SHORT REPORT

Balo’s concentric sclerosis: a clinical case study of brain MRI, biopsy, and proton magnetic resonance spectroscopic findings

Mee Okh Kim, Sang Ahm Lee, Choong Gon Choi, Joo Ryoung Huh, Myoung Chong Lee

Abstract
The antemortem diagnosis of Balo’s concentric sclerosis was made in a 52 year old woman with subacute right hemiparesis on the basis of brain MRI and stereotactic brain biopsy, which showed multiple ring-like lesions of lamellated demyelination alternating with spared white matter. Proton magnetic resonance spectroscopy (1H-MRS) was carried out one and nine months after the onset of illness. The first 1H-MRS showed a decreased N-acetyl aspartate peak, an increased choline peak, presence of large lipid peaks, and high resonance at 1-4 ppm. The second 1H-MRS disclosed changes such as a decrease of lipid signal, a decrease of resonance at 1-4 ppm, and an increase in the myoinositol peak. These findings are similar to those reported for multiple sclerosis. It seems that this is the first report of 1H-MRS findings in Balo’s concentric sclerosis.

(J Neurol Neurosurg Psychiatry 1997;62:655–658)

Keywords: Balo’s concentric sclerosis; brain MRI; brain biopsy; proton magnetic resonance spectroscopy

Balo’s concentric sclerosis is a rare demyelinating disorder. It is pathologically characterised by alternating bands of demyelinated and myelinated white matter.1–3 The pathogenesis of these banded patterns are still poorly understood. Proton magnetic resonance spectroscopy (1H-MRS) offers a non-invasive way to study biochemical changes that may help to understand the pathogenesis.

Recently, some patients with Balo’s concentric sclerosis have been diagnosed at antemortem examination using brain MRI and biopsy.4,5 However, in vivo 1H-MRS findings in Balo’s concentric sclerosis has not yet been described.

We present 1H-MRS findings in a patient with Balo’s concentric sclerosis, diagnosed by MRI and stereotactic brain biopsy.

Case report
A 52 year old woman presented with subacute right hemiparesis. She was well until two weeks before admission, when she noted right leg weakness. By the next week she had developed weakness of her right arm.

Neurological examination disclosed a right hemiparesis with right facial weakness of a central type. The deep tendon reflexes were increased with extensor plantar response on the right side. A snout reflex was found.

Analysis of CSF showed mild pleocytosis (9 white blood cells/mm3), normal total protein, and no oligoclonal band. The IgG index in CSF was 0.79. Brainstem auditory evoked potentials, pattern shift visual evoked potentials, and somatosensory evoked potentials were all normal. Routine haematological and biochemical tests were normal.

Brain MRI showed multiple lamellated white matter lesions with marginal gadolinium enhancement in the right frontal lobe and bilateral centrum semiovale (fig 1A). Treatment was started with intravenous dexamethasone. The right hemiparesis began to improve during the first week of treatment. Brain MRI was repeated 14 days after treatment. The ring-like pattern of lesions was not changed but the right frontal lesion was considerably increased in size despite clinical improvement. The brain biopsy was done on the 22nd day in hospital. The right hemiparesis gradually resolved over the next five months. She has done well without relapse for more than 17 months.

Pathological findings
An MR guided stereotactic needle biopsy was obtained from the left centrum semiovale lesion. Haematoxylin-eosin and luxol fast blue staining of biopsied brain sections showed the alternation of bright and dark bands under light microscopy (× 40); the dark bands represented preserved myelinated areas and the bright bands demyelinated ones. An immunohistochemical study with HAM 56 antibody disclosed much increased numbers of macro-
phages, but only in the demyelinated areas (fig 1B). In the demyelinated areas, reactive astrocytosis, foamy macrophages, and perivascular mononuclear cells infiltration were noted under light microscopy (× 200).

Proton magnetic resonance spectroscopy (1H-MRS)
In vivo 1H-MRS using a GE 1.5T Signa MR system was performed one and nine months after the onset of illness. Local proton spectra were recorded with stimulated echo acquisition mode (STEAM) sequences (TR 3-0 s, TE 30 ms, 64 AVG, volume 8-9 ml) before gadolinium enhancement.

The first 1H-MRS was obtained from the left enhancing centrum semiovale lesion. It showed a decrease in N-acetyl aspartate:creatine ratio and an increase in choline:creatine ratio. The more striking findings were the presence of large resonances at 0-9, 1-3, and 1-4 ppm (fig 2A).

The second 1H-MRS was obtained from the right non-enhancing centrum semiovale lesion because brain biopsy was done at the left side. Compared with the first 1H-MRS, the large resonances at 0-9, 1-3, and 1-4 ppm were very reduced and the myoinositol peak was significantly increased (fig 2B). There were no significant changes in N-acetyl aspartate and choline peaks.

