

# Hydrocephalus Revisited: New Insights into Dynamics of Neurofluids on Macro- and Microscales

Hans C. Ludwig<sup>1</sup>  Hans C. Bock<sup>1</sup>  Jutta Gärtner<sup>2</sup>  Stina Schiller<sup>2</sup> Jens Frahm<sup>3</sup>   
Steffi Dreha-Kulaczewski<sup>2</sup> 

<sup>1</sup> Division of Pediatric Neurosurgery, Department of Neurosurgery, University Medical Center Göttingen, Göttingen, Germany

<sup>2</sup> Division of Pediatric Neurology, Department of Pediatrics and Adolescent Medicine, University Medical Center Göttingen, Göttingen, Germany

<sup>3</sup> Biomedical NMR, Max Planck Institute for Biophysical Chemistry, Göttingen, Germany

**Address for correspondence** Hans C. Ludwig, MD, Division of Pediatric Neurosurgery, Department of Neurosurgery, University Medical Center Göttingen, Robert-Koch-Str. 40, 37075 Göttingen, Germany (e-mail: hludwig@med.uni-goettingen.de).

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## Abstract

### Keywords

- cerebrospinal fluid
- real-time MRI
- hydrocephalus
- space flight disease
- aquaporin
- spontaneous intracranial hypotension
- neural tube defect

New experimental and clinical findings question the historic view of hydrocephalus and its 100-year-old classification. In particular, real-time magnetic resonance imaging (MRI) evaluation of cerebrospinal fluid (CSF) flow and detailed insights into brain water regulation on the molecular scale indicate the existence of at least three main mechanisms that determine the dynamics of neurofluids: (1) inspiration is a major driving force; (2) adequate filling of brain ventricles by balanced CSF upsurge is sensed by cilia; and (3) the perivascular glial network connects the ependymal surface to the pericapillary Virchow–Robin spaces. Hitherto, these aspects have not been considered a common physiologic framework, improving knowledge and therapy for severe disorders of normal-pressure and posthemorrhagic hydrocephalus, spontaneous intracranial hypotension, and spaceflight disease.

## Introduction

### Challenging the Historic View

Brain disorders arising from disturbed cerebrospinal fluid (CSF) circulation constitute a major proportion of human diseases. Incidences of 49.5/100,000 have been found for prevalent forms such as isolated hydrocephalus (HC) and as high as 81.2/100,000 for HC related to prematurity.<sup>1</sup> In fact, children represent the majority of patients affected by HC, often in conjunction with congenital neural tube defects, brain malformations, and posthemorrhagic hydrocephalus (PHHC). As a consequence, HC is the most frequent reason for brain surgery during childhood. Epidemiologic studies demonstrate the influence of economic status; in high-income countries, children mostly suffer from PHHC, while in low- and medium-income countries, HC predominantly emerges as a sequela of neural tube defects and congenital anomalies.<sup>2</sup> On the other

hand, reflecting the demographic shift with the increasing incidence of neurodegenerative disorders, there has been an increase in elderly patients diagnosed with idiopathic normal-pressure hydrocephalus (iNPH) with a mortality rate of more than 80% over 5 years when untreated.<sup>3</sup>

Our current pathophysiologic concept of HC is largely based on the hypothesis of bulk flow and Dandy's classification of communicating and obstructive HC derived from a small number of animal experiments.<sup>4</sup> Decades later, the hydrodynamic theory was added by Greitz,<sup>5,6</sup> demonstrating pulsatile flow from the ventricles through the aqueduct. Klarica et al found aqueductal flow in cats to be close to zero over a period of 2 hours after open neurosurgical access to the aqueduct.<sup>7</sup> These results contradicted earlier assumptions of unidirectional CSF routes from the choroid plexus via extracerebral cisterns to pacchionian granulations, spinal

