

SCIENTIFIC REPORTS



OPEN

Choroidal fissure acts as an overflow device in cerebrospinal fluid drainage: morphological comparison between idiopathic and secondary normal-pressure hydrocephalus

Received: 23 May 2016
Accepted: 17 November 2016
Published: 12 December 2016

Shigeki Yamada^{1,2}, Masatsune Ishikawa^{1,2}, Yasushi Iwamuro² & Kazuo Yamamoto²

To clarify the pathogenesis of two different types of adult-onset normal-pressure hydrocephalus (NPH), we investigated cerebrospinal fluid distribution on the high-field three-dimensional MRI. The subarachnoid spaces in secondary NPH were smaller than those in the controls, whereas those in idiopathic NPH were of similar size to the controls. In idiopathic NPH, however, the basal cistern and Sylvian fissure were enlarged in concurrence with ventricular enlargement towards the z-direction, but the convexity subarachnoid space was severely diminished. In this article, we provide evidence that the key cause of the disproportionate cerebrospinal fluid distribution in idiopathic NPH is the compensatory direct CSF communication between the inferior horn of the lateral ventricles and the ambient cistern at the choroidal fissure. In contrast, all parts of the subarachnoid spaces were equally and severely decreased in secondary NPH. Blockage of CSF drainage from the subarachnoid spaces could cause the omnidirectional ventricular enlargement in secondary NPH.

Since 1965, adult-onset normal pressure hydrocephalus (NPH) has been largely classified into idiopathic NPH (iNPH) or secondary NPH (sNPH), which develops after subarachnoid haemorrhage, trauma or infection¹. The diagnosis and management of iNPH has made rapid progress since evidence-based guidelines were established^{2–6}. The most typical morphological characteristics of CSF distribution in iNPH were narrow sulci at the high convexity due to z-axial expansion of the lateral ventricles and enlargement of the basal cistern and Sylvian fissure^{7–11}. Subsequently, many MRI volumetric analyses proposed several parameters specific to iNPH^{9,10,12–14}. Nevertheless, little information is available on the CSF distribution in sNPH. Therefore, we quantitatively assessed the CSF distributions in patients with sNPH compared with those in patients with iNPH by using volumetric analysis based on the high-field T2-weighted 3D MRI. Recently, we provided new evidence for the comparison of the CSF volumes observed in patients with iNPH and those with Alzheimer disease¹⁰. Here, we reported that the basal cistern and Sylvian fissure in patients with iNPH was enlarged having the same mean volume as that in patients with Alzheimer disease, and, in addition the ventricles in iNPH patients were >40 mL larger than those in Alzheimer disease. These findings contribute to our understanding of the development of concurrent expansion of the ventricles and basal cistern and Sylvian fissure in iNPH. Compensatory direct CSF pathways between ventricles and subarachnoid spaces other than the foramina of Luschka and Magendie have been observed in hydrocephalic rat models^{15–18}. Therefore, we hypothesized that the choroidal fissure, which is formed from the medial wall of the lateral ventricles, was opened and directly mediated the flow of CSF between the lateral ventricles and the basal cistern, acting as an overflow device under conditions of increased intracranial CSF volume, such as in iNPH. To

¹Normal Pressure Hydrocephalus Center, Rakuwakai Otowa Hospital, Kyoto, Japan. ²Department of Neurosurgery and Stroke Center, Rakuwakai Otowa Hospital, Kyoto, Japan. Correspondence and requests for materials should be addressed to S.Y. (email: shigekiyamada39@gmail.com)

	iNPH	sNPH	Control	P1	P2	P3
Total number	52	15	31			
Mean age (years)	76.7 ± 6.9	72.3 ± 10.8	75.8 ± 7.4	0.220	0.546	0.534
Male	31 (60%)	8 (53%)	17 (55%)	0.891	0.844	1.000
Evans index >0.3	41 (79%)	12 (80%)	6 (19%)	1.000	<0.001	<0.001
Z-Evans index >0.4	45 (87%)	7 (47%)	3 (10%)	0.006	<0.001	0.014
Callosal angle <90°	45 (87%)	8 (53%)	5 (16%)	0.005	<0.001	0.023
High convexity tightness	46 (89%)	1 (7%)	1 (3%)	<0.001	<0.001	1.000
Enlarged Sylvian fissure	45 (87%)	2 (13%)	0	<0.001	<0.001	0.191
Opening of the inferior choroidal point	39 (75%)	2 (13%)	0	<0.001	<0.001	0.191
Severe periventricular hyperintensity	44 (85%)	11 (73%)	18 (58%)	0.534	0.015	0.497
Severe deep white matter hyperintensity	36 (69%)	9 (60%)	15 (48%)	0.720	0.098	0.671

Table 1. Morphological characteristics of the study population. The Evans index was measured as the maximal width of the frontal horns of the bilateral ventricles to the maximal width of the internal diameter of the cranium on the basis of the X dimension. The Z-Evans index was measured as the maximum z-axial length of the frontal horns of the lateral ventricles to the maximum z-axial length of the median skull. The callosal angle was measured as the angle of the roof of the bilateral ventricles on the coronal plane, which was perpendicular to the anteroposterior commissure plane on the posterior commissure. Opening of the inferior choroidal point was defined as direct communication of CSF spaces between the inferior horn of the lateral ventricles and the ambient cistern at the inferior choroidal point of the choroidal fissure. Severe periventricular hyperintensity was defined as large (Grade 2) or extension (Grade 3), or deep white matter hyperintensity according to the grading scales reported by Fazekas *et al.*²⁰. P1; probability value of iNPH vs. sNPH. P2; probability value of iNPH vs. control. P3; probability value of sNPH vs. control.

elucidate the pathogenesis of iNPH and sNPH, we investigated this direct communication between the ventricles and the basal cistern, focusing on the inferior choroidal point of the choroidal fissure.

Results

Clinical characteristics and morphological indices. Fifty-two patients were diagnosed with iNPH, 15 patients were diagnosed with sNPH, and 31 participants were allocated to the control group. The mean ages were not significantly different among the three groups (Table 1). The mean time of disease duration from the initial presentation of symptoms until the diagnosis of iNPH was 2 years. The mean duration from the initial event of subarachnoid haemorrhage or head injury until the diagnosis of sNPH was 5 months. Evans index >0.3, which is the most popular index of ventricular enlargement^{2–6}, was satisfied by approximately 80% of the patients diagnosed both with iNPH and sNPH. In contrast, z-Evans index >0.4⁹, callosal angle <90°¹⁹, high convexity tightness and enlarged Sylvian fissure^{7,8,11}, which have been reported to be specific indices of iNPH, were satisfied in ≥85% of patients with iNPH but in <55% of those with sNPH (Table 1); this difference between the two groups was statistically significant. Severe periventricular hyperintensity (smooth halo or expanding) and severe deep white matter hyperintensity (confluence of foci) on the Fazekas rating scale²⁰ were frequently observed in the patients with iNPH, whereas the patients with sNPH had similar percentages of hyperintensity when compared to the controls (Table 1). Additionally, an opening of the inferior choroidal point, which we here newly define as a direct pathway linking CSF spaces between the inferior horn of the lateral ventricles and the ambient cistern at the inferior choroidal point of the choroidal fissure, was observed in 39 of 52 patients (75%) with iNPH but in only 2 of 11 (13%) with sNPH and in none of the controls (Table 1).

