Percutaneous evacuation (PEVAC) of multivesicular echinococcal cysts with or without cystobiliary fistulas which contain non-drainable material: first results of a modified PAIR method

H G Schipper, J S Laméris, O M van Delden, E A Rauws, P A Kager

Background: Surgery is the treatment of choice in echinococcal cysts with cystobiliary fistulas. PAIR (puncture, aspiration, injection, and reaspiration of scolecidals) is contraindicated in these cases.

Aim: To evaluate a modified PAIR method for percutaneous treatment of multivesicular echinococcal cysts with or without cystobiliary fistulas which contain non-drainable material.

Patients: Twelve patients were treated: 10 patients with multivesicular cysts which contained non-drainable material and were complicated by spontaneous intrabiliary rupture, secondary cystobiliary fistulas, cyst infection, or obstructed portal or hepatic veins; and two patients with large univesicular cysts and a ruptured laminated membrane, one obstructing the portal and hepatic veins and one a suspected cystobiliary fistula.

Methods: The methods used, termed PEVAC (percutaneous evacuation of cyst content), involved the following steps: ultrasound guided cyst puncture and aspiration of cyst fluid to release intracystic pressure and thereby to avoid leakage; insertion of a large bore catheter; aspiration and evacuation of daughter cysts and endocyst by injection and reaspiration of isotonic saline; cystography; injection of scolecidals only if no cystobiliary fistula was present; external drainage of cystobiliary fistulas combined with endoprosthesis or sphincterotomy; catheter removal after complete cyst collapse and closure of the cystobiliary fistula.

Results: In all 12 patients initial cyst size was 13.1 (6–20) cm (mean [range]). At follow up 17.9 (4–30) months after PEVAC, seven cysts had disappeared and five cysts had decreased to 2.4 (1–4) cm (p=0.002). In eight patients with multivesicular cysts, a cystobiliary fistula, and infection, cyst size was 12.5 (6–20) cm, catheter time 72.3 (28–128) days, and hospital stay 38.1 (20–55) days. At 17.3 (4–28) months of follow up, six cysts had disappeared and in two cysts residual size was 1 and 2.9 cm, respectively (p=0.012). In four patients without a cystobiliary fistula, cyst size was 14.4 (12.7–16) cm, catheter time 8.8 (3–13) days, and hospital stay 11.5 (8–14) days. At 19.3 (9–30) months of follow up, one cyst had disappeared and three cysts were 85 (69–94)% smaller (2.2 (1–4) cm) (p=0.068).

Conclusion: PEVAC is a safe and effective method for percutaneous treatment of multivesicular echinococcal cysts with or without cystobiliary fistulas which contain non-drainable material.

PAIR (puncture, aspiration, injection, and reaspiration of a scolecidal agent) of echinococcal cysts in the liver is an invasive procedure with a low complication rate (10%), a high success rate (>95%), and a low relapse rate (<4%).

PAIR is safe and simple to perform even in poorly equipped clinics in developing countries. In comparison with albendazole treatment, PAIR was superior, and in experienced hands PAIR is an effective and safe alternative to surgery.

Usually, PAIR is advocated for uncomplicated univesicular cysts (Gharbi types 1 and 2) although experts also use it for multivesicular so-called “mother and daughter” cysts (Gharbi type 3). Each daughter cyst has to be punctured separately which is labourious and inconvenient for the patient. Therefore, an alternative method was developed, the catheterisation technique, in which multivesicular cysts are irrigated with scolecidals to destroy the daughter cysts and laminated membrane. Saremi described a percutaneous drainage method which is essentially different from PAIR and the catheterisation technique. In this technique, a special cutting instrument is used to fragment and evacuate daughter cysts and laminated membrane while the cavity is continuously irrigated with scolecidals. Both in PAIR and the catheterisation technique, the ruptured daughter cysts and laminated membrane remain inside the cavity. However, using scolecidals, none of these three methods can be safely used for the treatment of cysts with a cystobiliary communication.

