Dopa responsive dystonia

improved and within one month, abnormal neurological signs had resolved, apart from the hand posturing. He is now taking full part in school sports.

PATIENT 2
This 10 year old girl had already been extensively investigated for a progressive gait disorder and diagnosed with hereditary spastic paraparesis. The diagnosis of DRD was considered after a chance remark by her grandmother to the physiotherapist that she had no symptoms in the mornings. Further questioning revealed progressive worsening during the day, requiring assistance to walk home from school. This history had been recorded in the notes several times but its significance never appreciated. The family also reported facial grimacing and abnormal posturing of the hands occurring at the end of the day when she was tired.

Her symptoms had been slowly progressive from the age of 5 years. Investigations at age 9 years included a blood film (excluding acanthocytosis), amino and organic acid chromatography, liver function, ammonia, copper and zinc concentrations, white cell enzymes, computed tomography of the head, and magnetic resonance imaging of the spine; all gave normal results.

On examination at 10 years, she had a spastic gait with toe walking, a mild thoracic scoliosis with a right rib hump, expressionless face, markedly increased tone in her lower limbs with brisk reflexes, ankle clonus, and spontaneously up-going plantars. At the end of the day, walking induced fisting or clasping together of the hands (the latter possibly to increase truncal stability).

She responded in a similar fashion to levodopa (12-5 mg) and carbidopa (3-125 mg), and is now taking part in full normal sporting activities. The scoliosis has resolved.

Discussion
These two reports illustrate some of the difficulties in diagnosis of this treatable syndrome. Most accounts consider torsion dystonia and other extrapyramidal disorders as the main differential diagnosis, but DRD should be considered in any progressive gait disorder, in particular where there is no 'glued to the floor sign' despite significant disability. This sign has been considered pathognomonic of dystonia. Objective clinical signs of organic involvement of the central nervous system make a diagnosis of dystonia untenable but early in the course of DRD these may be 'soft'.

Hereditary spastic paraplegia is a heterogeneous disorder characterised by a particular 'dragging' type of gait, occasionally asymmetrical, which our second patient did not show. An association with choreoathetosis has been reported and the need to consider DRD previously emphasised. Our second patient's extrapyramidal signs were subtle and it was the grandmother's chance remark that led to the diagnosis.

A history of diurnal variation should always be sought, as it is a vital diagnostic clue. Patients with neurological disorders may become easily fatigued but loss of function at the end of the day is probably more significant. Sequential video filming might be useful in objectifying progressive loss of function.


Co-trimoxazole induced aseptic meningitis

Dinesh Pashankar, Maureen McArdle, Anthony Robinson

Duchess of York
Children's Hospital, Manchester
D Pashankar
M McArdle
A Robinson
Correspondence to: Dr Anthony Robinson, Department of Paediatrics, Duchess of York Children's Hospital, Neil Lane, West Didsbury, Manchester M20 2LR
Accepted 18 April 1995

Abstract
A 15 year old boy presented with two episodes of aseptic meningitis-like reactions after ingestion of co-trimoxazole. The diagnosis of co-trimoxazole induced aseptic meningitis was made. This syndrome should be considered in the differential diagnosis of aseptic meningitis.

(Arch Dis Child 1995; 73: 257–258)

Keywords: aseptic meningitis, co-trimoxazole.

Aseptic meningitis is relatively common in childhood and has many causes besides viral infections. In the adult literature there are several case reports of drug induced aseptic meningitis. After an extensive search Maigen et al found 11 cases after ingestion of co-trimoxazole and eight cases after trimethoprim. So far we are aware of only one case report of aseptic meningitis after co-trimoxazole in childhood. This drug is currently widely used in paediatric practice and therefore to increase

Downloaded from http://adc.bmj.com/ on September 7, 2017 - Published by group.bmj.com
awareness of this potential adverse effect we report another case.

Case report
A 15 year old boy with insulin dependent diabetes mellitus was admitted with fever, headache, and vomiting. Earlier on the same day he had received co-trimoxazole (Septrin, Wellcome) for the first time as treatment for an upper respiratory infection. His symptoms worsened over a few hours and he rapidly became unwell. On examination, his temperature was 38-2°C and he had marked photophobia. He had neck stiffness and Kernig’s sign was strongly positive. His fundi were normal. A lumbar puncture resulted in a traumatic tap and intravenous cefotaxime was begun with the presumptive diagnosis of bacterial meningitis. Computed tomography of the head was normal. He began to improve and was almost back to normal within 48 hours. There was no leucocytosis on peripheral blood film and blood cultures were negative. He was discharged after 10 days of antibiotic treatment.

