BRIEF REVIEW

The aging human lens: structure, growth, and physiological behaviour

George Duncan, I Michael Wormstone, Peter D Davies

The aging human lens has been the subject of intense research over the past 20 years, for a number of quite disparate reasons. The fact that the incidence of cataract rises exponentially with age after 50 years' provides the driving influence for much of the effort, but the unique accessibility, homogeneity, and basic simplicity of structure of the organ itself makes it a fruitful system for fundamental studies of tissue growth, development, and differentiation.1-4 Images of the whole human lens in vivo have been available for detailed analysis since the introduction of the quantitative slit lamp (or Scheimpflug) camera (Fig 1). This has been invaluable in providing an understanding of the changes in shape and optical properties both of the 'normal' and cataractous aging lens.5-7 The lens is also accessible as an in vitro system of study through the provision of donor eyes for corneal transplant and general research. Since the lens has no direct blood supply, it survives well both in the globe itself and in organ culture media.8-10 Whole cataractous lenses were also once routinely available before the advent of extracapsular cataract extraction (ECCE) with intraocular lens implantation and in the past there have been combined slit lamp and in vitro studies which have correlated changes in light scatter and absorbance with specific alterations in ion and protein levels (Fig 1 and Marcantonio et al11 and Hockwin et al12).

The optical homogeneity of the lens is also reflected at the cellular level. The entire organ is composed of cells of surface ectodermal origin. The single monolayer of anterior epithelial cells in the mature lens represents a very static population with little cell division and less cell death occurring.2,4,11 Only cells near the equator divide before elongating to form fully differentiated fibre cells that fill the bulk of the lens. The symmetry of growth of the fibre cells was appreciated by late 18th century anatomists such as Sir David Brewster13-15 and the cellular symmetry has an optical correlate as very beautiful birefringence patterns can be obtained from the radially symmetric arrangements of lens fibres (Figs 1 and 2). The symmetry and homogeneity of the patterns are maintained into old age, and only break down when cataract intervenes. This symmetry is maintained throughout the continuing, relatively linear, growth of the lens from the age of 10 to 80 years.

Although the advent of ECCE surgery8-10 has deprived lens researchers of many intact cataractous lenses, it has opened up a further, and potentially just as valuable, field of research into the behaviour of aging lens cells.16 This review is, therefore, devoted to recent advances in our knowledge of the changes that occur in the cell biology and physiology of the intact lens and also very recent data that have become available concerning the proliferation of cells within the capsular bag that gives rise to age related differences in the severity and incidence of posterior capsular opacification (PCO).

The intact lens—structural and cell biological aspects

The human lens continues to grow throughout life and at all decades from 10 to 70 years; the male lens is heavier than its female counterpart.16 These age related differences between males and females are interesting because not only do their relative susceptibilities to cataract change with age, but so does their response to physical trauma. Below the age of 50, women have a lower incidence of cataract than men, but above that age the incidence is greater.17 This suggests a hormonal influence and recent epidemiological studies show that women undergoing hormone replacement therapy have a reduced incidence of cortical cataract compared with a control cohort of the same age.18 There have recently been significant advances in our understanding of the possible mechanisms underlying the hormonal input into cataract. Transforming growth factor (TGF-β), which is present in the aqueous and vitreous humours,19 has been shown to induce cataract in organ cultured rat lenses exposed to relatively high concentrations of the factor.20 Interestingly, lenses from male rats are more susceptible than those from female rats and, furthermore, the latter receive added protection from TGF-β if oestrogen is also present in the medium.21 The molecular mechanisms underlying the cataractogenic effect of TGF-β are poorly understood, but TGF-β is known to induce transdifferentiation of lens cells so that they produce at least two types of foreign protein, smooth muscle actin and collagen types 1 and 3.22 Neither of these is synthesised in significant amounts by normal lens cells, but can be detected in certain cataracts23 and in cells giving rise to PCO.24 The TGF-β stimulated production of abnormal intracellular and extracellular proteins disrupts the homogeneous structure of the anterior epithelium and light scattering multilayered cell aggregates are produced.25 Interestingly, if TGF-β is injected into the vitreous chamber of the rat eye in vivo, then the most pronounced changes occur at the bow region of the lens initiating more typical cortical cataracts.26

