

Dementia & Cognitive Disorders: Disorders: The Clinical Playbook

A Visual Diagnostic & Therapeutic Pathway





2nd Leading Cause of Death in Australia

849,300

Projected by 2054

411,100

Australians living with dementia in 2023

\$36 Billion

Total annual cost

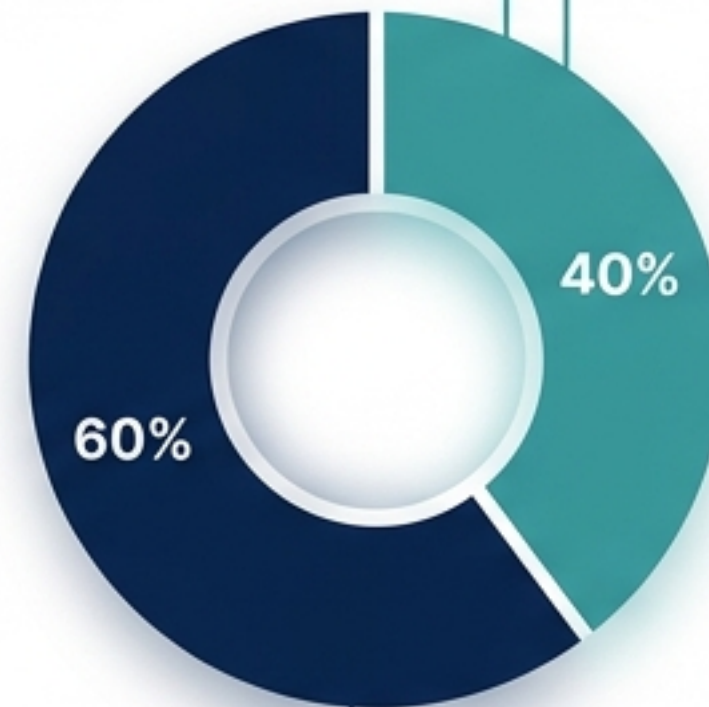
2023

2036

2040

2054

Modifiable Risk Factors



Hypertension

Diabetes

Hearing Loss

Physical Inactivity

Social Isolation

Non-Modifiable Risk Factors

Age (Strongest Factor)

APOE ε4 Allele

Family History



**Universal
Screening is
NOT
Recommended**



Triggers for Cognitive Screening

1



Expressed concern about memory or cognitive changes (Patient or Informant).

2

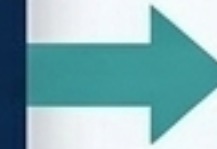


Observed functional decline (missed appointments, medication errors, financial mismanagement).

3



RACGP Health Assessment for patients aged ≥ 75 years (MBS Item 707).



**Proceed to
Cognitive
Screening**

Tool	Cut-off Score	Time	Clinical Utility
Mini-Mental State Examination (MMSE)	$\leq 23/30$	5–10 min	Widely validated, but copyrighted and heavily education-biased.
Montreal Cognitive Assessment (MoCA)	$\leq 25/30$	10–15 min	Superior sensitivity for Mild Cognitive Impairment (MCI); assesses executive function.
★ General Practitioner Assessment of Cognition (GPCOG)	$\leq 4/9$ (Patient)	5 min	Designed for Australian GPs. Includes informant component.
Rowland Universal Dementia Assessment Scale (RUDAS)	$\leq 22/30$	5–10 min	Crucial for Culturally and Linguistically Diverse (CALD) and Indigenous populations. Minimal education bias.

DIFFERENTIATING DELIRIUM & DEMENTIA

DELIRIUM (Medical Emergency)

Onset: Acute (hours to days)

Course: Fluctuating (often worse at night)

Consciousness: Altered, clouded, or drowsy

Attention: Markedly impaired (**Hallmark**)

Reversibility: Usually reversible with treatment

DEMENTIA (Progressive Syndrome)

Onset: Insidious (months to years)

