Inheritance of type 2 Crigler-Najjar hyperbilirubinemia

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SUMMARY The families of three patients with Crigler-Najjar hyperbilirubinemia, type 2, whose plasma bilirubin levels had responded to phenobarbitone treatment, were investigated. All the parents and several relatives had mildly raised bilirubin levels.

It is suggested that this condition may be an example of genetic heterogeneity and that the propositi had inherited two different abnormal genes. The separation of the Gilbert and type 2 Crigler-Najjar syndromes is at present arbitrary.

Patients with unconjugated hyperbilirubinemia not due to haemolysis or to acquired disease have been divided into three types. Those patients with only mildly raised plasma levels, usually less than 5 mg per 100 ml, are the commonest, and are said to have Gilbert's syndrome. Those with plasma levels greater than 5 mg per 100 ml have been placed in two groups (Arias, Gartner, Cohen, Ben Ezzer, and Levi, 1969). Type 1 patients are rare and were first described by Crigler and Najjar (1952).

Bilirubin levels are usually greater than 20 mg per 100 ml, kernicterus often occurs in infancy, the bile contains no conjugated bilirubin, and there is no detectable activity of bilirubin glucuronoyl-transferase in homogenates of hepatic tissue. Type 2 patients are less rare. Their bilirubin levels are usually less than 20 mg per 100 ml but are sometimes higher, kernicterus is believed not to occur, conjugated bilirubin is present in small amount in bile, and some activity of glucuronoyl-transferase is detectable.

Another difference between type 1 and 2 patients is their response to phenobarbitone, which has no effect in type 1, whereas it reduces the bilirubin levels of patients of type 2, as it also does in Gilbert's syndrome. Arias et al (1969) have suggested that the type 1 disorder is inherited as an autosomal recessive characteristic and type 2 as a dominant, but the frequencies of raised bilirubin levels and of abnormal menthol glucuronidation in the families of the propositi have varied greatly.

We report here studies of the bilirubin levels in the families of two further type 2 patients in addition to a family previously reported (Smith, Middleton, and Williams, 1967). We suggest that these patients do not inherit a dominant defect but abnormal genes from both parents.

Methods

Venous blood samples were taken from non-fasting subjects between 8.0 and 10.0 am. Plasma bilirubin concentrations were determined by a modification of Michaelsson's method (Thompson, 1969), normal levels being less than 0·80 mg per 100 ml for men and 0·65 for women (Thompson, Eddleston, and Williams, 1969a). Standard liver function tests were performed. Free N-acetyl aminophenol and its glucuronide were measured by the method of Brodie and Axelrod (1948) and Schmid and Hammaker (1959). The activity of hepatic bilirubin UDP-glucuronoyl-transferase was determined by the method of Van Roy and Heirwegh (1968), slightly modified (Pilcher, Thompson, and Williams, 1972).

Findings

The families came from different parts of the country, and there was no consanguinity.

D.B.

He was a 21-year-old university student who had been jaundiced since childhood (Smith et al, 1967) with bilirubin levels up to 11·0 mg per 100 ml all
unconjugated. Other liver function tests, including the clearance of bromsulphthalein, were normal, as was the histology of a needle biopsy specimen of the liver.

Phenobarbitone and then dicophane (p,p'-DDT) therapy reduced the hyperbilirubinaemia (Thompson, Stathers, Pilcher, McLean, Robinson, and Williams, 1969b), the bilirubin levels falling to 1.2 mg per 100 ml.

In 1967 his parents, two sibs, maternal aunt, and one of two maternal cousins were found to have mildly raised bilirubin levels of 0.7 to 2.8 mg per 100 ml (Fig. 1).

S.M.

A male, was the product of a normal pregnancy, and delivery at 39 weeks’ gestation. Birth weight was 2.7 kg. He was breast fed, until he was found to have marked unconjugated hyperbilirubinaemia in the first week. There was no hepatic or splenic enlargement, and no kernicterus. Subsequently, at King’s College Hospital, at 6-8 months of age, plasma bilirubin levels of 9.5 to 13.8 mg per 100 ml (mean of seven readings: 10.1 mg) were found, with no direct-reacting pigment; other liver function tests were normal. The histology of a needle biopsy specimen of liver was normal, and no bilirubin glucuronyl-transferase activity was detected in the specimen.

Treatment with phenobarbitone, 1.9 mg kg, failed to lower his bilirubin levels below 6.1 to 8.1 mg per 100 ml (mean of four readings: 6.6 mg), but 3.5 mg kg then maintained it between 1.5 and 5.0 mg per 100 ml for one year. Mental and motor development have been normal, with the body weight between the 75th and 95th percentiles.

Both parents and several relatives have had mildly raised levels (0.8-1.8 mg per 100 ml) of unconjugated plasma bilirubin (Fig. 1).

