



6° CONVEGNO NAZIONALE SUI CENTRI DIURNI ALZHEIMER


FONDAZIONE
CASSA DI RISPARMIO
DI PISTOIA E PESCIA

Centro Monteliveto

"Casa dell'Inziano"

ANTICHI E NUOVI INTERVENTI TERAPEUTICI IN
CENTRO DIURNO ALZHEIMER

12.10

La demenza nel disabile intellettivo che invecchia
Luc De Vreese (Modena)

Introduce: *Nicola Cariglia (Pistoia)*

15 - 16 Maggio 2015

Auditorium

Via Panconi, 14 - Pistoia

Research Article

Prevalence of Depression and Dementia among Adults with Developmental Disabilities in Manitoba, Canada

Shahin Shooshtari,¹ Patricia Joan Martens,² Charles A. Burchill,² Natalia Dik,²
and Saba Naghipur³

TABLE 2: Dementia prevalence (per 100 population), 2000/01–2004/05.

Study population and age groupings	Total N	Number with dementia	Prevalence (95% CI)
DD cohort			
20–54 Yrs	1,401	73	5.21 (4.05, 6.37)
55+ Yrs	218	30	13.76 (9.19, 18.33)
Matched comparison group (no DD)			
20–54 Yrs	2,798	39	1.39(0.96, 1.83)
55+ Yrs	433	15	3.46 (1.74, 5.19)

La prevalenza di demenza è da 4 a 5 volte superiore rispetto alla popolazione generale

La nuova longevità

Changes in mortality and causes of death in the Swedish Down syndrome population.

- All individuals with DS that died between 1969 and 2003 in Sweden, and all individuals born with DS in Sweden between 1974 and 2003 were included.
- Data were obtained from the Swedish Medical Birth Register, the Swedish Birth Defects Register, and the National Cause of Death Register.
- Median age at death has increased by **1.8 years per year**

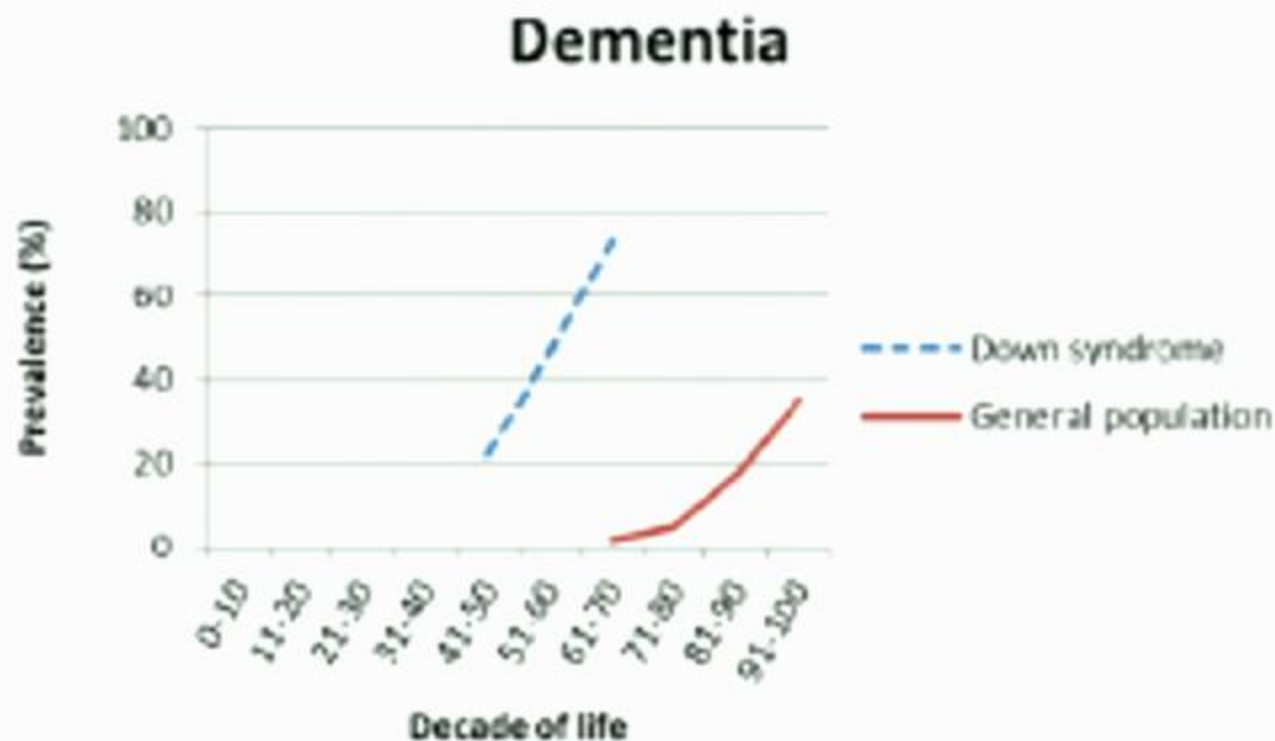
Syndrome	40-50	50-60	60-70	70-80	80-90	years
Angelman					72	
Cerebral Palsy			55			
Cornelia de Lange				71		
Down syndrome					82	
Fragile X						87
Phenylketonuria (without diet)				70		
Prader-Willi				72		
Rett			55			
Sanfilippo				69		
Tuberous sclerosis			50			
Williams			55			
						life expectancy with maximum age

