Plasma concentrations of L-dopa and 3-methoxydopa and improvement in clinical ratings and motor performance in patients with Parkinsonism treated with L-dopa alone or in combination with amantadine

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SYNOPSIS  Six patients with idiopathic Parkinsonism were treated with a combination of amantadine and L-dopa and after 12 to 24 weeks amantadine was replaced by placebo for a six week period in a double-blind trial. Although there was a tendency for clinical disability ratings and scores on objective ratings of motor skills to deteriorate initially after amantadine removal, there was no significant deterioration in clinical improvement or motor performance during the period of amantadine withdrawal. Amantadine withdrawal also failed to cause any significant change in plasma concentrations of L-dopa or its metabolite 3-methoxy-dopa in these patients. In a group of 27 patients seen regularly as outpatients measurements of plasma L-dopa failed to correlate significantly with either oral dose or with clinical improvement scores. The plasma concentration of 3-methoxy-dopa, however, was on average 2.8 times higher than that of L-dopa, and there was a significant correlation between plasma levels of this metabolite and clinical improvement. It is suggested that 3-methoxy-dopa may contribute significantly to the therapeutic actions of L-dopa in Parkinsonism.

Amantadine and L-dopa are in widespread use, either singly or in combination, in the treatment of patients with Parkinsonism. The combination of amantadine and L-dopa has been claimed to be more effective than L-dopa alone, although not all trials have confirmed this synergism (for review see Parkes et al., 1973). Amantadine has also been shown to decrease the daily L-dopa therapeutic dosage requirements and it has been suggested that amantadine may decrease the extracerebral metabolism of L-dopa, thus rendering more drug available to the central nervous system (Peaston et al., 1973). In the present study we present clinical and biochemical findings in a group of patients treated with the combination of L-dopa and amantadine, or with L-dopa alone which do not appear to support the latter conclusions. In addition, we report that only a poor correlation exists between plasma concentrations of L-dopa and clinical improvement in patients with Parkinsonism, although there is a significant correlation with the plasma concentration of the metabolite 3-methoxy-dopa.

METHODS

ISOLATION AND ESTIMATION OF L-DOPA AND 3-METHOXYPYDOPA IN PLASMA SAMPLES  Plasma samples were stored at −20°C for up to two weeks before isolation of the amino acids and fluorimetric assay; preliminary experiments showed that both compounds were stable under these storage conditions. Plasma samples (4 ml) were deproteinized by addition of 4 ml ice-cold 0.4N perchloric acid, and stand-

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(Accepted 16 August 1974.)
ing for at least 15 minutes at ice temperature. The samples were centrifuged and the supernatant fluid adjusted to pH 2.0 by addition of potassium hydroxide. The supernatants were then passed through columns (2.5 cm x 0.6 cm diameter) of the ion exchange resin Amberlite CG-120-Type II-H+ form. The resin was washed with 10 ml distilled water, and both amino acids eluted in 10 ml of a 0.1M sodium phosphate buffer pH 6.5. In experiments in which known amounts of L-dopa or 3-methoxydopa were added to plasma samples the average recovery of L-dopa was found to be 90 ± 2% and of 3-methoxydopa 73 ± 3% (mean ± SE n = 12). All values were corrected for these recoveries.

L-dopa was assayed in 1 ml samples of the eluate by addition of 0.05 ml 0.25% (w/v) potassium ferricyanide, and after three minutes addition of 1.5 ml of alkaline ascorbic acid (9 vol 5N sodium hydroxide: 1 vol 2% ascorbic acid). After mixing and standing at room temperature for 30 minutes samples were read in a spectrofluorimeter with excitation wavelength = 370 nm and recording = 520 nm. Blank samples of eluate were treated in the same way except that sodium hydroxide was added three minutes before the addition of the ascorbic acid; ‘faded’ blanks of this type gave readings not significantly different from those of reagent blanks, not significant from which 1 ml of phosphate buffer was used in place of an eluate sample. The sensitivity of this assay—that is, the amount of amino acid needed to give readings twice as great as those of blank samples—was 16 ng L-dopa. Concentrations in plasma down to 0.05 µg/ml could be accurately measured.