Course: Gradual, progressive decline

Consciousness: Normal until late stages

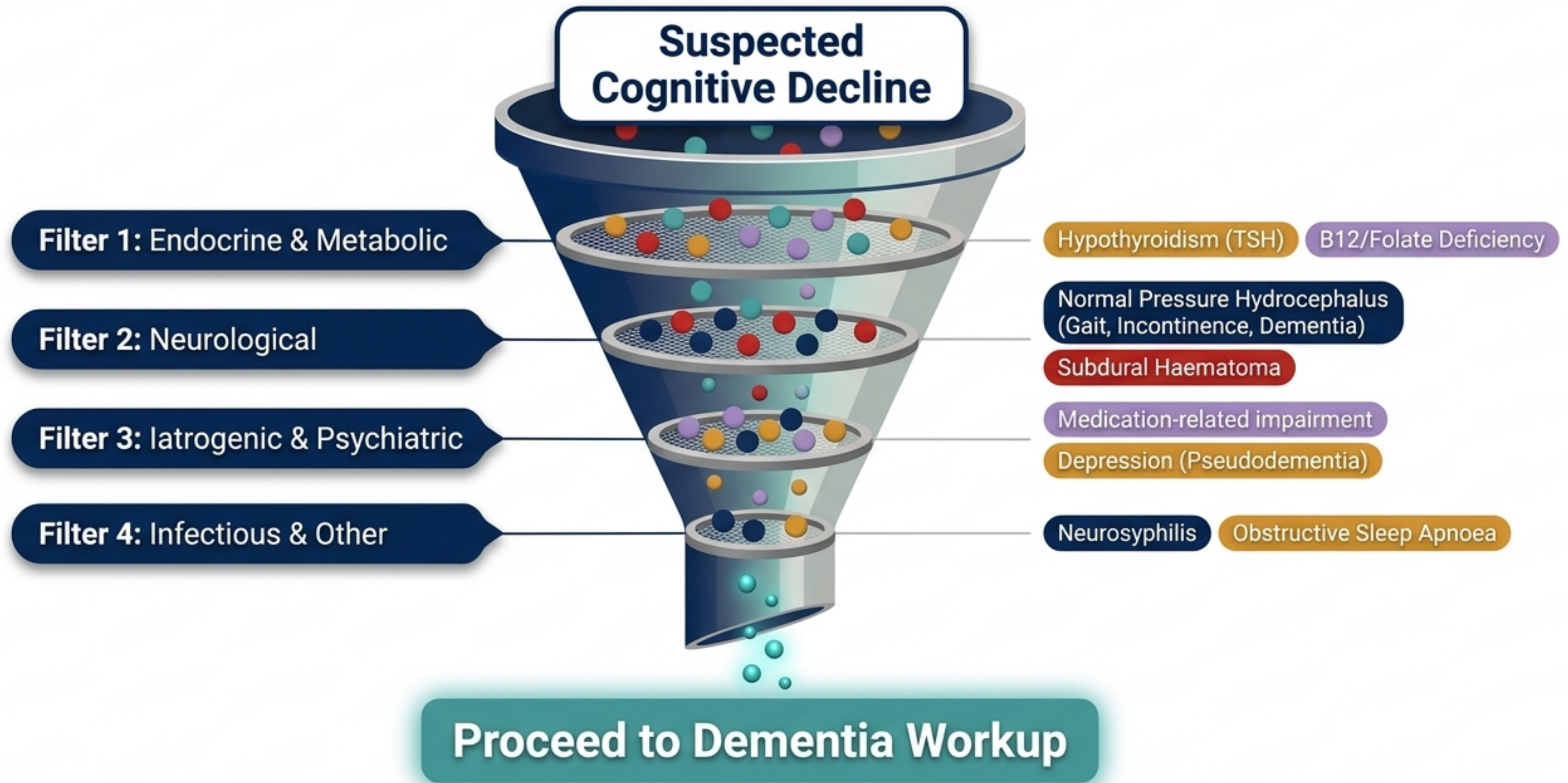
Attention: Relatively preserved until moderate-severe

Reversibility: Generally irreversible



Caution: Delirium is frequently superimposed on pre-existing dementia.
Assess for acute changes from baseline.

