Metrizamide shuntography for evaluation of shunt malfunction in hydrocephalus.

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Metrizamide Shuntography for Evaluation of Shunt Malfunction in Hydrocephalus

Ventricular shunts were evaluated by a shuntographic technique using metrizamide for the contrast material and delayed spot films and computed tomography (CT) to evaluate shunt function. Thirty hydrocephalic patients were studied who had clinical presentations of shunt malfunction. Fifty-two shuntograms were obtained; most demonstrated the cause of shunt malfunction. No contrast-related complications resulted from the procedure. Diagnostic accuracy was greatly improved by this method, since both anatomic and functional evaluations could be performed. In patients with normal studies, unnecessary surgery was eliminated; in abnormal shuntograms, the duration of surgery and the likelihood of complications were reduced because the surgeons knew the exact anatomic derangements to be corrected.

Since July 1981, patients with clinical and computed tomographic (CT) findings suggesting ventricular shunt malfunction have been evaluated by shuntography using metrizamide as the contrast agent. Except for a few cases, CT showed hydrocephalus in all cases. The shuntogram was obtained to delineate the exact cause of shunt malfunction, thereby directing the surgical approach to the specific area of abnormality. This obviated total revision of the shunt and shortened the surgical procedure. A few symptomatic patients with normal or slightly enlarged ventricles were also evaluated, and it was shown that the patients’ problems were not related to shunt malfunction. Shuntograms also delineated some rare causes of shunt malfunction that would have been missed by clinical or CT evaluation alone. We describe our technique and report our experience in 30 patients.

Materials and Methods

The patient population comprised 30 patients with clinical presentation of shunt malfunction and/or CT documentation of hydrocephalus. Most (26) were infants and young children. A total of 52 shuntograms were obtained, since some patients required more than one revision.

Shunt Anatomy

The shunt is a mechanical conduit designed to carry excess cerebrospinal fluid (CSF) from the ventricular system or subarachnoid or cystic spaces into an absorbing cavity such as the peritoneal or pleural space, ureter, or cardiovascular system. Ventriculoperitoneal or ventriculopleural shunts were used predominantly. Although other types of shunts are available, we prefer the Pudenz valve system. The Pudenz system, like the other shunt systems, has three major internal components: (1) the ventricular or proximal catheter, (2) the reservoir and valve, and (3) the peritoneal or distal catheter.

Ventricular catheter. This is a flexible silicone catheter with a small radiopaque tip and multiple side holes in its distal segment (fig. 1A). The position of the tip of the ventricular catheter outside of the ventricles does not necessarily imply shunt malfunction, since the ventricle is drained by the side holes rather than by the tip. The proximal segment of the ventricular catheter is attached to the shunt reservoir by a plastic joint.
Reservoir chamber. The reservoir is divided into proximal and distal compartments by a thin perforated diaphragm. The proximal chamber is in continuity with the CSF, while the distal compartment is connected to the absorbing space (fig. 1B).

Distal catheter. This is a relatively long, usually radiopaque catheter with either a single distal opening or multiple slits (fig. 1C).

Shuntographic Technique

The patient is placed in a supine position with the head in a lateral position. The skin over the shunt reservoir is shaved, prepared, and draped in sterile fashion. In infants, moderate sedation is required to prevent patient motion. The patient must be kept in the supine position for some time before the study so that the intraventricular pressure exceeds the opening pressure of the shunt system at the time of the shuntogram. If this precaution is not observed, the results of the dynamic study will be inaccurate [1]. A small skin incision is made to prevent introduction of epithelium into the shunt system, and the center of the reservoir is punctured by a 25-gauge butterfly needle held perpendicular to the reservoir dome to avoid damage to the diaphragm seat; such damage could result in loss of pressure control function [2].

The tip of the needle is then gently advanced further through the diaphragm into the proximal compartment, and CSF return is noted in the butterfly tubing. For beginners, it may be desirable to position the patient’s head in such a way that the reservoir is in a tangential position to the x-ray beam so that intracranial entrance is avoided.

