The stimulant effect of drugs on indocyanine green clearance by the liver

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SUMMARY The rate of removal of a standard dose of 25 mg indocyanine green was studied in 67 patients at the bedside using dichromatic ear densitometry. The determination of the percentage disappearance rate per minute and the half life of the dye permitted separation of patients into three groups: those with normal liver function, those with liver damage, and a group taking opiates (9), anticonvulsants (8), phenylbutazone (2), haloperidol (1), and nitrofurantoin (1). The last group showed enhanced clearance of indocyanine green from the circulation.

The dye indocyanine green has been shown to be an excellent agent for studying hepatic function (Fox, Brooker, Heseltine, Essex, and Wood, 1957; Wheeler, Cranston, and Meltzer, 1958; Rapaport, Ketterer, and Wiegand, 1959; Cherrick, Stein, Leevy, and Davidson, 1960; Hunton, Bollman, and Hoffman, 1960; Caesar, Shaldon, Chiantussi, Guevara, and Sherlock, 1961). It is rapidly bound to plasma albumin after intravenous injection (Cherrick et al., 1960), undergoes no enterohepatic or extrahepatic circulation, and is taken up exclusively by the liver (Wheeler et al., 1958; Rapaport et al., 1959; Cherrick et al., 1960; Hunton et al., 1960). It is secreted almost entirely into the bile but not in conjugated form (Wheeler et al., 1958; Rapaport et al., 1959; Cherrick et al., 1960). Plasma measurements are unaffected by the level of serum bilirubin. This and its lack of toxicity make indocyanine green valuable for serial studies of liver function both in health and disease.

The use of a dichromatic ear piece densitometer (Reed and Wood, 1964) for detecting plasma indocyanine green obviates the need for biochemical estimations (Howard, Senyszyn, and Leevy, 1965; Leevy, Smith, Longueville, Paumgartner, and Howard, 1967) and results obtained for percentage disappearance rate per minute and half life (t/2) show good correlation with those determined by plasma dye measurements (Howard et al., 1965; Leevy et al., 1967; Andersen and Kuchiba, 1970). Since the test is carried out at the patient's bedside, it provides a rapid assessment of liver function. While employing this method in drug addicts we were struck by their rapid removal of indocyanine green, and we have therefore studied the effect of drugs on the test. The results are compared with those obtained in normal subjects and in patients with liver damage.

Patients

Sixty-seven patients (40 male, 27 female) between the ages of 21 and 85 were studied, three of them on two occasions each.

GROUP I

These 11 patients had normal liver function as judged by clinical and biochemical evidence.

GROUP II

Of 38 patients with liver dysfunction, 27 had biochemical and/or histological evidence of hepatitis or cirrhosis. These have been labelled 'liver disease'. The remaining 11 consisted of patients with lymphomas, liver metastases, and one taking an oral contraceptive, all of whom had abnormal liver function tests. This subgroup has been labelled 'liver involvement'. None were on enzyme-inducing drugs (Levi, Sherlock, and Walker, 1968).

GROUP III

These were 12 patients receiving anticonvulsants (8), phenylbutazone (2), haloperidol (1), nitrofurantoin

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Methods

Indocyanine green clearance was measured by a Waters dichromatic ear densitometer which had a compensatory photoelectric cell to correct for variations in blood volume, haematocrit, oxygen saturation, and pCO₂ and a detection photocell to register levels of the dye in the circulation. With the patient recumbent and the ear piece attached to the pinna, indocyanine green was injected into an antecubital vein as a bolus, and the densitometer readings were recorded continuously on a chart recorder.

A standard dose of 25 mg indocyanine green was used in each patient. In three normal subjects repeated clearance tests were carried out using doses of 25 mg and 0·5 mg/kg body weight and the results agreed to within 5% provided the studies were not carried out within an hour of each other. Only the results of the 25 mg clearance tests were included in the overall analysis.

Readings were taken at 20-second intervals from the dye decay curve and the method of least squares was used to establish the regression line between the logarithm of the densitometer reading and time. The half life (t/2) of the dye was calculated, and the percentage disappearance rate (PDR) per minute was obtained from the formula (Leevy et al, 1967):

\[ \text{PDR} = \frac{0.693}{t/2} \times 100 \]

Results

Table I summarizes the mean percentage disappearance rate per minute and half life of indocyanine green in the three groups of subjects studied, and representative clearances plotted on semilogarithmic paper are shown in Figure 1.

<table>
<thead>
<tr>
<th>No.</th>
<th>Group</th>
<th>Percentage Disappearance Rate per Minute</th>
<th>Half Life in Minutes</th>
</tr>
</thead>
<tbody>
<tr>
<td>11</td>
<td>Normal</td>
<td>50·6 ± 1·98</td>
<td>1·3 ± 0·05</td>
</tr>
<tr>
<td>38</td>
<td>Liver dysfunction</td>
<td>24·8 ± 1·76</td>
<td>3·9 ± 0·53</td>
</tr>
<tr>
<td>21</td>
<td>Drugs</td>
<td>115·8 ± 8·04</td>
<td>0·65 ± 0·04</td>
</tr>
</tbody>
</table>

Table I Mean percentage disappearance rate per minute and half life of indocyanine green measured by dichromatic ear densitometry (70 observations in 67 subjects)¹

¹Figures indicate mean ± standard error

The mean percentage disappearance rate was 50·6 per minute in subjects without liver disease, while patients with abnormal liver function had a mean of 24·8 per minute. The mean percentage disappearance rate for the patients who were taking drugs but had normal liver function was 115·8 per minute.

