

DEMENTIA

- The diagnosis of dementia types
- Recognise the behavioural and psychological symptoms associated with dementia
- Identify the risks of poly-pharmacy in a patient with dementia
- Compare pharmacological and non-pharmacological interventions for patients with dementia

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Consultant Physician Geriatrician SALHN

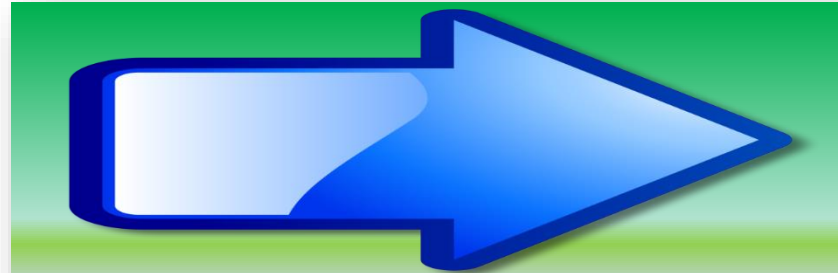
Dementia Prevalence Epidemic

2020

38,400 South Australians

2028

47,300



Dementia in Australia

459,000
living with
dementia



27,800
living with
younger onset
dementia

WHY?

Multiple Potential Pathways to Dementia

LIFESTYLE FACTORS

Physical Activity

Diet

Drug/Alcohol Abuse

ENVIRONMENTAL FACTORS

Education

Head Trauma

Toxins/Other

PSYCHOSOCIAL FACTORS

PTSD

Depression/Anxiety

OTHER MEDICAL RISKS

Metabolic / Obesity / Diabetes

Hypertension / Heart Disease / Stroke

Inflammation

Certain Infectious Diseases

Certain Medications



HEALTH DISPARITIES FACTORS

AGING

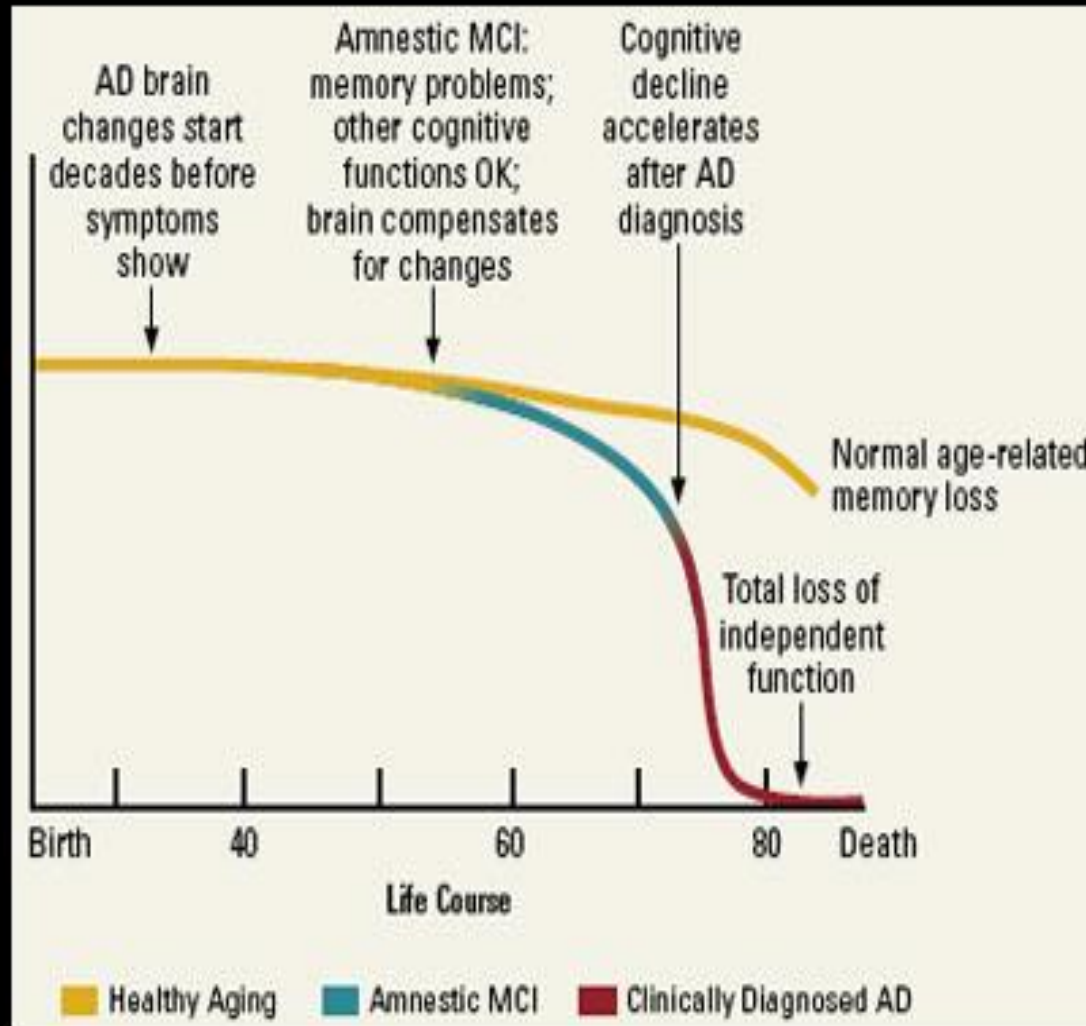
GENETIC FACTORS APOE4 PSEN1 MAPT TDP-43

SEX F>M

DEMENTIA TYPES



Disease Progression



Rate of progression

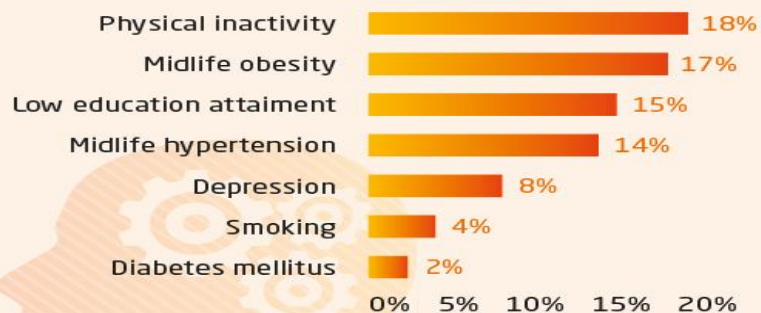


■ slow progress

■ fast progress

Risk Factors for Dementia

Proportion of dementia cases attributable to different risk factors



Source: Adapted from Ashby-Mitchell et al. (2017)

Resilience to Dementia

Factors that predict resilience to dementia

Female only

- Living with someone
- Being married
- Lower pulse pressure
- Higher peak expiratory flow
- Faster walking time
- Faster turning time
- Volunteering more often

Male only

- Less depressive symptoms

Both genders

- Younger age
- Higher education
- Stronger grip
- More cognitive activity

Source: Adapted from McDermott et al. (2016)

Rising Cost of Dementia

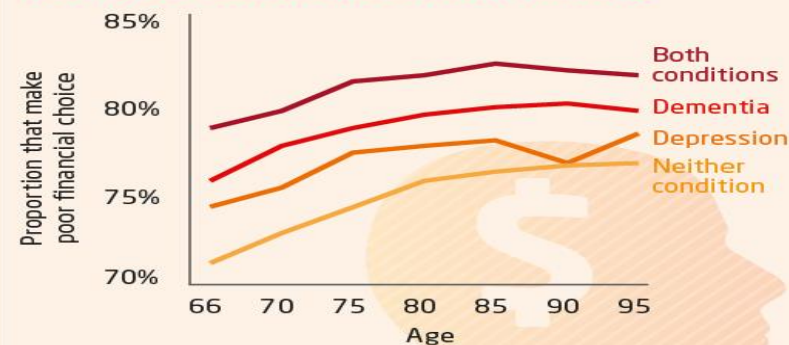
Total direct and indirect costs of dementia



Source: NATSEM calculation

Financial Fragility

Dementia and depression are correlated with poor financial decision-making

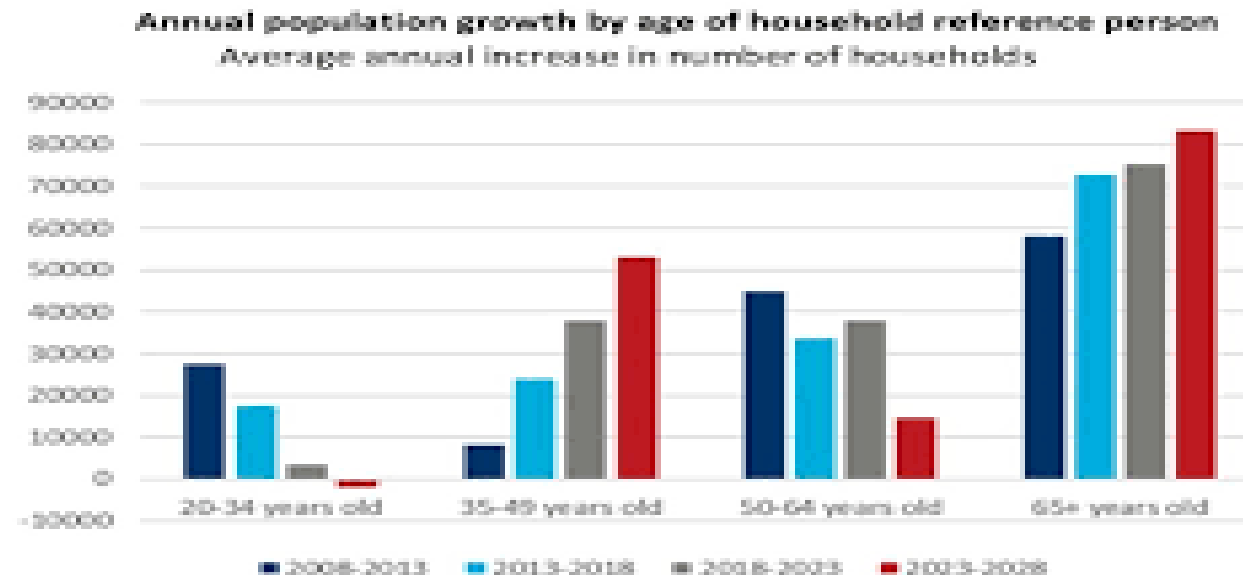


Data displayed at 5-year intervals. Source: Keane et al. (2017)

Hospital care for people with dementia 2016–17

- 43% of hospitalisations and 50% of patient days were for people aged 65 and over

In 2017, over 1 in 7 Australians were aged 65 years and over



Who is most likely to be hospitalised with dementia?

- Male hospitalisation rates were 1.3 times higher Female
- People aged 95+ were 5 times of 75–79yrs.
- Dementia overall fell by 23% (408 to 313 hospitalisations per 100,000 population)
- **Vascular dementia rose by 37%**
- **Delirium superimposed on dementia rose by 416%**

Australian Journal of **DementiaCare**

For all who work with people with dementia

Vol 4 No 2 April/May 2015

The quest for person-centred care

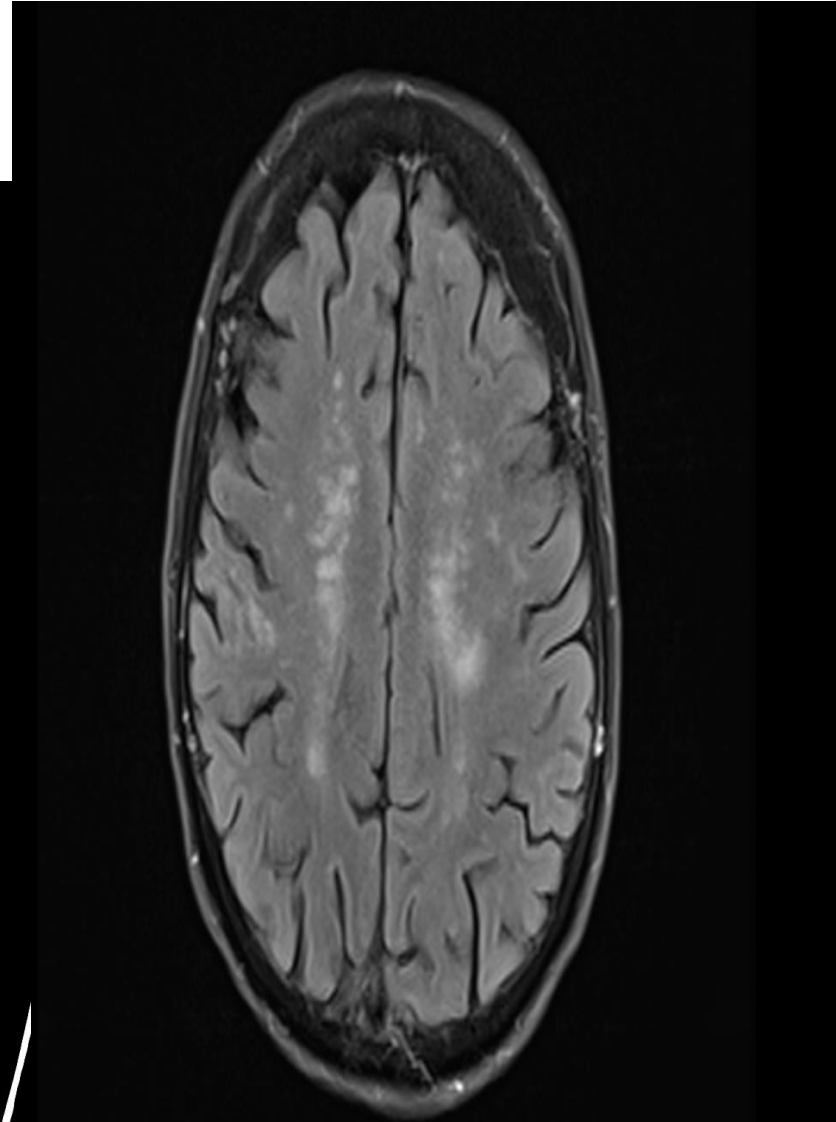


**Also inside
this issue:**

- Fun and playfulness ■ A new communication tool
- Frontotemporal dementia ■ One-page profiles

Diagnosis + Type of Dementia

- Clinical Symptomatology
- Cognitive testing
- Imaging
- Genetic
- Autopsy
- Blood test



Structural Scan

MRI

Volumetric figures

Diffusion tensor imaging

Stroke SOL detection

white matter tracts

Common measurements :

Global brain volume,

White/gray matter volume,

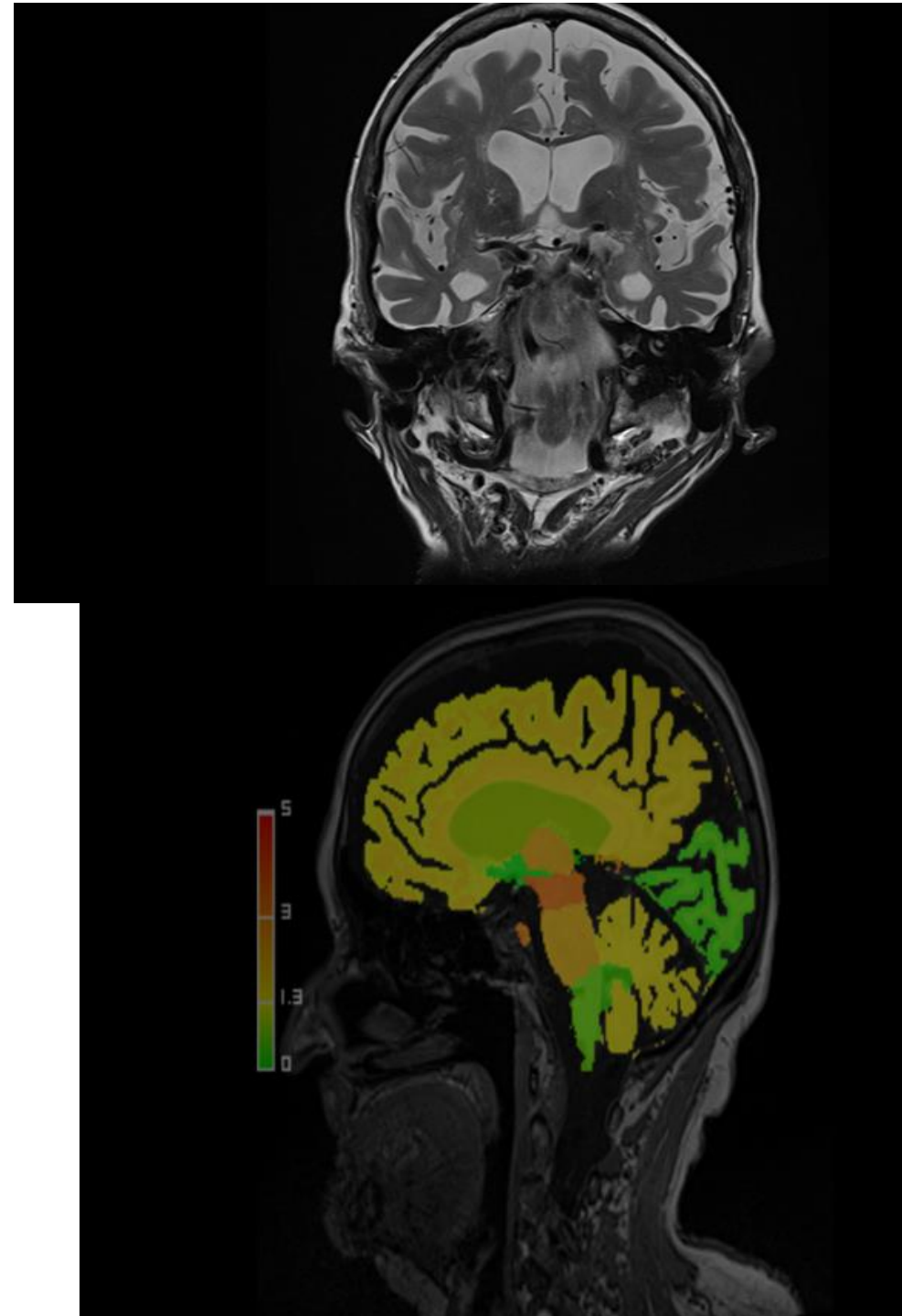
Cortical thickness,

Amygdala

cingulate cortex volume,

hippocampal volume

(decreased in MDD and schizophrenia)



