pulsatility and peak pressure to increase. In the presence of a valvular mechanism, the communicating pouch would remain at a higher pressure compared with the CSF pressure and so cause spinal cord compression. This led 10 months later to progressive spinal cord compression, although continued filling and enlargement of the pouch resulting in greater pressure on the cord by virtue of its increased surface area.

Histologically, in this case there was absence of the inner arachnoid membrane noted in previous studies and no evidence of arachnoitid or haemosiderin within the cyst making subarachnoid haemorrhage from the cyst traumatic aetiology unlikely. This is by contrast with arachnoid cysts that have been described after trauma, inflammation, or haemorrhage, where the cyst wall consists of a delicate connective tissue with a coating of meningotheil cells. This case demonstrates the rare association of an intradural meningeal cyst with painless thoracic cord compression. It supports previous studies suggesting the congenital nature of these lesions and the possibility of fluctuating cord compression caused by volume changes in the cyst. We found no evidence of a direct association with subarachnoid haemorrhage, although changes in the CSF dynamics after haemorrhage may lead to pressure changes within the cyst. Our histological evidence points to the presence of a single arachnoid layer as being an inconsistent finding in making the diagnosis of a meningeal cyst.

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2 Spiegelman R, Rappaport ZH, Sahar A. Spinal arachnoid cyst with unusual presen-

Bilateral chronic subdural haematoma: an unusual presentation with isolated oculomotor nerve palsy

Isolated third nerve palsy is a common presentation of intracranial aneurysms, diabetes mellitus, chronic lymphocytic meningeval inlammation, and cavernous sinus lesions. Bilateral subdural haematomas presenting with an isolated oculomotor paralysis, however, without any other notable symptoms or signs except for mild headache, are unusual. We report a 60 year old man referred to us with a three week history of mild generalised headache, two weeks of visual blurring, and diplopia for two days. He was known to be hypertensive and on treatment with metoprolol. Relevant medical history included two episodes of transient ischaemic attacks in the form of transient left hemiparesis in 1982 and 1989, for which he was taking warfarin (5 mg per day). The patient was alert and orientated and the only deficit was a complete right oculomotor nerve paralysis. Clinically an aneurysm of the right internal carotid artery was suspected. Surprisingly, CT showed bilateral chronic subdural haematoma (fig. 1). The haematomas were evacuated through bilateral frontal and parietal burr holes. Immediately after the operation the ptosis recovered partially, the pupil reacted sluggishly to light, and six hours later resolution of the third nerve palsy was complete. After the operation he had a transient left hemiparesis that was presumed to be caused by a transient ischaemic attack. Cerebral angiograms performed before discharge did not show any abnormality.

One of us (MMC) previously reported on 114 cases of chronic subdural haematomas and in that series no patient presented with an isolated oculomotor palsy.

One of the most common pathogenic mechanisms of isolated oculomotor palsy is microvascular infarction of the nerve, which may be associated with diabetes mellitus, hypertension, atherosclerosis, or collagen vascular disease. Under these circumstances there is usually partial or complete sparing of the pupil. Our case did not have pupillary sparing. When mydriasis is present, compression of the nerve must be considered, as it is the earliest sign of compression. The cause of the oculomotor paralysis in our case was presumably pressure of the herniating uncus of the right temporal lobe, a false localising sign, common in raised intracranial pressure due to head injuries and intracranial tumours causing brain shift. Chronic subdural haematomas may also present this way, usually with other localising signs, impairment of higher mental functions, or a deteriorating sensorium. The fact that only the right third nerve was paralysed led us to believe that the right side subdural haematoma was larger. In fact, the CT and findings at operation showed that both were of similar size. Perhaps slight anatomical variation in the position of the third nerve in relation to the tentorial edges and unci, and also minor asymmetry of the perimesencephalic cistern explains the lateralisation to the right. Rapid recovery of the third nerve after evacuation of the subdural haematoma lends support to our contention that the palsy was due to distortion of the nerve, and not from another cause.

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Intrathecal baclofen pump infection treated by adjunct intrareservoir antibiotic instillation

The delivery of intrathecal baclofen via subcutaneous pumps is gaining increasing use in the management of intractable spasticity of spinal origin. An uncommon but potentially fatal complication is infection within the pump of the catheter that contacts the pump to the intraspacial space. We report the case of one such infected pump that was successfully sterilised in situ by the combined use of systemic and intrareservoir injection of antibiotics.

A 68 year old man had been receiving intrathecal baclofen via a manually controlled subcutaneous Cordis Secor pump for severe bilateral spasticity and muscle spasms secondary to multiple sclerosis. The pump was operated by carers and medical staff at the nursing home where the patient lived. He was admitted from the nursing home with a severe spastic tetraparesis with power in his left arm only (grade 3/5). He had no meningism and no obvious source of infection.

