Dementia Management: Medication and More

PRIMARY CARE ROUNDS, MAY 27, 2021

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Learning Objectives

- 1. Know when to start, stop, or switch cholinesterase inhibitors and/or memantine
- 2. Understand risks and benefits of other medications and nonpharmacological alternatives
- 3. Know how to holistically and systematically assess treatment effectiveness

Dementia prevention, intervention, and care: 2020 report of the Lancet Commission



Figure 2: Possible brain mechanisms for enhancing or maintaining cognitive reserve and risk reduction of potentially modifiable risk factors in dementia

thelancet.com Vol 396 August 8, 2020

DOI: 10.1002/alz12105

PERSPECTIVE

Alzheimer's & Dementia® THE JOURNAL OF THE ALZHEIMER'S ASSOCIATION

Recommendations of the 5th Canadian Consensus Conference on the diagnosis and treatment of dementia

Zahinoor Ismail¹ | Sandra E. Black² | Richard Camicioli³ | Howard Chertkow⁴ | Nathan Herrmann⁵ | Robert Laforce Jr.⁶ | Manuel Montero-Odasso^{7,8} | Kenneth Rockwood⁹ | Pedro Rosa-Neto¹⁰ | Dallas Seitz¹¹ | Saskia Sivananthan¹² | Eric E. Smith¹¹ | Jean-Paul Soucy¹³ | Isabelle Vedel¹⁴ | Serge Gauthier¹⁵ | the CCCDTD5 participants

► 1st CCCDTD was in 1989

▶ 5th CCCDTD is the first to include non-pharmacological treatments

Dementia risk reduction, 5th CCCDTD

- Mediterranean diet; avoid saturated fatty acids; increase fruit and vegetable intake
- Physical activity of at least moderate intensity (aerobic exercise and/or resistance training)
- Dance interventions and mind-body exercise (e.g. Tai Chi, Qigong)
- Screen for hearing loss; audiologic rehab +/- hearing devices
- Sleep history; target 7-8 hours sleep per night; insufficient evidence to recommend insomnia treatment
- Computer-based or group cognitive training; cognitively stimulating pastimes, volunteering, life-long learning
- Minimize medications with anticholinergic properties

Antihypertensives

- Treating HTN may reduce risk of dementia
- Assess, diagnose, and treat HTN according to guidelines from Hypertension Canada
- If a vascular contribution is suspected, treat if diastolic > 90 mmHg and if systolic > 140 mmHg (consider target < 120 mmHg if associated vascular risk factors)

Aspirin

- Not recommended for patients with MCI or dementia who have brain imaging evidence of covert white matter lesions of presumed vascular origin without Hx of stroke or brain infarcts
- If there are covert brain infarcts on neuroimaging, but no Hx of stroke, the use of ASA is reasonable, but the benefit is unclear





Dementia management: Why?

There is no cure. There is no "slowing of progression."

- 1. Enhance / maintain cognition
- 2. Enhance / maintain functional status
- 3. Treat symptoms
- 4. Consider the caregiver



Dementia Management: Evidence or Art?

Medications

Nonpharmacological approaches DOI: 10.1002/alz.12105

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► Comprehensive

Informed by both the Evidence and the Art

Is the treatment working?

- Tracking treatment response and change over time should be individualized
- Requires a multi-dimensional approach plus caregiver or reliable informant input.
- Do not rely on a single tool or clinical domain (e.g. MMSE score)
- Reassess every 6 to 12 months (more frequently if there are behavioural symptoms)

The ABCC's of treatment response

Is there a particular target symptom or symptoms?

- A: ADL's and IADL's: is there improvement or stabilization of function?
- ▶ B: Behaviour: are the specific "target symptoms" better?
- C: Cognition: how do post-treatment cognitive tests compare to pretreatment?
- C: Caregiver health: burden / resilience

ADL's and IADL's

Assessing function is integral in the follow up of treated patients

- Disability Assessment in Dementia (DAD)
- Functional Assessment Staging Scale (FAST)
- Functional Activities Questionnaire (FAQ)
- The Older Americans Resources and Services Multidimensional Functional Assessment (OARS)
- The Barthel Index Score

Behaviour

"Response behaviours", behavioural and psychiatric symptoms of dementia (BPSD)

- Neuropsychiatric Inventory brief version (NPI-Q)
- Mild Behavioural Impairment Checklist (MBI-C); need reliable informant
- The Geriatric Depression Scale (GDS); less sensitive with disease progression
- The Cornell Scale for Depression in Dementia
- ► The Patient Health Questionnaire (PHQ-9)



Memory Impairment Screen (MIS) + Clock

<5 min

Mini-Cog

AD8

4-item MoCA

GP Assessment of Cognition







Caregiver

- Caregiver burden is a major determinant of hospitalization and long term care placement
- Zarit Burden Interview
- ► The Resilience Scale
- Informant Questionnaire on Cognitive Decline in the Elderly (IQCODE)
- ► The HABC-Monitor

"Start low, go slow!"