Once CSF return is established, pressure measurements can be made that reflect intracranial pressure. If there is no CSF return, contrast material may be injected to pinpoint the obstruction. Under fluoroscopic control, 1 or 2 ml of metrizamide (170 mg/ml) is injected.
Fig. 3.—Technique for studying distal catheter patency. A, Tip of butterfly needle is withdrawn and positioned within distal chamber. Contrast material is then injected, closing valve and filling distal catheter (arrow). B, Radiograph of abdomen. Spillage of contrast material outlines bowel loops. Tip of peritoneal catheter (arrow).

In the absence of loculation, contrast material mixes rapidly with CSF and may be seen in the dependent part of the ventricle. In cases of loculation, contrast material fills a cavity and does not diffuse into the rest of the ventricular system.

Injection of air must be avoided, since it may cause air embolism in cases of ventriculovenous shunt and because it may delay emptying, which would interfere with dynamic studies [3]. If the ventricular catheter is patent, contrast material will exit through the distal holes, and a jet phenomenon will be clearly seen (fig. 2).

The tip of the needle is now slowly withdrawn until it lies within the distal chamber. Contrast material is again injected, opacifying the entire length of the distal catheter into the absorptive cavity, while the valve prevents intraventricular reflux (fig. 3A). In the case of ventriculoperitoneal shunts, normal peritoneal spillage is seen fluoroscopically, outlining loops of bowel (fig. 3B).

In normal shuntograms, delayed films of reservoir clearance are made at 3 and 9 min. A cranial CT scan can may be obtained at 6, 12, and 24 hr after the shuntogram. Both of these methods display
the shunt's dynamic functions. In cases of anticipated complex intracranial pathology, a CT study is done immediately, since the amount of contrast material used is insufficient to evaluate the entire ventricular system with routine radiographic techniques. To avoid pooling of contrast material in the dependent part of the ventricular system before the CT scan, the patient's head must be repositioned to insure adequate CSF-metrizamide mixing.

**Results**

The shuntograms in all 30 patients were obtained without complications. In most, the cause of shunt malfunction was mechanical, including obstruction of proximal or distal catheters (figs. 4 and 5) or bending, breakage, or dislocation of the distal catheter (figs. 6 and 7). Cyst formation around the distal end of the peritoneal or pleural catheters contributed to shunt malfunction in some patients (figs. 8–10). The most important contributions of the shuntogram, however, were in those patients who had loculated hydrocephalus or intracranial cyst formation, which necessitated exact anatomic evaluation before shunt revision (figs. 11–13). The CT shuntogram demonstrated the location of the lesions and confirmed the presence or absence of communication between the cystic struc-
Fig. 10.—Pseudocyst formation around distal catheter tip within abdominal wall. A, Anteroposterior view of right abdomen after introduction of metrizamide in reservoir (R) of lumbar-peritoneal shunt. Cyst (C) around distal segment of peritoneal catheter (arrow). Suggestion of medial displacement of ascending colon by cyst. Cyst was believed to be either in extraperitoneal space or within peritoneal cavity. B, CT scan at level of cyst shows exact cyst location between external and internal oblique muscles. Minimal mass effect on ascending colon.

Fig. 11.—Suprasellar arachnoid cyst. A, Axial unenhanced cranial CT scan. Dilated frontal (straight solid arrow) and temporal (open arrow) horns. Cystic structure (C) in midline is larger than lateral ventricles, excluding possibility of its being third ventricle. Porencephalic changes in both temporal lobes (curved arrows). B, After introduction of metrizamide. Proximal catheter inadvertently entered cystic structure. No filling of lateral or third ventricle.