Not only do male and female lenses differ in their relative sensitivity to TGF-β, but they also respond differently to mechanical stress. Weale26,27 carried out a quantitative study of the birefringence of male and female lenses and although the overall pattern is the same (Fig 1) the effect of external stress on the birefringence pattern measured in vitro is different in males and females. Weale measured the greatest stress that could be given before an irreversible change in birefringence occurred and although in both cases the magnitude of the reversible stress declines with age, the rate of decline appears to be steeper with
female lenses. Furthermore he identified a number of female lenses in which the merest mechanical stress induced irreversible birefringence changes. He concluded that this pointed to a subtle structural difference between male and female lenses. It has previously been pointed out that the cytoplasm of lens fibres represents a remarkably stable gel-like structure and that calcium has a critical role to play in the maintenance of this stability. Interestingly, calcium also has a critical role in osteoporosis, which is not only more common in aging women than men, but it is a condition for which hormone replacement therapy also provides a good degree of protection.

Birefringence is an optical property arising from a high degree of order in a structure and it has been shown that α-crystallin, the major protein of the lens, is capable of a high degree of packing regularity and short range order at high concentrations. In fact, if this were not the case, then the lens optical density would be predicted to be much higher. The lens optical density does, indeed, increase with age and the rate of increase is much more apparent after the age of 40 years. The lens also becomes increasingly coloured (yellow) with age and the intrinsic fluorescence also increases. All of these changes tend to degrade the optical properties of the lens.

In cataract the changes in the optical properties of the lens are not generally uniformly distributed. Posterior polar cataracts can involve a very small volume of the lens, but as they lie directly on the visual axis, their effect on vision is great (Fig 1). Cortical cataracts can again involve only small areas of the cortex, while pure nuclear cataracts involve a post-translational change in the nuclear proteins alone while the cortex is quite clear and unaffected (Fig 1).

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**Figure 1** Images of normal human lenses (A, B, and C), posterior polar cataract (D, E, F), and pure nuclear cataract (G, H, I). Note that slit lamp camera images (A, D, G) all have a scattering reflect artefact (small white rectangle). The normal subject (A) was 40 years of age and the accompanying in vitro grid photographs (B) and polarising images (C) were obtained from a donor eye of similar age (42 years). The white arrow gives the direction of plane of polarisation of the major axis of the first order red plate. Note that (B) was photographed under fluid and so the grid is out of focus compared with (E) and (H), which were photographed in air. Also note that the polarising images of the cataractous lenses (F and I) maintain the blue/yellow radially symmetrical pattern of the normal lens (C) except where the opacities occur in highly-localised polar cataract (F) and where the brunescence is strongest in the nuclear cataract (I). The images (D), (E), (G), (H) are taken from Marcanotto et al while the additional images are unpublished.
fibres and in the mid sections of intact, isolated single fibres. \(^{39}\) It is possible that calpain also has a role to play in this process by removing space filling cytoskeletal elements which would normally prevent the fibre membranes from collapsing inwards and fusing. \(^{30, 41}\)

Pure nuclear cataracts with their brunescent, optically dense nucleus and clear cortex (Fig 1) probably represent a quite different uncoupling mechanism. The oxidative changes that have taken place in the nucleus are not present in the cortex. For example, protein disulphide bridges are formed exclusively in the nucleus and there is an accompanying oxidation of glutathione. The glutathione content of the clear outer cortex is normal and in the reduced form. \(^{32, 43}\) Furthermore, some post-translational modification of the nuclear proteins alone occurs which renders them fluorescent. \(^{39, 44}\) This functional uncoupling and selective oxidation of one region of the lens relative to another does not involve a calcium increase \(^{13}\) and does not involve massive structural reorganisation as the birefringence patterns from nuclear cataracts are relatively normal in the clear cortex regions but are obscured in the central regions only by the light absorbing brunescent nucleus (Fig 1).

