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# When the penny drops

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A 26-year-old woman presented after developing slurred speech and drooling overnight. She had a 6-month history of tiredness and poor concentration, and episodes of loss of consciousness accompanied by paraesthesia of her hands, feet and mouth. Occasionally, one arm would 'hang in a strange way'.

Her mother reported a preceding personality change with angry outbursts and anxiety, although these had largely subsided after starting antidepressants. Her only past history was a severe depressive episode in her teens and a difficult and occasionally abusive family life. There was no family history of neurological disease.

On examination, she was alert and orientated. There was mild dysarthria but normal motor, sensory and coordination examination.

## Question 1

What is the likely differential diagnosis?

## Comment

The pattern of neurological features does not immediately suggest any single lesion and, with no objective examination findings, might easily be construed as functional or psychosomatic. The initial clinical suspicion was of a psychiatric presentation, given her personal history; however, her mood had recently been more stable and there was no obvious psychosocial precipitant for this acute presentation.

The possibilities of reflex (vasovagal) syncope or hyperventilation syndrome were suggested to account for the loss of consciousness with facial paraesthesia. Vasovagal syncope seemed unlikely with no presyncopal symptoms or triggers, such as prolonged standing, dehydration or stress. Hyperventilation syndrome was also unlikely, as forced hyperventilation did not reproduce the symptoms.

On presentation, the patient herself thought she was having a stroke. Despite the acute onset and nature of her symptoms, her age, examination findings and lack of risk factors made this unlikely. Stroke mimics can occur in young people and form part of the differential diagnoses. These include space-occupying lesions, migraine, focal seizures, multiple sclerosis and periodic paralysis.

## Question 2

What would you do next?

## Comment

Routine blood tests showed only mild thrombocytopenia. There was no postural hypotension. She continued taking antidepressants and underwent cognitive behavioural therapy, which helped in stabilising her mood.

After she developed widespread arthralgia and dystonic tremor of the left hand, as well as persistent fatigue, slurred speech and episodes of loss of consciousness, her medical team requested a specialist neurological opinion.

The neurologist noted the above findings in addition to horizontal diplopia, worse in primary position, and arranged repeat bloods with 12-lead ECG, EEG and MR scan of brain (table 1).

## PROGRESSION

Over the next 18 months, she reported more prominent diplopia with occasional blurred vision, as well as frequent dystonic posturing, particularly of the left hand. She suffered extreme fatigue with variable confusion and irritability, exacerbated by chronic headaches. In addition, she developed widespread arthralgia. There was a marked decline in her handwriting, which became progressively smaller and more illegible.

On examination, she had a markedly ataxic gait and worsening coordination.



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## TEST YOURSELF

**Table 1** Investigation results

Investigation	Result
Full blood count	Normal, except thrombocytopenia
Liver function tests	Normal
Urea and electrolytes	Normal
Copper studies	Normal serum copper and caeruloplasmin Non-caeruloplasmin-associated copper 17 µg/dL (raised; target <10)
Serum creatine kinase	Normal
Amino acid analysis	Slightly raised tyrosine, indicative of possible liver involvement
12-Lead ECG	Normal
EEG	Normal
MR scan of brain	Altered signal in basal ganglia, pons and dorsal midbrain

Upper limb examination showed cogwheeling reinforced by synkinesis, and mild dysdiadochokinesis. Reflexes and muscle bulk remained largely intact. A speech and language therapist felt that her fluctuating slurred speech with mild word-finding problems was a 'Parkinsonian-like dysarthria'; there was also dysfunctional coordination of swallowing. The patient reported her speech and swallowing worsened with tiredness or anxiety. Her depression deepened, with frequent mood swings and agitation.

**Question 3**

How might these developments affect the differential diagnoses and what further investigations may help?

**Comment**

The widespread symptoms and their protracted onset might suggest a genetic, inflammatory or degenerative condition. We explored several differential diagnoses.

