A woman with forgetfulness and falls

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A 69 year old woman presented to her general practitioner with a six month history of occasional falls and fluctuating forgetfulness and attention. Although she reported no difficulties with names and dates she needed help with taking drugs and preparing meals. She had also had two episodes of apparent visual hallucinations of a woman standing at the foot of her bed. Her sleep behaviour had been poor for many years, with frequent strong physical jerks and motion while sleeping. She had no symptoms of altered or low mood. Her medical history included hypothyroidism, osteoporosis, and cholesteatoma, and she was being investigated for a mixed fibre peripheral sensory neuropathy of unknown cause. Current drugs included levothyroxine, calcitriol, calcium carbonate-colecalficferol, and lansoprazole. She did not drink alcohol and was a non-smoker.

On examination she was fully orientated in time and place and her AMTS (abbreviated mental test score) was 10/10 with an MMSE (mini-mental state examination) score of 26/30. Her blood pressure, temperature, and cardiovascular and respiratory examinations were normal. Cranial nerve examination was normal with no primitive reflexes or supranuclear gaze palsy. Tone and power were normal throughout all limbs, as were sensation and reflexes in the upper limbs. The lower limbs showed reduced vibration and pin prick sensation to the mid-shin bilaterally, and joint position sense was limited to large movements. She showed no evidence of bradykinesia or apraxia, but her gait was ataxic in keeping with her peripheral sensory impairment.

Questions
1 What is the differential diagnosis?
2 What investigations would you do?
3 How should this patient be managed?

Answers
1 What is the differential diagnosis?

Long answer
To differentiate between delirium or dementia as causes of our patient’s symptoms, it is important to clarify whether the onset of symptoms was gradual, acute, or fluctuating. These two syndromes can also be distinguished between on the basis of changes in consciousness, attention, and psychomotor activity, all of which tend to be disturbed or fluctuating in delirium but minimally affected in dementia, although they are common in dementia with Lewy bodies. The simple CAM tool (confusion assessment method) can be used to screen for delirium and is easily administered within a general practice consultation, although it also needs a supporting history. The tool identifies an acute onset and fluctuating course, inattention, disorganised thinking, and an altered level of consciousness. A diagnosis of delirium requires the first two criteria and either the third or fourth. The diagnosis is complicated by the strongest risk factor for delirium being dementia, with a fivefold increase in risk of delirium in patients with dementia.

Common infective causes of delirium in older people include pneumonia and urinary tract infection, which can present atypically in older people. Fluctuating cognitive effects can be caused by hypoglycaemia as a result of poorly controlled diabetes or insulinomas, although these tumours are relatively rare. Hyperparathyroidism can also present with a mixed picture of fluctuating symptoms associated with diurnal variation but on a background of gradual decline. Common drugs that can cause delirium include anticholinergic drugs, anticonvulsants, antidepressants, antipsychotics, antiparkinsonian agents, opioid analgesics, and sedatives, in addition to multipharmacy.
The most likely cause of this patient’s protracted symptoms of forgetfulness is a dementia syndrome. Dementia is characterised by a progressive decline that causes memory loss and impairment in at least one other cognitive function, such as recognition, language, speech, or higher executive functions that affect activities of daily living. The prevalence of the various subtypes of dementia is unclear, but nearly two thirds (62%) of patients with dementia in the United Kingdom are thought to have Alzheimer’s disease and around a quarter (27%) to have vascular dementia or mixed Alzheimer’s disease and vascular dementia pathology. Most of the remainder (around 8%) have dementia with Lewy bodies, frontotemporal, Parkinson’s disease dementia, and alcoholic dementia in decreasing frequency.

Our patient’s symptoms were most typical of dementia with Lewy bodies—her cognitive symptoms fluctuated, often on an hourly basis, and included visual hallucinations. She also had supporting features of frequent falls and rapid eye movement sleep behaviour disorder (characterised by complex motor activity during rapid eye movement sleep). Dementia with Lewy bodies—first described in 1990—has symptoms similar to both Alzheimer’s disease and Parkinson’s disease, but the parkinsonian motor features present after the cognitive symptoms. Dementia with Lewy bodies, idiopathic Parkinson’s disease, and Parkinson’s disease dementia are different but overlapping phenotypes, and it is unclear whether they are distinct but convergent clinical syndromes or divergent common neurobiological pathologies. Memory impairment occurs later in dementia with Lewy bodies and is less pronounced than in Alzheimer’s dementia, although both dementia subtypes seem to have a similar rate of progression. It is important to distinguish between these two types of dementia because the clinical progression can be different and patients with dementia with Lewy bodies are exquisitely sensitive to neuroleptic drugs, with 50-80% having side effects such as confusion, sedation, rigidity, and most importantly increased mortality.