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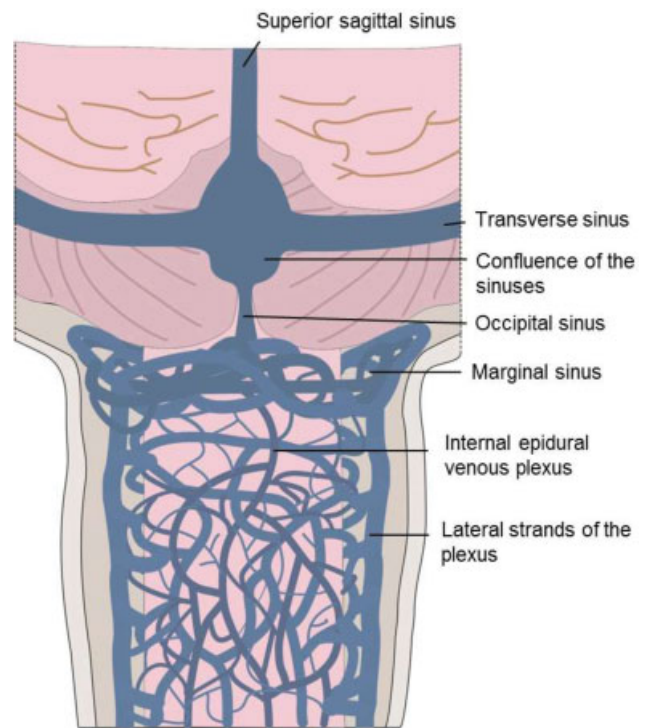
dural nerve roots, and cranial nerve arachnoid resorption. In later studies,<sup>8,9</sup> the same authors concluded CSF movements to be “random” without directional preference, driven by cardiac action and respiration. Brinker et al<sup>10</sup> treated CSF dynamics as a diffusion process that leads to passive mixing and distribution into the different ventricular and subarachnoid compartments, while Buishas et al postulated CSF exchange by osmosis.<sup>11</sup>

Recent findings challenge these views about CSF circulation and call for a physiologic concept that takes the interplay of macroscopic fluid flow and microscopic drainage pathways into account.<sup>12</sup> New insights particularly apply to (1) the discovery of mainly respiration-induced CSF dynamics by real-time magnetic resonance imaging (MRI),<sup>13,14</sup> (2) novel details about the regulatory cascades of cerebral aquaporin water channels,<sup>15</sup> and (3) increasing clinical expertise regarding disorders of perturbed CSF circulation.<sup>16</sup> This review brings together the current thoughts on the multicompartamental dynamics of neurofluids to foster a better understanding of CSF disorders such as HC accompanying Chiari malformations, rhinoliquorrhea, and spontaneous intracranial hypotension (SIH). Moreover, aspects about cerebral fluid dynamics in zero gravity are included. The important findings about changes of brain volumes during space flights just recently been published are perfectly in line with the concept of HC discussed here. Hence, their discussion will contribute greatly to the understanding of the new point of view.

### Breathing, CSF Dynamics, and Venous Flow

Noninvasive flow MRI techniques play a pivotal role in the research of human HC because they offer quantitative evaluations of both CSF movement and blood flow.<sup>17</sup> Extending conventional techniques, real-time MRI developments now allow for dynamic assessments of the functional anatomy and fluid flow at high spatial and temporal resolution.<sup>13</sup> Most importantly, real-time MRI studies may be performed without the need for physiologic gating. Physiologic gating to date affects almost all flow MRI studies by restricting the results to processes governed by cardiac periodicity. Without such prejudice, real-time MRI has facilitated the discovery of respiration-induced CSF transport<sup>14</sup> and elucidated the tightly connected interplay between the venous fluid systems and the observed CSF upward surge.<sup>18</sup> Similar results of a CSF flow through the aqueduct and into the ventricles could be shown by time-SLIP MRI techniques first presented by Yamada et al,<sup>19</sup> although these purely qualitative measurements preclude quantitative assessments of flow direction, velocities, and volumes. In fact, CSF movement into the head during inspiration is accompanied by downward blood flow within the veins of the head/neck region, as demonstrated in spinal epidural veins (→Fig. 1). The pooling of venous blood in large extracranial collecting veins ensures cardiac preload, which then leads to adequate systolic ejection according to the Frank-Starling law.

In contrast to that of extraspinal cervical veins, which are equipped with valves, the high-capacity system of the epidural venous plexus remains valveless.<sup>20,21</sup> Epidural veins



**Fig. 1** Coronal view of the posterior fossa with the major dural sinuses (bilateral transverse sinus and singular sagittal sinus) and their confluence. The smaller occipital sinus connects to the marginal sinus lining the foramen magnum. From here, direct connections to the epidural venous plexus of the upper spinal canal exist. Thick strands at both sides of the spinal cord form the lateral parts of the usually rope ladder-like cervical venous plexus, which can be detected by flow MRI. Abundance of connections between intracranial and intraspinal venous compartments through the craniocervical junction are modified from figures by Henriques<sup>20</sup> showing resin casts of vertebral veins.

