

# Migration Of The Distal Catheter Of A Ventriculoperitoneal Shunt Into The Abdominal Wall

## Case Report

ZEKİ ŞEKERCİ, ÖMER İYİGÜN, RIZA RIZALAR, GÖKHAN BOZKURT,  
CENGİZ ÇOKLUK, CEMİL RAKUNT, FAHRETTİN ÇELİK

Ondokuz Mayıs University, Faculty of Medicine, Departments of Neurosurgery and Pediatric Surgery, Samsun - Türkiye

**Abstract :** A case of migration of a Pudenz type peritoneal shunt catheter into the abdominal wall diagnosed with computed tomography is presented. This is a rare complication of ventriculoperitoneal shunt. It seems that a foreign body in the

abdominal cavity and infection are the most important causal factors.

**Key words :** Abdominal wall, infection, ventriculoperitoneal shunt complication.

Many complications secondary to ventriculoperitoneal (VP) shunt placement have been reported (9, 16, 20, 22, 31). With infections and obstruction being the most common (10, 22, 29, 31). Intraabdominal complications are relatively rare (13), but one of is extrusion of the shunt catheter from the abdominal cavity (7, 11, 14, 15, 17, 29, 30, 34, 35).

The Raimondi spring wire catheter and less flexible silicon catheters have been implicated in most cases of extrusion or perforation (1, 8, 15, 17, 19, 21, 24, 26, 27, 28, 31). However in several cases Holter and Pudenz catheters were reported to cause such complications (11). Other causal factors have also been implicated in extrusion of the shunt from the abdominal cavity (18, 29, 33).

In this report a case of migration of a Pudenz type peritoneal catheter into the abdominal wall at the right flank is presented, and the mechanism of this rare complication is discussed.

### CASE REPORT

A 4-year-old-girl was admitted on September 17, 1992 with episodes of fever and swelling at the right flank.

A medium pressure ventriculoperitoneal shunt with a Pudenz valve had been inserted for treatment of communicating hydrocephalus secondary to tuberculous meningitis 3 years previously and antituberculosis drugs had been used for 1.5 years.

At admission, physical examination a painful cystic mass with fluctuation at the right flank was palpated. Right parietal scalp and subcostal abdominal skin incision scars secondary to V-P shunt placement were also observed.

Neurological examination was normal except mental retardation.

Initial laboratory evaluation included: Haemoglobin 12.1 gram /100 ml, leucocyte count 13 800 /mm<sup>3</sup>. Chest X-ray was normal. Cerebrospinal fluid (CSF) contained 56 mg of glucose, 187 mg protein and 110 mg of chloride per 100 ml. Gram stain of the cerebrospinal fluid revealed no organisms. Electrolytes were also within normal limits. Body temperature was 38.5 0C and sedimentation rate 20 mm/ hour.

Abdominal CT revealed a cystic mass in the abdominal wall at the right flank containing the tip

of the distal catheter of a V-P shunt (Fig. 1, Fig. 2), located about 5 cm from the subcostal incision posteriorly. When the wall of the cystic mass was incised infected material and the tip of the V-P shunt were encountered. After draining the infected material CSF dropping was seen at the tip of the distal catheter. Microscopic examination of the cystic fluid showed pleocytosis. Staining for Koch bacilli (Zeil-Nielsen) was negative as was the culture of infected material from the cystic cavity.

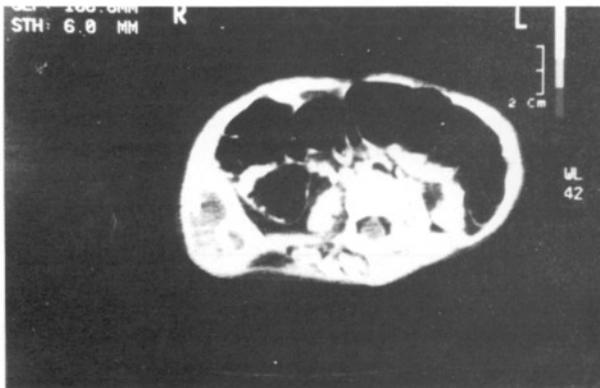


Fig. 1. CT of the cystic maas in the abdominal wall at the right flank containing distal tip of the peritoneal catheter.

