Assessment of Cerebrospinal Fluid Hydrodynamics Using Magnetic Resonance Imaging in Postcraniospinal Surgery Patients

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Abstract

Objective  Aim of this study is to evaluate the effect of craniospinal interventions on cerebrospinal fluid (CSF) flow hydrodynamics and study the correlation of postoperative changes in flow alteration with clinical outcome.

Materials and Methods  Fifty patients who underwent various craniospinal procedures were studied using conventional and phase-contrast magnetic resonance imaging (PCMRI) protocol. CSF flow quantification was performed at cerebral aqueduct, foramen magnum, C2–3, and D12–L1 vertebral levels with site showing maximal alteration of CSF flow dynamics considered as the region of interest. Velocity encoding was kept at 20 cm/s. Patients with pathology at craniovertebral junction were considered separately (group I) from others (group II) due to different flow dynamics. Follow-up scans were performed after an interval of 1 month for temporal evaluation of changes in CSF flow dynamics.

Results  Patients in both groups showed a significant change in peak CSF velocity postoperatively (mean change of 1.34 cm/s in group I and 0.28 cm/s in group II) with bidirectional improvement in flow on cine-phase-contrast qualitative images. Regional pain (82%) and headache (46%) were seen in most of the patients preoperatively. Postoperatively clinical symptoms improved in 59.5%, static in 26.2%, and worsened in 14.3%. In both the groups, an improvement in clinical symptomatology had significant correlation with mean changes in peak CSF velocity postoperatively (p = 0.04 in both groups).

Conclusion  PCMRI can effectively evaluate changes in CSF flow noninvasively both pre- and postoperatively. This may have potential role in determining clinical outcome and prognosis of patients undergoing procedures in craniospinal axis.

Keywords
- cerebrospinal fluid
- craniospinal
- PCMRI

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Introduction

Cerebrospinal fluid (CSF) shows complex oscillatory flow during the cardiac cycle across different levels with areas of relative change in velocity and flow patterns.1 CSF flow may be obstructed to a varying degree in different pathologies leading to a wide array of clinical and neuroradiological presentations requiring various surgical interventions.2 It is imperative to develop noninvasive predictors of surgical outcome by imaging so as to prognosticate about the potential benefits. Ongoing advancements in magnetic resonance imaging (MRI) and development of phase-contrast MRI (PCMRI) have allowed a qualitative and quantitative assessment of CSF flow which is accurate and reproducible.

There is only limited literature available outlining the clinical applications of PCMRI in determining the changes of CSF flow in postsurgical scenario with no recognized literature on temporal evaluation in such patients.3,4 The present study evaluated the alteration of the CSF flow hydrodynamics postsurgically and its correlation with clinical outcome.

Materials and Methods

The study was a longitudinal study with a case-control design comprising 50 patients and 25 controls, conducted over a period of 2 years in a tertiary-care hospital in India. Patients with various craniospinal pathologies causing CSF flow alterations undergoing surgeries irrespective of age and gender comprised the study group. Prior approval of the Institutional Ethical Committee (IEC approval date: May 28, 2018) was taken. Healthy subjects formed the control group.

Data Collection Techniques and Tools

Conventional magnetic resonance imaging of the brain was performed before PCMRI. For qualitative PCMRI, a two-dimensional fast low-angle shot (FLASH) sequence was used. T2-weighted sagittal images, phase, rephase, and magnitude images were acquired. Imaging parameters were repetition time/echo time (TR/TE) = 34.45/9.73 ms, field of view (FOV) = 240 mm, flip angle = 10 degrees, and slice thickness = 4.5 mm. CSF flow dynamics were quantitatively studied by using a prospective cardiac-gated (pulse trigger being used for cardiac synchronization) high-resolution axial phase-contrast protocol with an optimal imaging plane (MRI) and development of phase-contrast MRI (PCMRI) have allowed a qualitative and quantitative assessment of CSF flow which is accurate and reproducible.

Results

Fifty patients and 25 controls were included in the study. The mean age of patients was 42.36 ± 21.38 and of controls was 38.56 ± 18.69 years.