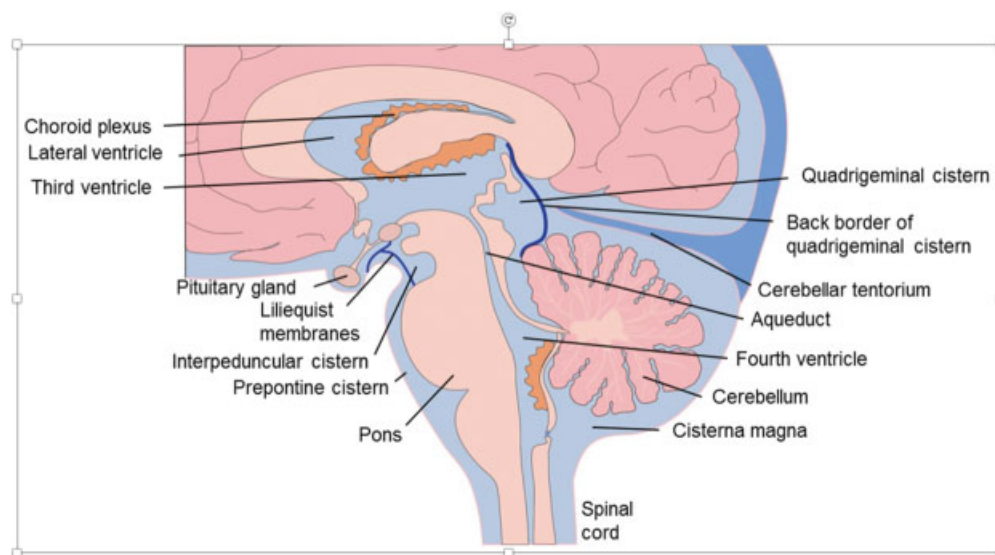
are kept open by negative pleural and epidural transmural venous pressure even under mechanical ventilation.<sup>22</sup> This well-known phenomenon can be observed during lumbar puncture when the pendant saline drop at the needle opening gets aspirated as soon as the tip reaches and penetrates into the epidural space regardless of posture.<sup>23</sup> This observation is valuable for excluding any suspected bias from CSF dynamic studies performed in the supine position. The epidural venous plexus accompanies the dural sac throughout its entire length. It forms manifold connections between intra- and extraspinal spaces and thoracic and abdominal cavities. Dreha-Kulaczewski et al<sup>18,24</sup> demonstrated the enhanced filling of spinal epidural veins during inspiration followed by draining during expiration. The respiratory-induced flow changes in this important venous drainage pathway can be transmitted into the filling and emptying of the directly connected venous compartments of the posterior fossa (→Fig. 1). Following the Monroe and Kellie doctrine of constant intracranial volume, the volume of fluid leaving the cranial vault must be counterbalanced. Thus, the upward movement of CSF and the associated venous outflow compensate for each other during respiration. There is long-standing discussion in the literature regarding the dynamic of net CSF flow, whether by cardiac pulsatility or fluid

displacement by respiration and each individual contribution. To separate cardiac- and respiratory-driven CSF motion, Takizawa et al performed asynchronous 2D phase-contrast MRI experiments under respiratory guidance. They found that for velocity, the cardiac component was greater than the respiratory component. In contrast, for the amount of volume displacement, the respiratory component was greater.<sup>25</sup> This pattern of fast and small cardiac-driven modulations but slow and high displacement by respiration was confirmed by the study of Yatsushiro et al.<sup>26</sup> Yildiz et al were able to demonstrate a comparable contribution of respiration and cardiac pulsations on CSF dynamics through the aqueduct during deep breathing but not during natural breathing.<sup>27</sup> This pattern could resemble the situation during deep non-rapid eye movement (NREM) sleep.<sup>28</sup>

From this perspective, the posterior fossa of the skull might be considered a transfer unit. Further upward, the aqueductal tube connecting the fourth and third ventricles constitutes a singular gate into the supratentorial brain CSF spaces. It might even be hypothesized that it serves as a control valve for the passage of large flow volumes during inspiration. Given that respiration-related CSF flow is continuously subjected to more rapid and low-amplitude modulating fluctuations caused by cardiac pulsatility,<sup>25</sup> the thin diameter of the aqueduct results in laminar flow and enables instantaneous control of CSF flow volumes according to the law of Hagen–Poiseuille.<sup>16</sup> It accounts for the radial diameter of the aqueduct to the fourth power, which renders the structure predestined as a fine-tuned system for CSF dynamic flow into and out of the brain ventricles.<sup>29</sup> This hypothesis is supported by the anatomic details of the aqueductal environment, in which the frontal prepontine Liliequist membrane and the dense arachnoid membranes of the dorsal cisterna quadrigemina<sup>30</sup> prevent any bypassing sub-arachnoid para-aqueductal exchange of CSF, which would

jeopardize the tuning capabilities (►Fig. 2). Any flow of CSF through the cisterns bypassing the aqueduct, as reported by studies based on experiments in rats,<sup>31</sup> would prevent aqueductal stenosis and obstructive HC in humans. The cisterna quadrigemina itself, as a protrusion of the third ventricle, might be able to promote and transmit intraventricular pressure immediately to the aqueduct, thereby influencing the tube diameter (►Fig. 2).<sup>32</sup>

