Cerebrospinal fluid dynamics between the intracranial and the subarachnoid space of the optic nerve. Is it always bidirectional?


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CSF is thought to flow continuously from the site of production in the ventricles into interconnected spaces; i.e. cisterns and subarachnoid spaces (SASs). Since the SAS of the optic nerve is defined by a cul-de-sac anatomy, it is not evident how local CSF might recycle from that region to the general SAS. The concept of free communication of CSF has recently been challenged by the description of a concentration gradient of beta-trace protein, a lipocalin-like prostaglandin D-synthase (L-PGDS), between the spinal CSF and that in the SAS of the optic nerve, indicating diminished local clearance or local overproduction of L-PGDS here. In fact, computed cisternography with a contrast agent in three patients with idiopathic intracranial hypertension and asymmetric papilloedema demonstrate a lack of contrast-loaded CSF in the SAS of the optic nerve despite it being present in the intracranial SAS, thus suggesting compartmentation of the SAS of the optic nerve. The concept of an optic nerve compartment syndrome is further supported by a concentration gradient of brain-derived L-PGDS between the spinal CSF and the CSF from the optic nerve SAS in the same patients.

Keywords: beta-trace protein; cerebrospinal fluid dynamics; computed cisternography; idiopathic intracranial hypertension; optic nerve subarachnoid space

Abbreviations: ANTG = atypical normal tension glaucoma; CLCSF = contrast-loaded cerebrospinal fluid; ICP = intracranial pressure; IIH = idiopathic intracranial hypertension; L-PGDS = lipocalin-like prostaglandin D-synthase; SAS = subarachnoid space


Introduction

CSF is thought to be fairly homogeneous in composition and to be distributed evenly with a continuous flow through all CSF spaces, such as ventricles, cisterns and subarachnoid spaces (SASs) including the SAS of the optic nerve (Fig. 1). It is generally agreed that there is a bulk circulation from the site of origin to the site of absorption; i.e. from the ventricles to the arachnoid villi in the cranial SAS (Dichiro, 1964; Bito and Davson, 1966; Milohart, 1972; Wood, 1982). CSF circulation and direction of flow in the large CSF spaces (ventricles and spinal CSF space) have been studied with radiocisternography and other tracers. (Dichiro, 1966; Greitz et al., 1992; Greitz and Hannerz, 1996). The mechanism by which CSF is propelled on its circulatory route is not fully understood but probably is influenced by the outpouring of newly produced CSF, postural effects, ventricular pulsations, the pulse pressure of the vascular choroid plexus, and a piston action of the brain (Bito and Davson, 1966; Milohart, 1972; Greitz et al., 1991, 1992). CSF is renewed within 24 h, thus indicating a fast turnover. The velocity of CSF can be calculated using its total volume \( V_{\text{tot}} \) and the recycle time \( R_t \) as follows:

\[
\text{velocity} = \frac{V_{\text{tot}}}{R_t}
\]

In addition to the well-established concept of CSF resorption in the arachnoid villi, in vivo research in animals strongly supports the concept of resorption via lymphatics (Johnston, 2000, 2003; Zakharov et al., 2003). Indeed, lymphatics have been demonstrated on light and transmission electron microscopy in human tissue, and Indian ink injected into the SAS of the optic
From intracranial (SAS to the orbital SAS. Gradient the direction of flow is directed from the intracranial characterized by small trabeculae (broad septae (surrounding the optic nerve (orbital CSF space). The optic chiasm (intracranial CSF space). CSF flows from intracranial (A) into the SAS of the optic nerve (B). The SAS of the optic nerve is most narrow in the canalicular region (C). The intraorbital segment of the SAS is characterized by broad septae (D), whereas the retrobulbar segment is characterized by small trabeculae (E). Due to the CSF volume gradient the direction of flow is directed from the intracranial SAS to the orbital SAS.

Nerve subsequently appears in dural lymphatics (Killer et al., 1999).

