This study has shown that amitriptyline and several related tricyclic antidepressants have higher fatal toxicity indices than the monoamine oxidase inhibitors, which are in turn more toxic than several newer antidepressants. All of the drugs introduced since 1973 have a favourable toxicity profile except maprotiline. If the newer drugs have as good a record of clinical effectiveness, combined with their apparent lower potential to cause fatal poisoning when taken in overdose, serious consideration should be given to preferentially prescribing the newer drugs, especially to patients who are considered at particular risk of suicide by ingestion of an overdose of their medication.

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References

1 Brewer C, Farmer R. Self poisoning in 1984: a prediction that didn't come true. *Br Med J* 1985;290:391.
2 Osselson MD, Blackmore RC, King LA, Moffat AC. Poisoning associated deaths between 1973 and 1980. *Hum Toxicol* 1984;3:201-21.
3 Barraclough BM. Are there safer hypnotics than barbiturates? *Lancet* 1974;i:57-8.
4 Girdwood RH. Death after taking medicaments. *Br Med J* 1974;i:501-4.
5 King LA, Moffat AC. A possible index of fatal drug toxicity in humans. *Med Sci Law* 1983;23:193-8.
6 Stead AH, Moffat AC, King LA. Quantitative structure-activity relationships in forensic toxicology. In: Dearden JC, ed. *Quantitative approaches to drug design*. Oxford: Elsevier, 1983:277-8.

7 British Medical Association and the Pharmaceutical Society of Great Britain. *British national formulary*. London: British Medical Association, 1975-84.
8 Association of the British Pharmaceutical Industry. *Data sheet compendium*. London: Association of the British Pharmaceutical Industry, 1975-84.
9 Office of Population Censuses and Surveys. *Mortality statistics—accidents and violence (England and Wales)*. London: HMSO, 1975-84. (Series DH4.)
10 Registrar General. *Annual report: mortality statistics*. Scotland: General Register Office, 1975-84.
11 Blackwell B. Antidepressant drugs. In: Dukes MNG, ed. *Meyler's side effects of drugs*. 10th ed. Oxford: Elsevier, 1984:24-61.
12 Vale JA, Buckley BM, Meredith TJ. Deaths from paracetamol and dextropropoxyphene (Distalgesic) poisoning in England and Wales in 1979. *Hum Toxicol* 1984;3(suppl):135-43S.
13 Callaham M, Kassel D. Epidemiology of fatal tricyclic antidepressant ingestion: implications for management. *Ann Emerg Med* 1985;14:1-9.
14 Crome P, Newman B. Why patients die from tricyclic antidepressant poisoning. *Vet Hum Toxicol* 1979;21(suppl):56-8.
15 Crome P, Newman B. The problem of tricyclic antidepressant poisoning. *Postgrad Med J* 1979;55:528-32.
16 Pentel PR, Benowitz NL. Tricyclic antidepressant poisoning: management of arrhythmias. *Medical Toxicology* 1986;1:101-21.
17 King LA. Fergusson's principle and the prediction of fatal drug levels in blood. *Hum Toxicol* 1985;4:273-8.
18 Wedin GP, Oderda GM, Klein-Schwartz W, Gorman RL. Relative toxicity of cyclic antidepressants. *Ann Emerg Med* 1986;15:797-804.
19 Nickander RC, Emmerson J, Hynes MD, Steinberg MI, Sullivan HR. Pharmacologic and toxicologic effects in animals of dextropropoxyphene and its major metabolite norpropoxyphene: a review. *Hum Toxicol* 1984;3(suppl):13-36S.
20 Zaman S, Lamb JM, Esberger DA, Pearson RM. Inhibition of sperm motility by opiate drugs. *Br J Clin Pharmacol* 1984;18:302P.
21 Yasuhara H, Tonooka M, Kamei K, Sakamoto K. Membrane effects of various drugs on isolated rat hepatocytes and erythrocytes. *Toxicol Appl Pharmacol* 1985;79:453-60.
22 Yasuhara H, Matsuo H, Sakamoto K, Ueda I. Mechanism of membrane stabilising and lytic effects of tricyclic antidepressants. *Jpn J Pharmacol* 1980;30:397-401.
23 Kulig K. Management of poisoning associated with "newer" antidepressant agents. *Ann Emerg Med* 1986;15:1039-44.
24 Wakeling A. Efficacy and side effects of mianserin, a tetracyclic antidepressant. *Postgrad Med J* 1983;59:229-31.
25 Ali CJ, Henry JA. Trazodone overdosage. Experience over five years. *Neuropsychobiology* 1986;15(suppl 1):44-5.
26 Rawls WN. Trazodone. *Drug Intell Clin Pharm* 1982;16:7-13.
27 Cassidy SL, Henry JA. Rapid in vitro techniques for the assessment of the membrane stabilizing activity of drugs. *Food Chem Toxicol* 1986;24:807-9.
28 Leonard BE. Toxicity of antidepressants. *Lancet* 1986;i:1105.

