Henry Brodaty



How you can understand, prevent & remedy behavioural and psychological symptoms of dementia (BPSD)

Never Stand Still

Medicine

Dementia Collaborative Research Centre

www.dementiaresearch.org.au

Centre for Healthy Brain Ageing

www.cheba.unsw.edu.au

University of New South Wales (UNSW Australia)







Potential conflict of interests

- Advisor, consultant, remunerated speaker and/or investigator for multiple drug companies over last 30 years
- Currently: advisory board for Nutricia & recently completed trial for Tau Therapeutics







Clinical scenario



"Dr, Mr Smith-Jones is hitting the nurses, disrupting the other residents and being impossible. Can you prescribe something?"







What are BPSD?

- Agitation
- Aggression
- Calling out/ screaming
- Disinhibition (sexual)
- Wandering
- Night time disturbance
- Shadowing
- Swearing

- Depression
- Anxiety
- Apathy
- Delusions
- Hallucinations
- Irritability
- Elation/euphoria







Why are BPSD important?

- Ubiquitous, >90% of PWD during course
- Distress to PWD and to caregivers
- Increase rate of institutionalisation
- Higher rate of complications in hospital
- Associated with:
 - Faster rate of decline
 - Increased mortality







Effects of BPSD

- Residents with BPSD are more likely to¹:
 - be physically restrained
 - receive antipsychotic medication
 - negatively influence other residents
- BPSD increase the cost of institutional care for persons with dementia²
- BPSD, especially aggression³ & calling out⁴, increase nurse stress

¹Maslow K 1994; ²O'Brien JA et al, 2000; ³Rodney, 2000; ⁴Draper et al, 2000







Prevalence of BPSD

- In community
 - 2/3 PWD have at least one behavioural Sx
 - 1/3 PWD have significant symptoms
- In developing countries similar rates
- In residential care, residents with dementia:
 - 40- 90% have BPSD
 - Rates in similar NHs vary >3-fold

¹Lyketsos et al, Am.J. Psychiatry, 2000; 157:708-714; ²Prince M et al 2004; ³Brodaty H et al, 2001;

⁴ Seitz et al, *Int Psychogeriatrics*, 2010; 22:1025–1039





PROPERTY.

The bio-psycho-social framework









Biological causes - intrinsic

- Frontal pathology (behavioural disturbance, disinhibition, depression)
- Basal ganglia lesions (delusions)
- Temporal lobe (delusions, hallucinations)
- Locus coeruleus (psychosis, depression)
- Chemical changes serotonin, NA, DA
- Genes serotonin, dopamine receptors
- Family history of psychiatric disorder







Biological causes - extrinsic

- Acute medical illness
- Medication
- Pain
- Constipation
- Sensory impairments
- Fatigue
- Fears
- Basic needs (hunger, thirst...)
- Psychiatric syndromes eg depression







The bio-psycho-social framework

Environmental vulnerability → ↓threshold for stress or stimuli ¹	Unmet needs; unable to comprehend or make needs known ²
Neurological	Behavioural: triggers
deterioration →	and feedback from
behavioural	others control
disinhibition ⁴	behaviours ³

¹Hall and Buckwalter 1987; ² Algase et al, 1996; ³ Teri & Logsdon 2000; ⁴ Cummings JL







Before intervening ...

- **1.Is the description accurate?**
- **2.Identification of target behaviour**
- **3.Does behaviour require intervention?**
- 4. Careful diary of behaviours
- **5.Exclude non-dementia causes**
- 6.Correct sensory impairment hearing, vision







Socio- environmental	Interpersonal
Biological	Psychological







How to intervene: Environment

- Modify environment rather than person
- Avoid too much or too little stimulation
- Adequate space
- Privacy available







How to intervene: Environment

- Secure grounds
- Personalised space
- Non-institutionalised environment
- Home-like

- Colour, furnishings, architecture
- Lighting
- Resident mix
- Size of residential facility







Enhanced Environment











Good evidence for ...

