UNUSUAL NEUROLOGICAL DISORDERS of an unusual cause with an unusual CT Scan picture are the main characteristics of the following case.

Case Report

A 64 year old right-handed manager was admitted on January 1st 1982 for 'right hemiplegia and aphasia'. On the eve of the New Year he was uncooking a bottle in the kitchen when his wife heard a thud. He was found on the ground, conscious, unable to move his right side and to speak. There was no evidence of headache and he did not vomit.

Two years previously, on August 17, 1979 while bending below the dashboard of his car he had felt a sharp non throbbing pain in the left parietal region. This lasted 2 minutes. Simultaneously he had experienced difficulty in moving his right lower limb for 10 minutes. Medical examination half an hour later was reported normal. From August 19 to September 2, once or twice a day, he had a transient motor deficit of his right lower limb for 5 to 10 minutes, with or without concomitant left parietal headache. No particular activity or exertion were noted at the onset of these episodes. On September 1, 1979 he felt pin and needles in his left thigh for 10 minutes. One of us (J.C.G) saw him on September 4, 1979. Neurological examination was considered normal. No bruit was heard in the neck or over the skull. Blood pressure was 150/90 mmHg. There was no evidence of subarachnoid hemorrhage.

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area persisted in the left frontal lobe. On March 18 the malformation was operated (Pr. Pertuiset). The abnormal veins were outside the brain and were excised. Post-operative angiography showed no abnormal vessels. The evolution was simple and the patient was discharged home.

He was seen again as an outpatient on June 6, 1982. Voice and speech were normal; bladder and rectal control were normal. The patient thought that his intellect was normal. He had not resumed his professional activities.

Comments

Several classifications of dural arterio-venous fistulae (AVF) have been proposed. The more recent one distinguishing among pure meningeal AVF those draining: 1) into a sinus or meningeal vein; 2) into a sinus with a significant reflux into veins that arrive at the sinus; 3) into cortical veins; 4) into large dural or subdural lakes, acting like space occupying lesions. Our case belongs to (3) a condition which is obviously liable to result in cerebral and/or subarachnoid hemorrhage.

Two years and 3 months prior to the cerebral hemorrhage there were during two weeks, once or twice a day, transient episodes of paresis of the right lower limb. Once the transient episode was of a sensory nature. There were never jerks. The first episode occurred while the patient was bent forward below the dashboard of his car but no particular exertion or posture was noticeable at the beginning of the numerous other episodes. Some of the latter were accompanied by a sharp non throbbing pain in the left parietal region but many were not. What were these episodes? Obviously their brief duration rules out hemorrhage. There were no jerks prior to the motor deficit and no march of pin and needles so epilepsy is unlikely. Were they transient ischemic episodes? Pain concomitant to TIA’s is rare indeed but the pathological condition of this patient, i.e. dural AVF draining into cortical veins is rare as well. It might be that we are here dealing with a particular kind of TIA in which pain in the head would be a particular feature.

Between September 1, 1979 and December 31, 1981 there were no symptoms. During this period 2 CT Scans with contrast showed an unusual picture of the superior sagittal sinus (fig. 1) to which several hyperdense nodes were appended. This picture didn’t lead to diagnosis and among (false) hypotheses a rare kind of meningiomatosis was mentioned by a neurosurgeon to whom the case was submitted. CT Scan appearances of dural AVF are not well known and this particular picture appears not to be included in a recent paper. Can this picture be explained a posteriori? It is likely to be due to reflux of contrast medium into veins adjacent to the superior sagittal sinus meaning high pressure in the sinus and this of course is suggestive of an AVF. Anyway the CT Scan is so particular as to be easily reminded with its particular cause in this particular patient.

The left frontal hematoma resulted in a rather typical syndrome of the frontal lobe: contralateral grasping reflex, disorders of sphincter control, laconic responses, abulia. The latter has been recently briefly commented. A striking feature of speech was whispering. Although the latter is listed in classical Textbooks in the anterior cerebral artery syndrome its physiopathology is not easily understood. In our patient (and maybe generally) it might represent a period
between mutism and normal speech or voice. It may be for voice what akinesia and hypokinesia are for movements. Is it present in right frontal lobe lesions as well as in left ones? If yes, are there some disimilarities? Additional cases deserve to be reported with more detailed studies.

References
A cerebrovascular accident with unusual features.
J C Gautier, A Awada and P Loron

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