Chronic disease self-management support for persons with dementia, in a clinical setting

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Abstract: The burden of chronic disease is greater in individuals with dementia, a patient group that is growing as the population is aging. The cornerstone of optimal management of chronic disease requires effective patient self-management. However, this is particularly challenging in older persons with a comorbid diagnosis of dementia. The impact of dementia on a person’s ability to self-manage his/her chronic disease (eg, diabetes mellitus or heart failure) varies according to the cognitive domain(s) affected, severity of impairment and complexity of self-care tasks. A framework is presented that describes how impairment in cognitive domains (attention and information processing, language, visuospatial ability and praxis, learning and memory and executive function) impacts on the five key processes of chronic disease self-management. Recognizing the presence of dementia in a patient with chronic disease may lead to better outcomes. Patients with dementia require individually tailored strategies that accommodate and adjust to the individual and the cognitive domains that are impaired, to optimize their capacity for self-management. Management strategies for clinicians to counter poor self-management due to differentially impaired cognitive domains are also detailed in the presented framework. Clinicians should work in collaboration with patients and care givers to assess a patient’s current capabilities, identify potential barriers to successful self-management and make efforts to adjust the provision of information according to the patient’s skill set. The increasing prevalence of age-related chronic illness along with a decline in the availability of informal caregivers calls for innovative programs to support self-management at a primary care level.

Keywords: chronic disease, dementia, self-management, cognitive domains

Background
Chronic disease is the leading cause of ill-health in developed nations and causes two-thirds of all deaths worldwide.1 The burden of chronic disease is greater in people with dementia,2,3 and this population is growing with an aging population. By 2050, the global population of those aged ≥80 years is projected to reach 379 million4 and there will also be 115.4 million people with dementia.5 Contemporary dementia care requires a positive and enabling approach that supports the person to live well and remain as involved in their care as is possible within the limitations of impaired cognitive domains.

Appropriate care and management may prevent or delay chronic disease and reduce progression and disability.6,7 Successful chronic disease intervention involves a coordinated multidisciplinary care team, with a central core of patient self-management.7,8 However, strategies to develop effective self-management, especially in persons with
dementia, appear to be lacking. There remain high rates of non-adherence to both medication and prescribed lifestyle change among patients with chronic diseases, with considerable associated economic and health burden as a result.\textsuperscript{9–12}

Lorig and Holman\textsuperscript{13} have characterized self-management into five core processes (Table 1) that each requires a complex set of physical, emotional and cognitive abilities.\textsuperscript{14,15} Conceptualizations of self-management tend to be based on self-efficacy or social cognitive theory.\textsuperscript{12,13} From a cognitive perspective, self-management involves a dynamic and continuous process of self-regulation and adaptation to feedback.

Therefore, decrements in cognitive functioning, level of insight into own abilities, previous coping mechanisms and the social context impede an individual’s ability to successfully self-manage chronic disease unless treatment and support approaches are adapted accordingly.\textsuperscript{16} However, the current health care system relies on the assumption that older populations have well-developed skills required for self-management of chronic disease. A person’s ability to self-manage is also prone to be overestimated as dementia often goes undiagnosed (Table 2) until significant deterioration has developed.\textsuperscript{17} Clinicians must acknowledge that in addition to the relationship between cognition and self-management, this ability is also modulated by a range of complex, dynamic and interlinked factors, including age, race, socioeconomic status, participation in self-management education programs, education level, doctor–patient interaction, disease knowledge, disease severity, experience with disease, social support and psychosocial well-being.\textsuperscript{18–20}

Multiple models that describe the cognitive domains required for general day-to-day adaptive functioning exist.\textsuperscript{12} Commonly described domains include attentional functions, speed of information processing, visuospatial and constructional skills, new learning and memory, receptive and expressive language, praxis, abstract reasoning and executive functions (which include self-monitoring, problem identification, planning, judgment, insight and response regulation).\textsuperscript{21} The impact of dementia on a patient’s ability to self-manage will vary according to the cognitive domain(s) affected, severity of the impairment and complexity of the self-care tasks.

A better understanding of how different forms of cognitive impairment may impact on the key skills of self-management will better enable clinicians’ identification and response to the causes of treatment non-adherence.

