

Dementia Echo

Determining Specific Dementia Diagnosis

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How to Diagnose Dementia

Clinical
Syndrome

History and
exam

Screening
Tests

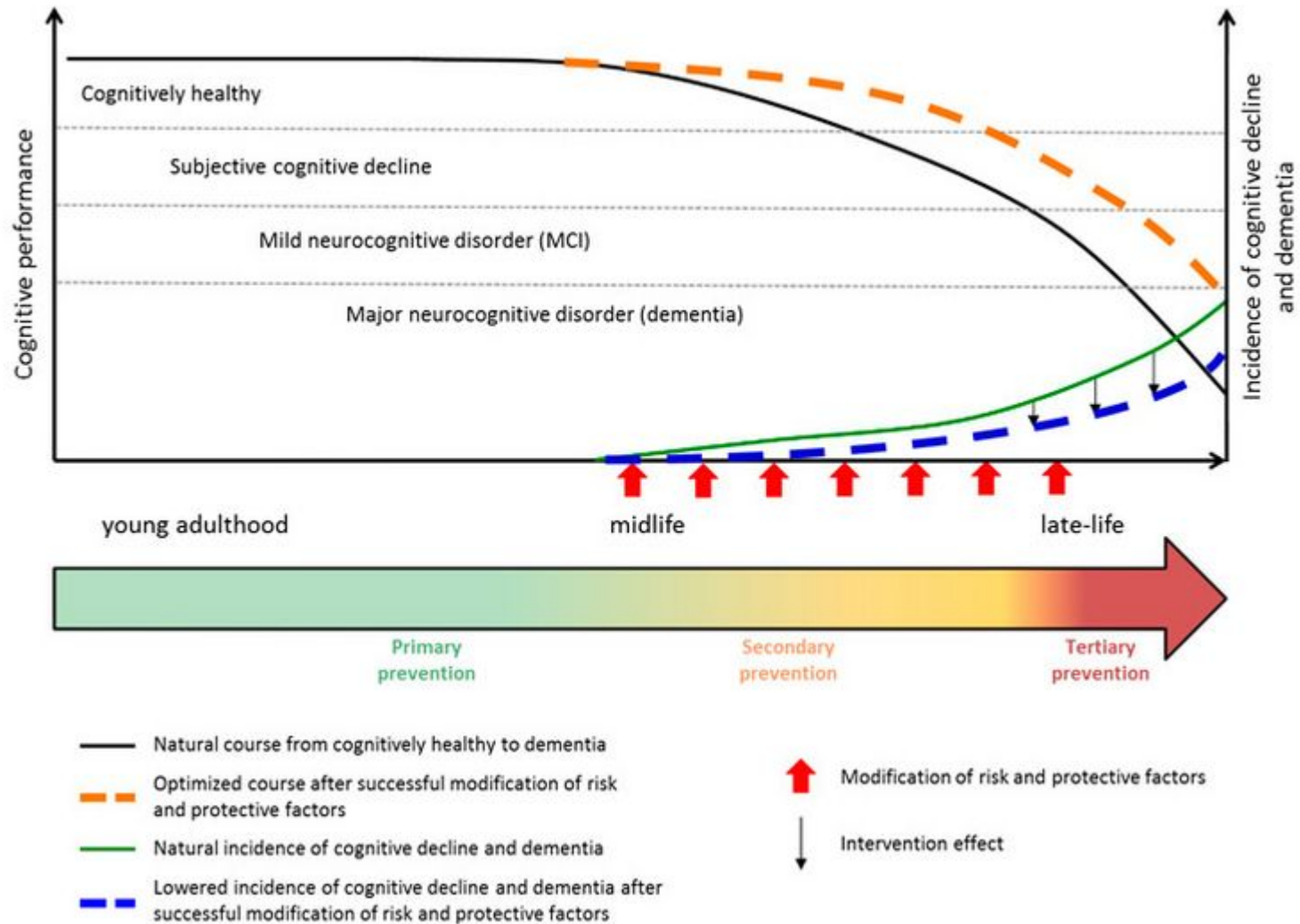
Investigations

