Are alpha-1-antichymotrypsin and inter-alpha-trypsin inhibitor peripheral markers of Alzheimer's disease?

The definite diagnosis of Alzheimer's disease (AD) requires both clinical criteria of probable AD and neuropathological evidence of AD lesions. At present there is no laboratory test for a premortem diagnosis. Recently, genetic and histochemical studies identified protease inhibitors as components that might be implicated in the formation of the amyloid substance in AD brains. First, Abraham et al. suggested a potential role of alpha-1-antichymotrypsin (ACT) in the pathogenesis of the lesions, moreover Matsubara et al. found an increased serum concentration of ACT in AD. Second, several authors showed that one transcript of A4 amyloid precursor contained an additional sequence similar to the active site of inter-alpha-trypsin inhibitor (ITI). The purpose of our study was to test the diagnostic value of ACT and ITI in serum and CSF from AD patients.

Sera and CSF were collected from eight men and 16 women with probable AD, mean (SD) age 66 (8.3) years, and from a control group of 19 men and six women aged 64 (8.3) years. Controls were volunteers free of any neurological disease, with a MMS score higher than 28, who had had a myelo- or radioculography for proven disk herniation. CSF was not collected especially for this study. The procedure was approved by the ethical committee of Lille. ACT and ITI contents were measured by electrophoresis-diffusion methods. Semi-quantitative determination was used for ITI in CSF because of its low concentration. Statistical assessment used non parametric tests (Mann and Whitney's U test and Spearman's rank correlation test).

In the control subjects there were 1) no difference in serum or CSF ACT and ITI contents between males and females, 2) no correlation between age and both serum ITI and CSF ACT contents, 3) a positive correlation between serum ACT contents and age (p < 0.02).

Between AD patients and controls, there were no difference in serum or CSF ACT and ITI contents, and no difference of the ACT/CSF ratio in serum (table).

Our results show that ACT and ITI are not useful markers of AD in serum and CSF. They don't confirm that Abraham et al. The ACT/CSF serum ratio was not significantly modified in AD patients, which is consistent with the hypothesis that the blood-brain barrier is not strongly affected in this disease. The correlation between serum ITI contents and the severity of the dementia could be explained by non specific metabolic disturbances.

**Table: Serum Alpha-1-antichymotrypsin (ACT) and Inter-alpha-trypsin inhibitor (ITI) contents, CSF ACT contents and ACT/CSF ratio in serum and CSF of Alzheimer's disease (AD) patients.**

<table>
<thead>
<tr>
<th>Control group</th>
<th>AD Patients</th>
</tr>
</thead>
<tbody>
<tr>
<td>Serum</td>
<td></td>
</tr>
<tr>
<td>ACT mean (SD)</td>
<td>6.07 (0.27)</td>
</tr>
<tr>
<td>CSF</td>
<td>6.97 (4.45)</td>
</tr>
<tr>
<td>ITI mean (SD)</td>
<td>10.71 (4.5)</td>
</tr>
<tr>
<td>Serum</td>
<td></td>
</tr>
<tr>
<td>ACT mean (SD)</td>
<td>0.71 (0.19)</td>
</tr>
</tbody>
</table>

Postiradiation motor neuron syndrome of the upper cervical region—a manifestation of the combined effect on cranial irradiation and intrathoracic chemotherapy?

CNS prophylaxis is now an integral part of the treatment of acute lymphoblastic leukaemia and was regarded as the United Kingdom Acute Lymphoblastic Leukaemia Trial 4 (UKALL 4) (intensive) schedule. This comprised induction with cyclophosphamide, cytosine arabinoside (ara-C), vincristine, prednisolone and intrathecal ara-C; consolidation with the same, together with adriamycin, asparaginase, 6-mercaptopurine, intrathecal methotrexate and cranial irradiation; and maintenance with vincristine, methotrexate, ara-C, 6-mercaptopurine and prednisolone. The total dose of irradiation was 2400 cGy (rads) and the field extended to the level of the C3 vertebral body.

Apart from an early bone marrow relapse in June 1977, he made a complete recovery. In particular, there was no evidence of CNS involvement at any time.