A very important aspect of the tightly connected and well-balanced CSF and venous systems is to ensure upward movements of CSF against hydrostatic pressure. This physiologic principle is a prerequisite for a sustained upright body position because it maintains brain turgor and thus adequate intracranial CSF volume. Above all, hydrostatic uplift or buoyant forces inside the ventricles and arachnoidal spaces must be ensured in all body positions as well as over long periods of significant skull growth, as brain volume increases from infancy until adolescence. In view of the physiologic consequences of an upright position, findings from quadrupeds may therefore be misleading, as their aqueducts are oriented differently with respect to gravity and hydrostatic pressure. Further differences might be due to the much smaller dimensions of intracranial fluid spaces and absorption structures. Sheep, for example, are able to absorb up to 50% of their CSF volume by cervical vessel structures.<sup>33</sup> It should also be noted that the majority of experimental studies have been conducted in very young animals in which CSF structures and dynamics are not yet fully developed—comparable with infants without pacchionian granulations during the first year of life.<sup>34</sup> The exact volumes of reabsorption in adults and children above 4 years of age are still under debate. The CSF circulates at a daily rate of around 500 mL. The concepts of the reabsorption process by perivascular aquaporin channels and Virchow–Robin spaces (VRS) have to be separated from respiration- and cardiac pulsatility-induced



**Fig. 2** Sagittal view of the internal CSF spaces and main cisterns constituting external CSF spaces. The Liliequist membranes (dark blue) block the prepontine pathways of CSF (modified from Hussein et al<sup>32</sup>). The line (dark blue) outlining the quadrigeminal cistern represents the complex webs of dense arachnoid membranes hampering CSF flux. Note the unique anatomical position of this cistern. Here, variations in size or pressure may foster the modulation of aqueductal diameter, thereby regulating CSF flow into or out of brain ventricles.

dynamics. Reabsorption might be dependent more on the lymphatic system and its molecular features.

### Ependymal Fluid Flow

Ependymal cells cover the outer surface of the choroid plexus and the inner wall of the brain ventricular system as well as the central canal of the spinal cord. These unique structures can be damaged by infections, tumors, and subarachnoid hemorrhage (SAH) or intraventricular hemorrhage (IVH) due to germinal matrix disruption in prematurity. The latter is described as a bleeding-induced N-cadherin-mediated disruption of the ependymal junctions and is usually followed by severe and catastrophic destruction of the ependymal lining of the ventricular wall.<sup>35</sup> The ventricular wall encloses, protects, and organizes one of the most delicate structures of the central nervous system, the subventricular zone, also called “the lake front property.”<sup>36</sup> It harbors valuable stem cells essential for repair and plasticity until adulthood. The disturbance of this zone arising from, e.g., germinal matrix bleeding from PHHC, prematurity, chronic hypoxia, and inflammation is one of the major causes of the lifelong disability associated with periventricular leucomalacia.<sup>37</sup>

The cilia of these ependymal cells provide a flow-sensing system that represents a regulatory mechanism serving a variety of purposes. In addition to promoting active CSF transport near the ventricular walls, these cilia are capable of detecting flow directions,<sup>38</sup> turbulence patterns, and disturbances in circadian rhythms.<sup>39</sup> For example, ventricular dilation without an increase in intracranial pressure (ICP) but with perturbation of CSF circulation, as accomplished by an inflatable balloon in lambs, supports the notion of ependymal fluid flow and pressure-sensing mechanisms as being responsible for the observed alterations of ventricular dimensions.<sup>16,40</sup> Disturbances of the sensing cilia system have been identified as being responsible for alterations in ventricular morphology and dimension.<sup>41</sup> The ability of the ependyma to sense CSF flow properties in conjunction with the aqueduct provides a means to fine-tune and adapt CSF circulation for acute physiologic demands.<sup>42</sup>

Notably, pediatric HC can be evoked not only by CSF-related pathologies but also by underlying structural anomalies such as schizencephaly, aplasia, or severe cortical dysplasia and migration deficits.

