Our patient had no features of phenytoin toxicity despite a concentration of 31.3 mg/l. After starting tolbutamide the phenytoin concentration decreased to 27.4 mg/l. The total concentration was measured, not the free level. Neither the Committee on Safety of Medicines nor the manufacturer knows of any other cases of phenytoin toxicity produced by tolbutamide.

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Convulsion and coma after intranasal desmopressin in cystic fibrosis

Drs E J SIMMONDS, M J MAHONY, and J M LITTLEWOOD (Regional Cystic Fibrosis Unit, St James's Hospital, Leeds LS9 7TF) write: Desmopressin has been advocated as a treatment for nocturnal enuresis.^{1,2} We report on a child with cystic fibrosis who developed water intoxication while being treated with desmopressin. The patient, a girl aged 13 years, had made good progress apart from recurrent nasal polyps requiring operative removal. By the age of 12, however, she had developed chronic *Pseudomonas aeruginosa* respiratory infection and began three monthly admissions for intravenous antibiotics.³ During one of these she started taking intravenous azlocillin (5 g three times a day) and tobramycin (130 mg three times a day). Pancreatic and vitamin supplements were continued. She also had primary nocturnal enuresis resistant to normal therapeutic interventions so we decided to try desmopressin nasal spray. She was given four doses on consecutive nights: 10 µg, 20 µg, 20 µg, 10 µg.

On the morning after her fourth dose of desmopressin the patient developed headache, nausea, and vomiting. She had remained dry for the first time the previous night. Review of her weight chart showed a sharp weight gain since treatment started: 41.35 kg to 43.15 kg. On the third day of treatment serum sodium concentration had been 125 mmol/l and on the fourth day this had fallen to 114 mmol/l. Later that day she had a convulsion lasting about five minutes and subsequently remained comatose. Her desmopressin was discontinued and she was managed with fluid restriction alone to prevent possible pontine myelinolysis.⁴ Twenty eight hours after her last dose of desmopressin she passed urine again and continued to pass good quantities of dilute urine. Forty hours after fluid restriction was started intravenous fluids with her normal daily sodium requirements were started. Her serum sodium concentration rose and five days later was normal (135 mmol/l).

Forty eight hours after her convulsion she was still deeply comatose, however. Physical examination showed response only to deep pain, poor corneal reflexes, absent cough reflex, and absent peripheral reflexes. Computed tomography of her head and lumbar puncture showed nothing abnormal. She gradually regained consciousness and over two weeks returned to normal with no neurological sequelae.

This patient with cystic fibrosis had a severe adverse response to intranasal desmopressin used in doses recommended by the manufacturer; several authors have reported that desmopressin by this route is safe for both short¹ and long term² treatment of nocturnal enuresis. There are no reports of water intoxication, although mild side effects were noted in a few patients treated with intranasal desmopressin for diabetes insipidus.⁵ It is not clear why our patient responded as she did. The duration of action of nasally administered desmopressin is variable,⁷ depending on its rate of clearance from the nose. Our patient's nasal mucosa had the abnormal physical and electrical properties associated with cystic fibrosis as well as nasal polyps, so the duration of action of the drug may have been prolonged by any of these factors. In the light of what is known about the basic defect in cystic fibrosis it is tempting to speculate that this patient's sensitivity to desmopressin, resulting in abnormality of sodium and water homeostasis, is linked to the chloride channel block described.⁸ We would recommend that desmopressin should be used with great caution in children with cystic fibrosis or nasal polyps. At the very least careful weighing and frequent electrolyte measurements should be done.

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Severe hypersensitivity reaction to fenbufen

Dr M M MUTHIAH (Bassetlaw District General Hospital, Worksop S81 0BD) writes: The common side effects of the non-steroidal anti-inflammatory drug fenbufen are mild gastrointestinal disturbance, rash, and neurological symptoms.^{1,2} The potentially serious side effects are peptic ulceration and gastrointestinal haemorrhage. I report here a serious systemic reaction.

A 41 year old previously healthy woman was given fenbufen 900 mg/day for a painful left knee. She took it for 10 days, and four days after stopping the drug she developed a generalised erythematous rash and became unwell with throat pain, fever, and rigors. There was no history of arthritis, drug allergy, asthma, or eczema. On examination she was ill (temperature 40°C) with generalised lymphadenopathy and an extensive erythematous exfoliative rash affecting the whole body and face. The left knee was swollen and tender. Investigations showed: haemoglobin 133 g/l, white cell count 42×10⁹/l, eosinophils 10.1×10⁹/l, neutrophils 14.7×10⁹/l, platelets 139×10⁹/l, prothrombin time 2.4 s, activated partial thromboplastin time 41 s (control 21-32), fibrinogen normal. Aspartate aminotransferase activity was 1015 IU/l (normal 0-40), lactate dehydrogenase 2190 IU/l (230-460), alkaline phosphatase 1233 IU/l (100-280), and albumin 30 g/l. A Paul-Bunnell test, autoantibody screen, and toxoplasma antibody titres were all negative. Erythrocyte sedimentation rate, chest radiograph, radiograph of the left knee, and calcium and phosphate concentrations were normal. Immunoglobulins showed a polyclonal rise. On admission the bone marrow showed a grossly reactive picture with eosinophilia and increased mature lymphocytes.