Comparison of ventricular shape and CSF volumes between iNPH and sNPH. Figure 1 shows the ventricles in the representative cases with iNPH and sNPH and in the control groups. The total size of all the ventricles in patients with iNPH and sNPH were 2.6 and 2.0 times larger, respectively, than those in the controls (Table 2). The mean volume and volume ratio of the total subarachnoid spaces in patients with iNPH and controls were almost identical (260 mL and 17% versus 272 mL and 18%, respectively), however, the CSF distribution in the subarachnoid spaces was different between these two groups, as shown in Fig. 2 and Table 2. The mean volume of the basal cistern and Sylvian fissure in patients with iNPH was 30 mL larger than that in the controls, whereas the volume of the convexity subarachnoid space was 50 mL smaller in patients with iNPH than in the controls. Furthermore, we observed that each of the three parts of the subarachnoid spaces in patients with sNPH was significantly smaller than each of those parts in the controls by equal proportions, and the mean volume of the total subarachnoid spaces in patients with sNPH was >100 mL smaller than that in the controls. Overall, the intracranial CSF spaces in patients with iNPH were significantly larger than those in the age-matched controls, whereas those in patients with sNPH were smaller (Fig. 2 and Table 2).

Patients with iNPH possess a direct CSF pathway at the inferior choroidal point of the choroidal fissure. Figure 3 shows key sections on 3D-constructive interference in steady state (CISS) MRI sequences from representative patients with iNPH or sNPH and control groups. Compared to the controls, all regions of subarachnoid space, including the ambient cistern and Sylvian fissure, were severely diminished in the patients with sNPH, although the ventricles were enormously enlarged. On the contrary, not only the ventricles but also

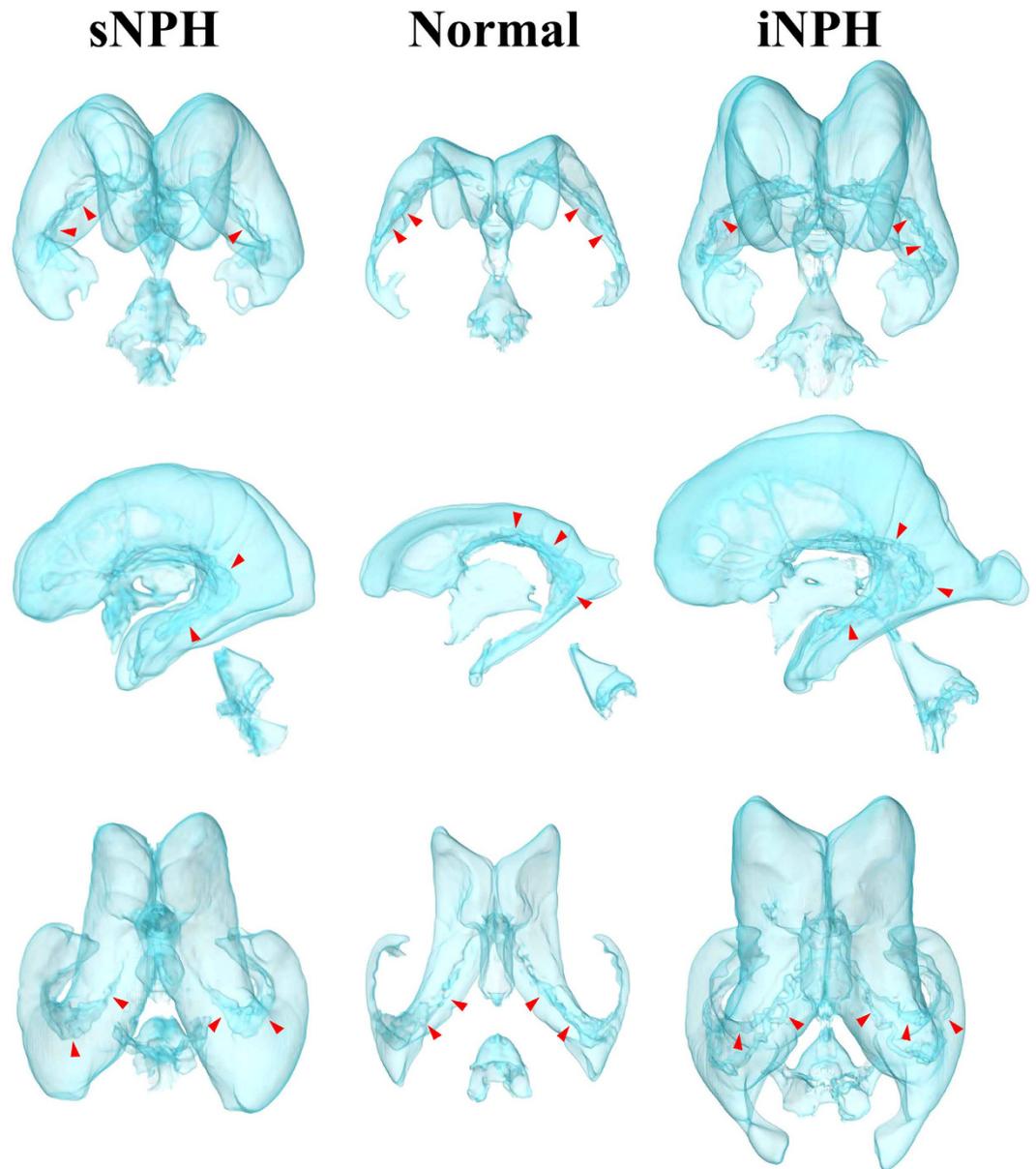


Figure 1. Three-dimensional views of the ventricles in iNPH, sNPH and normal brain. Compared to the controls (middle), the ventricles in sNPH patients expanded symmetrically in all directions, whereas the ventricles in iNPH patients expanded disproportionately towards the z-axis direction. The red arrow heads indicate the choroid plexus.

the Sylvian fissure and basal cistern, including the ambient cistern, were conspicuously enlarged in the patients with iNPH. Normally, the choroidal fissure at the medial wall of the lateral ventricles is constructed from the choroid plexus and a thin membrane without ependyma^{21,22}. At the inferior choroidal point of the choroidal fissure, the anterior choroid arteries and inferior choroidal veins run through the ambient cistern and the inferior horn of the lateral ventricles. In the patients with iNPH, however, the choroid plexus appeared to float in the CSF, and the border seemed to disappear due to the enlargement of the ambient cistern and inferior horn and the upward displacement and compression of the head of the hippocampus (Fig. 3, right).

Further evidence indicating a direct pathway of CSF flow at the inferior choroidal point of the choroidal fissure is presented in head CT images taken before and immediately after shuntography performed on a patient with iNPH with a suspected shunt malfunction (Fig. 4). On the CT image taken immediately after shuntography in the lower right lateral decubitus position, contrast medium was observed in both the inferior horn of the lateral ventricles and in the ambient cistern.