### The intact lens—physiological aspects

The membrane mechanisms underlying the ionic imbalances of cortical cataracts have been investigated in detail by applying a combination of electrophysiological and radioisotope techniques to lenses that have been removed by intracapsular surgery. \(^{43, 44}\) Since all of the cells within the lens are normally in good electrical communication with one another, the voltage measured by inserting an electrode into the lens is the same at all points throughout the lens. \(^{45}\) The membrane mechanisms underlying the ionic imbalances of the cortical cataracts can be investigated by inserting a membrane potential measuring electrode into lenses that have been removed by intracapsular surgery. The membrane potentials of lenses with cortical cataracts measured in this way are extremely low while those of the pure nuclear variety are relatively high and are similar to normal lenses of the same age (Fig 3). The latter statement is important since the membrane potential of the normal lens appears to decline with age, particularly after the age of 40 years. The overall resistance of the membranes to the passage of ions can be determined by inserting a second current passing electrode into the lens. The decline in voltage is accompanied by a decrease in membrane resistance, indicating that some channel mechanism is being activated in the aging lens. \(^{47}\) Since the membrane potential is depolarising during this activation, a channel involving an increased movement of sodium must be involved. Although pure sodium channels do appear to reside in lens membranes, \(^{48}\) they occur relatively infrequently and a much more likely candidate for the ageing activated process is a widely distributed species of non-specific cation channel. This channel is present in lens membranes and appears to permit Na\(^+\), K\(^+\), and Ca\(^{2+}\) to pass. \(^{47, 48}\) It is interesting in this respect that the lens sodium and free calcium content also appears to increase after the age of 40. \(^{49}\) It is possible to mimic all of the age related membrane permeability and ion content changes in the lens simply by complexing or oxidising membrane sulphydryl groups. \(^{50}\) For example, perfusing the isolated lens with very low concentrations of the non-permeating sulphydryl complexing agent PCMPS leads to a very rapid depolarisation of lens membrane potential with a concomitant increase in membrane conductance. These changes are accompanied by an increased influx of sodium and calcium into the lens. \(^{49}\)
with the nuclear proteins. This may help explain why the totally different aetiology and morphological appearance of senile cataracts are, in fact, mixed in form, having contributions from both nuclear and cortical changes. Recent epidemiological studies of cataract do suggest that a high intake of antioxidants either in the diet, or in the form of supplements, does confer a considerable protective effect. It is possible to compute for the aging human lens, the expected change in ion permeabilities that would accompany the age related voltage depolarisation and increase in sodium content. There is a remarkable agreement between the relative increase in permeability to sodium and the increase in lens optical density measured at the wavelength of peak sensitivity of the eye (Fig 4). Both increase more rapidly after the age of 40 and indicate once more a common mechanism between alterations in the ionic and structural protein contents of the human lens.

The above account of the ‘resting physiology’ of the normal and cataractous lens gives little information concerning the ability of the lens to respond to more natural external stimuli and evidence has accumulated over the past few years that the lens can, indeed, respond to a very wide range of agonists. Again, using electrophysiological methods it is possible to assess how this ability becomes modulated by age. The human lens responds to a surprising range of growth factors and other cytokines including adrenaline, ATP, histamine, and acetylcholine. The response to the latter is particularly interesting as agents that interfere with acetylcholine metabolism, such as cholinesterase inhibitors, are known to induce cataract.

However, no physiological role for acetylcholine has yet been found in the lens. Electrophysiological experiments on whole human lenses are, however, beginning to shed some light on this intriguing problem since it has recently been found that ACh induces a marked depolarisation of the membrane potential of the lens. As the response declines with age, it may first appear that the lens ACh receptors may become less active with age. However, more careful analysis indicates that this is not the case. Thomas et al argue that since the lens resting voltage also naturally declines with age (see Fig 3) then the extent to which ACh can change or depolarise the lens voltage must also decline with age. In fact, when the ACh induced depolarisation is plotted against resting voltage, then a linear relation is indeed obtained, implying that the ACh receptors remain fully functional throughout the life of the lens.

Additional fluorometric calcium imaging experiments carried out on tissue cultured human lens cells and on the isolated anterior epithelium show that ACh also stimulates the release of calcium from endoplasmic reticulum stores through the activation of specific muscarinic receptors. There are many potential sources of ACh in the tissues surrounding the lens, including the ciliary body and retina, and it now appears that increasing ACh in the vicinity of the lens would be expected both to depolarise the lens and increase the free calcium content. Interestingly, an increase in lens calcium and membrane depolarisation is associated with cortical cataracts (see Fig 3 and Duncan and Jacob and Duncan and Hightower).

