Discussion

Radiosynoviorthosis aims at satisfactory irradiation of the synovium with minimal irradiation of other tissues. Of the yttrium radiocolloids, the silicate is best retained in the knee⁸ and is associated with least chromosomal damage.⁴ Further attempts to reduce extra-articular leakage have included using longer periods of bed rest and encasing the limb in plaster-of-Paris after injection, which considerably increase the nursing work load and the patient's discomfort.

This study has shown that a firm, light splint on a mobilised patient minimises extra-articular spread. The chromosomal damage observed in mobilised patients is comparable with that found in other studies of fully immobilised patients.^{4, 5}

We thank Dr Norman Veall for his continuing encouragement and

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SHORT REPORTS

Air embolism after accidental removal of intravenous catheter

Serious and even fatal complications of central venous catheterisation have often been described but remain little appreciated. We report a case of air embolism that occurred in unusual circumstances.

Case report

A 38-year-old woman with a long history of depression was admitted to our intensive care unit after an overdose of drugs. She was deeply unconscious with an unstable cardiac rhythm and shortly after admission developed ventricular tachycardia leading to cardiac arrest. She was intubated and ventilated with oxygen and was successfully resuscitated after 15 minutes of external cardiac massage and electrical defibrillation. Ventilation was continued and over the next few days her consciousness improved. Initially she had a peripheral venous drip, but this tissued, so an intravenous catheter was substituted to permit measurement of central venous pressure. It was inserted via an introducer cannula into the right internal jugular. Both the catheter and the introducer were stitched firmly to the skin to prevent accidental removal.

Five days after the cardiac arrest we attempted to wean the patient off the ventilator. She was awake and co-operative, but after an hour sitting up and breathing spontaneously with her nasotracheal tube still in place she became confused and began pulling at the nasogastric tube, endotracheal tube, and central venous line. As her colour deteriorated we decided to resume ventilation. In her confusion she suddenly pulled out the venous catheter, leaving the introducer in the vein. Despite being immediately laid head down on her left side she again suffered cardiac arrest, presumably due to massive air embolus. The electrocardiogram showed acute right heart strain. She was again resuscitated but over three months later remained unconscious, breathing spontaneously but with all the other clinical signs of brain death.

Comment

This case demonstrates the dramatic effect of connecting the internal jugular vein to atmosphere via a 13-gauge introducer cannula; this has a bore of 1.6 mm. The catheter had a male Luer hub, which had originally been firmly engaged in the female hub of the introducer. Nevertheless, the patient disconnected them. The complication might have been prevented by using a different type of catheter. It is feasible to use a long (15 cm) cannula in the right internal jugular vein, passing down the right innominate vein into the superior vena cava. Alternatively, a catheter may be introduced over a guide by the Seldinger technique. In either case a stiff catheter with a sharp, tapered end must be used, with the serious risk of perforating the great veins. In our unit we usually use a soft, flexible 20 cm catheter with a rounded end and side holes, which is unlikely to damage veins.¹ This type of catheter is inserted through a plastic cannula which remains in place around the external end of the catheter, providing valuable reinforcement of the vulnerable junction of the catheter with its hub. Correct location of the catheter depends in part on a firm attachment to the hub of the cannula, so that the two will move only as a single unit. We strongly recommend withdrawing this combined unit so that

the tip of the introducer no longer lies within the vein, thus preventing the possibility of air entering if the catheter alone is removed. We also believe that catheters should always be firmly stitched in place and that all devices attached to them, such as drip sets or central venous pressure manometers, should have locking hubs.

Mr H G Wallace, who makes our venous catheters, has produced a modified design incorporating a locking ring on the catheter hub, which is also much easier to stitch down. This should prevent the catheter being removed accidentally without also removing the introducer (see figure).

Designs of catheter and introducer. Top: Present design, with simple male-female junction between catheter and introducer. Bottom: New design with addition of locking ring.

We are grateful to Mr H G Wallace for his prompt response in modifying the design of the catheter and introducer.

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Digoxin dosage in infants

In infants digoxin is a life-saving medication, but the demarcation line between therapeutic and toxic doses is narrow, and the clinical response to the drug varies greatly. Hence it is crucial to assess how the child has responded to the first dose before giving any more.

Regimen

At the Radcliffe Infirmary our practice is to use digoxin injection paediatric BP, 100 mg/l. The initial loading dose advised is 10 µg/kg (0.1 ml/kg) intramuscularly every four hours to a total of four to six doses—that is,

40-60 $\mu\text{g/kg}$ depending on the clinical response. It is important to reassess the child's response before each subsequent dose is given—for example, if the child vomits or the pulse rate falls below 120 beats/min, then the next dose is omitted. If urgent digitalisation is thought necessary the first two doses can be given together. Subsequent maintenance treatment should be 10 $\mu\text{g/kg}$ once or twice daily, or digoxin elixir paediatric BPC 50 mg/l by mouth once or twice daily.

For example, a baby weighing 4500 g will require a dose of 45 μg , or 0.45 ml, of intramuscular digoxin injection paediatric, BP, every four hours for four to six doses, with careful clinical reassessment before each dose. The same dose—that is, 0.9 ml of Lanoxin-PG Elixir—can then be given once or twice daily by mouth as maintenance treatment. A simple nomogram (see figure) acts as an aide mémoire to this regimen, and the use of a Mantoux syringe allows accurate and simple preparation and administration.