Coppus AM. Dev Disabil Res Rev 2013;18(1):6-16.

- Oggi in Italia le persone con SD sono circa 48.000, di cui il **13%** ha più di **44** anni
- La spettanza di vita alla nascita è di **61,6** anni per i maschi e di **57,8** anni per le femmine

Epidemiologia

Dementia in Alzheimer's disease (DAD)



Nonostante il verificarsi di AD patologia nel cervello di individui con SD a partire dall'età di 40 anni (Wisniewski et al., 1994), la DAD esordisce in media attorno ai 50-55 anni di età (con un *range* tra 38-70 anni) con tassi di incidenza crescente con l'avanzare dell'età (da 2,53/100 persone-anno a 13,31/100 persone-anno rispettivamente nelle fasce di età <50 e ≥ 65 anni).

Coppus A. et al. *J Intellect Disabil Res* 2006;50:768-77.

Glasson EJ et al. *J Intellect Disabil Res* 2014;58:393-8.

Nelle persone con SD la demenza nella Malattia Alzheimer (*Dementia in Alzheimer's Disease*; Prasher, 2005) è la causa più importante di fragilità e di mortalità (fino al **30%**).

Coppus AM. *Dev Disabil Res Rev* 2013;18(1):6-16.

Prasher V.P. *Alzheimer's disease and Dementia in Down Syndrome and Intellectual Disabilities*, Abingdon: Radcliffe Publishing, 2005.

Survival in Elderly Persons with Down Syndrome

Antonia M. W. Coppus, MD,^{†} Heleen M. Evenhuis, MD, PhD,[‡] Gert-Jan Verberne,[†] Frank E. Visser, MD, PhD,[§] Ben A. Oostra, PhD,^{||} Piet Eikelenboom, MD, PhD,[#] Willem A. van Gool, MD, PhD,[#] A. Cecile J. W. Janssens, PhD,^{**} and Cornelia M. van Duijn, PhD^{*}*

Relative preservation of cognitive and functional ability is associated with better survival in this study population. Clinically, the most important disorders in persons with DS that are related to mortality are dementia, mobility restrictions, visual impairment, and epilepsy but not cardiovascular diseases. Also, level of intellectual disability and institutionalization are associated with mortality. *J Am Geriatr Soc* 56:2311–2316, 2008.



