Abnormal arborisations of Purkinje cell dendrites in Creutzfeldt-Jakob disease: a manifestation of neuronal plasticity?

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SUMMARY A case is presented of the ataxic variety of Creutzfeldt-Jakob disease with particular reference to the cerebellar cortex. The main features were loss of granule cells, subtotal in the vermis, severe in the lateral lobes, mild to moderate loss of Purkinje cells and preservation of tangential and basket fibres. The Purkinje cell dendrites showed malorientation and hypertrophy of the primary and secondary branches, the so-called "antler" or "staghorn" deformity. These findings indicate that remodelling of the dendritic tree may start early in the course of the disease even in adults, the total length of history in this case being eight months. They do not throw any additional light on the pathogenesis of the dendritic abnormalities, in particular on the controversy whether they are a non-specific response of the Purkinje cell to a variety of noxious agents or a reaction to partial deafferentation. The authors favour the latter hypothesis.

Bizarre ramifications of Purkinje cell dendrites were described by Nagotte and Léon-Kindberg and by Cajal. Neither author provided sufficient data to throw light on the pathogenesis of these abnormalities. Subsequently similar lesions were described in a variety of unrelated conditions, most of them originating in infancy and early childhood. These include primary degeneration of the granular layer, both familial and sporadic, Menkes' disease and the "amaurotic idiocies", in particular Batten's disease or ceroid lipofuscinosis. More recently, similar observations have been made in other conditions, such as cerebello-brain stem leukodystrophy and ornithine carbamyl transferase deficiency. In addition, a small group of adult conditions with similar changes has been reported: industrial organic mercury poisoning, Minamata disease and crossed cerebellar atrophy. A common feature of all these conditions is severe rarefaction or subtotal loss of neurons in the granular layer of the cerebellar cortex. The widely held interpretation of the Purkinje cell abnormalities is that they represent a non-specific response to a variety of unrelated factors, genetic, metabolic, toxic or infective. An alternative hypothesis ascribes the changes in Purkinje cell dendrites to partial deafferentation. In the latter case these abnormalities would be an expression of neuronal plasticity which may be defined as an attempt by the neuron to establish new synaptic contacts when its normal connections have either failed to develop (developmental plasticity) or have been destroyed (reactive plasticity).

The main difference between the infantile and the adult cases appears to be in the tempo of evolution of the dendritic abnormalities. They seem to develop rapidly in the former group and may be observed at an early age, while, in the latter, they have been recorded only in cases of extreme chronicity. We have therefore considered it of interest to investigate an adult case with a relatively short clinical history. A case of the ataxic form of Creutzfeldt-Jakob disease appeared to be suitable for this purpose, since atrophy of the granular layer is a feature of this condition as well as of kuru.

Materials and methods

The material consisted of the brain of a case of the ataxic form of Creutzfeldt-Jakob disease. Representative blocks were taken from the cerebral cortex, basal ganglia, thalamus and brain stem, as well as multiple blocks of the cerebellum, both of the vermis and the lateral lobes. All blocks were embedded in paraffin and stained by standard neuro-
difficulty with 48-year-old Hispanic A report Case vertigo and by stained (PAS) and techniques, including periodic-acid pathological of tangential and basket fibres and “antler” deformation of Purkinje cell dendrite. (Bielschowsky’s silver impregnation x 400.)

pathological techniques, including periodic-acid Schiff (PAS) and a modification of Bielschowsky’s silver impregnation. In addition, frozen sections of the vermis were stained by Cajal’s silver nitrate-pyridine (SNP) method.

Case report

Clinical history

A 48-year-old Hispanic male school teacher was in good health until September 1983 when he started complaining of vertigo and diplopia. Subsequently he developed progressive difficulty with balance, fell frequently and had difficulty climbing stairs. On admission to hospital in December 1983 he was fully oriented with some intellectual impairment and short term memory deficits. Other signs included jerky ocular movements, cerebellar dysarthria, broad based gait, truncal ataxia and past pointing, but no myoclonus. CT scan showed atrophy of the cerebellar vermis. The diagnosis of Creutzfeldt-Jakob disease was finally confirmed by brain biopsy. The patient died in May 1984 after a progressive downhill course with increasing dementia.

Pathology

Cerebral biopsy Fine spongiform changes were seen in the neuropil largely confined to the deeper cortical layers (V and VI). Electron microscopy showed typical double membrane bound complex vacuoles in the neuropil.

Necropsy revealed no significant general findings. The brain after fixation weighed 1150 grams. Mild cortical atrophy, particularly of frontal and parietal lobes was present with severe cerebellar atrophy with shrunken folia, both in vermis and lateral lobes. Coronal sections revealed flattening of the caudate nucleus and marked atrophy of the putamen.

Microscopy The cerebral cortex showed marked spongiform changes in the neuropil, loss of neurons and proliferation of reactive astrocytes, mainly in the deeper layers, in all areas examined. The putamen and caudate nucleus were atrophic, with almost total neuronal loss, intense astrocytic proliferation and coarse status spongiosus. Focal spongiform change, neuronal degeneration and astrogliosis were also present in the thalamus. In the brain stem similar changes were seen in the tectum of the midbrain and basis pontis. In the cerebellum the most striking feature was loss of granule cells, subtotal in the vermis, severe in the lateral lobes, with partial preservation on their inferior surface. The tangential and basket fibres were generally well preserved; some empty baskets were seen in places. Sparse “kuru” plaques were found in the molecular layer. Loss of Purkinje cells was mild to moderate. Many of the surviving Purkinje cells showed abnormalities of their dendrites. Some primary dendrites showed malorientation and were running obliquely or horizontally. Extensive ramification of thick
cases of Minamata bellar atrophy after the poisoning which opinion disease, even in Shiraki was seen in the present they thematic remodelling and expansions stellate. These perikarya and diffuse fusiform ("torpedoes") are frequently by methods impregnations metallic obliquely. The dendritic tree, begins in some adults, and is already conspicuous in the total number of Purkinje cells, observed both in the inferior olive and terminal ramifications (baskets) is a conspicuous feature of Menkes' disease. On the other hand, they are remarkably well preserved in Creutzfeldt-Jakob disease as they were also in the present case. It is tempting to suggest that persistence or neoformation of perisomatic dendrites is associated with the absence or loss of axosomatic synapses between the basket fibres and the Purkinje cell soma.

Another abnormality associated with the absence of basket fibres is the malalignment of Purkinje cell perikarya due to their dislocation into the molecular layer. This may be tentatively ascribed to loss of tethering of the Purkinje cell by its afferent fibres. It was not a feature of the present case.

In conclusion, the observations on this case provide evidence of early remodelling of Purkinje cell dendrites in the direction of the "antler" deformity. The absence of other abnormalities does not contribute to our understanding of their pathogenesis, but does not contradict the current views on their evolution. Our findings do not resolve the controversy between the theories which ascribe the abnormalities of Purkinje dendrites to a direct effect of noxious agents on one hand, and to neuronal plasticity in response to loss of afferents on the other. Excessive, often bizarre, dendritic ramifications in the inferior olive are associated with lesions in the ipsilateral central tegmental tract or contralateral dentate nucleus. Similar dendritic abnormalities in the substantia nigra were observed in association with long-standing massive striatal infarction. The unifying hypothesis which ascribes all these dendritic abnormalities at various sites to a single mechanism, response to partial deafferentation, appears to us far more attractive.

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References