### Bergmann Glia and Astrocytic Fluid Flow

A crucial process of the ventricular system during embryonic development is the formation of a CSF-filled tube. This structure consists of neuroepithelial cells arranged in tubular forms conducting fluid and precursor CSF and providing steady turgor with a continuous flow of signaling peptides to control progenitor cells building up the Bergmann glia layer.<sup>43–45</sup> Bergmann glial cells are characterized by a unique polarity extending between the ventricular wall and the glia limitans, and they mature later to astrocytic layers. The glia limitans consists of extracellular matrix laminin and fibronectin<sup>46–48</sup> contacting and surrounding the capillary wall<sup>47</sup> as well as the basal membrane of the ependyma. The astrocytic layer is essential for the brain water balancing

system because the polarized astrocytic basal membrane is equipped with aquaporin-4 (AQP4), a water channel expressed in the astrocyte foot processes and ependymal cells lining the ventricles. AQP4 facilitates the transport of excess water out of the brain.<sup>49</sup>

The correct polarization of Bergman glia and later astrocytes between the capillary wall and subependymal glia is mandatory for water homeostasis in the brain. Disturbances of this system by hemoglobin<sup>50</sup> in preterm IVH lead to deleterious lesions by destroying the N-cadherin-dependent cell junctions.<sup>35</sup> This alteration is responsible for the regional destruction of the ependymal cell layer and activation of extracellular matrix proteins around the astrocyte membrane as well as for the reported loss of polarized AQP4.<sup>51</sup> The condition is not restricted to prematurity and could also be identified in congenital and even adult HC.<sup>52,53</sup>

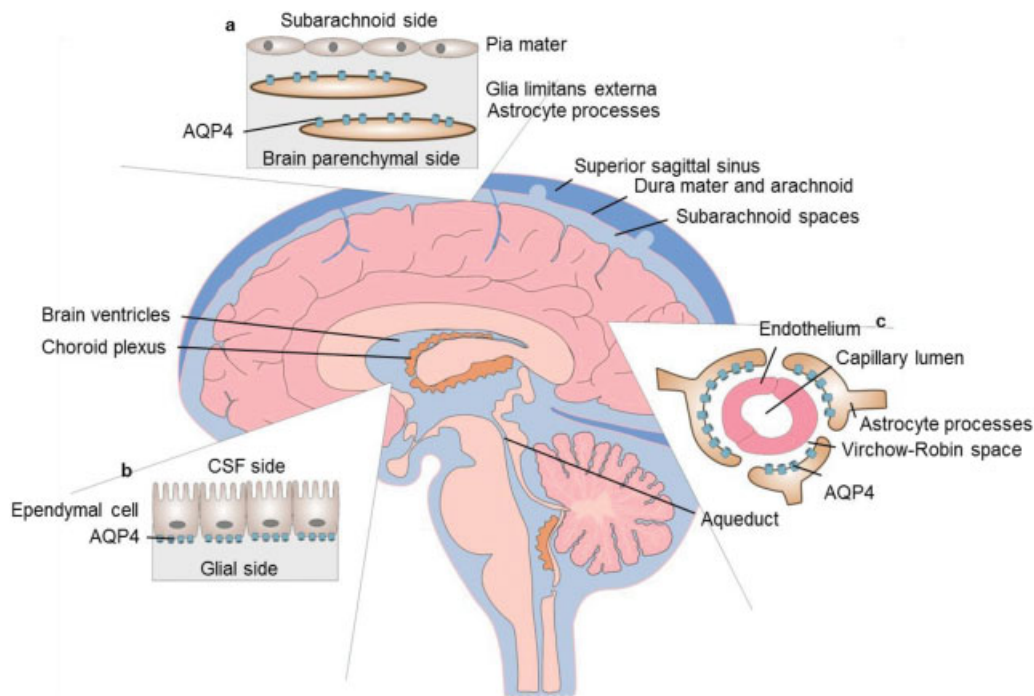
AQP4 is responsible for the recycling of brain water and cellular waste by connecting intercellular and glymphatic flow with CSF circulation. The system prevents cardiac- and respiration-related pulsatility from propagating to the interstitial fluid compartment of the brain. Exposure to those dominant forces jeopardizes the blood–brain barrier and precludes any autoregulative vasomotor control of capillary flow by the direct transduction of systemic blood pressure.<sup>48</sup> However, without an alternative hydrodynamic mechanism, the brain interstitial fluid remains stagnant. As noted by Nakada,<sup>54</sup> the AQP4-supported influx of water into the pericapillary VRS generates continuous interstitial flow, which constitutes an essential part of the clearance system for A $\beta$ -amyloid. Both adequate ventricular CSF circulation and interstitial flow in VRS are mandatory for proper neurofluid circulation (**► Fig. 3**). These interacting systems control the brain water balance under diverse conditions ranging from growth to standing and prone positions under variable systemic blood pressures as well as during physical exercise and sleep. Taken together, the combined understanding of functional neurofluids at the macroscopic, microscopic, and molecular levels may be exploited to re-evaluate the pathophysiology of conditions with disturbed CSF distribution and transport. At this point in time, aspects discussed here are still conceptual, derived from the novel *in vivo* insights facilitated by real-time flow MRI and supported by the literature.

