The problem of tricyclic antidepressant poisoning

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Summary

In the year May 1976 to April 1977, 489 enquiries about the management of tricyclic antidepressant poisoning received at the London Centre of the National Poisons Information Service were followed-up. One hundred and sixty-four patients (33.5%) were unconscious, convulsions occurred in 62 (12.7%), hypotension in 31 (6.3%), respiratory depression in 28 (5.7%), tachydysrhythmias in 17 (3.5%) and cardiac arrest in 12 patients (2.5%). Sixteen patients died (3.3%). No statistically significant differences were found between individual antidepressants although poisoning with amitriptyline-like drugs resulted in a significantly higher proportion of unconscious patients than poisoning with imipramine-like drugs ($P < 0.01$). There were more asymptomatic children than adults and more unconscious adults than children. Tricyclic antidepressant poisoning is a major clinical problem in general medical and paediatric practice.

Introduction

So-called tricyclic antidepressants were introduced into clinical practice 20 years ago (Kuhn, 1958). Since then the number of cases of overdose from these drugs has increased steadily so that today 10 000 patients (or 10% of all overdoses) are admitted each year to hospital in England and Wales for this reason. The relative frequencies of the various clinical features have been described in a number of series (Goel and Shanks, 1974; Brown, Dwyer and Stocks, 1971; Fournier, 1973; Bismuth *et al.*, 1969; Noble and Matthew, 1969; Thorstrand, 1976; Fréjaville *et al.*, 1965, 1966; Sedal *et al.*, 1972; Serafimovski *et al.*, 1975; Biggs *et al.*, 1977; Noto, Robert and Hanote, 1970; Christensen and Andersen, 1977). Anti-cholinergic effects, pyramidal signs and sinus tachycardia occur commonly but coma, convulsions, respiratory depression, electrocardiographic abnormalities and myocardial depression may all ensue.

There has been only one large study of tricyclic antidepressant poisoning in adults in the United Kingdom, by Noble and Matthew (1969). They reported 100 such consecutive admissions to the

Edinburgh Poisons Treatment Centre. Their conclusions were that most of these patients were not severely affected and that recovery was usually rapid and uneventful, supportive treatment being all that was necessary. In their study no patients required artificial ventilation and there were no deaths. It seemed possible to the present authors that those findings no longer held true, for since then both the number of different antidepressants and the number of admissions to hospital have increased. Support for this view comes in a study from the same unit which reported that the proportion of patients admitted with poisoning from these drugs who were in Grade IV coma had increased from 3.9% in the 5 years 1967-71 to 15.8% in the period 1972-76 (Proudfoot and Park, 1978). It was therefore decided that there was a need for a further survey to assess what proportion of patients were severely affected, whether age had any influence on the incidence of symptoms and whether any one antidepressant was more toxic than another.

Method

This study covered the year May 1976 to April 1977. During this period all enquiries received at the London Centre of the National Poisons Information Service at Guy's Hospital about the management of tricyclic antidepressant poisoning were followed-up, using a specially designed questionnaire. A second and third request for information was made when necessary. Any patient who had taken a tricyclic drug as only a minor part of a mixed drug overdose was excluded. Patients ingesting maprotiline were included in the study for, although described as a tetracyclic antidepressant, it is chemically and pharmacologically related to the tricyclic group (Pinder *et al.*, 1977).

The following information (wherever possible) was recorded on each patient: age and sex, drug and amount ingested, source of antidepressants, symptoms, treatment and outcome. This information was then coded to facilitate computer analysis but excluding any details which might have identified the patient. Differences between groups were compared using the χ^2 test.

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TABLE 1. Age and sex distribution in 485 patients with tricyclic antidepressant poisoning

Sex	Age (years)										Total
	< 5	5-9	10-14	15-19	20-29	30-39	40-49	50-59	60-69	70-79	
Male	49	11	6	10	53	33	12	9	3	2	188
Female	40	1	10	44	66	60	41	18	11	6	297
Total	89	12	16	54	119	93	53	27	14	8	485*

* Excludes 3 patients whose age was not stated and one patient whose sex was not stated.

Results

During the 12 months of the survey there was a total of 1083 enquiries about tricyclic antidepressant poisoning and satisfactory completed questionnaires were received on 489 patients (45.1%). These 489 patients are discussed below.

Age and sex distribution

The age and sex distribution of the patients is shown in Table 1. The 2 decades with most patients were 20-29 years old (24.5%) and 0-10 years old (20.8%). Boys outnumbered girls by 1.3 : 1 whereas in adults (> 15 years old) the position was reversed, with women outnumbering men by 2 : 1.

Drugs ingested

Amitriptyline was by far the commonest drug ingested (203 patients, 41.5%). The others, in order of frequency, were: dothiepin 70 patients (14.3%), imipramine 68 (13.9%), maprotiline 48 patients (9.8%), nortriptyline and trimipramine 27 patients each (5.5%), clomipramine 22 patients (4.5%), doxepin 21 patients (4.3%), dibenzepin 2 patients (0.4%) and protriptyline one patient (0.2%). Two patients took 2 different tricyclic drugs. None of them took butriptyline or desipramine, the other 2 antidepressants of this group available in the United Kingdom. Sixty-nine patients were reported to have ingested more than 1 g of drug; 160 between 250 mg and 1 g, and 104 < 250 mg. In 156 patients (31.9%), however, the amount ingested was not known.