In the video clip showing an amygdalohippocampectomy (Supplementary), it is evident that, before microsurgical dissection of the transparent membrane of the choroidal fissure, the CSF in the ambient cistern passed through this membrane and moved freely in and out of the inferior horn of the lateral ventricles. This to-and-fro movement of the CSF provides evidence that the fluid could freely penetrate the thin arachnoid membrane at the

	iNPH (52)	sNPH (15)	Control (31)	P1	P2	P3
Total intracranial spaces	1517 ± 156	1467 ± 147	1487 ± 138	0.310	0.369	0.639
Brain parenchyma	1096 ± 119 (72.4%)	1177 ± 134 (80.3%)	1154 ± 147 (77.5%)	0.061	0.017	0.746
Total CSF	420.4 ± 84.7 (27.6%)	290.0 ± 85.2 (19.7%)	333.1 ± 94.0 (22.5%)	<0.001	<0.001	0.075
Total ventricles	160.8 ± 45.2 (10.6%)	121.4 ± 26.8 (8.2%)	61.3 ± 31.9 (4.2%)	0.002	<0.001	<0.001
Bilateral ventricles	152.9 ± 41.7	112.2 ± 26.7	52.1 ± 26.4	<0.001	<0.001	<0.001
Third ventricle	5.3 ± 1.5	5.2 ± 1.2	3.5 ± 1.3	0.982	<0.001	<0.001
Fourth ventricle	4.1 ± 2.0	4.5 ± 1.1	2.8 ± 1.5	0.181	0.002	<0.001
Total subarachnoid spaces	259.6 ± 71.5 (17.1%)	168.4 ± 72.3 (11.5%)	271.7 ± 85.7 (18.3%)	<0.001	0.569	<0.001
Convexity subarachnoid space	76.4 ± 35.4	73.0 ± 35.9	125.8 ± 54.8	0.758	<0.001	<0.001
Basal cistern and Sylvian fissure	120.4 ± 32.7	51.6 ± 35.0	90.7 ± 34.0	<0.001	<0.001	0.001
Posterior fossa	62.8 ± 17.7	43.9 ± 10.2	55.2 ± 19.7	<0.001	0.057	0.038

Table 2. Mean volumes (mL) and volume ratios (%) of CSF and brain parenchyma. P1; probability value of iNPH vs. sNPH for the Mann-Whitney-Wilcoxon test. P2; probability value of iNPH vs. control for the Mann-Whitney-Wilcoxon test. P3; probability value of sNPH vs. control for the Mann-Whitney-Wilcoxon test.

inferior choroidal point of the choroidal fissure. However, these additional clinical findings are not sufficient to prove direct CSF communication at the inferior choroidal point of the choroidal fissure because each finding was verified in only one patient.

Discussion

This study shows that CSF distribution markedly differs between the two categories of NPH. Figure 5 shows schemata representing intracranial CSF distributions in iNPH-affected, sNPH-affected and normal brain. The most distinct morphological characteristics in the patients with iNPH were z-axial expansion of the bilateral ventricles in concurrence with enlargement of the basal cistern and Sylvian fissure and diminishment of the convexity subarachnoid spaces. In contrast, the patients with sNPH exhibited ventricles that were enlarged symmetrically in all directions in concurrence with each of the three subarachnoid spaces being severely diminished. The morphological discrepancies between iNPH and sNPH lead us to conclude that the pathogenesis of iNPH is markedly different from the developmental process of sNPH following subarachnoid haemorrhage or brain injury.

Recently, classical understanding of CSF bulk flow and the concept of the third circulation have changed^{23–31}. In the concept of third circulation^{32–34}, CSF mainly secreted by the choroid plexus in the ventricles moves as unidirectional bulk flow from the ventricles to subarachnoid spaces through the foramina of Luschka and Magendie and finally absorbs from the arachnoid villi at the superior sagittal sinus. In contrast, recent experimental data have provided evidence that the choroid plexus is not the major site of CSF production and parenchymal interstitial fluid is excreted into the ventricles and subarachnoid spaces as CSF^{25–27,29–31,35,36}. Ichimura *et al.* reported that the direction of the interstitial fluid flow towards the ventricles or subarachnoid spaces varies in an unpredictable manner³⁵. Furthermore, current research suggests that lymphatic drainage via the peri- and para-vascular space^{23,31,37–40}, perineural subarachnoid spaces of the cranial nerves or meninges^{25,26,41,42} is the major pathway for absorption of CSF and interstitial fluid, and venous drainage via the dural arachnoid villi is the minor^{23,24,28,43}. Newly developed MRI techniques for assessing CSF dynamics have revealed that CSF moves in a pulsatile fashion and not in a unidirectional bulk flow^{44–50}. CSF moves upward and downward around the cranio-cervical junction with cardiac pulsation and respiration as driving forces^{44–50}. These recent findings have negated the classical concept of the CSF bulk flow from the ventricles to subarachnoid spaces through the foramina of Luschka and Magendie.

Our observation of CSF distribution reduced subarachnoid spaces in the high parietal convexity in iNPH supports the current view of CSF absorption. The main pathogenic mechanism associated with iNPH is thought to be the disruption of the most downstream site of CSF drainage systems, because patients with iNPH have large volumes of intracranial CSF. The subarachnoid spaces might be harder to expand than the ventricles because there are many trabeculae and partitions in the subarachnoid spaces, as shown in Fig. 6. Additionally, the most downstream site for CSF absorption is likely to become the most enlarged CSF spaces which were the ventricles, basal cistern and Sylvian fissure, but not convexity subarachnoid space in iNPH. We have also produced new evidence that the ventricles and subarachnoid space possess several direct alternative connections other than the foramina of Luschka and Magendie that allow protection from overflow^{15–18}. The expansion of intracranial CSF volume in iNPH creates compensatory routes between the ventricles and subarachnoid spaces. Mainly, the inferior horn of the lateral ventricles directly communicates with the ambient cistern via the opening of the inferior choroidal point of the choroidal fissure, serving as an overflow management point for the ventricular drainage system. However, the choroidal fissure appears open in most, but not all, patients with iNPH. In a hydrocephalic rat model, Park *et al.* found that the interface between the third ventricle and the quadrigeminal cistern also serves as a compensatory direct CSF pathway in addition to the choroidal fissure¹⁷. We confirmed that the subarachnoid spaces around the third ventricle appear enlarged in patients with iNPH and diminished in patients with sNPH compared to a control population (Fig. 6). Further investigations are needed to confirm the other compensatory direct CSF pathways in iNPH patients with a closed choroidal fissure.

On the other hand, the main pathogenic factor for the omnidirectional expansion of ventricles in sNPH is thought to be the disruption of CSF drainage from subarachnoid spaces, because the subarachnoid spaces in