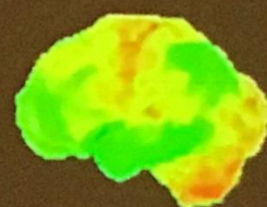
Multimodal Biomarker-PET/MR differential diagnosis of AD

Multiparametric MRI

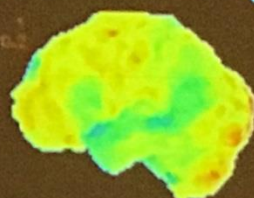
Tumors, vascular, inflammat.
& structural changes

Amyloid-PET

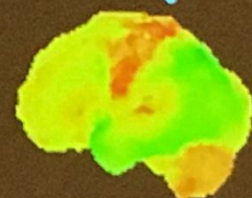
Neuronal injury



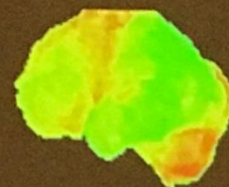
AD



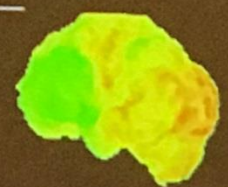
lvPPA



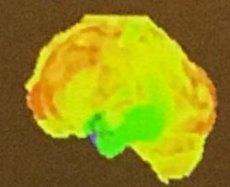
PCA/DLB



DLB

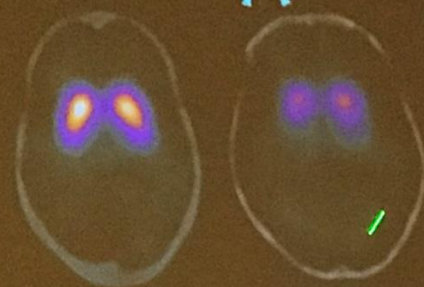


bvFTD

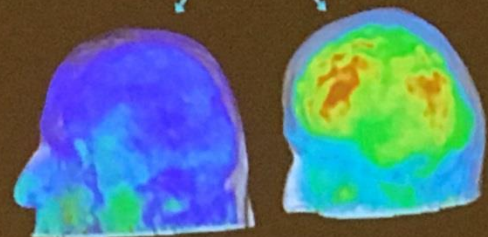


nfv/svPPA

ADD on



DaTSCAN



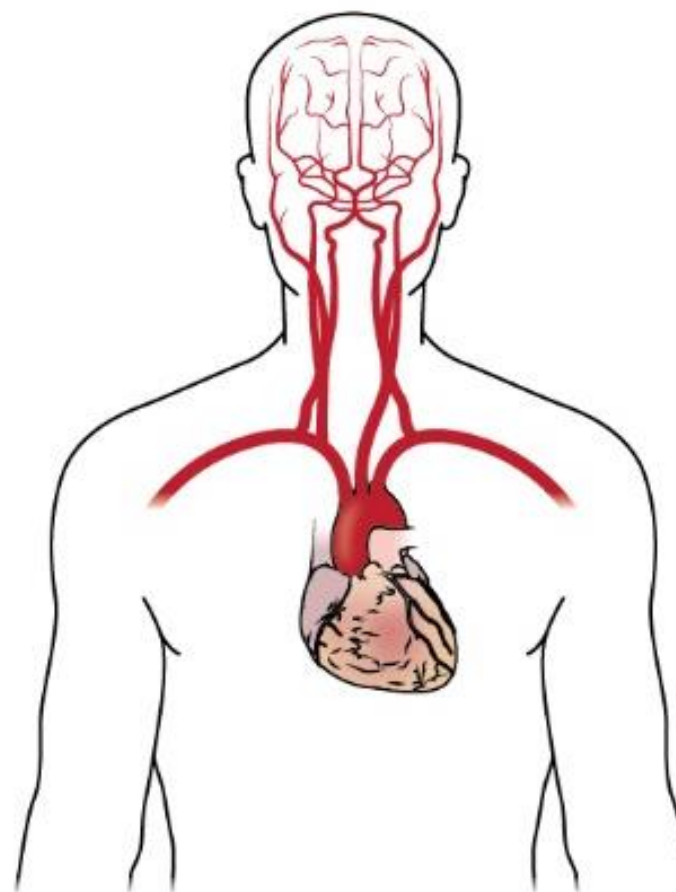
Tau-PET

AD, DLB

DLB, FTLD

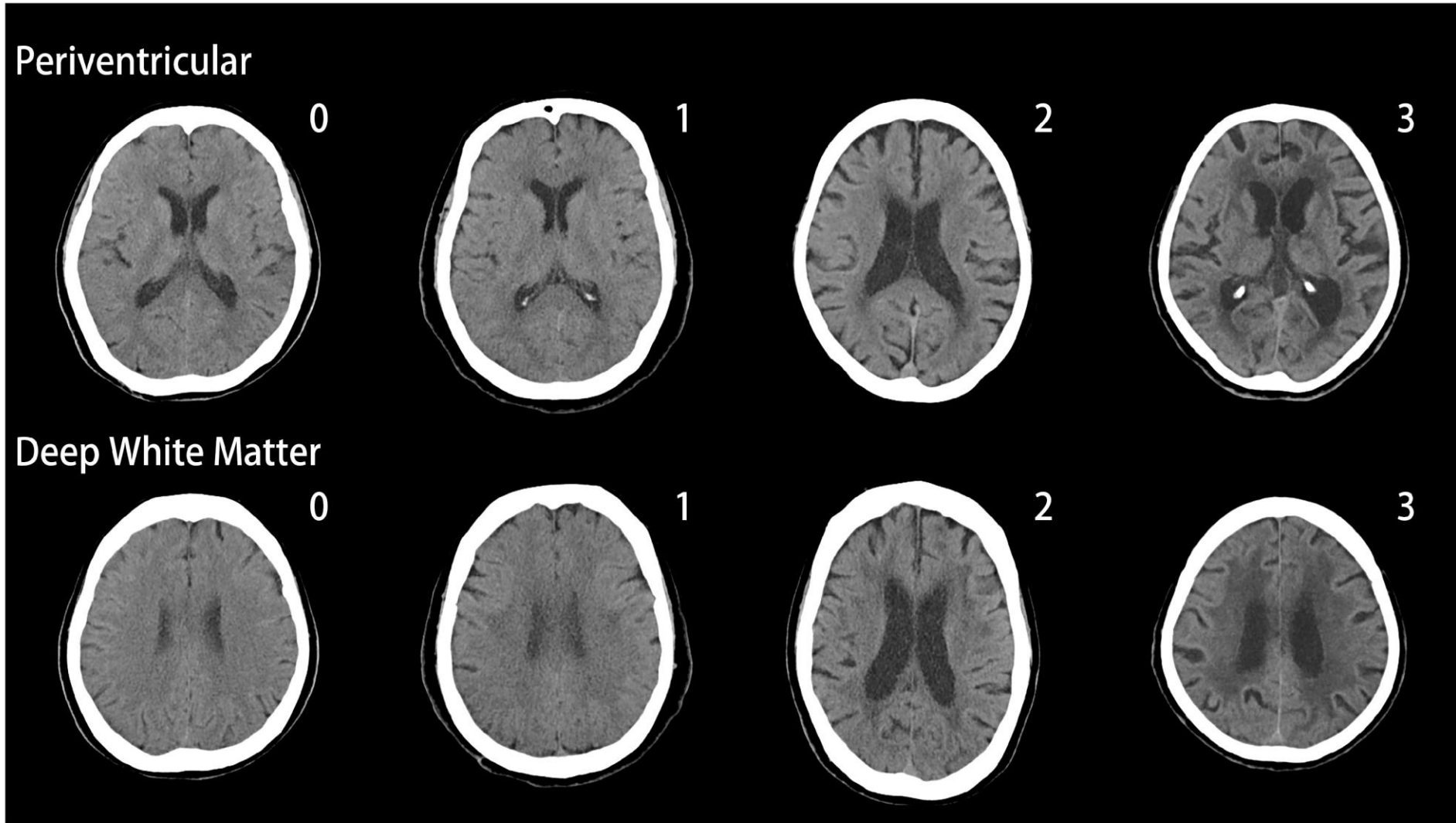
Vascular Contributions to Cognitive Impairment and Dementia (VCID)

Cognitive impairment	Micro-infarct
	Micro-bleed
	Silent stroke
	Cardiac disease
	Transient ischemic attack (TIA)
	Small vessel ischemic stroke
	CADASIL
	Small vessel hemorrhagic stroke
	Cerebral amyloid angiopathy (CAA)
	Large vessel ischemic stroke
Dementia	Large vessel hemorrhagic stroke

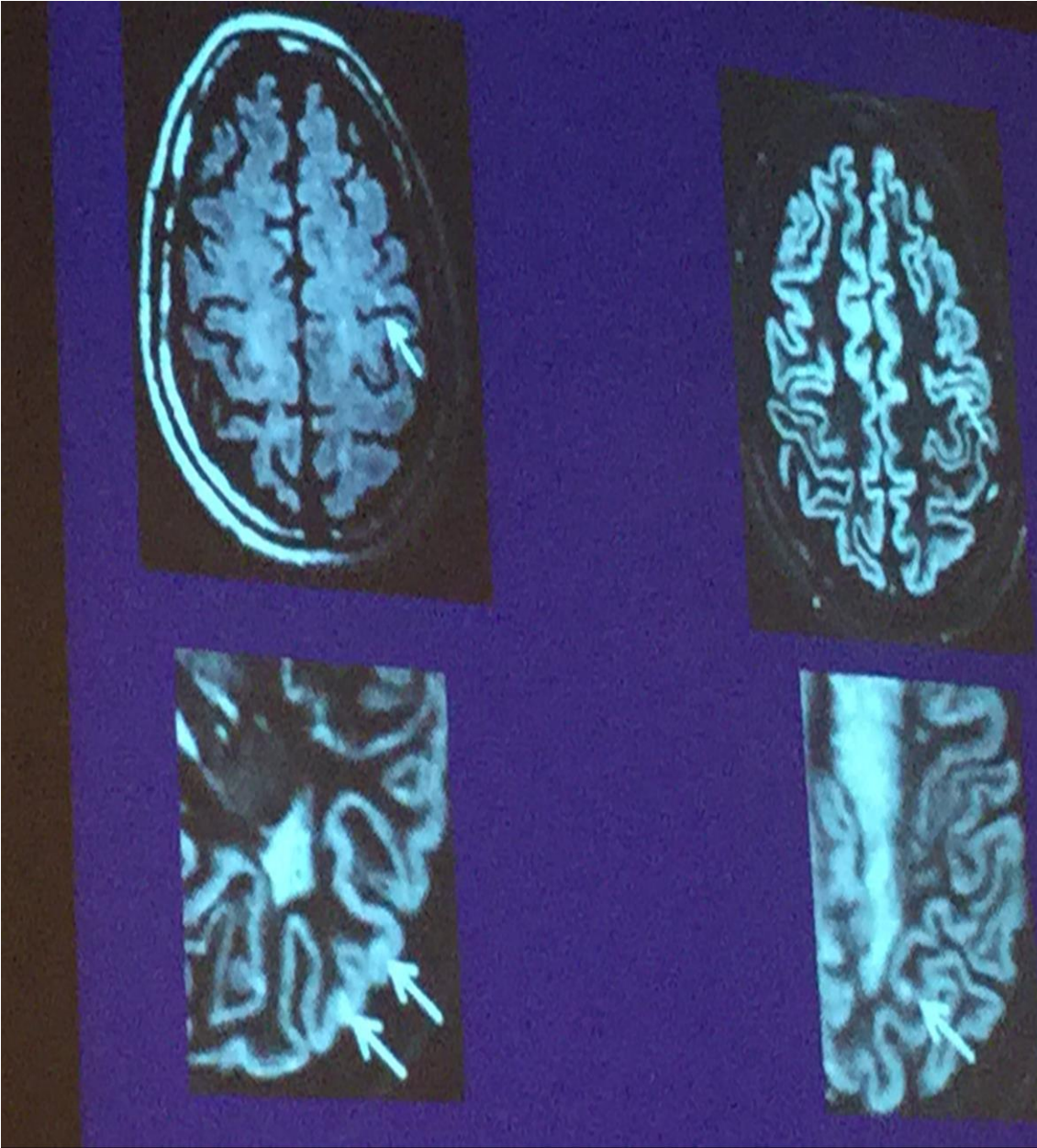


Small vessel disease- Subcortical degeneration

White matter changes modified Fazekas



The right diagnosis



- Suitability for Memory enhancers
- Planning for the future
- Affairs management
- Bucket list in order
- Genetic testing

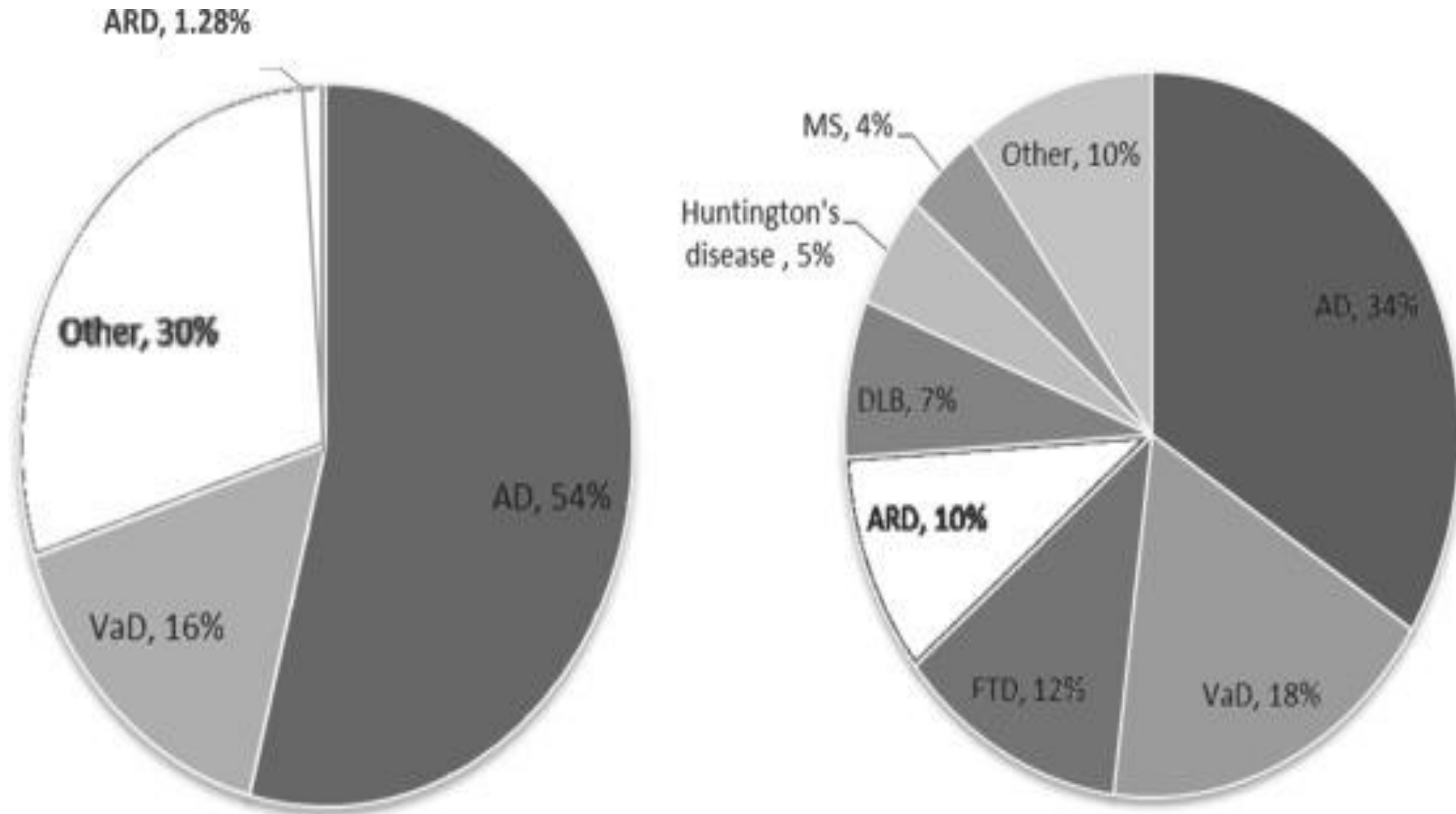
CN, 79 year-old, MMSE 30→29, interval 18.0 months