DIAGNOSTIC WORKUP: RULING OUT REVERSIBLE CAUSES



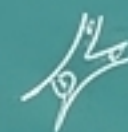
DEMENTIA SPECTRUM & DIAGNOSTIC MATRIX

Prevalence Spectrum of Dementia

Alzheimer Disease
(60-70%)



Vascular
Dementia
(15-20%)



Lewy Body
Dementia
(5-10%)



Frontotemporal
Dementia
(5-10%)



Diagnostic Matrix & Clinical Presentation

Alzheimer Disease



Clinical Features

- Insidious episodic memory decline
- Difficulty forming new memories

Imaging Findings

- Medial temporal atrophy
- Hippocampal atrophy (MRI)

Vascular Dementia



Clinical Features

- Stepwise or gradual decline
- Executive dysfunction
- Psychomotor slowing

Imaging Findings

- White matter hyperintensities (MRI)

Dementia with Lewy Bodies (DLB)



Core Clinical Features

- Fluctuating cognition
- Recurrent visual hallucinations
- Spontaneous parkinsonism

Suggestive Features

- REM sleep behaviour disorder

Frontotemporal Dementia (FTD)



Clinical Variants

- Behavioural variant: personality change
- Disinhibition
- Hyperorality
- Primary progressive aphasia

Imaging Findings

- Frontotemporal atrophy (MRI)

CLINICAL ALERTS & PRESENTATION NUANCES



Severe Neuroleptic Sensitivity in DLB

Patients with DLB can experience life-threatening neuroleptic malignant syndrome and marked worsening of parkinsonism if given typical (haloperidol) or atypical (risperidone) antipsychotics.

Avoid completely Haloperidol Risperidone antipsychoticaliness.

Use quetiapine under strict specialist guidance



FTD and Younger-Onset Presentations

Proportionally higher in those <65 years (up to 20-30% of FTD cases). Frequently negative on standard cognitive screens like MMSE, as memory is often preserved initially. MMSE negative

Misdiagnosed as psychiatric illness

Cholinesterase inhibitors ineffective and

Worsen behavioural symptoms

COGNITIVE IMPAIRMENT DIFFERENTIAL: PSEUDODEMENTIA VS. NEURODEGENERATION



Depression-Related Cognitive Impairment (Pseudodementia)

Subacute onset

History of prior depressive episodes

Clinical Presentation

Patient highlights and complains extensively about memory loss. Often replies 'I don't know' to questions but performs poorly on effort. Affect is flat or tearful. High distress over cognitive failures.

Screen with Geriatric Depression Scale (GDS-15 ≥ 5).
Treat mood and reassess cognition at 8-12 weeks.



True Neurodegeneration

Insidious onset

Gradual decline

Clinical Presentation

Patient often minimizes difficulties or is genuinely unaware of deficits. Frequently confabulates or tries to cover up mistakes. Shows less overt distress regarding the cognitive failure itself.



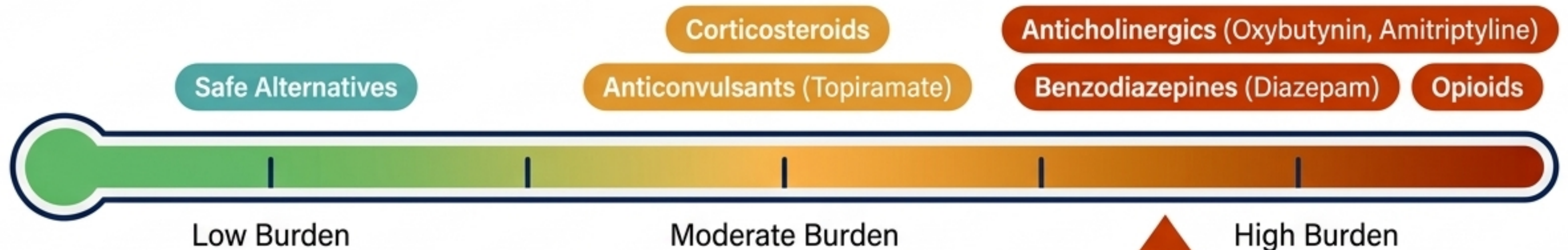
The Informant Interview

Assess Basic ADLs
(self-care, dressing)

Assess Instrumental ADLs
(finances, driving, medications)

Document timeline, trajectory,
and behavioural changes

Anticholinergic Cognitive Burden Scale



Deprescribing is Mandatory. Identify and taper high-burden medications.