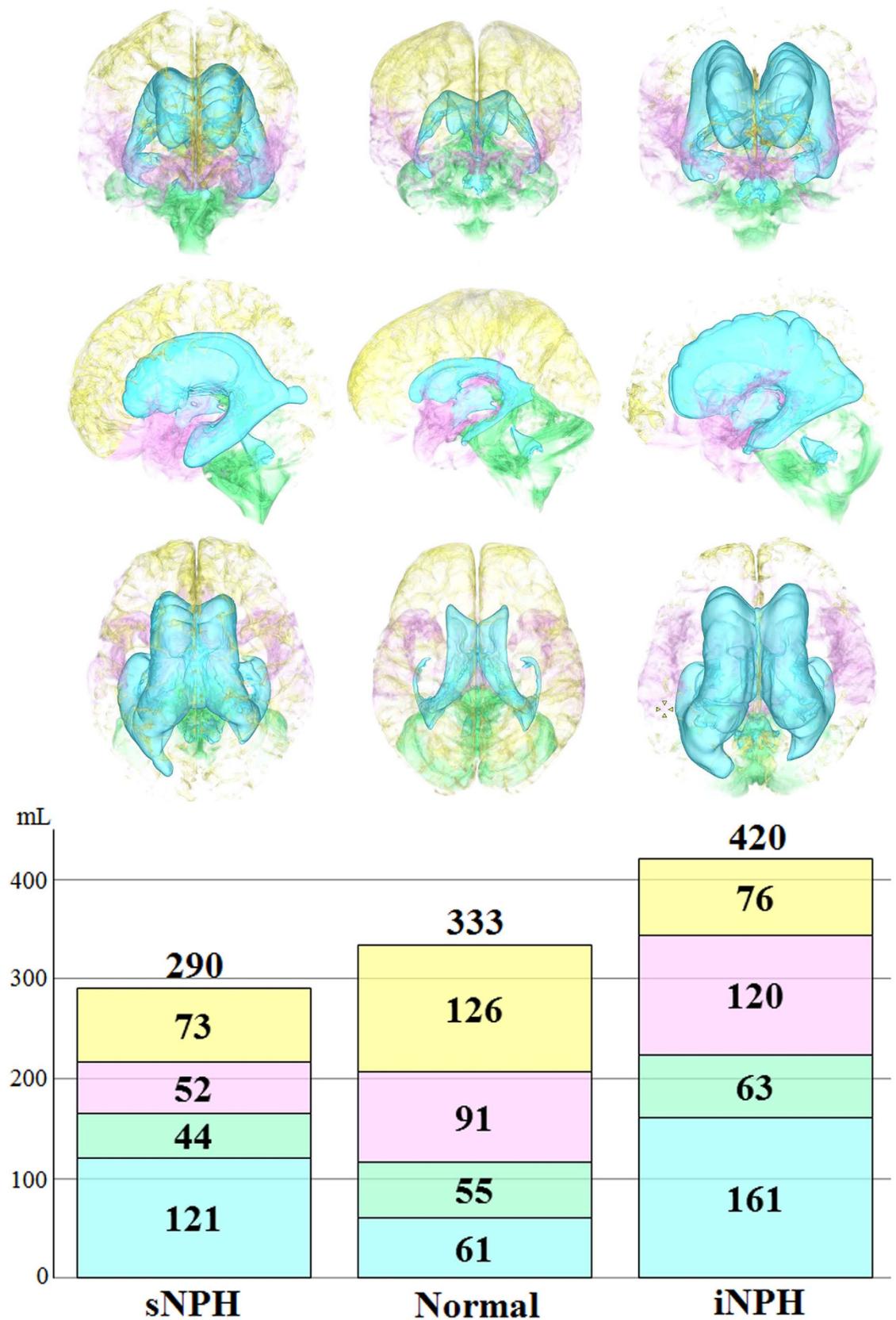


Figure 2. Mean volumes of the segmented intracranial cerebrospinal fluid spaces in iNPH, sNPH and normal brain. The mean volumes of ventricles (sky blue), basal cistern and Sylvian fissure (pink) and subarachnoid space in the posterior fossa (light green) were the largest in patients with iNPH. The mean volume of the convexity part of the subarachnoid space (light yellow) was equally decreased in patients with iNPH and those with sNPH.

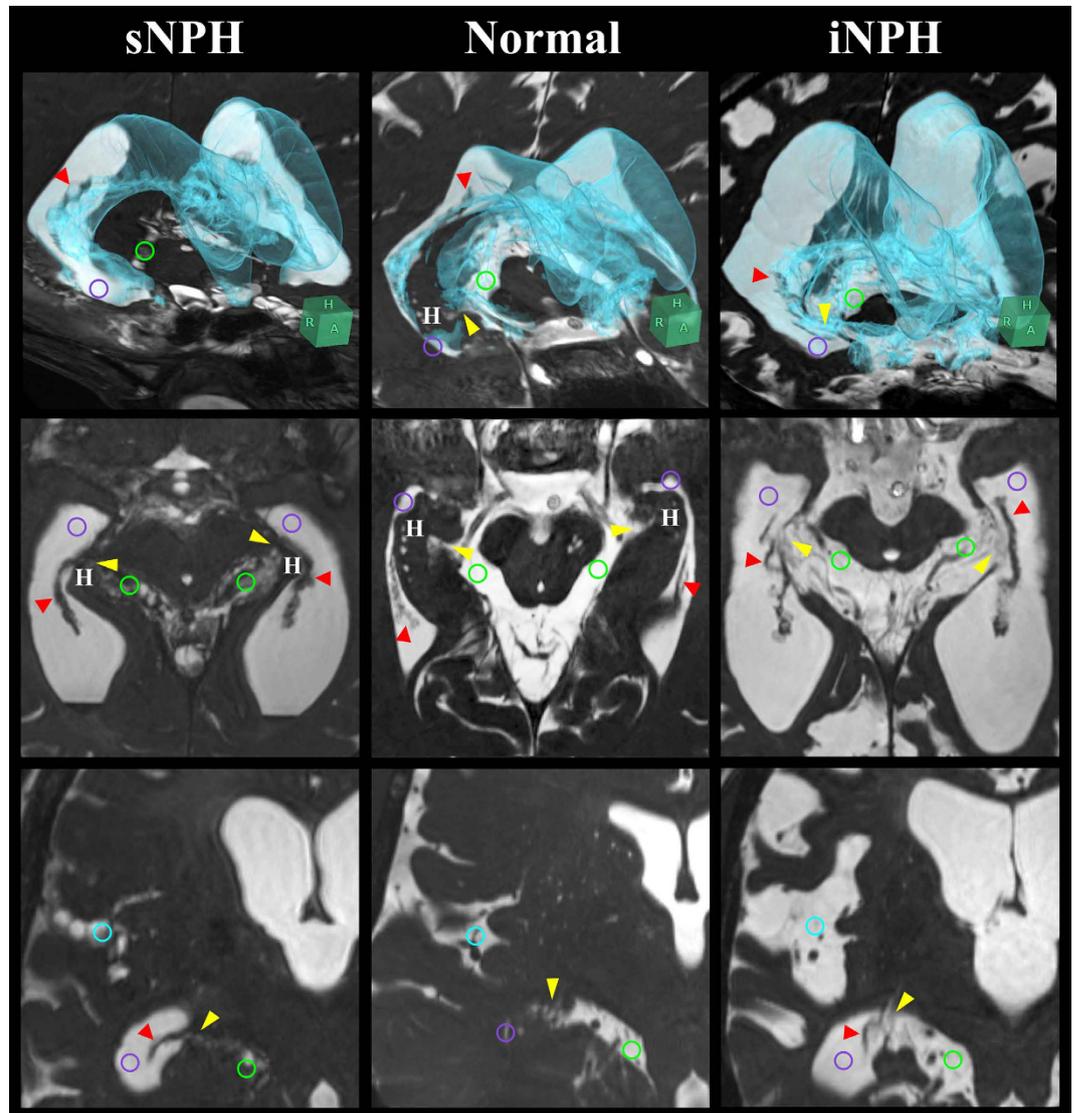


Figure 3. Fine structures of the lateral ventricles and the basal cistern around the inferior choroidal point of the choroidal fissure. The 3D views (upper) which combine axial (middle) and coronal (lower) sections were at the level of the inferior choroidal point of the choroidal fissure (yellow arrow head) between the ambient cistern (light green circle) and the inferior horn of the lateral ventricles (purple circle). In the normal situation, at the head of the hippocampus (H) and choroid plexus (red arrow head), the ambient cistern is separated from the inferior horn of the lateral ventricles. In patients with iNPH, the ventricles, basal cistern and Sylvian fissure (light blue circle) were enlarged with the concurrent opening of the inferior choroidal point of the choroidal fissure. However, in patients with sNPH, only the ventricle was enlarged and all of the subarachnoid spaces, including the ambient cistern and Sylvian fissure, were severely diminished.