Common
subtypes

Common
pitfalls

Questions

Cognitive Reserve Theory



Younger Onset Dementia

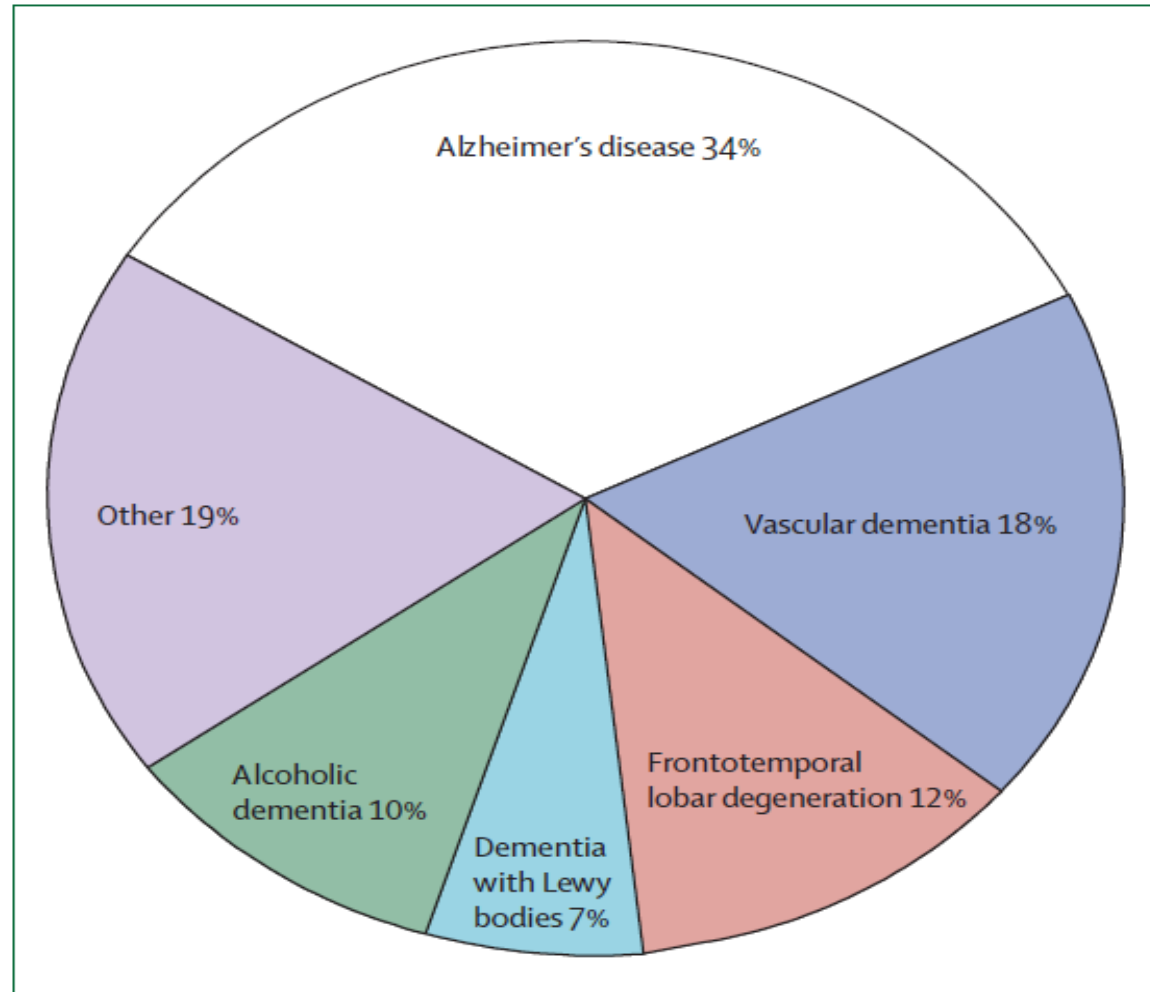


Figure 1: Epidemiology of young-onset dementia
Data from the community study of Harvey and colleagues.⁶