Discussion
The antemortem diagnosis of Balo’s concentric sclerosis was made in our patient on the basis of brain MRI and stereotactic biopsy, which showed multiple ring-like lesions of lamellated demyelination alternating with spared white matter. These findings are the hallmarks of Balo’s concentric sclerosis. The pathogenesis of these banded patterns in large cerebral lesions are still poorly understood. Recently, Yao et al considered that oligodendroglial loss is important in the pathogenesis of demyelination and that partially myelinated areas probably represent stages of ongoing myelin breakdown rather than remyelination of previously demyelinated areas.10

1H-MRS may provide new insights into various brain diseases, especially into demyelinating diseases. There are several reports of the value of 1H-MRS in diagnosing multiple sclerosis, monitoring its progression, and evaluating its response to treatment.11-18 Balo’s concentric sclerosis has apparently not yet been studied with 1H-MRS.

In our case 1H-MRS was initially studied at the acute enhancing lesion one month after the onset of symptom. The characteristic findings were a decrease in the N-acetyl aspartate:creatine ratio, an increase in choline:creatine ratio, and a presence of high resonances at 0-9, 1-3, and 1-4 ppm. These spectroscopic findings are very similar to those found in acute plaques of multiple sclerosis.11-13 15 N-acetyl aspartate has been implicated as a neuronal marker and its decrease may be explained by neuronal or axonal loss and gliosis.11-15 The increase in choline may reflect the breakdown of myelin membrane phospholipids.11-13 15 The short echo 1H-MRS can detect myelin breakdown products, which probably cause an increase in lipid resonances. Short echo spectra from acute enhancing multiple sclerosis plaques invariably show the presence of large lipid resonances at 0-9 and 1-3 ppm. These resonances are not due to extra voxel contamination from the scalp19 and have been assigned to the methyl and methylen groups of lipid.19 A resonance at 1-4 ppm was very prominent in our case. It is not yet clear which metabolite produces this resonance. There is some evidence that the polypeptide thymosin 4 is found in macrophages and in a subset of oligodendrocytes, and produces resonances in this region of the spectrum.20 Davie et al showed that this resonance has a longer T2 than the methylene group at 1-3 ppm because it was more prominent at TEs of 20–30 ms.11

Follow up 1H-MRS showed changes in spectra from the previous one. The relevant differences were a remarkable decrease of lipid signal, the resonance at 1-4 ppm, and the increase in the myoinositol peak. There were no significant changes in N-acetyl aspartate and choline peaks. Some reports suggest that the time course of increased lipid resonance is compatible with the histologically determined time course of disappearance of lipid laden
Balo's concentric sclerosis: a clinical case study of brain MRI, biopsy, and proton magnetic resonance spectroscopic findings

Figure 2. (A) This first 1H-MRS (TR = 3 s, TE = 30 ms) shows large lipid signals at 0-9 and 1-3 ppm, a large resonance at 1-4 ppm (white arrow), a decrease in NAA:creatine ratio, and an increase in choline:creatine ratio. (B) The second 1H-MRS (TR = 3 s, TE = 30 ms), nine months after the initial symptoms shows changes such as a decrease in lipid signal, a decrease in the resonance at 1-4 ppm, and an increase in myoinositol peak. There are no significant changes in NAA and choline peaks. NAA = N-acetyl aspartate; ml = myoinositol; Cho = choline; Cr = creatine.

Macrophages from areas of acute myelin destruction. Myoinositol has been reported to be present only in glial cells and it was suggested that it may be used as a glial marker. This increase of myoinositol was attributed to the process of gliosis, suggesting the healing phase of demyelinating plaque. Increase of myoinositol in vivo have been seen in chronic as well as acute lesions. These characteristic 1H-MRS findings in our case, which are similar to those reported in multiple sclerosis, may be useful in understanding the pathogenesis of Balo's concentric sclerosis. Also, 1H-MRS may have additional diagnostic value in allowing brain biopsy to be avoided in the diagnosis of Balo's concentric sclerosis if 1H-MRS is combined with typical MRI findings.

1 Balo J. Encephalitis periaxialis concentrica. Archives of Neurology and Psychiatry 1928;19:242-64.
Balo’s concentric sclerosis: a clinical case study of brain MRI, biopsy, and proton magnetic resonance spectroscopic findings.
M O Kim, S A Lee, C G Choi, J R Huh and M C Lee

J Neurol Neurosurg Psychiatry 1997 62: 655-658
doi: 10.1136/jnnp.62.6.655

Updated information and services can be found at:
http://jnnp.bmj.com/content/62/6/655

Email alerting service

These include:
Receive free email alerts when new articles cite this article. Sign up in the box at the top right corner of the online article.

Notes

To request permissions go to:
http://group.bmj.com/group/rights-licensing/permissions

To order reprints go to:
http://journals.bmj.com/cgi/reprintform

To subscribe to BMJ go to:
http://group.bmj.com/subscribe/