## CSF Disorders and Beyond

### Chiari Malformations

The neural tube of the fetus usually closes around embryonic day 25. Incomplete closure results in open spina bifida or fetal meningocele (MMC), which forms a CSF fistula over the remaining months of pregnancy until surgical closure by either fetal intrauterine or postnatal surgery. A study of the effectiveness of intrauterine MMC repair demonstrated that intrauterine closure at approximately the 24th week of pregnancy significantly reduced the rate of Chiari II malformation with less tonsillar herniation and HC.<sup>55</sup> The lack of distention of the embryonic posterior fossa had already been postulated as the underlying mechanism decades ago.<sup>56</sup> After surgical closure of the MMC fistula,





**Fig. 3** Sagittal view of the brain and CSF spaces. Insets illustrate the main location of aquaporin-4 water channel (AQP4) expression (modified from Nakada et al<sup>49</sup>). Astrocytes harboring the AQP4 system in their end-feet processes are essential components of brain tissue–CSF and brain tissue–blood interfaces. (a) Astrocytes of the glia limitans externa (situated subpial) expressing AQP4 are part of the outer brain barrier between the subdural spaces and outer brain surface. (b) AQP4 is expressed on the glial/brain parenchymal side of the ependymal cell layer, which covers all internal CSF spaces. (c) Astrocyte end feet enclosing pericapillary spaces referred to as Virchow–Robin spaces (VRS) are equipped with AQP4 water channels and contribute to the blood–brain barrier. Here, the astrocytic AQP4 system regulates water influx into the VRS and generates interstitial flow.

sufficient upward movement of CSF ensures appropriate ventricular filling and growth of the posterior fossa. Remarkably, at approximately the 24th gestational week, fetal respiration emerges.<sup>57</sup> Assuming the same physiologic principles as those postnatally, the inspiratory extension of the thoracic cavity and lowering of the intrathoracic pressure probably induces venous outflow from intracranial venous compartments toward the heart, as the most important factor for cranial fluid flow.<sup>57</sup> Breathing movements of the thorax govern the transport of amniotic fluid instead of air in and out of the lungs. The venous fluid shifting out of the cranium has thus to be counterbalanced by upwardly directed CSF flow.

The establishment of adequate prenatal CSF dynamics is a prerequisite not only for the development of brain structures and skull size but also for neuronal migratory processes. Fetuses with spina bifida aperta without prenatal surgical treatment present significantly impaired neuronal migration of the subventricular zone with consecutive heterotopia, damage of the ependymal layers with regional loss of the ependyma, desynchronized ciliary beating, and resultant astrocytic scarring.<sup>58,59</sup> All of these factors are also known to promote the formation of HC.

### Spontaneous Intracranial Hypotension and Rhinorrhea

SIH is a severe pathophysiologic condition affecting adults up to the age of 60 years, but it very rarely affects children. Commonly, osseous spurs inside the spinal canal cause dural leakage and epidural collection of CSF inside the canal below

the cerebellar tentorium. Depending on the loss of CSF, cranial hypotension may ensue, leading to subdural effusion and chronic subdural bleeding. Subsequently, characteristic signs of brain sagging, such as a lowered cerebellum and tonsillar herniation, point out the reduced brain turgor and impaired buoyant force. The condition, termed *nongeriatric chronic subdural hematoma*, can cause severe morbidity and even mortality. Therapy includes the neurosurgical application of spinal blood patches, bed rest, and, if possible, closure of the leakage by an intraspinal intradural approach.<sup>60,61</sup>

In contrast to infratentorial leakage, CSF loss due to traumatic rhinorrhea, for example, at the petrous bone or in frontobasal locations, occurs within the supratentorial subarachnoid CSF spaces. It is important to note that although severely complicated by local infections or meningitis, brain sagging alluding to SIH with loss of buoyancy has not been observed. Symptoms such as severe headache and vomiting exaggerated in an upright position are similar to those after lumbar puncture and consecutive temporary shortage of CSF. Therapeutic measures with aminophylline and caffeine have been proven effective.<sup>62</sup> Notably, both substances elevate breathing frequency, which, considering it the main driving force, could result in more profound CSF being moved upward into the cranial vault.

The important difference between loss of CSF in rhinorrhea permanent spinal leakage is the relative location to the aqueduct. The capacity to counterbalance continuous fluid loss below the aqueduct by breathing-induced upward movements of CSF is limited by the shortage of available CSF

volume; thus, SIH and chronic subdural hematoma may evolve over time. On the other hand, CSF loss due to subarachnoid leakage above the aqueduct obviously does not induce SIH and has not been reported. Extending our hypothesis about the potential valve function of the aqueduct in conjunction with dense sealing of the para-aqueductal pathways (► **Fig. 2**), the exclusive fluid passage seems to be able to prevent intracerebral hypotension.