Two hundred and five patients (41.9%) took other drugs in addition to tricyclic antidepressants, benzodiazepines (105 patients) being the most common.

Source of drugs

Table 2 shows the relationship between age and source of tablets. Overall, 303 patients were reported to have taken drugs prescribed for themselves or for a relative. Adults most often took drugs intended for themselves, whilst children under the age of 5 years most frequently took tablets intended for a relative.

Symptoms

Symptoms were reported in 390 patients (79.8%). These were coma 164 patients (33.5%), drowsiness 192 patients (39.3%), delirium 19 patients (3.9%),

TABLE 2. Relationship between age and source of tablets*

Source of tablets	Age (years)			Total
	≤ 5	6-14	≥ 15	
Prescribed for patient	9	10	211	230
Prescribed for relative	52	7	14	73
Neither	19	3	17	39
Not known	12	5	127	144
Total	92	25	369	486

* Excludes 3 patients whose age was not stated.

anti-cholinergic effects 47 patients (9.6%), respiratory depression 28 patients (5.7%), respiratory arrest 6 patients (1.2%), convulsions 62 patients (12.7%), repeated convulsions (3 or more) 39 patients (8%), cardiac complications 141 patients (28%) including sinus tachycardia 96 patients (19.6%), tachydysrhythmias 17 patients (3.5%), bradycardia 9 patients (1.8%), hypotension 31 patients (6.3%), and cardiac arrest 12 patients (2.5%). Convulsions, cardiac complications and either respiratory depression or coma were seen together in 27 patients (5.5%).

The relationship between age and the frequency of symptoms is shown in Table 3. The only major

TABLE 3. Relationship between age and incidence of symptoms

	Age (years)			Total
	≤ 5	6-14	≥ 15	
No symptoms	39	7	53	99
Coma	10	9	144	163
Drowsiness	33	9	149	191
Delirium	3	2	14	19
Anti-cholinergic effects	7	1	37	45
Respiratory depression	2	1	25	28
Respiratory arrest	1	0	5	6
Convulsions				
Present	14	3	45	62
Repeated	9	1	29	39
Cardiac complications				
Present	26	4	111	141
Tachycardia	17	4	75	96
Tachydysrhythmia	4	0	13	17
Bradycardia	2	0	7	9
Hypotension	4	1	26	31
Cardiac arrest	3	0	9	12
Deaths	3	0	13	16
Total no. of patients	92	25	369	486*

* Excludes 3 patients whose age was not stated.

TABLE 4. Relationship between drug ingested and incidence of symptoms*

Drug	No. of patients	Symptoms									Deaths
		Convul- sions	Coma	Cardiac complications					Hypo- tension		
				Present	Tachy- cardia	Tachy- dysrhythmia	Brady- cardia	Cardiac arrest			
Amitriptyline	(203)	16	89	59	39	9	4	5	14	9	
Dothiepin	(70)	12	24	18	17	1	0	0	2	1	
Imipramine	(68)	14	20	22	14	2	2	2	5	2	
Maprotiline	(48)	12	12	11	6	2	2	2	2	1	
Nortriptyline	(27)	6	11	9	5	2	0	1	2	0	
Trimipramine	(27)	0	8	6	5	0	1	0	2	0	
Clomipramine	(22)	0	4	6	4	0	0	1	1	1	
Doxepin	(21)	0	6	7	4	0	0	7	4	1	

* Excludes 2 patients who took dibenzepin and one patient who took protriptyline.

differences were that patients aged more than 15 years had a higher incidence of coma (39%), compared with those under 5 years (10.9%), and that 42.4% of children were asymptomatic compared with only 14.4% of adults.

The relationship between the antidepressant ingested and the frequency of symptoms is shown in Table 4. The only major difference was seen in the incidence of convulsions where 25% of patients ingesting maprotiline developed this complication but none of the patients taking clomipramine, trimipramine or doxepin did so. Overall, however, there were no statistically significant differences between individual antidepressants.

Treatment

Anticonvulsants were used in 52 instances (diazepam 33 patients, phenobarbitone 8 patients, phenytoin 5 patients, paraldehyde 4 patients, chlormethiazole 2 patients); antiarrhythmic agents were used in 10 cases (lignocaine 7 patients, atropine 2 patients, practolol one patient); catecholamines were used in 9 patients (isoprenaline 5 patients, adrenaline 2 patients and dopamine 2 patients); cholinesterase-inhibitors were used in 21 cases (pyridostigmine 10 patients, neostigmine 6 patients, physostigmine 5 patients). Eighteen patients received artificial ventilation. The treatment listed above exclude those used during cardiac arrest procedures. In no less than 357 patients, however, no treatment was recorded as having been given.