- ▶ **Multiple sclerosis.** Her age, sex and dissemination of neurological symptoms in space and time indicate the need to consider multiple sclerosis in the differential diagnosis; however, psychiatric symptoms are unusual as presenting features and arthralgia is not a feature.
- ▶ **Wilson's disease.** The combination of neurological symptoms with arthralgia might suggest Wilson's disease, although the onset is relatively late and the fluctuating dystonic posturing would be atypical. In addition, there was no relevant family history and copper studies and liver function tests were both reported as normal.
- ▶ **Juvenile Parkinson's disease.** This was highlighted given the Parkinsonian-like dysarthria and the later development of micrographia and cogwheeling with synkinesis. However, there was no resting tremor or other physical features of Parkinsonism.
- ▶ **Mitochondrial disorder, for example, MELAS, adult-onset Leigh's syndrome.** These are characterised by maternal inheritance, poor growth, myopathy and learning difficulties. An example we considered was MELAS, an umbrella

disorder comprising mitochondrial myopathy, encephalomyopathy, lactic acidosis and stroke-like symptoms. Leigh's syndrome, a subacute sclerosing encephalopathy, usually presents in infancy but clinical manifestations depend on the site of neuropathological involvement. It may manifest initially in adults as a movement disorder, including multifocal dystonia and pyramidal weakness, irritability, fatigability and cognitive decline.

- ▶ **Variant Creutzfeldt-Jakob disease (vCJD).** This presentation would be atypical for vCJD, which produces myoclonic jerks and rapidly-progressive dementia over 12–14 months. Nevertheless, dysphasia, anxiety, depression, seizures and ataxia are prominent features, with psychiatric symptoms often preceding neurological ones.
- ▶ **Postencephalitic Parkinsonism (Von Economo's disease).** This is rare diagnosis; outbreaks followed influenza pandemics, the last being in the early 20th century. In these cases, viral encephalitis probably triggers degeneration of the substantia nigra, causing Parkinsonism. There are occasional sporadic cases.
- ▶ **Myeloma.** Although classically affecting older people, up to 2% of cases occur between 20 and 40 years. The symptoms vary but those suggesting myeloma in this case include fatigue and weakness (hypercalcaemia), headache and visual changes (hyperviscosity), and paraesthesia (amyloid infiltration of peripheral nerves).
- ▶ **Autoimmune encephalitides.** Several recently described autoimmune and paraneoplastic disorders, particularly anti-NMDA receptor encephalitis, have features in common with our patient. This presents with neuropsychiatric symptoms, dyskinesia, seizures and dysautonomia. Anti-AMPA receptor encephalitis typically presents with limbic encephalitis or with isolated psychiatric symptoms. Antiglycine disease can cause ataxia, often with prominent stiffness and other features, and mGluR1 disease may also show ataxia. These conditions are important to consider early as outcome is better with early treatment and many are associated with underlying neoplasms.<sup>1</sup>

Five years after her initial presentation, there was still no definitive diagnosis. We performed some further investigations (table 2).

Three years after her initial presentation, during which time her vision had continued to deteriorate, an optician documented 'yellow-brown deposits circumferentially in the inner layer of the peripheral cornea'.

**Question 4**

What is the diagnosis and how would you confirm it?

**Comment**

The diagnosis is Wilson's disease. The corneal deposits represent Kayser-Fleischer rings which, although no longer considered pathognomic of Wilson's

**Table 2** Further investigations

Investigation	Result
B <sub>12</sub> and folate	Normal
Leucocyte and plasma enzymes	
Serum antinuclear antibody	
Serum protein electrophoresis	
Urine porphyrin screen	
Urine organic acids	
Cerebrospinal fluid	No oligoclonal bands
Evoked potentials	Bilateral delay of tibial sensory evoked potentials, particularly on the left. Visual evoked potentials were normal. Right brainstem auditory evoked potentials were mildly delayed
Repeat MR scan of brain	Axial MR scan of brain (see <a href="#">figure 1</a> ) showed signal change in the brainstem and cerebellar white matter with bilateral symmetrical high signal in both thalami. T2-weighted low signal in the globi pallidi and cerebral peduncle, consistent with mineralisation. Also, symmetrical signal change in the lentiform nuclei, pons, cerebellar hemispheres, dentate nuclei with some cystic component and infratentorial atrophy. Appearance favours Wilson's disease or Leigh's syndrome
Muscle biopsy	Non-specific fat storage abnormality
Amino acid levels	Raised tyrosine (may indicate liver involvement)

disease, classically occur in affected patients. They are especially prevalent (up to 95%) in those with neurological manifestations. She was referred to a specialist centre where repeat caeruloplasmin, serum and urinary copper, and a D-penicillamine challenge test confirmed the diagnosis. Subsequent DNA analysis showed a homozygous H1069Q mutation in the *ATB7B* Wilson's gene.