We report the first short term results of percutaneous evacuation of multivesicular echinococcal cysts with a cystobiliary communication, using a modified PAIR method. This innovative method has two major advantages. Firstly, it can be used safely in cystobiliary fistulas because scolecidals are avoided. Secondly, it is not necessary to puncture each individual daughter cyst. Cyst content is simply aspirated and evacuated via a large bore catheter. We call this method PEVAC (percutaneous evacuation of cyst content).

METHODS
In all patients, the diagnosis of echinococcal cyst was based on history, physical examination, ultrasound (US), computed tomography; US, ultrasound.

Abbreviations: PAIR, puncture, aspiration, injection, reaspiration; PEVAC, percutaneous evacuation; CB, cystobiliary; CBD, common bile duct; ERCP, endoscopic retrograde cholangiopancreatography; CT, computed tomography; US, ultrasound.
4) or a calcified wall (Gharbi type 5).

characterized by a heterogeneous complex mass (Gharbi type

tion for the time of albendazole treatment; and cysts

or albendazole; pregnancy or women who refused contracep-

coagulation abnormalities; known allergy to local anaesthetics

age <18 years or >75 years; no informed consent; serious

hepatic or portal veins or bile ducts. Exclusion criteria were:

there was a cystobiliary communication or compression of the

non-drainable debris were included (fig 1). Patients with uni-

multivesicular echinococcal cysts (Gharbi type 3) with or without a cystobiliary fistula and containing non-drainable debris were included (fig 1). Patients with uni-

vesicular cysts (Gharbi types 1 and 2) were included only if there was a cystobiliary communication or compression of the hepatic or portal veins or bile ducts. Exclusion criteria were: age <18 years or >75 years; no informed consent; serious coagulation abnormalities; known allergy to local anesthetics or albendazole; pregnancy or women who refused contraception for the time of albendazole treatment; and cysts characterized by a heterogeneous complex mass (Gharbi type 4) or a calcified wall (Gharbi type 5).

The drainage procedure is performed in an interventional radiology suite, appropriately equipped and staffed for emergency situations. The patient has an intravenous canula and is continuously monitored with pulse oximetry. After the procedure, the patient is admitted to the medical ward and checked at hourly intervals until the morning after the procedure.

The optimal route for puncture of the echinococcal cyst is determined by CT and US. After intravenous sedation with 2.5 mg midazolam and 0.1 mg fentanyl, and local anesthesia with 10–15 ml lidocaine 2%, the cyst is punctured under US guidance. Then, to prevent leakage, as much cyst fluid as possible is aspirated to decrease intracystic pressure. A sample of the aspirated cyst fluid is used for parasitological investigation and bacterial culture. Viability is assessed by observing protoscolices motility by direct microscopy and by neutral eosin staining. Subsequently, using the Seldinger technique, a 10–12 French gauge (F) catheter with multiple sideholes is inserted into the cyst cavity after dilating only the skin and proximal intrahepatic part of the puncture tract, again to prevent leakage. The catheter position is confirmed by fluoroscopy. By repetitive manual injection and reaspiration of small amounts (5–10 ml) of isotonic saline, the remaining liquefied part of the mother cyst is removed. The catheter is left in place for drainage. In a second session the catheter is replaced with a 14–18 F stiff Amplatz sheath (William Cook, Bjaeverskov, Denmark). A suction catheter is introduced into the cyst cavity through the sheath and the cyst content is evacuated by directing the catheter towards the daughter cysts, endocyst, and non-drainable material, and applying suction (fig 2). After removal of the sheath a catheter of the same French size as the sheath is placed and contrast is injected into the cyst cavity to detect a possible communication with the biliary tree. In the case of a cystobiliary fistula, an endoprosthesis is introduced endoscopically into the common bile duct (CBD). Sphincterotomy is performed when membrane fragments have to be removed from the CBD or when stenting does not adequately decrease intraductal pressure. Finally, the catheter is removed after complete cyst collapse and closure of a possible cystobiliary fistula, indicated by a daily catheter output of <10 ml.