Table 1  Cumulative disability scoring of clinical symptoms

<table>
<thead>
<tr>
<th>Group</th>
<th>Number of patients</th>
<th>Preoperative disability score</th>
<th>Postoperative disability score</th>
</tr>
</thead>
<tbody>
<tr>
<td>Improved</td>
<td>25</td>
<td>15.88</td>
<td>9.6</td>
</tr>
<tr>
<td>Static</td>
<td>11</td>
<td>14.04</td>
<td>13.04</td>
</tr>
<tr>
<td>Worsened</td>
<td>6</td>
<td>14.9</td>
<td>15.7</td>
</tr>
</tbody>
</table>

The primary presenting symptoms in study participants were regional pain (82%) and headache (46%). Other symptoms included limb weakness (24%), visual disturbance (12%), and gait disturbance (8%). Postoperative imaging follow-up...
was done in 44 patients after an interval of 1 month. Two infants did have postsurgical follow-up. However, accurate assessment of clinical symptomatology improvement, especially pain, was not achieved. Postoperatively, a majority (59.5%) of the patients showed improvement in clinical symptoms. Eleven patients (26.2%) showed no change, while six patients (14.3%) showed deterioration. A greater disability score reduction favored an improved clinical outcome (Table 1).

Phase-Contrast Magnetic Resonance Imaging
Quantitative parameters measured at cerebral aqueduct, foramen magnum, C2–3, and D12–L1 vertebral levels, in controls, showed no significant difference in peak CSF velocities. An overview of the calculated parameters of CSF peak velocity is compiled in Table 2. Qualitative analysis on cine MR images showed normodynamic pulsatile bidirectional flow with typical sinusoidal waveforms. However, in patients, a strikingly high variability of flow pattern and quantitative values was observed, particularly at ROI, with an irregular low-amplitude nonpulsatile waveform at site of alteration in most of the patients. The highest peak CSF velocity was noted in cervical SAS ventrally in both patients and controls, that is, 3.04 and 3.25 cm/s, respectively.

At ROI, an overall higher peak CSF velocity was obtained postoperatively (postoperatively: 2.51 cm/s vs. preoperatively: 2.16 cm/s). However, the comparative difference was insignificant owing to large range of altered flow parameters.

Patients were divided into two subgroups due to differences in pattern of flow alteration, that is, group I: ROI at craniocervical junction and group II: ROI at aqueduct, cervical, and dorsolumbar canal. Groups I and II both showed improvement in CSF flow at ROI on dynamic cine MR images in most of the patients. However, a varied response of change in CSF peak velocity was noted to surgical intervention. Group I showed an overall reduction in peak CSF velocity postoperatively (mean reduction of ~1.34 cm/s), and in group II, an overall increase (mean increase of ~0.28 cm/s) was observed.

Comparison of quantitative values in patients pre- and postoperatively at ROI showed significant differences in values of peak CSF velocity in both groups I and II which showed a significant correlation \( p = 0.04 \) in both with improvement in clinical outcome (improved versus others) as illustrated

<table>
<thead>
<tr>
<th>Region of interest</th>
<th>Control Peak systolic velocity (cm/s)</th>
<th>Patient (preoperative) Peak systolic velocity (cm/s)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Aqueduct</td>
<td>2.14</td>
<td>2.28</td>
</tr>
<tr>
<td>Foramen magnum (ventral)</td>
<td>2.74</td>
<td>2.14</td>
</tr>
<tr>
<td>Foramen magnum (dorsal)</td>
<td>2.80</td>
<td>2.74</td>
</tr>
<tr>
<td>C2–3 (ventral)</td>
<td>2.32</td>
<td>2.80</td>
</tr>
<tr>
<td>C2–3 (dorsal)</td>
<td>2.10</td>
<td>2.32</td>
</tr>
<tr>
<td>D12–L1 (ventral)</td>
<td>3.04</td>
<td>2.10</td>
</tr>
<tr>
<td>D12–L1 (dorsal)</td>
<td>3.25</td>
<td>3.04</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Patient group</th>
<th>Clinical outcome</th>
<th>n</th>
<th>Mean change in peak CSF velocity (cm/s)</th>
<th>SD</th>
<th>( p )</th>
</tr>
</thead>
<tbody>
<tr>
<td>Group I</td>
<td>Improved</td>
<td>2</td>
<td>2.24</td>
<td>0.08</td>
<td>0.04a (improved vs. others)</td>
</tr>
<tr>
<td></td>
<td>Others</td>
<td>2</td>
<td>0.55</td>
<td>0.01</td>
<td></td>
</tr>
<tr>
<td>Group II</td>
<td>Improved</td>
<td>21</td>
<td>0.28</td>
<td>0.49</td>
<td>0.04a (improved vs. others)</td>
</tr>
<tr>
<td></td>
<td>Others</td>
<td>16</td>
<td>0.02</td>
<td>0.10</td>
<td></td>
</tr>
</tbody>
</table>

Abbreviations: CSF, cerebrospinal fluid; SD, standard deviation.