- Careful optimisation of level of stimulation
 - Reduce unhelpful stimuli
 - eg noise, busy entry doors
 - Optimise helpful stimuli
 - eg light
- Good visual access to toilets
- Outdoor access with staff



Fleming R – www.dementiaresearch.org.au







Moderate evidence

Small unit size

 hard to differentiate effect of unit size from staff related factors

 Opportunity to engage in ordinary ADLs

 hard to differentiate from staff support/ engagement

Fleming R – www.dementiaresearch.org.au







Interesting – but little evidence

- Signage
- Display of personal memorabilia









Snoezelen: multisensory stimulation¹

- Significant treatment effect
 - Apathetic behaviour \downarrow
 - Loss of decorum \downarrow
 - Rebellious behaviour \downarrow
 - Aggressive behaviour \downarrow
 - Depression \downarrow



- Well-being during morning care↑
- Numbers small, methodology moderate

Van Weert et al, JAGS 2005;53: 24–33 Verkaik R et al, IJGP 2005; 20: 301–314

the Construction a research into practice





Aroma therapy

Lavender



Lemon Balm



moderate evidence from Cochrane review







Lemon balm (melissa officinalis)



- Antibacterial (eugenol)
- Antiviral (tannins)
- Mild sedative or calming agent (terpenes)
- Antioxidant activity







Light therapy





- Five studies met criteria; only 3 able to be included
- No adequate evidence of effectiveness of BLT

Forbes D, Morgan DG, Bangma J et al; Cochrane Review 2004, updated 2006







Review on animal-assisted therapy (AAT)¹

- 11 papers examining the impact of AAT on BPSD regarding their ability to
 - Reduce agitation and/or aggression
 - Promote social behaviour
 - Improve nutrition
 - Role of pet substitutes
- Small samples, short duration, few studies



¹Filan & Llewellyn-Jones (2006) Int. Psychogeriatr; 18:4, 597-611







Robotic pets, toys, dolls











The bio-psycho-social framework









Family caregivers

- Family carers as therapists for people living in the community
- Systematic review
 - ES 0.34 for decreasing BPSD
 - ES 0.15 for decreasing caregiver "stress"

Brodaty H & Arasaratnam C, Am J Psychiatry, 2012







CGs administer behavioural treatments for depression to patients with AD¹

- Behaviour therapies (pleasurable events schedule or problem solving techniques) → pt depression Sx & Dx better than controls
- Improvements maintained @ 6 months
- Bonus: CGs' depression better

¹Teri et al, J. Gerontol. 1997; 52B:159-166







Dementia Care Mapping & Person Centred Care for agitation









The bio-psycho-social framework

Socio-
environmentalInterpersonalBiologicalPsychological







Psychological Mx approaches to BPSD

- 1632 studies identified → 162 met inclusion criteria → 9 studies with Level 1 evidence
- Psycho-education for CGs effective
- Benefits lasted months
- Other CG interventions not effective
- Behaviour Mx techniques centering on individual pts' or CG behaviours → similar benefits
- Residential care staff education beneficial
- Cognitive stimulation similar effects

Livingston G et al Am J Psychiatry 2005; 162:1996-2021







Psychological approaches to BPSD

- Music therapy
- Snoezelen
- ? Sensory stimulation
- Interventions that changed visual environment looked promising, but ...
 - ... ⇒ research required

Useful during treatment but not long term

> ¹Livingston G et al Am J Psychiatry 2005; 162:1996-2021







Individualised music¹



¹Gerdner L et al, Int Psychogeriatr 2000, 12, 49-65







Calming music and/or hand massage



Remington, Nursing Research, 2002







Novel strategies

- Humour therapy
- Volunteers
- Music, singing, dance therapy
- Integrating kindergarten/ babies











Humor therapy: SMILE study

- 20% reduction in agitation
- Effect size = antipsychotic medications for agitation
- Adjusting for dose of humour therapy
 - Decreased depression
 - Improved quality of life

Low LF et al BMJ Open 2013 Brodaty et al Am J Ger Psych 2014 Low LF et al JAMDA 2014










Key elements

- Engagement
- Understanding
- Time

Barriers

- Time
- Money
- Staff
- Attitudes
- Training









Pharmacological interventions







Anti-Alzheimer medications







ChEls & BPSD

- Some benefit, statistically significant in some reviews but questionable clinical significance
- Individual Sx may be more susceptible: apathy, hallucinations, aberrant motor behaviour, delusions, anxiety, depression
- Trinh N-H et al, 2003
- Rodda et al, 2009
- Campbell et al, 2008

www.ipa-online.org







Memantine on BPSD

- Mixed results
 - Several negative results ¹⁻²
 - Some positive results ³⁻⁴
- Specific benefits reported for cluster of aggression, hallucinations & delusions