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SHORT REPORTS

Secretor state of patients with insulin dependent or non-insulin-dependent diabetes mellitus

The inability to secrete the water soluble glycoprotein form of the ABO blood group antigens is associated with increased susceptibility to several infections, particularly among children in the vulnerable period between losing maternal antibodies and developing their own active immunity.¹ This is a characteristic that does not alter with age or environmental influences. The secretor (Se) gene is located on chromosome 19; it is not linked to sex or to the histocompatibility locus antigen (HLA) markers. We have also found a considerable increase in the proportion of non-secretors (se) among patients who have ankylosing spondylitis, a rheumatic condition that has a close association (about 95% of cases) with the HLA B27 marker and for which an infectious aetiology has been postulated.² As viral infections have been suggested to contribute to the development of type I diabetes,³ a recent

Patients, methods, and results

During six weeks 205 diabetic patients were examined (102 insulin dependent (56 men and 46 women); 103 non-insulin-dependent (60 men and 43 women)). The patients were recruited from those attending routine clinics; the first five to 12 patients at each session were asked to participate. None of the patients was admitted to hospital. Patients who had type I disease were classified by insulin dependence, their clinical history, and family history of the disease. The patients who had type II disease were not dependent on insulin. The ABO and Lewis blood groups were determined by agglutination and the secretor state by the haemagglutination inhibition method⁴ from saliva. The Lewis group was used to confirm the results obtained from the saliva sample. The results were compared with those found for local blood donors by χ^2 test. There was no significant difference in the distribution of the ABO blood groups between the insulin dependent and non-insulin-dependent diabetic patients ($\chi^2=0.5911$, df=3, $p>0.50$), and the results were similar to those found for local blood donors (type I: $\chi^2=3.702$, df=3, $p>0.10$; type II: $\chi^2=1.266$, df=3, $p>0.50$). Among the patients who had type II disease there was no significant difference in the proportion of non-secretors (25%) compared with the controls. Among the patients who had type I disease the proportion of non-secretors was significantly increased (40%) ($\chi^2=6.223$, $p<0.02$) (table).

Blood group and secretor state of patients with type I and type II diabetes mellitus. All figures are numbers (percentages) of patients

	ABO blood group				Secretor state		Significance
	A	B	O	AB	Secretor (Se)	Non-secretor (se)	
Diabetic patients:							
Type I (n=102)	35 (34)	11 (11)	56 (55)		61 (60)	41 (40)	p<0.02 NS
Type II (n=103)	37 (36)	11 (11)	53 (51)	2 (2)	77 (75)	26 (25)	
Controls (n=334)	104 (31)	42 (13)	173 (52)	15 (4)	245 (73)	89 (27)	

editorial on the genetics of this disorder⁴ prompted us to compare the secretor state of patients who have type I disease with those who have type II disease. If there were a parallel with ankylosing spondylitis we predicted that there would be a higher proportion of non-secretors among the patients who had type I diabetes.

Comment

Among the patients studied there were probably clinical and genetic heterogeneities within the insulin dependent and non-insulin-dependent groups; secretor state, however, is an additional, easily determined genetic

marker that can be used further to investigate differences in the two main aetiological types of diabetes.

The parallel of our results for type I diabetes with those for ankylosing spondylitis is striking. In both there are strong associations with particular HLA markers and a significant increase in the proportion of non-secretors. This is further indirect evidence to support the suggestion that an infectious agent is implicated in the initiation of type I diabetes in some genetically predisposed patients. Determining the secretor state of patients who have various viral infections would be a useful first step in investigating this hypothesis. We suggest that the initial susceptibility to some viral infections might be greater in the non-immune, non-secretor host. The specific immune response to the putative infectious agent, controlled by the class II HLA DR3/4 genes, might contribute to pathogenic sequelae leading to the diabetic state.

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- 1 Blackwell CC, Jonsdottir K, Hanson M, *et al*. Non-secretion of ABO antigens predisposing to infection by *Neisseria meningitidis* and *Streptococcus pneumoniae*. *Lancet* 1986;iii:284-5.
- 2 Shinebaum R, Blackwell CC, Forster PJG, *et al*. Non-secretion of ABO blood group antigens as a host susceptibility factor in the spondyloarthropathies. *Br Med J* 1987;294:208-10.
- 3 Bottazzo GF. Death of a beta cell: homicide or suicide? *Diabetic Medicine* 1986;3:119-30.
- 4 Keen H. The genetics of diabetes: from nightmare to headache. *Br Med J* 1987;294:917-9.
- 5 Mollison PL. *Blood transfusion in clinical medicine*. Oxford: Blackwell Scientific Publications, 1979:414-82.

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Quality of haemofiltration fluids: a potential cause of severe electrolyte imbalance

Haemofiltration, which requires the infusion of large amounts of electrolyte replacement solutions, is finding increasing application in general hospitals. These solutions do not have a product licence, so responsibility for their quality, safety, and efficacy is taken by the prescriber. We report a case in which the wide limits of quality control of a solution had clinical implications and discuss a survey of the limits of quality control of some commonly used haemofiltration fluids.

