Hepatomegaly due to self-induced hyperinsulinism

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SUMMARY Repeated hypoglycaemic attacks, associated with transient hepatomegaly, in a 12-year-old insulin-dependent diabetic girl continued despite reduction in dose and, later, complete discontinuance of insulin. The attacks ceased while she was in hospital, necessitating reinstitution of insulin. The hepatomegaly resolved when surreptitious additional insulin injections were discovered and stopped. Hepatomegaly in diabetics should arouse suspicion of over dosage with insulin.

Hepatomegaly associated with diabetic hypoglycaemia was well known before long-acting insulin became available (Madison, 1969), but it has seldom been encountered recently (Rosenbloom and Giordano, 1977). We present a 12-year-old insulin-dependent diabetic with transient hepatomegaly associated with hypoglycaemia induced by surreptitious additional insulin injections, and discuss the role of over dosage with insulin in inducing hepatomegaly.

Case report

We have followed-up a 12-year-old girl since age 5 when insulin-dependent diabetes was diagnosed and treatment initiated. For 7 years control was good, but then frequent hypoglycaemic attacks of moderate severity were reported. The hypoglycaemic attacks persisted despite reduction in insulin dose and, later, its complete discontinuance. Marked hepatomegaly was noted, and the girl was admitted for investigation. She was then a lean, normotensive, healthy looking child with a firm, tender liver palpable 12 cm below the costal margin. SGOT 138 IU/ml (normal <40), LDH 156 IU/ml (normal <235), alkaline phosphatase 181 IU/ml (normal for age), acid phosphatase 1.5 U/ml (normal 0–1), bromosulphalein retention 2.4% after 45 min, serum ammonia 57 μg/100 ml (normal <85), antitrypsin 88 mg/100 ml (0.88 g/l) α-fetoprotein negative; normal bilirubin, electrolytes, and copper ceruloplasmin. Australian antigen tests, endocrinological studies including evaluation of pituitary and adrenal, and haematological study, including bone marrow aspiration, were all normal. Scan showed diffuse liver enlargement and a normal spleen (Fig. 1a).

After 3 days without treatment, the fasting serum insulin was 5.2 μU/ml. Because of hyperglycaemia (up to 405 mg/dl) and glycosuria (up to 4%), the insulin was restarted. The hypoglycaemic attacks then ceased, the liver size decreased to 6 cm below the costal margin, and she was discharged.

After 5 months of good control in which only the edge of the liver remained palpable (Fig. 1b) the hypoglycaemia and hepatomegaly recurred. Because of the history and laboratory findings, we suspected overinsulinism. The mother then discovered that the girl surreptitiously injected additional insulin, probably to attract notice by provoking hypoglycaemic spells.
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1967; Davidson and Berliner, 1974; Dershewitz et al., 1976). Insulin is known to affect hepatic glucose metabolism directly (Madison, 1969) by increasing the uptake and decreasing the output (Steele et al., 1965). Liver glucose output is largely determined by opposing effects of insulin and glucagon (Jefferson et al., 1968). Glucagon increases the hepatic cyclic-AMP, thus stimulating glycogenolysis and gluconeogenesis (Robison et al., 1971). Insulin opposes this effect by inhibiting the synthesis and increasing the degradation of cyclic-AMP (Jefferson et al., 1968), thus leading to increased glycogen storage.

The excessive insulin taken by our patient undoubtedly disturbed the balance between insulin and glucagon, leading to increased glycogen storage in the liver which presumably was responsible for the hepatomegaly. Stopping the insulin overdose, enabled the glucagon to increase the synthesis of cyclic-AMP, thus inducing glycogenolysis and resolution of the hepatomegaly.

References


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Figure  Liver scan. (a) Diffuse liver enlargement during overinsulinism; (b) Decreased liver size after adjustment of insulin dose.

We insisted that the extra insulin injections be stopped and referred the girl to a psychologist. The hypoglycaemic episodes then disappeared and the hepatomegaly resolved within a few weeks.

Discussion

After excluding diseases associated with hepatomegaly, the diagnosis lay between uncontrolled diabetes and over dosage with insulin. The long periods of good control and disappearance of hypoglycaemia during observation in hospital favoured the latter.

Rosenbloom and Giordano (1977) reviewing overtreatment in diabetic children noted findings similar to those of our patient. Hepatomegaly associated with protracted hypoglycaemia was noted before the availability of long-acting insulin, mainly among poorly controlled juvenile diabetics treated with daily, multiple injections of regular insulin (Mandell and Berenberg, 1974). The hepatomegaly of hyperinsulinism is thought to be secondary to accumulation of glycogen in the liver by inhibition of cyclic-AMP synthesis (Bishop et al., 1965; Mortimore et al.,
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