COGNITIVE IMPAIRMENT DIFFERENTIAL: LABORATORY WORKUP

REVERSIBLE & TREATABLE CAUSES SCREEN



FBC (Full Blood Count)

Rationale: Rule out anaemia, macrocytosis, infection.



UEC & LFTs

(Urea, Electrolytes, Creatinine & Liver Function Tests)

Rationale: Rule out renal failure, hyponatraemia, hepatic encephalopathy.



TSH

(Thyroid Stimulating Hormone)

Rationale: Rule out Hypothyroidism (Reversible).



B12 & Folate

Rationale: Rule out subacute combined degeneration (check methylmalonic acid if borderline).



Glucose/HbA1c & Calcium

Rationale: Assess vascular risk; rule out hypercalcaemia.

Targeted Tests (Specific Clinical Indication)

RPR/VDRL (Neurosyphilis)

HIV Serology (HAND)

ESR/CRP (Vasculitis)

COGNITIVE IMPAIRMENT DIFFERENTIAL: NEUROIMAGING ASSESSMENT

CT Brain (Non-contrast)

Primary Care Baseline

Adequate for excluding space-occupying lesions, hydrocephalus, subdural haematomas, and large infarcts (MBS Item 56000).

MRI Brain (Preferred)

Structural Detail

Superior for assessing Medial Temporal Atrophy (MTA scale for AD) and White Matter hyperintensities (Fazekas scale for Vascular). Requires specialist referral.

FDG-PET / Amyloid PET

Metabolic / Pathological Profiling

Shows frontotemporal hypometabolism (FTD) or temporoparietal (AD). Detects amyloid plaque burden to confirm AD pathology.

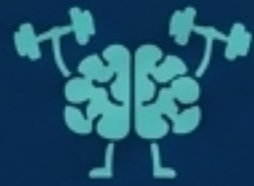
DAT-SPECT (DaTSCAN)

Receptor Imaging

Shows reduced dopamine transporter uptake in putamen. Used specifically to distinguish Dementia with Lewy Bodies (DLB) from Alzheimer Disease.



COGNITIVE IMPAIRMENT: NON-PHARMACOLOGICAL INTERVENTIONS



Cognitive Stimulation Therapy

Level A evidence group programme involving themed activities to stimulate thinking, concentration, and memory. Focuses on maintaining cognitive function.



Physical Exercise

≥150 mins/week of aerobic and resistance training.
Improves cognition, reduces BPSD, and slows functional decline. Promotes overall well-being.



Music Therapy

Personalised playlists and structured sessions significantly reduce agitation, anxiety, and depression. Enhances mood and engagement.



Environmental Modification

Simplify the environment, use visual cues, ensure adequate lighting, and maintain consistent daily routines. Reduces confusion and falls risk.



Carer Skills Training

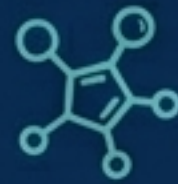
Equip carers with communication strategies and behavior management techniques (e.g., Dementia Australia's Living with Dementia). Reduces carer burden.



Occupational Therapy

Functional assessment, activity adaptation, assistive technology, and home modification via My Aged Care. Maximizes independence.

CHOLINESTERASE INHIBITORS (ChEIs): PBS AUTHORITY REQUIRED FOR MILD-TO-MODERATE AD

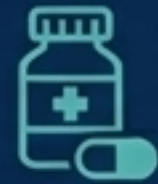


Donepezil

5mg nocte → 10mg

Hepatically Cleared

No renal adjustment required.
Key side effects include
nausea, diarrhoea,
bradycardia, and vivid dreams.



Rivastigmine

1.5mg BD → 6mg BD

Transdermal Patch Available

Patch formulation (4.6mg →
9.5mg/24h) offers better GI
tolerability. Also indicated for
Parkinson Disease Dementia.



Galantamine

4mg BD → 12mg BD

Renal Cautions

Avoid in severe renal
impairment (eGFR <30).
Watch for CYP2D6 and
CYP3A4 drug interactions.