**Aim**

We describe the challenges that older patients with a comorbid dementia syndrome and clinicians may face in chronic disease management and the potential interventions aimed at improving self-management. A framework is presented for the purpose of identifying the impact of specific cognitive domains on self-management processes as well as the measures clinicians may take to address these and hence improve management of these persons.

**Describing the effect of dementia syndromes on the domains of chronic disease self-management**

Lorig and Holman\textsuperscript{13} described the five key processes of disease self-management as problem identification and solution generation, decision-making, utilizing appropriate resources, working with clinicians and “taking action”. Impairment of specific cognitive domains may impact each process differently (Table 1). For instance, mild cognitive impairment and Alzheimer’s disease predominantly impact learning and memory, frontotemporal dementia impacts language and executive function, while vascular dementia impacts domains of attention, information processing as well as visuospatial ability and praxis.\textsuperscript{22} Dementia syndromes also vary in their profile of impairment across cognitive domains; thus, clinicians must be able to acknowledge that patients require individually tailored strategies to optimize their capacity for self-management (Table 1).

**Process 1: identifying problems and generating realistic solutions**

Process 1 is identifying problems and generating realistic solutions. This is a cognitively demanding process and likely to be overwhelming for an individual with declining cognitive capabilities. Patients with poor executive functions may be confused or overly agitated and combative as it is difficult for them to shift their thinking or consider alternate options presented to them. Conversely, these patients may appear passive and submissive to all recommendations that a clinician suggests, as they are unable to generate ideas or solve problems independently (Table 1).\textsuperscript{23,24} Patients with mild-to-moderate dementia may have some awareness of their declining health and/or cognitive functioning, but they may be reluctant to acknowledge this for fear that it will result in a loss of their independence. This scenario makes it difficult for the clinician to be certain that patients have a sound understanding of their condition and its implications.\textsuperscript{25} Many forms of dementia are characterized by “rapid forgetting”, which means that even patients who have attended to important information during a visit are likely to have forgotten it by the time they return home (Table 1).\textsuperscript{26,27}
### Table 1 Effect of cognitive impairment due to dementia on chronic disease self-management

