Structured approach to patients with memory difficulties in family practice

Linda Lee MD MCIsc(FM) CCFP FCFP W. Wayne Weston MD CCFP FCFP George Heckman MD MSc MMATH FRCPC Micheline Gagnon MD MEd FRCP FACP F. Joseph Lee MD MCIsc(FM) CCFP FCFP Scott Sloka MD FRCP

Abstract
Objective To provide family physicians with a structured approach to patients presenting with memory difficulties.
Sources of information The approach is based on an accredited memory clinic training program developed by the Centre for Family Medicine Memory Clinic in partnership with the Ontario College of Family Physicians.
Main message Use of a structured clinical reasoning approach can assist physicians in achieving an accurate diagnosis in patients presenting with memory difficulties. Delirium, depression, and reversible causes need to be excluded, followed by differentiation among normal cognitive aging, mild cognitive impairment, and dementia. Obtaining collateral history and accurate functional assessment are critical. Common forms of dementia can be clinically differentiated by the order in which symptoms appear and by how cognitive deficits evolve over time. Typically, early signs of Alzheimer dementia involve impairment in episodic memory, whereas dementia involving predominantly vascular causes might present with early loss of executive function and relatively preserved episodic memory. Frontotemporal dementia and Lewy body spectrum disorders might have early loss of executive function and visuospatial function, as well as characteristic clinical features.
Conclusion A clinical reasoning approach can help physicians achieve early, accurate diagnoses that can guide appropriate management and improve care for patients with memory difficulties.

While current Canadian consensus guidelines suggest that most patients with dementia can be adequately assessed and managed by their primary care physicians, the issues for patients with memory difficulties are complex and are often challenging to address in family practice. Indeed, for primary care physicians, studies have demonstrated that complexity of care and diagnostic uncertainty remain considerable barriers to the early diagnosis and management of dementia. Currently, most people living in the community with dementia remain undiagnosed and untreated yet early detection offers important benefits. These include the option of early initiation of appropriate medications and access to supports, the opportunity for patients to maximally participate in future care planning, and possible net fiscal benefits.

Achieving an early, accurate diagnosis is therefore an important first step in helping to guide appropriate further management for these patients. Medical education literature suggests that use of a clinically relevant framework or scaffolding can help in assessing patients with complex problems and applying a structured clinical reasoning approach might help family physicians to simplify the process of assessing patients who present with memory difficulties.

Case description
Mrs S. is a 73-year-old patient presenting with memory difficulties. Her medical history includes atrial fibrillation, transient ischemic attack 5 years ago, diabetes, hypertension, hyperlipidemia, osteoarthritis, urinary incontinence, and depression first noted 3 years ago. Her current medications include 20 mg of citalopram once daily; 1 mg of lorazepam at bedtime as needed; 2 mg of warfarin once daily; 2 tablets of a combination of 300 mg acetaminophen and 30 mg of codeine, 3 times daily; 5 mg of bisoprolol once daily; 25 mg of hydrochlorothiazide once daily.

KEY POINTS Diagnostic uncertainty and the complexity of the care required for patients with dementia remain considerable barriers to the early diagnosis and management of dementia in primary care. Early, accurate diagnosis is an important first step in helping to guide appropriate management for such patients. The authors present a 7-step clinical reasoning model to help physicians assess patients presenting with memory difficulties. Office-based cognitive testing can supplement patient history in sorting out the common types of dementia likely to be seen in primary care.

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La traduction en français de cet article se trouve à www.cfp.ca dans la table des matières du numéro de mars 2013 à la page c129.
daily; 10 mg of atorvastatin once daily; 0.125 mg of digoxin once daily; 5 mg of ramipril once daily; 5 mg of glyburide twice daily; and 2 mg of tolterodine twice daily. She is a retired bookkeeper and has 3 children. She is accompanied by her husband and daughter.

During the past 2 years, family members have noticed increasingly frequent missed bill payments and medication errors. Mrs S. now has difficulty with meal preparation and can no longer manage chores. She repeats questions and gets lost in familiar environments. Her daughter is concerned about Mrs S.’s ability to drive.

Sources of information
The approach described below is based on a Mainpro-C accredited memory clinic training program developed by the Centre for Family Medicine (CFFM) Memory Clinic with the support of the Ontario College of Family Physicians.

Clinical reasoning approach (Figure 1)

When an elderly person presents with memory difficulties, the first step is to rule out delirium—a common, potentially fatal condition that is often missed. The Confusion Assessment Method is a validated, easy-to-use screening tool with high sensitivity and specificity for detection of delirium (Table 1).