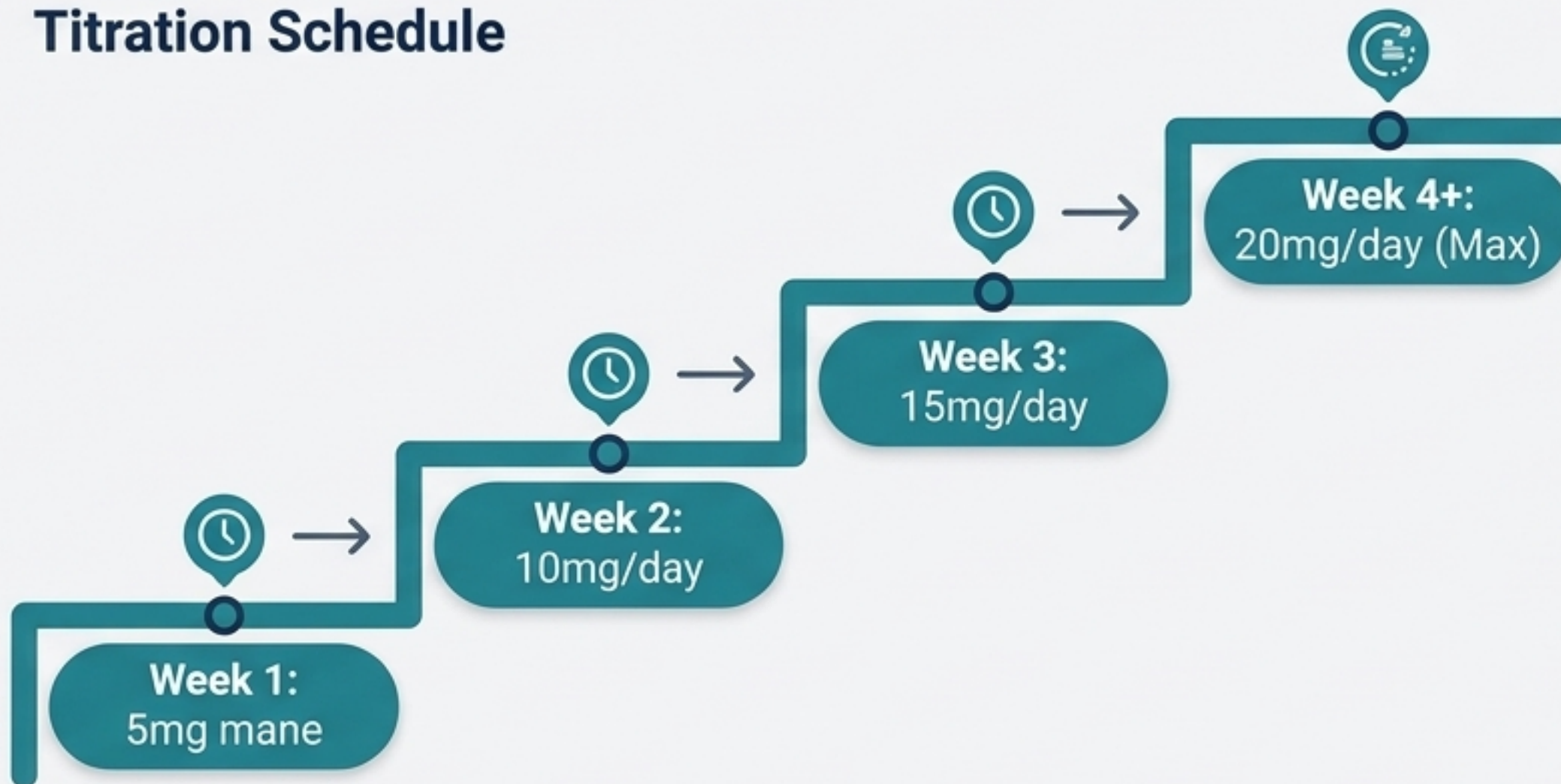


RULE OF THUMB: Start low, go slow. Prescribe with food to minimize GI side effects. Ineffective in bvFTD.

Memantine (NMDA Receptor Antagonist)

Indication: Moderate-to-severe AD (PBS Authority Required). Monotherapy or combined with ChEI.