<table>
<thead>
<tr>
<th>Tasks and subtasks of self-management</th>
<th>Cognitive domain</th>
<th>Impact of impairment, possible presentations</th>
<th>Suggested strategies</th>
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</thead>
<tbody>
<tr>
<td>1. Problem solving: identifying problems and generating solutions</td>
<td>Attention and information processing: requires adequate orientation and alertness, capacity to sustain attention and adequate processing speed (to “keep up” with presentation of info). Reduced processing speed is a common aspect of aging. However, if the rate of decline significantly exceeds normal rates of aging, the patient may struggle to process verbal information and will become overwhelmed, making it difficult to attend to new information.</td>
<td>Patient may talk over the top of the clinician or ask repetitive questions. Patients may “shut down” and become disengaged from the conversation, with passive agreement, minimal responses or a lack of asking meaningful questions. At follow-up appointments, the patient may recognize information previously presented but will not be able to freely recall it on their own. Rapid forgetting: important information attended to during visit likely to be forgotten in a short period of time.</td>
<td>Asking a patient at the end of a session to summarize the key points of information that were discussed will give an indication of understanding and assist the patient to further encode and rehearse important information. Encouraging carers or family members to attend appointments may be appropriate. Separating information into manageable chunks and checking understanding at each step. Written notes or reminders of important information. Such notes should be composed in such a way that does not require the patient to rely on recall of previously conveyed information in order to understand or act upon them. Ask the clinician to summarize, and check for understanding at the end of each session. Use of tables and flowcharts as memory aids. Explain importance of medication. Use repetition to re-emphasize medication importance. Practice telling the clinician the medication name and the positive or negative effects.</td>
</tr>
<tr>
<td>Patient acquires information</td>
<td>Patient understands significance of information</td>
<td>Patient may be unable to acknowledge the extent of their health issues. They may disregard the implications of ill-health. They may be dismissive of proposed solutions.</td>
<td>Explanation of disease process in very simple terms and provide concrete evidence of the patient’s pathology (eg, imaging results). Provide clear examples how the disease will impact ability to enjoy personally meaningful activities (eg, “unless we get this under control, it will mean that you will have to stop driving. That will make it hard to keep playing bowls and visiting your grandchildren”).</td>
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<tr>
<td>Patient generates solutions</td>
<td>Executive functions and abstract reasoning</td>
<td>Patient unable to generate simple solutions to disease management problems (eg, patient fails to visit a GP or specialist for important review because person they usually obtain lift from is not available. They are unable to generate the idea of using a taxi or calling somebody for assistance).</td>
<td>Teach the patient a limited variety of response options before the need arises. These options will provide a baseline to refer to when they are in a situation where a response is required spontaneously.</td>
</tr>
<tr>
<td>2. Decision-making: acting in response to changes in disease condition</td>
<td>Executive functions: requires the ability to think abstractly, predict consequences and formulate plans. Requires the ability to self-monitor and to shift from previously learned patterns of behavior in the face of novel problems</td>
<td>Patients may respond concretely (eg, repeatedly presenting to hospital for non-life-threatening changes in condition) if they are unable to plan and predict (eg, to self-titrate their medication or to escalate problems to their GP).</td>
<td>Regular monitoring and reassurance to increase patients and family’s confidence that problems can be self-managed successfully. Access to home-based supports such as District Nursing, home help and regular GP contact.</td>
</tr>
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<td><strong>Memory</strong>: patients must be able to remember and understand information about their condition and treatment options in order to make informed decisions.</td>
<td></td>
<td>Patients with impaired recall may be agreeable to any suggestion made to them and may appear motivated, but due to their poor retention of why and how they are implementing changes will find it difficult to guide their own management.</td>
<td>Individually tailored flowcharts, decision trees or checklists may help to prompt patients to put known solutions into practice in response to changed condition. Regular monitoring or scheduled appointments may also be helpful in supporting patients to monitor and proactively monitor their own health over time.</td>
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<td><strong>3. Finding and utilizing appropriate resources</strong></td>
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<td>Use medical devices</td>
<td>Praxis, visuospatial and constructional abilities: requires high-level motor control and to be able to coordinate complex motor movements. Requires ability to process the visuospatial elements of equipment and construct or modify devices to enable use.</td>
<td>Failure to adhere to medication and lifestyle regimen that is dependent upon practical skills, such as comprehension of written instructions/health information, setting up and using equipment or monitoring devices (eg, inhalers for pulmonary disease, glucometers for diabetes mellitus monitoring, use of CPAP for severe sleep apnea).</td>
<td>Direct observation of patient's use of equipment. Use of tailored equipment. Allied health input from physiotherapy, occupational therapy, speech pathology, diabetes educators, etc.</td>
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<td>Requires ability to comprehend written instructions and health information for appropriate use of devices (language).</td>
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<td>Attends clinical appointments</td>
<td>Planning and response regulation (executive functions): requires the ability to plan and carry out plan without being waylaid by competing goals or demands. Visuospatial ability: requires adequate topographical orientation.</td>
<td>Appears overwhelmed and anxious or fails to attend appointments, particularly if they are not consistently scheduled for the same time or if there are multiple appointments scheduled with different, but similar service providers. Gets lost on way to appointments, particularly in hospital outpatient clinics where they may have other appointments in different locations. Unable to access transport or find their way to an emergency department or clinic.</td>
<td>Provision of call back or reminder notices or similar prompts. Establish routine that does not vary. Provision of escort. Where possible provide home visits.</td>
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<tr>
<td><strong>4. Working with health care professionals to make decisions about treatment</strong></td>
<td>Executive functions: requires insight, judgment and abstract reasoning.</td>
<td>May be unable to agree upon mutually acceptable goals of care and appear “stubborn” due to lack of insight build.</td>
<td>Rapport building to develop a patient’s trust. Clinician may present a small selection of management options at a health literacy level (brief, plain language) that the patient may understand. Efforts should be made to identify the reason for unwillingness to take on new goals (ie, whether past habits serve a protective psychological purpose). Illustrated how health care goals may serve the patient’s personally valued goals/priorities in life.</td>
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Communicates with services and negotiates interpersonal relationships

Expressive and receptive language. Deterioration in language skills, often characterized by word-finding difficulties, makes it difficult for these patients to accurately express the nature of changes in their symptoms to clinicians. They may be unable to describe symptoms clearly or appear to delay seeking appropriate help. If comprehension and expressive language skills are limited, patients may become argumentative and combative, particularly if they are frustrated with the management of their symptoms or feel they are not being understood.

