

PERSPECTIVE

Mechanosensitivity and the eye: cells coping with the pressure

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The cells of the various organ systems in humans are subject to mechanical forces to which they must respond. Here the authors review what is known of the ways in which the cells of animals, ranging from the prokaryotic to humans, sense and transduce mechanical forces to respond to such stimuli. In what way this pertains to the eye, especially with respect to axial myopia and the pressure related disease of glaucoma, is then surveyed.

across different organ systems, and indeed even across different species.

For a cell to be mechanosensitive it must be able to respond to forces acting within its immediate vicinity. The sensing of these forces and conversion into signals that promote a response is termed mechanotransduction. Aortic endothelial cells elongate and their long axis and microtubules align perpendicular to flow induced shear stress²; aortic smooth muscles,³ cardiac myocytes,⁴ and skin and scar fibroblasts⁵ orientate perpendicular to the direction of stress; when strained, arterial smooth muscles develop more prominent actin cables and lay down extracellular matrix (ECM) proteoglycans⁶; intermittently stretched skeletal muscles bearing static loads increase protein and collagen synthesis and prostaglandin F (PGF-2 α)^{7,8}; new bone formation occurs after a load cycle⁹ and osteoblasts produce more PGE-2 and cAMP¹⁰; lung type II alveolar cells, when stretched, release more surfactant and phosphatidylcholine, surfactant's major component¹¹; cardiac and skeletal muscle mass are affected by external load in vivo¹²; when the median nerve is subjected to pressures of 30 mm Hg or more in carpal tunnel syndrome, its nerve cell bodies and axonal transport are altered^{13,14}; trabecular meshwork (TM) cells express the protein myocilin in response to stretch and raised hydrostatic pressure.¹⁵ Some cells mediating sensation are specialised for mechanotransduction: minute deflections of auditory hair cells are transduced into signals that allow us to hear, and pacinian receptors in the skin transduce pressure into electrical signals. Other specialised baroreceptor cells in the myocardium, arteries, and kidney have a feedback role on cardiac load, serum osmolarity, and blood pressure to allow physiological regulation.

Early laboratory studies of the biological consequences of extreme hydrostatic pressure were inspired by deep sea exploration, chief of which was the *Talisman* dredging expedition in the 19th century, in which various baroduric or barophilic organisms were found living in depths exceeding 6000 metres under pressures of more than 600 atmospheres (4.6 \times 10⁵ mm Hg; 1 atm = 760 mm Hg). It is estimated that over half the earth's biosphere by area and nearly 90% of the seabed is under at least 1 km of water and pressures exceeding 100 atm. The Marianas Trench, the deepest known part of our oceans, is over 11 000 metres deep with its floor under at least 1100 atm pressure. Entire ecosystems exist in these places¹ and organisms living here must be able to survive under pressure.

Our own cells, singly or in tissues, are subject to mechanical forces that, though nowhere as extreme as at the ocean bed, must be responded and adapted to. Forces are exerted on the cell from without—hydrostatic pressure, tension, compression (squeezing), torsion (twisting), vibration, shear and stretch (tensile stress) that can deform the cell—and from within, such as changes in osmotic pressure and swelling. By this the cell receives vital information about its physical world.

It stands to reason that if such stresses are excessive or not responded to optimally, they may damage cells, cause disease, and even kill the organism. How do cells detect mechanical changes in their environment and signal responses? What is the nature of such responses? Cells of our various body systems—for example, heart and blood vessel, lung, kidney, bone, neurological and eye, are constantly subjected to forces which may well overwhelm their capacity to respond accordingly. What events transpire when this happens? Are damaged cells able to repair themselves? It is certainly possible that some basic mechanisms mediating sensitivity to mechanical forces are common to cells

Much fundamental research has sought to define the cellular mechanisms mediating mechanosensitivity. One such mechanism implicates membrane bound mechanically gated channels which are sensitive, speedy in response, and allow huge ion influxes and amplified signals. Yet another mechanism is the coupled