Three weeks later, he was well when reviewed. However, he was complaining of dysuria and was prescribed co-trimoxazole for mild balanitis. Fifteen minutes after ingestion of the first dose (480 mg) he had fever with rigors, severe headache, and vomiting. On admission to hospital he looked unwell and was feverish (temperature 39.1°C). He had photophobia and meningeal signs were strongly positive. Examination of his cerebrospinal fluid showed 122 white blood cells/μl (100% polymorphonuclear leucocytes), protein 1.16 g/l, and a normal glucose concentration. Intravenous antibiotics were started and his symptoms settled within 48 hours.

The following tests on the cerebrospinal fluid were negative: Gram stain; latex agglutination tests for the antigens of Haemophilus influenzae, Neisseria meningitidis, and Streptococcus pneumoniae; mycobacterium polymerase chain reaction; cytospin preparation for malignant cells; cultures for bacteria, viruses, fungi, and tubercle bacilli. Blood tests were as follows: white cell count 4.6×10⁹/l; there was no growth on culture; serum viral and mycoplasma titres not raised; and tests for Coxsackie virus type B IgM and Borrelia burgdorferi IgG were negative. Serum immunoglobulins and complements were normal except for a slightly raised C3 (1.35 g/l, normal range 0.6–1.2 g/l), thought to be due to acute phase reaction. Urine culture was negative and no viruses were isolated from his throat. Chest radiography and repeat computed tomography of the head were normal. Radiography of the skull and spine were also normal with an incidental finding of spina bifida occulta at S1 level.

We diagnosed co-trimoxazole induced aseptic meningitis. Rechallenge was considered unethical and the patient was asked to avoid the drug. He made an uneventful recovery and there has been no recurrence over a one year period.

Discussion
After the second episode of meningitis, an extensive search for infectious agents, immunodeficiency, and possible anatomic defects along the craniospinal axis was negative in our patient. The diagnosis of co-trimoxazole induced aseptic meningitis was based on the close temporal association with ingestion of co-trimoxazole and rapid recovery after withdrawal. Unfortunately we could not obtain cerebrospinal fluid in the first episode of ‘meningitis’ but even normal findings in the cerebrospinal fluid would be compatible with a reaction to co-trimoxazole as a similar neurological illness has been reported after co-trimoxazole with normal findings.3 However, usually the cerebrospinal fluid picture is abnormal and consists of polymorphonuclear leucocytosis, raised protein concentration, and normal glucose concentrations, which is similar to our findings.1 3 Another feature of this condition, also observed by ourselves, is a trend towards increasing severity and rapidity of onset with each successive exposure to the drug.

Aseptic meningitis has been reported in adults after various drugs, for example, ibuprofen, naproxen, co-trimoxazole, trimethoprim, carbamazepine, and azathioprine.1 4 Most cases occurred in young women with connective tissue disorders. The mechanism is unknown, most authors favouring an acute hypersensitivity reaction involving the meninges.1 3 Reports of meningitis after trimethoprim alone suggest that this drug could be the offending agent in co-trimoxazole.3

This condition may be more common than is currently recognised and should be considered in the differential diagnosis of obscure recurrent meningitis or partially treated meningitis in children. A careful drug history may therefore give an important clue to the diagnosis and avoid unnecessary extensive investigations.

Co-trimoxazole induced aseptic meningitis.

D Pashankar, M McArdle and A Robinson

Arch Dis Child 1995 73: 257-258
doi: 10.1136/adc.73.3.257

Updated information and services can be found at:
http://adc.bmj.com/content/73/3/257

Email alerting service

These include:
Receive free email alerts when new articles cite this article. Sign up in the box at the top right corner of the online article.

Notes

To request permissions go to:
http://group.bmj.com/group/rights-licensing/permissions

To order reprints go to:
http://journals.bmj.com/cgi/reprintform

To subscribe to BMJ go to:
http://group.bmj.com/subscribe/