Primary Cerebral Lymphoma Causing Remitting and Relapsing Neurological Symptoms

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Abstract

Primary CNS lymphoma is a rare variant of non-Hodgkin’s lymphoma. Incidence has increased over the past 3 decades, but the optimum treatment protocol is yet to be established. We report a 44-year old man who presented with left-sided numbness. Magnetic resonance imaging of the brain revealed a right hemisphere ill-defined mass. Within four months the patient experienced a spontaneous near-full recovery. From this time the patient deteriorated with increasing left hemiparesis. After several months, the patient developed lethargy and dyspnoea, with episodes of epistaxis and per rectal bleeding. Full blood count revealed a pancytopaenia. Trephine biopsy demonstrated evidence of marrow involvement by diffuse large B-cell lymphoma. The patient was diagnosed with primary CNS lymphoma with systemic involvement. We conclude that the spontaneous remission of symptoms should not discourage a diagnosis of PCNSL if consistent with clinical and radiological findings.

Keywords: Primary central nervous system lymphoma; Diffuse large B cell lymphoma; Non-Hodgkin’s lymphoma

Introduction

Primary central nervous system lymphoma (PCNSL) is a rare variant of non-Hodgkin’s lymphoma, accounting for 3-5% of brain tumours [1]. Immunodeficiency is the only well-documented risk factor. Diffuse large B cell lymphoma (DLBCL) accounts for the vast majority of cases of PCNSL in immunocompetent patients. Although the overall incidence remains low, there has been an increase in the number of cases over the past 3 decades, particularly in immunocompetent individuals [2]. Owing to the low incidence of PCNSL, it has been difficult to establish the optimum treatment protocol for this disease [3].

Case Report

A 44-year-old man with no significant past medical history presented in April 2009 to his general practitioner with numbness in his left arm, and subsequently his left leg, causing him to limp with no significant discomfort. A brain MRI scan revealed a right hemisphere small ill-defined mass in the posterior centrum ovale and extending inferiorly towards the basal ganglia (Fig. 1A), which was considered to be inflammatory.

Within four months the patient experienced a spontaneous near-full recovery, enabling him to perform heavy exercise. An MRI scan at that time demonstrated that the ring-enhancing lesion had increased in size and developed central cavitation (Fig. 1B). From this time the patient began to clinically deteriorate with increasing left hemiparesis. On examination he had marked left upper motor neuron facial weakness, and power was 3/5 and 4/5 in the left arm and leg respectively.

In November 2009, the patient suddenly deteriorated, with an abrupt loss of sensation in the left arm and impairment of coordination in the left leg. There was no headache, dizziness, visual or hearing impairment. Clinical examination revealed fixed-flexion of the left arm, and brisk reflexes in the left lower limb accompanied by an up-going plantar reflex. Cranial nerve examination was normal, except for the previously reported left-sided weakness of the face, which was now accompanied by a loss of sensation. A third MRI scan demonstrated further progression, with increased size of the mass (2 cm), cavitation, and irregular thick ring en-
At this time routine haematological examination was normal. Cerebrospinal fluid (CSF) examination was also within normal limits with an opening pressure of 12 cm (10 - 25 cm), 1.33 g/L protein (< 0.4g/L), 3.9 mmol/L glucose (2.8 - 4.4 mmol/L), normal cytology, negative bacterial culture and gram stain. No monoclonal or oligoclonal bands were detected and polymerase chain reaction for Toxoplasma, Cryptococcus, syphilis, hepatitis B and C viruses were negative. Anti-nuclear, anti-neutrophilic cytoplasmic, anti-mitochondrial, anti-smooth muscle, anti-liver-kidney microsomal antibodies were all negative. CSF angiotensin-

Figure 1. (A). Coronal FLAIR-MRI showing an ill-defined T2 hyperintense 1cm mass in the right posterior centrum semiovale extending inferiorly towards the basal ganglia. Surrounding vasogenic oedema extends throughout adjacent white matter. (B). Axial post contrast T1-weighted MRI showing greater definition of the same enhancing 1.4 cm mass in the right posterior centrum semiovale, now with some central cavitation. (C). Axial post contrast T1-weighted MRI showing further definition, increase in size (2 cm) and central cavitation of the mass in the right posterior centrum semiovale. There is irregular thick ring enhancement of the mass, but no significant mass effect. (D). Axial diffusion-weighted MRI showing restricted diffusion of the centrally cavitating mass in the right posterior centrum semiovale. (E). Axial post-contrast T1-weighted MRI showing poor definition and enhancement of the centrally cavitating mass in the right posterior centrum semiovale. (F). Axial T2-weighted MRI showing the extensive vasogenic oedema surrounding the hyperintense mass in the right posterior centrum semiovale.
remitting and relapsing neurological symptoms

several episodes of neutropenic sepsis, from one of which
teriorated to 0/5 throughout. Treatment was complicated by
dtential for treatment of PCNSL with bone marrow involve

Dexamethasone 6 mg TDS was initiated leading to a
ificant clinical improvement. Due to the development of
on of intolerable side effects, the steroid dosage was gradually
inished by 2 mg BD, and azathioprine therapy was initiated.
oved to adverse effects (hepatic dysfunction), azathioprine was discontinued after one week.