After the procedure is completed, cyst size is monitored by US at two weeks and 1, 3, 6, 12, 18, and 24 months after PEVAC. If clinically indicated, US is repeated at shorter intervals. CT scans are performed 12 and 24 months after drainage. The primary end points were defined as complete cyst collapse at US at the end of the procedure, disappearance of cyst cavity or at least 50% reduction in cyst size at follow up imaging, and disappearance of complications such as pain, cystobiliary fistulas, vascular or biliary compression, and infection. Direct treatment results were evaluated at the end of the procedure: early results at six months and late results two years after PEVAC. The secondary end points of the study were recurrence of cyst cavity to >50% of its initial size, vascular or biliary compression, fistulas, pain and infection within two years after PEVAC, and death, withdrawal from the study, or loss to follow up. Three examples of cyst appearance before and after PEVAC are shown in figs 3–5.

PEVAC treatment was approved by the Institutional Review Board of the Academic Medical Centre, Amsterdam. Informed consent was obtained from all patients.

Statistics

The Mann-Whitney rank sum test was used to compare results between patient groups and the Wilcoxon signed ranks test for paired observations within patient groups (SPSS for Windows).

PATIENTS AND RESULTS

Patients

Twelve patients with hepatic echinococcosis, mean age 38 years (range 22–61), immigrants from Morocco (five), Turkey (three), Pakistan (one), Syria (one), Afghanistan (one), and Greece (one) were treated for recurrent and severe upper abdominal pain. Ten patients had multivesicular so-called “mother and daughter” cysts (Gharbi type 3) containing non-drainable material. Two patients had univesicular cysts...
(Gharbi type 1) with a ruptured laminated membrane. In one, the right hepatic and portal veins were compressed by the cyst. In the other patient, the univesicular cyst, which did contain one daughter cysts, was suspected of having a cysto-biliary communication.

Complications

Cystobiliary fistulas and infections were the main complications, which occurred only in patients with multivesicular cysts. Less frequently observed complications were significant obstruction of portal and/or hepatic veins in three patients and perforation of a gastric ulcer into the cyst in one patient. Allergic reactions (transient fever, skin rash, eosinophilia) due to leakage of cyst fluid were observed in three patients following changing or removal of the catheter.

Three patients presented with spontaneous intrabiliary rupture. In five other patients the cystobiliary fistula became radiologically apparent on days 8, 17, 20, 25, and 53, respectively, after starting PEVAC. All but one cystobiliary fistula were endoscopically treated by introducing an endoprosthesis into the CBD or by sphincterotomy. In one patient a small cystobiliary fistula closed spontaneously (table 1).

A primary cyst infection was diagnosed in two patients and a secondary cyst infection in seven patients. All patients were successfully treated with antibiotics. In one patient with a primary infection, *S milleri* and anaerobes were cultured from the cyst cavity. In the other patient, culture was negative due to antibiotic treatment. *S morbillorum, S epidermidis*, and *C freundii* were cultured from the cyst cavity in two patients who were readmitted with cyst infection one and five months, respectively, after a PAIR procedure. Remarkably, in both patients bacterial culture of cyst fluid at the end of PAIR was negative.

Contrast injection at endoscopic retrograde cholangiopancreatography (ERCP) caused cholangitis and secondary cyst infection in five patients; three also developed a mild transient pancreatitis. *S milleri, C freundii, C albicans, S sanguis, K pneumoniae, H influenzae*, anaerobes, and enterococci were cultured from their blood, cyst cavity, or bile.

Cyst viability

In the initially sampled cyst fluid of nine patients, viable protoscolices were diagnosed in addition to fragments of the laminated membrane or hooklets. Remarkably, in both patients with multivesicular cysts which became infected following a PAIR procedure, protoscolices were still viable. In two patients with spontaneously infected cysts, viability of daughter cysts could not be assessed because cyst content was too purulent. In one patient with a multivesicular cyst, no protoscolices or hooklets were diagnosed.

Catheter time

In patients with a cystobiliary fistula, catheters were removed 72.3 (28–128) days after PEVAC, 53.6 (7–120) days after endoscopic treatment of the cystobiliary fistulas (mean (range) (table 1). Hospital stay was 38.1 (20–55) days. In patients without a cystobiliary fistula, catheters were removed after 8.8 (3–13) days and hospital stay was 11.5 (8–14) days. Catheter time and hospital stay were significantly longer in patients with a cystobiliary fistula (p=0.007).