Psychological and emotional adjustment

Requires intact response regulation (including emotional control) in order to behave appropriately in social situations. Patients can often appear overwhelmed and anxious even by seemingly small changes in their health or care regime. Patients may also become increasingly combative with care providers as an expression of feelings of helplessness in the context of worsening symptoms. Helplessness and frustration frequently escalate to verbal and physical aggression in individuals with impaired executive functions.

5. Taking action

Adheres to monitoring, medication and lifestyle change

Planning and response regulation (executive functions): requires the ability to plan and organize daily activities and then regulate their behavior in order to execute their plan.

Memory: must be able to recall planned activities and health care tasks (medication regimes, appointments, etc).

Motivation: must remain motivated to adhere to regimes that may be complex and inconvenient.

Patients with impaired executive functions may be impulsive or unable to override ingrained habits in order to maintain agreed patterns of behavior changes. Failure to adhere to and monitor prescribed management may result in fluctuations in health and increasing reliance upon clinical and care staff to monitor treatment regimens. Missed medication doses, appointments or investigations.

Patients with low mood may present as unmotivated as well as having slowed cognition that may mimic dementia symptoms.

Careful clarification of symptoms, which may require the use of closed questions. Collateral history from family members and carers may help identify concerns.

Encourage patients to attend clinic with a supportive family member or friend. Clinicians can undertake basic training to respond to patients who are aggressive or emotionally labile. Written behavior support plans may be useful for patients who are known to be persistently aggressive.

Without further investigation may initially appear as non-compliance or a nonchalant approach to medication management. Implement behavior change to circumvent entrenched behavior patterns. If repeated consistently, new routines may develop into habits that require less conscious thought. May need to involve family members or carers.

 Provision of prompts or cues in the home (eg, wall calendar listing appointments; dose administration aids (Blister packs)). Additional supports from community nursing. Use of electronic diaries with automatic reminders (eg, phone, tablet).

Alertness to and screening for symptoms of depression in the elderly population, particularly those with early cognitive decline, is important, followed by appropriate treatment.

Note: “Self-management skills as described by Lorig and Holman.”

Abbreviations: CPAP, continuous positive airway pressure; GP, general practitioner.
An example of a skill required for this process is an understanding of the significance of important information. If individuals with dementia or cognitive impairment have deficits in the domain of executive function, they may not be able to acknowledge conceptually the extent of their health issues. Additionally, they may be dismissive of proposed solutions as it is difficult to comprehend alternate options to what their current course of management may be. Clinicians may circumvent these short comings by explaining the disease in simple terms and providing solid evidence of the patients' pathology (eg, imaging results) as well as discussing how the disease will impact their ability to enjoy meaningful activities (Table 1).

**Process 2: decision-making – acting in response to changes in disease condition**

Process 2 of self-management of chronic diseases requires decision-making by the patients on a daily basis to respond to changes in their condition and the environment. This requires the ability to think abstractly, conceptualize plans and predict consequences. It also requires patients to have intact learning and memory systems in order to use frameworks that aid decision-making (Table 1).

Patients with impaired memory or executive functions may frequently present to the emergency department when they notice subtle, non-life-threatening changes in their health. This pattern of behavior indicates that the patients are able to monitor their health and effectively identify changes but lacks the capacity to 1) retain the information provided to them by treating staff about how to best respond to such changes in the future and/or 2) generate different ideas about how to act accordingly. An example of an ability required to carry out this process is to choose appropriate solutions among a variety of solutions generated by the individual (if they are able to generate solutions in the first instance [Table 1, subtask {Process 1}]) according to changes in their condition/environment. Patients who have impairments in memory may not be able to remember or understand information about their condition and, hence, may not be able to make efficacious and informed decisions to choose the appropriate solution that addresses their situation. Although these persons may be agreeable to clinician suggestions and may appear motivated, due to poor information retention, they will find it difficult to guide their own management when required. They may benefit from management strategies including tailored flowcharts, decision trees or checklists to prompt implementation of known solutions, in response to a changed condition. Regular monitoring or scheduled appointments may be helpful in supporting these patients to monitor and proactively monitor their health over time (Table 1).