Despite her relatively late and atypical presentation, clinicians should consider rigorous exclusion of Wilson's disease in all young people with the combination of a movement disorder, psychiatric disturbance and arthralgia, because the condition is so treatable. Early treatment is essential before there is irreversible brain damage through copper deposition.

In their seminal series of 136 patients with neurological signs of Wilson's disease, Walshe and Yealland<sup>2</sup> found that the age at onset of neurological features was 9–40 years (median 16). Overall, 23% of patients had already an episode of liver damage as early as 3 years of age. In a further 114 cases, the diagnosis was already made before the onset of neurological symptoms. They divided the neurological group into four predominant clinical pictures according to the nature of the movement disorder: 45% were Parkinsonian, 24% 'pseudosclerotic', 15% dystonic and 11% choreic. Dysarthria, tremor, personality change and drooling were common initial symptoms. All patients with neurological disorders had corneal pigmentation, although not necessarily complete rings.

In our case, other pointers to the diagnosis included the initial serum copper level, which, although deemed normal at the time, was at the lower limit of normal. The serum caeruloplasmin level was within limits but had been measured using an immunological assay, as performed by many UK laboratories. Importantly, there is growing evidence to suggest that

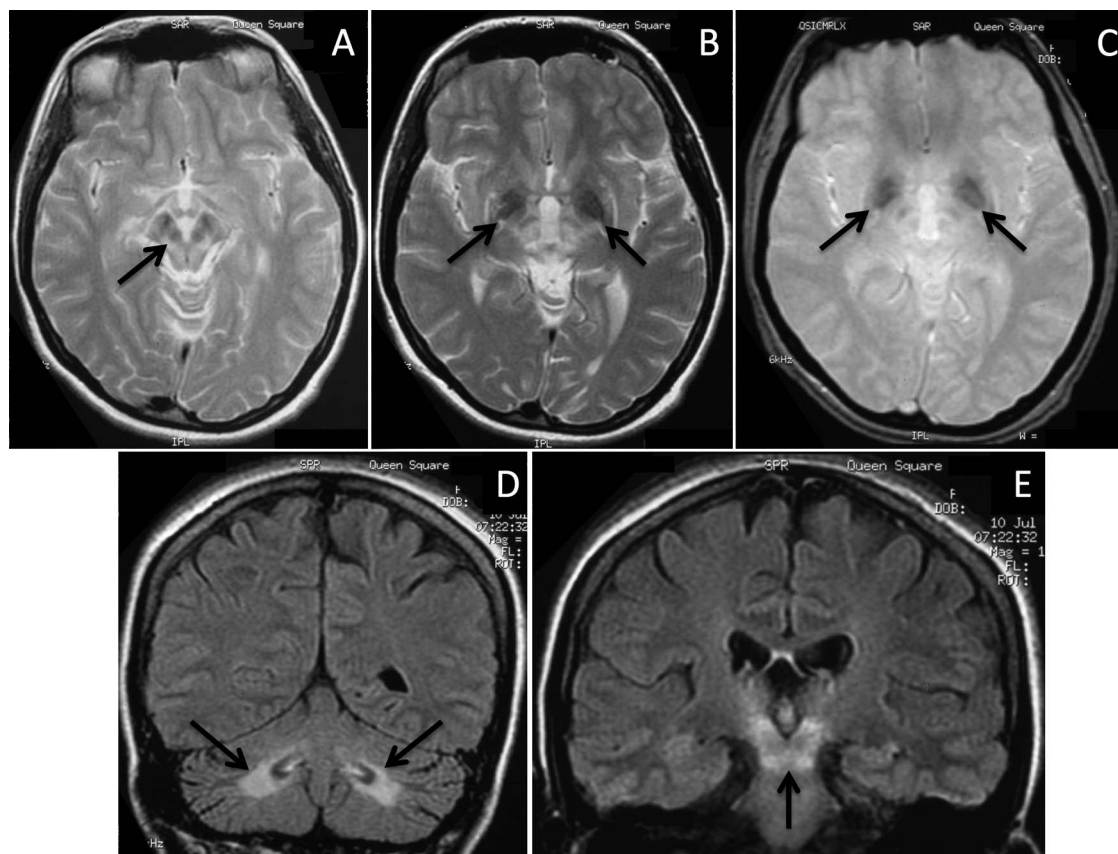
immunological assays overestimate caeruloplasmin levels, and that enzymatic assays may be preferable.<sup>3</sup>