Cyst size

Before treatment, cyst size was 12.5 (6–20) cm in patients with and 14.4 (12.7–16) cm in patients without a cystobiliary fistula (p=0.570). At the end of the procedure, when the catheters were removed, all complaints and complications had disappeared. Cavities had collapsed in all but one patient in whom the daughter cysts were partially evacuated. At follow up, 17.3
(4–28) months after treatment, cyst cavities had disappeared in six of eight patients with a cystobiliary fistula and were reduced to 1 and 2.9 cm, respectively, in the other two (p=0.012). In four patients without a fistula, cyst cavity had disappeared in one and cyst size reduced to 2.2 (1–4) cm in the other three, 19.3 (9–30) months after treatment. This 89 (69–100) % reduction in cyst size was not significant for this small number of patients (p=0.068). When all 12 patients were considered as a group, cyst size was significantly smaller 17.9 (4–30) months after treatment than before: 1.6 (0–7) cm versus 13.1 (6–20) cm (p=0.002). In none of the patients had complaints or complications recurred. No patient died, withdrew from the study, or was lost to follow up.

**Serology**

In comparison with cyst size, the response of total IgG echinococcal antibody titres was slow and started to occur

---

**Figure 4** Computed tomography (CT) scan before percutaneous evacuation (PEVAC) [A]. Cystography after partial evacuation of cyst content on day 1 [B] and CT scan on day 2 [C] showing partial cyst collapse. Four months after PEVAC, ultrasound shows remnant cavity near the gall bladder (GBL) and inferior caval vein (VCI) [D].

**Figure 5** Ultrasound before percutaneous evacuation (PEVAC) of multivesicular cyst which had spontaneously ruptured into the biliary tree (left). Computed tomography scan 13 months after PEVAC shows (arrow) partially calcified cyst remnant (right).
and mean hospital stay 19.8–34.6 days. Most infections were secondary to a prior intervention. Two patients were readmitted with a cyst infection at one and five months, respectively, after a PAIR procedure. Despite the infection both cysts were still viable. Contrast injection at ERCP also appeared to be a risk for cyst infection. All five patients who underwent ERCP developed cholangitis and cyst infection. In two spontaneously infected cysts, viability and possible infection of daughter cysts could not be diagnosed accurately. PEVAC was performed to support antibiotic treatment.

Our policy was to remove the catheter only when the cyst cavity had collapsed and catheter output was <10 ml/day. We reasoned that closure of the cystobiliary fistula and collapse of the cyst was achieved at surgical endocyst removal. Unlike surgery, it takes at least two sessions to complete the procedure. PEVAC was performed to support antibiotic treatment.

Three patients presented with a spontaneous intrabiliary rupture and an overt cystobiliary fistula. Of note, in five other patients the cystobiliary fistula became apparent only 1–7 weeks after evacuation of cyst content. Cyst fluid was initially clear but became bile stained in the course of the procedure, in two patients even after about 100 ml were drained. Cystography at the beginning of the procedure could not reliably reveal a cystobiliary communication. The daughter cysts prevented the contrast from reaching the outer limits of the mother cyst. Therefore, care has to be taken with early injection of scleroidal agents into the cavity of multivesicular cysts.

Most infections were secondary to a prior intervention. Two patients were readmitted with a cyst infection at one and five months, respectively, after a PAIR procedure. Despite the infection both cysts were still viable. Contrast injection at ERCP also appeared to be a risk for cyst infection. All five patients who underwent ERCP developed cholangitis and cyst infection. In two spontaneously infected cysts, viability and possible infection of daughter cysts could not be diagnosed accurately. PEVAC was performed to support antibiotic treatment.