### Table 2 Reasons for under-recognition of dementia and chronic disease

<table>
<thead>
<tr>
<th>Reason</th>
<th>Issues related to under-recognition</th>
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<tr>
<td>Health system is under-prepared for a relatively new area of medicine</td>
<td>Rapid surge in prevalence of dementia in recent decades. Dementia only recognized by the Australian government as a National Health Priority Initiative (Australian Institute of Health and Welfare) in 2005. Chronic disease management and self-management models are a relatively new field.</td>
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<tr>
<td>Subtle onset of disease</td>
<td>Subtle difficulties in the performance of everyday activities are common in individuals with mild cognitive impairment 2 years before a diagnosis of dementia.</td>
</tr>
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<tr>
<td>Health professionals lack skills to recognize dementia</td>
<td>Often physicians may not recognize subtle cognitive deficits during routine office visits. Increased awareness is needed in the health care sector. Rate and functional impact of cognitive decline differ across patients. Therefore, establishing what is normal age-related decline as opposed to severe decline for a patient might be difficult. Patients/caregivers may not recognize early symptoms of cognitive decline as a pertinent health issue and/or may choose not to report these to their general practitioner.</td>
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<tr>
<td>Limited awareness of issue by health professionals</td>
<td>Increasing sub-specialization leads to clinicians focusing on single organs or systems, leading to reduced awareness of the patient’s overall functioning, including signs of cognitive impairment. Health professional awareness of dementia may be limited.</td>
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<tr>
<td>Fear and social stigma of disease</td>
<td>Ongoing social stigma of dementia is also present among clinicians. Patients may be fearful of a loss of independence, which may hinder patient willingness to disclose symptoms/abilities to the general practitioner.</td>
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<tr>
<td>Assumptions about dementia</td>
<td>Belief that dementia will present as severe and obvious impairment. Belief that the need to manage chronic disease is not relevant in persons with dementia as the perception of the overall prognosis is poor. Attitude that the relative absence of therapy for dementia is justification for not making a diagnosis. Belief that the carer is able to manage all health-related issues.</td>
</tr>
<tr>
<td>Social isolation</td>
<td>Patients without family or carers can hinder diagnosis as some screening tools (eg, functional assessments) rely on caregiver observation and report. The patient with dementia may lack insight and be too cognitively impaired to provide an accurate history or attend a medical appointment independently.</td>
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</table>
Process 3: finding and utilizing appropriate resources

Process 3 requires the ability to find and use appropriate resources. The treating team needs to consider whether the patient has the necessary motor skills to manage therapeutic devices (eg, medication aids, glucometers, inhalers, home oxygen). The team also needs to evaluate whether the patient has the cognitive skills to access vital health care information (eg, seeking advice via telephone services, accessing a nutrition guide for diabetes mellitus through the Internet) (Table 1). Process 3 also requires the ability to process the visuospatial elements of equipment and construct or modify devices to enable their use, as well as remembering how visuospatial mechanisms work (eg, which way to turn the dial on their insulin pen to increase or decrease their dose) (Table 1).

Patients with dementia often experience symptoms of apraxia in the early stages and may have difficulty with these tasks and therefore appear clumsy. If the impairment is one of higher level praxis (ideational or conceptual praxis), the patients may be unable to perform action sequences in the appropriate order, even for well-learned procedures that they can explain verbally (eg, a patient starts blood pressure monitoring before placing cuff over arm) (Table 1).

Progressive impairment of language function in dementia (eg, primary progressive aphasia) may eventually result in dyslexia, which will impact a patient’s comprehension of written instructions and important health information (Table 1). Additionally, patients with executive function deficits may have impairments in planning and response regulation that may hinder their ability to find their way to appointments or access the necessary transport to the clinic.25,29

As mentioned previously, a skill required for this process is the ability for abstract thought, insight and planning (executive functions), as well as adequate communication skills to work collaboratively (Table 1). Impaired executive functions may be evidenced by a patient’s resistance to implementing lifestyle or treatment changes recommended by clinicians and appearing unwilling to consider suggested options, due to a lack of insight into disease progression.22 Deterioration in language skills, often characterized by word-finding difficulties, is common in cognitive decline, and as such it may be difficult for these patients to accurately express the nature of changes in their symptoms to clinicians.31 A skill required for this process is the ability to communicate with services and negotiate interpersonal relationships. Impairments in the expressive and receptive language domains may manifest as word-finding difficulties, which results in these persons having issues with explaining the nature of their symptoms, changes in their symptoms, management preferences, circumstances and seeking help in a timely manner.21 Furthermore, limited comprehension and expressive language skills may result in patients becoming argumentative and combative especially if they are frustrated with the management of their symptoms or feel misunderstood by clinicians/allied health staff.31 Management strategies to counter this involve careful clarification of symptoms using closed questions and obtaining a collateral history from carers or family members (Table 1).