The concept of a homogeneous CSF composition has been challenged based on the measurement of a marked concentration gradient of the brain-derived protein lipocalin-like prostaglandin D-synthase (L-PGDS) between the spinal CSF and that in the SAS surrounding the optic nerve in patients with idiopathic intracranial hypertension (IIH), optic nerve sheath meningioma and non-arteritic anterior ischaemic optic neuropathy (NAION) (Killer et al., 2006). This concentration gradient does however not elucidate the CSF dynamics between the intracranial optic nerve SAS, nor does it explain how the CSF in the SAS of the optic nerve could recycle back to the intracranial site of resorption, apparently in violation of the laws of hydrodynamics.

Persistent papilloedema and visual loss in patients with IIH despite a functioning CSF shunt—a phenomenon that may be more common than generally appreciated—poses a conundrum that cannot be explained by the current concept of free CSF circulation throughout all CSF spaces (Guy et al., 1990; Kelman et al., 1991; Ramsey et al., 2006). Compartmentation of the SAS of the optic nerve providing a barrier to free flow of CSF and the nerve offers a possible pathophysiological explanation for this phenomenon.

**Material and methods**

Three patients with IIH diagnosed by previous MR imaging of the brain and the orbits and lumbar puncture (LP) underwent complete neurological and ophthalmological examination. In all patients, bilateral asymmetric papilloedema was present on fundoscopy. All patients underwent CT-cisternography after intrathecal injection of 10 ml of contrast medium (Iopamidol, molecular weight 778 Da) by LP (Mironov et al., 1993).

CT-cisternography was performed in the same room as LP and contrast application. The time interval between contrast application and CT-cisternography measured on average 2–5 min.

In order to compare the results from patients with elevated intracranial pressure (ICP) with those from patients with normal ICP, we also performed CT-cisternography in two patients who were already enrolled in another study of atypical normal tension glaucoma (ANTG).

In all patients, CT-cisternography was performed while the patients were positioned on their knees and elbows. They were then asked to turn to the left and to the right with their heads facing the floor. In order to elevate the ICP, the patients were instructed to perform a Valsalva manoeuvre. In all patients CT-cisternography was performed prior to optic nerve sheath fenestration (ONSF).

CSF was obtained during the procedure and subsequently analysed for cells, glucose, immunoglobulin G antibodies, albumin and L-PGDS. In all patients (IIH and ANTG) optic nerve sheath decompression was performed under general anaesthesia via a medial transconjunctival orbitotomy. Special care was taken not to use fluids to moisten the cornea after the orbitotomy in order not to dilute the CSF. Multiple incisions were performed in the dura of the optic nerve with a 19-gauge blade, and the trabeculae in the SAS were loosened with a tenotomy hook. After the first incision, a gush of CSF was observed, followed by further slow outflow. CSF then was sampled with a syringe using a 37-gauge needle (Killer et al., 2006).

**Results**

No intracranial lesions were present in any of the patients on MRI; however, in the patients with IIH, orbital MRI revealed a variable degree of distension of the optic nerve sheath, most prominent in the bulbar segment adjacent to the posterior sclera (Figs 2–4). The amount of CSF surrounding the optic nerve was best seen on axial and coronal T2-weighted sequences. Flattening of the posterior sclera with protrusion of the optic disc into the vitreous body was present in all patients with IIH to a variable degree. CT-cisternography in the IIH patients demonstrated an impaired contrast-loaded CSF (CLCSF) communication between the intracranial SAS and the SAS of the optic nerve. In two patients with IIH, CLCSF entered the SAS of the optic nerve only within the optic canal, not in the SAS of the intraorbital optic nerve, suggesting blockage of CLCSF within the canalicular portion of the optic nerve (Figs 2 and 3). In another patient with IIH, CLCSF entered the SAS of the optic nerve up to 10 mm within the orbit on the right side, whereas CLCSF on the left side entered ~3 mm (Fig. 4).