There was also thrombocytopenia in the initial bloods. Although non-specific, there has long been a recognised minor association between Wilson's disease and thrombocytopenia.<sup>4</sup> This patient may be considered as having a late presentation, being diagnosed at age 31 years, although the literature generally regards onset of Wilson's disease after the age of 40 years as being of 'late onset'.

MR brain scan findings in Wilson's disease are widespread and variable, involving most structures in varying combinations but most commonly involving the basal ganglia, midbrain, pons and cerebellum. Most typically, there is symmetrically increased signal in the putamen, globus pallidus, thalamus, caudate head, dorsal midbrain and pons. There can also be more widespread involvement, including of the cortical white matter with diffuse brain atrophy and ventricular enlargement, but rarely the medulla.<sup>5</sup> Signal change together in the basal ganglia, thalamus and brainstem strongly suggests Wilson's disease. The so-called 'face of the giant panda and her cub' and 'double panda' signs are rare, but very typical of Wilson's disease.<sup>6</sup>

Skeletal changes on plain X-ray are common but not diagnostic. They include evidence of osteomalacia, osteochondritis, osteoarthritis and bone fragmentation, particularly of the lumbar and cervical spine, hands, wrists, elbows, knees and feet.

The difficulty in diagnosing this case arose from the atypical presentation and from lack of clarity regarding the exclusion of Wilson's disease. Copper studies are commonly used in clinical practice to deem a diagnosis of Wilson's disease unlikely, as done here ([table 3](#)).<sup>7</sup> Unfortunately, this practice may lead to a missed diagnosis. Although rapidly and cheaply obtained in most hospitals, the sensitivity of these



**Figure 1** . MR scan of brain (1.5 Tesla), patient aged 30 years. (A) Axial T2-weighted sequences at midbrain level show disproportionate volume loss and signal hyperintensity surrounding the red nucleus and substantia nigra, simulating the ‘giant panda’ sign (arrow). (B) Axial T2-weighted sequences at corpus striatum level show hypointensity of the globi pallidi and the putamen shows volume loss and a hyperintense rim (arrows). (C) Axial T2\*-weighted gradient-echo sequence confirms mineralisation of the globi pallidi (arrows). (D and E) Coronal FLAIR sequences through the dentate nuclei and midbrain show hyperintense signal change within the cerebellar white matter and surrounding the mineralised dentate nuclei (arrows in panel D) and midbrain (arrow in panel E).

tests for Wilson’s disease may be inadequate. A recent study identified the sensitivities for detecting Wilson’s disease were 85% for serum copper and 83% for

serum caeruloplasmin. Thus, using these tests alone leaves a significant proportion of people with undetected disease.<sup>8</sup>

**Table 3** Routine tests for diagnosing Wilson’s disease

Test	Typical finding	False ‘negative’	False ‘positive’
Serum caeruloplasmin	Decreased by 50% of lower normal value	Normal levels in patients with marked hepatic inflammation Overestimation by immunological assay, pregnancy and oestrogen therapy	Low levels in: ► malabsorption ► acaeruloplasminaemia ► heterozygotes
24-h urinary copper	>1.6 µmol/24 h in adults>0.64 µmol/24 h in children	Normal levels if: ► incorrect collection ► children without liver disease	Increased levels if: ► hepatocellular necrosis ► cholestasis ► contamination
Serum ‘free’ copper	>1.6 µmol/L	Normal if caeruloplasmin overestimated by immunological assay	
Hepatic copper	>4 µmol/g dry weight	Due to regional variation in: ► patients with active liver disease ► patients with regenerative nodules	Cholestatic syndromes
Kayser–Fleischer rings by slit-lamp examination	Present	Absent ► in up to 50% of patients with hepatic Wilson’s disease ► in most asymptomatic siblings	Primary biliary cirrhosis

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**Table 4** Scoring system for diagnosis of Wilson's disease