¹ <u>Reisberg B et al, 2003; ² Van Dyck et al, 2007;</u> ³ Tariot P et al, 2004 ; ⁴ Gauthier et al (2005), IJGP, 20, 459-464







Antidepressants







Sertraline for treatment of depression in AD: Wk-24 Outcomes (DIADS-2)

- 67 Sertraline, 64 placebo; 12 wk RCT + 12 wk
- No between-groups diff. in depression response
 - in CSDD score
 - remission rates
 - secondary outcomes
- SSRI associated > adverse events of diarrhoea, dizziness, dry mouth, pulmonary SAE (pneumonia)

Weintraub D et al. Am J Ger Psych, 2010;18:332-340









Effects of citalopram on BPSD

- Improve hallucinations and delusions (= antipsychotics)
- Improve agitation
- 60% ↓ irritability and apathy (but n.s.)
- ↓ hallucinations (statistical but ?clinical significance)

Pollock et al. (2002). Am J Psych 159: 460-465 Pollock et al. (2007). Am J Geriatr Psych 15: 1-11 Siddique et al. (2009) J Clin Psychiatry 70(6):915-918







CitAD RCT – citalopram & agitation



- Significant better with citalopram
 - Cognitive & cardiac adverse
 effects may limit
 effectiveness at
 30mg/day

Porsteinsson et al. JAMA. 2014;311(7):682-691. doi:10.1001/jama.2014.93







Anticonvulsants for BPSD¹

- Literature review of 7 RCT (2 carbamazepine & 5 valproate)
- Results (treatment vs placebo):
 - −1 study: sig. ↓ BPSD
 - 5 studies: no sig. difference
 - -1 study: sig. ☆ BPSD
 - AEs more frequent in treatment groups
- Might be beneficial for some patients
- Not recommended for routine use

¹ Kanovalov et al (2008). Int Psychogeriatr, 20:2







Antipsychotics











Drugs

O Home O Drugs O Drug Safety and Availability O Postmarket Drug Safety Information for Patients and Providers.

	Drug Safety and Availability	Public Health Advisory: Deaths with Antipsychotics in
	Postmarket Drug Safety Information for Patients and Providers	Elderly Patients with Behavioral Disturbances The issues described in this communication have been addressed in product labeling (see Drugs@FDA).
	Drug Safety Information for Healthcare Professionals	
		A/11/1000E



Effects of antipsychotics



- Meta-analysis from 13 studies¹:
 - Mean ES in Rx = 0.45
 - Mean ES in placebo = 0.32
- Effect sizes of atypical antipsychotics for BPSD are medium, not statistically better than placebo
- Increased rate of stroke²
- Increased mortality³
- Increased AEs in general

¹ Yury C & Fisher J, Psychotherapy and Psychosomatics 2007
 ² BrodatyH et al, J Clin Psychiatry 2003
 ³ Schneider L, 2005







Side effects of antipsychotics

- Sedation
- Dizziness
- Extra pyramidal symptoms
- Falls
- Metabolic syndrome
- Weight gain
- Orthostatic hypotension

- ↑ Prolactin gynaecomastia
- Anticholinergic side effects (e.g. glaucoma, urinary outflow)
- Stroke
- Death







Continuing vs stopping antipsychotics in dementia patients?

Ballard 2008

- 12 months RCT, continuous use vs placebo
- For most AD patients withdrawal no detriment
- Subgroup of pts with more severe symptoms (NPI ≥ 15) might benefit from continued Rx
- Devanand 2012
- Pts who responded for psychosis or agitation
- Discontinuation → higher rate of relapse

Ballard et al 2008 PLOS Medicine, 5:587-599; Devanand DP_NEJM, 2012

DART-AD – mortality associated with continuous Rx A Modified intention-to-treat (mITT) population



