Chronic Subdural Hematoma after Lumboperitoneal Shunt Replacement: A Case Report from Iran

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Abstract

Background and Importance: Lumboperitoneal shunts are commonly used to treat several conditions, but their use can result in significant complications. We discussed a complication of these shunts that has not been reported in the literature to date.

Case Presentation: An obese, 41-year-old woman admitted to Rasool Akram Hospital complained of a severe headache and blurred vision. The physical examination indicated that she had papilledema, and a lumbar puncture demonstrating cerebrospinal fluid opening pressure of 33 cm of H₂O. Pseudotumor cerebi was diagnosed, and the patient underwent surgery to insert a lumboperitoneal shunt. She suffered from severe headaches and nausea during the post-operative period. The post-operative syndrome was misdiagnosed as shunt-induced cerebellar tonsillar ptoosis; however, further evaluation indicated that intracranial chronic subdural hematoma, a very rare complication subsequent to lumboperitoneal shunting, was the cause of the post-operative syndrome. Subdural hematoma was evacuated, and her symptoms disappeared thereafter.

Conclusion: Persistent symptoms, such as headache, nausea, and vomiting, after placement of a lumboperitoneal shunt should be taken seriously, and imaging, such as a CT of the brain, should be done to rule out subdural hematoma.

Keywords: Benign Intracranial Hypertension; Chronic Subdural Hematoma; Lumboperitoneal Shunt

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Background and Importance

Lumboperitoneal (LP) shunting is a method of diverting cerebrospinal fluid (CSF) from the subarachnoid space to the abdominal cavity. It has an overall incidence of 1.6/100,000 per annum [1], an unclear pathogenesis, although it has significantly higher incidence in obese females aged 20-44 at 19/100,000 annually [2]. The advantages of LP shunting include avoidance of brain penetration with the shunt catheter, access to a large CSF space in the thecal sac, and the potential (for good or ill) of a large amount of CSF drainage [3,4]. For the past 50 years, the mainstay of CSF shunting for idiopathic intracranial hypertension (IIH), also known as pseudotumor cerebi (PTC) syndrome, has been LP shunt surgery [5,6]. Patients who require a LP shunt have a pathology that predisposes to the obstruction of CSF absorption or elevates the CSF pressure but must have communication of the spinal CSF with the cranial compartment. Specific indications include communicating hydrocephalus, normal pressure hydrocephalus, and IIH (PTC), though they are not limited to it [7-9]. Although the risk of intracranial complications after LP shunt insertion surgery is minimal, various complications have already been described, including shunt obstruction (14%), shunt infection (1%), radicular leg pain (5%), dyspnea, and disturbance of consciousness that necessitates conversion to a ventriculoperitoneal (VP) shunt (1%) in patients who subsequently were noted to have Chiari malformations, shunt migration, and intracranial bleeding (2%) [3,10].

Clinical Presentation and History

An obese 41-year-old woman was admitted to Rasool Akram Hospital presenting with headache, blurred vision, anxiety, and mental depression. She had been treated with acetazolamide for management of papilledema, which was unsuccessful, so she was referred to us. Her body mass index (BMI) was 38 kg/m². Her medical history showed consumption of fluoxetine, Inderal, haloperidol, desipramine, and Artan, which were prescribed by her previous psychologist.

Diagnostic Measures

A neuro-ophthalmologist observed swelling in her bilateral optic disc. Visual field perimetry and visual acuity were in the normal range. Other general and neurological examinations did not reveal any abnormalities. The laboratory analysis of her blood showed hyperglycemia due to unmanaged diabetes mellitus (FBS: 180 mg/dl, normal range: 70-100 mg/dl) and hyperprolactinemia (Prolactin: 98 ng/mL, normal range: 2-29 ng/mL). A lumbar puncture was performed to evaluate her CSF. Although she consumed a 250-mg tablet of acetazolamide every eight hours, her lumbar puncture opening pressure was 33 cm H₂O. Biochemistry and cytological analysis of CSF were within normal limits (Protein: 10 mg/dl, Sugar: 125 mg/dl, WBC: 0/hpf, RBC: 0/hpf). Her brain MRI before and after the administration of gadolinium had no abnormal findings except for the empty sella (Figures 1 & 2).

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Treatment
As a certain case of primary spontaneous PTC (intractable to medical therapy), the patient became a candidate for LP shunt surgery to channel the CSF from the lumbar thecal sac into the peritoneal cavity. During her hospitalization, she expressed no complaints except for low back pain (LBP) at the site of surgery, which was managed with regular analgesics, such as acetaminophen. The patient was discharged three days after the procedure, and she took a trip out of the country soon thereafter.

