Cerebrospinal fluid pressure and the eye

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ABSTRACT
Cerebrospinal fluid pressure (CSFP) interacts with intraocular pressure (IOP) and blood pressure to exert a major influence upon the eye, particularly the optic nerve head region. There is increased interest regarding the influence of CSFP upon disorders affecting this region, in particular glaucoma and idiopathic intracranial hypertension. Additionally, a high proportion of astronauts develop features similar to idiopathic intracranial hypertension that persist for years after returning to Earth. The factors that affect the CSFP influence upon the optic nerve and globe are likely to influence the outcome of various ophthalmic disorders.

INTRODUCTION
The optic nerve separates the central nervous system and cerebrospinal fluid pressure (CSFP) from the intraocular structures and intraocular pressure (IOP). The lamina cribrosa and its constituent neural, vascular and connective tissue elements are very sensitive to alterations in pressure across this region, with follow-up effects into the rest of the eye. Such changes influence the occurrence of glaucoma, idiopathic intracranial hypertension and venous occlusions.

ANATOMY
The optic nerve subarachnoid space (ONSAS) surrounds the optic nerve as it leaves the cranial cavity up to the posterior globe where it is terminated anteriorly by the sclera. Along its path there are numerous trabeculae and septae whose shape varies with some thickening towards the termination of the subarachnoid space (figure 1). Internal to the ONSAS lies the pia mater and then the optic nerve proper. Externally lies the dura mater and intracranial orbital tissue. On closer examination using tracer and specialised immunohistochemical stains, lymphatic structures containing CSF material can be identified in the tissue surrounding the termination of the ONSAS. These lymphatic structures appear more prominent in lower mammals, such as rodents.

PRESSURES AND DIFFERENCES
IOP averages 15 mm Hg with a normal range of 10–21 mm Hg in most populations and is generated by the balance between production from the ciliary body and egress through the trabecular meshwork and uvea. There is a small increase with age from 15.8 mm Hg at age 50 years to 16.1 mm Hg at age 80 years, although this effect is linked to the rise in systemic blood pressure.

CSF is formed in the choroid plexus and flows through the ventricular system into the subarachnoid space where some is thought to flow down the ONSAS traversing the trabeculae and septae present in it. The bulk of CSF is thought to drain out through the arachnoid granulations, however, up to 30% of CSF drains out via lymphatics in the olfactory bulb and ONSAS in lower mammals with an unknown proportion flowing out through these systems in primates and humans (figure 1). The CSFP, like IOP, has pulsatile having a small phase difference with peak IOP lagging behind peak CSFP but their trough pressures coincide. Unlike IOP, the CSFP is markedly altered by postural changes, with the normal range of CSFP in the lateral decubitus position being 5 to 15 mm Hg but in the sitting position being −10 to 0 mm Hg at eye level.

Although IOP doesn’t change as much as CSFP with postural change it does increase by 2–4 mm Hg when lying down from sitting. Hence, one should take into account the posture when interpreting results of CSFP measurements. CSFP tends to reduce with age in adulthood, from mean 11.5 mm Hg in less than 50 year-olds to 9.0 mm Hg in 70 year-olds. Production and drainage of CSF and aqueous are influenced by hydroxysteroid dehydrogenase with increased CSFP effect in obesity. Stimulation of certain regions of the rat brain increases IOP, CSFP and blood pressure to varying degrees and varying latency. It is likely that other humeral and neural stimuli influence CSFP and IOP.

MEASUREMENT OF CSFP
Currently, invasive techniques involving lumbar puncture or cranial burr holes are required for CSFP measurement. The development of non-invasive techniques for CSFP measurement would be a major advance for diagnosis and monitoring of various disorders. Previous attempts using transcranial Doppler, tympanic membrane reflectivity and optic nerve sheath diameter by ultrasound or MRI have given largely inaccurate measurements. The two most promising avenues being currently explored are refinement of ophthalmodynamometric measures of venous pulsation pressure and balancing intracranial/orbital ophthalmic artery flow velocities. Some groups have begun using multivariate algorithms incorporating blood pressure and body mass index as indirect indicators of CSFP, however one should remain aware of the limitations and dependencies of this technique.