### 2 What investigations would you do?

#### Short answer

No simple blood test is available to screen for dementia. Blood tests are important in helping to rule out delirium but are also used in the investigation of dementia—to look for reversible causes of dementia—and specialist memory clinics should undertake full neuropsychological profiling along with appropriate brain imaging.

#### Long answer

Investigations appropriate to primary care should include a screen for risk factors such as diabetes and hypercholesterolaemia that might contribute to a vascular component of dementia. Commonly requested blood tests include a full blood count; renal and liver profile; thyroid function; and concentrations of calcium, vitamin B₁₂, folate, fasting glucose, and cholesterol. Urinalysis should be performed to screen for infection as a cause of delirium. Electrocardiography and chest radiography are also advised by the UK guidelines. In addition, if the history or physical examination point to specific risk factors, request HIV, syphilis, and *Borrelia* serology blood tests.

Several cognitive screening tests are available for use in primary care. The most well known is the MMSE. However, this test is difficult to administer in the short time allowed for general practice consultations and it may show cultural and language bias. Newer screening tests have been developed and validated for use in the community setting. They have been shown to be as robust as the MMSE but can be administered in five minutes or less; these tests include the general practitioner assessment of cognition (GPCOG), mini-cog, and memory impairment screen, with sensitivity varying from 69% to 80% and specificity from 86% to 96%.

Another screening tool, the 6 item cognitive impairment test (6CIT) has also been shown to be at least equivocal to the MMSE in selected secondary care populations. GPCOG (www.gpcog.com.au/prep.php) and 6CIT (www.falklandsurgery.co.uk/6cit/index.asp) are available electronically for ease of administration in the surgery. Most cognitive screening tools are developed for typical symptoms of Alzheimer’s disease, and patients with dementia with Lewy bodies may fall within the normal range. These screening tests are used to help decide on the need for referral to a specialist memory clinic for more thorough cognitive and neuropsychological assessment, including structural neuroimaging. When a diagnosis of dementia is suspected, this possibility must be discussed candidly and sensitively with the patient and carer to gauge their expectations and wishes before making a referral to specialist services.

The battery of specific tests included in neuropsychological testing varies by clinic and also depends on the type of dementia and cognitive and functional disturbances that are suspected. Patients with dementia with Lewy bodies usually show prominent early impairment in attention, visuospatial skills, and visuoconstructive skills—as detected by clock drawing and figure copying tests—whereas those with Alzheimer’s dementia show greater impairments in memory.

Structural and physiological neuroimaging are increasingly being used to aid earlier distinction between dementia with Lewy bodies and Alzheimer’s disease. Computed tomography and magnetic resonance imaging tend to show more advanced hippocampal and mediotemporal lobe atrophy in Alzheimer’s disease than in dementia with Lewy bodies, but this is not specific, especially in early disease.

The dopaminergic system can be physiologically imaged using single photon emission computed tomography (SPECT) and positron emission tomography (PET) and the ligands FP-CIT labelled with iodine-123 (N-fluoropropyl-2-carbomethoxy-3b-4-123I-lodophenyl tropane) and dopa labelled with fluorine-18 (18F-fluorodopa), respectively. Lewy bodies cause disruption of the presynaptic dopaminergic neurones in the corpus striata, as shown by reduced ligand uptake in these areas with SPECT. Ongoing multicentre trials are assessing the concordance between clinical diagnoses, SPECT neuroimaging diagnoses, and postmortem diagnoses for dementia with Lewy bodies. PET is used much less often in clinical practice, but group studies have shown good discrimination between dementia with Lewy bodies and Alzheimer’s disease, with occipital hypoperfusion being seen in dementia with Lewy bodies; one group has suggested that occipital hypoperfusion may be responsible for the visual hallucinations in this condition. Guidelines now reflect the use of neuroimaging in early diagnosis and it is no longer necessary to have two core clinical features of dementia with Lewy bodies, but instead one core clinical feature with supportive neuroimaging can be used to make a probable diagnosis (box). Some clinics also perform analysis of cerebrospinal fluid to look for Creutzfeld-Jakob disease and to search for specific tau and amyloid β proteins seen in Alzheimer’s disease. More recently, cardiac imaging with 123I labelled meta-iiodobenzylguanidine (123I-MIBG) has shown a reduction in autonomic postganglionic sympathetic fibres in the cardiac plexus in Lewy body degenerative diseases, such as dementia with Lewy bodies and idiopathic Parkinson’s disease. This test
Revised criteria for the diagnosis of dementia with Lewy bodies

Core features *
- Fluctuating cognition with pronounced variations in attention and alertness
- Recurrent visual hallucinations that are typically well formed and detailed
- Spontaneous features of parkinsonism

