Cerebrospinal fluid (CSF) shunt malfunction is one of the most common life-threatening neurosurgical conditions. In the emergency department (ED), imaging techniques to identify shunt malfunction include the shunt series (SS) and CT scanning of the head. We sought to determine the test characteristics of the SS and CT scan for identifying children with shunt malfunction.

Methods: We retrospectively reviewed the medical records of children with a CSF shunt who presented to our tertiary care pediatric emergency department and received an SS during a 2-year period from Jan. 1, 2001, to Dec. 31, 2002. A pediatric neuroradiologist reviewed all SS and CT scans. We defined shunt malfunction as present if the child underwent operative shunt revision.

Results: We identified 437 ED visits by 280 children. Forty-seven SS were read as abnormal. A CT scan was performed in 386 (88.3%) cases and 80 were abnormal. Shunt malfunction was identified in 131 (30.0%) children. Sensitivity, specificity, positive predictive value, negative predictive value, positive likelihood ratio and negative likelihood ratio of the SS for identifying cases of shunt malfunction were 30.0%, 95.8%, 72.3%, 75.1%, 7.1 and 0.7, respectively; for the CT scan, they were 61.0%, 82.7%, 75.1%, 72.3%, 80.5%, 3.5 and 0.5, respectively.

Conclusion: Neuroimaging has a low sensitivity for identifying shunt malfunction. Neurosurgical consultation should be sought if shunt malfunction is clinically suspected, despite normal imaging.

Keywords: shunt series, CT scan, hydrocephalus, emergency department

Résumé

Le dysfonctionnement du dispositif de dérivation appelé « shunt » du liquide céphalorachidien (LCR) est l'une des principales causes de complications neurochirurgicales mettant la vie du patient en danger. Dans les urgences, les techniques d'imagerie médicale utilisées pour repérer...
Introduction

Cerebrospinal fluid (CSF) shunting is one of the most commonly performed neurosurgical operations. For individuals without normal CSF drainage, CSF shunting allows for the egress of fluid out of the head. Thus CSF shunting protects the patient from a life-threatening build up of intracranial pressure by diverting CSF to an alternative location (e.g., the peritoneal cavity) through a synthetic tube. Complications that can lead to life-threatening shunt malfunction and presentation to the emergency department (ED) include disconnection, calcification, migration of the shunt from its intended position and blockage.1,2 Identifying patients with shunt malfunction and obtaining prompt neurosurgical consultation for shunt revision is an important task for emergency physicians.

Neuroimaging is typically obtained to help in identifying patients with shunt malfunction.3-5 Imaging usually includes a shunt series (SS) and a CT scan of the head. An SS is a set of plain radiographs of the entire course of the shunt tubing (e.g., skull, chest and abdominal radiographs). Possible causes of shunt malfunction, such as tube disconnection, fracture, calcification or migration of the tip from the intended end point may be identified on the SS. If shunt malfunction is present, an expected finding on CT scan of the head is absolute or relative ventriculomegaly. Currently, there is little evidence to support the use of this neuroimaging to identify children with shunt malfunction.5-7

We sought to determine the test characteristics of the SS and CT scan for identifying children with shunt malfunction.

Methods

Our facility is a tertiary care children’s hospital that receives an average of 50,000 emergency visits annually. It is a common practice in our ED to evaluate a child with suspected shunt blockage by first performing an SS. Then, if deemed necessary by the ED physician and the neurosurgical team, a CT scan is obtained. We retrospectively reviewed the medical records of all patients who presented to our pediatric ED with a shunting device and had an SS obtained between Jan. 1, 2001, and Dec. 31, 2002. The patient’s hard copy or electronic charts or both were reviewed by 2 investigators (A.M. and S.A.). We defined a case as any visit to our ED during the study period during which a child underwent an SS. Therefore, owing to multiple visits during the study period by some children, a single child could contribute more than 1 case in our study. Patients were excluded if they had received an incomplete SS in the ED or if clinical data were missing from the chart. Patients were also excluded if they visited the ED within 2 weeks of a previous visit for the same chief complaint.

Collected data included demographic data, a history of the present illness and pertinent past medical history. Data regarding patient disposition, clinical outcome during hospital admission and surgical procedures were also included.