discussion

The introduction of ventricular shunting has greatly improved the life span of patients with hydrocephalus. The new shunt systems with unidirectional valves are very flexible, increasing both their efficiency and dramatically decreasing postshunting complications. But shunt malfunction does occur. Aside from the patient's symptoms and clinical signs, the simplest examination of shunt patency is digital compression of the shunt reservoir. In a normally functioning shunt, the reservoir chamber compresses easily and reexpands in a matter of seconds. Inability to compress the reservoir indicates obstruction of the distal catheter; failure to reexpand indicates obstruction of the proximal catheter. This test, however, is not entirely reliable. It has been shown that the reservoir may pump well despite shunt obstruction, and it may not pump well when it is working satisfactorily. A shunt may work well in the presence of increased intracranial pressure secondary to subdural hematoma or hygroma or multiloculated or isolated unilateral hydrocephalus [4–7].

A second mode of evaluation is CT, which is now widely used. Although CT demonstrates dilated ventricles or other causes of increased intracranial pressure, if CT alone is used, a small but significant number of patients with increased intracranial pressure will be missed. The size of the ventricles...
does not necessarily indicate the presence or absence of shunt malfunction.

Large ventricles may be seen in patients with working shunts who have a thin cortical mantle [8]. Likewise, malfunctioning shunts in the presence of small or normal ventricles have been observed [9-14]. It is extremely important that such patients be recognized so that proper management can be instituted. In such patients, the ventricles cannot expand despite the presence of increased intraventricular pressure. The mechanism of this phenomenon is not well understood; however, subependymal gliosis and granulation have been seen and may account for the decreased compliance of the ependyma [10, 14].

CT is also unreliable in diagnosing or accurately demonstrating some intraventricular loculations (multiloculated hydrocephalus) or intracranial cystic structures (i.e., suprasellar arachnoid cysts). Although CT may show some of the septations, because of the averaging effect it does not demonstrate thin membranes well [15-20]. A ventriculogram is needed to document the presence of loculations or cystic lesions.

Because of these limitations, other methods have been used to evaluate shunt function. These include radionuclide imaging, sonography, heat and cold transfer determination, subarachnoid infusion studies, and contrast shuntography [1, 3, 6, 21-28]. The radionuclide and contrast shuntograms can
evaluate shunt patency and function, but metrizamide CT shuntograms provide the best anatomic detail.

The documentation of shunt patency does not rule out shunt malfunction. Even with gentle injection, contrast material is introduced into the system at a pressure higher than CSF pressure; therefore, contrast material could overcome incomplete obstruction. The measurement of opening pressure before introduction of metrizamide is useful to estimate intracranial pressure. However, according to Early and Fink [29], this measurement is not reliable since the ventricular system may enlarge despite normal pressure and continue to enlarge despite the presence of a patent shunt.

It metrizamide shuntograms show an abnormality, no further functional tests are needed. However, if the shuntogram is negative, functional evaluation of shunt performance must be made to rule out shunt malfunction. So-called "deceptive patency," a patent shunt with inadequate capacity to keep the patient free of symptoms, was observed in 40% of the patients in the series of French and Swanson [1]. Functional evaluation can be performed by obtaining spot films of the shunt reservoir 3 and 9 min after completion of the shuntogram [3, 27]. Another method is to perform serial cranial CT examinations at 6, 12, and 24 hr after intraventricular introduction of metrizamide. In functioning shunts, the reservoir should be clear of any residual contrast material in 3 min. Persistence of residual contrast material at 9 min and beyond denotes shunt malfunction. Also, serial cranial CT studies should show complete clearing of intraventricular metrizamide at 24 hr [30, 31]. The presence of transepidermal absorption of CSF is another finding indicative of shunt malfunction.
Our technique combines anatomic and functional studies of the shunt system by using metrizamide as the contrast agent, delayed filming, and CT. This combination greatly improved our diagnostic accuracy and eliminated unnecessary surgery. In cases of shunt malfunction, the duration of surgery was shortened significantly by informing the surgeons of the exact anatomic derangement to be corrected.

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REFERENCES