57



58



59



59.9

**Stadiazione compressa per
adulti con Sindrome di
Down**

**Primo Stadio o
Lieve**
da 2 a 4 anni o più

**Stadio Medio- o
Moderato**
aa 2 a 10 anni

**Ultimo stadio o
Severo**
da 1 a 3 anni o più

Stages of Alzheimer's Disease

Research Article

Factors Determining Disease Duration in Alzheimer's Disease: A Postmortem Study of 103 Cases Using the Kaplan-Meier Estimator and Cox Regression

R. A. Armstrong

TABLE 1: Demographic details of the 103 cases used in the study (N : number of cases; FAD: familial Alzheimer's disease; SAD: sporadic Alzheimer's disease). Data for age at death, duration, and disease onset are means with range and standard deviations in parentheses.

Patient group	N	Age at death (years)	Duration (years)	Onset (years)
Early-onset FAD	19	61.9 (46–74, 10.8)	11.1 (3–20, 6.5)	50.7 (38–59, 6.5)
Early-onset SAD	22	70.4 (57–88, 11.1)	16.0 (6–30, 9.3)	54.4 (49–58, 4.2)
Late-onset FAD	12	77.4 (70–85, 5.1)	7.0 (1–15, 4.6)	70.4 (61–84, 7.6)
Late-onset SAD	50	80.1 (70–98, 6.6)	6.8 (1–21, 4.5)	73.5 (62–93, 7.2)



Incidence of dementia in older adults with intellectual disabilities



Andre Strydom^{a,*}, Trevor Chan^b, Michael King^a, Angela Hassiotis^a,
Gill Livingston^a

Anche nelle altre forme di DI, la demenza è molto frequente: quasi un **terzo** degli ultra60enni sviluppa una demenza conclamata, con tassi di incidenza di **circa cinque** volte superiore rispetto a quelli della popolazione generale (54,6/1000 persone-anno con un picco di 97,8/1000 persone- anno nelle fasce di età tra 70-74 anni). Prevala la demenza Alzheimer ($\approx 50\%$) come nella popolazione generale.

Screening

ARTICOLO ORIGINALE

Uno studio di attendibilità della versione italiana della scala Dementia Questionnaire for Persons with Intellectual disabilities (DMR)

A reliability study of the Italian version of Dementia Questionnaire for Persons with Intellectual disabilities (DMR)

LUC PIETER DE VREESE¹, ULRICO MANTESSO², MARCO SCARAZZINI³,
CRISTINA MENEGATTI⁵, TIZIANO GOMIERO⁴

Tab. 2 Gli *item*, le sottoscale e somma dei punteggi del DMR.

Sottoscale	N item	Punteggio (minimo-massimo)
1. Memoria episodica recente	8	0 - 16
2. Memoria episodica remota e autobiografica	7	0 - 14
3. Orientamento spaziale e temporale	7	0 - 14
Somma degli Scores Cognitivi (SOS)	22	0 - 44
4. Comunicazione verbale	4	0 - 8
5. Abilità di base	8	0 - 16
6. Umore	6	0 - 12
7. Attività e interessi	6	0 - 12
8. Comportamenti problematici	6	0 - 10
Somma degli Scores Sociali (SCS)	30	0 - 60



Contents lists available at ScienceDirect

Research in Developmental Disabilities



Review

A systematic review on assessment instruments for dementia in persons with intellectual disabilities



Elisabeth L. Zeilinger*, Katharina A.M. Stiehl, Germain Weber

As an example for the group of informant-based assessment instruments, the *National Task Group on Intellectual Disabilities and Dementia Practices—Early Detection Screen for Dementia* (NTG—EDSD) can be mentioned. The assessment of dementia needs to focus on the individual changes in the respective person. To detect early signs of dementia it is widely recommended to compare an early assessed baseline with periodic re-assessments (e.g., Aylward, Burt, Thorpe, Lai, & Dalton, 1997; Deb & McHugh, 2010; Kalsy & Oliver, 2005). The NTG—EDSD incorporated this recommendation and is available in several languages, including English, French, German, Greek, Italian, and Spanish. Especially in Europe a multi-lingual availability of instruments is important in order to make data comparable. Combined with the advantage of being a baseline assessment the NTG—EDSD has the potential of becoming an agreed upon procedure for the assessment of dementia in persons with ID.