Process 4: work with health care providers to make informed choices about their treatment

Process 4 requires achieving agreement on overarching goals of care, as well as negotiating specific treatment strategies with health care providers. This relies on the ability for abstract thought, insight and planning (executive functions), as well as negotiation, working with services, and making decisions about their treatment (Table 1). Patients with executive function deficits may have impaired decision-making skills and therefore find it difficult to make choices or discuss trade-offs with their clinician.31 A skill required for this process is the ability to accurately express the nature of changes in their symptoms, changes in their regimen that is dependent on practical skills such as comprehension of written instructions/health information, setting up and using equipment or monitoring devices.30 Potential solutions may include direct observation of the patient’s use of equipment, use of tailored equipment and allied health input by physiotherapy, occupational therapy and speech pathology to allow better use of these devices tailored to patient capabilities (Table 1).

Process 5: taking action

Process 5 requires implementing agreed-upon courses of action that involves the culmination of many of the above-described cognitive skills and processes and is dependent on successfully establishing the previous steps. If patients have successfully attended to and understood information, negotiated care plans and identified when to act, they must still be able to remember to take action and remain motivated and physically able to complete tasks (Table 1).

When patients have an awareness of their own physical and cognitive decline, one significant consequence is deterioration in mood and motivation. Mood issues associated with adjusting
to the vast array of age-related changes are often an overlooked obstacle to developing good self-care. In particular, depressive symptoms often have a “dampening effect” upon cognition and may negatively impact upon motivation, concentration and memory, social engagement and personality—often mimicking the onset of dementia.32 Patients with low mood states may present as unwilling or incapable of following multistep instructions or with a lack of confidence in their ability to independently follow such steps. They may also be preoccupied and have difficulty attending to, and thus learning and retaining, important information discussed.

Patients with impaired executive functions may have difficulty applying a process designed to take over from deeply entrenched behavior patterns—eg, dietary habits developed over a lifetime (Table 1).33 An example of a skill required to fulfill this process is adherence to monitoring, medication and lifestyle. These are well-established, critical features of self-management in chronic disease that, if not complied with, may result in poor outcomes.9–12 If patients are unable to recall planned activities and health care tasks due to memory impairments, they may miss medication doses, appointments or scheduled investigations.21,34 In order to improve self-management in this subset of cognitively impaired persons, prompts or cues in the home (eg, wall calendar listing appointments, dose administration aids) or use of electronic diaries with automatic reminders may be employed. Additional supports from community nursing may be utilized to ensure adherence to recommended monitoring, medication and lifestyle, particularly for those patients who have severe memory deficits.

Implications for clinicians: recognizing cognitive decline and supporting self-management
Recognizing the presence of dementia in a person with chronic disease may lead to better patient outcomes. Optimal disease control through self-management requires a complex set of skills that may be impacted by declining physical, emotional and cognitive health.

A key step for clinicians is an understanding that non-adherence or poor disease control could, in part, be due to an unrecognized comorbid dementia (Table 2). Some deterioration of cognitive skills with age is normal and expected; it is when the level of deterioration is over and above typical rates that it can start to have a negative impact upon a patient’s functional capacity, and thus potentially impede efforts to effectively implement self-management care plans.

The complex composition of different cognitive domains and how they interact is difficult to understand and assess, even for skilled clinicians.35 Additionally, the identification of mild dementia and other indicators of a neurodegenerative process in a patient with multiple comorbidities is a difficult one that can often be overlooked in favor of managing more pressing medical issues.8,17 This is particularly pertinent when considering reductions in higher order executive functions, as this level of thinking requires coordination of multiple cognitive skills. Clinicians often have limited opportunity to observe this level of thinking skills in the short space of time in which they are with their patients, unless specifically looking for and testing these skills.17,35 Formal cognitive testing is required to determine the presence or seriousness of any decline. Dementia is under-recognized with 15%–25% of patients aged over 75 years in general practices meeting criteria for cognitive impairment.36,37 Timely recognition of undiagnosed dementia or cognitive impairment symptoms can aid clinicians in deciding which persons would be suitable for comprehensive neuropsychological assessment and identifying those who would benefit from implementation of recommended techniques for continued self-management. The Mini-Mental State Examination is a commonly used screening tool for dementia; however, it has significant limitations and does not adequately assess executive functioning, making it an unreliable screening measure for early stages of dementia. Other brief screening tests designed to assess executive functioning also have shortcomings.35

