Primary orbital amyloidosis localised to an extraocular muscle

GERD E HOLMSTRÖM AND KARL G NYMAN

From the Department of Ophthalmology, Danderyds Hospital, Stockholm, Sweden

SUMMARY Primary, localised orbital amyloidosis has been reported previously in only 15 patients. A case with amyloidosis strictly localised to an extraocular muscle is presented.

Ophthalmic involvement frequently occurs in generalised amyloidosis.12 Localised orbital amyloidosis is very rare and has been reported in only 15 patients. A patient with amyloidosis strictly localised to an extraocular muscle is presented. This has not been previously reported.

Case report

For the last 10 years a 60-year-old woman had a tendency to diplopia when looking quickly to one side. In 1984 a clinical examination revealed a right exotropia, and a Lees screen showed reduced motility of her right eye, particularly a reduction of the adduction—that is, underaction of the medial rectus muscle. There was also a strong overaction of the lateral rectus muscle of the left eye. A forced abduction test of the right eye revealed great resistance in both adduction and abduction. The patient had normal visual acuity and only a slight hypermetropia (+0.5 right and left eyes). Her fundi were normal and no visual field defects were noticed. Computed tomographic (CT) scanning showed a normal brain configuration, but in the right orbit the medial rectus muscle had a fusiform thickening, being three times as thick as the left medial rectus muscle (Figs. 1, 2). No infiltration or destruction of the surrounding structures was seen.

In January 1985 the patient underwent surgery for her right exotropia (resection of the right medial rectus muscle and recession of the right lateral rectus muscle, with a traction suture in adduction). A biopsy of her right medial rectus muscle was also

Correspondence to G E Holmström, Department of Ophthalmology, Danderyds Hospital, S-182 88 Danderyd, Sweden.

Fig. 1  CT scanning of the orbits, 4 mm thick section parallel with the optic nerves.

Fig. 2  Reconstruction of sections retrobulbarly through the orbits, demonstrating the optic nerves and the extraocular muscles in frontal projection.
Primary orbital amyloidosis localised to an extraocular muscle

performed. This muscle appeared waxy and showed ordinary muscle structure only in the 2 mm closest to its insertion. Pathological examination revealed an amorphous structure which, after staining with Congo red, showed a green birefringence. Only a few fragments of the ordinary muscle structure could be identified. The histopathological diagnosis was amyloidosis.

A pulmonary x-ray was normal. Phonocardiography and ultracardiography did not show any abnormality. Biopsies of the subcutaneous fat from the abdomen and the cheek showed no amyloid deposits. No proteinuria was found, and the results of serum and urine electrophoresis were normal. No rheumatoid factor or antibodies against cell nuclei, smooth muscle, glomeruli, and mitochondria could be found. There was therefore no sign of generalised amyloidosis.

In April 1985 the patient underwent supplementary surgery for a slight esotropia with intermittent diplopia (recession of the right medial rectus muscle with an adjustable suture). A CT scan in July 1985 showed the same picture as before.

The patient will be followed up with ophthalmic examinations, tests of visual fields, and CT scanning.

Discussion

Amyloid is an extracellular, proteinaceous material that can be deposited in various tissues. Histopathological identification is best made with Congo red staining. Whenever with a polarising microscope all forms of amyloid show a green birefringence.

Amyloidosis is the term for a heterogeneous group of diseases characterised by accumulation of amyloid in different tissues. It has been subjected to many classifications. A clinically widely used classification is: (1) primary generalised amyloidosis (without any underlying disease); (2) secondary amyloidosis (associated with, for example, chronic infection, rheumatoid arthritis); (3) amyloidosis associated with myelomatosis; and (4) tumour-forming amyloidosis, characterised by localised amyloid masses. Some authors also note a fifth, hereditary, form of amyloidosis. Analyses of the chemical structure of amyloid are now making new classifications possible.

Ophthalmic involvement has been reported in all subgroups of amyloidosis described above. The case presented here is of localised orbital amyloidosis. Apart from amyloid tumours of the conjunctiva and lid, localised orbital amyloidosis is rare, and only 15 patients with primary localised orbital amyloidosis have been reported so far. Four of the previously reported 15 cases were bilateral. Three patients had their lesions localised in the lacrimal gland. In most of the remaining patients the amyloid tumour extended deep and infiltrated the orbit. No case has previously been reported in which the amyloid tumour is localised only to an extraocular muscle.

In conclusion, localised orbital amyloidosis seems to be rare. This is the first case reported with amyloid entirely localised to an extraocular muscle.

Dr A Moberg, Department of Pathology, Danderyds Hospital, kindly helped with the histopathological investigation.

References


Accepted for publication 3 April 1986.