Outcome

Sixteen (3.3%) of the patients died. Psychiatric assessment was made in 99 of those who recovered, 34 (7.0%) being admitted to psychiatric units and 21 patients (4.3%) discharging themselves. The drugs involved in the fatal cases were amitriptyline

(9 patients) imipramine (2 patients) and clomipramine, maprotiline, doxepin, dothiepin and dibenzepin (one patient each).

Discussion

All the published series of tricyclic antidepressant poisoning have been in patients admitted either to Poisons Treatment Centres or to Units with special interest in this problem. Comparison with these results is difficult because the authors believe their sample represented a cross-section of patients in terms of age, severity of poisoning, drugs ingested and hospital where treated. The epidemiological findings in this series were no different from the general pattern of poisoning with peak incidences amongst young children and women aged 15–35 years. The high proportion of older children (5–15 years) who took their own tablets was surprising. In recent years, deliberate poisoning has increased in this age group and in one children's hospital the number of admissions for this reason has increased by 7.5 times over the last 10 years (Broadhead, 1978). Clearly, if antidepressants are to be prescribed for depression in childhood then parents must be warned of their dangers and must supervise their administration.

Over 40% of the children in this survey could have ingested only trivial amounts of drug for they were reported to have remained symptom free. The incidence of significant toxicity in childhood poisoning as a whole is low, Broadhead reporting that less than one in 10 of children admitted to hospital for poisoning was in fact suffering from an adverse reaction (Broadhead, 1978). It seems likely that a considerable number of children could have been observed at home and thus spared hospital admission. In the case of tricyclic antidepressants it is probably safe to allow a child home if he remains unaffected

6 hr after an alleged overdose. However, if a slow-release preparation is implicated then a 12-hr observation is desirable.

The authors were unable to show any statistically significant differences between the toxicity of individual antidepressants although overdose of amitriptyline-like drugs resulted in a significantly higher incidence of coma than overdose of imipramine and its analogues ($P < 0.01$). This difference may have been because patients taking amitriptyline-like drugs ingest larger quantities than patients taking imipramine-like drugs, or because amitriptyline and its congeners are intrinsically more toxic. Unfortunately the information on the amounts of drug ingested was insufficiently detailed to allow the authors to conclude which of these explanations was correct. In only one of the published series has it been suggested that there were differences between the individual antidepressants. Among 153 patients reported by Thorstrand (1976), all 4 patients taking dibenzepin died. In the present series only 2 patients took this drug, so that definite conclusions are not possible, although one patient did die. It has also been suggested that doxepin is less cardiotoxic than other drugs because it does not prolong intraventricular conduction (Burrows *et al.*, 1976) although a similar incidence was found of cardiotoxicity with this drug compared with the others.

Park and Proudfoot (1977) have reported a high incidence of coma and convulsions in maprotiline overdosage and in the present series 25% of the patients taking this drug developed these complications. Recently, there have been reports of convulsions occurring during the therapeutic use of this drug (Shepherd and Kerr, 1978; Hall and Russell, 1978). Because of the high incidence of convulsions in their series, the authors decided to extend their survey. Up to July 1978, a total of 75 cases had been documented of which 22 (29.4%) patients were unconscious, 15 (20%) had convulsions, 9 (12%) sinus tachycardia, 4 (5.3%) bradycardia, 3 (4%) cardiac arrest, 2 (2.3%) hypotension and 2 patients (2.3%) died. The proportion of patients exhibiting convulsions is lower than in the original study (25%) but still higher than the overall percentage in the 489 patients (12.7%). Maprotiline is available in 5 tablet strengths ranging from 10 to 150 mg and this made it impossible to determine whether the high incidence of convulsions was due to the patients ingesting excessively large amounts of drug, since it has been suggested that convulsions may result from therapeutic use of high doses of this drug (Burley, Jukes and Steen, 1978; Hall and Russell, 1978; Shepherd and Kerr, 1978). The authors are unable to say whether maprotiline is intrinsically more toxic than the other tricyclic antidepressants but their study of this drug is continuing.

The number of patients reported to have received no treatment was higher than expected (357 patients, 73%). This probably reflects the attitude of doctors who do not regard the supportive care these patients receive as 'treatment'.

In a recent survey of the use of drugs in general practice (Skegg, Doll and Perry, 1977) it was found that in a single year, 11.2% of middle-aged women received antidepressant medication. With this number of prescriptions it is not surprising that overdose is a common problem. The present study shows that it is also a significant clinical problem with a substantial proportion of both adults and children developing serious symptoms such as coma, convulsions, and myocardial depression. It would be prudent for doctors prescribing these drugs to warn patients and relatives of their dangers, to monitor response to treatment at regular intervals and to issue repeat prescriptions only if the drug is still needed and the patient is actually taking it. For someone with suicidal inclinations or with a past history of parasuicide, it may prove wiser to prescribe one of the more recently introduced antidepressants such as mianserin, nomifensine and flupenthixol if further studies confirm the initial reports suggesting that they are less toxic when taken in overdose. (Crome *et al.*, 1978; Montgomery, Crome and Braithwaite, 1978).

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