Patient, method, and results

A 73 year old woman was admitted to hospital because of worsening breathlessness and peripheral oedema. She had a history of two mitral valve replacements nine years and two months previously. The clinical findings were severe tricuspid regurgitation with competent aortic and mitral valves. She progressively gained weight in hospital despite diuretics and infusion of dopamine. We decided to perform further tricuspid valve surgery, and to improve her clinical state haemofiltration was performed for 48 hours before surgery.

She underwent venovenous haemofiltration with a BSM-22 (Hospal UK), with vascular access through a right internal jugular haemofiltration catheter; a Biospal SCU/CAVH filter was used. About 16 litres of haemofiltrate was obtained on each day, with a negative balance of four litres. Fluid replacement through the BSM-22 was with Hemofiltrasol-21, with 8 mmol potassium chloride added to each three litre bag. In addition, she also received two units of human plasma protein fraction for the first 24 hours, 20% mannitol 100 ml intravenously twice and frusemide 80 mg intravenously once in the second 24 hours, and spironolactone 100 mg once daily.

Her clinical state improved. To our surprise her plasma sodium concentration decreased progressively from 138 mmol/l at the start of the procedure to 133 mmol/l after 48 hours of haemofiltration. Forty eight hours after haemofiltration

had stopped her plasma sodium concentration remained at 134 mmol/l. Blood glucose concentration rose from 5.2 mmol/l (random level) before treatment to 8.4 mmol/l during haemofiltration. The plasma potassium concentration, supplemented as described above, remained at 3.8 mmol/l. The serum calcium concentration rose from 2.37 mmol/l to 2.60 mmol/l. There was no appreciable change in the magnesium concentration.

We were surprised by the progressive decrease in plasma sodium concentration during haemofiltration. The Hemofiltrasol-21 bag quoted the sodium concentration as being 140 mmol/l. When a sample of this was analysed on the same flame photometer as the plasma it was found to contain 133 mmol sodium/l. This was confirmed in two other bags of the same batch.

The sodium concentration of 133 mmol/l is within the limits of quality control of Hemofiltrasol-21. We therefore investigated the limits of quality control fluids from four suppliers in this country. The limits varied widely between manufacturers—for example, the limits of quality control for sodium with one manufacturer were +3% to -3%, while with another they were +5% to -7%. Those of magnesium were +5% to -5% with one manufacturer and +13% to -13% with another. The table shows the stated concentration and the limits of some ions.

Concentration (mmol/l) of some common ions in haemofiltration fluids (figures in parentheses are limits of quality control)

Supplier	Sodium	Calcium	Magnesium	Chloride
1	142 (138-146)	2.00 (1.90-2.10)	0.75 (0.71-0.79)	103.0 (98.0-108.0)
2	140 (130-147)	1.60 (1.40-1.80)	0.75 (0.68-0.83)	100.0 (90.0-110.0)
3	135 (128-142)	1.88 (1.79-1.97)	0.75 (0.71-0.79)	106.5 (101.2-111.8)
4	135 (128-142)	1.20 (1.10-1.30)	0.75 (0.65-0.85)	103.0 (98.0-108.0)

Comment

The development of hyponatraemia in heart failure is associated with poor prognosis.¹ The wide limits of quality control applied to some ion concentrations make control of electrolyte balance during haemofiltration unpredictable and potentially hazardous, especially when large volumes are infused. Fluids from suppliers 3 and 4 are within 2% of their intended ion concentrations at the time of manufacture, though they apply wider limits of quality control at the end of the shelf life owing to the evaporation of water from polyvinylchloride bags.

Doctors must remain alert to electrolyte imbalance caused by variations in fluid constituents during large intravenous infusions. Medicinal products without a product licence are prepared as "specials," made to the clinician's specification. Until a manufacturer obtains a product licence doctors who require haemofiltration fluids should specify not only the required concentration of ions but also the acceptable limits of quality control.

- 1 Lee WH, Packer M. Prognostic importance of serum sodium concentration and its modifications by converting enzyme inhibition in patients with severe chronic heart failure. *Circulation* 1986;73:257-67.

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Increased cough reflex associated with angiotensin converting enzyme inhibitor cough

Angiotensin converting enzyme inhibitors are being used increasingly to treat hypertension and congestive cardiac failure as they are one of the few treatments that improve life expectancy in these conditions.¹ In general, these drugs are well tolerated, but recently unexpected and troublesome cough, without obvious pulmonary abnormality, has been reported in as many as 5-10% of patients taking both captopril and enalapril.² Until recently cough has been difficult to assess, but we have developed a method for measuring the sensitivity of the cough reflex to a standard inhaled stimulus: capsaicin (red pepper).³ Using a modification of this method, we assessed the sensitivity of the cough reflex in five patients with cough associated with treatment with angiotensin converting enzyme inhibitors.