RELATIONSHIP BETWEEN ONSAS PRESSURE AND INTRACRANIAL PRESSURE
It has been assumed that ONSAS pressure is equivalent to intracranial pressure and is likely to be the case in healthy animals. However, the work of Killer and coauthors clearly demonstrates the

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presence of significantly large trabeculae, resembling fenestrated sheets along the ONSAS pathway (figure 1)26 and that in some conditions, such as normal tension glaucoma and others, there is relative stasis of CSF flow along this pathway.27 They may alter fluid flow along the ONSAS although the flexibility of the septae may not impede pressure transmittance. However, significant trabecular changes such as following meningitis or other diseases may effectively block CSF flow and cause a disruption of pressure transmittance. In the normal situation we tend to assume that the ONSAS pressure is equivalent to intracranial pressure, however, another factor must be considered and this is the orbital tissue pressure found to be 2.6–4.4 mm Hg in the supine position using needles inserted into the orbit.28 29 Tissue pressure measurement using needle insertion tends to give a slightly higher value than when less invasive micropipette techniques are used,30 so true orbital tissue pressure may be lower than these values. When a person is sitting or standing and the intracranial pressure falls to zero or below at eye level, this is likely to be transmitted along the ONSAS. If orbital tissue pressure is greater than the intracranial pressure then it will tend to compress and, hence, collapse the ONSAS and force CSF fluid back into the intracranial compartment. It may also compress regions of the ONSAS causing loculation with lack of pressure transmission. This presumed influence of orbital tissue pressure is supported by experimental data (figure 2) in the dog where below CSFP 0 mm Hg the ONSAS pressure remains at 0 mm Hg.

**THE OPTIC NERVE RETROLAMINAR TISSUE PRESSURE**

The retrolaminar tissue pressure (RLTP) is the interstitial pressure within the optic nerve neural tissue just posterior to the lamina cribrosa. It is not always equivalent to ONSAS pressure with a variable pressure difference occurring across the pia mater indicating some pial elastic effect upon the RLTP.25 When there is a low external pressure the elasticity of the pia mater tends to elevate RLTP and this has been demonstrated experimentally whereby at ONSAS pressures of 0 mm Hg in the dog, the retrolaminar pressure is 3 mm Hg and remains at this level until the ONSAS and intracranial pressures rise to 3 mm Hg or above (figure 2).25 Above this pressure, the tension exerted by

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**Figure 1** Key anatomical features of the optic nerve and eye in relation to the optic nerve subarachnoid space demonstrating the path to the intracranial cavity. The variation in trabeculae is shown and adapted from Killer et al [2]. Glial lamina cribrosa (G) and scleral (collagenous) lamina cribrosa (S) are shown. Central retinal artery is omitted for clarity.

**Figure 2** The relationship of key pressures (±SEM) to intracranial cerebrospinal fluid pressure (CSFP) in the dog. A strong linear relationship is seen between optic nerve subarachnoid space pressure (ONSASP ●), retrolaminar tissue pressure (RLTP ○), venous pulsation pressure (VPP ▲) by ophthalmodynamometry (▲) and intracranial cerebrospinal fluid pressure (CSFP — —) above certain levels. Below these CSFP levels the pressures are constant. VPP is the minimum intraocular pressure (IOP) required to induce retinal vein pulsation. Lines of best fit are included. The difference between ONSAS and RLTP occurs across the pia mater (pia) and the difference between intracranial CSFP and ONSAS pressure is probably due to orbital pressure (orbit).
the pia mater reduces and the RLTP tends to equilibrate directly with the ONSAS. When the intracranial pressure is low there are likely to be two mechanisms that buffer the RLTP: the orbital tissue pressure tending to compress the ONSAS and the elasticity of the pia mater.

**PRESSURE DIFFERENCE AND PRESSURE GRADIENT ACROSS THE LAMINA CRIBROSAA**

Tissue pressure measurements across the optic nerve into the retrolaminar region demonstrate a pressure gradient across the lamina cribrosa (figure 3).25 31 In the dog, having an IOP of 15 mm Hg and RLTP of 3 mm Hg at CSFP 0 mm Hg with a lamina thickness of 531 μm, the average translaminar pressure gradient (TLPG) is 23 mm Hg/mm tissue and this doubles to 47 mm Hg/mm tissue and this doubles to 47 mm Hg/mm tissue as the IOP rises to 28 mm Hg.32 33 In the human, the posterior (collagenous) lamina thickness measures mean 341 μm compared with 339 μm for the combined glial and collagenous lamina. The normal human TLPG is estimated at being between 20 mm Hg/mm and 33 mm Hg/mm tissue, with the true value likely being closer to 33 mm Hg/mm given indirect evidence that the gradient exists across the collagenous lamina.32 33

Forces which act upon tissues are caused directly by pressure gradients and so the magnitude of force and its effect upon a tissue can be minimised by reducing the gradient.34 Practically speaking this means that a thicker lamina cribrosa can spread out the gradient over a greater distance and hence reduce the shear stress acting across this tissue. The corollary is that a thinner lamina cribrosa amplifies the pressure gradient. Patients with established glaucoma have lamina thinning and tend to progress at a greater rate for a given IOP.35

**PHYSIOLOGICAL FUNCTIONS ALTERED BY PRESSURE GRADIENTS**

In primate models of raised IOP and CSFP, orthograde axonal transport and retrograde axonal transport are inhibited at the level of the lamina cribrosa with accumulation of intracellular material including mitochondria and other vesicles. Axonal transport accumulation occurs mainly at the posterior lamina region in animal glaucoma models.36 and is inhibited in rabbit vagus nerve when the pressure difference exceeds 30 mm Hg and the gradient exceeds 45 mm Hg/mm.37 It has not been shown experimentally whether the pressure difference or gradient has greatest effect upon axonal transport.