Three months later the patient was admitted with weak
ness in the left side of the face, left arm (particularly the fin
ers) and left leg. On examination, there was marked weak
ness and hypertonia, and reflexes were brisk throughout the
left side. Sensation was impaired in both the upper and lower
left limbs, particularly light touch and pin-prick. The MRI at
this stage revealed that the pre-existing lesion was of similar
size but had become less discrete, and with less contrast en
hancement but greater restricted diffusion (Fig. 1D).

In addition to the neurological features mentioned, the
atient also complained of lethargy and dyspnoea on mild
xertion together with episodes of epistaxis and per rectal
leeding. He did not report any weight loss or night sweats
nd there was no clinical evidence of lymphadenopathy. The
ull blood count demonstrated a pancytopenia with 8.4g/
L haemoglobin (13 - 18 g/dL), 105.3 fL mean corpuscular
olume (76 - 96 fL), 86 × 10⁹/L platelets (150 - 400 × 10⁹/L)
nd 4.8 × 10⁹/L leukocytes (4 - 11 × 10⁹/L).

In view of the patient’s deteriorating haematological
rofile bone marrow aspiration and trephine biopsy were per
formed. The bone marrow aspirate demonstrated abnormal
ononuclear cells, the morphology favouring a haematopoietic
rocess (for example, large cell lymphoma). There was
so evidence of hemophagocytosis and erythroid hyperplasia,
ccounting for the anaemia. There was a grossly abnormal
ncellular infiltrate raising the possibility of haematological
alignancy. Trephine biopsy revealed small clusters of
cells in keeping with marrow involvement by DLBCL.

The patient was initiated on rituximab, cyclophospham
ide, hydroxydaunorubicin, vincristine and prednisolone
otherapy for treatment of PCNSL with bone marrow involve
ment. Two further brain MRI scans in May and July 2010
howed poorer definition of the lesion, which had remained
constant in size, but with greater surrounding oedema (Fig.
1E, F). Unfortunately, the patient’s symptoms progressed to
clude visual impairment, and power on the left side de
teriorated to 0/5 throughout. Treatment was complicated by
veral episodes of neutropenic sepsis, from one of which
the patient died, two months later.

Discussion

This case demonstrates an unusual presentation of a rare,
but increasingly encountered condition in immunocompe
tent hosts [3]. Owing to the relative rarity of PCNSL, it has
proved difficult to study, and the optimum treatment strategy
has yet to be determined [3].

Clinical manifestations of PCNSL include neuropsychiatric signs, raised intracranial pressure, seizures, ocular
symptoms and other focal deficits [1, 4]. The so-called, ‘B
symptoms’ include weight loss, fever and night sweats, and
are rarely encountered at initial presentation [4].

Neuroimaging is important in the diagnosis of PCNSL,
with contrast-enhanced MRI being the modality of choice
[4]. Lesions may be multiple, and may resemble inflammatory and infectious diseases, as well as other neoplastic
processes [1]. PCNSL masses have acquired the pseudonym
of “ghost tumours”, as radiological changes may disappear,
particularly after corticosteroid therapy [1]. Definitive di
agnosis is achieved by stereotactic biopsy, although this is
sometimes unnecessary owing to the presence of lymphomato
us cells in the CSF [1]. Glucocorticoid therapy may obscure
the histological diagnosis, so steroids should not be adminis
tered (pre-biopsy) if PCNSL is likely [5].

The development of effective chemotherapy regimens
has meant that a proportion of patients may achieve remis
sion. Relapse however, is a common occurrence, and only
20-30% of patients are cured [1]. The spontaneous remis
sion of symptoms that occurred initially in this case is an
extremely unusual occurrence, and may impose a significant
diagnostic challenge in cases with PCNSL. Consequently,
this patient’s symptoms were initially considered to be due
to an inflammatory or demyelinating process, culminating in
a delay in the initiation of appropriate cytotoxic therapy.

The diagnosis of PCNSL in the case described was not
established until symptoms of bone marrow involvement
had developed. The incidence of systemic involvement
following PCNSL is controversial, with some studies sug
gesting that extensive staging is unnecessary [1]. However,
investigation for occult systemic disease is increasingly un
dertaken, with full body CT scanning and bone marrow bi
opsy now recommended for staging of PCNSL [3]. Testicu
lar ultrasonography may be performed in elderly males, as
testicular lymphoma frequently disseminates to the brain [3].

Conclusion

The incidence of PCNSL in immunocompetent hosts is in
creasing. Spontaneous remission of symptoms is unusual,
but can occur, and should not discourage a differential di
agnosis of PCNSL if consistent with clinical and radiologi
cal findings. Extra-neural disease is increasingly recognised
in PCNSL, and patients should undergo thorough staging to
exclude systemic disease.
Declaration

The authors declare no competing interest.

References