Significant.
in Table 3. In addition, patients with clinical improvement showed a higher percentage increase in peak CSF velocity (28.9%) versus those who demonstrated worsening or no change (7%). Fig. 1 illustrates reduced flow across cerebral aqueduct in a 78-year-old male with subdural hematoma showing improvement in flow in cine MR across cerebral aqueduct postoperatively and corresponding improvement clinical symptoms. In addition, the changes in CSF flow dynamics in a 25-year-old female with cerebellar tonsillar herniation and cervical syrinx is shown in Fig. 2.

Furthermore, an increase was seen in area of SAS postoperatively at ROI (mean area of SAS preoperatively: 1.10 ± 1.13 cm² and postoperatively: 1.19 ± 1.18 cm²) which was found to be statistically significant (p = 0.03). The postoperative increase in area of SAS in a 64-year-old female with a dorsal intraspinal extramedullary lesion with resultant normalization of flow and increase in CSF peak velocity (preoperatively, −2.76 cm/s and postoperatively, −3.57 cm/s) and increase in area of SAS (−0.64 cm²) has been shown in Fig. 3. Similarly, postoperative increase in area of SAS in a 44-year-old male patient with degenerative disc bulge and corresponding improvement in CSF flow dynamics is shown in Fig. 4.

Discussion

The increasingly widespread capabilities and availability of MR imaging over the past decade have allowed better assessment of craniospinal disorders. PCMRI allows noninvasive evaluation of quantitative biomarkers, as well as dynamic information, pertaining to CSF flow alterations allowing postsurgical assessment. To the best of our knowledge, assessment of CSF flow dynamics at multiple craniospinal levels and role of temporal assessment in such patients is as yet unevaulated, wherein lies the strength of our methodology.

Increased emphasis has been placed on evaluation of CSF peak velocities in healthy volunteers; however, the reference standards at spinal levels are lacking. In this study, in control patients, the CSF velocities in the SAS across various levels were relatively uniform and did not present with any relevant differences. Similar parameters were reported in a study done by Wang et al inclusive of intraspinal CSF dynamics with measurements at foramen magnum, C7, and T7 levels. The small differences in CSF flow parameters at various levels are multivariate and can be attributed to anatomy of the CSF spaces, blood pressure, heart rate, compliance of surrounding tissues, and possible systematic errors like selecting the ROI.
which can be subject to interobserver variations. To reduce interobservers’ variability, the averages of mean quantitative values were taken and in case of significant inconsistencies, the values were confirmed by another investigator. Further peak CSF velocity was used for comparisons among the different flow parameters, as it has shown less interobserver variations and less sensitive to systematic errors as mentioned in previous studies.8

The changes of metrics in at ROI in site of pathology corroborate favorably with alteration of parameters reported by previous investigators.6 Twenty-seven of our patients had intracranial pathology, while 23 had intraspinal pathology. Intracranial flow alteration in our study included patients with intracranial/intraventricular masses altering CSF flow. We observed reduced/absent flow on cine MR images with reduced CSF quantitative parameters and irregular low-amplitude peak velocity waveforms in both groups. Postoperatively, there was increase in pulsatile CSF flow with increase in quantitative values. Mass lesions, whether infratentorial or supratentorial, both indirectly caused aqueduct obstruction by virtue of compression of the posterior third ventricle or cerebral aqueduct. The degree of CSF flow compression showed relative diminution of flow on cine MR images. Increased pulsatile flow was noted from the inferior portion of the fourth ventricle into the cisterna magna suggesting flow redistribution. This was consistent with study by Quencer et al which showed similar changes allowing assessment of level of obstruction based on changes in flow patterns.9

Fig. 3 (A-F) A 64-year-old female with a dorsal intraspinal extra- medullary lesion. Intraspinal extramedullary lesion in dorsal region preoperatively (elbow arrow) in T2-WI sagittal (A) and axial (B). Qualitative magnitude images showing reduced flow ventrally and dorsally (C). Postoperative resolution of lesion (D) with increase in SAS (E) and improvement in flow on magnitude images (F) (arrows). SAS, subarachnoid spaces; T2-WI, T2-weighted image.