Titration Schedule



Renal Adjustments & Tolerability

Renally cleared. Dose reduction essential to prevent toxicity.



eGFR 30–49: Max 10mg/day



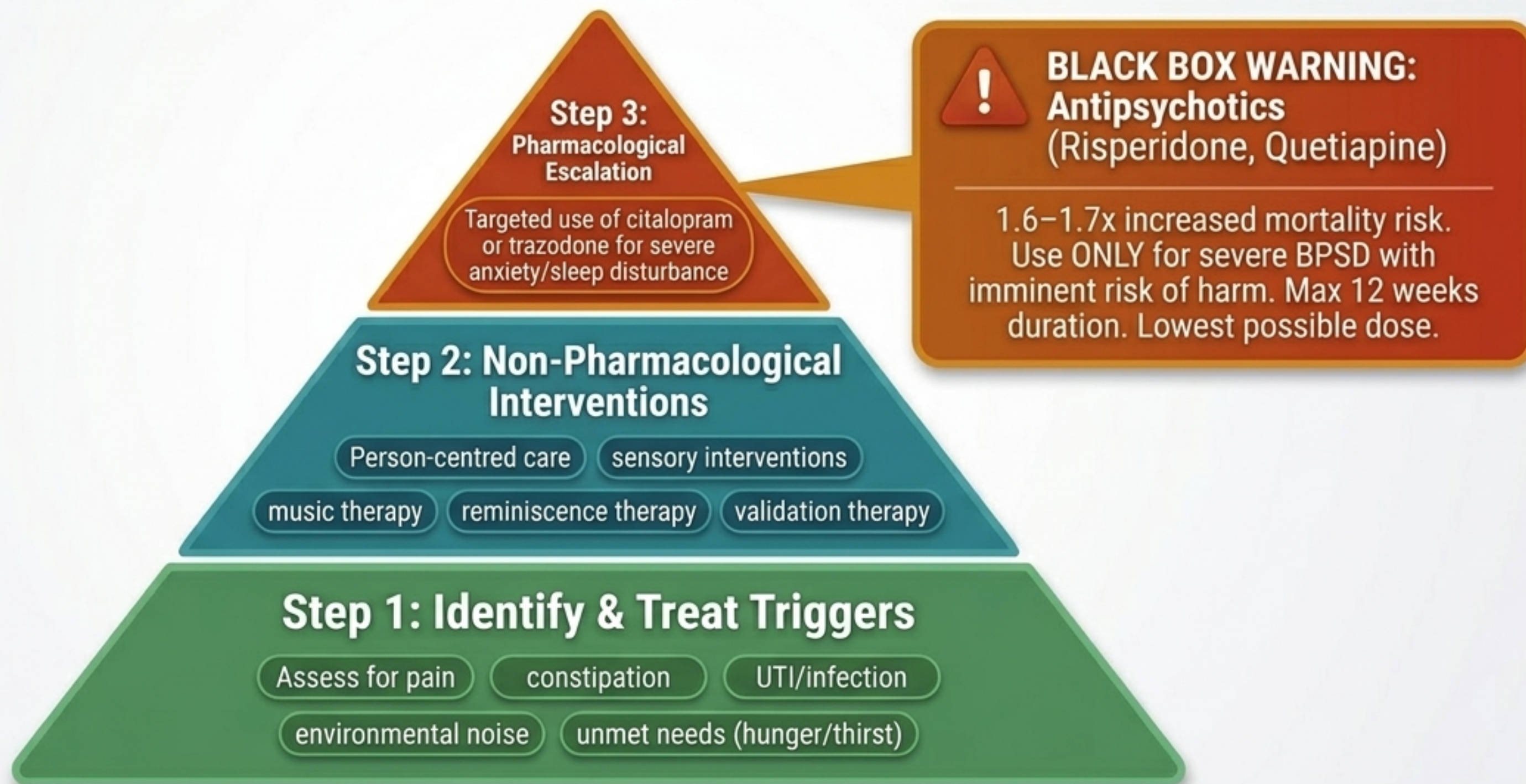
eGFR 5–29: Max 5mg/day



eGFR <5: Avoid completely

Fewer GI side effects than ChEIs. Watch for dizziness, headache, and somnolence.

