Diabetes insipidus with impaired osmotic regulation in septo-optic dysplasia and agenesis of the corpus callosum

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Abstract
The clinical and endocrinological findings in 24 children with septo-optic dysplasia and/or agenesis of the corpus callosum are described with particular reference to posterior pituitary function. Nine had diabetes insipidus. The prevalence of diabetes insipidus was similar in children with complete and incomplete forms of septo-optic dysplasia. Maintenance of normal osmotic balance was very difficult in six of these children, even after the introduction of treatment with vasopressin, either as desmopressin, or lysine vasopressin spray in one of the early cases.

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The association between congenital optic nerve hypoplasia and absence of the septum pellucidum was first described by Reeves in a 7 month old child.1 de Morsier confirmed the association in 1956 and suggested the term septo-optic dysplasia which has now been widely adopted.2 Hoyt et al reported the association of optic nerve hypoplasia, abnormal septum pellucidum, and deficiency of both anterior and posterior pituitary function.3 It has become apparent that the three individual defects are variably related to each other in the syndrome of septo-optic dysplasia represent one extreme of a wider spectrum of abnormalities of holoprosencephaly with single cerebral ventricle and absence of the corpus callosum, among other midline defects.

While anterior pituitary deficiency in septo-optic dysplasia has been well documented,4–7 diabetes insipidus has been reported less often and there is little information on the control of osmotic equilibrium in such patients. As our own observations have suggested that disordered osmotic homeostasis may be quite common in children with septo-optic dysplasia, we have carried out a retrospective study on 24 children seen in the endocrine clinics at this hospital between 1971 and 1992.

Subjects and methods

Subjects
The casenotes of 24 children with septo-optic dysplasia and/or agenesis of the corpus callosum were available for review. Eight children were considered to have the complete form of the syndrome (optic nerve hypoplasia, abnormal septum pellucidum, and pituitary deficiency) and 13 to have incomplete forms (two elements of the three). One child with complete septo-optic dysplasia and one with incomplete septo-optic dysplasia also had agenesis of the corpus callosum and three further children had isolated agenesis of the corpus callosum with pituitary deficiency. One child with septo-optic dysplasia was also a mosaic for Turner's syndrome (85% XO; 15% XX). The age at diagnosis ranged from 2 weeks to 8-8 years (mean 2-0 years) and the age at last follow up ranged from 1 to 17 years. Seven children were completely blind and eight had severe visual impairment. Thirteen showed either moderate or severe developmental delay.

METHODS

Pituitary function was assessed using standard methods.8 Insulin induced hypoglycaemia or glucagon tests were carried out to evaluate growth hormone and adrenocorticotropic hormone (ACTH) secretion in all cases. Estimation of plasma thyroxine and thyroid stimulating hormone (TSH) in all children (with thyroid releasing hormone stimulation in 15 cases) was used to assess pituitary TSH secretion; gonadotrophin secretion was assessed after injection of gonadotrophin releasing hormone (GnRH) in 14 cases. In 12 cases standard water deprivation tests were carried out.8

In all cases cranial computed tomography was performed to assess the midline structural abnormality.

Results

Overall, growth hormone insufficiency was present in 20 of the 24 cases, and 14 of these children have been treated with human pituitary or recombinant growth hormone. ACTH deficiency was documented in 15 cases. Gonadotrophin deficiency was demonstrated in 10 of the 14 cases tested with GnRH.

Nine children had diabetes insipidus. Seven of these children were completely blind or had severely restricted vision, seven had moderate or severe mental retardation, and four had one or more episodes of hypoglycaemia in early infancy (table). Details of the computed tomography findings, osmolar impairment, and other endocrine deficits are also given in the table. Three children presented with failure to thrive and hypernatraemia within two months of birth, and four more with hypernatraemia before the age of 1 year. In one child (case 8) there was no clinical evidence of...
While we are aware of 15 cases of diabetes insipidus reported in children with septo-optic dysplasia or agenesis of the corpus callosum,4 6 7 10-15 there is little information on either the prevalence of diabetes insipidus or on the maintenance of normal osmotic equilibrium in such patients. In the present study we found that the overall prevalence of diabetes insipidus was 38% and that disturbance of fluid balance was a significant problem in several of the cases. There was little correlation between the clinical features and the structural central nervous system lesions; 78% of the cases with diabetes insipidus had incomplete forms of septo-optic dysplasia, as compared with 60% of those without diabetes insipidus.

The management of diabetes insipidus was difficult in several cases and five required two or more admissions to hospital in an attempt to stabilise their daily fluid balance with appropriate doses of desmopressin or lysine vasopressin. A number of factors probably contributed to this difficulty. First, several children appeared to have impairment of their sense of thirst. This, together with their dependence on their parents for water and food as a result of their blindness and developmental delay, probably accounts for much of the difficulty.

It is not clear whether the mental retardation commonly seen in septo-optic dysplasia is a consequence of abnormal brain development or whether it is due to other factors, for example, hypoglycaemia or repeated episodes of electrolyte imbalance that may contribute to further neurological damage. In particular, rapid correction of hypopituitarism may be associated with central pontine myelinolysis that can also involve the basal nuclei and central white matter causing quadriplegia, mental retardation, and sometimes cranial nerve dysfunction.16 Psychosocial deprivation, which in our experience is quite common in such children, may also play a part in the developmental delay, and possibly in the abnormal growth hormone secretion.17

In conclusion, in our experience diabetes insipidus is a relatively common finding in patients with the complete and incomplete forms of septo-optic dysplasia and management of fluid balance can be difficult in such patients, particularly if they have an impaired sense of thirst.1
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