Clinical assessment may be able to more effectively identify executive dysfunction by systematically observing and recording their patient’s understanding and ability to undertake complex daily actions.39 For example, asking simple questions such as the date, time and knowledge of current world or local events can provide the clinician an insight into the patient’s awareness of their environment.

Consistent errors or confusion can indicate executive dysfunction, even if the patient provides reasoning for such confusion. Similarly, patients who are highly tangential in conversation, or conversely perseverate on particular topics, may be experiencing some executive dysfunction. Clinicians should take note of how difficult it is to redirect patients in these situations; if a pattern of this behavior is established, executive dysfunction is likely.

It is not realistic or appropriate to refer all patients for a comprehensive neuropsychological assessment to determine the likelihood of dementia.35,39 Clinicians should instead focus on developing an understanding of the functional representation of these areas of cognitive functioning and assess (by means of making clinical observations and having discussions with their patients) their capability of undertaking the tasks required for self-management.
Supportive and adaptive strategies to assist patients to integrate and consolidate new information can be employed to help compensate for cognitive decline.\textsuperscript{14,40} There is also a need for health professionals providing care for patients with chronic disease to develop a flexible and response approach to establishing care protocols to match patient needs and capabilities. Hence, we propose a framework that describes the effect of cognitive impairment on self-care processes. In this framework, we suggest how clinicians may address a patient’s individual cognitive deficits to circumvent self-management issues and, therefore, improve overall chronic disease management for these persons. This approach must be modified as cognitive function continues to deteriorate.

The inevitable progression and functional decline experienced by patients with dementia makes ongoing monitoring of their health and capacity for chronic disease self-management essential. It is vital to continue supporting independence and empowering patients within their capabilities. It is all too easy to strip all self-management tasks from persons with dementia and instead to rely on carers. This approach may be counterproductive, increasing dependence and caregiver stress, and is disrespectful to the patient.\textsuperscript{13–15} The increasing prevalence of age-related chronic illnesses along with a decline in the availability of informal carers increases the importance of primary care providers and a need for innovative programs to support self-management.\textsuperscript{41} Broader education and training for clinicians about dementia and key functional indicators for different types of dementia would be valuable for primary care providers and specialists who regularly manage patients with chronic disease.\textsuperscript{17,42} Another strategy is development of clinical practice guidelines and tools for tailoring management of specific chronic diseases for patients with coexisting dementia. Optimal management of common chronic diseases such as diabetes mellitus, heart failure and chronic pulmonary disease requires a different emphasis on each of the five self-management tasks.

Development of more accurate, functional and behavior-based cognitive screening tools for use by clinicians to identify possible dementia would be helpful. Not only would these assist clinicians who are not specifically trained in neuropsychology to efficiently characterize their patient’s cognitive profile, but they would also facilitate long-term care planning and assist in monitoring of cognitive functioning over time. Involvement of geriatric medicine expertise, occupational therapist support and other specialized support services (eg, aged care assessment teams) as required by the patient’s needs would also be valuable in providing holistic care to this growing patient group.

Conclusion
Managing chronic diseases in the context of cognitive impairment is complex, and patients with dementia are vulnerable to poorer outcomes if they are unable to negotiate self-management. Further to recognizing cognitive impairment, a holistic view encompassing personal capabilities, previous coping mechanisms and the individual’s social context should be considered when formulating management strategies for this population. A greater focus needs to be placed on care services for older adults that empower consumers, support independence and improve the viability and sustainability of the health and aged care sectors.

Providing education and relevant care information to the patient is not sufficient for implementing a self-management strategy. In reality, self-management differs from patient education, in that it incorporates cognitive tasks and strategies that enable patients to change their behavior and manage their condition.\textsuperscript{13}

Prudent practice requires the clinician to take responsibility for assessing the patient’s current capabilities, identify potential barriers to successful self-management and make efforts to adjust the provision of information according to the patient’s skill set.

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References


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