He received his last dose of vincristine in May 1979 and completed his chemotherapy by June 1979. The period of cranial irradiation spanned 19 days in April 1977.

In January 1981 he was referred to the neurology clinic with a three month history of progressive painless wasting and weakness of the shoulder girdles. There was marked bilateral winging of the scapulae, left worse than right. The trapezi, rhomboids, supra- and infraspinati, deltoids, teres major and both sternocostal and clavicular heads of the pectoralis major muscles were wasted, more on the left, and power was reduced to grade 4 on the left and 4+ on the right. There was a minimal weakness in the biceps, triceps and flexor carpi ulnaris. The thenar, hypothenar, and lumbricals were intact. The triceps muscles were spared as were the distal upper limb muscles and lower limbs. There was questionable weakness of the orbicularis oculi and frontalis to maintain elevation of the eyelids. Although his face was thin there was no focal wasting or demonstrable weakness of the other facial muscles. There were no sensory symptoms or signs. Tendon reflexes were well preserved and symmetrical. Plantar responses were flexor.

Investigations at this stage including muscle enzymes, thyroid function, cerebral spine radiographs, haematocrit screening and bone marrow were normal. Electroencephalograph EMG studies revealed reduced amplitude and arrhythmic action potentials. Evidence of chronic partial denervation of both deltoids, more on the left.

Thereafter the condition appeared to arrest with no objective progression noted during eight years of follow up (1981–9). Serial EMGs showed evidence of chronic partial denervation and reinnervation in the brachioradialis, biceps, deltoids, supraspinatus and trapezius muscles without pathological activity at rest. No significant abnormality was demonstrated in the quadriceps. In the right tibialis anterior a full interference pattern contained occasional polyphasic units of normal amplitude and duration which were not felt to be of clinical significance. Muscles
sampled serially showed little change with time.

Nerve conduction studies showed normal motor latencies, conduction velocities and F wave latencies. The ulnar sensory nerve action potential was increased, and mixed action potentials of the median nerve were of reduced amplitude but also had normal conduction times, suggesting an axonal degeneration.

A biopsy of the deltoid and quadriceps muscles showed no specific changes only and no dystrophic features.

In February 1989 an MRI scan of the cervical spine showed the upper cervical cord to be of normal size with increased signal intensity on the T2 weighted spin echo sequence, the significance of which was unclear.

This patient developed an asymmetrical and patchy wasting and weakness of the shoulder girdle muscles involving several myotomes from C3 to C7. The explanation for the symptomless minimal weakness of the orbicularis oculi and frontalis muscles is uncertain. It may reflect patchy involvement of a similar, symptomless involvement of the other cranial nerves such as the bulbar muscles (which escape detection) or might merely be constitutional.

We postulated that the neurological abnormalities in our patient are a manifestation of augmentation of irradiation by combination intrathecal and systemic chemotherapy with two potentially neurotoxic agents, methotrexate and cytosine arabinoside, in the context of the intensive UKALL 4 regimen. The predominant features of this case are in keeping with a postradiation motor neuron syndrome. The minor ulnar sensory abnormalities noted on EMG and without associated clinical symptoms or signs were probably secondary to vincristine.

Four types of radiation myelopathy have been described, but the least common being low-grade motor neuron syndrome. The mechanism underlying this phenomenon is unclear but there is evidence from clinical and pathological studies that radiation injury to vascular endothelium produces ischaemia which leads to selective anterior horn cell degeneration.

Greenfield and Stark observed this phenomenon in three patients, and Sadowsky et al reported it in a fourth. All four cases comprised of a selective lower motor neuron disease confined to lower limb muscles starting three to eight months after radiation of the spinal axis. All followed a subacute and self-limiting course. In our patient the latent period was longer (2½ years), in keeping with other studies where the average symptom free interval was 14 months, but otherwise the disease followed a similar course. Unlike the other cases described, he did not receive direct irradiation to the spinal cord apart from that part of the upper cervical cord included in the field during cranial irradiation.

Byfield et al noted that in the infant who developed radiation myelitis of the cervical cord after receiving routine doses of vincristine and radiation therapy and postulated a synergistic effect. In 1975 a histopathological study by Price et al suggested a similar synergism between irradiation and intravenous methotrexate in the development of fatal leukoencephalopathy in children with ALL. Intrathecal therapy has a distinct neurotoxic effect ranging from a chemical arachnoiditis to transient/permanent paresis and encephalopathy.