When the IOP is raised leading to an elevated translaminar pressure gradient, significant neurofilament changes occur within several hours followed by changes in the microtubules which are structures vital for axonal transport conductance.38 These cytoskeletal changes are likely to be fundamental to the cause of the axonal transport changes. Cytochrome oxidase activity also increases in the collagenous lamina region with increased TLPG indicating increased metabolic requirements in this region.39

The central retinal vein traverses the lamina cribrosa and is subject to the translaminar pressure gradient partially because the venous transmural pressure is close to zero.40 41 42 meaning that the surrounding tissue pressure generally affects the pressure within the vessel lumen. It is likely that the pressure gradient falls along the central retinal vein as it traverses the lamina cribrosa. Assuming such, a gradient of 33 mm Hg/mm is much higher than the typical 0.7 mm Hg/mm found in venules.43 Venous endothelial cells within the posterior (collagenous) laminar region are closer, morphologically speaking, to arterial endothelial cells than to other venous endothelial cells (figure 4), demonstrating that they are subject to higher shear stresses as a result of the pressure gradient.33 44 It is possible that these shear stresses and morphological changes are exacerbated when the pressure gradient is increased in glaucoma.

**INTERACTIONS WITHIN THIS REGION**

IOP and CSFP are pulsatile, being influenced by the cardiac cycle with CSFP having an additional influence from the respiratory cycle (figure 3). IOP and CSFP pulsatility affect optic nerve tissue pressure pulsatility (figure 3). One important clinical observation is the phenomenon of spontaneous venous pulsation close to the central retinal vein exit into the optic nerve, found in 95% of normal subjects. We have found that venous pulsation phase is in time with the IOP and CSFP phase.15 Classical theories had taught that pulsation would be counter phase to IOP and that the systolic peak of IOP that would lead to venous compression,45 46 however recent work suggests that CSFP pulse drives venous pulse timing and so has a more dominant effect upon key venous haemodynamic properties.12 The reasons for spontaneous venous pulsation are complicated and probably relate largely to the eye resembling a Starling resistor.47 Monkey and dog experiments demonstrate that an average 8 mm Hg pressure difference between IOP and CSFP is required for the induction of venous pulsation.48–50

**ABNORMAL ANATOMY**

**Enlargement of the ONSAS**

The ONSAS width is increased in many patients with raised CSFP. This is a significant relationship and has been used to estimate intracranial pressure non-invasively with several algorithms being published.51 52 However, enlargement of the ONSAS may be due to other factors such as loculation of CSF within the ONSAS. These may be part of the explanation for the observation that ONSAS width is increased in some patients with normal tension glaucoma.

**Myopia**

Persons with high myopia tend to have a thin lamina cribrosa (207 μm) compared with a normal 458 μm. Lamina thickness decreases in patients with moderately advanced glaucoma (201 μm) and patients with myopic glaucoma (78 μm).53 Such reductions in laminar thickness will double or quadruple the gradient effect of the pressure difference. This is likely to influence progression rates of glaucoma and may explain why patients with severe glaucoma and myopia tend to progress even at lower IOP levels.33 54 55 Persons with myopia also have an increased width of the ONSAS termination and a thin posterior sclera. Recent optical coherence tomography (OCT) scanning using swept source demonstrates that the posterior staphylomata frequently seen in persons with high myopia tend to overlie the region of the enlarged ONSAS termination (figure 5).16 The presumed reason for the staphyloma is the pressure difference across the thinned sclera between the IOP compartment and the ONSAS compartment leading to posterior distortion of the sclera just as posterior distortion of the lamina cribrosa occurs in glaucoma. The smaller degrees of optic disc excavation seen in persons with myopia with glaucoma may in part be due to the fact that the gradient is occurring across and causing distortion of the sclera and lamina cribrosa.

**Possible septal changes in ONSAS**

In patients with idiopathic intracranial hypertension and normal tension glaucoma it has been observed that there is a lower flow of CSF from the intracranial compartment along the ONSAS and this may be due to thickened trabeculae leading to increased
flow resistance along the ONSAS. This reduced flow may have metabolic consequences by leading to stagnation and reduced removal of toxic metabolites as well as reduced delivery of necessary metabolites to regions, particularly near the termination of the ONSAS. It also may impact upon functioning of the lymphatic and neural tissues within this region.