Dosage nomogram for the use of digoxin injection paediatric, BP (100 mg/l).

Comment

We have already shown that the initial phase of this regimen results in satisfactory plasma concentrations¹ in both preterm and full-term infants, although the wide variability found underlines the need for repeated clinical evaluation during initial loading. Our subsequent experience with this regimen suggests that it makes both drug prescribing and administration simple.

I acknowledge the help and encouragement given at all stages by the Department of Clinical Pharmacology, University of Oxford.

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Allopurinol hypersensitivity

Allopurinol (4-hydroxypyrazolo(3,4-d)pyrimidine) is the mainstay of treatment for gout. A common side effect is rash or gastrointestinal upset, but xanthine stones,¹ hepatic granulomas,² agranulocytosis,³ reversible leucopenia, and alopecia may occur. Hypersensitivity with vasculitis has been reported, mostly in patients with hyperuricaemia secondary to diuretic treatment—usually thiazides for hypertension—and often with mild renal impairment.⁴ The following case probably occurred in primary gout, though secondary hyperuricaemia cannot be excluded.

Case report

A West Indian aged 34 was diagnosed by his general practitioner as having gout with nephropathy because of acute pain in the first metatarsophalangeal joint, a serum urate concentration of 0.52 mmol/l (8.7 mg/100 ml), and a serum urea concentration of 8.5 mmol/l (51.2 mg/100 ml). Results of urine analysis are not available. Blood count was normal but erythrocyte sedimentation rate (ESR) was 41 mm in the first hour. On 4 August 1978 he began allopurinol 300 mg daily. On 13 September he noticed an intensely pruritic rash in the groins, which progressed over the next week associated with fever, sweats, lethargy, and anorexia. He was admitted on 21 September.

Examination showed an ill man with fever (38.5°C), generalised maculopapular rash, and cervical and axillary lymphadenopathy. Blood pressure was 120/70 mm Hg. Haemoglobin concentration was 13.4 g/dl; white cell count $13.1 \times 10^9/\text{l}$ (13 100/mm³), 15% eosinophils; ESR 42 mm in first hour; serum albumin concentration 29 g/l; serum aspartate aminotransferase activity 2778 IU/l; serum alanine aminotransferase activity 2222 IU/l; serum lactate dehydrogenase activity 6213 IU/l; serum alkaline phosphatase activity 133 IU/l; serum bilirubin concentration 16 $\mu\text{mol/l}$ (0.9 mg/100 ml); serum urea concentration 22 mmol/l (133 mg/100 ml); serum creatinine concentration 280 $\mu\text{mol/l}$ (3.2 mg/100 ml); and serum urate concentration 0.54 mmol/l (9.1 mg/100 ml). Urine contained 60×10^6 red cells and 60×10^6 white cells/l with granular casts; 24-hour collection yielded 3 g protein. Chest radiograph and electrocardiogram were normal. Blood cultures grew no pathogens. Serum IgA concentration was raised (6.5 g/l); IgG and IgM concentrations were normal. Serum C3 concentration was 1.60 g/l (normal 0.70-1.61) and C4 concentration 0.73 g/l (0.15-0.45). Antinuclear factor was not detected. Tests for soluble immune complexes and allopurinol sensitivity by lymphocyte transformation and migration inhibition were negative. Biopsy showed dermal vasculitis, and immunofluorescence staining disclosed granular deposits of IgG and IgM with complement around small dermal blood vessels.

Prednisolone 60 mg daily was instituted on 25 September. The patient improved, but on 13 October, when the dose was reduced to 20 mg daily, rash and fever recurred. He was discharged on 23 October taking 30 mg daily. Blood count and liver enzyme values were normal. ESR was 26 mm in first hour; serum albumin concentration 28 g/l; serum urea concentration 9.0 mmol/l (54.2 mg/100 ml); and serum creatinine concentration 190 $\mu\text{mol/l}$ (2.1 mg/100 ml).

Renal biopsy performed in February 1979 (Dr B H B Robinson) because of persistent renal impairment (creatinine clearance 36 ml/min) demonstrated glomerular fibrosis with extensive tubular atrophy and chronic inflammation of the interstitium consistent with an "allergic" glomerulonephritis.

Comment

The syndrome of allopurinol hypersensitivity consists of maculopapular rash, nephritis, hepatitis, fever, and eosinophilia. Although probably mediated by immune complexes, its precise mechanism is not understood. Allopurinol has a serum half life of 1.25 hours but its active metabolite oxypurinol has a half life of 18-30 hours. Oxypurinol is handled in a similar way to uric acid, being retained in patients taking thiazide diuretics and with renal impairment; thus sensitivity to this metabolite or possibly allopurinol ribonucleotide, whose fate is unknown, may be the cause. Alternatively, a persistent immunological reaction, sometimes necessitating prolonged steroid treatment, may reflect cross-reaction with normal purines, ribonucleotides, or nucleic acids.

In other reported cases symptoms occurred four to six weeks after starting allopurinol 200-400 mg daily, and in one case after only seven days.⁵ Over a third of these patients died from renal failure, infection, and gastrointestinal haemorrhage. Treated promptly most patients recover, although prolonged steroid treatment may be required. The hazards of allopurinol should be considered, particularly before prescribing for secondary hyperuricaemia without clinical gout, as both thiazides and renal impairment appear to be predisposing factors.

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