In one patient from the ANTG study group, CLCSF reached the mid-orbital portion of the optic nerve on both sides. A discrete CLCSF signal could be demonstrated only on the nasal side of the bulbar region of the right optic nerve, whereas no CLCSF reached the bulbar segment of
the left optic nerve (Fig. 5). In another ANTG patient, the CLCSF demonstrated a modest signal on the right with a patchy distribution and no CLCSF on the left (Fig. 5). In contrast to the patients with elevated ICP, an inversion of the L-PGDS ratio (L-PGDS optic nerve/L-PGDS LP) was measured (Table 1).

Discussion

In contrast to all other cranial nerves, the optic nerve is—to start with—by histological definition not really a nerve, but a white-matter tract of the CNS that extends into the orbit where it is surrounded by CSF throughout its entire length. As a consequence of this unique anatomy, the optic nerve can become involved in CNS disorders characterized by elevated ICP; e.g. IIH, intracranial masses, and inflammatory and infectious disease. The cul-de-sac structure of the covering of the optic nerve is the anatomical hallmark of the optic nerve and may be the crucial issue in understanding the CSF dynamics between the intracranial SAS and the SAS of the optic nerve. Free bidirectional communication between these areas has previously been taken for granted; however, this concept has recently been challenged by the description of a marked concentration gradient of L-PGDS (beta-trace protein) between the spinal CSF and the CSF in the SAS of the optic nerve that was measured in patients.
Table 1  Clinical characteristics of the patients studied

<table>
<thead>
<tr>
<th>Patient</th>
<th>C.M.</th>
<th>T.Z.</th>
<th>M.F.</th>
<th>H.E.</th>
<th>N.E.</th>
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<tr>
<td>Age (years)</td>
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<td>30</td>
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<td>IIH</td>
<td>IIH</td>
<td>ANTG</td>
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</tr>
<tr>
<td>Side</td>
<td>R</td>
<td>L</td>
<td>L</td>
<td>R</td>
<td>R</td>
</tr>
<tr>
<td>BMI (kg/m²)</td>
<td>31</td>
<td>34</td>
<td>35</td>
<td>22</td>
<td>25</td>
</tr>
<tr>
<td>LP pressure (cm H₂O)</td>
<td>30</td>
<td>31</td>
<td>30</td>
<td>18</td>
<td>16</td>
</tr>
<tr>
<td>L-PGDS LP (mg/l)</td>
<td>32.20</td>
<td>18.00</td>
<td>17.00</td>
<td>73.20</td>
<td>34.10</td>
</tr>
<tr>
<td>L-PGDS ON (mg/l)</td>
<td>70.80</td>
<td>132.00</td>
<td>47.80</td>
<td>27.00</td>
<td>30.60</td>
</tr>
<tr>
<td>L-PGDS ratio ON/LP (%)</td>
<td>220</td>
<td>733</td>
<td>281</td>
<td>37</td>
<td>90</td>
</tr>
</tbody>
</table>

M = male, F = female, L = left, R = right. ANTG = atypical normal tension glaucoma; BMI = body mass index; IIH = idiopathic intracranial hypertension (pseudotumour cerebri); LP = lumbar puncture; L-PGDS = lipocalin-like prostaglandin D-synthase (beta-trace); ON = optic nerve.