Abbreviations: BDNF, brain derived neurotrophic factor; cAMP, cyclic adenosine monophosphate; DAG, diacyl glycerol; ECM, extracellular matrix; EGF, epidermal growth factor; IOP, intraocular pressure; IP3, inositol-3-phosphate; MAP-kinase, mitogen activated protein kinase; MMPs, matrix metalloproteinases; PDGF, platelet derived growth factor; PGF, prostaglandin F; PKA, protein kinase A; PKC, protein kinase C; RGC, retinal ganglion cells; TIMP, tissue inhibitor of MMP; TM, trabecular meshwork; VEGF, vascular endothelium growth factor

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and interconnected cytoskeleton-ECM complex (see fig 1), which has critical roles in intracellular signalling.¹⁶ Mechanisms underlying mechanosensitivity in animals ranging from unicellular prokaryotes to invertebrates to humans have been described and we have sought to survey this body of information in a way that is relevant to the eye.

MECHANICALLY GATED ION CHANNELS

Mechanically gated channels open when the cell membrane receives stimuli such as stretch, shear, and displacement. Some channels are permeable to anions (for example, Cl^-) while others are permeable to cations (for example, Ca^{2+} , K^+).¹⁷ These channels are the most widely studied mechanosensitive structures in cell membranes. They were first conceived in whole cell studies of specialised mechanosensory neurons,^{18, 19} but first discovered by patch clamping in cultured chick skeletal muscle²⁰ at almost the same time as their discovery in embryonic *Xenopus* muscle.²¹ They have now been identified in many cell types including those within the eye.

Bacteria mostly have rigid cell walls that protect them from excessive swelling and deformation, although this also renders them less sensitive to mechanical stimuli. How bacteria might respond to external forces has been suggested by the discovery and cloning of mechanically gated channels in *Escherichia coli*²²⁻²⁴ and subsequently other bacteria, both Gram positive and negative having different cell wall structures.²⁵⁻²⁷ A class of *E coli* receptors called MscL are postulated to protect the bacterium from osmotic damage as they open just below pressures that would otherwise disrupt the bacterial cell membrane. Mechanically gated channels have also been demonstrated in *archaea*,^{28, 29} the other major domain of the phylogenetic tree. These micro-organisms live in harsh environments such as extremely hot ocean hydrothermal vents or the salty Dead Sea. The mechanically gated channels of prokaryotes are generally activated by high tensions, have high ion conductance, and lack ionic specificity when compared with those of eukaryotes. It is known however that genetically mutating prokaryotic channels can confer upon them the gating characteristics of eukaryotic channels.³⁰

Eukaryotic cells lack rigid walls but their plasma membranes are internally supported by an elaborate cytoskeleton. The actin cytoskeleton resists membrane deformability, while providing internal scaffolding for linking or tethering various proteins such as signalling molecules and mechanosensitive channels. Disrupting actin increases the conductance of mechanosensitive channels.³¹ The actin cytoskeleton is a dynamic structure that constantly rearranges itself and is interconnected with the ECM and cytoskeleton of adjacent cells by specialised membrane bound proteins. Any changes in the cytoskeleton, plasma membrane and ECM might thus be expected to influence the mechanically gated channels.

Mechanically gated channels have been identified in diverse cells.³² There are several classes of channels, many of which are permeable to cations, not uncommonly Ca^{2+} , a ubiquitous intracellular messenger. Ca^{2+} rushing into the cell can itself induce Ca^{2+} release from internal stores and trigger molecular switches such as protein kinase C (PKC) which in turn phosphorylates target and gene regulatory proteins. For instance, bending of auditory hair cells opens a non-selective mechanically gated cation channel that depolarises the cell and activates voltage sensitive Ca^{2+} channels to induce synaptic vesicle exocytosis.^{33, 34} A recently identified family of channels with weak inwardly rectifying K^+ conductance³⁵⁻³⁷ includes TREK-1 and TRAAK, which are found widely in brain and spinal cord.³⁸ In vitro, TREK-1 is sensitive to suction and osmotic swelling and shrinkage. TRAAK is sensitive to suction and pressure but also to arachidonic acid. It is present in the retina in ganglion cells, amacrine, horizontal and rod bipolar cells, and outer segments of photoreceptors.³⁸⁻⁴⁰