SUVr: 1.0

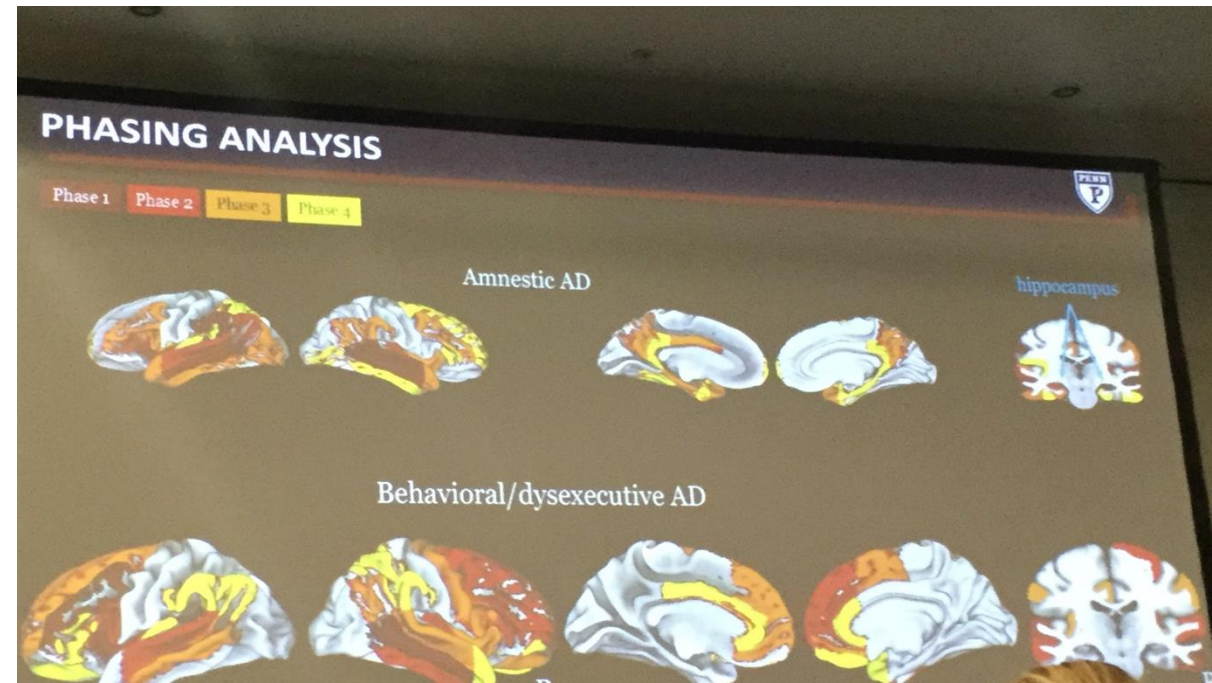
2.0

Alcohol-Related Dementia: A Systemic Review of Epidemiological Studies Jul2017



The near future Diagnosis

Tammy, said the PrecivityAD™ blood test would have saved her a decade of “futile trips to doctors who chalked up her symptoms to depression, anxiety or menopause before a \$5,000 brain scan last year finally showed she had Alzheimer’s.”



Pseudo-dementia

Pseudo-pseudo-dementia

Pseudo-Depression

- Overlapping Low Mood
- Beliefs of worthless # Indifference
- Behavioural passivity/ refusal
- Appetite, Sleep, Libido changes
- Anxiety
- Response to antidepressant
- Fluctuation of inactivity

Atypical Alzheimer's

Cholinergic deficiency +

Posterior Cortical atrophy

- Visual clarity
- Verbal and written language deficits
- Sit to stand
- Complex tasks problems

Frontal variant AD TDP-43

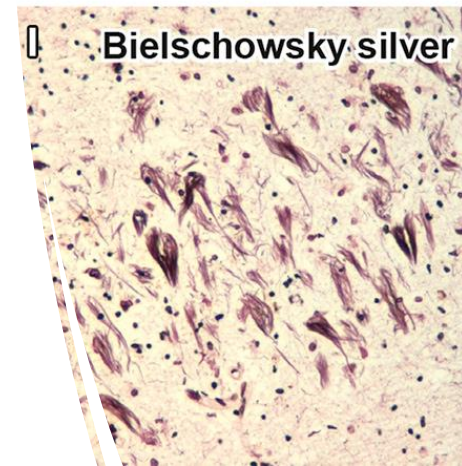
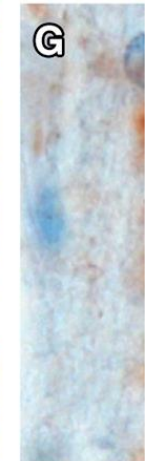
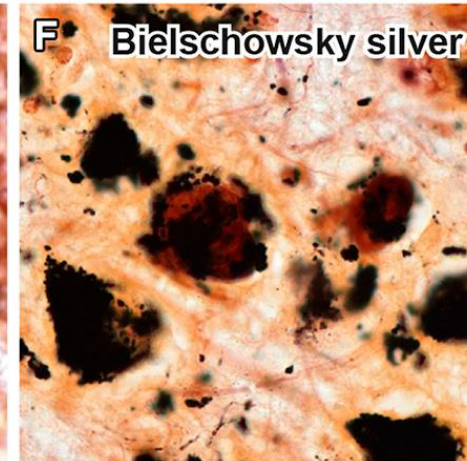
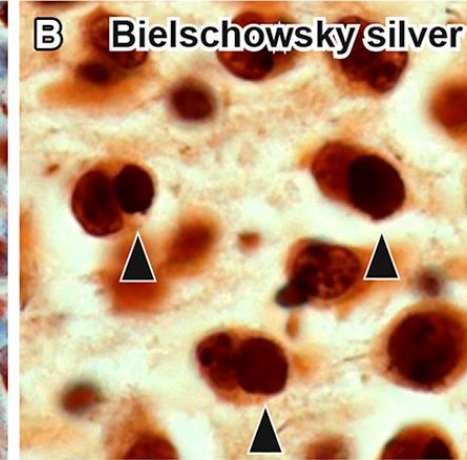
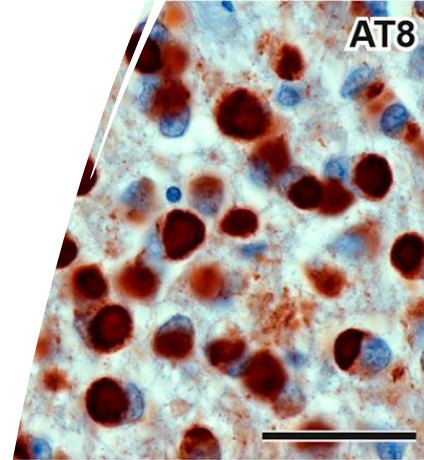
- Aberrant motor behaviour
- Apathy

Lewy Body D

- Anxiety
- Irritability
- Hallucinations
- Sleep disturbance

FTLD-tau types

- bvFTD
- IFTD
- PiD
- CBS
- PSP:P, RS, C types
- Pure Akinesia Gait failure
PAGF
- Progressive Non Fluent
Aphasia nfvPPA
- SvPPA
- IvPPA



Pathology of FTLD syndromes

Relationship between Clinical Features, Pathology and Proteinopathies in FTLD sub-syndromes

[modified from Cairns et al, 2007 and Josephs 2008]

	Frontotemporal Lobar Degeneration Syndromes							
Sub-syndrome	bvFTD		SD	PNFA		FTD-MND	FTDP-17	
							[MAPT or GRN mutations]	
Clinical features	Predominantly behavioural		Predom. language	Predominantly language		Behavioural Language + MND	Behavioural Language +/- EPS	
Histopathology	PiD-CBD-PSP-types	FTLD-U types 1-3	FTLD-U types 1-3	PiD-CBD-PSP-types	FTLD-U types 1-3	FTLD-U types 1-3	Tau +ve neuronal / glial inclusions	FTLD-U type 3
Molecular pathology	Tau	TDP-43	TDP-43	Tau	TDP-43	TDP-43	Tau	TDP-43

Pick's disease



Atypical Cognitive Symptoms

- Behavioural Psychological symptoms

Bradyphrenia

Slow responses

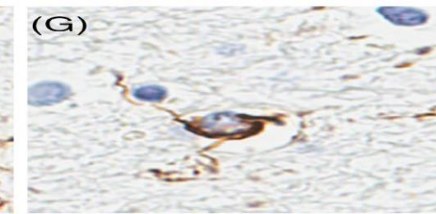
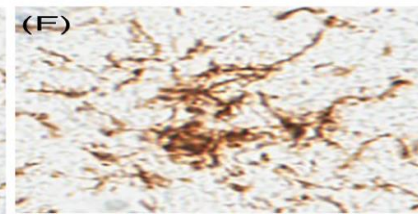
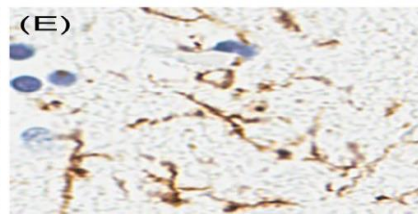
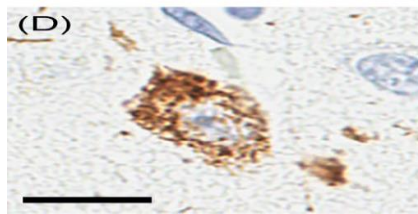
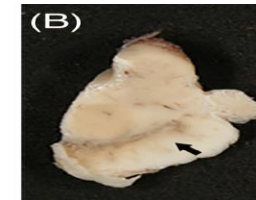
Memory recall

Language disorder

Disinhibition

Apathy -Inertia

Motor deficits/



COGNITIVE + NON-COGNITIVE SYMPTOMS

Short
term

Memory

Calculation

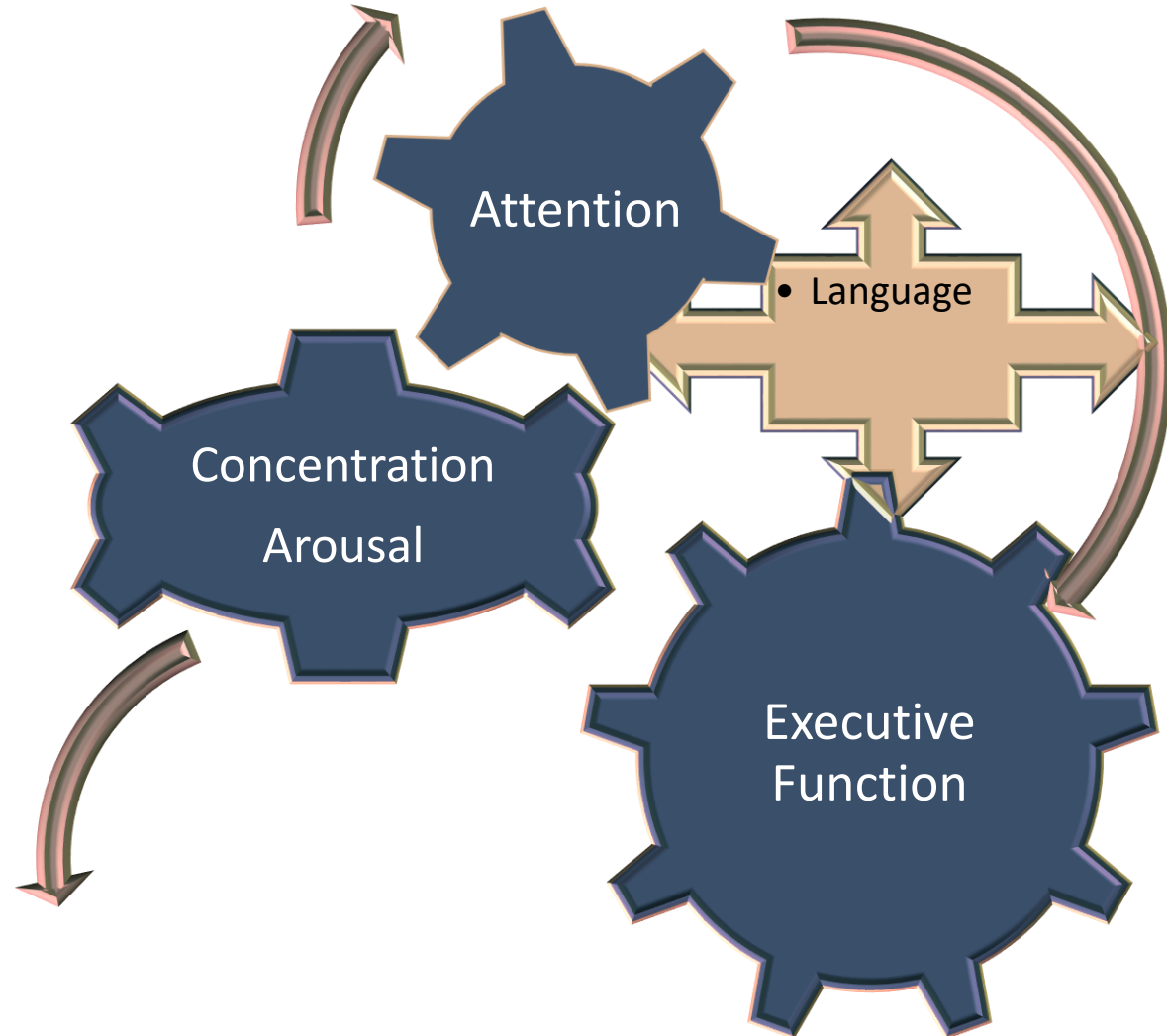
Multitasking

Visuo-spatial

Verbal recall

Abstract thinking

Verbal fluency



Cognition variability

- Characteristics of a person actions and reactions with the environment

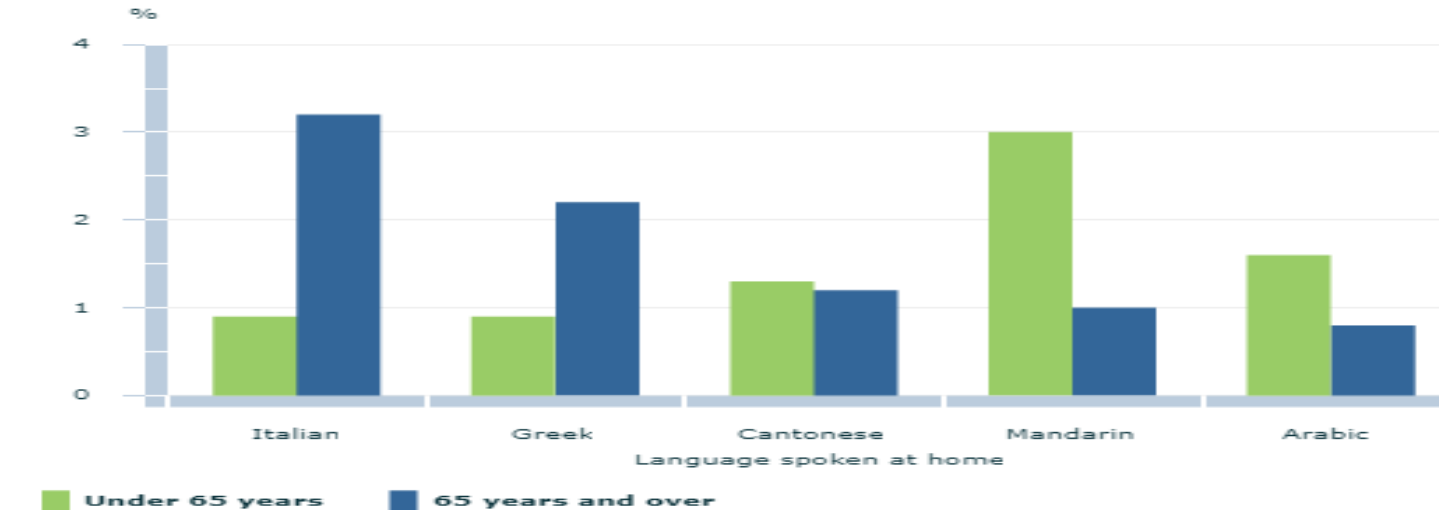
Genetics

Developmental opportunities

Culture

Era

Top five languages spoken at home(a), Older people, 2016



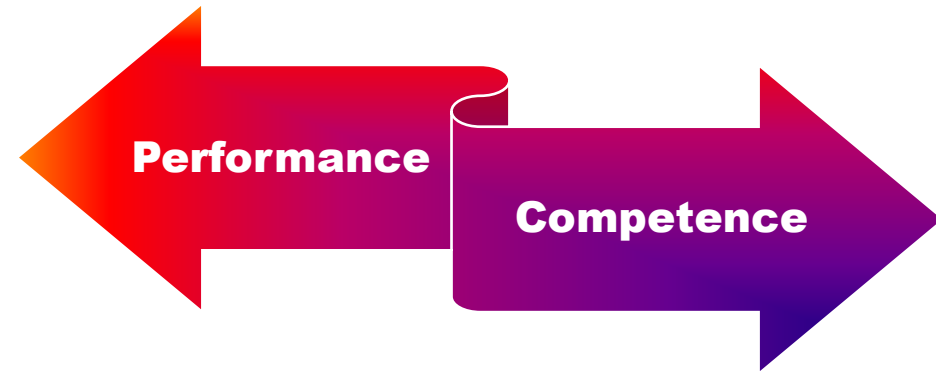
Save Chart Image

Australian Bureau of Statistics

© Commonwealth of Australia 2019.

