capillary forms (Robbins, 1963). Thrombosis of the cavernous spaces and any stage of organization of the thrombi may be present. Some degree of fibrosis is usual and calcification can occur as was seen in our patient on radiological examination.

Considering the pathological anatomy of a haemangioma, it is possible to surmise that red-cell fragmentation could occur in such a situation as a result of coagulation in vascular channels especially when the tumour is very extensive.

References


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Autonomic dysfunction in syringomyelia

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Since the clinical syndrome of postural hypotension was first described by Bradbury & Eggleston in 1925, numerous reports have appeared in the literature recording the association of this syndrome with disease of the nervous or endocrine systems. In this report, we describe a patient who developed autonomic dysfunction, with postural hypotension, in association with syringomyelia.

Case report

Mrs A.T., a 56-year-old housewife with syringomyelia was admitted to the Middlesex Hospital (No. G61625) under the care of Professor R. W. Gilliatt in April 1970, complaining of increasing weakness of the left leg.

The family history was non-contributory. The patient first noticed slight swelling and flexion deformity of the fingers of both hands at the age of 14 years and 2 years later found she could not feel heat normally with her hands. These symptoms gradually progressed and she noticed bruises and blisters on her hands, especially the left, from painless injuries and burns. At 25 years of age she developed weakness of the left arm and hand. She led an active life as a housewife until she was 50 years old, when her legs became weak. At this time she also developed urinary frequency, with nocturia, urgency, occasional urinary incontinence and less frequently faecal incontinence, without loss of bladder or rectal sensation. In the last 4 years she experienced several episodes of unsteadiness which were suggestive of postural hypotension as they occurred immediately after standing. She did not remember having sweated in hot weather for many years.

Examination revealed a dorsal kyphosis and scoliosis. There was no intellectual deficit and the cranial nerves were intact. In the arms there was wasting of the intrinsic muscles of both hands and of the left shoulder muscles, a painless arthropathy of the metacarpophalangeal joints of the right hand and clawing of the fourth and fifth digits of the left hand. Several scars from past injuries were present over both arms and hands. She had generalized weakness of the left arm with distal muscle weakness in the right arm, and loss of all tendon reflexes in the upper limbs. The abdominal muscles were weak and she was unable to sit up without support; the abdominal reflexes were absent. There was a spastic paraparesis.
with hyper-reflexia and bilateral extensor plantar responses. On sensory testing, pin-prick and temperature appreciation were impaired between the fourth cervical and eighth thoracic segments on the right, and between the second cervical and eighth thoracic segments on the left, with preservation of light touch, vibration and joint position sense. In the legs there was some impairment of joint position sense at the metatarso-phalangeal joints and of vibration at the ankles. She had a lax anal sphincter.

The following investigations were within normal limits: full blood count and sedimentation rate; blood urea, electrolytes and sugar; and serum calcium, phosphate and alkaline phosphatase. Serological tests for syphilis and for rheumatoid disease were negative. An electrocardiogram was normal. Three MSUs were sterile on culture.

A chest X-ray was normal. X-rays of the cervical and thoracic spine showed a gross kyphoscoliosis convex to the right with some anterior wedging of the mid-dorsal vertebral bodies. Tomography of the base of the skull showed no platybasia. X-rays of the right hand revealed partial subluxation of the second and third metacarpo-phalangeal joints and of the proximal interphalangeal joints of the ring and little fingers, with new bone formation at the head of the third metacarpal; in the left hand osteoarthritic changes were present in the terminal interphalangeal joint of the middle finger.