Sensory elements showing specific responses to mechanical stimulation have been identified in the cornea of rabbits, and cornea, sclera, bulbar conjunctiva, and uvea of cats.^{41, 42} Mammalian corneal epithelium, non-pigmented ciliary epithelium, and TM cells also have volume regulated outwardly rectifying chloride channels which respond to hypotonic cellular swelling and potentially help maintain the clarity of the ocular media and the secretion and outflow of the aqueous humour.⁴³⁻⁴⁸

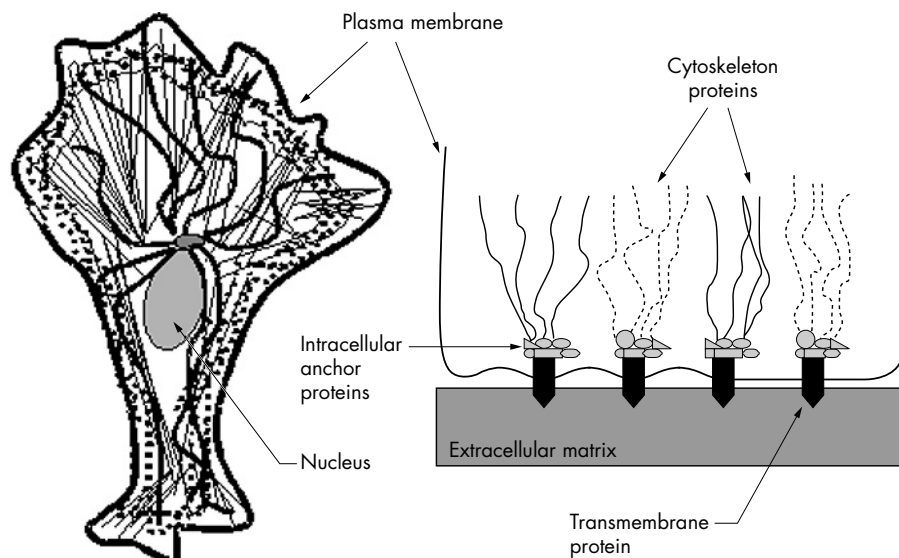


Figure 1 The cell cytoskeleton (left) and its relation to cell-extracellular matrix adhesions (right). (Left) The cytoskeleton consists of three main proteins: actin microfilaments (thin lines), microtubules (bold wavy lines), and intermediate filaments (dotted lines). Microtubules meet at the centromere next to the nucleus. (Right) The family of integrins are important transmembrane proteins that anchor the cell to the extracellular matrix. Cell-matrix contacts are called focal adhesions if they connect with actin microfilaments intracellularly, and hemidesmosomes if they connect with intermediate filaments. Integrins link with actin via various proteins such as talin, α -actinin, and filamin at focal adhesion complexes.

TRANSMISSION OF FORCES BETWEEN THE CYTOSKELETON AND EXTRACELLULAR MATRIX

The eukaryotic cell can be thought of as a physical structure in which mechanical stresses are distributed across its interconnected elements.^{49–50} The integration and coupling of the cell cytoskeleton to its plasma membrane (and its channels), the ECM via focal adhesions, and adjacent cells by cell-cell contacts such as adherens junctions provides a means for mediating mechanosensitivity and transduction.

The main cytoskeletal proteins are actin, intermediate filaments, and microtubules. Actin filaments, which determine cell shape and movement, are distributed throughout the cell, though especially just beneath the plasma membrane as the cortical cytoskeleton. Intermediate filaments are the cell's cable scaffolding. They give it mechanical strength, help it withstand shear stress, and assist with motility. Microtubules direct intracellular traffic and determine where organelles lie. Integrins are membrane spanning proteins that help cells grip their ECM at sites called focal adhesions (for actin) or hemidesmosomes (for intermediate filaments). Integrin intracellular domains link to the cytoskeleton by proteins such as talin, α -actinin, and filamin as well as many other intervening proteins, as illustrated in figure 1. The cytoskeleton is linked to adjacent cells at cell-cell junctions via other intervening proteins such as cadherin. These linkages facilitate the forming of an interconnected structural unit, allowing concerted regulation of cell shape, orientation and movement,^{49–50} and ECM organisation.^{51–52}

