Syringomyelia in association with a neurofibroma of the filum terminale

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SUMMARY A case of neurofibroma of the filum terminale associated with syringomyelia is described. The relationship of spinal cord tumours and syringomyelia is discussed and a possible mode of formation of the syrinx in the present case is suggested.

The association of syringomyelia with intraspinal tumours has been well established during the past 100 years. The present case is being reported because of the unusual findings of a large syringomyelic cavity of the cord together with an apparently anatomically unconnected neurofibroma of the filum terminale.

CASE REPORT

A 15 year old girl was admitted to the Wessex Neurological Unit in April 1970 (case no. 10327) with a three year history of lower lumbar pain. This pain was constant and aggravated by standing, walking, and coughing. In January 1970 she became aware of the gradual onset of an inconstant burning pain in the upper lumbar region radiating around both loins into the left hypochondrium, which was aggravated by the same factors as before. During the next four months she developed a progressive weakness affecting first the left and soon after the right lower limbs. Two weeks before her admission her urinary flow became impaired with an intermittent and variable stream.

EXAMINATION

The configuration and range of movements of the spinal column were normal. Abdominal reflexes were absent and there was moderate weakness of the left and slight weakness of the right lower limbs such that she could stand and walk only with assistance (grades 3 and 4 respectively). Tone was symmetrically increased in both lower limbs and bilateral ankle clonus and extensor plantar responses were present. Pain and temperature sense on both sides of the trunk and of the right lower limb below the level of T7 dermatome were impaired, with the relative preservation of all sensory modalities of the left lower limb. General examination was normal; blood pressure was 120/80 mm Hg.

INVESTIGATIONS

Chest radiographs were normal. Radiographs of the whole spine showed generalized narrowing of the pedicles and widening of the interpeduncular distance from the T3 vertebral level to a transitional vertebra (L6). The thoracic and lumbar vertebral bodies also showed marked scalloping (Fig. 1). Cisternal gas myelography demonstrated a normal cervical cord but gas would not pass beyond the upper thoracic region. Cisternal positive contrast myelography using 3 ml. iophendylate (Ethiodol) confirmed the presence of a block at the level of T5 vertebral body. Analysis of the cisternal cerebrospinal fluid (CSF) obtained at this examination showed 120 mg protein, 62 mg glucose/100 ml., and 4 lymphocytes/c.mm.

Lumbar myelography using 6 ml. iophendylate outlined a continuous sac extending from T7 to L3 vertebra containing a 4 cm × 3 cm mass preventing filling of the lumbosacral space below this level, although later rescanning showed that contrast medium did pass below the lumbar lesion (Fig. 2). The radiological diagnosis was an expanded cord extending from T3 vertebra to the lower thoracic region. There was also an extramedullary, intradural mass at the L3 vertebral level.

Examination of the xanthochromic fluid obtained by lumbar puncture at this time showed a protein content of 4.7 g/100 ml. with a large excess of globulin, glucose 62 mg/100 ml., and 1 lymphocyte/c.mm.

FIRST OPERATION (May 1970: Mr. John Garfield)

In view of the clinical and radiographic findings it
was decided to perform a laminectomy from T4–6 inclusive. A fullness of the dura mater extending downwards from T5 vertebra could be seen and when the dura mater was opened marked expansion of the cord was confirmed. A sharp needle was inserted in the midline of the cord at the level of maximum swelling and 25 ml. clear yellow fluid obtained with resultant collapse of the cord.

This fluid was reported upon by Professor J. N. Cumings (Institute of Neurology, Queen Square) to contain (normal CSF values in parentheses):

- Total protein (15–20 mg/100 ml.), 4-6 g/100 ml.
- Alkaline phosphatase (0 i.u./l.), 44 i.u./l.
- Cholesterol (less than 1 mg/100 ml.), 88 mg/100 ml.
- Phosphohexose isomerase (0 i.u./l.), 83 i.u./l.
- Lactic dehydrogenase (0–10 i.u./l.), 35 i.u./l.

Professor Cumings suggested that these findings were consistent with a cyst within a relatively benign tumour and would not be obtained from cerebrospinal fluid.

**Progress** The patient's immediate postoperative recovery was satisfactory, but on the fifth day she complained of weakness and paraesthesiae of both hands; examination revealed severe weakness of both lower limbs (grade 0 on the left and grade 1 on the right). There was weakness throughout the right upper limb with sensory loss from C6–T3 dermatomes on that side, there being a band from T3–7 dermatomes in which sensation was normal and below that the sensory level was as previously
described. Thereafter there was only gradual and barely significant improvement in the power of her lower limbs.

SECOND OPERATION (5 June 1970: Mr. John Garfield) The deterioration of the patient's condition was attributed to a block occurring at the level of the previously demonstrated intradural lumbar mass and accordingly a laminectomy was made from L2-4 inclusive. On opening the dura mater, a firm solid tumour adherent to the nerve roots was seen. The tumour was freed from the surrounding nerve roots and it became apparent that it was arising from the filum terminale. This structure was divided above and below the mass, which was then completely excised. It was clear that the tumour lay in a cystic cavity, which contained yellow fluid and droplets of iophendylate, the upper part of which was removed establishing free communication between the cavity and the sub-arachnoid space.

