Marked delay in indocyanine green plasma clearance with a near-normal bromosulphophthalein retention test: a constitutional abnormality?

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SUMMARY Five patients showing a normal to near-normal BSP test and a marked delay in ICG plasma clearance, and two families with clustering of the same abnormality, are described. Two had Gilbert's syndrome, one was convalescing from acute hepatitis, and the other two had no detectable liver abnormality. Measured indices of ICG metabolism indicated a marked reduction in the hepatic uptake, storage capacity, biliary transport maximum, and an increased reflux into plasma. Biochemical studies on the binding of ICG by plasma proteins failed to demonstrate any difference from normal controls. Thus, the primary defect seems to be in the transport of ICG by the hepatocyte. Although the relationship of the defect in ICG metabolism to other constitutional hyperbilirubinaemias is not clear, it is possible that the defect in these patients is a constitutional one involving some steps in the hepatic disposal of organic anions.

Indocyanine green (ICG) has supplanted bromosulphophthalein (BSP) as a liver function test mainly because of the occasional severe reactions to the latter. Unlike BSP, ICG is not conjugated in the liver nor reabsorbed from the gut, and serves as a more direct indicator of the excretory capability of hepatocytes (Cherrick *et al.*, 1960; Caesar *et al.*, 1961). During the past few years, we have encountered five patients, two with mild hyperbilirubinaemia and three without, in whom a striking retention of intravenous ICG was noted in the absence of grossly abnormal BSP test. Study of the families of two propoiti revealed a high incidence of the same abnormality. A number of similar cases with markedly disparate BSP and ICG tests are being seen in Japan, including several familial occurrences and some with Gilbert's syndrome (Nambu and Namihisa, 1975; Namihisa *et al.*, 1975; Ohkubo *et al.*, 1975). The abnormality is always in the clearance of ICG, while the BSP test is normal or near normal, and there is no liver pathology. Thus, this selective abnormality in ICG metabolism seems to reflect some constitutional defect.

Patients

Of the five patients found to have selective ICG abnormality, two (cases 1 and 2) had clinical features of Gilbert's syndrome. Case 3 had a chronic cervical lymphadenitis. Case 4 had an acute, HBsAg negative hepatitis and ICG retention was noted after recovery. Case 5 had a prostate carcinoma, and had been treated with oestrogen (200 mg/day). Biopsy histology was available in all and was unremarkable. The laboratory data are shown in Table 1.

Methods

Serum bilirubin was determined by Michaëlsson's modification (1961) of the Jendrassik method, and the direct bilirubin was measured at 15 minutes. BSP and ICG in serum or plasma were determined spectrophotometrically at 594 nm and 805 nm, respectively. Plasma disappearance curves were drawn after intravenous administration of 0.5 mg/kg ICG or 5 mg/kg BSP under fasting conditions. The kinetics of each dye was studied according to the method of Barber-Riley *et al.* (1961), based on a two compartments model. Data were plotted on a semilogarithmic scale, and 'a', 'b', and 'h' were

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Table 1 Major laboratory tests in five patients with dissociated ICG and BSP tests

Test	Case				
	1 (22 yr, M)	2 (34 yr, M)	3 (25 yr, M)	4 (35 yr, F)	5 (70 yr, M)
Erythrocyte count (10 ⁶ /cu mm)	487	510	440	405	375
Haemoglobin (g/dl)	14.4	16.0	13.4	13.0	11.0
Reticulocytes (%)	0.6	1.0	0.6	0.5	1.1
Serum bilirubin (mg/100 ml)*	2.8	3.4	0.6	0.9	0.5
Direct (15 min direct)*	0.6	1.2	0.1	0.4	0.1
SGOT (IU) (normal, 7-40)	24	26	29	32	34
SGPT (IU) (normal, 5-40)	41	19	30	36	23
Alkaline phosphatase (Bessey U) (0.9-2.9)	1.9	1.0	1.0	2.3	5.9
Albumin (g/100 ml)	3.9	5.2	4.9	5.1	3.7
γ -globulin (g/100 ml)	1.2	1.1	0.7	1.5	1.3
Serum iron (μ g/100 ml)	85	165	105	120	180
Haptoglobin (mg/100 ml)	195	254	264	205	120
Coombs test, direct and indirect	(—)	(—)	(—)	(—)	(—)
Hepatitis B surface antigen	(—)	(—)	(—)	(—)	(—)

*By Michaëlsson's (1961) modification of Jendrassik method.

calculated from the exponential components of the curve. They represent the fractional transfer rates from plasma to liver, liver to plasma, and liver to bile, respectively. Disappearance rate 'k' was

calculated from $k = \frac{0.693}{t_{\frac{1}{2}}}$, where $t_{\frac{1}{2}}$ is half time

of the initial slope. Secretory transport maximum (T_m) and relative storage capacity (S) were determined according to the procedure of Wheeler *et al.* (1960). ICG, 75 mg in 500 ml 5% glucose, was infused at 0.6 mg/min, 0.1 mg/min, and 0.3 mg/min; BSP, 900 mg in 500 ml saline, was infused at 6.0 mg/min, 1.0 mg/min, and 3.0 mg/min.