Three patients presented with a spontaneous intrabiliary rupture and an overt cystobiliary fistula. Of note, in five other patients the cystobiliary fistula became apparent only 1–7 weeks after evacuation of cyst content. Cyst fluid was initially clear but became bile stained in the course of the procedure, in two patients even after about 100 ml were drained. Cystography at the beginning of the procedure could not reliably reveal a cystobiliary communication. The daughter cysts prevented the contrast from reaching the outer limits of the mother cyst. Therefore, care has to be taken with early injection of scleroidal agents into the cavity of multivesicular cysts.

Most infections were secondary to a prior intervention. Two patients were readmitted with a cyst infection at one and five months, respectively, after a PAIR procedure. Despite the infection both cysts were still viable. Contrast injection at ERCP also appeared to be a risk for cyst infection. All five patients who underwent ERCP developed cholangitis and cyst infection. In two spontaneously infected cysts, viability and possible infection of daughter cysts could not be diagnosed accurately. PEVAC was performed to support antibiotic treatment.

Three patients presented with a spontaneous intrabiliary rupture and an overt cystobiliary fistula. Of note, in five other patients the cystobiliary fistula became apparent only 1–7 weeks after evacuation of cyst content. Cyst fluid was initially clear but became bile stained in the course of the procedure, in two patients even after about 100 ml were drained. Cystography at the beginning of the procedure could not reliably reveal a cystobiliary communication. The daughter cysts prevented the contrast from reaching the outer limits of the mother cyst. Therefore, care has to be taken with early injection of scleroidal agents into the cavity of multivesicular cysts.
the cyst cavity would be closely related. Therefore, catheter time and hospital stay were long. In the case of a cystobiliary fistula, catheter time was 72.3 (28–128) days and hospital stay 38.1 (20–55) days; without a fistula, these times were 8.8 (3–13) days and 11.5 (8–14) days, respectively (p=0.007 and 0.007, respectively). This is comparable with a mean hospital stay reported in surgically treated patients with cystobiliary fistulas of 19.8–34.6 days.11–15 Hospital stay in our patients with cystobiliary fistulas might have been shorter if we had accepted some residual cyst size by removing the catheter earlier. Another option is to discharge the patient and monitor closely in the outpatient department, removing the catheter when daily output is <10 ml.

The final result at follow up, 17.9 (4–30) months after PEVAC, was encouraging. Cyst cavities had disappeared or became significantly smaller (p=0.002). The decrease in echinococcal antibody titles was slow, less remarkable, and a significant increase was observed both with and without a marked allergic reaction.

In summary, PEVAC is a safe and effective method for percutaneous treatment of multivesicular echinococcal cysts with or without cystobiliary fistulas which contain non-drainable content, especially in cases of vascular or biliary obstruction. In univesicular cysts, PEVAC is only indicated in selected cases with a cystobiliary communication, or vascular or biliary obstruction. Following PEVAC, cysts disappeared completely or became >60% smaller. Compression of bile ducts and portal or hepatic veins resolved. Cyst infections and cystobiliary fistulas were the main complications. In patients with a cystobiliary fistula, pretreatment and post-treatment morbidity was high and hospital stay was long but comparable with that of surgery. PEVAC may be improved to reduce morbidity and hospital stay. PEVAC will not replace surgery but may simply create access to a less invasive treatment for more patients. Whether PEVAC reduces the relapse rate will need to be demonstrated in future studies. The observation period of our study was too short to draw any conclusions regarding the recurrence rate following PEVAC.

ACKNOWLEDGEMENT
We thank Professor C Filice (University of Pavia, Italy) for his support; Dr T van Gool (Laboratory of Parasitology, AMC) for parasitological examination of the cyst fluid; and Dr AM Polderman (Laboratory of Parasitology, Leiden University Medical Centre) for the serological tests.

REFERENCES

Authors’ affiliations
H G Schipper, P A Koger, Department of Infectious Diseases, Tropical Medicine, and AIDS, Academic Medical Centre, Meibergdreef 9, 1105 AZ, Amsterdam, the Netherlands
J S Laméris, Department of Radiology, Academic Medical Centre, Meibergdreef 9, 1105 AZ, Amsterdam, the Netherlands
E A Rauws, Department of Gastroenterology, Academic Medical Centre, Meibergdreef 9, 1105 AZ, Amsterdam, the Netherlands

Acknowledgements