Short report

Basal cell naevus syndrome and intracranial meningioma

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SUMMARY Two cases of the basal cell naevus syndrome in association with an intracranial meningioma are described and the relationship discussed.

The basal cell naevus syndrome is a condition with autosomal dominant inheritance of high penetrance and variable expressivity.¹ It is characterised by multiple basal cell naevi, jaw cysts, cutaneous pits of the hands and feet, vertebral and rib abnormalities, and intracranial calcification.² The basal cell naevi usually appear in childhood, particularly over the face, neck, trunk and upper limbs. There may be few to hundreds varying in size and appearance from small papules to larger nodules, sometimes with pigmentation. Clinically the tumours resemble pigmented moles (naevi). There may be a gross resemblance to neurofibromatosis. However, microscopically, the tumours cannot be differentiated from ordinary basal cell carcinomas. In addition, central nervous system abnormalities have been described since the first case report, in which agenesis of the corpus callosum was found.³ We now describe two cases of basal cell naevus syndrome associated with an intracranial meningioma.

Case 1

A girl aged 14 years developed cysts in her jaw requiring numerous operations for their removal. Two years later she had a basal cell carcinoma removed from her nose. Subsequently, she had five more identical lesions excised from different sites on her face. When aged 47, she developed simple partial seizures affecting the left upper limb with secondary generalisation. Examination revealed ocular hypertelorism, frontal bossing, palmar pits and multiple basal cell naevi on the face (fig 1). In addition, she had

a mild left hemiparesis. Skull radiographs demonstrated heavy calcification of the falx cerebri, flecks of calcium in the parasagittal region and an osteolytic lesion in the right fronto-parietal region. Right carotid angiography confirmed the presence of a right fronto-parietal space-occupying lesion with no pathological circulation. At craniotomy, a tumour 6 cm in diameter was excised which histology proved to be a meningioma. Her son also has basal cell naevus syndrome and has had multiple basal cell carcinoma lesions excised.

Case 2

This patient first developed a basal cell carcinoma on the left temple at the age of 40 years. The tumour was excised and the site treated with superficial radiotherapy. Two years later, he presented with four similar lesions close together which were treated by a radium implant. At the age of 44 he was given a further course of superficial radiotherapy to a basal cell carcinoma of the left inner canthus. Later that year he gradually developed difficulty in concentration, slurred speech and weakness and numbness of the right hand. Examination revealed post-irradiation changes in the skin of the temple with adjacent hair loss, mild aphasia and dysarthria, and weakness and numbness of the right hand. Skull radiographs showed calcification of the falx cerebri. Left carotid angiogram demonstrated a left parietal space occupying lesion with some vascular supply from the external carotid artery. At craniotomy, a tumour 6.5 cm in diameter was removed. Histology showed the tumour to be a meningioma of transitional type with marked cellularity and numerous mitotic figures suggestive of malignancy. At the age of 45 years he had developed several further basal cell carcinoma lesions. It was noted that he had palmar pits, frontal bossing and the diagnosis of basal cell naevus syndrome was made. Radiographs of the chest and cervico-thoracic spine demonstrated multiple rib anomalies and extensive spina bifida from the second to the fourth thoracic vertebrae. To

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Fig 1 Case 1 showing three naevoid basal cell carcinomas.

date, at least 25 basal cell carcinomas have been treated. His mother has also been treated for two such lesions.

Discussion

Basal cell naevus syndrome has been compared to other neurocutaneous syndromes such as neurofibromatosis and tuberous sclerosis.^{3,4} There have been no further reports of associated agenesis of the corpus callosum since the initial case report, but various other neurological abnormalities have been described. Intracranial calcification occurring most frequently in the falx cerebri and less commonly in the tentorium cerebelli and petroclinoid ligaments, is found in more than 80% of cases.² Cases of associated intracerebral cysts⁵ and choroid plexus cysts⁶ and communicating hydrocephalus⁷ have been reported. Intracranial neoplasms have also been described, particularly medulloblastoma, and development of this tumour may actually precede the appearance of the skin lesions.⁸ A single case associated with a cerebellar astrocytoma⁹ and two cases with a meningioma have been described.^{10,11} Both meningiomas were incidental post-mortem findings, and in one case there was also a craniopharyngioma.

The finding of four meningiomas, including the two reported here, suggests that development of the tumours is part of the syndrome. Unlike medulloblastomas, meningiomas in the basal cell naevus syndrome appear to present relatively late, the four patients described all being over 40 years old. It is

interesting that our second case had extensive radiotherapy to the region of the left temple prior to the discovery of the left parietal meningioma. Patients with basal cell naevus syndrome appear to be particularly predisposed to tumour formation following radiotherapy. Children with basal cell naevus syndrome and medulloblastoma who, following surgery, were treated with radiotherapy to the neuraxis, all developed multiple basal cell carcinomas in the irradiated areas within six months to three years of radiotherapy.¹² Fibrosarcoma of the ovary has also been described in one of these children following neuraxis irradiation, and fibrosarcoma of the jaw following local radiotherapy to mandibular cysts in two adult patients.^{3,11} Meningioma development in the basal cell naevus syndrome is probably a reflection of a genetic propensity for tumour formation which may be augmented by local radiotherapy.

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