Report NTG

'My Thinker's Not Working': A National Strategy for Enabling Adults with Intellectual Disabilities Affected by Dementia to Remain in Their Community and Receive Quality Supports



- ▣ Fattori relativi alla popolazione
- ▣ Le sfide per le persone affette da demenza e DI
- ▣ Supporti della comunità
- ▣ Educazione e formazione
- ▣ Finanziamento
- ▣ Possibili soluzioni
- ▣ Piano d'azione e (raccomandazioni) per le persone con DI e demenza

Available from: www.aadmd.org/ntg
www.psicogeriatra.it

BADL - ATTIVITÀ DI BASE DELLA VITA QUOTIDIANA

KATZ INDEX OF INDEPENDENCE IN ACTIVITIES OF DAILY LIVING

1. Fare il bagno (vasca, doccia, spugnature)

- 1. Fa il bagno da solo (entra ed esce dalla vasca da solo)
- 2. Ha bisogno di assistenza soltanto nella pulizia di una parte del corpo (es. schiena)
- 3. Ha bisogno di assistenza per più di una parte del corpo

DSQIID. Dementia Screening Questionnaire for Individuals with Intellectual Disabilities.

DEB, S., HARE, M., PRIOR, L. and BHAUMIK, S. (2007), BRITISH JOURNAL OF PSYCHIATRY, 190, 440-444. doi: 10.1192/bjp.bp.106.024984

Questionario Parte 2

Esempio: Non Riesce a lavarsi/fare il bagno senza aiuto.

Sempre stato così.

Se la persona ha sempre avuto bisogno di aiuto

Sempre stato così, ma peggiora.

Se la persona sembra aver avuto un declino delle sue capacità in quest'area

Sintomo nuovo

Se la persona aveva acquisito questa abilità nell'età adulta e l'ha persa recentemente

Non è il suo caso (non applicabile)

la persona si lava senza nessun aiuto ed è sempre stato così

1	Non riesce a lavarsi e/o a fare il bagno senza aiuto	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
2	Non riesce a vestirsi senza aiuto	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
3	Si veste in modo inappropriato (es., confonde avanti e dietro, o incompleto)	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
4	Si spoglia in contesti inappropriati (es., in pubblico)	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
5	Necessita di assistenza per mangiare	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
6	Necessita di assistenza per usare i servizi igienici	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
7	Incontinente (incluso anche incidenti occasionali)	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
8	Non inizia una conversazione	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
9	Non riesce a trovare le parole	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
10	Non riesce a seguire semplici istruzioni	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
11	Non riesce a seguire più di una istruzione per volta	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
12	Si interrompe a metà di un compito	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
13	Non riesce a leggere	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
14	Non riesce a scrivere (nemmeno il proprio nome)	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
15	Cambiamenti nel ritmo del sonno (dorme di più o di meno)	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
16	Si sveglia frequentemente durante la notte	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
17	Di notte è confuso	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
18	Dorme durante il giorno	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
19	Vagabonda di notte	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
20	Si disorienta in ambienti familiari	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
21	Vagabonda (incluso fuori dall'abitazione)	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
22	Perde il senso del tempo (ora del giorno, giorno della settimana, stagione)	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
23*	Non sicuro nel camminare su piccole fessure, linee sul pavimento o superfici irregolari.	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
24	Il cammino instabile, perde l'equilibrio	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
25	Non riesce a camminare senza aiuto	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
26	Non riconosce persone familiari (operatori/parenti)	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
27	Non ricorda i nomi di persone familiari	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
28	Non ricorda eventi recenti	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
29	Ritiro dalle attività sociali	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
30	Ritiro dai contatti personali	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
31	Perdita di interesse in hobbies/attività	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
32	Sembra vivere in un mondo tutto suo	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
33	Comportamenti ossessivi o ripetitivi (es., svuota ripetutamente gli armadi)	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
34	Nasconde o accumula oggetti	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
35	Perde oggetti	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
36	Ripone oggetti familiari in luoghi impropri	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
37	Non sa cosa fare con oggetti familiari	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
38	Sembra insicuro/a	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
39	Sembra ansioso/a o nervoso/a	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
40	Sembra depresso/a	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
41	Mostra aggressività (verbale o fisica)	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
42	Convulsioni/Epilessia	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
43	Parla da solo/a	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Dementia Screening Questionnaire for Individuals with Intellectual Disabilities-Italian version (DSQIID-I): a multicentre validation study in aging adults with Down syndrome and other forms of Intellectual Disabilities