Typical clinical symptoms and signs		Other tests	
Kayser–Fleischer rings		Liver copper (in the absence of cholestasis)	2
Present	2	>5×upper limit of normal (>4 µmol/g)	1
Absent	0	0.8–4.0 µmol/g	–
		Normal (<0.8 µmol/g)	1
		Rhodanine-positive granules*	1
Neurological symptoms†		Urinary copper (in the absence of acute hepatitis)	
Severe	2	Normal	0
Mild	1	1–2×upper limit of normal	1
Absent	0	>2×upper limit of normal	2
		Normal but >5×upper limit of normal after D-penicillamine	2
Serum caeruloplasmin		Mutation analysis	
Normal (>0.2 g/L)	0	On both chromosomes detected	4
0.1–0.2 g/L	1	On one chromosome detected	1
<0.1 g/L	2	No mutations detected	0
Coombs'–negative haemolytic anaemia			
Present	1		
Absent	0		
Total score		Evaluation:	
4 or more		Diagnosis established	
3		Diagnosis possible, more tests needed	
2 or fewer		Diagnosis very unlikely	

\*If no quantified liver copper available.

†Typical abnormalities at MR scan of brain.

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Various tests may support a diagnosis of Wilson's disease, including 24-h urinary copper (53% sensitivity in the above study), D-penicillamine challenge, DNA analysis and hepatic parenchymal copper content by liver biopsy. There are over 300 reported mutations in the *ATP7B* gene; most patients are compound heterozygotes. Gene sequencing is commonly used in populations with well-defined mutations, to test family members where the mutation is already known or in cases where conventional tests have not led to a clear diagnosis.<sup>9</sup> A recent study of 181 clinically and biochemically confirmed cases of Wilson's disease from the UK identified mutations in the *ATP7B* gene in 98%. The commonest mutation was H1069Q (seen in our patient) in 19%, a mutation known to be common in those of European descent.<sup>10</sup>

Recent practice has moved away from reliance on individual investigations and towards scoring systems based on clinical and laboratory findings.<sup>11</sup> Table 4 outlines one widely used example of these, developed at the 8th international meeting on Wilson's disease.<sup>7</sup>

### Question 5

How would you manage this patient?

### Comment

Several drugs are commonly used to treat Wilson's disease. These are maintained lifelong once the

diagnosis is established, although there are few high-quality trials supporting these medications.

D-penicillamine and its newer alternative trientine each acts by chelating copper and increasing urinary copper excretion. Patients should receive 25–50 mg of supplemental pyridoxine along with D-penicillamine

### Practice points

- ▶ Neuropsychiatric presentations with arthralgia or movement disorder should suggest the possibility of Wilson's disease.
- ▶ Clinicians should have a low threshold for testing for Wilson's disease due to its potential treatability and better outcomes with early treatment.
- ▶ Serum copper and caeruloplasmin may be within normal ranges, particularly early on, and so they do not completely exclude the diagnosis; enzymatic caeruloplasmin assays are better than immunological tests.
- ▶ We advise involving an ophthalmologist early to assess for Kayser–Fleischer rings with a slit lamp, and seeking advice from a liver specialist.
- ▶ Laboratory results correlate with the clinical symptoms; if there is doubt, then consider a 24-h urinary copper, liver function tests and, if necessary, liver biopsy or genetic testing.
- ▶ Scoring systems are a more reliable tool than individual tests in diagnosing Wilson's disease.

to minimise its interference with pyridoxine action. Clinical improvement may occur after 2–6 months in those with symptomatic liver disease and after up to 3 years in patients with neurological features. In all, 10%–50% of patients experience worsening of neurological symptoms when starting D-penicillamine treatment. Other side effects include nephrotoxicity, bone marrow suppression and a lupus-like syndrome.

Trientene, developed for those intolerant to D-penicillamine, is also an effective initial treatment with fewer adverse effects. However, because it is an iron chelator, patients must avoid co-prescription of iron. A 24-h urinary copper excretion can assess the effectiveness of both D-penicillamine and trientine.

Ammonium tetrathiomolybdate, zinc and various antioxidants may potentially help in Wilson's disease, but there is sparse evidence to support them. People with advanced disease may need liver transplantation because of severe cirrhosis or hepatic failure.

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