Fig. 4 (A-F) A 44-year-old male with degenerative disc bulge. (A) Degenerative disc bulge at C5–6 intervertebral level (black arrow) on sagittal T2-WI causing spinal canal narrowing and cervical cord compression with reduced flow on qualitative magnitude images (B). Reduced quantitative flow parameters. (C) Postoperative reduction in disc bulge (D) and improvement in CSF flow on qualitative magnitude images (E) and increase in peak CSF velocity (F). CSF, cerebrospinal fluid; T2-WI, T2-weighted image.
Different pathogenetic mechanisms have been proposed causing varying alteration of CSF flow at cranio cervical junction and rest of the SAS. It was suggested by Bhadelia et al and Oldfield et al that the rapid caudal tonsillar movements in Chiari’s malformation act as a piston leading to increased systolic velocities and in contrast elsewhere a more obstructive nature of flow is seen leading to reduction in flow velocity.\textsuperscript{10,11} Postoperatively, in our study following surgical intervention, a different response in quantitative biomarkers was seen in group-I patients. The complex CSF flow dynamics at the foramen magnum, the anterior, and posterior cervical spinal canal in patients with tonsillar herniation plausibly alters the dynamics in a different manner, probably due to abnormal brainstem motion and tonsillar piston pressure effect. Adhesions obstructing or redirecting flow may also alter CSF dynamics in addition to tonsillar descent. The discrepancies in postoperative changes in quantitative values across the levels might be attributed to the regional variability in CSF dynamics instead of degree of CSF flow alteration.\textsuperscript{12,13}

Intraspinal flow alterations in our study comprised those with flow alteration sequelae to injury/collection or mass lesion. In all such cases, we observed pattern of CSF flow alteration similar to intracranial pathologies both qualitatively and quantitatively at ROI with postoperative elevation in CSF peak velocities. The narrowing or obliteration of flow at spinal SAS leads to a nonlaminar/absent CSF flow with redistribution of pressure dynamics. In such patients, surgical decompression increases the spinal canal volume and causes widening of SAS with resultant augmentation of flow at the region of compression with regularization of pulsatility of temporal CSF velocity waveform. Qualitative dynamic cine images demonstrated a return to near-normal CSF dynamics on follow-up in most of the patients, reflecting the predicted mechanical response of CSF flow pulsations to decompressive surgery.

**Clinical Symptomatology and Flow Dynamics**

Postoperatively, there was a significant correlation in improvement in clinical outcome with change in CSF velocity in groups I and II, increasing with degree of mean change in CSF velocity. Improvement of disability score was seen in these patients. An improved CSF velocity profile following surgery in such patients may be a useful guide to anticipate a symptomatic improvement. Given the multifactorial etiology of symptoms in our patients, the improvement in surgical response observed in patients may be due to greater alteration of CSF flow as a more direct result of anatomical flow disturbance. In addition, the preoperative symptomatology may vary as per the chronicity of the lesion most commonly due to development of fibroglial adhesions leading to difference in outcome.

Further, a majority of the patients with improvement in symptoms showed a combined ventral and dorsal CSF flow abnormality on preoperative MRI. The higher improvement in patients with both ventral and dorsal flow alterations is possibly due to volumetric displacement of flow in singular flow (i.e., only either ventral or dorsal flow) alteration.

Findings were consistent with a previous study done by du Boulay.\textsuperscript{14} This may have implication in selection of patients that are preferred candidates for surgical decompression and are more likely to have a better clinical end result.

Our study had few limitations. Manual segmentation technique is prone to technical errors resulting in inaccurate estimation of CSF flow parameters. Quantitative values are underrepresented in regions of complex flow alterations. Further, spinal region is subject to pulsation artifacts and a higher signal-to-noise ratio altering value parameters.

**Conclusion**

Cine MR images provide dynamic information about level and degree of CSF flow obstruction and may be used in conjunction with conventional MR sequences for better evaluation of craniospinal flow alterations. Regional variability in CSF dynamics is seen at cranio cervical junction (CCJ) and rest of SAS with distinct change in quantitative flow values following surgical correction. Improvement in CSF flow postsurgically correspondingly reflects in clinical outcome. However, future studies inclusive of a long-term follow-up and with technical improvements in PCMR technique will help further investigate our results. Further, a larger sample size and separate evaluation of CCJ and rest of spine may improve our understanding of flow dynamics in such cases.

**Declaration of Patient Consent**

The authors certify that they have obtained all appropriate patient consent forms. In the form, the patient(s) has/have given his/her/their consent for his/her/their images and other clinical information to be reported in the journal. The patients understand that their names and initials will not be published and due efforts will be made to conceal their identity, but anonymity cannot be guaranteed.

**Financial Support and Sponsorship**

None.

**Conflicts of Interest**

There are no conflicts of interest.

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