A pediatric neuroradiologist (M.S.) simultaneously reviewed all SS and CT scan films for each study subject and compared these studies with prior radiographic images, if available. The pediatric neuroradiologist was blinded to previous radiologic reports of the SS and CT and to the
outcome of the patients. Findings on SS and CT scans were categorized. We grouped the SS findings into 4 categories: normal, discontinuity in the shunt tubing, a kink in the shunt tubing or some “other” abnormality. We grouped the CT findings into 6 categories: normal, hydrocephalus without previous CT for comparison, hydrocephalus that was unchanged from a prior comparison CT scan, increased hydrocephalus when compared with a prior CT scan, improved hydrocephalus compared with a prior CT scan or some “other” abnormality. Subsequently, we included hydrocephalus that was unchanged from a prior comparison CT scan and improved hydrocephalus compared with a prior CT scan in the “normal” category.

Our main outcome measure was the presence of shunt malfunction. If the neurosurgeon performed a shunt revision, we defined this as shunt malfunction. The neurosurgeons were not blinded to the findings on SS and CT scan when making their decision about performing a shunt revision.

Data were entered into Microsoft Excel 2003 (Microsoft Corporation, Redmond, Washington). SPSS for windows (version 13.0, SPSS, Inc. Chicago, Illinois) was used for data analysis. Sensitivity, specificity, positive and negative predictive values and likelihood ratios were calculated. Our study was approved by the Research Ethics Board at the Hospital for Sick Children in Toronto, Ontario.

Table 1. Study subject enrolment by number of visits

<table>
<thead>
<tr>
<th>No. of visits per child</th>
<th>No. of children</th>
<th>Total no. of visits</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>189</td>
<td>189</td>
</tr>
<tr>
<td>2</td>
<td>56</td>
<td>112</td>
</tr>
<tr>
<td>3</td>
<td>20</td>
<td>60</td>
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<td>20</td>
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<td>5</td>
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<td>30</td>
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<tr>
<td>6</td>
<td>2</td>
<td>12</td>
</tr>
<tr>
<td>7</td>
<td>2</td>
<td>14</td>
</tr>
<tr>
<td>Total</td>
<td>280</td>
<td>437</td>
</tr>
</tbody>
</table>

Table 2. Shunt malfunction by results of the shunt series

<table>
<thead>
<tr>
<th>Shunt series results; n = 437</th>
<th>Shunt malfunction</th>
<th>No shunt malfunction</th>
</tr>
</thead>
<tbody>
<tr>
<td>Normal</td>
<td>97</td>
<td>293</td>
</tr>
<tr>
<td>Discontinuity of the tubing</td>
<td>30</td>
<td>3</td>
</tr>
<tr>
<td>Kink in the tubing</td>
<td>4</td>
<td>6</td>
</tr>
<tr>
<td>Other abnormalities*</td>
<td>0</td>
<td>4</td>
</tr>
<tr>
<td>Total</td>
<td>131</td>
<td>306</td>
</tr>
</tbody>
</table>

*Other abnormalities included migration of the distal end of the tubing into the scrotum (1 case), migration of the distal end of the tubing into the sigmoid colon (1 case) and inconclusive results (2 cases).

Results

We identified 458 instances in which an SS was obtained during the study period. Of these, 11 were excluded because the medical record or images were incomplete or unavailable, 9 because the child had returned within 2 weeks of a previous visit for the same chief complaint and 1 case was excluded because the child was transferred from another hospital after neuroimaging had been obtained. Therefore, our main study group consisted of the remaining 437 visits. A total of 280 children were responsible for these 437 visits, and the number of visits by a single child during the study period ranged from 1 to 7 (Table 1). The majority of children enrolled in our study (186, 66.4%) visited the ED once during the study period. Of the 280 children in our study, 123 (43.9%) were girls. Their mean age at presentation to the ED was 8.4 years (range 1.8 mo to 18.8 yr). The etiology for hydrocephalus was congenital in 75 (26.8%) children, myelomeningocele in 68 (24.3%), neonatal intraventricular hemorrhage in 53 (18.9%), brain tumors in 26 (9.3%), meningitis in 13 (4.6%), trauma in 9 (3.2%) and other etiologies in 36 (12.9%).

At least 3 different types of shunts were identified in our study population. Ventriculoperitoneal shunts were seen in 420 cases (96.1%), ventriculopleural shunts in 6 (1.1%) and cistoperitoneal shunts in 5 (1.1%). Six cases (1.1%) had more than 1 type of shunt.