sNPH extensively diminished and the total intracranial CSF volume was smaller than that in the age-matched controls. Some clinical studies have reported that significantly increased inflammatory fibrogenic cytokines or ferritin levels in CSF are associated with sNPH after subarachnoid haemorrhage^{51–53}. Additionally, a rat model of communicating hydrocephalus by injecting kaolin in the subarachnoid spaces mimics human sNPH^{54–56}. In a kaolin-induced sNPH model, the subarachnoid spaces were obstructed with the number of macrophages and severe fibrosis^{55,56}. Inflammation in CSF is known to be associated with arachnoid and pia matter fibrosis⁵⁷, which might lead to the closure of subarachnoid spaces. The present findings serve as a foundation for further investigation into the pathophysiological mechanisms underlying the development of iNPH and sNPH.

In conclusion, the present study revealed the characteristics of CSF distribution in patients with iNPH, which strongly differed from those in patients with sNPH. Patients with iNPH had a substantially larger total CSF volume than control subjects, whereas patients with sNPH had a smaller total CSF volume. Under normal conditions, the ventricles directly communicate with the subarachnoid spaces via the foramina of Luschka and Magendie. However, in iNPH, the ventricles and basal cistern are enlarged, along with the opening of the inferior choroidal point of the choroidal fissure. This provides a direct compensatory CSF communication route

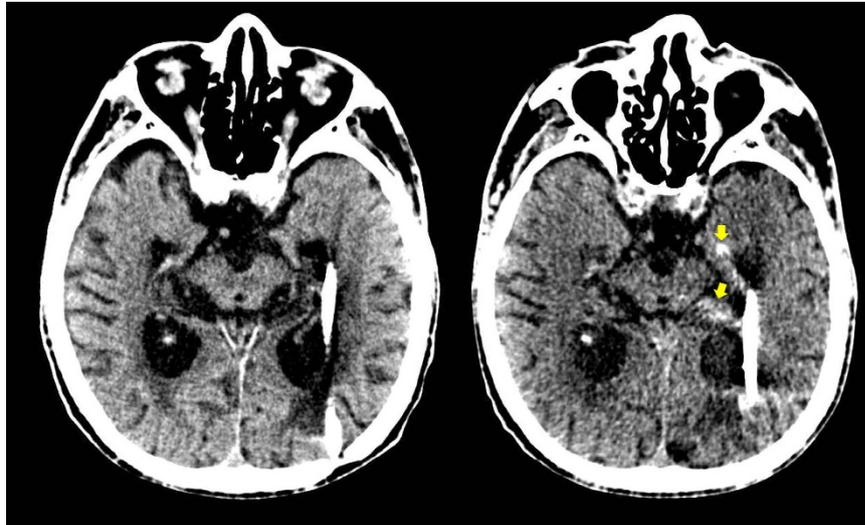


Figure 4. Confirmation of the direct CSF communication between the lateral ventricles and the ambient cistern by shuntography in patients with iNPH with a suspected shunt malfunction. Contrast medium (yellow arrow head) was observed not only in the left inferior horn of the lateral ventricles but also in the left ambient cistern on CT scan immediately after shuntography (right), compared to that before shuntography (left).

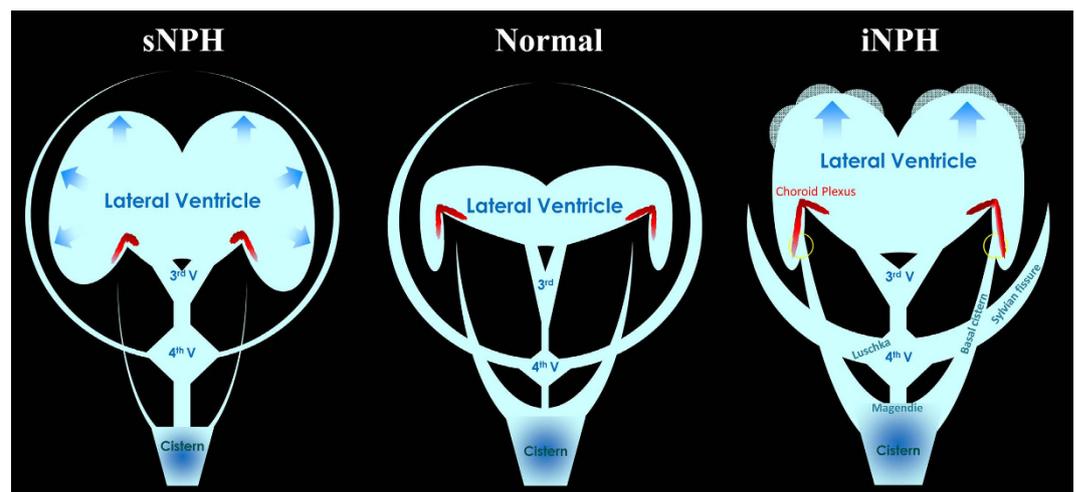


Figure 5. Schematic diagram showing the distribution of CSF in patients with iNPH and sNPH, compared to normal brains. In iNPH patients, the inferior horn of the lateral ventricles is directly connected with the basal cistern at the inferior choroidal point of the choroidal fissure (yellow circle) inside the choroid plexus.

between the ventricles and subarachnoid spaces. Thus, this is a key cause of the disproportionate CSF distribution in iNPH, named “Disproportionately Enlarged Subarachnoid-space Hydrocephalus (DESH)”⁷, which is z-axial expansion of the bilateral ventricles in concurrence with enlargement of the basal cistern and Sylvian fissure and diminishment of the convexity subarachnoid spaces. Conversely, in sNPH all parts of the subarachnoid spaces are equally and severely reduced in size. We postulate that fibrosis of arachnoid and pia matter reduces the subarachnoid spaces leading to expansion of the ventricles. These novel findings may contribute to future studies of the mechanisms that drive CSF movement and absorption.

Methods

Ethical approval and patient consent. The study protocol was designed in accordance with guidelines outlined in the Declaration of Helsinki and was approved by the ethics committee for human research at Rakuwakai Otowa Hospital, Kyoto, Japan (approval number; 14-003). The methods were performed in accordance with the approved guidelines. Written informed consent was obtained from all patients or their family members.