Microscopy of the urine and three sets of blood cultures were negative. A chest radiograph was normal. Aspiration, microscopy, and culture of the residual baclofen in his reservoir confirmed the presence of a Staphylococcus aureus infection within the reservoir.

His clinical condition was such that immediate removal of the device was not considered mandatory. An attempt was made to sterilise the pump while in situ as the patient was unwilling to undergo surgery to replace the pump if the present one had to be removed. The delivery of intrathecal baclofen was stopped and oral baclofen was started to prevent troublesome spasms in the legs, but this was ineffective. Despite receiving treatment with high dose intravenous fluocinol (1g four times daily) and fucidin (580 mg three times
daily) for two weeks, he remained pyrexial but still without meningism. The microbiology laboratory later confirmed bacterial sensitivity to these antibiotics.

Further aspiration and culture of reservoir washout fluid confirmed the persistence of the Staphylococcus aureus. Analysis of CSF showed the presence of 3 erythrocytes/mm³, 165 leucocytes/mm³ (105 were neutrophils), protein 1·18 g/l, CSF glucose 1·9 mmol/l, (plasma glucose 4·8 mmol/l) and an opening pressure of 11·0 cm of water, consistent with a diagnosis of a partially treated meningitis.

The oral baclofen was discontinued. Ten milligrams (in 10 ml) of intrathecal gentamicin were aseptically instilled into the reservoir, and 2 ml of this (equivalent to the volume of the catheter) were flushed from the reservoir into the catheter. Three hours later, the patient’s course was complicated by a decrease in his level of consciousness, respiratory depression, and extreme flaccidity of all his limbs. A diagnosis of intrathecal baclofen overdose was made. He was transferred to the intensive therapy unit where he required observation but no artificial ventilation. He regained normal consciousness and a spastic tone in his limbs within 48 hours.

After 10 days of simultaneous treatment with intravenous and intrareservoir antibiotics, he was afebrile and his condition had improved to his preadmission clinical state. The systemic antibiotics were successfully withdrawn and the reservoir was aspirated to dryness. Microscopy of this aspiration fluid showed the presence of Gram positive cocci but these now failed to grow in culture, confirming their death.

Undiluted intrathecal baclofen (11 mg in 11 ml) was reintroduced into the reservoir and bolus doses of 100 µg (0·1 ml) were cautiously pumped into the intrathecal space, producing a therapeutic reduction of the patient’s spasticity and spasms without any deleterious effect on his level of consciousness. Initially, he developed a transient (24 hours) low grade pyrexia, but recovered spontaneously and was discharged after one week of further observation. Review at one month revealed no recurrence of sepsis and continuing benefit from the intrathecal administration of baclofen.

Subcutaneous pumps differ from other artificial implants (for example, prosthetic hips, heart valves, and pacemakers) in that direct instillation of antibiotics into the reservoir allows the antibiotic to reach the site of an infection. Thus intrareservoir gentamicin may be used to sterilise the interior of the pump, whereas systemic antibiotics combat the meningitis. Our patient’s meningitis was mild at presentation, but removal of any implant should be considered early in other cases.

This case shows other important aspects in management of subcutaneous intrathecal baclofen pumps. Firstly, strict asepsis during transdermal refilling is crucial to maintain internal sterility; the reservoirs should be refilled as infrequently as possible, with undiluted intrathecal baclofen to reduce the opportunity of introducing contaminants. Some programmable pumps have inbuilt bacterial filters that further reduce this risk.

Secondly, gentamicin was used as it is non-irritant to the meninges and is bactericidal. Flushing of gentamicin along the catheter is justified to expose the whole system to the gentamicin. Flushing should be performed in small aliquots that are unlikely to deliver an overdose of baclofen to the intrathecal space. Excessive flushing of gentamicin may displace baclofen already residual in the catheter into the intrathecal space, leading to respiratory depression and flaccidity. This is one of the disadvantages of the Secor pump. Unlike the Medtronic and the Infusaid pumps, it has no side port to enable the catheter to be aspirated independently. This complication should be anticipated, even if oral baclofen is stopped, and the patient should be observed closely, preferably on a unit where respiratory support is rapidly available.

Thirdly, as there are no leucocytes within the pump reservoir, dead bacteria will not be phagocytosed until they are flushed into the intrathecal space. The bacterial wall endotoxin may have caused the transient pyrexia that we noted. The presence of bacteria on microscopy does not imply their vitality, as they may not grow in culture. Further antibiotic treatment at this stage would be premature unless the pyrexia persists or the patient’s condition deteriorates.

In summary, mild sepsis within Cordis Secor pumps may be treated by gentamicin instillation into the reservoir with cautious flushing of the catheter, in addition to systemically administered antibiotics. If successful, this would allay the need for removal and would enable the patients to continue to benefit from an effective and generally safe treatment.

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