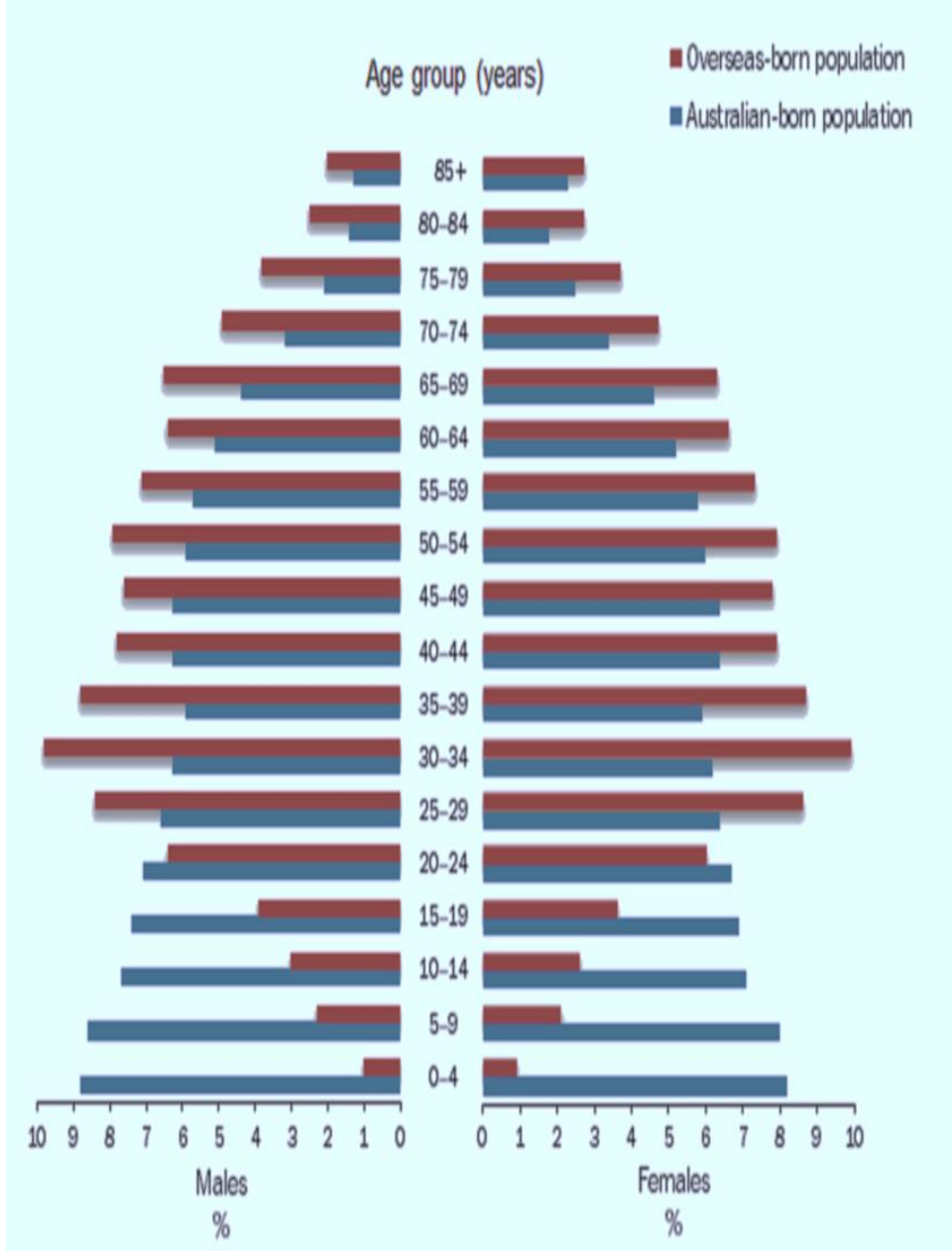
Diagnostic Tools for Cognition

- MMSE CLOCK ADAS-Cog Adenbrook MOCA CAMCOG Digit test
- Reliable
- Easily Applicable
- **Quick**
- **Sensitive**
- **Specific**
- **Simple training**

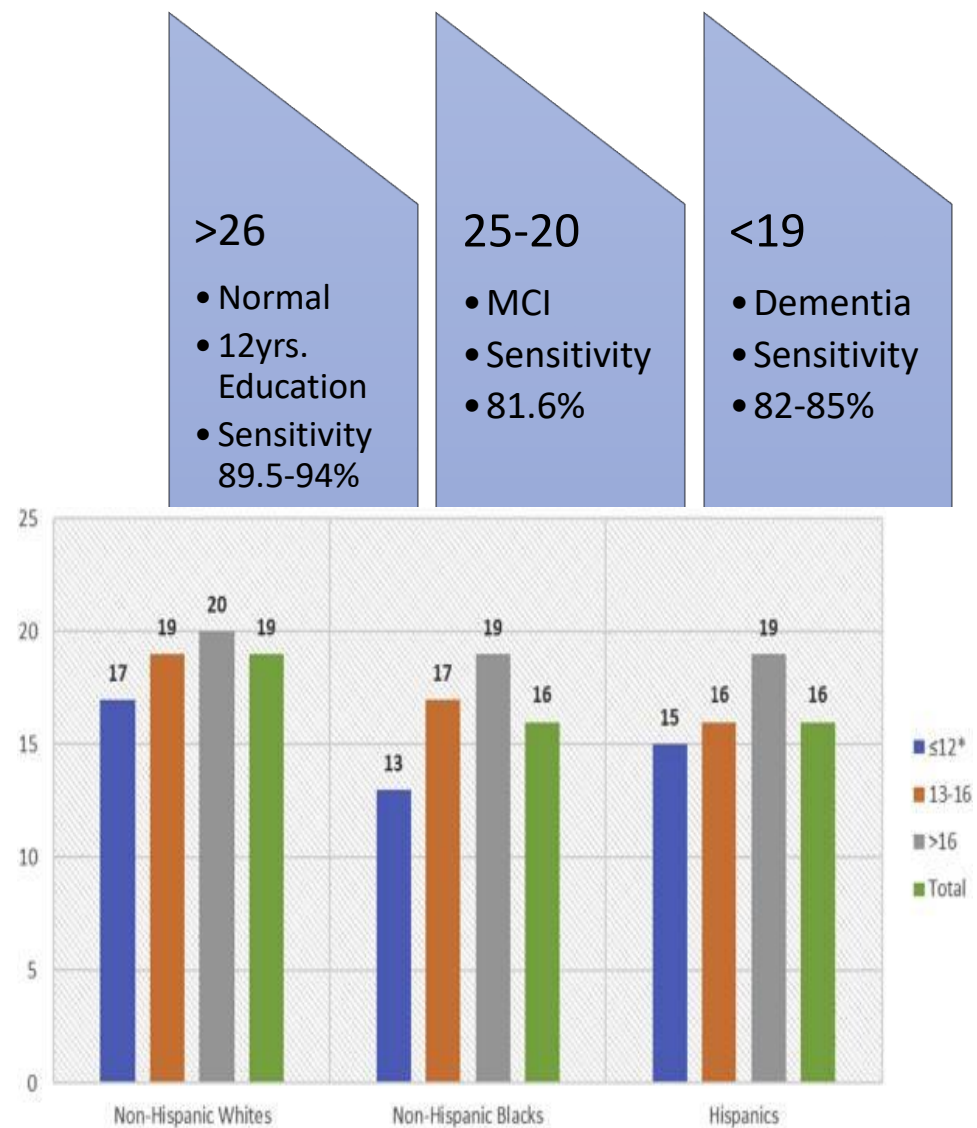


Scoring / Performance

- Educational Level
- Cultural setting
- Skills
- Emotional status
- Sensory problems
- Physical problems
- Neurological problems



SCORE	SEVERITY	STAGE
30-26	Could be normal	Could be normal
25-20	Mild deficits	Early Dementia
19-10	Moderate to severe	Middle to later
9-0	severe	late



7

Stages of Alzheimer's Disease and its Symptoms

Stage One

- No noticeable signs of deficit or impairment
- An evaluation with a physician would likely result in no diagnosis of the disease



Stage Two

- Mild loss of short-term memory. Small but noticeable lapses may present as age-related (momentarily forgetting where you parked your car)
- Minor loss of language skills – choosing incorrect words, trouble recalling words or names of familiar places

Stage Three

- Increased language deficit – difficulty choosing the correct word and remembering people's names
- Difficulty performing social or work-related tasks that were previously very easy or routine
- Forgetfulness – frequently losing objects and valuables, forgetting important appointments and dates
- Trouble planning and organizing
- Difficulty with directions when driving

Stage Four

- Heightened forgetfulness – trouble recalling recent events
- Loss of ability to perform mental math
- Trouble paying bills, performing daily social tasks, planning, and organizing events
- Forgetting personal history and past events
- Changes in mood – may exhibit antisocial behaviors, depression, anxiety, or mild irritability



Stage Five

- Noticeable lapses in memory and functioning
- Inability to recall important personal details such as personal phone number, address, place of work, alma mater, etc
- Confusion in day-to-day life – such as forgetting where they are and what day it is
- Trouble computing simple arithmetic



Stage Six

- Loss of awareness of surroundings and experiences
- Changes in sleep patterns – sleeping during the day and being awake at night
- Inability to dress themselves or choose appropriate clothing
- Trouble remembering names of relatives and caregivers
- Forgetting personal history
- Major changes in personality and behavior
- Delusions and hallucinations such as believing someone is out to get them
- Tendency to become lost

Stage Seven

- Loss of verbal skills
- Loss of motor skills and ability to control movement
- Inability to dress, bathe, or feed themselves
- Difficulty sitting or holding up head without support
- Rigid muscles
- Trouble swallowing



Time line of disease progression

Mild short term
memory loss

Severe memory loss

Less independent

Forgetting timing days

Trouble doing simple arithmetic's

No day-to day recall

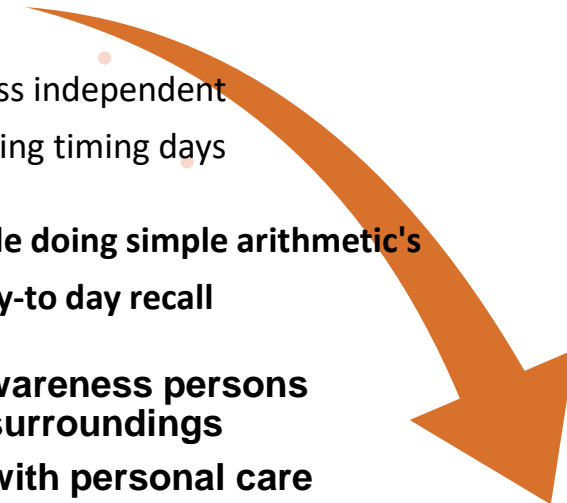
No Awareness persons
surroundings

Help with personal care
Behaviours

Non Verbal

Non mobile or frequent falls

Impaired swallowing



Drug therapies
Cholinergic Replacement
NMDA control
Symptoms control with Psychotropics

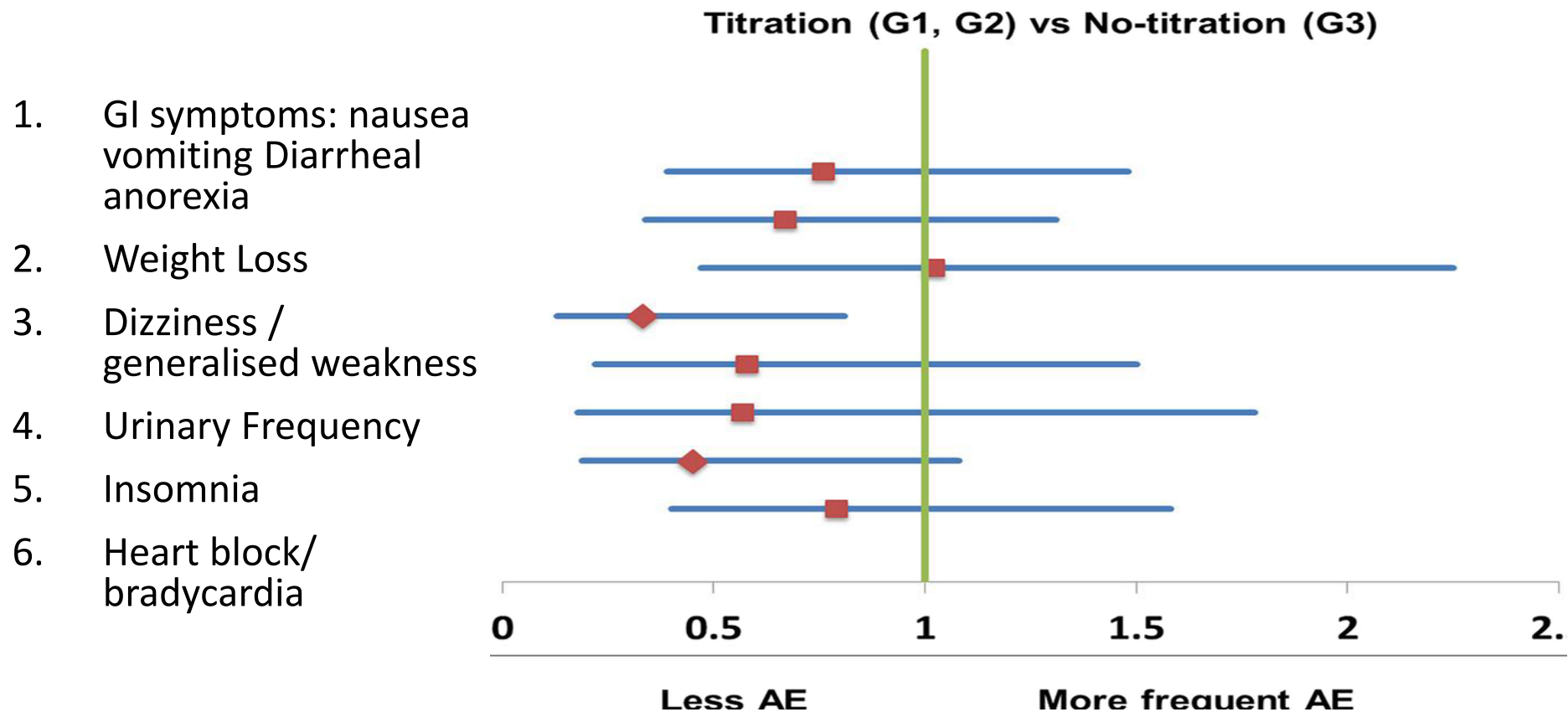


Memory Enhancers

Cholinesterase inhibitors

NMDA Inh Memantine





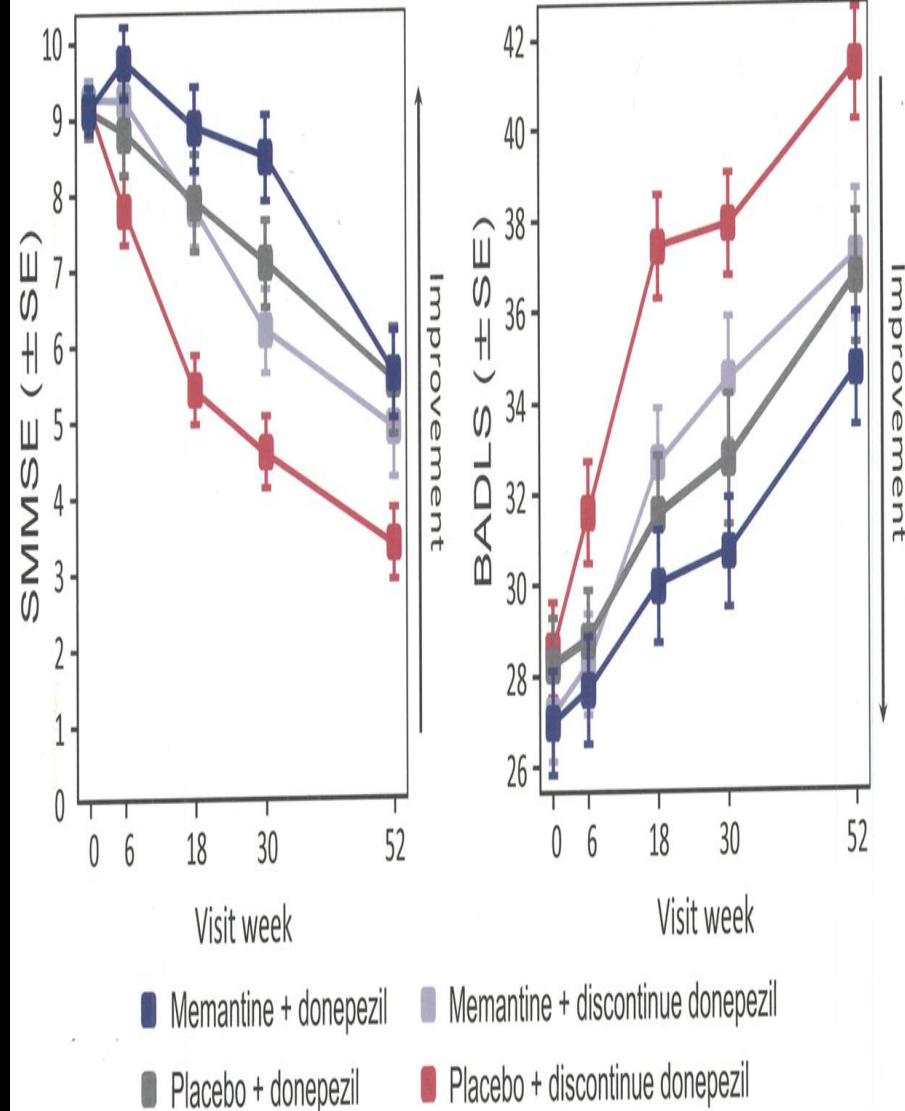
Cholinh +/- Memantine? DOMINO Trial

Original Article

Donepezil and Memantine for Moderate-to-Severe Alzheimer's Disease

Robert Howard, M.D., Rupert McShane, F.R.C.Psych., James Lindesay, D.M., Craig Ritchie, M.D., Ph.D., Ashley Baldwin, M.R.C.Psych., Robert Barber, M.D., Alistair Burns, F.R.C.Psych., Tom Denning, F.R.C.Psych., David Findlay, M.B., Ch.B., Clive Holmes, Ph.D., Alan Hughes, M.B., Ch.B., Robin Jacoby, D.M., Rob Jones, M.B., Ch.B., Roy Jones, M.B., Ian McKeith, F.Med.Sc., Ajay Macharouthu, M.R.C.Psych., John O'Brien, D.M., Peter Passmore, M.D., Bart Sheehan, M.D., Edmund Juszcak, M.Sc., Cornelius Katona, M.D., Robert Hills, D.Phil., Martin Knapp, Ph.D., Clive Ballard, M.D., Richard Brown, Ph.D., Sube Banerjee, M.D., Caroline Onions, P.G.Dip., Mary Griffin, R.G.N., Jessica Adams, B.Sc., Richard Gray, M.Sc., Tony Johnson, Ph.D., Peter Bentham, M.B., Ch.B., and Patrick Phillips, Ph.D.