Cholinesterase inhibitors (ChEI)

- Donepezil (Aricept)
- Galantamine (Reminyl)
- Rivastigmine (Exelon)
- Indicated for Alzheimer's disease, Parkinson's Disease Dementia, Dementia with Lewy Bodies, Vascular Dementia
- Not indicated for subjective cognitive decline, Mild Cognitive Impairment (MCI), frontotemporal dementia, or other neurodegenerative conditions
- Assess anticholinergic burden of other medications

When to avoid cholinesterase inhibitors?

- Not wanted
- Absolute contraindications (e.g. 2nd or 3rd degree AV block, asthma)
- Relative contraindications (e.g. seizure disorder, anorexia, frailty)
- Life expectancy less than 6 months
- Frontotemporal dementias unlikely to have an Alzheimer component (e.g. Pick's disease, or behavioural variant FTD)

Switching cholinesterase inhibitors

- Switching not indicated for perceived lack of benefit
- Consider switching if side effects limit treatment
- Donepezil covered by Pharmacare with special authority
- Galantamine covered by Pharmacare if donepezil caused side effects
- Rivastigmine (oral) covered by Pharmacare if donepezil caused side effects
- Rivastigmine patch not covered but better tolerated

Memantine

- Indicated for Alzheimer's disease, Parkinson's Disease Dementia, Dementia with Lewy Bodies, Vascular Dementia
- Indicated for moderate to severe stage dementias listed above
- Not indicated for subjective cognitive decline, Mild Cognitive Impairment (MCI), frontotemporal dementia, or other neurodegenerative conditions
- Not covered by Pharmacare

Behavioural and psychiatric symptoms of dementia (BPSD)



Behavioural and psychiatric symptoms of dementia (BPSD)

Table 2- Examples of BPSD Usually not Amenable to Antipsychotic Treatment

 wandering 	 vocally disruptive behaviour 	 inappropriate voiding
 hiding and hoarding 	 inappropriate dressing /undressing 	 eating inedible objects
repetitive activity	 tugging at seatbelts 	 pushing wheel chair bound residents

Note: Try to avoid use of antipsychotics if possible for residents with dementia due to Parkinson's disease or Lewy Body dementia. Cholinesterase inhibitors are the first line of treatment for residents with psychosis and aggression associated with these type of dementias. Cholinesterase inhibitor drugs are covered by the Ministry of Health through the Alzheimer Drug Therapy Initiative.

Managing behavioural symptoms

Nonpharmacological Strategies	Cognitive Enhancers	 Sedative/hypnotics Melatonin Trazodone Nozinan Benzodiazepines
Antidepressants • Trazodone • SSRI's	Antiepileptics • Gabapentin • Pregabalin	Atypical antipsychotics • Haloperidol • Risperidone • Loxapine • Olanzapine • Quetiapine



Best Practice Guideline for Accommodating and Managing Behavioural and Psychological Symptoms of Dementia in Residential Care

A Person-Centered Interdisciplinary Approach

Key considerations:

- Carefully weigh the potential benefits of pharmacological intervention versus the potential for harm.
- Recognize that the evidence base for drug therapy is modest.
 - (Number needed to treat that ranges from 5-14)[†]
- Engage the resident/family/substitute decision-maker in the health care planning and decision-making process.
- Obtain consent for health care treatment from the appropriate decision-maker before administering antipsychotic medication.
- Regularly review the need (or not) for ongoing antipsychotic therapy for behavioural psychological symptoms of dementia and trial withdrawal.

Physical restraints exacerbate behavioural symptoms.



When to stop ChEI and/or memantine?

- Clinically meaningful worsening of dementia as reflected in changes in cognition, functioning, or global assessment
- Worsening cannot be attributed to other medical conditions (e.g. delirium) or environmental factors (e.g. moving into LTC)
- No meaningful benefit (improvement in target symptom, stabilization, or decreased rate of decline) was ever seen
- Severe or end-stage dementia
- Intolerable side effects
 - ChEI: nausea, vomiting, anorexia, weight loss, falls
 - ► Memantine: confusion, dizziness, falls
- Poor medication adherence precludes safe ongoing use or inability to assess medication effectiveness

How to Deprescribe

- Reduce gradually
- Reduce dose by 50% every 4 weeks
- Discontinue after 4 weeks on the recommended starting dose
- Reinitiate treatment if patient shows clinically meaningful worsening of cognition, function, behaviour that appears to relate to cessation of therapy
- Do not deprescribe ChEI in patients with psychosis, agitation, or aggression until these symptoms have stabilized (unless symptoms appear to have worsened with ChEI initiation or increase in dose)

Psychosocial interventions



Individual

- Exercise
- Group cognitive
 stimulation therapy
- Caregiver psychoeducational interventions



Community

- Dementia friendly
 organizations/communities
- Case management

Case History



Dementia Management Key Points

- Strongly consider non-pharmacological interventions
- Avoid anticholinergics if possible
- Cholinesterase inhibitors and memantine are the only "cognitive enhancers"
- Use sedative/hypnotics, antidepressants, anticonvulsants, antipsychotics sparingly and with caution
- "Start low, go slow"
- Evaluation of treatment response is multidimensional ... ABCC's

Questions?



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