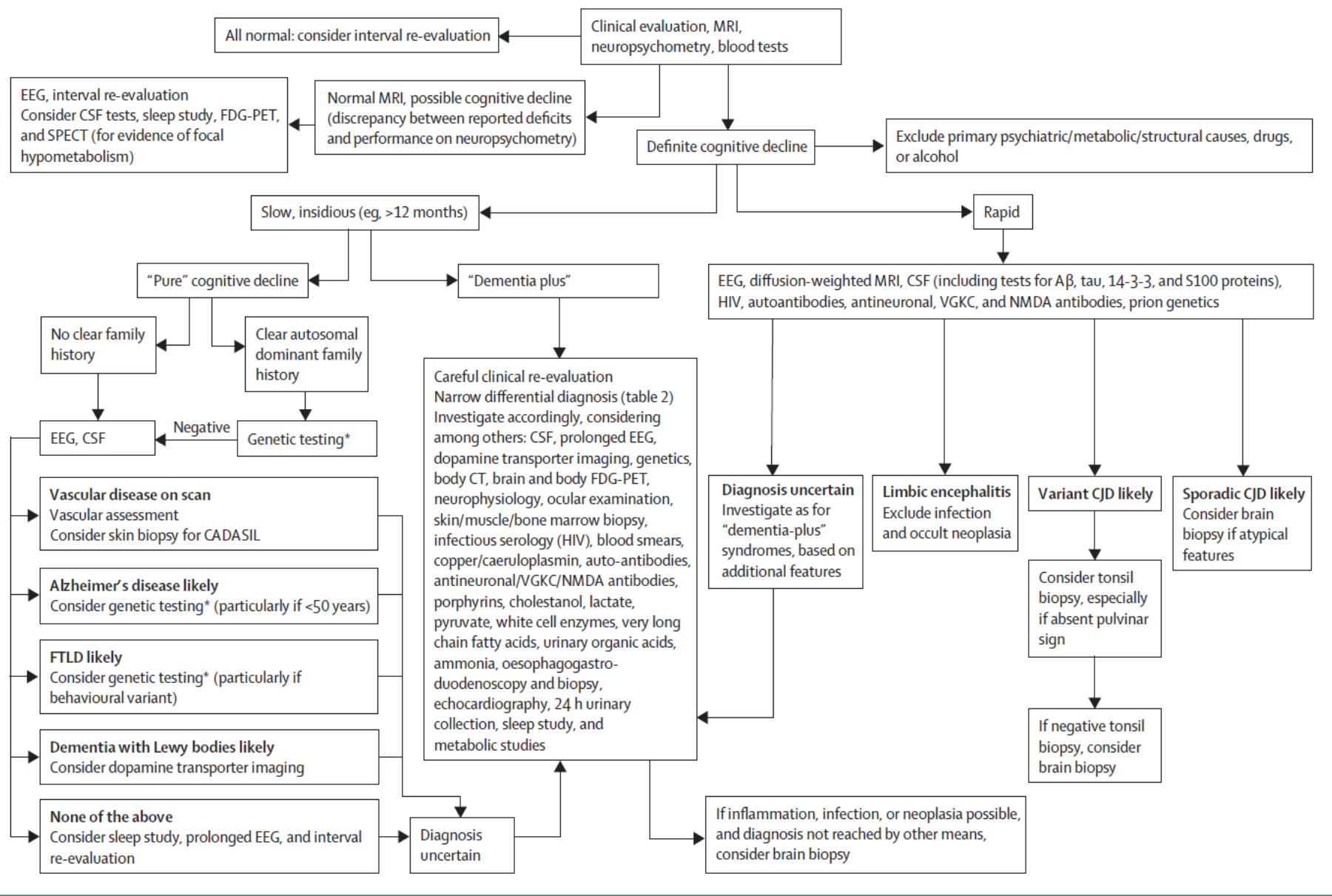


Figure 3: Flow chart for assessment and investigation of young-onset dementia

This algorithm provides an overview of the diagnostic approach to patients with young-onset dementia. Given the many causes, this can only act as a general guide. *In amnesic young-onset dementia, first-line genetic testing is for *APP*, *PSEN1*, *PSEN2*, and prion. In behavioural cases, first-line testing is for *MAPT* (particularly if symmetrical atrophy on MRI) and *GRN* (particularly if asymmetric pattern of atrophy). EEG=electroencephalogram. FDG=fluorodeoxyglucose. SPECT=single photon emission computed tomography. Aβ=amyloid β. VGKC=voltage-gated potassium channel. FTLD=frontotemporal lobar degeneration. CADASIL=cerebral autosomal dominant arteriopathy with subcortical infarcts and leukoencephalopathy. CJD=Creutzfeldt-Jakob disease.

Dementia Diagnosis

- A. Evidence of significant cognitive decline from a previous baseline in one or more cognitive domains
- B. Cognitive deficits interfere with independence in everyday activities
- C. Deficits are not exclusively due to delirium
- D. Not better explained by another mental disorder

Diagnosis

- Patient and informant history
- Brain imaging- CT, MRI, PET MRI
- Labs- FBC, Chem 20, B12, TFT, HIV, Syphilis, ESR
- Screening tools- AMT, MMSE, MOCA, RUDAS, KICA, ACE111
- Neuropsychology

- <https://qheps.health.qld.gov.au/caru/networks/cognitive-impairment-toolkit>
- QHEPS- Cognitive impairment toolkit

Taking the History

- Begins in the waiting room!
 - Patient and a collateral history
 - ? Structured questionnaire
 - Begin with a conversation- open ended questions...
-
- Baseline
 - Cognition
 - Function
 - Medications
 - Psychiatric/ D&A hx
 - PMH

Cognitive Domains

Orientation and concentration

Working/ episodic memory

Remote memory

Language

Visuospatial

Executive function

BPSD

- Aphasia- loss or impairment of language due to brain damage
- Dysarthria – disorders of articulation (central or peripheral)
- Dyslexia- disorders of reading
- Agnosia – non recognition eg visual agnosia

Alzheimer's Disease(AD)

- Neurodegenerative disease