N Engl J Med
Volume 366(10):893-903
March 8, 2012



Course of **neuropsychiatric** symptoms in dementia a 5 year longitudinal study

Geriatric Psychiatry. Vol33, Is10,Oct 2018 Pg1361-69

Neuropsychiatric symptoms were common at baseline, and only a moderate increase in total score

97% scored ≥ 16 , 49% scored ≥ 36

Most common symptoms: **Apathy 83%**

Depression 63%

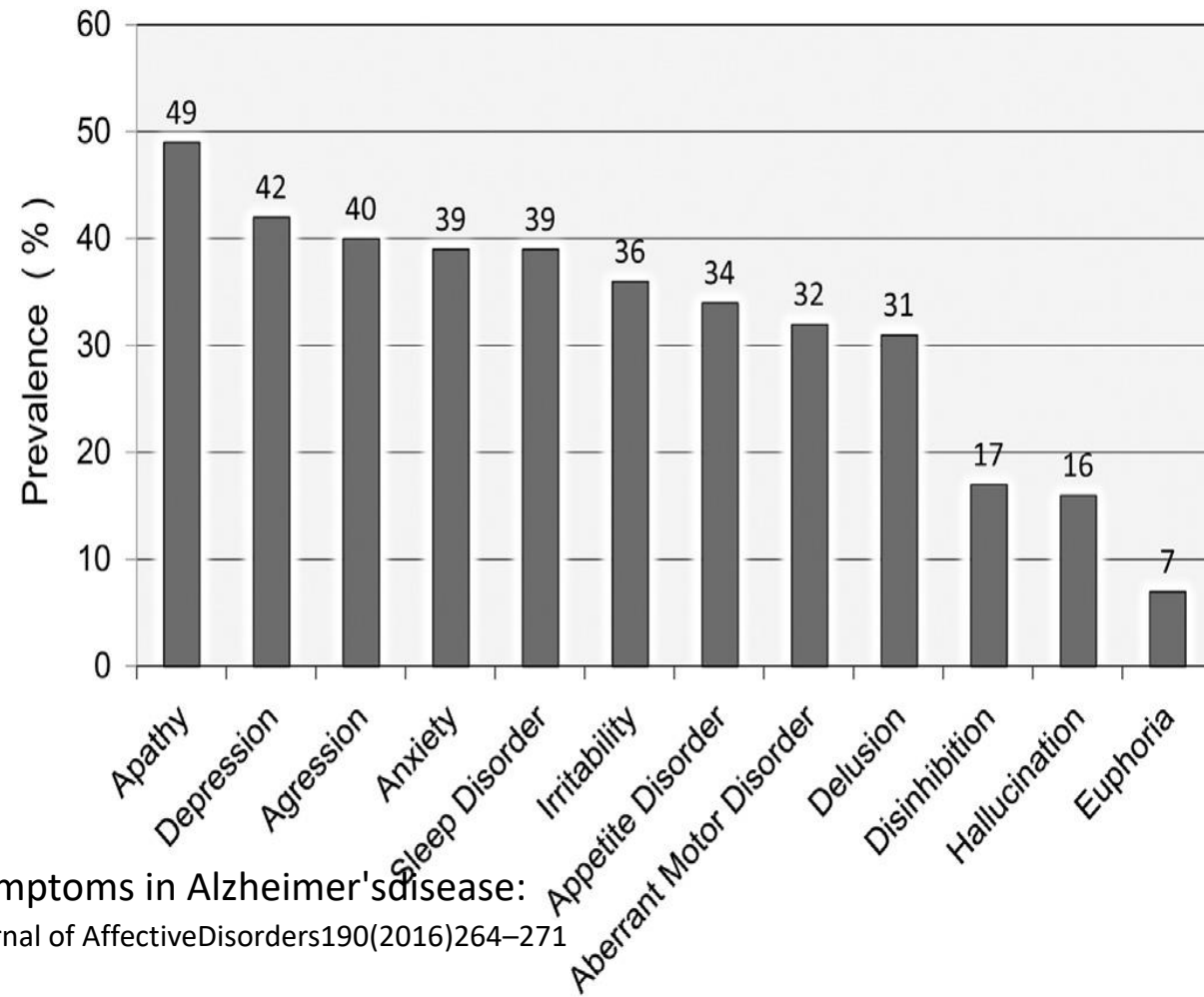
Appetite 63%

Aberrant motor behaviour 60%

- ◆ Cognitive decline was associated with higher NPI tot. score and several NPI items
- ◆ Lewy body dementia was associated with higher NPI total score and psychotic symptoms.
- ◆ Alzheimer's disease was associated with increase in apathy.

HIGH PREVALENCE BEHAVIOURS

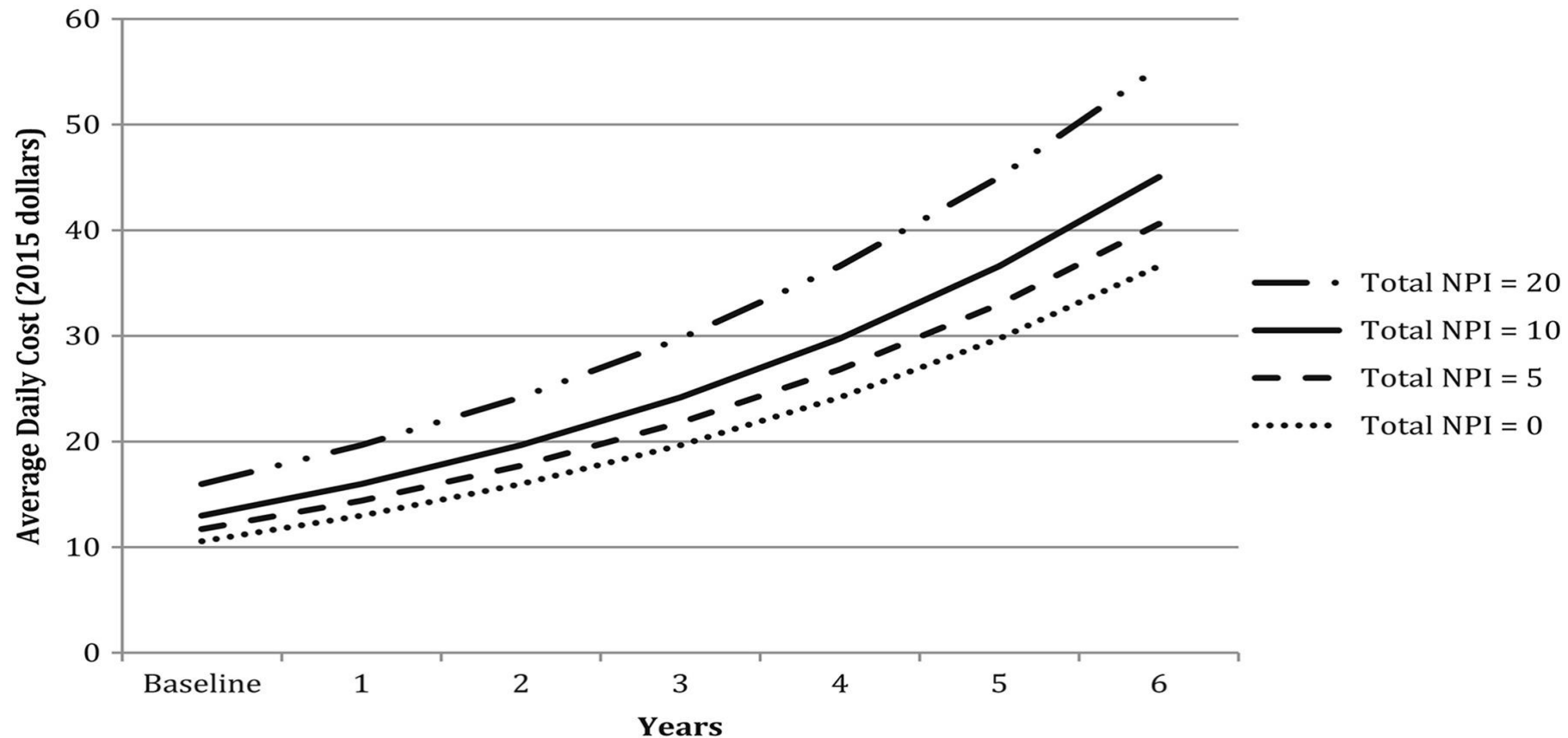
Depression
Anxiety
Apathy
Irritability
Agitation
Aggression
Wandering



The prevalence of neuropsychiatric symptoms in Alzheimer's disease:
Systematic review and meta-analysis Journal of Affective Disorders 190(2016)264–271

increasing costs over time and with increasing dementia severity
(decreasing Mini-Mental State Examination [MMSE] score each year

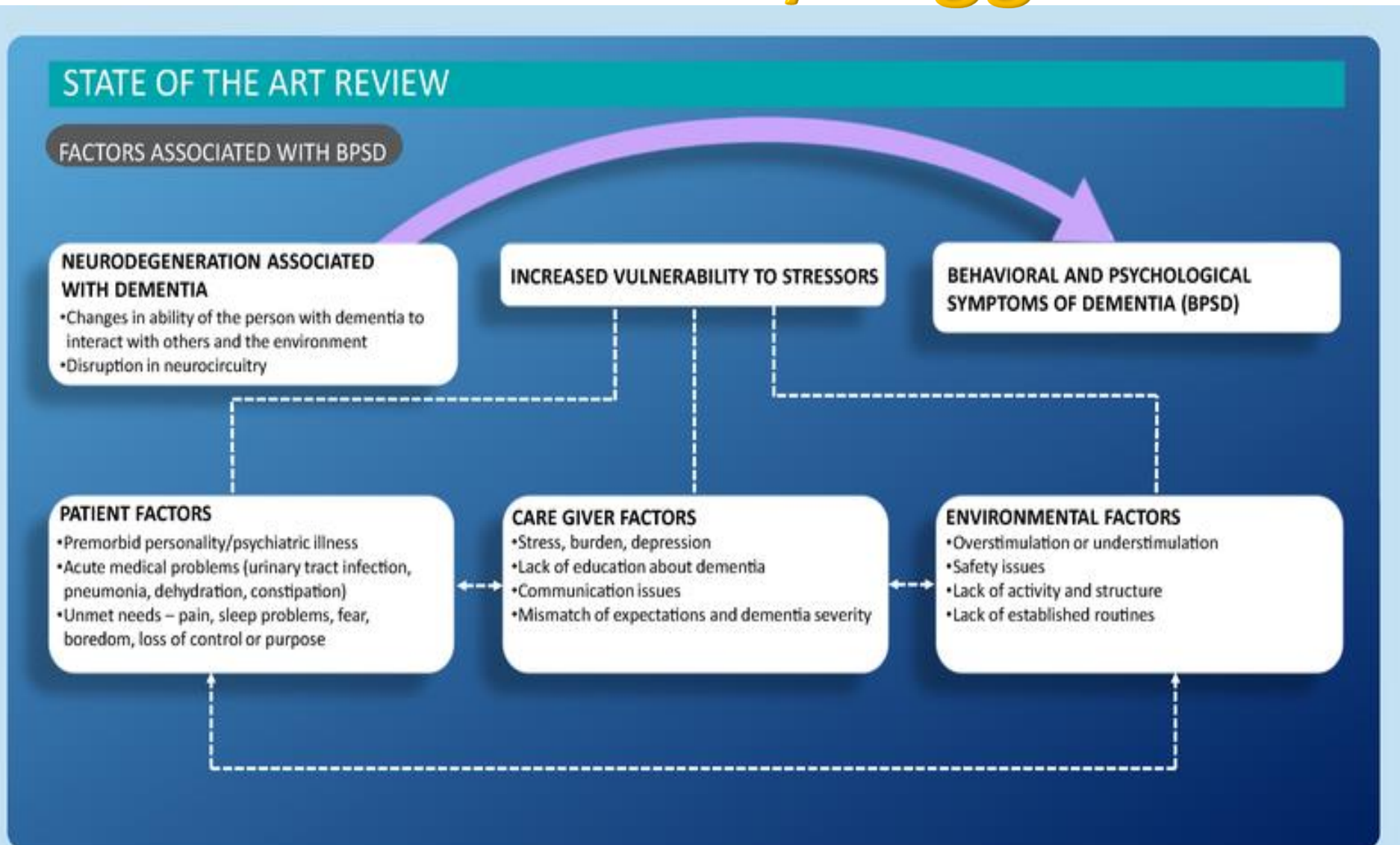
**Estimates of Informal Costs for Selected NPI -
12 Scores**



Neuropsychiatric symptoms in patients with dementia and the
longitudinal costs of informal care in the Cache County population

G B Rattinger et al. 2016

Factors associated/trigger BPSD



Emotions

- Primary Reinforcers

Taste Touch Smell

Facial Expression

Facial Beauty

Auditory Consonance

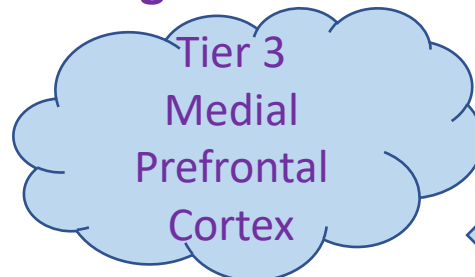
Pleasantness

Orbitofrontal Cortex
Tier 2

Amygdala

Value computation

Decision
making



Aggression

- *Personality trait*

Uncontrolled temperament

Impulsiveness

Low empathy

Agreeableness

Conscientiousness

- *Bipolar I disorder*

- *Borderline personality disorder 1.6-6%*

- *ADHD disorder*

- *Autism spectrum RBFOX1 gene Chr 16*

- *Heritability 50%*

- *Alzheimer Disease presents a high occurrence of aggression, estimated to occur in around 40%*



Agitation /Aggression

- Pacing nervously
- Rummaging
- Restlessness
- Repetitive movements
- Wandering trying to leave

Excessive
Motor activity



- Argumentative
- Bad tempered
- Open conflict with others
- Verbally abusive
- Yelling Screaming
- Using profanity
- Insulting
- Overly critical
- Shut up to others

Verbal
aggression



- Resisting care
- Grabbing
- Pushing hitting
- Slamming doors
- Taking things from others
- destroying property
- Scratching
- Biting
- Spitting Shoving slapping
- Kicking

Physical
aggression



Aggression

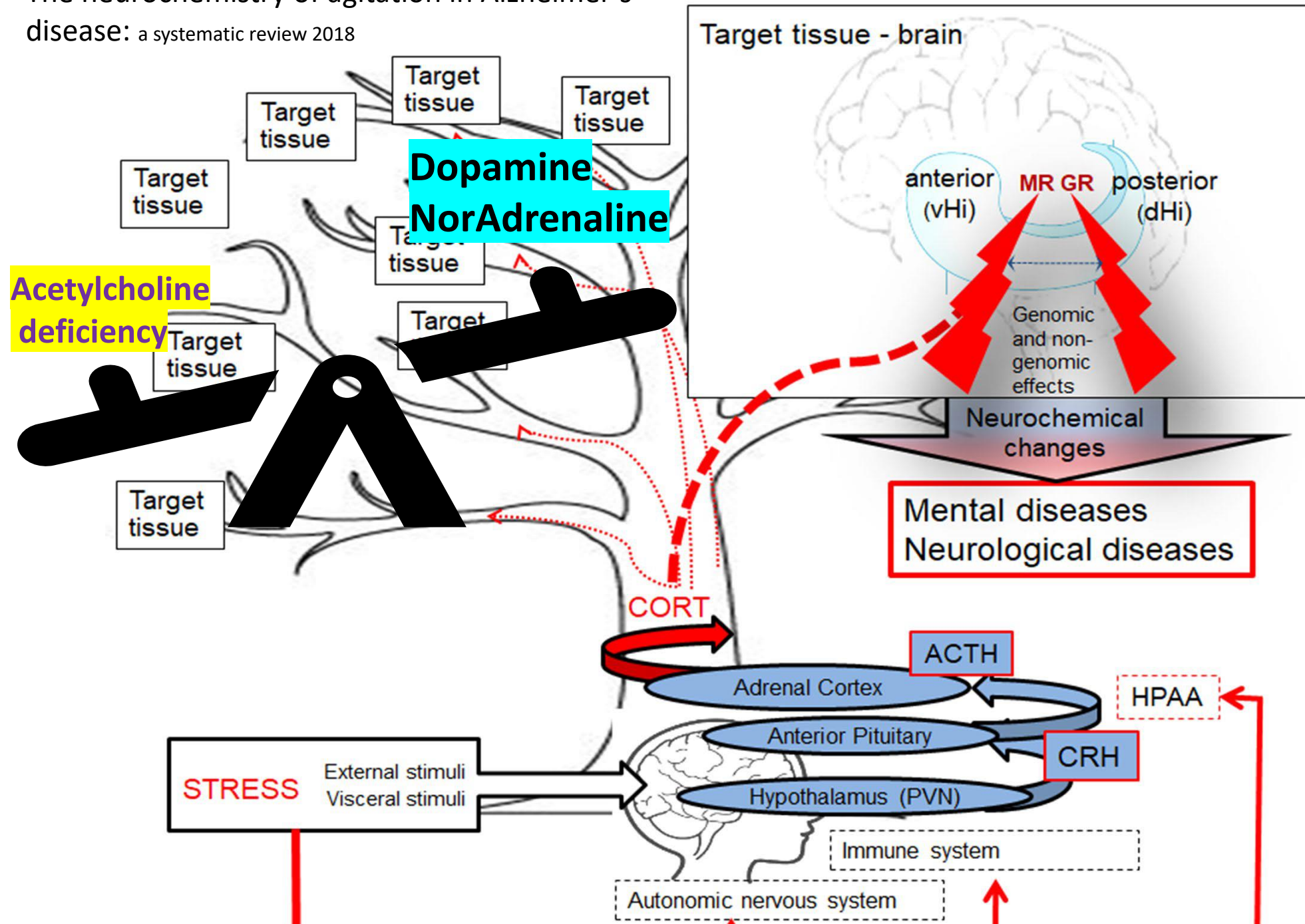
- Specific genes influence aggressive behaviours:
- ***Opioid Mu receptor*** Less physical aggressive actions
- Brain activation during aggressive decision making
- Reactivity to social provocation
- Activation of Cortico-limbic structures: Insular cortex
Orbito-frontal cortex
- Neuro excitability Anterior Cingulate Pre-frontal dorso
lateral Cortex