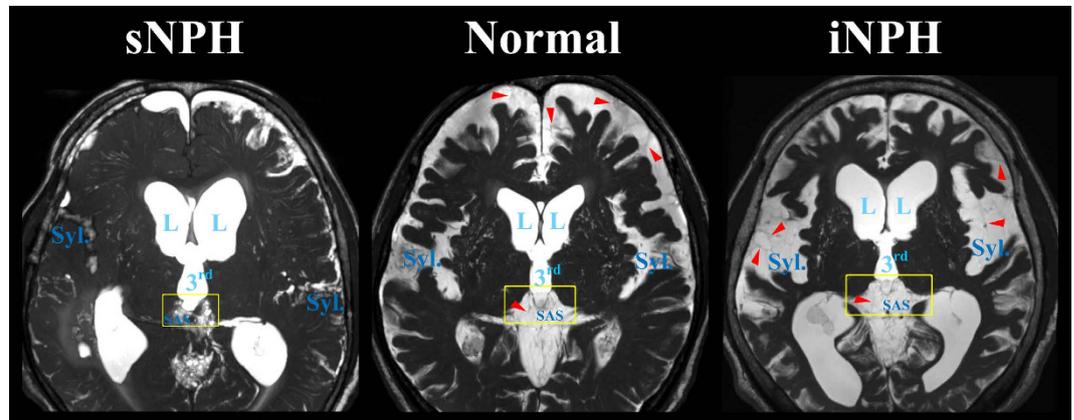


Figure 6. Fine structures of the subarachnoid spaces around the third ventricle. Patients with iNPH had the enlarged Sylvian fissure (Syl.) in concurrent with enlargement of the lateral (L) and third (3rd) ventricles, whereas patients with sNPH had the diminished Sylvian fissure. The border between third ventricle and surrounding subarachnoid spaces (SAS) was thinning in iNPH patients (yellow square). The red arrow heads indicate the trabeculas and partitions in the subarachnoid spaces.

Study population. Details regarding clinical data collection, image acquisition, and segmentation and quantification of the ventricles and subarachnoid spaces were described in our prior publications^{9,10}. In brief, 72 patients diagnosed with NPH based on the enlarged ventricles under normal CSF pressure and the symptoms of gait disturbance, cognitive impairment and urinary disturbance participated in this study. All of them underwent MRI examinations and the CSF tap-test, which consisted of removing ≥ 30 mL of CSF via a lumbar tap after written informed consent. Eight patients were diagnosed with sNPH after subarachnoid haemorrhage, 7 patients developed sNPH after head injury, and 52 patients were diagnosed with iNPH because a known antecedent cause was absent. Five patients who were diagnosed with congenital aetiology NPH based on the morphological features were excluded from this study. There were relatively few patients with sNPH for statistical analysis, because our NPH centre was consulted for the diagnosis and treatment of a patient once a patient was suspected to have iNPH, but rarely consulted for patients diagnosed with sNPH following subarachnoid haemorrhage. Thirty-one participants ≥ 60 years old were recruited as controls since they did not have any symptoms of short-stepped gait and/or cognitive impairment, and these controls were confirmed via MRI to be without ventricular dilatation or fluid accumulation, such as subdural haematoma. Controls included patients who were diagnosed with unruptured small intracranial aneurysms or who had a history of small intracerebral haemorrhage or lacunar infarction. Additionally, patients with a history of subarachnoid haemorrhage, head injury, suspected dementia, or neurodegenerative disorders such as Alzheimer's disease and Parkinson's disease were excluded from the control group.

Image acquisition for measurement of segmented CSF volume and parameters. All MRI examinations were performed with a 64-channel 3-Tesla MRI system (MAGNETOM Skyra, Siemens AG, Muenchen, Germany). The sagittal source images of the T2-weighted 3D-SPACE were automatically processed to create 3D volume-rendering reconstruction and multiplanar reconstruction images by using an independent 3D volume analyser workstation (SYNAPSE 3D; FUJIFILM Medical Systems; Tokyo, Japan). The intracranial volumes were segmented by the utilization of the combined techniques of the edge-guided nonlinear interpolation and user-steered live-wire segmentation^{58,59}. After that, the CSF spaces were automatically segmented from brain parenchyma by using a simple threshold algorithm⁶⁰. The intracranial CSF spaces were manually segmented into the bilateral, third, and fourth ventricles, convexity subarachnoid space, basal cistern and Sylvian fissure, and subarachnoid space in the posterior fossa. Each segmented volume was automatically measured by counting the number of voxels. Brain parenchymal volume was calculated as intracranial volume minus total CSF volume. The volume ratios (%) were calculated as ratios of target volumes to the intracranial volume. The validity, reliability and reproducibility of 3D segmentation and quantification of CSF volumes were assessed and described in the previous article⁹. The Evans Index was measured as the maximal width of the frontal horns of the bilateral ventricles to the maximal width of the internal diameter of the cranium on the basis of the X dimension; an Evans index >0.3 was defined as conventional ventricular dilatation, according to the diagnostic guidelines for iNPH²⁻⁶. Alternatively, the z-Evans index was defined as the maximum z-axial length of the frontal horns of the lateral ventricles to the maximum cranial z-axial length; a z-Evans index >0.4 is considered a useful index for the diagnosis of iNPH⁹. In addition, a callosal angle $<90^\circ$, measured as the angle of the roof of the bilateral ventricles on the coronal plane, which was perpendicular to the anteroposterior commissure plane on the posterior commissure, was also used as an index specific to iNPH¹⁹. The fluid attenuated inversion recovery (FLAIR) sequence was also conducted for evaluating the periventricular hyperintensity and deep white matter hyperintensity on the Fazekas rating scale²⁰.

Evidence of direct CSF communication between the lateral ventricles and basal cistern via the choroidal fissure. We subjectively assessed whether there was an opening at the inferior choroidal point of the choroidal fissure using 3D-CISS sequences to evaluate the fine structures in the CSF spaces. Additionally, one patient who was diagnosed with iNPH and who had received a ventriculo-peritoneal shunt four years ago at another hospital needed to undergo shuntography to test the patency of his shunt system because his gait and cognitive function had worsened gradually. A CT scan was performed before and immediately after shuntography, in which a small amount of OMNIPAQUE-240 (iohexol, GE Healthcare) was injected into the left inferior horn of the lateral ventricles via reservoir and ventricular catheter of the shunt system. Furthermore, the actual CSF movement through the thin arachnoid membrane between the inferior horn of the lateral ventricles and the ambient cistern was observed *in situ* during the amygdalohippocampectomy of one patient diagnosed with hippocampal glioma.

Statistics. Mean values and standard deviations for age, sex, and several MRI parameters were calculated and compared among patients with iNPH, sNPH, and control groups by a Mann-Whitney-Wilcoxon test. The Chi-square test was used to compare the two proportions. Statistical significance was assumed at a probability value (p) of less than 0.05. All missing data were treated as deficit data that did not affect other variables. Statistical analysis was performed using R software (version 3.0.1; R Foundation for Statistical Computing, Vienna, Austria; <http://www.R-project.org>).