BEHAVIORAL & PSYCHOLOGICAL SYMPTOMS OF DEMENTIA (BPSD) MANAGEMENT: STEPPED APPROACH





THE DRIVING MANDATE



AUSTROADS GUIDELINES:

All patients diagnosed with dementia must legally cease driving.



ACTION STEPS:

Advise patient and family, document discussion clearly in notes, advise transport authority notification.



IF DIAGNOSIS IS UNCERTAIN OR EARLY MCI:

Refer for formal Occupational Therapy driving assessment.



HOME SAFETY & ENVIRONMENTAL MODIFICATIONS



KITCHEN SAFETY:

Auto-shutoff stoves, hot water temperature limiters (50°C).



EXITS & WANDERING:

Door alarms, GPS tracking devices (Safe Return program) for wandering risk.



FALLS PREVENTION:

Comprehensive falls risk assessment, grab rails, removal of trip hazards.



MEDICATION MANAGEMENT:

Webster-paks or dosette boxes for secure medication management.

DEMENTIA SUPPORT NETWORK & ADVANCE CARE PLANNING

COMMUNITY RESOURCES & LEGAL FRAMEWORKS FOR DEMENTIA CARE



DEMENTIA AUSTRALIA

1800 100 500

Living with Dementia program

Counselling & Advocacy

Carer mental health screening (Zarit burden interview).



MY AGED CARE

1800 200 422

Home Care Packages

Centre-Based Day Respite

Residential Respite & Care.



ADVANCE CARE PLANNING

Initiate early while capacity remains

Appoint Enduring Power of Attorney (financial)

Appoint Enduring Guardian (health)

Register Advance Care Directive.

ABORIGINAL & TORRES STRAIT ISLANDER HEALTH & CULTURALLY SAFE CARE

ABORIGINAL & TORRES STRAIT ISLANDER HEALTH



RISK FACTORS & ACTION

High prevalence of modifiable cardiovascular risk factors (diabetes, hypertension, RHD).

Use Close the Gap PBS Co-payments.

CULTURALLY SAFE CARE



ASSESSMENT TOOLS:

Use KICA (Kimberley Indigenous Cognitive Assessment) or RUDAS.
Avoid standard Western cognitive tests that may contain cultural or educational bias.



CLINICAL APPROACH:

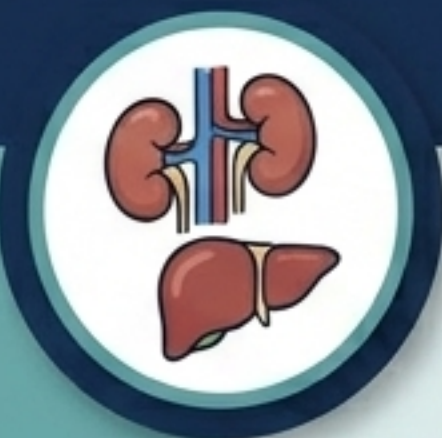
Utilize MBS Item 715 (Health Assessments) annually.
Engage Aboriginal Health Workers and liaison officers.



COMMUNITY CARE:

Support keeping Elders on Country rather than institutional care, which may be culturally inappropriate.

SPECIAL CONSIDERATIONS FOR COMPLEX CASES



RENAL & HEPATIC IMPAIRMENT

Donepezil preferred in renal impairment.
Reduce Memantine strictly in severe CKD.
Assess for uraemic or hepatic encephalopathy as reversible cognitive factors.



IMMUNOCOMPROMISED PATIENTS

Screen for HIV-associated neurocognitive disorder (HAND).
Monitor for progressive multifocal leukoencephalopathy (PML) or CNS infections in transplant or autoimmune patients.



PREGNANCY & YOUNGER ONSET

Category B3.
Cholinesterase inhibitors and Memantine are not recommended.
Provide genetic counselling for autosomal dominant familial dementias.