Suggestive features
- Low dopamine transporter uptake in the basal ganglia demonstrated by single photon emission computed tomography and positron emission tomography
- Severe neuroleptic sensitivity
- Rapid eye movement sleep behaviour disorder

*Two core features needed for a probable diagnosis in a cognitively impaired patient, one for a possible diagnosis
†One core feature plus one or more suggestive features in a cognitively impaired patient sufficient to make a probable diagnosis

has been used to discriminate between Alzheimer’s disease and dementia with Lewy bodies in people with moderate dementia. These changes may support the early diagnosis of dementia with Lewy bodies, but they lack specificity because reduced uptake on MIBG scans is also seen in cardiovascular diseases that are common in this typically older age group, such as ischaemic heart disease, cardiomyopathy, and their risk factor, diabetes. A postmortem brain biopsy is needed for a definitive diagnosis, with cortical or brainstem Lewy body pathology being essential criteria. Other changes are also common, such as Alzheimer’s pathology and spongiform pathology. Lewy bodies are accumulations of α synuclein protein that form strongly eosinophilic intracytoplasmic neuronal inclusions. As with Alzheimer’s disease, these protein accumulations are thought to be associated with the clinical symptoms of Lewy body diseases through neuronal cell loss and disruption of normal brain architecture.

3 How should this patient be managed?

Short answer
After baseline investigations in primary care, we referred our patient to local specialist memory services for neuroimaging, neuropsychological assessment, and consideration of pharmacotherapy. A thorough assessment of current needs and future goals should be made with the patient and carers, and appropriate referral should be made to members of the multidisciplinary care team, with planned regular review.

Long answer
UK dementia clinical guidelines from 2006 describe overarching priorities of care for patients with dementia that should guide decision making between patients, carers, and health and social care practitioners. The recommendations include non-discrimination; seeking valid consent using the provisions of the Mental Capacity Act 2005; assessing and providing for the needs of carers; integrating health and social care; providing specialist memory services for all people with a possible diagnosis of dementia; and comprehensively assessing and managing non-cognitive and behavioural symptoms that can distress patients and carers. The Department of Health built on this guidance with its National Dementia Strategy (2009), and most recently with its publication of risk guidance for dementia (2010) to maintain function and to promote and support personal freedoms and independent living in people with dementia. Patients and carers should have early signposting to support groups such as the Alzheimer’s Society (http://alzheimers.org.uk/), Lewy Body Dementia Association (www.lbda.org/), the Lewy Body Society (www.lewybody.co.uk/), Dementia UK (www.dementiauk.org/), and MIND. Carers of patients with dementia will need support—such as psychological services, support groups, and respite care—to help them in their role. Planning should be encouraged, particularly with regard to advocacy, decisions on medical treatment that may include advanced statements; living arrangements; and finances, including appointing a lasting power of attorney. The guiding theme of all management decisions taken with the patient and carers should be one of supporting and maintaining the patient’s independence and quality of life.

Specialist memory services are primarily responsible for balancing the treatment of the many facets of this clinical syndrome—cognitive, behavioural, psychiatric, mood related, and extrapyramidal. First line drugs for cognitive and psychiatric symptoms are the cholinesterase inhibitors—donepezil, galantamine, and rivastigmine—which rarely exacerbate the associated extrapyramidal symptoms; rivastigmine has the most supportive data for this indication. Treatment parkinsonian motor features with dopaminergic agents, commonly carbidopa and levodopa, and titrate them to the optimum dose to balance any improvement in motor function against possible worsening of cognitive and psychiatric symptoms. Depression is common in dementia with Lewy bodies and despite the lack of clinical trials assessing the efficacy of antidepressants in this group, selective serotonin reuptake inhibitors are advised as first line agents, with avoidance of tricyclic antidepressants in case of antimuscarinic side effects. The management of psychotic symptoms, such as visual hallucinations and delusions that can lead to pronounced agitation, are particularly challenging. Patients with Lewy body dementia have marked neuroleptic sensitivity, with increased morbidity and mortality, which abrogates the use of typical antipsychotics. Non-pharmacological methods should be instituted first, and only in the case of severe and recalcitrant symptoms should specialist teams consider small doses of the atypical antipsychotics; clozapine has been consistently more effective than quetiapine for this indication.

Patient outcome
Our patient developed symptoms of depression and was followed up jointly by the local neurology and memory services and mental health teams for older people. She was given mirtazapine for depressive symptoms and the cholinesterase inhibitor, rivastigmine, for first line treatment of cognitive symptoms and visual hallucinations. Physiotherapy was started for her coexistent polyneuropathy symptoms, although frank parkinsonian motor features were mild except for syncopal attacks. Ongoing coordinated review was organised between the general practitioner and secondary care teams.