A female, born after a normal pregnancy, was delivered at 39 weeks’ gestation. Birth weight was 2.5 kg. She was bottle-fed and jaundice was noticed on the third day. At eight days serum bilirubin was 16.5 mg per 100 ml, and 21.5 mg the next day with no direct-reacting pigment. Other liver function tests were normal. The glucuronidation of oral N-acetyl p-aminophenol was markedly impaired compared to a normal infant of the same age. A liver biopsy was not done. Treatment with light therapy and phenobarbitone, starting with 45 mg daily (16 mg per kg), rapidly lowered the bilirubin level, which rose when treatment was stopped. Since then phenobarbitone alone has maintained it between 4 and 8 mg per 100 ml (30 readings) for one year. Development has been normal.

The father had bilirubin levels of 1.7 and 1.5 mg per 100 ml and the mother had one abnormal level of 1.2 mg with three normal values. There are no sibs.

Discussion

Previous studies of the bilirubin levels in relatives of patients with unconjugated hyperbilirubinaemia have been given widely different results. There may be several reasons for this. Thus methods of measuring bilirubin levels close to the normal range are inaccurate. It is significant that among the previous studies, those in which variations of the Jendrassik-Grof method have been used have more frequently found abnormal bilirubin levels in relatives (Alwall, Laurell, and Nilsby, 1946; Powell, Hemingway, Billing, and Sherlock, 1967) than have those studies (Arias, 1962; Sleisenger, Kohn, Barniville, Rubin, Ben Ezzer, and Arias, 1967; Arias et al, 1969) in which the Malloy-Evelyn method (Malloy and Evelyn, 1937) has been used. The latter method is especially unreliable at low bilirubin concentrations (Michaëllsson, Nosslin, and Sjölén, 1966). Many reports do not state the time of day when samples were taken in spite of the large fluctuations of bilirubin levels during the day (With, 1968), and
this may also allow abnormal levels to be missed. The upper limit of normal for bilirubin has frequently been taken to be 1-0 mg per 100 ml or greater, but careful studies have shown a skew distribution of the few values above 0-8 mg per 100 ml in men, and a pronounced sex difference (Powell et al, 1967; Thompson et al, 1969a); again abnormal levels could thus be missed.

Further sources of error are differences in food intake which affect bilirubin levels (With, 1968), and the ingestion of enzyme-inducing drugs, such as sleeping tablets and anticonvulsant drugs, which reduce bilirubin levels (Thompson et al, 1969a). Thus the paternal grandmother of S.M. had a normal bilirubin level but was taking phenylbutazone which is known to induce hepatic microsomal enzymes (Conney, 1967). Bilirubin levels are probably lower in children, so that levels in younger relatives, such as in one of the cousins of D.B., may later become higher. All the parents of the propositi had mildly but definitely abnormal bilirubin levels, the only two siblings of the propositi (both of case D.B.) also had raised levels, and the levels of several other relatives were abnormal. The prevalence of abnormal bilirubin levels in the families of these patients is therefore probably higher than has been reported (Arias et al, 1969).

The inheritance of this condition has also been studied by testing the relatives for the in-vivo glucuronidation of substrates such as menthol, and abnormal results have frequently been found (Szabó and Ėbrey, 1963; Sleisenger et al, 1967; Arias et al, 1969). These may, however, be unreliable, for the glucuronidation of bilirubin could be different from that of other substrates. In the Gunn strain of rat, which, like type 1 patients, has a total bilirubin glucuronyl-transferase deficiency, the glucuronidation of some substrates is selectively impaired; this may be so in these patients and their families.

It is possible that our propositi are homozygous for a common abnormal gene for which all patients with mildly raised bilirubin levels are heterozygous. If so, owing to the high frequency (perhaps 1%) of abnormal bilirubin levels in the population, the prevalence of type 2 patients should be higher than it is. Alternatively, the wide range of bilirubin levels found in type 2 patients may indicate that there is a range of allelomorphs for each of the genes that control the processes of uptake and conjugation of bilirubin. The enzymes synthesized by these genes need only differ in one amino acid, and therefore in stability or specific activity (Harris, 1970). Combinations of mildly abnormal genes would result in the observed range of phenotypes with abnormal bilirubin levels, so our propositi with bilirubin levels much higher than those of their parents would be heterozygous for each of two different, abnormal genes. Twin studies would be helpful (Wetstone and Honeymoon, 1969) in deciding this.

There is a continuous spectrum of abnormality among those patients with non-haemolytic hyper-bilirubinaemia who respond to phenobarbitone. Their separation into Crigler-Najjar type 2 and Gilbert's syndrome is therefore arbitrary, especially since the activity of glucuronyl-transferase is probably greatly reduced in the latter also (Black and Billing, 1969). The position is different in patients with severe hyperbilirubinaemia presumed unresponsive to phenobarbitone who have been reported from the United States (Crigler and Najjar, 1952; Childs and Najjar, 1965; Arias et al, 1969), Canada (Huang, Rozdilsky, Gerrard, Goluboff, and Holman, 1970), and Hungary (Szabó, Kovács, and Ėbrey, 1962). Their parents, for instance, may not have raised bilirubin levels, and from the reports the condition seems to be recessively inherited. The bilirubin level of 21·5 mg per 100 ml of H.P. does, however, suggest that kernicterus could occur in type 2 patients as well as type 1, so that the response to phenobarbitone is at present the best test to differentiate them.

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References
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