References

- Hakim, S. & Adams, R. D. The special clinical problem of symptomatic hydrocephalus with normal cerebrospinal fluid pressure. Observations on cerebrospinal fluid hydrodynamics. *Journal of the neurological sciences* **2**, 307–327 (1965).
- Ishikawa, M. Guideline Committee for Idiopathic Normal Pressure Hydrocephalus, J.S.o.N.P.H. Clinical guidelines for idiopathic normal pressure hydrocephalus. *Neurologia medico-chirurgica* **44**, 222–223 (2004).
- Ishikawa, M. *et al.* Guidelines for management of idiopathic normal pressure hydrocephalus. *Neurologia medico-chirurgica* **48** Suppl, S1–23 (2008).
- Marmarou, A., Bergsneider, M., Relkin, N., Klinge, P. & Black, P. M. Development of guidelines for idiopathic normal-pressure hydrocephalus: introduction. *Neurosurgery* **57**, S1–3 (2005).
- Mori, E. *et al.* Guidelines for management of idiopathic normal pressure hydrocephalus: second edition. *Neurologia medico-chirurgica* **52**, 775–809 (2012).
- Relkin, N., Marmarou, A., Klinge, P., Bergsneider, M. & Black, P. M. Diagnosing idiopathic normal-pressure hydrocephalus. *Neurosurgery* **57**, 4–16 (2005).
- Hashimoto, M., Ishikawa, M., Mori, E. & Kuwana, N. Study of, I.o.n.i. Diagnosis of idiopathic normal pressure hydrocephalus is supported by MRI-based scheme: a prospective cohort study. *Cerebrospinal fluid research* **7**, 18 (2010).
- Sasaki, M. *et al.* Narrow CSF space at high convexity and high midline areas in idiopathic normal pressure hydrocephalus detected by axial and coronal MRI. *Neuroradiology* **50**, 117–122 (2008).
- Yamada, S., Ishikawa, M. & Yamamoto, K. Optimal Diagnostic Indices for Idiopathic Normal Pressure Hydrocephalus Based on the 3D Quantitative Volumetric Analysis for the Cerebral Ventricle and Subarachnoid Space. *AJNR American journal of neuroradiology* **36**, 2262–2269 (2015).
- Yamada, S., Ishikawa, M. & Yamamoto, K. Comparison of CSF Distribution between Idiopathic Normal Pressure Hydrocephalus and Alzheimer Disease. *AJNR American journal of neuroradiology* **37**, 1249–1255 (2016).
- Yamashita, F. *et al.* Detection of changes in cerebrospinal fluid space in idiopathic normal pressure hydrocephalus using voxel-based morphometry. *Neuroradiology* **52**, 381–386 (2010).
- Kang, K., Yoon, U., Lee, J. M. & Lee, H. W. Idiopathic normal-pressure hydrocephalus, cortical thinning, and the cerebrospinal fluid tap test. *Journal of the neurological sciences* **334**, 55–62 (2013).
- Moore, D. W. *et al.* A pilot study of quantitative MRI measurements of ventricular volume and cortical atrophy for the differential diagnosis of normal pressure hydrocephalus. *Neurology research international* **2012**, 718150 (2012).
- Serulle, Y. *et al.* Differentiating shunt-responsive normal pressure hydrocephalus from Alzheimer disease and normal aging: pilot study using automated MRI brain tissue segmentation. *Journal of neurology* **261**, 1994–2002 (2014).
- Fenstermacher, J. D., Gherzi-Egea, J. F., Finnegan, W. & Chen, J. L. The rapid flow of cerebrospinal fluid from ventricles to cisterns via subarachnoid velae in the normal rat. *Acta neurochirurgica Supplement* **70**, 285–287 (1997).
- Gherzi-Egea, J. F., Finnegan, W., Chen, J. L. & Fenstermacher, J. D. Rapid distribution of intravenously administered sucrose into cerebrospinal fluid cisterns via subarachnoid velae in rat. *Neuroscience* **75**, 1271–1288 (1996).
- Park, J. H. *et al.* Cerebrospinal fluid pathways from cisterns to ventricles in N-butyl cyanoacrylate-induced hydrocephalic rats. *Journal of neurosurgery Pediatrics* **8**, 640–646 (2011).
- Yoon, J. S., Nam, T. K., Kwon, J. T., Park, S. W. & Park, Y. S. CSF flow pathways through the ventricle-cistern interfaces in kaolin-induced hydrocephalus rats-laboratory investigation. *Child's nervous system : ChNS : official journal of the International Society for Pediatric Neurosurgery* (2015).
- Ishii, K. *et al.* Clinical impact of the callosal angle in the diagnosis of idiopathic normal pressure hydrocephalus. *European radiology* **18**, 2678–2683 (2008).
- Fazekas, F., Chawluk, J. B., Alavi, A., Hurtig, H. I. & Zimmerman, R. A. MR signal abnormalities at 1.5 T in Alzheimer's dementia and normal aging. *AJR American journal of roentgenology* **149**, 351–356 (1987).
- Nagata, S., Rhoton, A. L. Jr. & Barry, M. Microsurgical anatomy of the choroidal fissure. *Surgical neurology* **30**, 3–59 (1988).
- Zemmoura, I., Velut, S. & Francois, P. The choroidal fissure: anatomy and surgical implications. *Advances and technical standards in neurosurgery* **38**, 97–113 (2012).
- Bakker, E. N. *et al.* Lymphatic Clearance of the Brain: Perivascular, Paravascular and Significance for Neurodegenerative Diseases. *Cellular and molecular neurobiology* **36**, 181–194 (2016).
- Brinker, T., Stopa, E., Morrison, J. & Klinge, P. A new look at cerebrospinal fluid circulation. *Fluids and barriers of the CNS* **11**, 10 (2014).
- Kida, S., Pantazis, A. & Weller, R. O. CSF drains directly from the subarachnoid space into nasal lymphatics in the rat. Anatomy, histology and immunological significance. *Neuropathology and applied neurobiology* **19**, 480–488 (1993).
- Koh, L., Zakharov, A. & Johnston, M. Integration of the subarachnoid space and lymphatics: is it time to embrace a new concept of cerebrospinal fluid absorption? *Cerebrospinal fluid research* **2**, 6 (2005).
- McComb, J. G. Recent research into the nature of cerebrospinal fluid formation and absorption. *Journal of neurosurgery* **59**, 369–383 (1983).
- Tarasoff-Conway, J. M. *et al.* Clearance systems in the brain-implications for Alzheimer disease. *Nature reviews Neurology* **11**, 457–470 (2015).