Similarly, the mean t/2 for the three groups was 1·3, 3·9, and 0·65 minutes respectively. There thus appears to be a definite range for each group with some overlap both for percentage disappearance rate (Fig. 2) and t/2, despite the fact that a constant dose of dye was used.

Fig. 1 Examples of indocyanine green clearance in each of the three groups of patients. ICG Conc. = indocyanine green concentrations as recorded by densitometer.
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In general, patients with 'liver disease' had a rate below the mean for group II, while patients with 'liver involvement' tended to have higher rates (Fig. 2). One patient, who had a carcinoma of the pancreas and was included in the 'liver involvement' group, was taking probenecid for gout, a drug which has been shown to decrease plasma indocyanine green clearance in the dog (Vogin, Scott, Boyd, Bear, and Mattis, 1966). Patient S, taking an oral contraceptive, was later re-tested after treatment with phenylbutazone for a superficial thrombophlebitis (see below).

All patients who were either drug addicts or were receiving drugs for therapeutic purposes showed increased plasma indocyanine green clearance (Fig. 2). The patient with the highest percentage disappearance rate (210 per minute) was taking heroin and opium and had a normal liver biopsy, while the two lowest rates (67 and 69 per minute) occurred in the two addicts with liver damage. In spite of this, their plasma indocyanine green clearance was above the range for normal subjects.

Patients on anticonvulsants were receiving either phenobarbitone, a combination of phenobarbitone and phenytoin, or a mixture of phenobarbitone, primidone, and sulthiame. All showed an increased percentage disappearance rate as did the two patients who were receiving haloperidol and nitrofurantoin respectively. Patients C and S (Fig. 2) were treated with phenylbutazone for superficial thrombophlebitis. Indocyanine green clearance was measured before and after two weeks' therapy and a marked increase in percentage disappearance rate per minute was demonstrated. A similar result was obtained in patient J, who received phenobarbitone during treatment for a gastric ulcer.

Discussion

Dichromatic ear densitometry with indocyanine green has been used by many workers (Howard et al, 1965; Leevy et al, 1967; Andersen and Kuchiba, 1970) as a test of liver function, since Fox et al (1957) reported that the dye was removed from the circulation at a constant rate and Reed and Wood (1964) showed that its disappearance could be recorded by a photoelectric cell.

A constant dose (25 mg) of indocyanine green was used in the present study. Several authors, however, have used a dose of 0.5 mg/kg body weight, while Leevy et al (1967) have suggested that larger doses (5 mg/kg body weight) should be used in order to detect mild liver disease. Recently, Andersen and Kuchiba (1970) have shown that reliable and reproducible hepatic decay curves can be produced with doses as low as 0.25 mg/kg body weight. We were unable to find any significant difference in disappearance rate at two different dose levels in three individuals.

In our hands, the results of the test fell into three distinct categories, differentiating normal subjects from patients with liver dysfunction and those taking drugs. Using a constant dose (25 mg), which represented a mean figure of 0.4 mg/kg body weight (range 0.27-0.64 mg/kg body weight; SD ± 0.08), our normal range for percentage disappearance rate was 38-58 per minute, figures which are greater than the 18-25 per minute obtained by other authors (Rapaport et al, 1959; Cherrick et al, 1960; Hunton et al, 1960; Cooke, Harrison, and Skyring, 1963; Leevy et al, 1967). Nevertheless, it was still possible to assign individuals according to the percentage disappearance rate (and t/2) into one of three groups, which corresponded with the clinical, biochemical, and histological findings.

Patients with cirrhosis or hepatitis, as was to be
expected, had a much reduced percentage disappearance rate and increased t/2. Those with slight to moderate liver damage had values which merged with the range found in normal subjects. This agrees with the findings of Hunton et al (1960) and Leevy et al (1967).

Indocyanine green is rapidly removed from the plasma, stored in the liver, and secreted slowly into bile (Wheeler et al, 1958; Rapaport et al, 1959; Cherrick et al, 1960; Hunton et al, 1960). The most striking finding in the present study was the above normal clearance of the dye by patients receiving drugs and addicts, and presumably this could be due to increased uptake by the liver cell, enhanced binding by specific hepatic carrier proteins, or more rapid excretion into bile. Since some of the processes are enzyme-mediated, it would be tempting to postulate that rapid clearance is a reflection of enzyme induction. However, this phenomenon usually refers to drug-metabolizing microsomal enzymes, and, since indocyanine green is not conjugated it seems unlikely that enzyme induction is responsible. Alternative possibilities are increased synthesis of hepatic carrier proteins or increased bile flow, both of which have been demonstrated in animal experiments following drug administration (Reyes, Levi, Gatmaitan, and Arias, 1971; Conklin and Wagner 1971). There is as yet no information that these mechanisms are stimulated in man by drugs and the present findings cannot be explained. There is some evidence from patient S and the two addicts with liver damage that drugs are still capable of enhancing indocyanine green clearance in the diseased liver. If this implies improved hepatic function indocyanine green clearance could be used as a simple and safe procedure for repeated assessment.

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References


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