Those with suspected delirium require a thorough history and physical examination to identify underlying causes. Common precipitants include medications (particularly highly anticholinergic drugs, benzodiazepines, and narcotic analgesics), metabolic imbalance, infection, and occult organ failure such as myocardial infarction or respiratory failure.

The second step is to rule out depression, which might mimic or coexist with dementia. In elderly persons, depression can present atypically with unexplainable physical symptoms, social withdrawal, anxiety, or memory difficulties. In patients with dementia, depression might present as cognitive deterioration, apathy, irritability, or lack of interest in previously enjoyed activities. The Geriatric Depression Scale or SIG E CAPS mnemonic list of symptoms can be used to screen for depression in elderly persons with normal cognitive functioning or mild to moderate cognitive loss. The Cornell Scale for Depression in Dementia is the most validated screen for depression in patients with moderate to severe dementia but it might be less practical for use in busy family practice. If depression is suspected, a trial of therapy with antidepressant medication with low anticholinergic load and few drug interactions (eg, citalopram) might be warranted, with reassessment of cognitive functioning after the depression is adequately treated. Of note, patients experiencing a first episode of depression after age 60 often go on to develop dementia.

It is important to exclude reversible causes as outlined in Figure 1. If none is identified, the next step

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**Figure 1. CFFM Memory Clinic clinical reasoning model**

1. **Is it delirium?**
   - Use the Confusion Assessment Method:
     - Acute onset and fluctuating course
     - Inattention
     - Disorganized thinking or altered level of consciousness

2. **Is it depression?**
   - Consider atypical presentations: anxiety, irritability, unexplained physical complaints, worsening cognition

3. **Is there a reversible cause?**
   - Measure CBC, TSH, creatinine, electrolytes, calcium, glucose, and vitamin B12; consider cranial imaging

4. **Is it dementia, MCI, or normal aging?**
   - Dementia: objective findings of cognitive loss with impairment of ADLs
   - MCI: objective findings of cognitive loss without impairment of ADLs
   - Normal cognitive aging: no objective findings of cognitive loss

5. **If it is dementia, what type or types?**
   - AD: initial short-term memory loss
   - VaD: vascular risk factors; neuroimaging evidence of cerebrovascular involvement
   - FTD: younger age, behavioural symptoms, or language impairment
   - DLB: bradykinesia or features of parkinsonism, fluctuating cognition, visual hallucinations
   - PDD: dementia occurring >1 y after onset of Parkinson disease motor symptoms

6. **How will you manage this?**

7. **Is driving a concern?**

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AD—Alzheimer dementia, ADLs—activities of daily living, CBC—complete blood count, CFFM—Centre for Family Medicine, CT—computed tomography, DLB—Lewy body dementia, FTD—frontotemporal dementia, MCI—mild cognitive impairment, MRI—magnetic resonance imaging, PDD—Parkinson disease dementia, TSH—thyroid-stimulating hormone, VaD—vascular dementia.
is to determine whether the patient's memory difficulties represent "normal" cognitive aging, mild cognitive impairment (MCI), or dementia. These represent a continuum of cognitive states in the elderly. Normal aging is characterized by patients' subjective complaints of memory loss but no substantially abnormal findings on cognitive testing. Slower processing speed and naming difficulties particularly with proper nouns can be common manifestations of normal cognitive aging, but, typically, forgotten memories are recalled with cues. Mild cognitive impairment refers to the symptomatic predementia stage and has a prevalence of 16.8% in those 65 years of age and older and a lifetime conversion rate to dementia of 60% to 80%. In MCI, there are subjective complaints of memory loss and objective evidence of impairment on cognitive testing, but no substantial decline in functional abilities. Validated tools such as the Montreal Cognitive Assessment (MoCA) can be used to screen for impairment in various domains of cognitive functioning. Compared with the Mini-Mental State Examination (MMSE), the MoCA is a more sensitive screening tool for the milder stages of cognitive impairment commonly seen in ambulatory care. The MoCA is validated for use in various medical conditions, in many different languages, and in several international settings; currently, the Canadian Stroke Consortium and the US National Institutes of Health recommended that the MoCA be used for cognitive assessment.