3-Methoxydopa was assayed in parallel 1 ml aliquots of the eluates by a method similar to that described for homovanillic acid by Andén et al. (1963), for 3-methoxydopamine by Guldberg et al. (1971) and for 3-methoxydopa by Fahn et al. (1972). To a 1 ml sample eluate, 1 ml of concentrated (33%) ammonia containing 20 µg potassium ferricyanide/ml; two minutes later 0.1 ml of cysteine solution (1 mg/ml) was added. After five minutes the samples were read in a spectrofluorimeter at excitation wavelength = 315 nm and recording = 430 nm. The sensitivity of this assay was 20 ng 3-methoxydopa, and concentrations down to 0.10 µg/ml in plasma could be measured. The presence of L-dopa did not interfere with this assay procedure, nor did the presence of 3-methoxydopa interfere with the fluorimetric assay of L-dopa.

CLINICAL ASSESSMENTS In a group of 27 patients seen regularly as outpatients an assessment of disability was made on the basis of clinicians' notes of the interview; this gave a five-point disability rating as follows: 0 = normal activities; 1 = slight dis-
3. Alternate tapping  (i) Tapping between two large (5.5 cm) discs placed 50 cm apart on the board. (ii) Similar tapping between two small (3.5 cm) discs 50 cm apart. (iii) Repeated with the large discs 25 cm apart. (iv) Repeated with the small discs 25 cm apart.

For the remaining tests four large (5.5 cm) discs were arranged on the board as follows:

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<th>1</th>
<th>2</th>
<th>3</th>
<th>4</th>
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<tr>
<td>25 cm</td>
<td>25 cm</td>
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4. Bimanual parallel  A second stylus was provided and the patient asked to tap with both hands between pairs of discs 25 cm apart, placed to the left and right side of the board—that is, to alternate simultaneously between 1 and 2 with one hand and between 3 and 4 with the other. Starting positions were 1 and 3.

5. Converge/diverge  The discs remained in the same positions and the patient was asked to use both hands and to tap simultaneously between 2 and 1 and 3 and 4, starting at positions 2 and 3.

6. Patterned tapping  The preferred hand only was used and the patient asked to tap between 1 and 3 until a verbal signal was given (after five seconds), whereupon he was required to change as quickly as possible to tapping between 2 and 4. The trial then continued until 10 seconds had elapsed.

The patients were tested on this battery of tests on each hospital visit immediately after the clinical assessment had been made. Testing continued when the L-dopa dose had reached its clinical maximum and during the double blind experiment with amantadine withdrawal.

STATISTICAL ANALYSIS  Values for dosage of L-dopa (in mg/kg), in absolute dose per day, size of last dose, and time interval from last dose to time of blood sampling, and plasma concentrations of L-dopa and 3-methoxydopa, together with clinical rating scores were assembled in a table and a FORTRAN programme written to calculate the matrix of partial correlation coefficients, using the method of Draper and Smith (1966). These coefficients were tested for significance by reference to a table of the distribution of r (Dixon and Massey, 1969, Table 30). The Kendall rank correlation coefficient (Conover, 1971) was used for analysis of the relationships between these variables and the ordinal data of the disability rating scales. A FORTRAN programme was written to compute and test the significance of the relationships between the disability scale and each of the other variables, including also the number of days since starting L-dopa therapy.

RESULTS

COMBINATION OF L-DOPA AND AMANTADINE  Five of the six patients examined showed a marked improvement in clinical state during treatment with the drug combination, this improvement was seen by a significant reduction in both the ‘disability’ and ‘activities of daily life’ ratings, which tended to continue for the entire period of the trial—up to 40 weeks after commencing treatment (see for example, Fig. 1). The dose of L-dopa was stabilized after eight to 12 weeks and

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<th>TABLE 1</th>
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<td>CLINICAL RATING AND PLASMA L-DOPA AND 3-METHOXYDOPA CONCENTRATION IN PATIENTS TREATED WITH L-DOPA OR COMBINATION OF L-DOPA AND AMANTADINE</td>
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<td>-----------------</td>
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<tr>
<td></td>
</tr>
<tr>
<td>Disability</td>
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<tr>
<td>Pre-therapy</td>
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<tr>
<td>After stabilizing treatment with L-dopa + amantadine</td>
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<tr>
<td>During six week period of amantadine removal</td>
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</table>

All values are means ± SEM for six patients, each value is the average for a six week period (three measurements) before commencing any drug therapy after stabilization on L-dopa + amantadine (12–24 weeks after commencement) and during a six week period in which placebo tablets were substituted for amantadine in a double blind manner. Disability and daily activity ratings are explained in detail in text in the Methods section; improvement in each case is shown by a decreased rating when compared with pre-therapy values. None of the values measured during the period in which amantadine was removed were significantly different from those obtained with the combination of drugs.

