contents also might cause the dark cells to attain a strong positive PAS reaction while the other pale ones attained a moderate positive reaction. These findings were in agreement with Ahmed (1983), who described two types of cells in the rabbit lacri-
mal acini; type I cells appeared pale while type II cells were dark. Allen, M, Wright, p. and Bron. et al. (1997) also reported that there were two types of cells lining the acini of the human lacrimal gland, one type was serous and the other was mucous. They added that the serous cells contained lyso- somes and were similar to paneth cells. On the other hand, Kalleny, (1995), described only one type of cells in the acini of lacrimal gland in the rat. This controversy might be related to species differences.

Draper. et al, (1997) in their study of age related morphological changes of male rat lacrimal gland, stated that the differences between young and aged rat glands included deposition of connective tissue, inflammatory cell infiltration, periductal fibrosis, acinar atrophy and reduced innervations. They estimated the percentage distribution of the acinar type of gland in different age groups, in young gland the majority of acini were serous (81%), few seromucous acini (17%), and pure mucous acini (2%). In aged gland their were significantly reduction in serous acini over all distribution was 23% when compared with young gland and marked increase in presence of seromucous acini over all distribution 53%.

In the present study, myoepithelial cells were seen around the secretory lacrimal acini. This in agreement with similar cells were noticed in lacrimal acini of one humped camels, Mongolian gerbils, mice and albino rats Prince, J.H, Diesem, C.D, Eglitis, I, Fahmy, M. F, Shain, Y.M, Sakai, yohro, (1981) Johnston:et.al (1985) and Yaseen. et al, (1994). These myoepithelial cells were not noticed around the lacrimal intralobular or interlobular ducts, in this study. However, Gillette. et al. (1980) and Kalleny, (1995) described myoepithelial cells around these ducts, in the human lacrimal glands. They explained that these cells might have a role in squeezing the lacrimal secretion in the beginning of its flow.

In present study the lacrimal gland of the rabbits, have a well developed duct system in which the intralobular ducts are lined by a single layer of cuboidal or low columnar epithelial lining. These findings were in agreement with Gillette. et al, (1980) and Kalleny, (1995).
The increase in the size of the lacrimal acini in senile rabbits could be attributed to the accumulation of the secretory granules in these cells. These findings were in agreement with Yaseen et al., (1994).

The increase of connective tissue fibers, inflammatory cells in senile lacrimal glands, might be attributed to the repeated episodes of inflammation of these glands throughout life. These inflammatory reactions might be followed by fibrosis with subsequent ischemia and degeneration of acinar cells. This was in agreement with Demato, B.E., Allen et al. (1984), Roen et al. (1985), Yaseen et al., (1994) and Draper et al. (1998). On the other hand, periductal areas as well as the structural damage of the acini. In senile age Makinodan and Kay, (1980), reported that repeated episodes of some systemic diseases such as rheumatoid arthritis were thought to involve the lacrimal gland.

The histological changes, in senile lacrimal glands, could be also due to recurrent episodes of vasculitis and arteriosclerosis and as aging lead to ischemia of the lacrimal acini. Parish, (1980) and Friedlander, (1992) postulated similar explanation.

The presence of periodic acid Schiff reactive substance in the acini and connective tissue stroma of rabbits lacrimal glands, noticed in this study, came in agreement with Jensen, O.A., Hensen, F., Kallen, (1995) and Garginlo, A.M., Aglio, C.D., Coliolio, P., Ceccarell, P. and Pedini et al., (1999) who found polysaccharides in the Lacrimal acini of mouse, man, rabbits and sheep’s. On the other hand, Yashimora et al., (1980), and Khater, (1984), denied the presence of polysaccharides in the lacrimal acini and ducts of most mammals except the rabbits.

The presence of a moderate positive PAS reaction, in the lacrimal acinar cells of the young animals, might be due to the accumulation of glycogen droplets in these cells Nagato, (1993). This moderate positive PAS reaction, in the lacrimal acinar cells of both young and adult animals, and strong positive PAS reaction in ductal cells were noticed also by Draper et al., (1998), in the rat lacrimal gland. However, the weak positive PAS re-
action, in the rat lacrimal acini of senile animals, might denote a decrease of the secretory capacity in the exhausted senile tissue. Furthermore, the thickening of the basement membrane, of the lacrimal acini in adult and senile animals, could be the result of recurrent addition of basement membrane materials from the surrounding connective tissue cells. This, in turn, aggravates the ischemic condition of the acini. These findings were documented by Yaseen. et al, (1994).