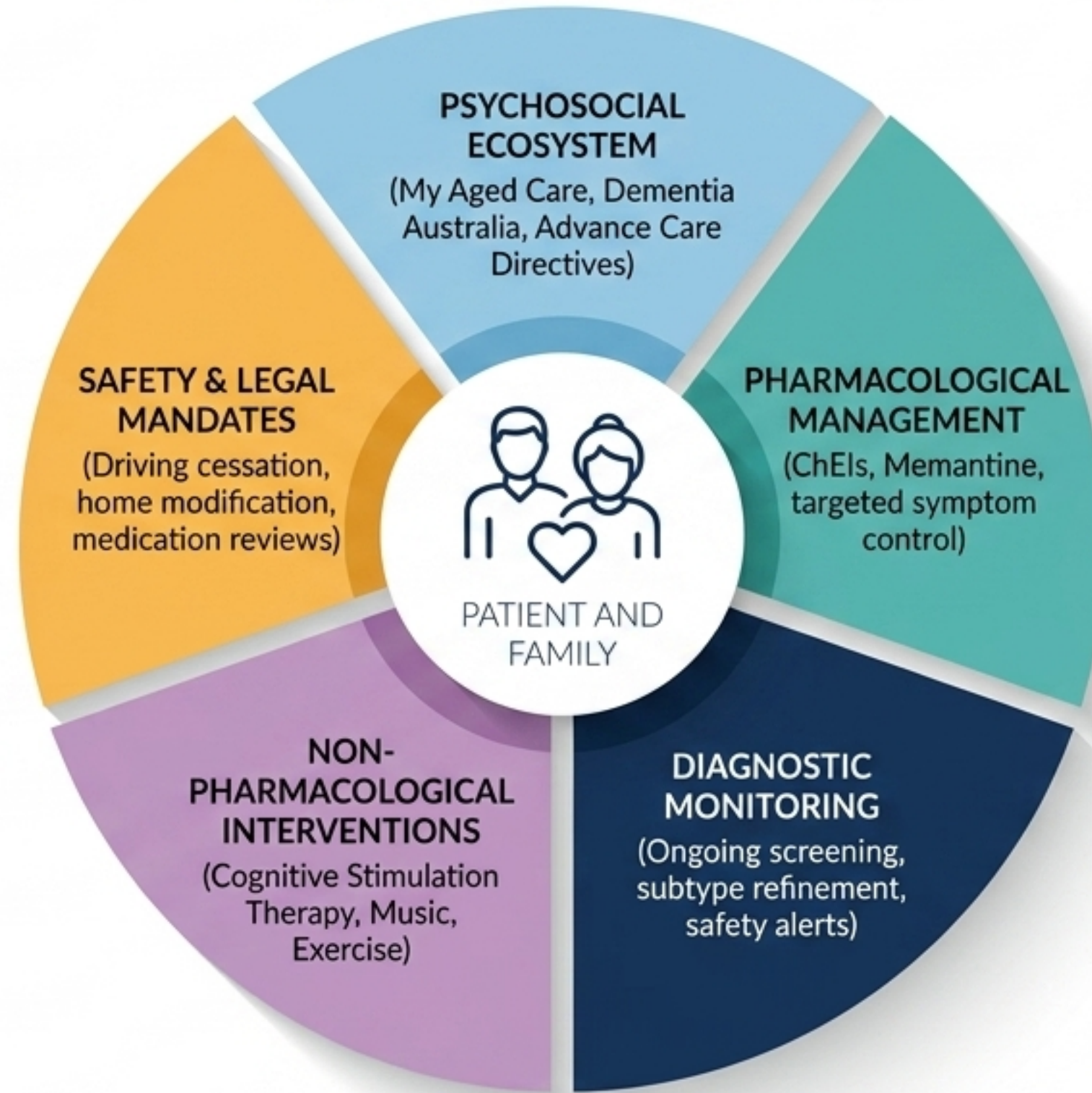


PAEDIATRIC DEMENTIA

Extremely rare (e.g., lysosomal storage disorders).
Requires urgent paediatric neurology referral.
Standard adult dementia medications are not routinely indicated.

DEMENTIA CARE ECOSYSTEM: COMPREHENSIVE, PATIENT-CENTRED MANAGEMENT

ECOSYSTEM OF CARE



DEMENTIA CARE EXTENDS FAR BEYOND A PRESCRIPTION. THE CLINICIAN'S ROLE IS TO ACT AS THE ARCHITECT OF A COMPREHENSIVE, PATIENT-CENTRED ECOSYSTEM.