Cytosine arabinoside, the other intrathecal agent in our regimen, can cause disseminated multifocal coagulation necrosis of white matter and has been implicated in enhancing radiation induced central nervous abnormalities. A recent study has shown that intrathecal ara-C significantly reduces the isoeffect doses required for the development of radiation damage in rat spinal cord.

We are not aware of any previous reports describing an upper cervical cord motor neuron syndrome occurring following cranial irradiation. Would it be interesting to hear if others have encountered this feature in leukemic patients treated similarly.

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Optic nerve cysticercosis: a case report

Cysticercosis is caused by infection from the larval form of Taenia solium and humans are an intermediate host. The subcutaneous tissue, brain, eye, muscle, heart, liver and peritoneum are common sites of encystment. In the eye, the conjuctiva is the most common site. Subconjunctival or subretinal location have also been described. Cysticercosis of the intracranial portion of the optic nerve is rare. A 15 year old school girl presented with deteriorating vision in her right eye which had progressed over a seven month period. There was no history of pain or inflammation in the eye, trauma, headache, vomiting, seizures, or any other general physical or systemic examinations were normal. There was no subcutaneous nodule or face-au-lait spots. Neurological examination was also normal except for the patient’s inability to count fingers at less than 30 cm with the right eye. The pupils were equal and reacting. Fundus ocular examination revealed disc pallor in the upper half, disc margins elevated in the lower half, and a small white lesion near the optic disc for an area of two dipters below the disc. Vessels and macula were normal. Pigmentary changes were present. The tension was 17.3 mm in both eyes. Other cranial nerves were normal. The rest of the neurological examination did not reveal any abnormality.

Haematological and biochemical parameters were normal. The erythrocyte sedimentation (ESR) was 26 mm and the VDRL was negative. Skull radiographs of the optic foramen and superior orbit fissures were normal. CT scan of the head and orbit with contrast enhancement showed a retrobulbar segment of the right optic nerve thickened with a small area of low attenuation in the thickened portion of the optic nerve. Retrobulbar fat was preserved and the muscle cone was normal. The optic nerve at the orbital apex appeared to be of normal width. The brain parenchyma was normal as was the left optic nerve (fig). Perimetry revealed superior altitudinous right hemianopia. Ultrasonomography showed a mass in the region of the right optic nerve. The Casoni test was negative.

A diagnosis of optic nerve glioma or granuloma was considered. A right frontal craniotomy and extradural frontal orbitotomy was performed. After incising the tenon’s fascia normal retrobulbar fat protruded. The optic nerve was exposed by microdissection and was found to have fusiform thickening. A small portion of nerve just behind the optic globe was normal as was nerve near the apex. A tenuous incision was made over the maximum bulge. There was intense fibrosis. On deeper incision a sago grain like cyst was found and excised. Histopathology revealed it to be cysticercosis. Postoperatively the patient’s vision fully recovered but there was IIIrd nerve paresis.

Cysticercosis is one of the most serious public health problems in the developing countries. Any part of the neuraxis can be involved, except the peripheral nerves, resulting in protein features. Ocular cysticercosis occurs in 3% of cases and may be single, unilateral or bilateral. Subretinal involvement of the eye usually occurs initially through the posterior ciliary arteries but migration of the parasite is common. The nasal side of the eye is more commonly involved than the lateral side. This is due to an anatomical peculiarity of the ophthalmic artery which after giving rise the lacrimal branch runs along the medial side of the orbit and divides into its terminal branches. The optic nerve obtains its blood supply from the branch of the central artery of the retina and retinal blood vessels may thus be involved.

The usual symptoms are of pain, irritation of the eyes due to iridocyclitis and diminness of vision. The eye may be involved alone or may be associated with other clinical features of neuro-cysticercosis when the brain is also involved.

Optic nerve involvement is rare in any kind of cyst or granuloma. As a result, in our

Figure CT scan shows thickened retrobulbar segment of right optic nerve with small area of low attenuation in the thickened portion.
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