Of our 437 cases, one-half (219, 50.1%) were discharged home from the ED and the remainder were admitted to the hospital. Shunt malfunction was diagnosed in 131 (30.0%) cases.

Forty-seven (10.8%) SS were abnormal (Table 2). A CT scan of the head was performed in 386 (88.3%) cases. Of those children who underwent CT scanning, 41 (10.6%)...
underwent shunt revision despite having hydrocephalus that was either unchanged or improved compared with prior studies (Table 3).

Nine cases had abnormal findings on SS but not on the CT scan. Of these, 6 showed a kink in the shunt, 2 showed that the tubing had migrated into the scrotum and 1 revealed a disconnection in the shunt tubing. This child also had an abscess at the shunt site. Of these 9 cases, only this child had shunt malfunction.

The test characteristics for neuroimaging were calculated. Sensitivity, specificity, positive predictive value, negative predictive value, positive likelihood ratio and negative likelihood ratio of the SS for identifying cases of shunt malfunction were 30.0%, 95.8%, 72.3%, 75.1%, 7.1 and 0.7, respectively; for the CT scan they were 61.0%, 82.7%, 64.5%, 80.5%, 3.5 and 0.5, respectively.

**Discussion**

The traditional imaging studies used to assist in diagnosing shunt malfunction are insensitive and have relatively poor test characteristics. Our results support those of prior studies with regard to both the SS and head CT scanning. Out of 67 SS in 1 study, none were found to be abnormal, while one-third of the children had shunt malfunction. Another study found that SS had a sensitivity of 20% and a negative predictive value of 22%. In this study, the yield of routine shunt series in detecting unsuspected abnormalities was low (0.8%, 95% confidence interval 0.1%–3.0%).

With regard to head CT scanning, it is intuitively appealing to think that shunt malfunction should lead to absolute or relative ventriculomegaly in nearly all cases. As intuitive as this concept is, it is wrong. Normal CT scans, those demonstrating stable ventriculomegaly and even those with decreased ventricular size compared with prior CT scans may be seen in children experiencing shunt malfunction. In one report of 100 children who had shunt malfunction, approximately 11% of the shunt failures had brain imaging studies (CT and magnetic resonance imaging scans) showing small ventricles. Among 84 patients in another series, the false negative rate was 4% (small ventricles with a nonfunctioning shunt) and the false positive rate was 13% (large ventricles with a functioning shunt). Estimates of sensitivity of the CT scan reported in previous studies have ranged from 64% to 92%. These findings are similar to ours.

**Limitations**

Limitations of our study include its retrospective methodology, which potentially limits the generalizability of our results. We did not perform interrater reliability testing to assess the reproducibility of our findings. Also, the SS and CT scans were reviewed by a pediatric neuroradiologist. This may be different from actual practice. During the clinical care of our study subjects, a general radiologist typically provided the interpretations that impacted patient care. These readings may differ somewhat from those of a pediatric neuroradiologist. Our subject list was generated through the radiology database and it is possible that some children with shunt malfunctions came to the ED during the study period and did not have a shunt series performed.

Our main study group included several visits by individual children. If there were some unique features to the children who had multiple visits during the study period, this may overrepresent these features in our results. This could adversely affect the generalizability of our results. In addition, the neurosurgeons were not blinded to the results of the neuroimaging. Since we defined shunt malfunction as the decision by the neurosurgeon to perform a shunt revision, we are at risk for introducing incorporation bias. However, incorporation bias would lead to an overestimation of the sensitivity of the SS and CT scanning. We demonstrated relatively poor sensitivity of the SS and head CT in detecting shunt malfunction. It is conceivable that owing to incorporation bias we have overestimated the sensitivity and that the true sensitivity is even worse than found in our study.

**Conclusion**

Our findings show that diagnostic neuroimaging has a low sensitivity for shunt malfunction. Neurosurgical consultation should be sought if shunt malfunction is clinically suspected, despite normal neuroimaging.

**Competing interests:** None declared.

**References**


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Erratum

In the November 2007 issue of CJEM, Clinton T. Forsythe’s surname was mis-spelled on the cover and table of contents pages. We apologize for this error and any inconvenience it may have caused.

Reference