Functional impairment is determined by the patient's decline in ability to perform 1 or more activities of daily living, which can be assessed with the CFFM Memory Clinic Brain Map, available from CFPlus.*

If a decline in functional abilities is identified and there is evidence of considerable impairment in at least 1 domain on cognitive testing, then a diagnosis of dementia can be considered. The Diagnostic and Statistical Manual of Mental Disorders, 4th edition, criteria for dementia include memory loss, decline in at least 1 other cognitive domain (aphasia, apraxia, agnosia, or executive function), and substantial functional, social, or occupational impairment with decline from the previous level of functioning. Revisions are proposed for the fifth edition, in which dementia criteria will be redefined. The prevalence of dementia in Canadians older than 60 years of age is estimated to be 7%. Common types of dementia include Alzheimer disease, vascular dementia, and mixed dementia (Alzheimer plus vascular pathology); in community-based studies, mixed dementia seems to be most common. Less frequently encountered dementia types include Lewy body spectrum disorders (Lewy body dementia and Parkinson disease dementia) and frontotemporal dementia. It is important to diagnose the type of dementia involved, if possible, because treatment and prognosis for patients can differ depending on the underlying cause. The differentiation between the various types of dementia remains a clinical exercise based on the order in which symptoms appeared and the evolution of specific cognitive deficits over time.

**Clinical features of common types of dementia.** Typical Alzheimer disease is characterized by early episodic memory loss (deficits in recall of recently learned information, often referred to as *short-term memory loss*), followed by later involvement of executive dysfunction and visuospatial impairment as the pathology spreads from the medial temporal lobe and hippocampus to other areas of the brain. Much less commonly, Alzheimer

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*The Centre for Family Medicine Memory Clinic Brain Map is available at www.cfp.ca. Go to the full text of the article online, then click on CFPlus in the top right-hand side of the page.*

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<tr>
<th>CRITERIA</th>
<th>DESCRIPTION</th>
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<tbody>
<tr>
<td>1. Acute onset and fluctuating course</td>
<td>Is there evidence of an acute change in the patient's mental status? Did this behaviour fluctuate during the past day—ie, come and go or increase and decrease in severity? (Usually requires information from family members or caregivers)</td>
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<tr>
<td>2. Inattention</td>
<td>Does the patient have difficulty focusing?—eg, are they easily distracted or do they have difficulty keeping track of what is being said? (Inattention can be detected by the digit span test or asking for the days of the week to be recited backward)</td>
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<td>3. Disorganized thinking</td>
<td>Is the patient's speech disorganized or incoherent, such as rambling or irrelevant conversation, unclear or illogical flow of ideas, or unpredictable switching between subjects? (Disorganized thinking and sleepiness can also be detected during conversation with the patient)</td>
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<tr>
<td>4. Altered level of consciousness</td>
<td>Overall, would you rate this patient's level of consciousness as alert (normal), vigilant (hyperalert), lethargic (drowsy, easily aroused), stupor (difficult to arouse), or coma (cannot be roused)? (All ratings except alert are scored as abnormal)</td>
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Data from Inouye et al.21
disease can present atypically with prominent and early behavioural problems, executive dysfunction, or other focal cortical syndromes. On cognitive testing, patients with Alzheimer disease usually demonstrate deficits in episodic memory early in the disease, with impaired 3-word delayed recall on the MMSE or 5-word delayed recall on the MoCA.

Currently there is a lack of consensus about the diagnostic criteria for vascular and mixed dementias. These conditions might represent a continuous spectrum of conditions with relatively pure vascular-type dementia on one end, relatively pure Alzheimer disease on the other end, and a combination of pathologies (mixed dementia) representing the largest group in between.

At present, vascular dementia remains a clinical-radiologic diagnosis: in addition to vascular risk factors, there must be neuroimaging evidence of cerebrovascular involvement. Typically, there is early loss of executive function owing to vascular disease affecting primarily the frontal lobes and their subcortical connections. Executive function refers to cognitive processes that are responsible for planning, initiating, sequencing, and monitoring complex goal-directed behaviours.

Examples of findings suggestive of executive dysfunction include abnormalities in the Trail Making Test Part B (which tests attention switching between letters and numbers), phonemic verbal fluency (eg, f words on the MoCA), or the Luria Fist-Edge-Palm test.

Mixed dementia remains an enigma with no currently accepted, validated clinical guidelines for diagnosis. Patients with mixed dementia will have features of both Alzheimer disease and vascular dementia. They will have vascular risk factors but neuroimaging evidence of cerebrovascular infarcts might or might not be present.

Frontotemporal dementias typically present in the middle years of life with early progressive changes in behaviour, personality, or language functioning. Behavioural changes include loss of social skills, emotional blunting, loss of insight, and lack of concern. Patients presenting with language forms of frontotemporal dementia might have word-finding difficulties and speech that is nonfluent, agrammatic, containing primarily nouns, with frequent word-finding pauses or fluent but with impaired comprehension. On cognitive testing, there is often early loss of executive function with relative preservation of memory and visuospatial skills.

