Fatal digoxin overdose

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Summary
Overdosage with cardiac glycosides is well documented, but not usually fatal. A case is described in which cardiac arrest in asystole was associated with a serum digoxin level of 50.4 µg/l, the highest concentration yet reported after oral administration.

Introduction
In various series of cases of overdosage with cardiac glycosides, the overall mortality has been 13% (Bergy, Fergus and Bruce, 1957), 20% (Gaultier et al., 1968), and 22% (Bismuth et al., 1973). Another series has included measurement of serum digoxin levels (Somogyi, Kaldor and Jankovics, 1972). The highest level previously recorded after oral administration was 42.0 µg/l (Smith and Willerson, 1971).

Case report
A 54-year-old woman was admitted to the Tameside General Hospital about 90 min after taking an unknown quantity of digoxin BP 0.25 mg tablets, and a diuretic (hydrochlorothiazide 50 mg, with amiloride 5 mg). Seven months previously, she had presented with acute pulmonary oedema, which was found to be due to mitral stenosis, and treatment with digoxin 0.25 mg b.d. and the diuretic two tablets daily was commenced. In addition, she had recurrent depression for which she had received ECT and tricyclic antidepressive drugs in the past. At the time of admission, her only other medication was chloral hydrate 15 ml at night.

On admission, she was semi-conscious, pale, sweating and vomiting, with a bradycardia of 26 beats/min and an unrecordable blood pressure. Monitoring showed prolongation of the P-R interval, and an Elecath transthoracic bipolar pacing catheter was inserted into the right ventricle. Ventricular fibrillation was produced, and sinus rhythm restored by DC defibrillation; demand pacing was established at a rate of 80 beats/min but required maximum output (25 mA). Sodium bicarbonate, 200 mmol, was infused. There was some clinical improvement, and the monitor showed atrial fibrillation with a ventricular rate of 130 beats/min.

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and consequent leakage of potassium from the myocardial cells (Glynn, 1964). Attempts to control the hyperkalaemia, including using haemodialysis, may be unsuccessful (Reza et al., 1974; Smith and Willerson, 1971). Little attention was paid to potassium balance in the present patient owing to the rapidity of events, but this probably made little difference to the fatal outcome. Other unfavourable factors in this case include the previous maintenance therapy, which may have saturated the drug-binding sites, and the relative intolerance of diseased myocardium to digoxin.

One of the commonest cardiac arrhythmias is bradycardia with varying degrees of atrioventricular block. Atropine may be tried, but pacing is usually required. In the presence of hyperkalaemia, a pacemaker should be inserted as a prophylactic measure even if the ECG is normal (Bismuth et al., 1973). However, in fatal cases there is often terminal resistance to pacing. In the present case, there was complete cardiac standstill despite direct electrical stimulation of the myocardium. This too seems to be due to failure of the myocardial cell membrane pump (Reza et al., 1974).

The recent description of digoxin-specific antibodies and their use in severe digoxin poisoning may be a significant advance (Smith et al., 1976); in the case they report, hyperkalaemia and pacing resistance were both reversed and the patient survived.

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References


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Leiomyosarcomatosis of probable uterine origin with long survival—a case report

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Summary

A case of leiomyosarcomatosis is presented. Over a period of 15 years the patient underwent seven operations to remove eleven tumours, the largest as big as a football, before dying of widespread metastases: between operations the patient was remarkably well. The disease was almost certainly of uterine origin from apparently benign fibroids, and photomicrographs are provided as supportive evidence. The various modes of presentation of abdominal leiomyosarcoma—mass, pain, obstruction, fistula, anaemia—are illustrated in the case report. The value of repeated palliative surgery in such cases is emphasized.

Case report

At the age of 53, in 1960, the patient presented at

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