was between 1.5 and 3.0 g per day. The maximum dose of L-dopa was limited by the appearance of involuntary movements in two subjects, gastrointestinal upset in two subjects, and good therapeutic response in two. The dose of amantadine was constant at 300 mg per day. During the period of placebo substitution for amantadine, keeping the dose of L-dopa constant, there was a tendency for the clinical disability rating to deteriorate at least initially but this finding was not statistically significant, nor were there any
notable changes in the `activities of daily life' ratings (Table 1). During this period there were also no significant changes in plasma concentrations of L-dopa or of 3-methoxydopa (Table 1, Fig. 1). The concentrations of L-dopa and 3-methoxydopa in plasma were determined one to four hours after the last dose of L-dopa, at 14 day intervals during the period of the study. Plasma concentrations of L-dopa varied considerably between patients and in samples taken at different times from the same patient, although an approximately constant level was reached in most patients soon after the dose of L-dopa was stabilized (Fig. 1). This concentration ranged however, from values of 1 µg/ml or more (as in Fig. 1) to values as low as 0.2–0.3 µg/ml in other patients. There was little or no obvious correlation between clinical improvement and plasma concentration of L-dopa, the one patient who failed to respond to drug treatment had plasma concentrations of amino acid in the normal range.

PLASMA CONCENTRATIONS OF L-DOPA AND 3-METHOXYDOPA IN OUTPATIENT GROUP There was again a wide variation in plasma concentrations of L-dopa and its metabolite 3-methoxydopa between patients, and between samples from the same patient taken at different times. Part of this variation stems from the difficulty of obtaining plasma samples at a fixed time interval after the last dose of L-dopa in an outpatient population. In a small number of hospitalized patients serial blood samples were obtained after a single oral dose of 500 mg L-dopa, and the results from these showed that plasma concentrations of L-dopa reach a maximum within two hours after a dose of the drug and decline in the following two hours (Fig. 2). In the samples from the small number of patients studied in the drug combination experiment, average values of

![Graph showing plasma L-dopa concentration and disability rating score](image)

**FIG. 1** Plasma L-dopa concentration (○), disability rating score (●), and oral dose of L-dopa in one patient during a 33 week period after onset of treatment with L-dopa and amantadine (300 mg per day). Amantadine replaced by placebo for a six week period from week 24.

![Graph showing plasma concentrations of L-dopa](image)

**FIG. 2** Plasma concentrations of L-dopa in serial samples of blood taken after an oral dose of 500 mg L-dopa. Mean values ± SEM for four patients.

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**TABLE 2**

<table>
<thead>
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<th>Parameter</th>
<th>Correlation coefficient with</th>
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<tr>
<td></td>
<td>Clinical rating</td>
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<tr>
<td>L-dopa dose (mg/kg)</td>
<td>—</td>
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<tr>
<td>L-dopa daily dose (g)</td>
<td>—</td>
</tr>
<tr>
<td>L-dopa last dose (mg)</td>
<td>—0.267*</td>
</tr>
<tr>
<td>Time after last dose</td>
<td>0.066</td>
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<tr>
<td>Plasma L-dopa</td>
<td>0.014</td>
</tr>
<tr>
<td>Plasma methoxydopa</td>
<td>—0.325*</td>
</tr>
<tr>
<td>Time after start of therapy</td>
<td>—0.210†</td>
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Correlation coefficients were computed as described in text, using the values obtained from all patients involved in the study.

* P < 0.05. † P < 0.01. ‡ P < 0.001.
Patients with Parkinsonism treated with L-dopa alone or in combination with amantadine

The data acquired during the trial showed a similar decrease in L-dopa concentrations with time after last dose (Fig. 3), although plasma concentrations of 3-methoxydopa did not seem to fluctuate as markedly with time interval after last dose (Fig. 3).

Plasma concentrations of 3-methoxydopa were consistently higher than those of L-dopa. In 27 patients in which the plasma concentrations of the two metabolites were estimated two to three hours after the last dose of L-dopa the concentration of 3-methoxydopa was greater than that of L-dopa in every case, the mean ratio 3-methoxydopa/dopa in these 27 patients being $2.84 \pm 1.59$ (SD). Absolute values for 3-methoxydopa ranged from 0.4-4.0 ml (Table 2).