Raised translaminar pressure gradient
The TLPG is most simply described by the difference between IOP and RLTP divided by the distance between the two pressure compartments. Physiologically speaking this means the difference between IOP and ONSAS pressure, allowing for the buffering effects of orbital pressure and the pia mater divided by the lamina cribrosa thickness. One must also take into account changes in posture if one is using CSFP measurements from the lumbar spine. The caveats concerning general assumptions that the pressures are equivalent between the intracranial compartment, lumbar spine and ONSAS must be considered and weighed carefully at each set of observations. So, an increased TLPG can theoretically be caused by increased IOP and reduced CSFP and also by reduced buffering, that is, a lower orbital tissue pressure and reduced elasticity of the pia mater.

Figure 3  Micropipette tissue pressure measurements in dog (B) as the micropipette is passed from the vitreous region, the prelaminar region, the lamina and into the retrolaminar optic nerve. Oscillations from the respiratory and cardiac cycles can be seen. Intraocular pressure (A), cerebrospinal fluid pressure (C) and blood pressure (D). Lamina cribrosa was from 558 μm to 919 μm.

Figure 4  Central retinal arterial (CRA) (A–C) and central retinal venous (CRV) (D–G) endothelial morphology. Confocal microscope images and schematic outlines demonstrate endothelial morphology in the prelaminar, anterior lamina cribrosa, posterior lamina cribrosa and retrolaminar regions. In the posterior lamina cribrosa region, venous endothelial cells (F) appeared similar to the arterial endothelium displaying spindle-shaped morphology. Scale bar, 50 μm.
monkey model demonstrates glaucoma-like changes to the nerve. Prostaglandin analogues appear to hasten orbital fat atrophy. It is of some concern that the RL TP and exacerbate the TLPG. It is of some concern that the pressures around the optic nerve, all of which would tend to lower changes may occur leading to reduced pressure buffering properties. As one ages the IOP tends to rise gradually and CSFP tends to drop slightly. Orbital tissues tend to atrophy with age leading to a rise in CSFP are well known causes of papilloedema. Reduced or reversed translaminar pressure gradient Idiopathic intracranial hypertension and other disorders that lead to a rise in CSFP are well known causes of papilloedema.

Reduced or reversed translaminar pressure gradient

Idiopathic intracranial hypertension and other disorders that lead to a rise in CSFP are well known causes of papilloedema.58 59 Additionally, recent work using a chronic low CSFP elevation may be a risk factor for diabetic retinopathy development.23

Figure 5 Swept source OCT of eye with mild glaucoma (A) having a collagenous lamina cribrosa (LC) thickness approximating 300 μm. Some retrolaminar tissue (RLT) is shown. The choroid (C) and sclera (S) are well developed. In the highly myopic eye the sclera (S) is thin, particularly under the posterior staphyloma region, which overlies the large optic nerve subarachnoid space (ONSAS). One trabecula (T) is clearly seen. The collagenous LC is very thin and the RLT is clearly seen. Scale bars shown.

Prolonged space flight

Prolonged space flight by astronauts in space for more than 6 months frequently (20%) leads to a clinical syndrome bearing marked similarities to idiopathic intracranial hypertension with papilloedema, also some cotton wool spots as well as choroidal folds and a hyperopic refractive shift.57 MRI scans show elevated optic disks with an anteriorly shifted macula and dilated ONSAS.68 These subjects have increased intracranial pressure, which persists for months and years after their return to earth. The explanation for this phenomenon is likely to involve the fact that in microgravity there is a cephalad fluid shift with increased venous pressures and impaired lymphatic flow around the head.69 70 Both factors may conspire to impede CSF outflow and lead to an increased average CSFP which is not reduced or ameliorated by postural changes, that is, by standing. It is difficult to explain why the elevated CSFP persists for so long upon returning to earth but the most likely explanation involves a permanent change in the outflow system, either involving the arachnoidal granulations or the lymphatic outflow apparatus around the olfactory bulb, optic nerve and other regions. It is possible that in space with CSF stagnation along the ONSAS, toxic changes are occurring within the anterior optic nerve involving lymphatics leading to some permanent change in CSF outflow in this region.

CONCLUSION

We hope to have demonstrated that the influence of CSFP upon the eye and in particular the anterior optic nerve is significant and relevant to common ocular diseases. There is a rapidly growing body of literature reporting significant effects of CSFP upon glaucoma as well as diseases causing papilloedema and other disorders like diabetes. A simplistic understanding incorporating only CSFP and IOP without considering lamina cribrosa properties, orbital tissue, pia mater and subarachnoid space properties is unlikely to give a complete understanding of these
common disorders. Some of our understanding is hindered by the lack of an accurate non-invasive measurement method for CSFP.

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