In view of her symptoms of orthostatic hypotension, her inability to sweat and her bladder and bowel disturbances, the patient was investigated for evidence of autonomic dysfunction. With the patient supine, her blood pressure varied between 130/85 and 95/50 mmHg; on standing there was a postural drop that varied between 0 and 35 mmHg in the systolic level, and 0–25 mmHg in the diastolic. For example, on one occasion her blood pressure fell from 115/70 mmHg supine to 80/40 mmHg on standing and her pulse rate increased from 78/min to 88/min; this was accompanied by transient dizziness but the patient did not become pale. With continuous blood-pressure recording via an indwelling radial artery needle the patient performed a Valsalva manoeuvre as described by Sharpes, Schafer (1953). During the forced expiration there was a continual fall in blood pressure without tachycardia, so that the reflex latent period (Gross, 1970) could not be measured; at the end of the manoeuvre there was no 'overshoot' of the blood pressure or bradycardia. This type of record is seen in patients with severely impaired circulatory reflexes. The afferent limb of this reflex arc in the ninth and tenth cranial nerves cannot be tested directly (Bannister, Ardill & Fenem, 1967) but chemoreceptor fibres in these nerves were shown to be intact by the rapid intravenous injection of lobeline 2 mg; an involuntary cough was produced after a 14-sec latent period by stimulation of the chemoreceptors (Berliner, 1940). The vasomotor response of the patient to a deep inspiration was recorded in the index fingers of both hands using a plethysmograph; a marked inspiratory vasoconstriction occurred as a spinal reflex (Gilliat, Guttmann & Whitteridge, 1948). The patient was warmed with a radiant heat cradle (800 W) over the trunk until her sublingual temperature had risen by 2.5°C, and quininizarin powder (Guttmann, 1940) used to detect sweating. Sweating occurred in the groins, axillae, palms and soles, and over the eyebrows, but there was a complete absence of thermo-regulatory sweating over the trunk, neck and limbs. However, the intradermal injection of acetylcholine (0.1 ml of 1% solution) in both thighs and forearms produced local piloerection and sweating through an axonal reflex in functionally intact postganglionic sympathetic fibres (Coon & Rothman, 1941; Janowitz & Grossman, 1950; Barany & Cooper, 1956). An intravenous pyelogram was normal. A micturating cinecystogram showed that the bladder held 400 ml; the rise of pressure in the cystometrogram was steep and there was spontaneous leakage around the catheter, leaving approximately 150 ml. The bladder emptied at a gross pressure of 70 cm water by straining. No spontaneous detrusor contraction was seen but the outflow tract appeared normal and there was no residue or reflux. The appearances were those of a 'systolic' bladder (Bates, 1971).

Discussion

The only previous report that we have found in the literature of an association between orthostatic hypotension and syringomyelia is by Ellis & Haynes, 1936, and their patient clearly had co-existent syringobulbia. Although they reached no definite conclusions about the site of the lesion responsible for the vasomotor disturbance, these authors suggested that it might be in the hypothalamus.

This patient with syringomyelia and symptoms of autonomic dysfunction was found to have postural hypotension, impairment of circulatory reflexes and thermo-regulatory sweating, and a 'systolic' bladder. The functional integrity of autonomic afferent fibres and of sympathetic ganglia and postganglionic fibres has been demonstrated by the normal evoked responses to intravenous lobeline and intradermal acetylcholine respectively. This implies that the autonomic nervous system is affected at an intermediate point. An abnormality at foramen magnum level, such as that which is frequently associated with syringomyelia (Gardner, 1965), may have interrupted descending autonomic fibres above the syninx in the upper cervical cord. Alternatively, clinical assessment indicates that the patient has...
Addison’s disease, vitiligo and multiple autoantibodies

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Case report

In 1959 a married English housewife aged 29 was seen because of increasing weight loss, weakness and craving for salt for 1 year with amenorrhoea for 3 months. Previously, she had noticed bald patches on her scalp which had completely disappeared without treatment.

She had not been seriously ill before and had married 5 years previously but not become pregnant. Her father had died of rheumatic heart disease and pulmonary tuberculosis, her mother had been thyrotoxic but was well as were two siblings. She appeared thin and wasted and her blood pressure was 95/70 mmHg. There was brown pigmentation of the buccal mucosa and exposed areas of her skin but there were white patches over the knuckles. There was no poliosis or premature greying of the hair.

Haemoglobin 11.3 g/100 ml, ESR 48 mm/hr, plasma Na+ 133, K+ 6.1 and Cl− 100.5 meq/l.


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References


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a syrinx extending from the second and fourth cervical to the eighth thoracic segments and it may be presumed that this is involving the descending autonomic pathways. The intermedio-lateral columns or autonomic outflow paths in the spinal cord may also be affected, in which case the absence of Horner’s syndrome and the preservation of the inspiratory vaso-constrictor reflex implies that this lesion is incomplete.

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