The lens capsule—PCO mechanisms and age

The most common surgical procedure currently performed to treat cataract is extracapsular cataract surgery (ECCE), which consists of lens substance removal and irrigation/aspiration of residual lens fibres. This procedure leaves the capsular bag in situ and permits an intraocular lens (IOL) to be implanted into this natural holder. However, a proportion of the lens anterior and equatorial epithelial cell population survives the various surgical manipulations. These viable cells do not remain static on the capsule, but proliferate most notably across the previously cell free posterior surface, subsequently
entering the visual axis. This aberrant growth gives rise to a secondary visual impairment known as posterior capsule opacification (PCO) or secondary cataract, affecting 20–50% of patients to the point where further surgery is required. A study by Moissiev et al. gives an overall PCO incidence, 4 years after ECCE surgery, of 41% from a pool of 94 patients. In addition, it was shown that patients aged over 40 years were found to have a lower incidence (37%) than those under 40 years (70%), which clearly demonstrated PCO is inversely related to age. PCO is of particular importance in infantile eyes where it is nearly universal if the posterior capsule is left intact. As PCO is a major clinical problem a great deal of effort has been directed towards developing both in vivo and in vitro techniques to investigate the mechanisms of PCO. However, despite the well established relation between age and PCO, very few experimental investigations have been carried out to understand this phenomenon.

In the normal lens the mitotic index is low and in vitro studies show that the dividing cells are strictly confined to the equatorial region of the epithelium. However, once the capsule has been breached and the fibres removed, the mitotic index is dramatically increased. Interestingly, the increase is first apparent in the equatorial zone rather than at the injured rhexis region. Within a few days, dividing cells can be detected throughout the anterior epithelium and cells can also be seen moving across the posterior capsule. Modern imaging techniques permit a detailed study of the characteristics of PCO in vivo and it is now apparent that cell growth can also be observed within a few weeks of the operation. Growth on the posterior capsule is more rapid and prolific than that on the IOL surface. This is precisely the pattern seen in vitro when a serum free medium is used to culture the capsular bag. Other clinical features of PCO are also reproduced in vitro and they include wrinkling and tensioning of the bag and light scattering and multilayering of cells. We can be confident, therefore, that the behaviour of cells in vitro faithfully reflect their behaviour in vivo.

Probably the most important finding of the in vitro studies has been the impressive growth of human lens cells in the capsular bag in an entirely growth factor free environment. This in itself helps to explain why PCO is such a common problem. Growth within capsular bags from younger donors (<40 years) was threefold faster than growth within older capsular bags (>60 years), again reflecting clinical findings. Interestingly, serum stimulation had little effect on younger cells, but older cells could be driven at rates approaching those of their younger counterparts (Fig 5). This suggests that the difference observed in the absence of serum is due to an age related slowing down in the production of growth factors, rather than a lack of appropriate receptors. Furthermore, some aged capsules showed only partial cover of the posterior capsule and addition of high serum induced a rapid proliferation until confluency was attained (Fig 6). There are reports that PCO in vivo can be greatly stimulated by inflammatory episodes.

The capsular bag system has shown that lens cells have a great, inherent capacity to proliferate and yet cells within the intact lens divide only slowly and in a very defined region. It will, therefore, be necessary in the future to investigate the factors that limit cell growth in the lens as well as those which stimulate proliferation. Most studies in the past have been limited to investigating the external factors, such as FGF, that might control growth. The recent data reviewed here suggest that much more attention should be focused on the factors produced by the lens itself both when growth is limited and when it is stimulated. The necessary in vitro systems are now in place for the whole lens and capsular bag for these age related studies to proceed apace.

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GEORGE DUNCAN
I MICHAEL WORMSTONE
School of Biological Sciences, University of East Anglia, Norwich NR4 7TJ

PETER D DAVIES
Department of Ophthalmology, West Norwich Hospital, Bowthorpe Road, Norwich

Correspondence to: G Duncan.
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GEORGE DUNCAN, I MICHAEL WORMSTONE and PETER D DAVIES

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