Post-operative Complications
The patient was referred to the hospital 20 days after the surgery and complained about nausea, vomiting, and headache. Because of the misdiagnosis of LP shunt-induced cerebellar tonsillar ptosis, she was conservatively treated with medications, such as promethazine and trifluoperazine. Her nausea and headache were exacerbated, and she was readmitted. Her computed tomography (CT) scan of the brain indicated left frontotemporoparietal subacute subdural hematoma with subfalcine herniation (Figure 3). In her second surgery, the left parietal burr-hole was performed, and the subdural hematoma was evacuated. The post-operative period was uneventful, and her symptoms disappeared. Her delayed brain CT showed relief of hematoma and herniation (Figure 4).

Discussion
PTC, also called benign intracranial hypertension (BIH), is characterized by elevated intracranial pressure that is unrelated to hydrocephalus, tumor, brain edema, or any other structural entities associated with normal CSF analysis [11]. Usually, PTC presents with headache, and often there are changes in the vision of obese women of childbearing age [12]. Almost all patients with PTC have headaches, i.e., 90-94%. Their headaches are associated with a feeling of pressure and throbbing. The headaches are usually continual and occur with retro-ocular pain, and the patient may become nauseated [12,13].
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is the most-feared result of PTC, but it is usually temporary. Approximately 68-85% of such patients experience vision loss [12-15]. There are usually impairments in the visual field, the most common of which is tunnel vision [12]. The vision changes are usually explained as transient ischemia of the optic nerve due to pressure [15]. Photopsia (54%) and eye pain (44%) are other common symptoms [15]. More severe symptoms occur infrequently, but diplopia (38%) and vision loss (30%) occur in a significant number of patients [15].

There is a wide variety of treatment options for BIH. Initially, it may be treated with medications, such as acetazolamide. Subsequently, surgical and non-surgical treatments may be necessary. More aggressive measures for preventing sequelae of PTC are typically used for two groups of patients, i.e., those who continue to experience vision loss despite conservative management and those who initially have rapid loss of vision. LP shunt surgery is a method in which the elevated CSF pressure is referred to the peritoneal cavity, but it entails serious risk of complications that can range from obstruction of the shunt, shunt-related meningitis, abdominal infections, tonsillar herniation, and even death [13]. Shunts fail in about 50% of patients, and approximately 10% of patients have worsening vision after the shunt surgery. Thus, frequent neurologic follow-up is required after shunts are placed [15,16].

Many ophthalmologists advocate decompression as an alternative surgical approach, because it has fewer complications and better outcomes [13,17]. The LP shunt is inserted into the subarachnoid cavity between two of the vertebrae in the lumbar region of the spine. This cavity is also referred to as the subarachnoid space. The cavity is a spongy, tissue-filled area that surrounds the brain and spinal cord, and it contains the CSF. The shunt is placed under the skin, and it continues around the oblique muscles on one side of the body. The shunt ends at the peritoneal cavity in the abdominal area. After the LP shunt is in place, it is used to drain excess CSF from the brain via the subarachnoid cavity. The fluid is transported to the peritoneal cavity, where it is absorbed and eliminated in the urine [18].

In the presented case, there was an obese 41-year-old woman who complained of severe headache, anxiety, and blurred vision. In her physical examination, bilateral papilledema was detected. The only abnormal finding in the patient’s imaging was empty sella, and no subarachnoid space widening was observed. The diagnosis was PTC, and a LP shunt was placed. The patient was discharged three days after the procedure. She was referred to hospital 20 days later and complained of headache, nausea, and vomiting. The patient was treated conservatively with medications, such as promethazine and trifluoperazine. LP shunt-induced cerebellar tonsillar ptosis was suspected, but after the exacerbation of her symptoms, a CT of the brain was performed, and it indicated the presence of a left frontotemporoparietal, subacute, subdural hematoma with subfalcine herniation. The subdural hematoma was evacuated in the second surgery, and her symptoms disappeared thereafter.

**Conclusion**

No history of head trauma of any kind was reported in our patient. Contrary to hydrocephalic patients, the
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Authors’ Contribution
Conception and Design: All authors. Data Collection: All authors. Drafting the Article: All authors. Critical Revising the Article: All authors. Reviewing Submitted Version of Manuscript: All authors. Approving the Final Version of the Manuscript: All authors.

References