Lewy body spectrum disorders represent a continuum of diseases associated with Lewy body pathology and include Parkinson disease dementia and Lewy body dementia. These conditions share common clinical features of bradykinesia and other symptoms of parkinsonism, fluctuating alertness and cognition, and often the presence of well-formed visual hallucinations.

Although it is an arbitrary distinction, Parkinson disease dementia and Lewy body dementia have been differentiated by the “1-year rule”: Parkinson disease dementia is suspected if onset of dementia occurs after a year or more of parkinsonism; in Lewy body dementia, dementia onset is before or within a year of the development of parkinsonism symptoms. Often, findings on cognitive testing demonstrate early loss of executive and visuospatial function with relative sparing of memory and language functions until the later stages of illness.

Screening tests for visuospatial function (constructional praxis) can include pen-and-paper reproduction of intersecting pentagons on the MMSE or the 3-dimensional cube on the MoCA. Clock drawing tests assess both visuospatial and executive functioning. Visuospatial tasks rely primarily on parietal lobe functioning but also involve frontal and occipital lobes.

Steps 5 and 6 of the CFFM Memory Clinic clinical reasoning model address management and driving issues that have been well documented elsewhere. Molnar et al provide a practical approach to assessing fitness to drive in dementia.

Brain map. Common types of dementia can be clinically differentiated by the order in which symptoms and signs appear over time. This requires careful history taking with corroborated information from family members, as well as supporting evidence on cognitive testing and targeted physical examination for focal neurologic deficits or features of parkinsonism. Areas of deficits on cognitive test performance can help to identify the parts of the brain affected by the pathologic process, thus providing important clues to the type of dementia involved. To facilitate localization of these deficits, we have developed a CFFM Memory Clinic Brain Map* to assist the busy primary care physician in mapping elements of cognitive screening tests that can help identify the type of dementia involved. This approach to education is consistent with the theory of cognitive apprenticeship, in which the Brain Map, along with the clinical reasoning model, serves as a scaffold to guide thinking processes as physicians become more skilled in assessing patients with memory problems.

Case resolution

On cognitive testing, Mrs S.’s MoCA score is 14 out of 30, with 0 out of 5 for delayed recall, impairment on the Trail Making Test and cube drawing, and verbal fluency of just 5 f words. Performance on the clock drawing test is grossly impaired, as is performance on the Trail Making Test Part B, which has numerous errors and requires 5 minutes to complete. She is unable to perform the Luria Fist-Edge-Palm test. She does not appear depressed. A cranial computed
tomography scan reveals moderate periventricular microangiopathic changes.

Working through the CFFM clinical reasoning model, Mrs S. does not meet the criteria for delirium based on the Confusion Assessment Method; however, her cognition might be adversely affected by lorazepam, codeine, and tolterodine (which has high anticholinergic load), and these drugs should be discontinued and her cognitive functioning reassessed. Her depression is appropriately treated with citalopram. If, after medication adjustment, her cognitive deficits persist, she might have dementia, as shown by impairment in activities of daily living and objective findings of deficits on cognitive testing in episodic memory (delayed recall), executive functioning (trails testing, verbal fluency, Luria test), and visuospatial functioning (clock and cube drawing).

The likely diagnosis is mixed dementia, given features of both Alzheimer disease (predominance of episodic memory loss) and vascular dementia (transient ischemic attack and vascular risk factors). Appropriate management can be initiated, and fitness to drive must be assessed.

Conclusion
A 7-step clinical reasoning approach and the use of a Brain Map can assist primary care physicians in assessing patients presenting with memory difficulties. After excluding delirium, depression, and reversible causes, a distinction must be made between normal cognitive aging, MCI, and dementia. Common forms of dementia can often be clinically differentiated by the order in which cognitive deficits appear and by how symptoms evolve over time. Office-based cognitive testing can supplement patient history in sorting out symptoms evolve over time. Office-based cognitive testing can supplement patient history in sorting out symptoms evolve over time. Office-based cognitive testing can supplement patient history in sorting out symptoms evolve over time. Office-based cognitive testing can supplement patient history in sorting out symptoms evolve over time. Office-based cognitive testing can supplement patient history in sorting out symptoms evolve over time. Office-based cognitive testing can supplement patient history in sorting out symptoms evolve over time. Office-based cognitive testing can supplement patient history in sorting out symptoms evolve over time.

Appropriate management can be initiated, and fitness to drive must be assessed.

Competing interests
None declared.

Correspondence
Dr Linda Lee, Centre for Family Medicine, 10 B Victoria St S, Kitchener, ON N2G 1C5; e-mail joellinda@rogers.com

References