Sephadex G-200 gel filtration and diethylaminoethyl cellulose (DEAE) chromatography (Sober and Peterson, 1958) were carried out to study binding of ICG by serum proteins. One millilitre of serum was filtered through a 16 $\times$ 700 mm Sephadex column using 0.01 M phosphate buffer of pH 7.4 in 0.5 M NaCl at a speed of 5 ml/h; DEAE chromatography was carried out using a 10 $\times$ 100 mm column and 0.01 M phosphate buffer of pH 7.0, eluted by changing the phosphate buffer to 0.05 M, 0.10 M, 0.15 M, 0.20 M, 0.25 M, and 0.3 M NaCl, and finally to 0.5 M NaOH.

Bilirubin load test was that of Billing *et al.* (1964). From the clearance curve, transfer rates 'a', 'b', and 'm', were calculated; where 'a' is the rate of uptake of bilirubin by the liver, 'b' rate at which unconjugated bilirubin is being returned to plasma, and 'm' rate of transfer of unconjugated bilirubin in the liver to the conjugated bilirubin compartment.

Results

PLASMA DISAPPEARANCE CURVES FOR ICG AND BSP

Attempts were made to determine a, b and h for

ICG and BSP. In case 1, in which k for ICG was extremely low (0.022) and 15 minute retention ($R_{15,ICG}$) 67.3%, plasma ICG declined linearly and calculation of a, b and h was impossible. $R_{45,BSP}$ was 4.9% and only minimally delayed (Fig. 1; Table 2). Very similar results were obtained for ICG in case 2. The k for BSP was 0.041, smaller than normal, and an abnormality in the initial decline was suggested as described by Berk *et al.* (1972). In case 3, k for ICG was even smaller, and a deviation (step) from the initial decline curve (Namiyama *et al.*, 1974) was noted, making the calculation of a, b,

Fig. 1 Plasma disappearance curves for ICG and BSP in case 1. Retention of BSP at 45 min was 4.9%, and of ICG at 15 min, 67.3%. Calculation of transfer rates a, b, and h was not possible because of the linear decay of plasma ICG.

Table 2 Plasma disappearance rate (*K*), retention rate (*R*), and transfer rate constants (*a*, *b*, *h*) for ICG and BSP in five patients

Case	Disappearance rate		Retention rate		Transfer rate constants for					
	<i>K</i> _{ICG}	<i>K</i> _{BSP}	<i>R</i> _{15-ICG}	<i>R</i> _{45-BSP}	ICG (0.25 mg/kg load)			BSP (5.0 mg/kg load)		
					<i>a</i>	<i>b</i>	<i>h</i>	<i>a</i>	<i>b</i>	<i>h</i>
1	0.022		67.3	4.9						
2	0.025	0.041	68.4	4.7	0.003	0.039	0.025			
3	0.014	0.099	88.2	6.7	0.079	0.086	0.015	0.120	0.013	0.033
4	0.014		76.0	4.8						
5	0.016		64.0	5.5						
Normal value	0.180 ± 0.024	0.104 ± 0.026	4.2 ± 1.6	2.9 ± 1.3	0.2080 ± 0.0145	0.0073 ± 0.0012	0.0324 ± 0.0106	0.1350 ± 0.0214	0.0123 ± 0.0045	0.0295 ± 0.0125

Normal values are based on the data in our laboratory. Figures are mean ± standard deviation.

and *h* impossible. The *k* for BSP was 0.099 and nearly normal.

When ICG load was reduced to 0.25 mg/kg, a change in the slope of plasma disappearance curve occurred around 20 minutes. The calculated values of *a*, *b*, and *h* for case 2 were 0.033, 0.039, and 0.025, and for case 3, 0.079, 0.086, and 0.015 respectively (Table 2). In both, *a* was extremely small, *b* greater than normal, and *h* decreased mildly.

The *Tm* and *S* for cases 1-3 (Table 3) indicated marked reduction of both for ICG; reduction of *S* was particularly pronounced in the two cases with unconjugated hyperbilirubinaemia. The *Tm* for BSP was relatively low in cases 1 and 2, but control values varied considerably and the degree of reduction was smaller compared with that for ICG.