Behavioral Escalation

- 💣 Restless Behaviours\Irritable
- 💣 Wandering
- 💣 Rummaging
- 💣 Agitation
- 💣 Non physical aggression
- 💣 Verbal aggression
- 💣 **Physical Aggression**

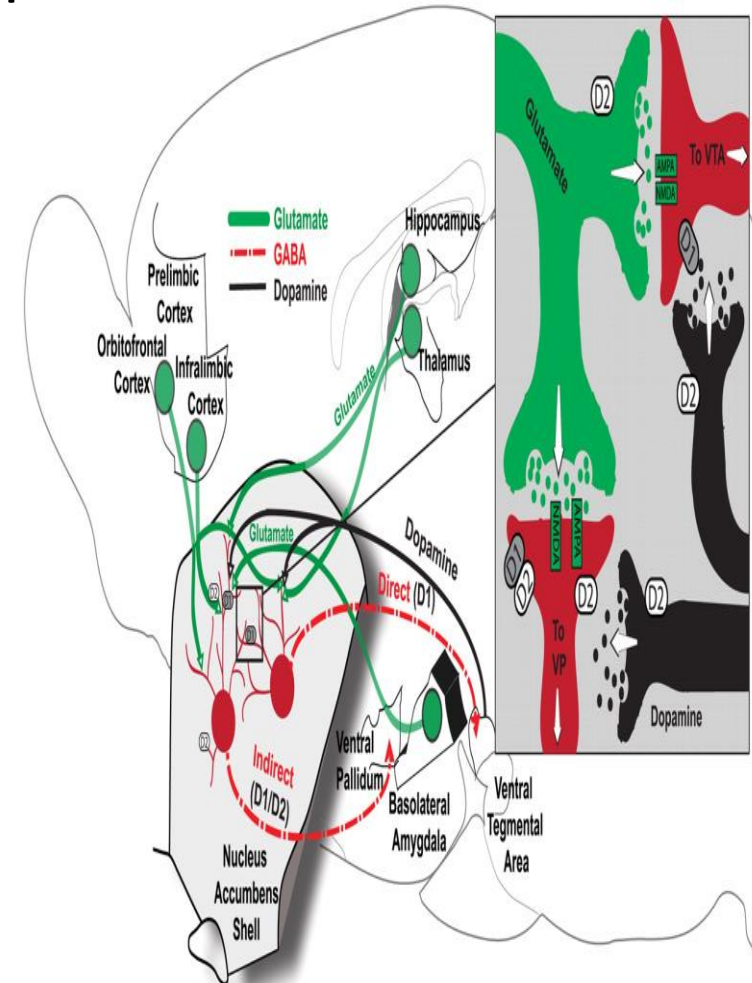


The neurochemistry of agitation in Alzheimer's disease: a systematic review 2018



Neuro-excitation in dementia

- Glutamate/ GABA pathways disruption : Learning
- Memory processing
- Memory consolidation
- Neuroplasticity
- Reduced Energy production
- Tau phosphorylation
- and toxicity



Are premorbid abnormal personality traits associated with behavioural and psychological symptoms in dementia?

J Prior et al, Inter.J.Geriatric psychiatry. Vol31, Is9, Pg 1050-55, Sept 2016

The presence of Clusters A (solitary/paranoid) and C (anxious/dependent) abnormal premorbid personality traits seems to affect the expression of certain behavioural and psychological symptoms in dementia, depression in particular

Personality and behavioural and psychological Symptoms in dementia: Results from the PACO study

The Journal of the Alzheimer's Association, 2018, Vol 14, Iss7, Pg1468

Premorbid Neuroticism was positively correlated with depression, anxiety, irritability, sleep disorders, and the total NPI score. Premorbid conscientiousness was negatively correlated with delusions, apathy, and total NPI score

Premorbid Mood Disorders

- Anxiety 15%
 - Depression 6-20%
 - Obsessive Compulsive symptoms 1-3%
 - Intermittent Explosive Disorder 1-2%
-
- Co existence of more entities
 - Overlapping symptoms
 - No insight into the magnitude of symptoms

Symptomatology

Action Plan

DESCRIBE

INVESTIGATE

CREATE

EVALUATE




NPI	Neuropsychiatric Inventory Worksheet	NPI	Neuropsychiatric Inventory Worksheet
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Directions: Read all items from the NPI "Instructions for Administration of the NPI". Mark Caregiver responses on this worksheet before scoring the Frequency, Severity, and Caregiver Distress for each item.

A. DELUSIONS: <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> N/A Frequency_____Severity_____Distress_____ <input type="checkbox"/> 1. Fear of harm <input type="checkbox"/> 2. Fear of theft <input type="checkbox"/> 3. Spousal affair <input type="checkbox"/> 4. Phantom boarder <input type="checkbox"/> 5. Spouse imposter <input type="checkbox"/> 6. House not home <input type="checkbox"/> 7. Fear of abandonment <input type="checkbox"/> 8. Talks to TV, etc. <input type="checkbox"/> 9. Other _____	B. HALLUCINATIONS: <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> N/A Frequency_____Severity_____Distress_____ <input type="checkbox"/> 1. Hears voices <input type="checkbox"/> 2. Talks to people not there <input type="checkbox"/> 3. Sees things not there <input type="checkbox"/> 4. Smells things not there <input type="checkbox"/> 5. Feels things not there <input type="checkbox"/> 6. Unusual taste sensations <input type="checkbox"/> 7. Other _____	G. APATHY/INDIFFERENCE: <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> N/A Frequency_____Severity_____Distress_____ <input type="checkbox"/> 1. Less spontaneous or active <input type="checkbox"/> 2. Less likely to initiate conversation <input type="checkbox"/> 3. Less affectionate, lacking emotions <input type="checkbox"/> 4. Contributes less to household chores <input type="checkbox"/> 5. Less interested in others <input type="checkbox"/> 6. Lost interest in friends or family <input type="checkbox"/> 7. Less enthusiastic about interests <input type="checkbox"/> 8. Other _____	H. DISINHIBITION: <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> N/A Frequency_____Severity_____Distress_____ <input type="checkbox"/> 1. Acts impulsively <input type="checkbox"/> 2. Excessively familiar with strangers <input type="checkbox"/> 3. Insensitive or hurtful remarks <input type="checkbox"/> 4. Crude or sexual remarks <input type="checkbox"/> 5. Talks openly of private matters <input type="checkbox"/> 6. Inappropriate touching of others <input type="checkbox"/> 7. Other _____
C. AGITATION/AGGRESSION: <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> N/A Frequency_____Severity_____Distress_____ <input type="checkbox"/> 1. Upset with caregiver; resists ADL's <input type="checkbox"/> 2. Stubbornness <input type="checkbox"/> 3. Uncooperative; resists help <input type="checkbox"/> 4. Hard to handle <input type="checkbox"/> 5. Cursing or shouting angrily <input type="checkbox"/> 6. Slams doors; kicks, throws things <input type="checkbox"/> 7. Hits, harms others <input type="checkbox"/> 8. Other _____	D. DEPRESSION/DYSPHORIA: <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> N/A Frequency_____Severity_____Distress_____ <input type="checkbox"/> 1. Tearful and sobbing <input type="checkbox"/> 2. States, acts as if sad <input type="checkbox"/> 3. Puts self down, feels like failure <input type="checkbox"/> 4. "Bad person", deserves punishment <input type="checkbox"/> 5. Discouraged, no future <input type="checkbox"/> 6. Burden to family <input type="checkbox"/> 7. Talks about dying, killing self <input type="checkbox"/> 8. Other _____	I. IRRITABILITY/LIBILITY: <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> N/A Frequency_____Severity_____Distress_____ <input type="checkbox"/> 1. Bad temper, "flies off handle" easily <input type="checkbox"/> 2. Rapid changes in mood <input type="checkbox"/> 3. Sudden flashes of anger <input type="checkbox"/> 4. Impatient, trouble coping with delays <input type="checkbox"/> 5. Cranky, irritable <input type="checkbox"/> 6. Argues, difficult to get along with <input type="checkbox"/> 7. Other _____	J. ABERRANT MOTOR BEHAVIOR: <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> N/A Frequency_____Severity_____Distress_____ <input type="checkbox"/> 1. Paces without purpose <input type="checkbox"/> 2. Opens or unpacks closets or drawers <input type="checkbox"/> 3. Repeatedly dresses and undresses <input type="checkbox"/> 4. Repetitive activities or "habits" <input type="checkbox"/> 5. Handling, picking, wrapping behavior <input type="checkbox"/> 6. Excessively fidgety <input type="checkbox"/> 7. Other _____
E. ANXIETY: <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> N/A Frequency_____Severity_____Distress_____ <input type="checkbox"/> 1. Worries about planned events <input type="checkbox"/> 2. Feels shaky, tense <input type="checkbox"/> 3. Sobs, sighs, gasps <input type="checkbox"/> 4. Racing heart, "butterflies" <input type="checkbox"/> 5. Phobic avoidance <input type="checkbox"/> 6. Separation anxiety <input type="checkbox"/> 7. Other _____	F. ELATION/EUPHORIA: <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> N/A Frequency_____Severity_____Distress_____ <input type="checkbox"/> 1. Feels too good, too happy <input type="checkbox"/> 2. Abnormal humor <input type="checkbox"/> 3. Childish, laughs inappropriately <input type="checkbox"/> 4. Jokes or remarks not funny to others <input type="checkbox"/> 5. Childish pranks <input type="checkbox"/> 6. Talks "big", grandiose <input type="checkbox"/> 7. Other _____	K. SLEEP AND NIGHTTIME BEHAVIOR DISORDERS: <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> N/A Frequency_____Severity_____Distress_____ <input type="checkbox"/> 1. Difficulty falling asleep <input type="checkbox"/> 2. Up during the night <input type="checkbox"/> 3. Wanders, paces, inappropriate activity <input type="checkbox"/> 4. Awakens others at night <input type="checkbox"/> 5. Wakes and dresses to go out at night <input type="checkbox"/> 6. Early morning awakening <input type="checkbox"/> 7. Sleeps excessively during the day <input type="checkbox"/> 8. Other _____	L. APPETITE/EATING CHANGES: <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> N/A Frequency_____Severity_____Distress_____ <input type="checkbox"/> 1. Loss of appetite <input type="checkbox"/> 2. Increased appetite <input type="checkbox"/> 3. Weight loss <input type="checkbox"/> 4. Weight gain <input type="checkbox"/> 5. Change in eating habits <input type="checkbox"/> 6. Change in food preferences <input type="checkbox"/> 7. Eating rituals <input type="checkbox"/> 8. Other _____

NPI scoring

- Distress is scored as: Frequency  Severity
- 0. Not at all
- 1. Minimally (almost no change in work routine)
- 2. Mildly (some change in work routine but little time re-budgeting required)
- 3. Moderately (disrupts work routine, requires time re-budgeting)
- 4. Severely (disruptive, upsetting to staff and other residents, major time infringement)
- 5. Very Severely or Extremely (very disruptive, major source of distress for staff and other residents, requires time usually devoted to other residents)

1	Short attention span, easy distractible, unable to concentrate			
2	Impulsive, impatient, low tolerance for pain frustration			
3	Uncooperative, resistance to care, demanding			
4	Violent/threatening violence people property			
5	Explosive and/or unpredictable anger			
6	Rocking, rubbing, moaning/other stimulating behaviour			
7	Pulling at tubes restrains blankets			
8	Wandering form treatment areas			
9	Restless, pacing, excessive movements			
10	Repetitive behaviours motor/verbal			
11	Rapid loud/ excessive talking			
12	Sudden change of mood			
13	Easily initiated/ excessive crying/ laughing			
14	Sub-abusiveness physical/verbal			
	Total scores	Disinhibition score	Aggression score	Lability score

Souvenaid

3 trials

Prodromal mild Ad
and MCI 24 weeks
little or no difference
in global or specific
cognitive functions



Behavioural Intervention

The **DICE** Approach



Describe

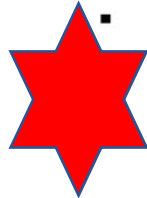
Investigate

Create

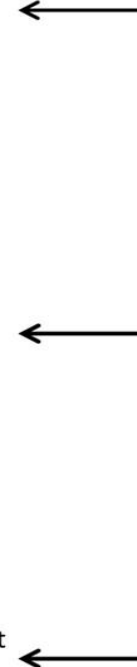
Evaluate

Kales
et al,
JAGS,
2014

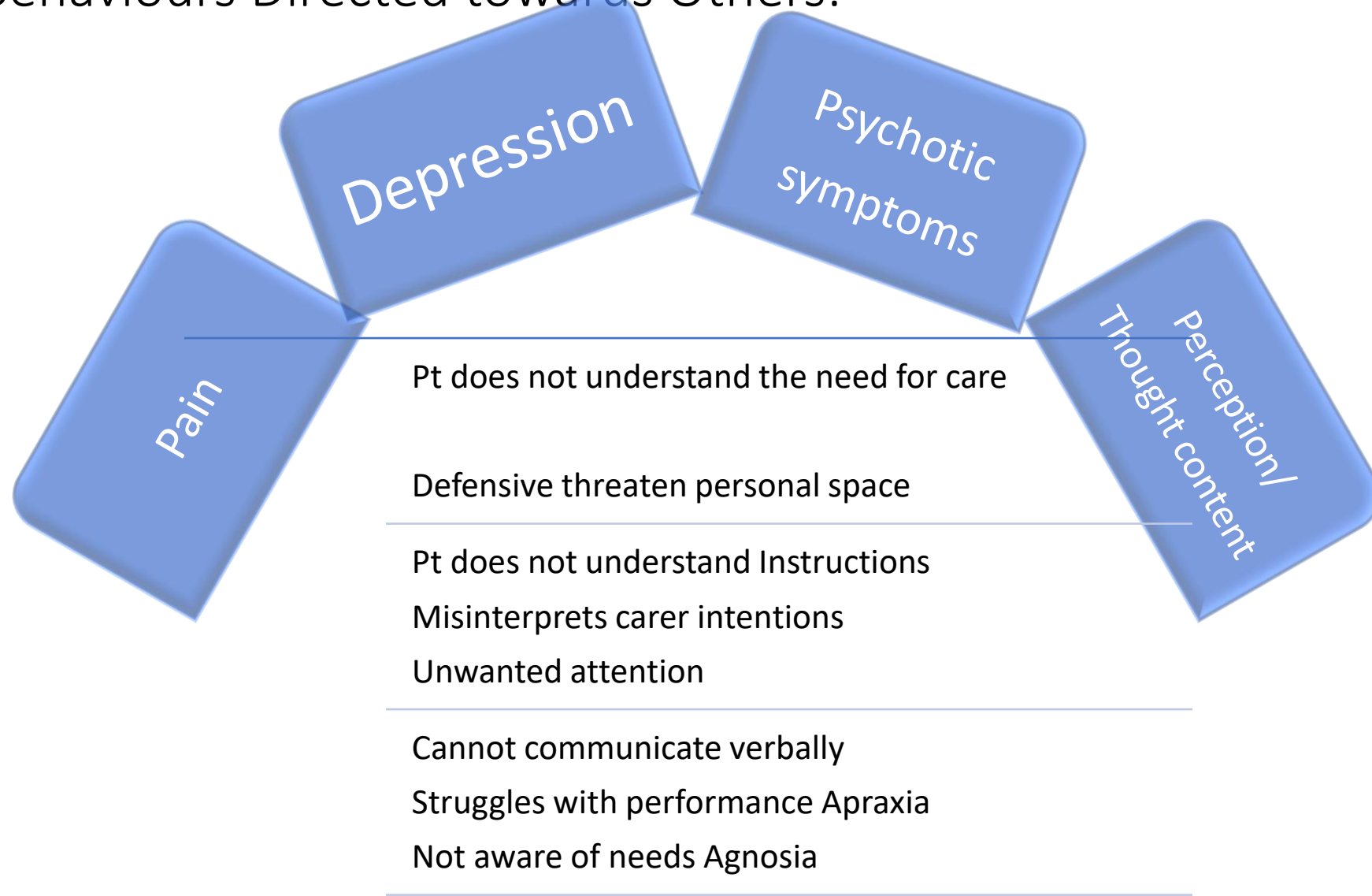
- Caregiver **describes** problematic behavior
 - Context (who, what, when and where)
 - Social and physical environment
 - Patient perspective
 - Degree of distress to patient and caregiver
- Provider **investigates** possible causes of problem behavior
 - Patient
 - Medication side effects
 - Pain
 - Functional limitations
 - Medical conditions
 - Psychiatric comorbidity
 - Severity of cognitive impairment, executive dysfunction
 - Poor sleep hygiene
 - Sensory changes
 - Fear, sense of loss of control, boredom
 - Caregiver effects/expectations
 - Social and physical environment
 - Cultural factors
- Provider, caregiver and team **collaborate to create** and implement treatment plan
 - Respond to physical problems
 - Strategize behavioral interventions
 - Providing caregiver education and support
 - Enhancing communication with the patient
 - Creating meaningful activities for the patient
 - Simplifying tasks
 - Ensuring the environment is safe
 - Increasing or decreasing stimulation in the environment
- Provider **evaluates** whether “CREATE” interventions have been implemented by caregiver and are safe and effective