Histology The tumour measured 2-8 cm × 1-8 cm and was enclosed in a thin intact capsule. On section it was largely solid but contained some necrotic areas. It was vascular with numerous thin- and thick-walled blood vessels distributed irregularly through a cellular stroma which was sometimes reticular and elsewhere arranged in interweaving bundles. Nuclear palisading was occasionally a feature of the latter. There was no evidence of rapid growth or malignant change.

The stromal cells produced reticulin.

A pathological diagnosis of neurofibroma was made.

After the second operation she made good progress and within two weeks she had regained full bladder control, increased power of her lower limbs, and she was beginning to walk. The appreciation of sensation below the level of T6 dermatome was improving, while her upper limbs had returned to normal.

Examination six months later showed that she had continued to improve, bladder control remained normal, and her gait was excellent. Sensation was normal apart from a band of hypoaesthesia over the T10 and T11 segments on the left side of her trunk. The plantar responses were still extensor.

DISCUSSION

The generally accepted pathological description of syringomyelia is of a longitudinal cavitation of the spinal cord or brain-stem in which the cavity may be unrelated to the central canal and is lined at least in part by glial tissue. Recent work (Gardner, McCormack, and Dohn, 1960; Gardner, 1965) relates the aetiology of syringomyelia to the failure of the exit foramina of the fourth ventricle to develop during embryonic life. In these circumstances there is a water hammer effect on the cerebrospinal fluid in the central canal of the spinal cord, which subsequently dilates. Tamaki and Lubin (1938) suggested that syringomyelia was the result of cavitation in glial tissue following the imperfect development of the central canal. A vascular basis has been put forward by some authors; Russell (1932) considered that there was evidence of anaemia and haemolytic softening in her cases, as did Wolf and Wilens (1934), whereas Tauber and Langworthy (1935) ascribed the syrinx to vascular insufficiency due to vascular anomalies.

In their reports of syringomyelia and intramedullary tumours many authors have discussed the pathological significance of the frequent association of the two conditions, Simon (1875) was perhaps the first to suggest that syringomyelia was possibly the result of softening occurring in the spinal cord 'gliomas'.

Baumler (1887) in his series of 96 cases of syringomyelia found that 17 of these had an intramedullary tumour. Dimitroff (1897) described 12 cases of tumour in 84 cases of syringomyelia that came to necropsy, these figures receiving support from Poser (1956) who collected 245 cases of syringomyelia from the literature and found that 40 (16-4%) of these were proven at necropsy to have an intramedullary tumour.

In their description of 301 cases of primary intramedullary tumour, Slooff, Kernohan, and MacCarty (1964) found that in the 33 cases which came to necropsy, 19 were associated with syringomyelia, as were six of Hamby's (1944) cases and two of Russell's (1932) three patients. Poser (1956) reviewed 209 cases of intramedullary tumour reported in the literature and found that 65 (31%) were associated with syringomyelia. He concluded that the discrepancy between the incidence of intramedullary tumour in syringomyelia and of syringomyelia in cases of intramedullary tumour might be because, where intramedullary tumour was diagnosed at operation, these cases were
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The tumour types associated with syringomyelia are neurofibroma, meningioma, epidermoid, and glioma (Kernohan, 1932). Multiple heterotopic tumours have been described (Tamaki and Lubin, 1938; Poser, 1956), these being usually epidermoids and neurofibromas or meningiomas and Poser and Sytre, 1922; Poser, 1956; Slooff et al., (1964) less frequently lipoma, chordoma, and haemangiopericytoma. Lennert's disease (1967) has also been described (Tamaki and Lubin, 1938). The authors have commented on the apparent continuity of the two lesions. Slooff et al. (1964) described one case (no. 24) where there was an ependymoma of the cauda equina with an anatomically separate cord extending from T4-T12 segments. Poser (1956) could find only 21 (11.3%) of the 185 cases in which the tumour and the syringomyelia appeared to be in direct communication with the subarachnoid space, as the protein content of this study is manifest in as much as 81% of the cases. Suggested mechanism of development of syringomyelia in the present case:

Assuming that the pathogenesis of these two lesions is connected, it would seem probable that the initial formative process is a neurofibroma forming below the conus medullaris, this secondary cyst formation causing the lower lumbar pain which persisted for three years. The onset of the upper lumbar pain radiating into the left hypochondrium, which was of a different character and associated with a progressive parapresis and micturition difficulties, may be attributed to the upper cyst.
extension of the cyst into the cord substance from the conus to the T5 segmental level when the block occurred at the lumbar level (Fig. 3).

It is possible that the syrinx in the present patient was due to the presence of an unproven ependymoma, but the occurrence of multiple heterotopic tumours with an associated syringomyelia is extremely uncommon. The continued improvement of the patient after surgery is encouraging but it will be several years before an ependymoma can be excluded on clinical grounds.

I wish to thank Mr. John Garfield and Dr. P. K. Robinson, under whose care this patient was admitted, for their permission to report this case.

REFERENCES