- Incidence increases with age
- Prominent working memory issues- rapid forgetting
- Cannot learn new information but remembers info from long ago
- Loses remote memory with time
- Language +/- visuospatial impairment
- Progresses over time 8-12 years

- MMSE

Vascular Dementia(VaD)

- More prominently effects executive function/ decision making
- Stepwise progression
- Can present as “flat”/ depressed or emotional labile
- In the presence of vascular risks- HTN, Stroke, IHD, DM etc
- Often mixed with AD
- Vascular gait in later stages
- MOCA

Lewy Body Dementia/ Parkinson's Dementia

- Less common
- All pts with PD eventually develop dementia
- Sleep disorders, constipation
- Fluctuations are prominent
- Early hallucinations – visual or auditory esp
- Early movement disorder- unilateral tremor, rigidity, bradykinesia (Slowing down) and falls
- Neuroleptic sensitivity
- Memory can be relatively preserved- screening tests can be N

Frontotemporal Dementia – Behavioural variant

- Less common
- Early change in behaviour and personality
- Loss of empathy or disinhibition
- Apathy and inertia
- Neglect and poor self care
- Stereotypic behaviours
- Change in eating habits
- Can look like psychiatric presentation
- Memory can be relatively preserved in early stages
- Cross over with MND

Frontotemporal Dementia- Language Variant

- Early language decline with function and memory preserved
- Testing looks worse than the patient presents- MMSE <15 but functional and continent
- Testing and dx difficult
- Semantic dementia- loss of memory for words
- Progressive non-fluent aphasia- loss of output, grammar, telegraphic speech
- Logopenic progressive aphasia- AD variant

Dementia Mimics

- Depression
- Untreated psychiatric disease
- Intellectual impairment
- Poor educational achievement
- CALD
- Medications
- Delirium
- Acquired brain injury

Things to look out for- Specialist review suggested

- Younger onset <65 year
- Rapidly progressive
- Early BPSD
- Early Language involvement or movement disorder
- Positive family hx- >2 relatives <65 years

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