with IIH, NAION, and optic nerve sheath meningioma (Killer et al., 2006). Whether or not this concentration gradient of L-PGDS is due to a local surplus production of CSF or a lack of clearance from the SAS of the optic nerve is not yet understood. Although orbital MR imaging in patients with IIH demonstrates changes consistent with CSF congestion and bulging of the meningeal sheath, it does not provide either qualitative or quantitative information regarding the biochemical composition of the local CSF, nor does the appearance of CSF in the SAS of the optic nerve differ from that of the intracranial CSF. Furthermore, there is no hydrodynamic theory that would explain how the cranial CSF that enters the SAS of the optic nerve could change its direction of flow against the volume gradient that directs it from the site of production and higher volume (intracranially) towards the SAS of the optic nerve. Two possible outflow routes from the SAS of the optic nerve have been suggested. Outflow from the SAS of the distal portion of the optic nerve into the orbit was demonstrated in animal studies via leakage of contrast agent and radioisotopes (Weed, 1914; Field and Brierly, 1949; McComb et al., 1982; Shen et al., 1985; Fogt et al., 2004; Lüdemann et al., 2005). Drainage via lymphatics is a new powerful concept that has demonstrated a great capacity for CSF outflow from the CNS (Johnston, 2000, 2003). Lymphatics in the dura of the human optic nerve may also offer an outflow pathway (Gausas et al., 1999; Killer et al., 1999).

The possibility of impaired exchange between the intracranial CSF and that in the SAS of optic nerve has only recently become of interest because of reports about persistent papilloedema and visual loss in patients with IIH despite a functioning lumboperitoneal shunt (Kelman et al., 1992; Huna-Baron et al., 2003b). Considering the current understanding of CSF communication, patients with IIH without papilloedema are even more intriguing than asymmetric and unilateral papilloedema (Lipton and Michelson, 1972; Seggia and De Menezes, 1993). Although papilloedema is defined as disc swelling due to elevated ICP (Hayreh, 1968), typical radiological and fundoscopy features have been reported in patients without elevated ICP that presented with swollen discs resembling papilloedema due to retrobulbar neuritis and arachnoid cysts (Gass et al., 1996; Brodsky and Vaphiades, 1998; Killer and Flammer, 2001; Killer et al., 2003b; Hickman et al., 2005). Partial or total compartmentation of the SAS of the optic nerve would offer a pathophysiological explanation for such cases; however, other than biological evidence (L-PGDS concentration ratio), there have been no data to support the concept of an optic nerve compartment syndrome.

One might expect studies of CSF dynamics in vivo to provide information about the location of impeded CSF flow and probably about mechanisms resulting in impaired flow. However, although radionuclide cisternography is often used to provide information regarding CSF flow, the SAS of the optic nerve is too small for this technique to render reliable information about the CSF in this area. It is for this reason that we have used CT-cisternography with contrast. This technique has been used in the past for the diagnosis of CSF leaks in patients with fractures of the skull base (Mironov et al., 1993). The molecular size of the contrast agent (Iopamidol; 778 Da) allows easy access to the SAS of the optic nerve (molecular weight of L-PGDS 28,000 Da). In the present study, CT-cisternography with contrast demonstrated blockage (stasis) and impaired influx of CLCSF in patients with IIH as well as to some extent in the ANTG study group patients. The information from this technique, combined with the finding of L-PGDS concentration gradients, provides strong evidence for optic nerve sheath compartmentation in patients with IIH.

If a total block of the CSF occurs in some situations, it is feasible to assume that a partial block may occur in other conditions. An interesting candidate is ANTG. In glaucomatous optic neuropathy the astrocytes are activated. This leads to an upregulation of factors such as matrix...
metallo–proteinases, tumour necrosis factor α or endothelin. While on one hand, these factors may lead to arachnoiditis and thereby slow down CSF renewal, an increased concentration of these factors in the CSF on the other hand may contribute to the optic nerve damage. Endothelin, for example, is known to be increased in glaucoma patients—both locally and systemically. It not only reduces blood flow in the microcirculation but also interferes with the axoplasmatic transport (Hernández, 2000). Ongoing studies will clarify whether such physiological molecules reach local concentrations that may be toxic to the optic nerve.