The cytoskeleton imparts rigidity to cells and the fluid membrane bilayer and helps the cell withstand deforming forces and shear stress. Having this internal scaffolding means that eukaryotic cells do not need rigid walls, can receive feedback from their physical surroundings, and are able to assume non-spherical shapes. For example, microvilli can stably maintain excess plasma membrane to increase surface area, and better cope with intracellular volume changes; erythrocytes are concave and have 40% extra membrane surface area than if they were spherical, allowing volume increases to be easily accommodated.³⁰ The erythrocyte's interconnected cortical cytoskeleton of spectrin actin tropomyosin is somewhat "coiled" at rest, permitting structural expansion as needed.⁵³ Such variation in membrane curvature, surface area and tension might also be expected to influence the characteristics and gating of mechanosensitive channels. The Rho family of small GTPases, comprising signalling proteins such as Rho, Rac, and Cdc42, has a particularly vital role in regulating the response of cytoskeletal and cell contacts to mechanical forces.

PLASMA MEMBRANE MECHANOTRANSDUCTION AND REPAIR

The plasma membrane encapsulates the cell, is relatively impermeable to water soluble molecules, and maintains its internal environment as different from outside. It is a fluid, continuous lipid bilayer held together by non-covalent forces. Proteins embedded in or spanning the membrane are responsible for nearly all its functions: ionic and molecular transport, catalysing reactions, acting as receptors for signal transduction, and linking the membrane to the cytoskeleton, ECM, and adjacent cells. Each has a postulated role in mechanotransduction. For example, stretching the plasma membrane can activate several membrane bound intracellular signalling proteins such as receptor tyrosine kinases and G proteins, phospholipase C and phospholipase A2, the inositol phospholipid signalling pathway (via IP3 and Ca^{2+}), eicosanoids (via DAG and PKC), and cAMP (via PKA). Stretched lung alveolar epithelial cells activate G-proteins and MAP kinase pathways⁵²; mechanically stressed

osteocytes in culture increase PGE-2 by Ca^{2+} and cytoskeleton mediated mechanisms.⁵⁴

Cells have mechanisms to repair or reseal mechanically damaged or ruptured membranes.⁵⁵ It is proposed that both the fusion of endocytotic vesicles, a slow process taking minutes to hours, and exocytosis taking seconds can help plasma membranes reseal by incorporating freshly recruited vesicular membrane.^{56–57} Both are triggered by high intracellular Ca^{2+} concentrations and are under the regulation of the Rho family of GTPases. Fused larger vesicles traffic from the Golgi apparatus along microtubules towards the plasma membrane where bursts of exocytosis seal punctures.⁵⁵ Cytochalasin D affects actin polymerisation and appears to promote resealing possibly because it reduces membrane tension.⁵⁸ In erythrocytes, the spectrin network facilitates membrane healing after the plasma membrane is ruptured mechanically.⁵⁹

LIFE AND DEATH AFTER MECHANICAL STRESS

Cells can be injured and die if mechanical stresses become overwhelming. Disruption of the plasma membrane can cause influx of ions and alter cell osmolarity and volume, affecting ionic gradients, membrane potential, and the functioning of membrane proteins such as channels. Mechanically gated channels, by permitting Ca^{2+} influx, can elevate intracellular $[\text{Ca}^{2+}]$ and stimulate further Ca^{2+} release from cytoplasmic stores. Outward pumping of Ca^{2+} may be impeded if other membrane proteins stop working. Changes in intracellular $[\text{Ca}^{2+}]$ are needed for cell growth, differentiation, movement, intracellular signalling and synaptic transmission, but too much Ca^{2+} can cause necrosis and apoptosis. Necrotic cells swell and burst, but in apoptosis—a tightly regulated programme of cell death—the cell shrinks, its cytoskeleton involutes, nuclear envelope dismantles, nuclear DNA fragments and surface alters, inviting phagocytosis.