Statistical analysis of these results showed that there was a highly significant correlation of plasma L-dopa and methoxydopa concentrations with one another (Table 2). Neither concentration, however, correlated significantly with the dosage of L-dopa used. There was a significant correlation between plasma L-dopa concentration and time interval between administration of last dose and blood sampling. Clinical rating scores correlated significantly with time after start of therapy, and surprisingly with plasma methoxydopa but not with plasma L-dopa values (Table 2).

MOTOR PERFORMANCE WITH COMBINATION OF L-DOPA AND AMANTADINE Five of the six patients studied showed a steady improvement on the various motor tests during the weeks that the combination treatment was given and the L-dopa dose increased. The performance in one patient on four of the tasks (J.P. for whom the plasma L-dopa/disability rating correlation is illustrated in Fig. 1), is presented to illustrate the gradual improvement in formal motor acts (Fig. 4).

![Figure 3: Mean values for plasma L-dopa and 3-methoxydopa at various times after last dose for six patients assessed at regular intervals during treatment with 1.5-3.5 g L-dopa per day. Values were collected at various times after last dose during the course of the trial, and are not serial samples as in Fig. 2; means ± SE for six to 18 values.]

![Figure 4: Gradual improvement of a patient (J.P., cf Fig. 1) in motor performance during treatment.]

![Graph 1](http://jnnp.bmj.com/)
DOUBLE-BLIND AMANTADINE WITHDRAWAL. During amantadine withdrawal deterioration in performance was initially noted in some patients but this recovered during the weeks of the trial. When the full set of test results over the six sessions was subjected to a split plot analysis of variance, amantadine withdrawal was shown not to have a significantly detrimental effect when compared with placebo control treatment on the performance level on the formal motor tests (F < 1.00 NS).

DISCUSSION

The present results do not support previous conclusions that the combination of L-dopa and amantadine is more effective than L-dopa alone in the treatment of patients with Parkinson's disease. We also failed to find any significant alterations in plasma concentrations of L-dopa or its metabolite 3-methoxydopa in patients treated with L-dopa alone or in combination with amantadine. It is possible that our sample of patients was not sufficiently large, however, to be truly representative, since there was some tendency for clinical disability rating scores to deteriorate during a period of amantadine removal in patients treated with the drug combination.

In a larger group of patients with Parkinsonism treated with L-dopa we were unable to detect significant correlations between plasma concentrations of L-dopa and clinical improvement. Plasma concentrations of 3-methoxydopa, however, were high relative to those of L-dopa in all patients, and there was a significant correlation between these values and clinical improvement. 3-methoxydopa is known to be an important urinary metabolite of L-dopa in man (for review see Allen, 1973), and the present results confirm that it is also an important circulating metabolite. The presence of a substantial pool of 3-methoxydopa in the circulation and in body tissues may be of neuropharmacological importance in the treatment of Parkinsonism with L-dopa, since it has been suggested that 3-methoxydopa may be metabolized in part by 0-demethylation to give rise to the parent drug (Bartholini et al., 1972).

The lack of correlation between plasma L-dopa concentrations and clinical improvement ratings has been reported by others (see Morris, 1973). This is perhaps related to the possibility that clinical improvement on L-dopa treatment is dependent on the variable degree of degenerative damage to dopaminergic systems in the basal ganglia of patients with Parkinsonism. Jéquier and Dufresne (1972) and Curzon (1973) reported that patients with the most severe impairment of normal dopaminergic function, as assessed by reduced concentrations of the dopamine metabolite homovanillic acid in the cerebrospinal fluid, were more likely to show significant improvement on treatment with L-dopa than those with less severe damage. Thus, because of the variability of the biochemical deficit in different patients, no simple correlation between oral dose or plasma concentration and therapeutic response could be expected.

We are grateful to Dr G. Bartholini, F. Hoffman La Roche and Co, for supplies of 3-0-methyl-dopa, and to Dr F. Geissbuhler for advice on the assay of this substance in plasma. The help of Dr A. W. Galbraith, of Geigy Pharmaceuticals Ltd, is also acknowledged in advising on the structure of the double-blind trial and in providing placebo Amantadine preparation.

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*J Neurol Neurosurg Psychiatry* 1975 38: 129-135
doi: 10.1136/jnnp.38.2.129

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