### Shunt-Mediated CSF Overdrainage

An upright body position and concomitant hydrostatic forces represent a challenge for the treatment of CSF overdrainage using ventricular shunt systems. CSF overdrainage into the extracranial compartments is known to be a common side effect eventually associated with severe and acutely exacerbating complications, particularly during periods of fast growth in childhood and adolescence. Slit ventricles detected by MRI and complaints of severe headaches that escalate during movement into a standing position are typical symptoms. Their increase over the course of the day might be attributed to a diminution in systemic hydration and the consequently increasing imbalance of the venous–CSF fluid system. Affected children usually develop learning difficulties and complain about headaches and dizziness when moving from a horizontal position.

It is compelling to note that fluid overdrainage from the ventricles apparently does not provoke the characteristic sequela of SIH, as brain sagging has never been observed. Moreover, headaches early in the morning are comparatively rare, which may be attributed to the night-long filling effect due to breathing-related upward CSF movement from the spinal canal into the ventricles, i.e., during sleep in a horizontal position and under hydrostatic equilibrium. Over time, however, ventricular narrowing occurs and may lead to the encasement of shunt catheters between the walls. Chronic overdrainage may even cause subdural effusion and bleeding. The treatment of this condition requires the integration of a gravity control system into the shunt diversion to attenuate hydrostatic forces.

### Space Flight Disease

Recently, Van Ombergen et al.<sup>63,64</sup> reported persistent enlargement of the brain CSF spaces in astronauts after long-duration space flights. Postflight volume changes in the lateral ventricle were shown to be positively correlated with visual acuity loss—a well-known sequela of space flight–associated neuro-ocular syndrome (SANS). The underlying mechanisms of SANS, which refer to various ocular abnormalities such as globe flattening and optic disc edema, are still not fully understood. The current findings demonstrate not only long-lasting enlargement of ventricular and subarachnoid CSF spaces but also postflight reductions in white and gray matter volumes. This observation seems to link changes in brain CSF spaces to the formation of SANS. In fact, the observations of increased ventricular volume and decreased CSF space below the vertex highly support disturbances of CSF circulation as the underlying cause, possibly due to imbalances of breathing-induced CSF upward flow, as

hypothesized by Ludwig et al.<sup>65</sup> Interestingly, these alterations are sustained for as long as 7 months after space flight, implying that they eventually remain unaffected for a longer time by the return to gravity and the reestablishment of hydrostatic pressure conditions. This might indicate that after gravity is restored, the human system needs quite a long time for the excessive accumulation of CSF in the brain ventricles as well as the white and gray matter volumes to normalize and for the CSF flow system to reach physiological conditions.

Thus, long-term changes in ventricular volumes due to space flight might act as a model for the induction of HC.

Notably, the brain aquaporin system is able to adapt to different pressure situations<sup>66,67</sup> by adjusting the localization of aquaporin-1 (AQP1), an isoform of aquaporin water channels expressed at the ependyma of the choroid plexus,<sup>68</sup> and the expression density of AQP4 at astrocytic end feet. In mice experiencing space flights or head-down tilt (HDT), AQP1 was redistributed from apical to basal ependymal membranes.<sup>69</sup> AQP expression was altered by gravity.<sup>70</sup> In mice with kaolin-induced HC, AQP1 was redistributed from the apical to cytoplasmic vesicles and basal membranes of the ependyma.<sup>71</sup> Hence, alterations in the aquaporin system might be responsible for long-lasting or permanent adaptations of ventricular volumes after zero-gravity space flights and might be involved in many CSF disorders in which CSF dynamics are altered. A related pathway has been described for nitric oxide (NO),<sup>72</sup> which is delivered by neuronal or endothelial isozymes and is able to cross membranes<sup>73,74</sup> and AQP4 channels. It facilitates the motility and beating frequency of several ciliated epithelial cells.<sup>75</sup> A disturbance of the NO system by intraventricular hemoglobin in preterm infants or SAH<sup>76</sup> allows us to argue for a link between CSF flow, ependymal ciliary sensing function, endothelial relaxation, capillary dilation, and astrocytic clearance function. NO is also important for the structural maintenance of the periventricular subependymal zone.<sup>77</sup> It may therefore be worthwhile to focus on elevated NO expression in microgravity as well as in gravity simulation experiments by rat tail suspension or during human HDT.<sup>78</sup>