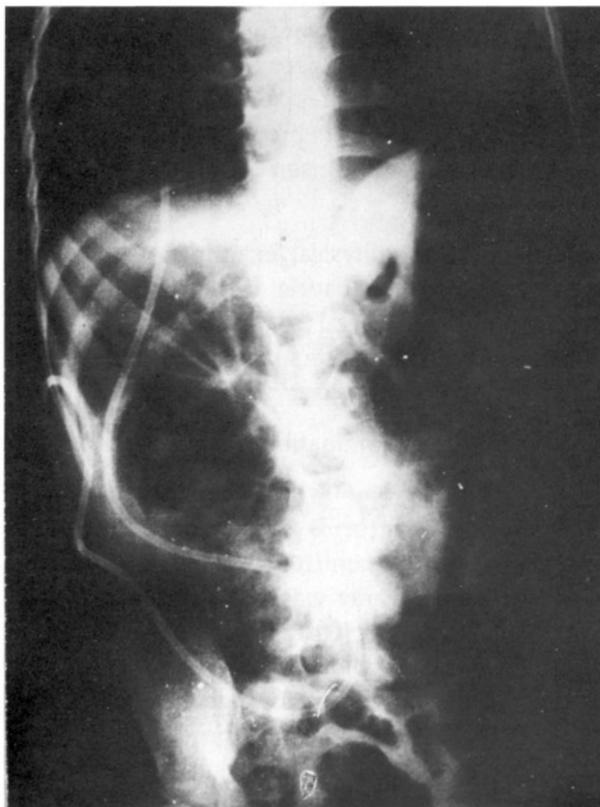


Fig. 2. Direct X-ray of peritoneal catheter. The tip of the catheter is seen at the right flank.

The peritoneal and ventricular catheters of the shunt were removed. Seftazidim was given 20 mg/kg for 20 days postoperatively. The postoperative course was uneventful.

## DISCUSSION

Intra abdominal complications associated with ventriculoperitoneal shunt are relatively rare. These include pseudocyst formation (3, 4, 13), bowel obstruction (9, 16, 23), migration of disconnected shunt (9, 12, 16, 23, 35), intractable acides (9, 29), perforation of viscera (1, 3, 13, 16, 23, 35), transdiaphragm migration (17), urethra obstruction (6), inguinal hernia (13), intrascrotal extrusion (25, 29), vaginal perforation (21, 26), umbilical perforation (2), and shunt extrusion (34).

Peritoneal catheters have been reported to extrude from the abdominal cavity through the flank (15), diaphragm (7, 11, 17), intestine (5, 29, 30), neck (34), umbilicus (2), uterus (14), vagina (21,26), abdominal incision or into the liver (28), or scrotum (20, 27). Less flexible catheters Raimondi spring wire or simple silastic have been implicated in most reported cases of extrusion (1, 15, 17, 19, 21, 28, 30) and extrusion of more flexible peritoneal catheters Holter and Pudenz from the abdominal cavity have also been reported occurred (11, 15, 29, 30).

Some authors indicate the effect of anatomical defects in the abdominal wall or diaphragm on extrusion of the shunt from the abdominal cavity (7, 33). The spring wire makes the catheter stiffer and less likely to kink, but it also seems to make it more likely to erode adjacent structures and extrude (19). Another factor in the mechanism of perforation of the abdominal wall in such cases is the presence of a foreign body in the abdominal cavity (18, 29).

Rubin suggested that the possible anchoring effect of fibrous encasement of the peritoneal end of the tube may result in repeated pressure by the tip at a fixed point on the bowel surface, eventually leading to organ perforation (29). Necrosis of the organ wall due to continuous contact with the peritoneal tube and infection at this site are other causes of perforation or extrusion (32). Diminished bowel peristaltic activity in elderly patients contributes to prolonged continuous contact of the tube with the abdominal wall (32).

In our case, the Pudenz type soft peritoneal catheter probably caused local infection and necrosis of the inner surface of the abdominal wall initially and the pulsatile effect of fluid and intestinal peristalsis eventually led to migration of the catheter into the abdominal wall and cystic dilatation.

**Correspondence :** Dr. Zeki Şekerci  
Ondokuz Mayıs Üniversitesi Tıp Fakültesi  
Nöroşirürji Anabilim dalı,  
55139, Samsun - Türkiye