Consideration of Psychotropic Use (Acuity/Safety)



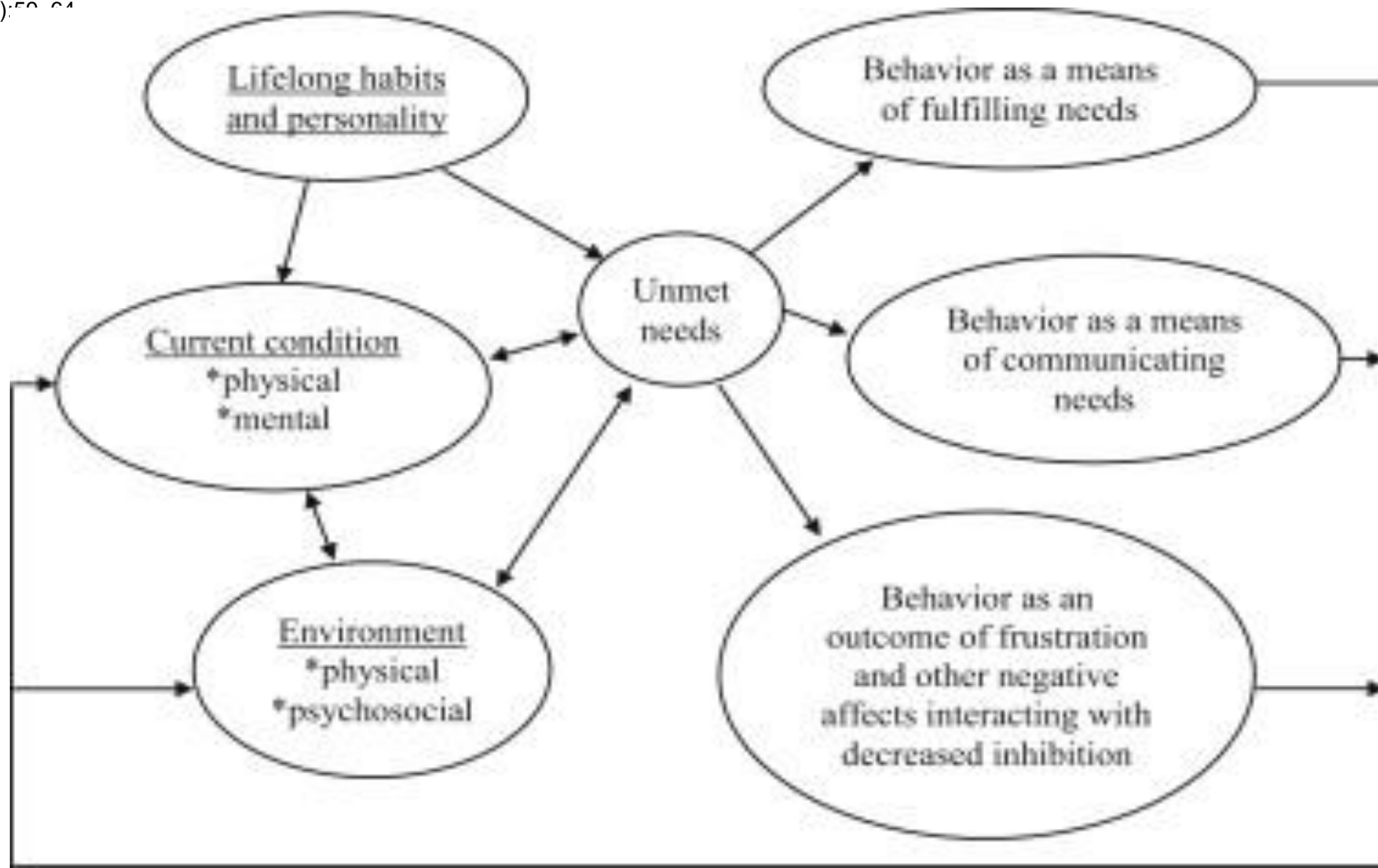
Reactive Factors Related to Rejection of Care and Behaviours Directed towards Others:



Which unmet needs contribute to behavior problems in persons with advanced dementia?

Cohen-Mansfield J, Dakheel-Ali M,

Marx MS, Thein K, Regier NG. Psychiatry Res.
2015;228(1):50-64



Which unmet needs contribute to behavior problems in persons with advanced dementia?

Cohen-Mansfield J, Dakheel-Ali M, Marx MS, Thein K, Regier NG. Psychiatry Res. 2015;228(1):59–64.

- On average, three unmet needs were identified per nursing home resident.
- The most common were needs for stimulation and for meaningful and social activities.
- The detection of PAIN

Lateral Thinking **Disruptive Behaviour**



Physical Discomfort
Emotional Distress

Associations between pain and behavioural and psychiatric symptoms of dementia, using generalised estimating equations in 230 older people with dementia and unplanned acute medical admission. PAINAD (pain during movement)(pain at rest)

2015International Association for the Study

of Pain ■

The Response of Agitated Behavior to Pain Management in Persons with Dementia

Husebo

B, Ballard C American Journal of Geriatric Psychiatry, 2014-07-01, Volume 22, Issue 7, Pages 708-717

- **Verbal Agitation:** $p < 0.001$

Complaining- Negativism Repetitious sentences -
Questioning-

Attention sicking- Cursing- Swearing

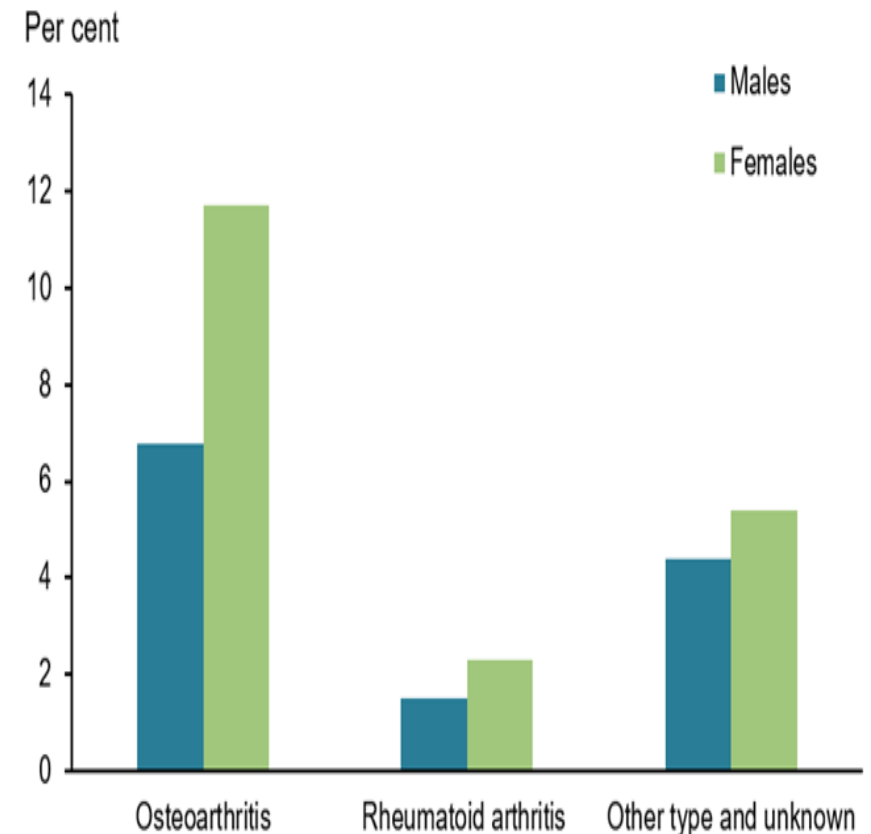
- **Physical Non-Aggressive behaviours** $p = 0.008$

Pacing- Restless-

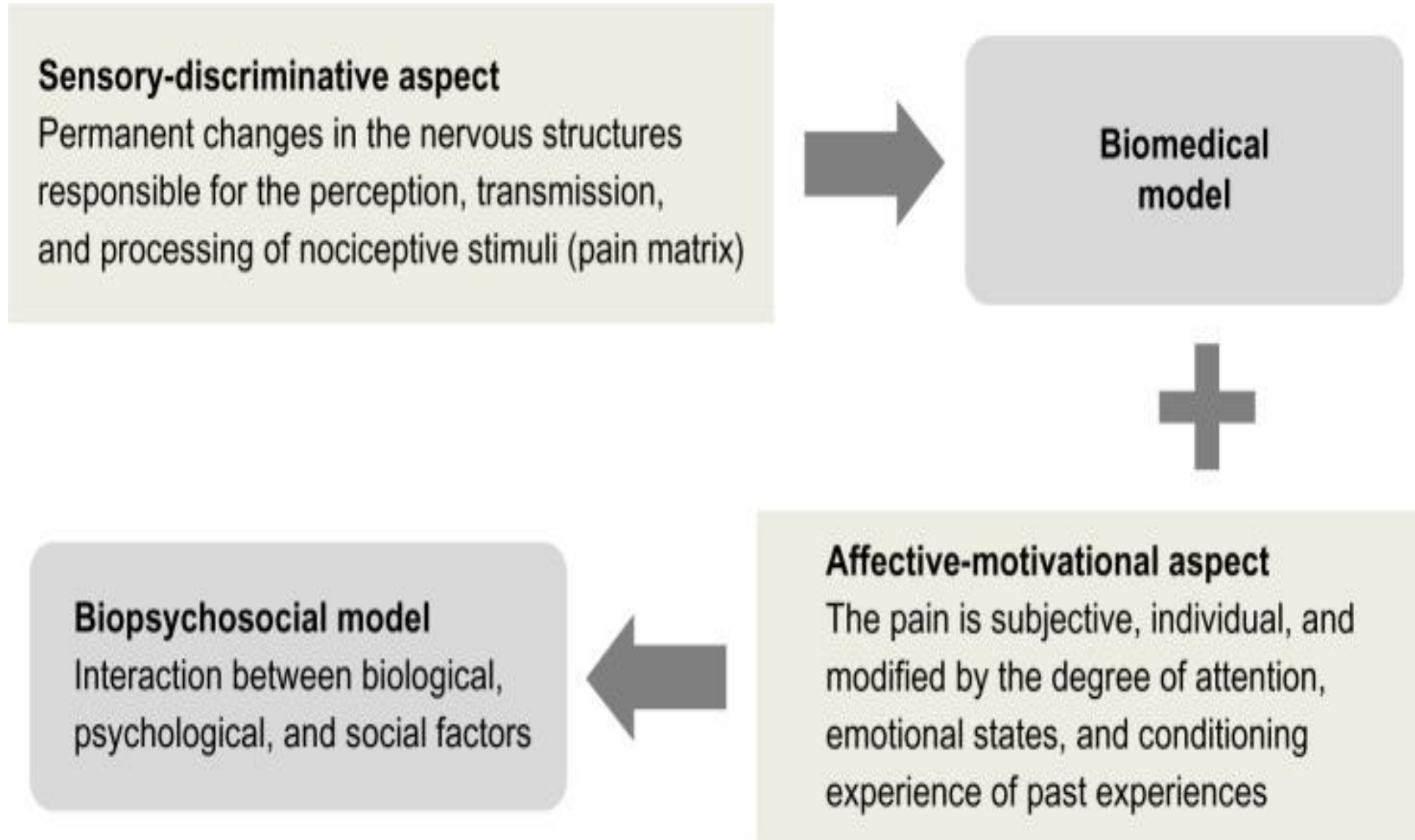
- **Physically aggressive behaviours** $p = 0.37$

Most popular pains

- Arthritis Shoulder knee hip pain >90%
- Complex neck Back pain >40%
- Osteoporosis >60%
- Stomach IBS
- Angina
- Peripheral Neuropathy
- Bladder// Pelvic pain



Pain in Older Adults



Survey of the most frequently used observational scales

Zwakhlen et al 2017

Observational scales	Available versions	Authors
The Abbey Pain Scale	English, Italian, Japanese	Abbey et al., ¹ 2004
Algoplus	English, French	Rat et al., ⁵³ 2011
Checklist of Nonverbal Pain Indicators (CNPI)	English, Norwegian	Feldt, 2000 ²⁸
CNA Pain Assessment Tool (CPAT)	English	Cervo et al., ¹³ 2009
Doloshort	French	Pautex et al., ⁴⁹ 2009
Doloplus-2	Chinese, Dutch, English, French, Italian, Japanese, Norwegian, Portuguese, Spanish	Levebre-Chapiro, 2001⁴³
Discomfort Scale for Dementia of the Alzheimer's Type (DS-DAT)	Dutch, English, Italian	Hurley et al., ³² 1992
Elderly Pain Caring Assessment (EPCA-2)	French	Morello et al., ⁴⁸ 2007
Mahoney Pain Scale (MPS)	English	Mahoney and Peters, 2008 ⁴⁷
Mobilization-Observation-Behavior-Intensity-Dementia (MOBID and MOBID-2)	Norwegian	Husebo et al., ³³ 2007
Noncommunicating Patient's Pain Assessment Instrument (NOPPAIN)	Brazilian, English, Italian, Portuguese	Snow et al., ⁶³ 2004
The Pain Assessment in the Cognitively Impaired (PACI)	English	Kaasalainen et al., ^{36,37} 2011
The Pain Assessment Scale for Seniors with Severe Dementia (PACSLAC)	Dutch, English, French, Japanese, Portuguese	Fuchs-Lacelle and Hadjistavropoulos, 2004³⁰
PACSLAC-D	Dutch	Zwakhlen et al., ⁷⁹ 2007
PACSLAC 2	English	Chan et al., ¹⁴ 2014
Pain Assessments in Dementing Elders (PADE)	English	Villanueva et al., ⁷⁴ 2003
The Pain Assessment in Advanced Dementia Scale (PAINAD)	Chinese, Dutch, English, German, Italian, Portuguese, Spanish	Warden et al.,⁷⁵ 2003
Pain Assessment in Noncommunicative Elderly Persons (PAINE)	English	Cohen-Mansfield, 2006 ¹⁶
Rotterdam Elderly Pain Observation Scale (REPOS)	Dutch, English	van Herk et al., ⁷² 2008
Bold denotes scales, which can be especially recommended according to others, ^{60,61} and one of the present authors (S.L.).		

PAINAD

	0	1	2	Score
Breathing, independent of vocalization	Normal	<ul style="list-style-type: none"> ▪Occasional labored breathing ▪Short period of hyperventilation 	<ul style="list-style-type: none"> ▪Noisy labored breathing ▪Long period of hyperventilation ▪Cheyne-Stokes respirations 	
Negative vocalization	None	<ul style="list-style-type: none"> ▪Occasional moan or groan ▪Low-level speech with a negative or disapproving quality 	<ul style="list-style-type: none"> ▪Repeated troubled calling out ▪Loud moaning or groaning ▪Crying 	
Facial expression	Smiling or inexpressive	<ul style="list-style-type: none"> ▪Sad, frightened ▪Frown 	<ul style="list-style-type: none"> ▪Facial grimacing 	
Body language	Relaxed	<ul style="list-style-type: none"> ▪Tense ▪Distressed pacing ▪Fidgeting 	<ul style="list-style-type: none"> ▪Rigid, fists clenched ▪Knees pulled up ▪Pulling or pushing away ▪Striking out 	
Consolability	No need to console	<ul style="list-style-type: none"> ▪Distracted or reassured by voice or touch 	<ul style="list-style-type: none"> ▪Unable to console, distract, or reassure 	
				TOTAL

Abbey Pain Scale

For measurement of pain in people with dementia who cannot verbalise.

How to use scale : While observing the resident, score questions 1 to 6.

Name of resident :

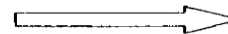
Name and designation of person completing the scale :

Date : **Time :**

Latest pain relief given was.....at.....hrs.