### Sleep and Head-down Tilt

Based on the observation that inspiration elicits an upward cranial motion of CSF, it can be assumed that sleep in a horizontal body position with zero hydrostatic pressure enhances fluid flow, while irregular breathing during sleep may lead to CSF-related disorders. For example, obstructive sleep apnea has been reported to be frequently associated with the development of iNPH.<sup>79</sup> This may be due to a disturbance in the well-orchestrated interplay between the venous and CSF systems. During sleep apnea, repeated attempts to breathe induce strong respiratory efforts against a closed glottis, which severely lowers the intrathoracic pressure. The subsequently compromised venous return to the heart may result in retrograde intracranial venous hypertension. Perturbations of venous–CSF equilibrium hamper CSF exchange between the spinal canal and head and may thus contribute to an increasing ventricular diameter and reduced glymphatic circulation.<sup>80,81</sup>

Enhanced upward CSF flow by inspiration during sleep was observed in NREM sleep in healthy individuals using electrophysiologic methods and blood oxygenation–level dependent MRI.<sup>28</sup> During NREM phases of deep sleep,<sup>82</sup> the human respiration cycle is presumably slow, regular, deep, and sustainable enough to move a sufficient CSF volume into the ventricles, probably facilitated by a reduction in the hydrostatic gradient. Such an ample supply of CSF might be the reason for the enhanced glymphatic clearance of tau and A $\beta$ -amyloid during sleep.<sup>83,84</sup>

An HDT position reverses hydrostatic conditions and leads to higher ICP and may serve as an experimental simulation of long-duration space flight with zero gravity. Recent studies have reported decreased cerebral blood flow in patients in an HDT position compared with a horizontal position. Furthermore, exposure to 3% CO<sub>2</sub> during HDT enhances aqueductal CSF velocities and lateral ventricle CSF mean diffusivity.<sup>85,86</sup> Remarkably, a simultaneous increase in lateral ventricle volume and the formation of optic disc edema in the HDT position resemble observations in astronauts after space flight.<sup>70,85,87</sup> In accordance with animal models of HC,<sup>53</sup> altered ICP, fluid flow physiology, and dynamics during HDT may be hypothesized to interfere with choroidal depolarization and AQP dysfunction.<sup>88</sup>

## Discussion

The recent observation of breathing-induced CSF upward motion in association with intrathoracic venous pooling opens new horizons for a profound understanding of CSF flow–related disorders. These insights apply to CSF homeostasis in pediatric diseases such as preterm PHHC and congenital obstructive HC as well as in adult intracranial spontaneous hypertension. Additional information may be gathered by observations of altered CSF physiology during space flight. Zero gravity offers a model to more precisely understand conditions that are related to the counterbalancing effects of gravity, venous outflow, and CSF upward flow. On the other hand, CSF exchange with interstitial fluid and VRS relies on proper expression of AQP4 water channels in astrocytic end feet. Thus, CSF regulation comprises at least three main regulatory mechanisms: (1) inspiration-induced CSF supply upward into the skull and control of CSF flow into brain CSF spaces by aqueductal function, (2) ependymal flow and its pressure-sensing properties, and (3) AQP1 and AQP4 functions regulating interstitial water secretion and resorption on a molecular scale, respectively. Such knowledge is mandatory to explain iNPH, PHHC, SIH, and spina bifida–related disorders as well as space flight disease. Furthermore, this knowledge may help us to further understand conditions associated with CSF disturbances and, above all, to develop and improve therapeutic approaches.

## Search Strategy and Selection Criteria

We searched PubMed, Crossref Search, and Web of Science for reports published in English from January 1, 1995, to March 31, 2020. The search terms “hydrocephalus” and “hydrocephalic” were combined with many terms such as

“epidemiology,” “pathophysiology,” “etiology,” “diagnosis,” “management,” “aquaporin,” “cerebro spinal fluid,” and “glymphatic.” In addition to the search results, we also hand searched the references of relevant articles retrieved by search strategy. The final reference list was generated on the basis of relevance to the topics covered in this review.

## Author Contributions

H. C. L., S. D.-K., and J. F. edited the manuscript; H. C. L., S. D.-K., H. C. B., and J. G. defined the scope, objective, and clinical details. H. C. L., S. D.-K., H. C. B., J. F., J.-G., and S. S. reviewed the manuscript.

## Competing Interests

The authors declare no competing financial interests.

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## Conflict of Interest

J. D. reports personal fees from Bayer, personal fees from Biogen, grants and personal fees from Novartis, and personal fees from Sanofi, outside the submitted work. In addition, J. D. has a patent “Treatment of Multiple Sulfatase Deficiency,” US Patent No 16591051 licensed.

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