Four days later she was deeply jaundiced with persistent fever. Repeat blood cultures were negative. Her haemoglobin started to drop and was 5 g/l 10 days after admission. Investigations confirmed a Coombs positive autoimmune haemolytic anaemia. Prednisolone 60 mg/day was started, with considerable improvement in the jaundice and liver function within four days. At this stage a liver biopsy showed swollen liver cells with focal cholestasis. Portal areas contained polymorphs with proliferation of bile ductules. The appearances were those of a resolving hepatitis. Lymph node biopsy showed fatty infiltration with atrophy and no evidence of lymphoma. Liver function values continued to improve after two weeks of steroid therapy, and the haematological and liver function abnormalities completely resolved. She was discharged with advice against using non-steroidal anti-inflammatory drugs. Follow up over a year confirmed that she was well with normal full blood count and liver function values.

Fenbufen was first marketed in the United Kingdom in 1980 and was the most commonly reported suspect drug on yellow cards in 1986 and 1987.³ Eighty per cent of these reports concerned mucocutaneous reactions, including erythema multiforme and Stevens-Johnson syndrome. There have been reports of allergic vasculitis, hepatitis with rash, haemolytic anaemia, and lymphadenopathy. Skin reactions have been reported with other non-steroidal drugs such as ibuprofen.⁴

The onset of illness in relation to drug intake, eosinophilia, and other features such as rash, lymphadenopathy, and a hepatic picture raise the strong possibility of drug induced hypersensitivity in this patient. The rapid resolution of clinical and laboratory abnormalities with prednisolone was consistent with

this. The differential diagnosis is angioimmunoblastic lymphadenopathy, which is known to occur as a hypersensitive reaction to therapeutic agents.

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Azapropazone induced hepatitis

Drs T C N LO and I W DYMCK (Department of Medicine, Stepping Hill Hospital, Stockport) write: Hepatotoxicity is a rare but increasingly recognised side effect of most non-steroidal anti-inflammatory drugs,¹ and several cases of fatal hepatitis have been reported.^{2,3} Nevertheless, complete recovery after early withdrawal of the offending drug usually occurs if the possible link is recognised.⁴ We report a case of azapropazone induced hepatitis in which complete recovery followed withdrawal of the drug.

A 57 year old woman was given azapropazone (Rheumox) 600 mg twice daily for osteoarthritis of both hips. Two weeks later she developed generalised erythema multiforme with slight involvement of the mucus membranes of the lips and mouth. She was admitted a week later with slight jaundice. There was no lymphadenopathy, hepatosplenomegaly, or ascites. She had no history of recent travel, contact with jaundiced patients, blood transfusion, injections, or tattoos. She was not taking other medication and did not abuse alcohol. Laboratory investigations showed: haemoglobin 133 g/l, white cell count 11.3×10⁹/l with 4% eosinophils, plasma total bilirubin concentration 42 µmol/l, aspartate transaminase 500 IU/l, γ glutamyltransferase 22 IU/l, alkaline phosphatase 311 IU/l, total protein 62 g/l (albumin 35 g/l), prothrombin time 17 s (control 12). She was negative for antinuclear factor, antimitochondrial antibodies, and antismooth muscle antibodies and had had no recent infection with herpes simplex virus, Epstein-Barr virus, cytomegalovirus, varicella zoster, or rubella. Tests for hepatitis A and B were also negative.

Azapropazone was stopped and her jaundice and rash resolved within one week of admission. A percutaneous liver biopsy performed 10 days after admission showed mild centrilobular cholestasis, slight fatty change, and scanty inflammatory cells distributed widely throughout the liver parenchyma. The Kupffer cells were prominent and contained periodic acid Schiff positive, diastase resistant material. Several portal tracts showed moderate to severe chronic inflammatory change similar to that seen in chronic persistent hepatitis with occasional multinucleated giant cells and ill defined granuloma. The overall appearances were those of mild non-specific hepatitis with granuloma formation consistent with drug induced hepatitis. She remained well with normal liver function values one month and three months after discharge.

The manufacturers have received only two reports of "suspected" adverse reactions affecting the liver since azapropazone was marketed in 1976. In each case the link could not be substantiated because of lack of definite temporal relations between drug ingestion and the onset of symptoms and both patients were taking several drugs. The two week interval between the start of treatment and the onset of symptoms in our patient together with negative virological and immunological results suggested a drug induced hypersensitivity reaction, though the underlying mechanism is still unknown.⁴ We have reported this case to the Committee on Safety of Medicines.

We thank Dr N L Reeve for his help in reporting the histological findings in this paper.

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