Draper. et al, (1998) postulated that the strong positive acid phosphatase reaction was observed in the basal portions of the lacrimal acinar cells in both young and senile animals, might explain why the lacrimal gland, in these two vulnerable ages, had high protective mechanisms from exposure to infections. However, the decline of the enzyme activity in the lacrimal acini, of senile rabbits, might be due to either a reduction in the ability of the acini to synthesize the enzyme or the number of acinar plasma membrane receptors for secretagogues. This is in agreement with study in which strong positive acid phosphatase reaction was observed in the basal portions of the lacrimal acinar cells in both young and senile animals.

The presence of a strong positive acid phosphatase reaction, in the lacrimal ducts, in the present study, could be explained by Khater, (1984), who reported that the duct might secrete lysosomal enzymes. On the other hand, the appearance of acid phosphatase activity in the Intralobular and interlobular connective tissue of the lacrimal glands of adult and senile animals might be attributed to the appearance of the macrophage activity.

The lacrimal acinar cells had Supranuclear Golgi apparatus, mitochondria, with lamellar cristae and secretory granules. This might signify that these cells had an active secretory function. Iwamoto and Jackobiec, (1982) and Draper. et al, (1997).

The acinar cells contained numerous secretory granules either granular or homogenous in character. The difference in electron density of the secretory granules in these acinar
cells could be attributed to the nature of the secretory materials inside these granules. Iwamoto and Jackobiec, (1982) said that the lacrimal secretory granules contained both serous (proteins) and mucous (polysaccharides) substances in various proportions that may differ from cell to cell. Furthermore, Miyazaki. et al, (2001) reported that the increase in the electron density of these granules denoted an increase in their protein content.

In present study the apparent increase of homogeneous electron dense granules, in the lacrimal acinar cells of the young animals, might denote the predominance of their protein content. On the other hand, most of these granules, in the senile groups, have granular contents, which might denote the predominance of their mucous content. These findings were supported by Dilly, (1994) and Draper. et al, (1998), who reported that, the maximum protein content output of the lacrimal gland occurred during the young age.

Draper. et al, (1997) noticed, in rat lacrimal gland, a change in the type of secretory cells, and hence the type of acini, in different age stages. They found that the serous acini, in young rats were predominant while in growing and adult rats seromucous acini were predominant. They added that, in senile rats mucous acini became the predominant ones.

Draper. et al, (1998), noticed that signs of degeneration in the secretory acini of senile lacrimal gland were manifested by pyknosis of nuclei, and coalescence of the secretory granules. These degenerative changes might result from ischemia of the lacrimal acini, which in turn resulted from fibroses of the gland or vasculitis of its blood vessels. Deposition of collagen fibres as well as the presence of plasma cells, fibroblasts and fibrocytes were noticed inbetween the lacrimal acini, these changes might be attributed to the previous attacks of inflammation of the lacrimal gland throughout life. On the other hand, the degeneration of the cellular organelles and mitochondria was accompanied by decrease in the homogeneous electron dense granules of senile gland. This suggested that there is a reduction or an inability of the acini to
synthesize and secrete proteins and other substances as the animal proceeded in age.

The result of this study have demonstrated that lacrimal gland acini change with age both morphological and histological. The acinar cells in young glands were predominately serous (protein producing and secreting cells), change of acinar cells to seromucous (protein and mucous producing and secreting cells) in the adult age, and to mucous acini in senile age.

These results indicate progressive alteration in the specialized protein-secreting cells of the lacrimal gland with age. Initially from serous to seromucous to mucous acini, this progression from serous to mucous is important finding. However, the causes and mechanism of these changes are due to the normal process of cellular aging which involves loss of cellular homeostasis and maintenance over time Kirkwood, (1992), Rattan, (1995).

The major mode of cellular maintenance includes; processes of cellular repair, cell division, cell replacement, neuronal and hormonal responsiveness, immune response. Loss of one or a number of these mechanisms may be responsible for the changes in the acinar cells, Draper. et al, (1998).

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الخلاص العرفي
التغيرات في الفجوة الدموية للارانب مع تقدم السن
(دراسات بالمجهر الضوئي والكهرباء)

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استهدفت هذه الدراسة إيضاح التغيرات الهستوغلوكينيات والهستوكليميائية التي تحدث في الفجوة الدموية في الأرانب في الأراذل العمرية المختلفة مع ربط هذه التغييرات مع تلك التي تحدث في التركيب الدقيق لهذه الفجوة.

وقد استخدم في هذه الدراسة عدد ثلاثون أرنب على أعمار مختلفة مقسمة الى ثلاث مجموعات

المجموعة الأولى: عمرها من 3 - 5 شهور، المجموعة الثانية: عمرها من 6 - 9 شهور

المجموعة الأخيرة: عمرها من 12 - 15 شهور.