BINDING OF ICG BY PLASMA PROTEINS

One millilitre serum containing 0.2 mg ICG was applied on a Sephadex column. Most ICG was bound to protein peaks I and II (Flodin and Kilander, 1962) (Fig. 2), and there was no significant difference between patients and six normal subjects in the relative ICG content in each protein fraction. Stepwise DEAE chromatography was also carried out for the same material. The bulk of ICG was eluted at 0.2-0.3 M NaCl (Fig. 3), and no difference was noted between the patient and control. Sera

obtained at five, 10 and 15 minutes after intravenous injection of 0.5 mg/kg ICG, were then applied on Sephadex columns. Peaks I and II contained more than 70% of the ICG in serum and, although ICG concentration decreased with time, the relative proportion in each peak remained rather constant. No difference was noted in ICG distribution nor in its amount bound per unit protein between the patient and control regardless of time.

BILIRUBIN LOAD TEST

Test was given in three patients. In case 1, *a* was 0.0046, *b* 0.0047, and *m* 0.007, and in case 2, *a* 0.0125, *b* 0.0072, and *m* 0.0052, all being markedly reduced as reported by Billing *et al.* (1964) in Gilbert's syndrome. In case 3, *a* was 0.018, *b* 0.014, and *m* 0.0038, with a mild reduction in *a*.

FAMILY STUDY

We were able to study two families. In the family of case 4, an older sister showed *R*_{15-ICG} of 67.5%, and *R*_{45-BSP} 4.5%; a 26 year old brother showed *R*_{15-ICG} of 85.0% and *R*_{45-BSP} 7.2%. Mother and a younger sister had mildly abnormal ICG tests, and only one sister was normal in the dye study (Fig. 4, above). None had any indication of liver disease. In the family of case 5, three of five of his live children had abnormal ICG retention without abnormal

Table 3 Secretory transport maximum (*Tm*) and relative storage capacity (*S*) for ICG and BSP in three patients

Case	ICG		BSP	
	<i>Tm</i> (mg/min)	<i>S</i> (mg/mg %)	<i>Tm</i> (mg/min)	<i>S</i> (mg/mg %)
1	0.17	3	1.8	36
2	0.01	10	2.0	40
3	0.11	33	4.5	60
Average	0.10 ± 0.07	15 ± 13	2.8 ± 1.2	45 ± 11
Normal control*	0.80 ± 0.11	80 ± 12	8.0 ± 4.2	40 ± 8

*Data in our laboratory.
Figures are mean ± SD.

Fig. 2 Sephadex G-200 gel filtration pattern of patient's serum (case 1) mixed with ICG. The solid line represents protein (absorbance at wave length of 280 nm) and interrupted line, ICG (805 nm).

Fig. 3 DEAE cellulose chromatography pattern of patient's serum (case 2) mixed with ICG. The upper chart represents elution of proteins and the lower chart that of ICG. The bulk of protein bound ICG was eluted at NaCl concentrations of 0.2-0.3 M, and by 0.5 N NaOH.

chemistry (Fig. 4, below). No consanguinity was demonstrated in these families and pedigrees. Presuming that this defective ICG metabolism is transmitted, the mode of inheritance is compatible with being autosomal recessive or incomplete dominant.

INCIDENCE OF SELECTIVE ABNORMALITY OF ICG METABOLISM

Three cases were seen at the Liver Unit, Chiba University Hospital, where ICG test has been used in preference to BSP for the past five years. Since the beginning of 1973, patients exhibiting abnormal

ICG retention have been studied in comparison with BSP. In 1973 and 1974, 1259 patients, including eight with constitutional unconjugated hyperbilirubinaemia, had ICG tests. Approximately one-fifth showed a definite ICG retention because of overt liver involvement, masking any selective abnormality. By subtracting them from the total, the incidence of disparity between marked ICG retention and normal BSP test has been calculated as roughly three out of 1000, and two out of eight with Gilbert's syndrome. These figures are by no means accurate, but do indicate that this abnormality is more frequent in Gilbert's syndrome. There has been no

has been more frequent among patients with constitutional unconjugated hyperbilirubinaemia in our series as well as in the collected cases in Japan. It is possible, but remains to be proven, that Gilbert's syndrome diagnosed with the current criteria encompasses more than one metabolic defect, and that there are patients who share a common defect in ICG handling with those exhibiting abnormality in ICG in the absence of overt jaundice. It has been shown that bilirubin conjugates are excreted into bile through a pathway which differs from that for bile salt (Alpert *et al.*, 1969). There may be many steps in the hepatic transport and biliary excretion of organic anions that have not been elucidated; bilirubin, ICG and BSP share some but not all of them for their excretion. Another line of explanation is that our two patients with Gilbert's syndrome represent a hitherto unrecognized constitutional abnormality characterised by a grossly defective hepatic handling of ICG and bilirubin with near normal BSP metabolism. The argument that these cases represent an extreme variance within the normal quantitative relationship among the three organic anions may be refuted by the established close correlation between BSP and ICG tests, by the failure of Berk *et al.* (1972) to find abnormal ICG test in 26 Gilbert patients, and by the failure of 20 liver units in Japan which have reported this selective ICG defect, to find a single case of selective BSP handling defect with normal ICG test and normal liver histology (Nambu and Namiyama, 1975).

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