Cells usually need to be attached to underlying ECM to grow, divide, and survive, and these activities are modulated by various influences such as biochemical activity at cell-ECM adhesions and growth factors. Growth factors, by binding to cell surface receptors, can activate growth promoting second messengers, while some also promote mitogenesis. Examples of growth factors, which various cells have been shown to secrete when their substrata are mechanically stretched *in vitro*, are PDGF, EGF and VEGF (for example, fetal lung (PDGF), smooth muscles (EGF), and retinal pigment epithelium (VEGF)).⁶⁰

AXIAL MYOPIA, A MODEL OF OCULAR "BAROPATHY"

The development of high myopia is associated with axial elongation, primarily as a result of deepening of the vitreous chamber.⁶¹ Axial elongation in myopic eyes is associated with scleral remodelling which causes marked thinning of the sclera especially at the posterior pole. Loss of scleral tissue and scleral weakening precipitates local ectatic change or posterior staphylomata may form.^{61–63} The altered scleral morphology is associated with changed collagen fibril ultrastructure, greater numbers of small diameter collagen fibrils, and a more lamellar arrangement of collagen fibril bundles in the posterior sclera.⁶⁴ Myopic scleral remodelling also results in reduced glycosaminoglycan and collagen content.⁶⁵ These changes reduce scleral resistance to expansion in response to a range of intraocular pressures (IOP), resulting in axial lengthening.

A study by Phillips *et al*⁶⁶ suggests that the presence of specialised contractile fibroblasts called myofibroblasts protects against the development of axial myopia. In this study, IOP was experimentally raised in both chick and tree shrew eyes. On normalising IOP, the chick but not the tree shrew

eye showed residual axial elongation as only the tree shrew was able to recover from the induced axial myopia. Tree shrew eyes were found by immunohistochemistry to have myofibroblasts within the sclera and choroid, but chick eyes had myofibroblasts only in the choroid.⁶⁶ It has been proposed that myofibroblasts work as an interlinked syncytium to both sense and respond to changes in their mechanical environment, providing a mechanotransduction pathway that helps maintain ocular structure. It is probably by contractile myofibroblasts responding to altered force-length relations within the scleral ECM that the tree shrew is able to restore vitreous chamber depth.⁶⁹ Scleral fibroblasts are known to produce matrix metalloproteases (MMPs), which further contribute to scleral remodelling.⁶⁷

At the gene expression level, microarray studies of human scleral fibroblasts show differential changes in gene expression after short (30 minutes) and longer (24 hours) durations of stretching. Genes found to be significantly upregulated code for cell receptors, protein kinases, cell growth/differentiation factors, ECM proteins, lipid and protein metabolism, transcription factors and water channels, and could be involved in the scleral remodelling occurring during axial myopia.⁶⁸

GLAUCOMA, ANOTHER OCULAR "BAROPATHY"

Glaucoma is an optic neuropathy that has as its basis slowly progressive loss of retinal ganglion cells (RGC) and their axons, and as a major risk factor, IOP. At least some RGC death in glaucoma occurs by apoptosis,⁶⁹⁻⁷¹ which may be a mechanism by which putative insults such as elevated hydrostatic pressure,⁷²⁻⁷³ glutamate neurotoxicity,⁷⁴⁻⁷⁵ neurotrophic deprivation,⁷⁶ autoimmunity,⁷⁷ nitric oxide,⁷⁸ and intracellular calcium toxicity and ischaemia cause RGC death.

Several lines of evidence implicate the optic nerve head's lamina cribrosa, where axons exit the eye, as the primary site of RGC damage leading to death.⁷⁹⁻⁸¹ Quigley *et al*'s⁸² human postmortem studies point to the cribriform plates of glaucoma eyes becoming compressed with disease, then collapsing and bowing posteriorly. Similar changes are seen in primate models of glaucoma.⁸³ Experimental studies show that slow and rapid axonal transport can get blocked at the lamina cribrosa with raised IOP,⁷⁹⁻⁸⁰⁻⁸⁴⁻⁸⁵ possibly affecting the transport and availability of trophic factors such as BDNF which ganglion cells need to survive.⁸⁵ This indicates that mechanical changes at the lamina cribrosa physically impinge on and injure RGCs, depriving nerve cells of structural and nutritional support and promoting apoptosis.

