A 50-year-old man presented acutely to the hospital with behavioural disturbance, choreiform movements and profound nihilistic delusions. He reported recent drug and alcohol abuse, and also apparent involvement in several recent criminal activities, for which he felt he should be punished. He arrived alone at the hospital after a concerned neighbour had called an ambulance. His initial level of agitation prevented formal cognitive testing. However, he was alert, verbally responsive and could obey commands. He was afebrile with normal observations and normal plasma glucose. Although his examination was challenging, the only abnormal neurological findings were bilateral choreiform upper limb movements.

**Question 1**

What should you do now?

**Comments**

Obtaining a collateral history is essential in establishing baseline function in people presenting with neuropsychiatric disturbance. This can help us focus on the differential diagnoses and plan for investigation. It can also avoid repeating previous investigations.

The admitting team contacted his brother, his next of kin. The patient was an unemployed ex-builder. One year earlier, he had consulted a movement disorder doctor with a high-frequency tremor and brief involuntary movements. Investigations at this time included a normal MR scan of the brain and routine blood tests, including thyroid function tests and serum calcium. In the 2 months before admission, his family had noticed occasional repetitive conversation. However, he still lived independently and attended training courses. His agitation and disorientation had developed over the 24 hours before presentation. There was no family history of dementia or of involuntary movements.

**Question 2**

What is the differential diagnosis?

**Question 3**

What initial investigations would you plan?

**Causes of cognitive impairment and chorea (Box 1).**

Urine screening for illicit drugs was negative. The following were normal: blood film, renal function, serum calcium, thyroid function tests and serum C-reactive protein; serum antistreptolysin O titres were negative. Urinalysis was normal. MR scan of the brain, including diffusion-weighted and susceptibility sequences, showed only minor non-specific white matter abnormalities. Cerebral MR angiography was also normal.

**Comments**

Before lumbar puncture, he was noted to have a prolonged activated partial thromboplastin time but normal prothrombin time. This did not correct with vitamin K, prompting his medical team to look for the lupus anticoagulant and for serum antinuclear antibodies. His lumbar puncture showed $20 \times 10^6$ red cells/l (0) and $10 \times 10^6$ white cells/l ($\leq 5$) (lymphocytes) with normal cerebrospinal fluid (CSF) protein and glucose concentrations. Oligoclonal band testing was equivocal.
His serum was positive for antinuclear antibody (1/640), anti-double-stranded DNA (ELISA) at 840.4 U/mL (<35) and positive anti-DNA immunophoresis. He had a prolonged activated partial thromboplastin time and positive testing for the lupus anticoagulant (Dilute Russell’s viper venom time (DRWT)). He had positive anticardiolipin IgG 16.9 U/mL (normal <10), but normal IgM levels. His serum complement C3 levels were initially low at 0.71 g/l (normal >0.8), with normal C4 levels.

**Question 4**
What is the most likely diagnosis?

**Question 5**
How would you initially treat him?

**Comments**
We made the presumptive diagnosis of diffuse neuropsychiatric lupus. He was originally treated with 5 days of intravenous methylprednisolone, with clear initial improvement in his agitation and lessening of his involuntary movements. He scored 54/100 on the revised Addenbrooke’s Cognitive Assessment (ACE-R). The rheumatology team started him on tapering oral corticosteroids, followed by intravenous cyclophosphamide 500 mg weekly for 6 weeks. He did not improve and, if anything, deteriorated further with more intrusive depressive thoughts, worsening cognitive function and a return of the chorea. He began complaining of visual hallucinations of a dog in his room.

**Question 6**
How would you manage him now?

**Question 7**
Are there any other treatment options and how might you monitor their impact?

**Comments**
Given his emerging depressive psychotic symptoms, we arranged for psychiatric evaluation. He was treated with venlafaxine and then additionally quetiapine, with a gradual impact on his depressive symptoms and resolution of the hallucinations. At this stage, he had five plasma exchange treatments without clear effect. An hexamethylpropyleneamine oxime (HMPOA) single photon emission CT (SPECT) showed bilateral cortical hypoperfusion, most prominent over the left temporal lobe (figure 1). Given his static and severely impaired cognitive function, we started him on intravenous rituximab. He received four weekly doses of 600 mg, having been screened for relevant infections (HIV, hepatitis B and C, and tuberculosis). His cognitive function significantly improved, the ACE-R score rising to 80/100. Repeat HMPOA SPECT showed increased perfusion (figure 1). During this time, he also received low-molecular-weight heparin for 6 weeks, given the associated antiphospholipid antibodies although vascular imaging never showed evidence of thrombosis. He now takes hydroxychloroquine as maintenance immunosuppressive therapy along with the venlafaxine and quetiapine. He manages self-care, but 12 months after initial presentation he still cannot yet live independently, requiring some supervision. His depression has resolved and his personality has returned to normal.