وقد استخرجت الفجوة الدموية من كل حيوان على حدة ومولحت للحصول على:

1- قطعات شمعية صبغت بالهيماتوكسنين والأيونين وصبغة مالوير وتفاعل الشيف الحامضي.

2- قطعات تلبية طازجة لتحديد نشاط انزيم النوكينات الحامضي.

3- قطعات شه رقيقة صبغت بالتصويرين الأزرق.

4- قطعات منتظمة الرقة للفحص بالليكروسكوب الإلكتروني.

في الأرانب توجد فجوة دموية ظاهرة في المنطقة الخلفية الظهرية لقلعه المين قطرها حوالي

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كما توجد غدة أخرى وهي غدة هاردير وليس لها نظير بالإنسان توجد في المنطقة الأمامية البطنية لقلة العين وقطرها حوالي 6 مم.

وقد أظهرت هذه الدراسة أن الغدة الدمعية الظهرية في الأرارب من 5 شهور تتكون من وحدات أنتروبية حويصلة تتفصل عن بعضها ببط谯ات كثيفة من النسيج السام لتتجمعها إلى فصوص تتكون الحويصلات من خلايا مكعبة ولها أنوية قاعدية مستمرة وسيرت بلازماً قاعدية محجبة. أما في الأرارب البالغة فقد لوحظ زيادة في كثافة النسيج السام حول الفصوص الدمعية وحول الأورمية الدموية وأيضاً لوحظ انتشار خلايا التهابية ووجود تحلل خلوى وتحلل في جدار بعض الحويصلات الدمعية.

للحظ في تفاعل الشيف الحاضمي ان الحويصلات الدمعية قد اكتسبت درجة متساوية في تفاعل الشيف الحاضمي في كل من الأعمار النامية والبالغة بينما اكتسبت هذه الخلايا درجة ضعيفه من ذلك التفاعل في الحويصلات الدمعية للحيوانات البالغة كما زادت سمنة الفضاء القاعدية للحويصلات الدمعية في كل من الحيوانات البالغة والمسنة.

أما في تفاعل الفضفاضات الحاضمي لوحظ تفاعل قوي في الأجزاء السفلية من خلايا الحويصلات الدمعية بينما اكتسبت باقي السيتوبلازم في هذه الخلايا تفاعل متساعاً متسعاً للانزيم.

وقد قلت نسبة هذا التفاعل مع تقدم العمر.

عند الفحص بالبيكروسكوب الإلكتروني وجد أن خلايا الغدة الدمعية بها جهاز جوليئي على نواة الخلية بالإضافة إلى الشبكة الأندوبلازمية الحببية والغريبات الآفازية وحيدود أن هذه الخلايا في الغدة الدمعية للحيوانات البالغة قد احتررت على كمية زائدة من الشبكة الأندوبلازمية العريضة والميتوكبدية وظهرت بعض علامات التحلل الخلوى في بعض من خلايا الحويصلات الدمعية في الحيوانات الأذية مثل انكماش افنجية بعض هذه الخلايا وتشرد في الشبكة الأندوبلازمية والميتوكبدية وظهور بعض الحويصلات السيتوبلازمية.

غدة هاردير:

تتكون غدة هاردير من وحدات أنتروبية حويصلة تتناقص إلى فصول ببطائق من النسيج السام. يوجد نوعان من الحويصلات في غدة هاردير (نوع رقم 1) (نوع رقم 2).

تتميز حويصلات النوع رقم 1 انها تجاوي ضيقة وخلاياها بها فجوات صغيرة مع انوية

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قاعدية مستديرة ً. أما النوع رقم (2) يتميز بتجاويف واسعة وفجوات في الخلايا أكبر واحادية مستديرة قاعدية. في الآثار المنبعثة تكون الحويصلات أكثر تقاربًا وغير منتظمة الشكل مع وجود أنبعج خارجي لخلايا الحويصلات.

المستنتاج:

يستنتج من هذه الدراسة أنه قد تحدث تغيرات هستولوجية وهيموكيماوية واخرى دقيقة في الفذة الدموية للأذناب أثناء مراحل عمرهم المختلفة قد تساهم في تفسير نقص افراز الدم المالي في الأعمار المتقدمة. يوصى بالعناية الفائقة بالعينين محاولة تجنب الاصابة المتكررة بالالتهاب في الفذة الدموية حيث يقل افراز هذه الغذة للانزيمات الواقية في عمر الشيخوخة.