Some *in vitro* evidence also supports the possibility that RGCs and astroglia may themselves be directly affected by elevated hydrostatic pressure. Apoptosis is increased in cultured RGCs and other central nervous system cells subjected to hydrostatic pressures mimicking conditions in acute (100 mm Hg) and chronic (30 mm Hg) glaucoma. In these experiments, higher pressures were associated with more apoptosis.⁷²⁻⁷³⁻⁸⁶ Cultured lamina cribrosa astroglia, when exposed to raised hydrostatic pressure, release tumour necrosis factor α and nitric oxide, both pro-apoptotic substances; if RGCs are co-incubated with these pressurised glia, the RGCs undergo apoptosis.⁸⁷ *In vivo*, retinal astrocytes upregulate glial fibrillary acid protein expression when IOP is raised⁸⁸; the caspase cascade has been implicated in RGC death following exposure to raised hydrostatic pressure.⁸⁸ TRAAK, the mechanically gated K⁺ channel, which is also opened by pressure, membrane stretch and arachidonic acid, and blocked by gadolinium, is found in the cell bodies and axons of RGCs. *In vitro*, treating RGCs with arachidonic acid to open TRAAK channels causes apoptosis,⁸⁹ suggesting the

possibility that such mechanically gated channels have a role in neuronal death caused by pressure.

Elevated IOP is implicated in glaucomatous optic neuropathy. IOP is determined by the equilibration of aqueous humour production by the ciliary body and aqueous outflow by the TM and uveoscleral pathways. Stretched cultured TM cells are reported to elongate and rearrange their actin, phosphorylate paxillin in focal adhesion complexes, and activate tyrosine phosphorylation and MAP kinase signalling.⁹⁰ High conductance Ca²⁺ activated K⁺ channels are activated when cultured TM cells are stretched or undergo volume changes.⁹¹ When exposed to elevated hydrostatic pressure, cells from pigmented and non-pigmented ciliary epithelium and the TM become rounder, their processes retract, actin filaments are displaced mainly to the periphery of the cytoplasm, and adenylyl cyclase activity is increased⁹²; stretched TM cells upregulate genes controlling inflammation, secretion, ECM digestion, cell responses to oxidative stress, and cytoskeleton organisation⁹³; stretch decreases levels of α B-crystallin, which has a possible role in stabilising and regulating actin;⁹⁴ phacoemulsification ultrasound vibration induces signal transduction in TM cells *in vitro*⁹⁵; stretched cultured TM cells alter their secretion of MMPs and their inhibitors, suggesting that ECM turnover is altered and apparently influences fluid outflow through the TM.⁹⁶ Mechanosensors are thought to be present at the scleral spur and might serve as a form of regulatory feedback.⁹⁷ In the TM, myocilin expression is upregulated and its secretion increased with raised hydrostatic pressure and stretch.¹⁵⁻⁹³⁻⁹⁸⁻⁹⁹ *In vivo*, the TM is itself in a contracted state, which if relaxed can dramatically increase aqueous outflow.¹⁰⁰⁻¹⁰²

TM cells subjected to mechanical strain/stretching show increased production of MMPs.¹⁰³⁻¹⁰⁵ Mechanically strained bovine TM cells show increased stromelysin and gelatinase A activity, which is reversible with relaxation of mechanical stretch.¹⁰⁴ Stretched TM increases production of MMP-2 and MMP-14 while reducing levels of the tissue inhibitor of MMP (TIMP-2).¹⁰³⁻¹⁰⁵ Pressure induced alterations in MMP activity with resultant ECM degradation may affect TM outflow resistance and have an important role in IOP regulation.

Taken together, these findings indicate that cells within the outflow pathways exhibit features of mechanosensitivity which may well provide them with the means to respond to physical changes in their environment.

CONCLUSIONS

It appears that cells have a repertoire of strategies for dealing with mechanical stimuli. Many of these strategies are common to the cells of different mammalian tissues and some are universal to a wide range of living organisms. Several appear relevant to the eye and to the way ocular cells respond to their physical environment in health and disease.

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