¹ Ballard et al, 2009 Lancet Neurology, 8, 151–157







The HALT study Halting Antipsychotic use in Long-Term care









HALT Protocol

- A single arm 12-month longitudinal study in 24 aged care facilities of at least 60 beds in urban and rural NSW
- Resident participants assessed ≈1-4 weeks prior to deprescribing (T1 & T2)
- Re-assessed 3, 6 and 12 months later (T3 T5)







HALT results to date

- Deprescribing commenced for all 134 participants assessed at baseline (T2)
- Antipsychotics ceased for 125 by 1st follow-up
- 39 deprescribed prior to planned HALT timeline
- 15 recommenced regular a'psychotic before f-up
- A further 9 at 6 months (T4; data collection)







NPI and CMAI



CMAI Total Score

p > .05

p < .05







HALT - Challenges

- Difficult to recruit NHs, GPs, Families
- Lack of education around BPSD for care staff,
 GPs and families
- Task oriented nursing care, change process to implementing PCC, family expectations
- Presence of "nurse led" prescribing of antipsychotics







HALT - Discussion

- Champion management partnerships essential to success
- Knowledge, awareness and shared confidence in non-pharmacological approaches
- Deprescribing successful some started early!
- Small sub-group re-prescribed reason why??
- Σ BPSD stable 6 months after deprescribing







Antipsychotics for ...

- Screaming X
- Wandering X
- Intruding into other people's rooms X
- Aggression ?√ (but not first line)
- Delusions and hallucinations ?√ (but not 1st)

Cochrane: aim to discontinue antipsychotics¹

¹ Declercq T et al, Cochrane Review, 2013







Benzodiazepines

- PBO RCTs: BDZ decrease agitated behaviours during short-term use
- Short-acting BDZs eg oxazepam or lorazepam that do not accumulate are better
 - most effective if used for short periods at low doses (e.g., lorazepam 0.5–2.0 mg/day)
- AEs = Sedation, falls, confusion, amnesia
 (Chesrow et al., 1965; Kirven and Montero, 1973; Covington, 1975; Coccaro et al., 1990)







Analgesics

- Cluster RCT, 60 NHs, 352 residents, 8 + 4wks
- Mod-severe dementia, CMAI > 39
- Stepped analgesia vs usual care
- \approx 70% of residents paracetamol 1gm tds
- CMAI ↓17% (9.6 vs 3.4, p<.001)
- CMAI score [↑] 4 weeks after stop analgesia
- NPI & Pain scores significantly ↓

Husebo BS et al, BMJ, 2011;343:d4065 doi: 10.1136bmj.d0465







Legal consent for psychotropics

- Depending on jurisdiction a Person Responsible must give consent
- Survey of 3 NHs; 77 residents without capacity to give informed consent; on psychotropics¹
- Only 6.5% written consent
- + 6.5% partial or attempted consent

¹ Rendina N et al, 2009







Prevention of BPSD

- Person centred care and environment
- Right level of stimulation
- Attention to environment
- Treat physical disorders quickly









An Australian Government Initiative

Behaviour Management A Guide to Good Practice

Managing Behavioural and Psychological Symptoms of Dementia



dementia behaviour management advisory service





Helping Australians with dementia, and their carers

Screenshots





interventions





Agitation

Psychosocial/environmental interventions

Acupressure

Scientific quality of research: Moderate Outcomes: Positive; 1 large & 1 small pilot study

Animal-assisted therapy

Scientific quality of research: Limited

Outcomes: Positive; 1 small case series

Aromatherapy with lavender oil inhalation

Scientific quality of research: Moderate

Outcomes: Positive; 1 study

Bright light therapy

Scientific quality of research: Moderate

Outcomes: No benefit; 1 study *MAY INCREASE AGITATION

Closing Group intervention, small group, resident driven program

Scientific quality of research: Limited

Outcomes: Positive; 1 small study





Wandering

Clinical scenario

Presentation

Mr E is a 63 year old Aboriginal man who moved to Adelaide from a regional community when he was 16. He lived with his wife until she died several years ago. While raising their family of five children, they maintained strong community links with Aboriginal friends and family in Mr E's original community. His connection to Country has remained very important to him. Family and community members have been supporting Mr E in the family home with the assistance of an Aboriginal-specific community service and this arrangement has been working well until recently. On three occasions in the past month Mr E has been found after dark some distance from home. underdressed for the weather and distressed. On the most recent occasion, a concerned passer-by alerted police after Mr E was unable to provide his address or contact details for his family. When the police approached Mr E he became uncooperative and verbally aggressive. Police ultimately located Mr E's daughter who collected him from the local police station to take him home.