**DISCUSSION**
Most neuropsychiatric lupus events occur at onset or within an year of onset (50%–60%). The American College of Rheumatology criteria for classifying neuropsychiatric lupus is very broad and includes 19 potential neurological and psychiatric syndromes (table 1), including both central and peripheral
syndromes. The spectrum of events varies from headache, mood disorders and mild cognitive impairment to severe cases. Severe cognitive cases such as this comprise only 3%–9% of neuropsychiatric lupus. Chorea is the most common movement disorder and is associated with antiphospholipid antibodies, but occurs in less than 4% of cases. Severe neuropsychiatric cases have either focal or diffuse presentations. The focal cases have ischaemic, thrombotic or demyelinating lesions. Our patient had a diffuse presentation, similar to several previously described cases. However, these diffuse cases were variably described as having severe neuropsychiatric lupus or acute confusional state presentations, making it difficult to compare the reported cases. These diffuse cases are most likely immune mediated, possibly with several autoantibodies, including antiribosomal-P, anti-DNA, antiphospholipid, NMDA NR2 and GABA-B antibodies, supporting this assertion.

Patients with suspected neuropsychiatric lupus should have an MR scan of brain to exclude several differential diagnoses, including demyelination, ischaemia or thrombosis. Over half of neuropsychiatric lupus cases have normal MR brain scans. Non-specific white matter abnormalities are common in systemic lupus erythematosus and in neuropsychiatric lupus cases, and are non-discriminatory. Serial MR brain scans in diffuse neuropsychiatric lupus cases may show rapidly progressive generalised brain atrophy. Lumbar puncture is important to exclude central nervous system (CNS) infection; it can show an inflammatory response but can be normal. Some patients have positive CSF oligoclonal bands and other CSF immune biomarkers, such as anti-DNA antibodies, interleukin-6 and tumour necrosis factor alpha.

Several studies have shown altered cerebral metabolism in neuropsychiatric lupus. Thus, SPECT scans may be diagnostically useful in patients with normal MRI, especially those with a diffuse cognitive presentation (where it is 75%–93% sensitive). SPECT scans may be used as a biomarker for the immediate response to immunotherapy, but do not help in understanding the diffuse involvement of the brain.

**Table 1** American College of Rheumatology Neuropsychiatric Systemic Lupus Erythematosus classification (1999)

<table>
<thead>
<tr>
<th>Central nervous system</th>
<th>Peripheral nervous system</th>
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<tbody>
<tr>
<td>Aseptic meningitis</td>
<td>Acute inflammatory</td>
</tr>
<tr>
<td></td>
<td>demyelinating polyradiculopathy</td>
</tr>
<tr>
<td>Cerebrovascular disease</td>
<td>Autonomic disorder</td>
</tr>
<tr>
<td>Demyelinating syndrome</td>
<td>Mononeuropathy (single/multiple)</td>
</tr>
<tr>
<td>Headache</td>
<td>Myasthenia gravis</td>
</tr>
<tr>
<td>Movement disorder (chorea)*</td>
<td>Cranial neuropathy</td>
</tr>
<tr>
<td>Myelopathy</td>
<td>Plexopathy</td>
</tr>
<tr>
<td>Seizure disorder*</td>
<td>Polyneuropathy</td>
</tr>
<tr>
<td>Acute confusional state*</td>
<td></td>
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<tr>
<td>Anxiety disorder</td>
<td></td>
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<tr>
<td>Cognitive dysfunction*</td>
<td></td>
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<td>Mood disorder*</td>
<td></td>
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<tr>
<td>Psychosis*</td>
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</table>

*Presentations associated with diffuse neuropsychiatric lupus.
subsequent monitoring or predicting further neuropsychiatric events.13

Patients with psychiatric symptoms should receive antidepressant and antipsychotic agents.1 Electroconvulsive therapy was effective in three cases with severe psychosis.16 Patients with cognitive decline often respond to glucocorticoids and other immunosuppressants, although corticosteroids can aggravate the psychiatric symptoms. There is evidence for using cyclophosphamide and plasma exchange (sometimes synchronised) in refractory cases5 17 and for rituximab in severe refractory cases.9 13 14 18 Patients with either a long disease duration or with more than one of the American College of Rheumatology syndromes (table 1)13 have a poorer therapeutic prognosis.9

Although several CSF cytokine, chemokine and growth factors have been used as markers of the inflammatory process, a more specific biomarker would greatly improve diagnostic and therapeutic research into this rare condition.9

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REFERENCES
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