The development of compartmentation may depend on several pathophysiological mechanisms. The smallest diameter of the SAS of the optic nerve is within the optic canal. Although the osseous component of the canicular portion of the optic nerve could theoretically play a role in the development of compartmentation, most patients with fibrous dysplasia retain normal vision in spite of narrowing of the optic canal (Lee et al., 2002). It seems therefore more likely that the site of the pathophysiology is within the dura mater and the arachnoid and its adherent structures, including the trabeculae and septae (Killer et al., 1999, 2003a; Sens et al., 2003). Inflammatory changes in the SAS of the optic nerve leading to distension of the sheath and subsequent disc swelling has been suggested in patients with anterior optic neuritis (Killer et al., 2003b; Hickman et al., 2005). Anatomical studies of the SAS in post-mortem specimens from patients without neurological disease demonstrate polymorphonuclear leucocytes attached to the arachnoid layer (Killer et al., 2003a). In addition, studies in rats demonstrate large numbers of MHC type II cells on the arachnoid in the SAS (Braun et al., 1993). Leucocytes, macrophages, lymphoblasts and monocytes were detected following injection with bacillus Calmette-Guerin (Merchant and Low, 1977). It seems therefore feasible that even mild inflammatory stimuli may produce arachnoiditis and trabeculitis with secondary fibrosis, eventually leading to an optic nerve sheath compartment syndrome. The same process may occur in patients with subarachnoid haemorrhage (Juwul et al., 1979). Inflammatory processes might also contribute to the closing of the arachnoid apertures that drain CSF into the meningeal lymphatics thus adding to the CSF compartmentation.

Our study using CT-cisternography provides a pathophysiological explanation—optic nerve compartment syndrome—for persistent papilloedema and progressive visual loss in patients with apparently functioning CSF shunts as well as the occurrence of unilateral or asymmetric papilloedema. At the same time, our findings raise questions concerning the biological effects of accumulated CSF and its components on the optic nerve following the development of this syndrome. A possible toxic effect of reduced CSF clearance on brain tissue has recently been postulated (Rubenstein, 1998; Silverberg et al., 2003). As the biochemical effects of L-PGDS are several, including neuroprotection of astrocytes on one hand and apoptotic activity and modulation of inflammatory processes on the other, it is to be expected that high concentrations of this agent will have a significant effect on the optic nerve (Logdberg and Wester, 2000; Govoni et al., 2001; Ragolia et al., 2001, 2003; Taniike et al., 2002; Kagitani-Shimono et al., 2006). The immediate retrobulbar portion of the nerve—from where CSF was sampled during sheath decompression—may be particularly vulnerable to these toxic effects. Histochemical and immunocytochemical studies have demonstrated a striking inverse relationship between myelination and mitochondrial distribution with the highest concentration of unprotected mitochondria in the bulbar region of the optic nerve (Bristow et al., 2002). Damage to the mitochondria in this region—and to the pial vasculature—may contribute to the loss of visual function in IIH as well as other types of optic nerve disease. The crucial role of mitochondria for an intact optic nerve becomes evident from patients with Leber’s optic neuropathy (Biousse and Newman, 2001).

In conclusion, understanding unilateral and asymmetric papilloedema in patients with elevated ICP is not possible without focusing on the SAS of the optic nerve. Damage in this setting can only be understood in the context of the close relationship of the optic nerve with the surrounding CSF and its proximity to the axons, mitochondria and the pia septal blood supply. We believe that the best explanation for both (unilateral and asymmetric papilloedema in patients with elevated ICP) is that the CSF in the SAS of the optic nerve may become sequestered, thus producing a compartment syndrome. The lack of CLCSF in the SAS of the optic nerve found in this study combined with the marked concentration gradient of a mainly brain-derived component—L-PGDS—between the spinal CSF and the SAS of the involved optic nerve strongly supports this concept of an optic nerve sheath compartment containing highly biological active molecules.

Based on the concentration gradient of L-PGDS and the corresponding CT-cisternography studies there is strong evidence for compartmentation of the SAS of the optic nerve. As there are at present no data on the normal population, the link between compartmentation and disease of the optic nerve is therefore based on the analogy of disturbed CSF dynamics and disease of the CNS (Rubenstein, 1998; Silverberg et al., 2003).

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References