Assessment



In order to reduce the ereceptine hohouigue





Clinical conclusions about management of BPSD



"Dr,

Mrs Smith-Jones is hitting the nurses, disrupting the other residents and being impossible. Can you prescribe something?"







Clinical conclusions about management of BPSD



"Dr, Mr Smith-Jones is hitting the nurses, disrupting the other residents and being impossible. Can you prescribe something?"







Clinical practice 1

- Ask nurses to monitor behaviours what, when, what happens before, during and after?
- How often, when, what are precipitants?
- Exclude pain, UTI
- Determine cause
- Correct reversible factors eg stimulation level
- Start with psychological & environmental intervention(s)
 - except if urgent or sometimes concurrent
 - informed consent







II: Understand the person -Don't just label the behaviour

- Why is this person behaving this way now?
- Aetiological map \rightarrow management plan
- Different approaches often together
- Be creative
- Document
 - Monitor outcome
 - Partnership with family/ carers







Interpers

Psycholog

Soc-env

Biological
Clinical practice 2

- No cause can be found or correctable
- Try psychosocial treatments
 - not sure how?
 - \rightarrow BPSD Guide on your app
 - → call DBMAS or local psychogeriatric team
- Psychosocial treatment fail
- Consider pharmacological treatment
- 1st need informed consent from patient or proxy (*Person Responsible*, Guardianship Act)
- Start low and go slow







Rx for BPSD - summary

- Analgesic stepped approach
- Cholinesterase inhibitors for apathy
- Memantine ?benefit for agitation/aggression/ delusions/ hallucinations
- Antidepressants citalopram, sertraline, venlafaxine, mirtazapine
- Risperidone 0.5 2mg/day; modal = 1mg
- Olanzapine 5mg/day, up to 10mg/day
- Oxazpeam 7.5 15 mg as short term rescue Rx
- Carbamazepine, valproate titrate dose against response, SEs and blood level







Prescribing & Deprescribing Psychotropics

- Review regularly
 - At least after 3 months
 - Trial reduction, monitor behaviours
- Resident arrived from hospital on psychotropics
 - Find out why
 - If primary psychiatric diagnosis eg Sz, BAD
 - → seek psychiatric review
 - If not, trial reduction after pt. settled







When everything fails?

- You do everything right but BPSD continues
- Risk to other residents/ staff/ family
- Special care units
 - Medium term \rightarrow transfer back to mainstream
- Intensive care unit for very aggressive/ violent

Brodaty H, Draper B and Low LF Medical Journal of Australia 2003









Summary ... d'oh!

- Drug treatments limited benefit and

 → side effects yet 30% of residents
 in Australia are on antipsychotics
 and half on ≥1 psychotropic
- Most drug Rx given without required consent¹
- Psychosocial and environmental therapies beneficial with effect size <u>></u> drug Rx

Rendina N et al, IJGP, 2009







Summary ... d'oh!

- So why are nursing homes not engaging more?
- Why is the knowledge not being translated into practice?
 - Training too little?
 - Cost too much?
 - Time not enough?
 - Residents, families, system??



D'oh!









How to make good care Practice As Usual?

- Incentives for owners, managers, staff
- Accreditation standards
- Drive demand families, residents
- Show cost effectiveness
- Publicise, communicate
- Leadership, training







Conclusions

- BPSD common
- Prevent BPSD PCC, environment, titrate stimulation, CG and staff training
- Drugs have limited effects and AEs
- Psychosocial treatments have 1 evidence
- Problem is implementation
- Practical suggestions for working with facilities
- Need policy recognition too accreditation standards, government policy, research support







Conclusions

- Pharmacotherapy
 - modestly effective for BPSD
 - Prescribe judiciously
 - Need medico-legal informed consent
 - Start low and go slow
 - Importance of deprescribing
 - Review regularly, at least 3 monthly
 - Trial reductions







Thank you

- **Dementia Collaborative Research Centre**
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