29. Yamada, S., DePasquale, M., Patlak, C. S. & Cserr, H. F. Albumin outflow into deep cervical lymph from different regions of rabbit brain. *The American journal of physiology* **261**, H1197–1204 (1991).
30. Weller, R. O. Pathology of cerebrospinal fluid and interstitial fluid of the CNS: significance for Alzheimer disease, prion disorders and multiple sclerosis. *Journal of neuropathology and experimental neurology* **57**, 885–894 (1998).
31. Weller, R. O., Galea, I., Carare, R. O. & Minagar, A. Pathophysiology of the lymphatic drainage of the central nervous system: Implications for pathogenesis and therapy of multiple sclerosis. *Pathophysiology: the official journal of the International Society for Pathophysiology/ISP* **17**, 295–306 (2010).
32. Cushing, H. The third circulation and its channels. *Lancet* **2**, 851–857 (1925).
33. Davson, H., Domer, F. R. & Hollingsworth, J. R. The mechanism of drainage of the cerebrospinal fluid. *Brain: a journal of neurology* **96**, 329–336 (1973).
34. Milhorat, T. H. The third circulation revisited. *Journal of neurosurgery* **42**, 628–645 (1975).
35. Ichimura, T., Fraser, P. A. & Cserr, H. F. Distribution of extracellular tracers in perivascular spaces of the rat brain. *Brain research* **545**, 103–113 (1991).
36. Iliff, J. J. *et al.* Cerebral arterial pulsation drives paravascular CSF-interstitial fluid exchange in the murine brain. *The Journal of neuroscience: the official journal of the Society for Neuroscience* **33**, 18190–18199 (2013).
37. Carare, R. O. *et al.* Solutes, but not cells, drain from the brain parenchyma along basement membranes of capillaries and arteries: significance for cerebral amyloid angiopathy and neuroimmunology. *Neuropathology and applied neurobiology* **34**, 131–144 (2008).
38. Hawkes, C. A. *et al.* Perivascular drainage of solutes is impaired in the ageing mouse brain and in the presence of cerebral amyloid angiopathy. *Acta neuropathologica* **121**, 431–443 (2011).
39. Iliff, J. J. *et al.* A paravascular pathway facilitates CSF flow through the brain parenchyma and the clearance of interstitial solutes, including amyloid beta. *Science translational medicine* **4**, 147ra111 (2012).
40. Weller, R. O., Djuanda, E., Yow, H. Y. & Carare, R. O. Lymphatic drainage of the brain and the pathophysiology of neurological disease. *Acta neuropathologica* **117**, 1–14 (2009).
41. Johnston, M., Zakharov, A., Koh, L. & Armstrong, D. Subarachnoid injection of Microfil reveals connections between cerebrospinal fluid and nasal lymphatics in the non-human primate. *Neuropathology and applied neurobiology* **31**, 632–640 (2005).
42. Louveau, A. *et al.* Structural and functional features of central nervous system lymphatic vessels. *Nature* **523**, 337–341 (2015).
43. Iliff, J. J., Goldman, S. A. & Nedergaard, M. Implications of the discovery of brain lymphatic pathways. *The Lancet Neurology* **14**, 977–979 (2015).
44. Bradley, W. G. Jr. CSF Flow in the Brain in the Context of Normal Pressure Hydrocephalus. *AJNR American journal of neuroradiology* **36**, 831–838 (2014).
45. Bunck, A. C. *et al.* Magnetic resonance 4D flow characteristics of cerebrospinal fluid at the craniocervical junction and the cervical spinal canal. *European radiology* **21**, 1788–1796 (2011).
46. El Sankari, S. *et al.* Cerebrospinal fluid and blood flow in mild cognitive impairment and Alzheimer's disease: a differential diagnosis from idiopathic normal pressure hydrocephalus. *Fluids and barriers of the CNS* **8**, 12 (2011).
47. Gupta, S. *et al.* Cerebrospinal fluid dynamics in the human cranial subarachnoid space: an overlooked mediator of cerebral disease. I. Computational model. *Journal of the Royal Society, Interface / the Royal Society* **7**, 1195–1204 (2010).
48. Yamada, S. *et al.* Influence of respiration on cerebrospinal fluid movement using magnetic resonance spin labeling. *Fluids and barriers of the CNS* **10**, 36 (2013).
49. Yamada, S. *et al.* Current and emerging MR imaging techniques for the diagnosis and management of CSF flow disorders: a review of phase-contrast and time-spatial labeling inversion pulse. *AJNR American journal of neuroradiology* **36**, 623–630 (2015).
50. Yamada, S. & Kelly, E. Cerebrospinal Fluid Dynamics and the Pathophysiology of Hydrocephalus: New Concepts. *Seminars in ultrasound, CT, and MR* **37**, 84–91 (2016).
51. Flood, C. *et al.* Transforming growth factor-beta1 in the cerebrospinal fluid of patients with subarachnoid hemorrhage: titers derived from exogenous and endogenous sources. *Journal of cerebral blood flow and metabolism: official journal of the International Society of Cerebral Blood Flow and Metabolism* **21**, 157–162 (2001).
52. Suzuki, H. *et al.* Cerebrospinal fluid ferritin in chronic hydrocephalus after aneurysmal subarachnoid hemorrhage. *Journal of neurology* **253**, 1170–1176 (2006).
53. Takizawa, T. *et al.* Inflammatory cytokine cascade released by leukocytes in cerebrospinal fluid after subarachnoid hemorrhage. *Neurological research* **23**, 724–730 (2001).
54. Li, J. *et al.* Communicating hydrocephalus in adult rats with kaolin obstruction of the basal cisterns or the cortical subarachnoid space. *Experimental neurology* **211**, 351–361 (2008).
55. Nagra, G. *et al.* Impaired lymphatic cerebrospinal fluid absorption in a rat model of kaolin-induced communicating hydrocephalus. *American journal of physiology Regulatory, integrative and comparative physiology* **294**, R1752–1759 (2008).
56. Jusue-Torres, I. *et al.* A Novel Experimental Animal Model of Adult Chronic Hydrocephalus. *Neurosurgery* **79**, 746–756 (2016).
57. Motohashi, O. *et al.* Thrombin and TGF-beta promote human leptomeningeal cell proliferation *in vitro*. *Neuroscience letters* **190**, 105–108 (1995).
58. Falcao, A. X. & Udupa, J. K. A 3D generalization of user-steered live-wire segmentation. *Medical image analysis* **4**, 389–402 (2000).
59. Zhang, L. & Wu, X. An edge-guided image interpolation algorithm via directional filtering and data fusion. *IEEE transactions on image processing: a publication of the IEEE Signal Processing Society* **15**, 2226–2238 (2006).
60. Gao, K. C., Nair, G., Cortese, I. C., Koretsky, A. & Reich, D. S. Sub-millimeter imaging of brain-free water for rapid volume assessment in atrophic brains. *NeuroImage* **100**, 370–378 (2014).

Acknowledgements

We would like to thank the staff at the Department of Radiology and Rehabilitation in the Rakuwakai Otowa Hospital. We would like to thank Dr. Shinya Yamada, Department of Neurosurgery, Toshiba Rinkan Hospital, for his advice.

Author Contributions

S.Y. is the corresponding author for this study and the principal investigator. S.Y. designed the study, collected, analysed and interpreted the data, conducted literature searches, prepared figures, and wrote the manuscript. S.Y. takes responsibility for data management, the accuracy of statistical analysis, the conduct of the research, and drafting of the manuscript. M.I., Y.I. and K.Y. have made substantial contributions to the intellectual content of the paper, have approved the final manuscript, and agree with submission to this journal.

Additional Information

Supplementary information accompanies this paper at <http://www.nature.com/srep>

Competing financial interests: We do not have any grant or financial support for this manuscript. Dr. Ishikawa has a grant from Japan's Ministry of Health and Welfare as a member for the research for normal pressure

hydrocephalus (number: 2014-Nanchi-Genera 1-052), and a consulting fee or honorarium: Johnson and Johnson, Medtronic and Nihon-Medi-Physics, as a speaker at seminars. Drs Yamada, Iwamuro and Yamamoto report no disclosures and no conflicts of interest.

How to cite this article: Yamada, S. *et al.* Choroidal fissure acts as an overflow device in cerebrospinal fluid drainage: morphological comparison between idiopathic and secondary normal-pressure hydrocephalus. *Sci. Rep.* **6**, 39070; doi: 10.1038/srep39070 (2016).

Publisher's note: Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.



This work is licensed under a Creative Commons Attribution 4.0 International License. The images or other third party material in this article are included in the article's Creative Commons license, unless indicated otherwise in the credit line; if the material is not included under the Creative Commons license, users will need to obtain permission from the license holder to reproduce the material. To view a copy of this license, visit <http://creativecommons.org/licenses/by/4.0/>

© The Author(s) 2016