### REFERENCES

1. Abu-Dalu K, Pov D, Hadani M, Sahar A (1983). Colonic complications of ventriculoperitoneal shunts. *Neurosurg*, 13: 167-170
2. Adeloye A (1973). Spontaneous extrusion of the abdominal tube through the umbilicus complicating peritoneal shunt for hydrocephalus. *J. Neurosurg* 38: 758-759
3. Agha FP, Amendola MA, Shirazi KK, Amendola BE, Chandler WF (1983). Unusual abdominal complications of ventriculoperitoneal shunts. *Radiology* 146: 323-326
4. Alther E (1965). Das magenventil: Eine neue operations methode zur behandlung des kindlichenhydrocephalus. *Schweiz Med Wochenschr* 95: 234-236
5. Arico M, Beluffi G, Fiori P, Chiari G, Pezzotta S, Pedosta AF, Bianchi E (1985). Rectal extrusion of the catheter and air ventriculography following bowel perforation in ventriculoperitoneal shunt. *Pediatr Radiol* 15: 53-55
6. Clarke CE, Paul KS, Lye RH (1983). Ventriculoperitoneal shunt procedure complicated by ureter obstruction: Case Report. *J Neurosurg* 59: 542-544
7. Cooper JR (1978). Migration of ventriculoperitoneal shunt into chest. *J. Neurosurg* 48: 146-147
8. Danişmend N, Kaday C (1989) : Unusual complication of ventriculoperitoneal shunt. Correspondance. *J. Neurosurg* 22 (4): 798
9. Davidson RI (1976). Peritoneal by pass in the treatment of hydrocephalus: Historic review and abdominal complications. *J. Neurol Neurosurg Psychiatry*. 39: 640-646
10. Forward KR, Fewer DF, Stiver HG (1983). Cerebrospinal fluid shunt infections. *J. Neurosurg* 59: 389-394
11. Fukamachi A, Wada H, Toyoda O, Wakao T, Kawafuchi J (1982). Migration or extrusion of shunt catheters. *Acta Neurochir (Wien)* 64:159-166
12. Giuffre' R, Di Lorenzo N (1975). Two unusual complications of ventriculoperitoneal shunt in the same infant *Surg Neurol*, 3: 23-24
13. Grosfeld JL, Cooney DR, Smith J, Campbell RL (1974). Intra-abdominal complications following ventriculoperitoneal shunt procedures. *Pediatrics* 54: 791-796
14. Guertin SR (1987). Cerebrospinal fluid shunts. *Pediatr Clin Nort Am*. 34: 203-217
15. Joubert MJ, Stephanov S(1983). Extrusion of peritoneal catheter through the mid lumbar region. *Surg Neurol*, 19:120-121
16. Lee FA, Gwinn JL (1985). Complications of ventriculoperitoneal shunts. *Ann Radiol (Paris)* 18: 471-478
17. Lourie H, Bajva S (1985). Transdiaphragmatic migration of a ventriculoperitoneal catheter. *Neurosurgery* 17: 324-326
18. Maingot R (1961). Abdominal operations. New York, Appleton-9 Century-Crofts, p 67
19. Mc Comb JG (1983). Rewiewer's comments. *Neurosurg* 13:169
20. Moazam F, Glenn JD, Kaplan BJ, Talbert JL, Mickle JP (1984). Inguinal hernias after ventriculoperitoneal shunt procedures in pediatric patients. *Surg Gynecol Obstet* 159: 570-572
21. Mozingo JR, Cauthen JC (1974). Vaginal perforation by a Raimondi peritoneal catheter in an adult. *Surg Neurol* 2: 195-196
22. Neetzel MJ, Baker RP(1984). Shunt fluid examination: Risks and benefits in the evaluation of shunt malfunction and infection. *J. Neurosurg* 61: 328-334
23. Nishijima M, Endoh S, Ohyama H, Higuchi H (1982). Gastric perforation by a ventriculoperitoneal shunt. *Neurosurgery*, 10:754-756
24. Oi S, Shose Y, Asano N, Oshio T, Matsumoto S (1987). Intra-gastric migration of a ventriculoperitoneal shunt catheter. *Neurosurgery* 21: 255-257
25. Oshio T, Matsumura C, Oi S (1985). Treatment of shunt tube migration into processus vaginalis in hydrocephalic children. *Shoni No Noshinkei* 10: 103-109,(in Japanese)
26. Patel CD, Matloub H (1973). Vaginal perforation as a complication of ventriculoperitoneal shunt. *J. Neurosurgery* 38: 761-762
27. Ramani PS (1974). Extrusion of abdominal catheter of ventriculoperitoneal shunt into the scrotum. *J. Neurosurg* 40: 772-773
28. Rana SR, Quivers ES, Haddy TB (1985). Hepatic cyst associated with ventriculoperitoneal shunt in a child with brain tumor. *Child's Nerv Syst* 1: 349-351
29. Rubin RC, Chatak NR, Visudhipon P(1972) A symptomatic perforated viscus and gram-negative ventriculitis as a complication of valve regulated ventriculoperitoneal shunt. *J. Neurosurgery* 37: 616-618
30. Schullhof LA, Worth RM, Kalsbeck JE (1975). Bowel perforation due to peritoneal shunt. *Surg Neurol* 3: 265-269
31. Sekhar LN, Moosy J, Guthkelch AN (1982). Malfunctioning ventriculoperitoneal shunts. *J. Neurosurg* 56: 441-416
32. Touho H, Nakauchi M, Tasawa T, Nakagawa J, Karasawa J (1987). Intrahepatic migration of a peritoneal shunt catheter: Case report. *Neurosurgery* 21: 258-259
33. Turner MS , Goodman J (1989). Extrusion of a Raimondi peritoneal catheter from the thigh. *Neurosurgery* 25: 833-835
34. Whittle IR, Johnston IH: Extrusion of peritoneal catheter through neck incision (1983). A rare complication of ventriculoperitoneal shunting. *Aust NZ J. Surg* 53: 177-178
35. Wilson CB, Bertan V (1966). Perforation of the bowel complicating peritoneal shunt for hydrocephalus: Report of two cases. *Ann Surg* 32: 601-603