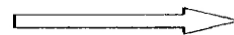
- | | | |
|---|-----------|----------------------|
| Q1. Vocalisation
eg whimpering, groaning, crying
<i>Absent 0 Mild 1 Moderate 2 Severe 3</i> | Q1 | <input type="text"/> |
| Q2. Facial expression
eg looking tense, frowning, grimacing, looking frightened
<i>Absent 0 Mild 1 Moderate 2 Severe 3</i> | Q2 | <input type="text"/> |
| Q3. Change in body language
eg fidgeting, rocking, guarding part of body, withdrawn
<i>Absent 0 Mild 1 Moderate 2 Severe 3</i> | Q3 | <input type="text"/> |
| Q4. Behavioural Change
eg increased confusion, refusing to eat, alteration in usual patterns
<i>Absent 0 Mild 1 Moderate 2 Severe 3</i> | Q4 | <input type="text"/> |
| Q5. Physiological change
eg temperature, pulse or blood pressure outside normal limits,
perspiring, flushing or pallor
<i>Absent 0 Mild 1 Moderate 2 Severe 3</i> | Q5 | <input type="text"/> |
| Q6. Physical changes
eg skin tears, pressure areas, arthritis, contractures,
previous injuries
<i>Absent 0 Mild 1 Moderate 2 Severe 3</i> | Q6 | <input type="text"/> |

Add scores for 1 - 6 and record here



Total Pain Score

**Now tick the box that matches the
Total Pain Score**



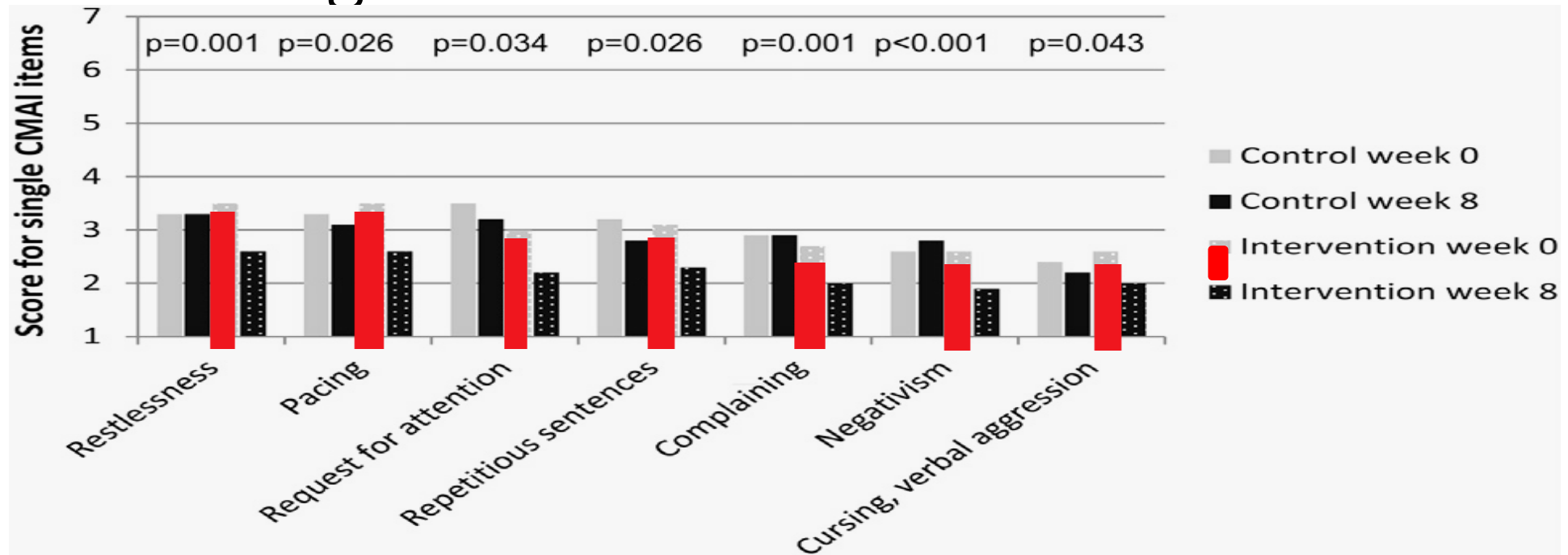
0 - 2 No pain	3 - 7 Mild	8 - 13 Moderate	14 + Severe
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**Finally, tick the box which matches
the type of pain**



Chronic	Acute	Acute on Chronic
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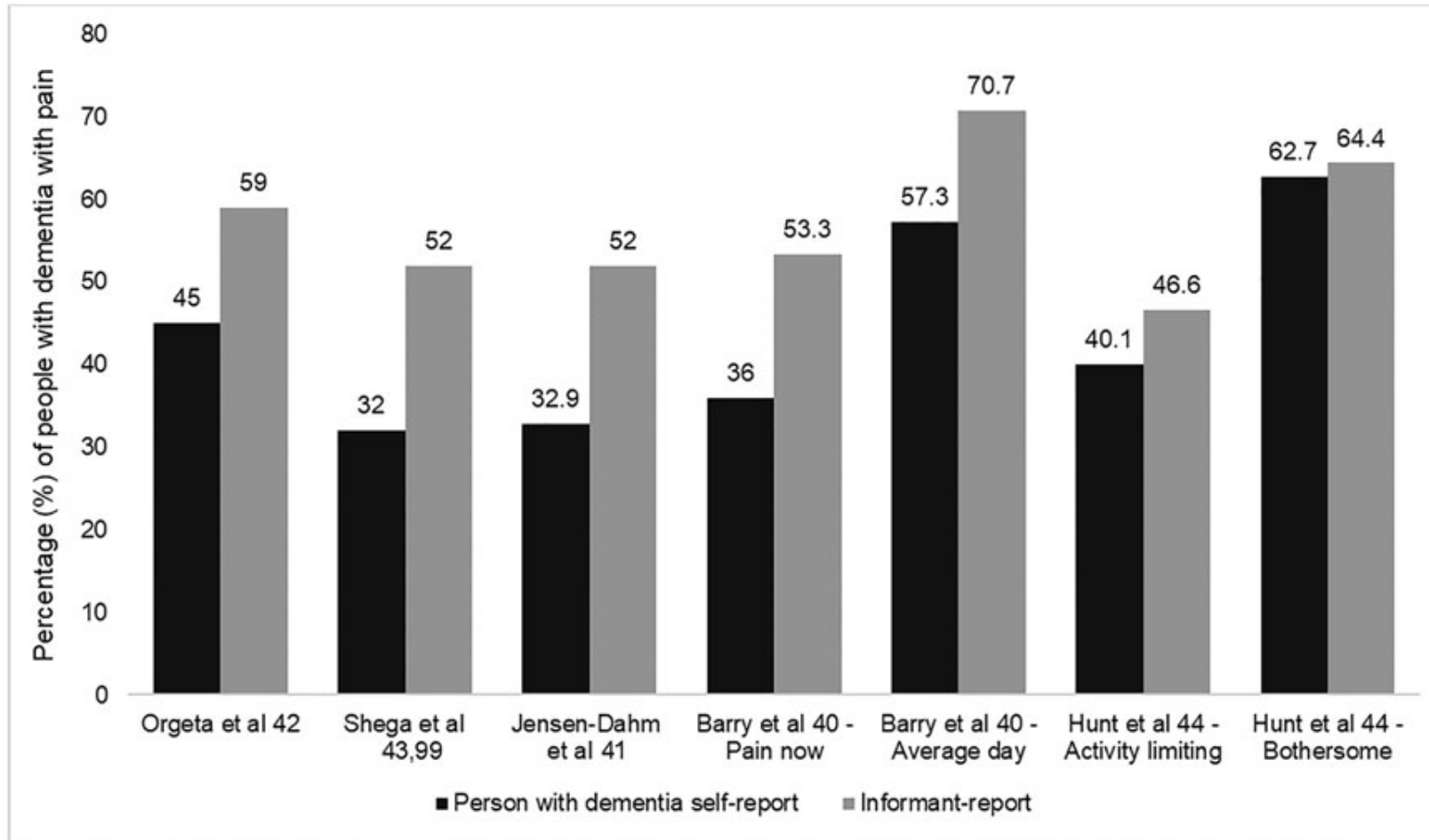
Prevalence of chronic pain in elderly nursing home residents



The response of Agitated Behavior to Pain

- Management of agitation in Persons with Dementia 2012
- Identifying and Managing Pain in People with Alzheimer's Disease and Other Types of Dementia: A Systematic Review
BS. Husebo, Wachterberg, E. FloCNS Drugs (2016) 30:481–497
- Husebo B.S., Ballard C., Sandvik R., et al: Efficacy of treating pain to reduce behavioural disturbances in residents of nursing homes with dementia: cluster randomised clinical trial. BMJ 2011; 343: pp. 1-10
- Husebo B.S., Strand L.I., Moe-Nilssen R., et al: Pain in older persons with severe dementia. Psychometric properties of the Mobilization-Observation-Behaviour-Intensity-Dementia (MOBID-2) Pain Scale in a clinical setting. Scand J Caring Sci 2010; 24: pp. 380-391
- Husebo B.S., Strand L.I., Moe-Nilssen R., et al: Who suffers most? Dementia and pain in nursing home patients: a cross-sectional study. JAMDA 2008; 9: pp. 427-433
- Husebo B.S., Strand L.I., Moe-Nilssen R., et al: Mobilization-Observation-Behavior-Intensity-Dementia Pain Scale (MOBID): development and validation of a nurse-administered pain assessment tool for use in dementia. J Pain Symp Manage 2007; 34: pp. 67-80

Pain self report/ Carers reporting

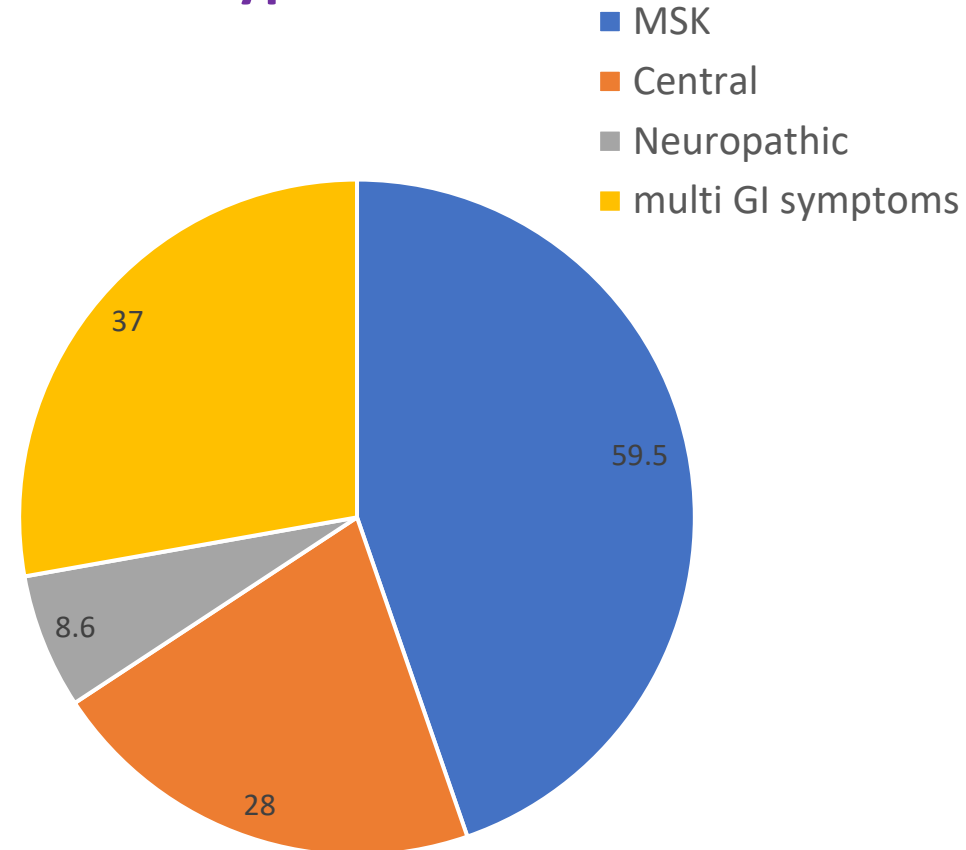


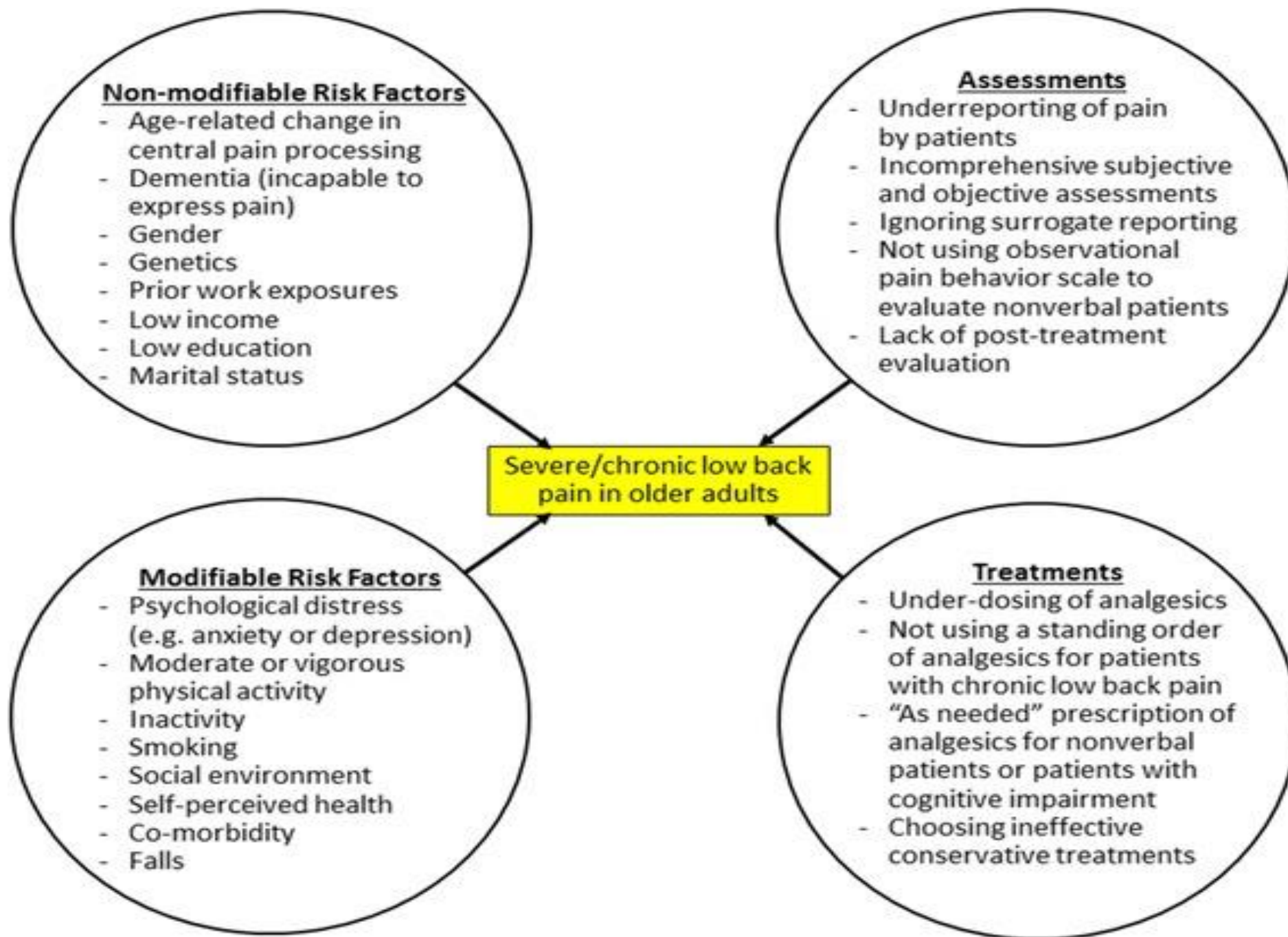
Pain assessment and pain treatment for community-dwelling people with dementia: A systematic review and narrative Synthesis Int J Geriatr Psychiatry. 2019;34:807–821

- **Prevalence of pain in atypical parkinsonism: a systematic review and meta-analysis** [*Journal of Neurology*](#) vol 266, pg2093–2102, 2019

Multiple system atrophy (MSA)
Progressive supranuclear palsy (PSP),
Corticobasal degeneration (CBD)
Dementia with Lewy bodies DLD

Pain types





Poly-pharmacy ?

- 5 > drugs vs >10 drugs
- Duration of therapy
- Appropriate vs Inappropriate
- Specific Co-morbidities



- **NNT**
- **NNH**

**3 in 4
people dispensed anti-
dementia medications
were also dispensed
medications for the
cardiovascular system
2019**

Opioids

MST

Sedating

Reduced
Clearance > 50s

Oxycontin

Less-sedating

2X stronger

Tramadol

Serotonin-
Noradrenaline
action

20% opioid action

Epileptic threshold
Digoxin toxicity

Fentanyl

Multiple
routes

100X
potent

Buprenorphine

Respiratory
Neutral

30X potent

Mu
receptors

Methadone

NMDA
action

Shorter
analgesic
action

DRUGS

- Absorption time
- Elimination= liver // kidneys
- Drug-drug interactions Amitriptyline
- Drug –disease interaction Fentanyl
- Side effects by age
- Target symptomatology

NNT

- **Statins**

>one year mostly at >4 years

Heart Protection Study,

4S study <80yrs old

more effective 65yrs old

NNT 67 intermediate CVrisk

NNH 21 Myalgia