Tiziano Gomiero^a, Marco Bertelli^b, Shoumitro Deb^c, Elisabeth Weger^a, Annachiara Marangoni^a, Elisa De Bastiani^a, Ulrico Mantesso^a, Luc Pieter De Vreese^d

a Project DAD (Down Alzheimer Dementia) ANFFAS Trentino Onlus, Trient, Italy

b Research and Clinical Centre (CREA) of San Sebastiano Foundation, Florence, Italy

c Imperial College London, Department of Medicine, Division of Brain Sciences, UK

d Local Health Agency, Dementia Project, Modena, Italy

	N	%
male	162	58.7
female	114	41.3

1.1.4. Dichotomic Diagnosis of ID (DS vs. non-DS)

	N	%
SD	85	30.8
non SD	191	69.2

1.1.1. Mean age

	Minimum	Maximum	Mean	SD
Age	39	80	54.61	7.559



Sintomi di esordio

Behavioral changes in Down syndrome diagnosed with Alzheimer's dementia.

Behavioral changes	Example reference
Apathy	Jervis (1948) and Deb et al. (2007)
"Episodic noisy excitement"	Jervis (1948)
Irritability	Jervis (1948) and Stanton and Coetzee (2004)
Wandering & confusion	Jervis (1948), Prasher and Filer (1995) and Deb et al. (2007)
Destructive, aggressive or difficult behavior	Jervis (1948), Prasher and Filer (1995), Urv, Zigman, and Silverman (2008) and Stanton and Coetzee (2004)
Lethargy, withdrawal, loss of interest	Crapper, Dalton, Skoptiz, Scott, and Hachinski (1975) and Stanton and Coetzee (2004)
"Silliness"	Crapper et al. (1975)
Limited response to people	Crapper et al. (1975)
Social inadequacy, isolation	Urv et al. (2008) and Stanton and Coetzee (2004)
Extreme changes in appetite (typically weight loss)	Crapper et al. (1975)
Restlessness	Prasher and Filer (1995) and Stanton and Coetzee (2004)
Sleep disturbance	Prasher and Filer (1995), Deb et al. (2007), Urv et al. (2008) and Stanton and Coetzee (2004)
Incontinence	Prasher and Filer (1995)
Excessively uncooperative	Cooper and Prasher (1998) and Stanton and Coetzee (2004)
Anxiety & fearfulness	Urv et al. (2008) and Stanton and Coetzee (2004)
Sadness	Urv et al. (2008)
Stealing & general regressive behavior	Urv et al. (2008)
Personality changes	Stanton and Coetzee (2004)
Increased dependence	Stanton and Coetzee (2004)

Behavioral changes in Down syndrome diagnosed with Alzheimer's dementia.

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<p>As an explanation for this "frontal signs first" hypothesis, Azizeh et al. (2000) suggested that the sequence of brain regions affected by Alzheimer's dementia may differ with earlier amyloid deposition in the frontal lobes in the Down population while in the general population the hippocampus (and thus episodic memory) would atrophy first. Holland et al. (1998) extended Mortimer's (1988) reserve capacity model by implying that early compromise was to be found in the frontal lobes in Down syndrome. Also relevant to this discussion is that the frontal lobes have been found to be underdeveloped in Down syndrome (Crome & Stern, 1972) and this could suggest that only a small amount of neuropathology may tip the threshold into impairments, especially on frontal tasks in these people.</p>	
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Prevalence of dementia in intellectual disability using different diagnostic criteria

A. STRYDOM, G. LIVINGSTON, M. KING and A. HASSIOTIS

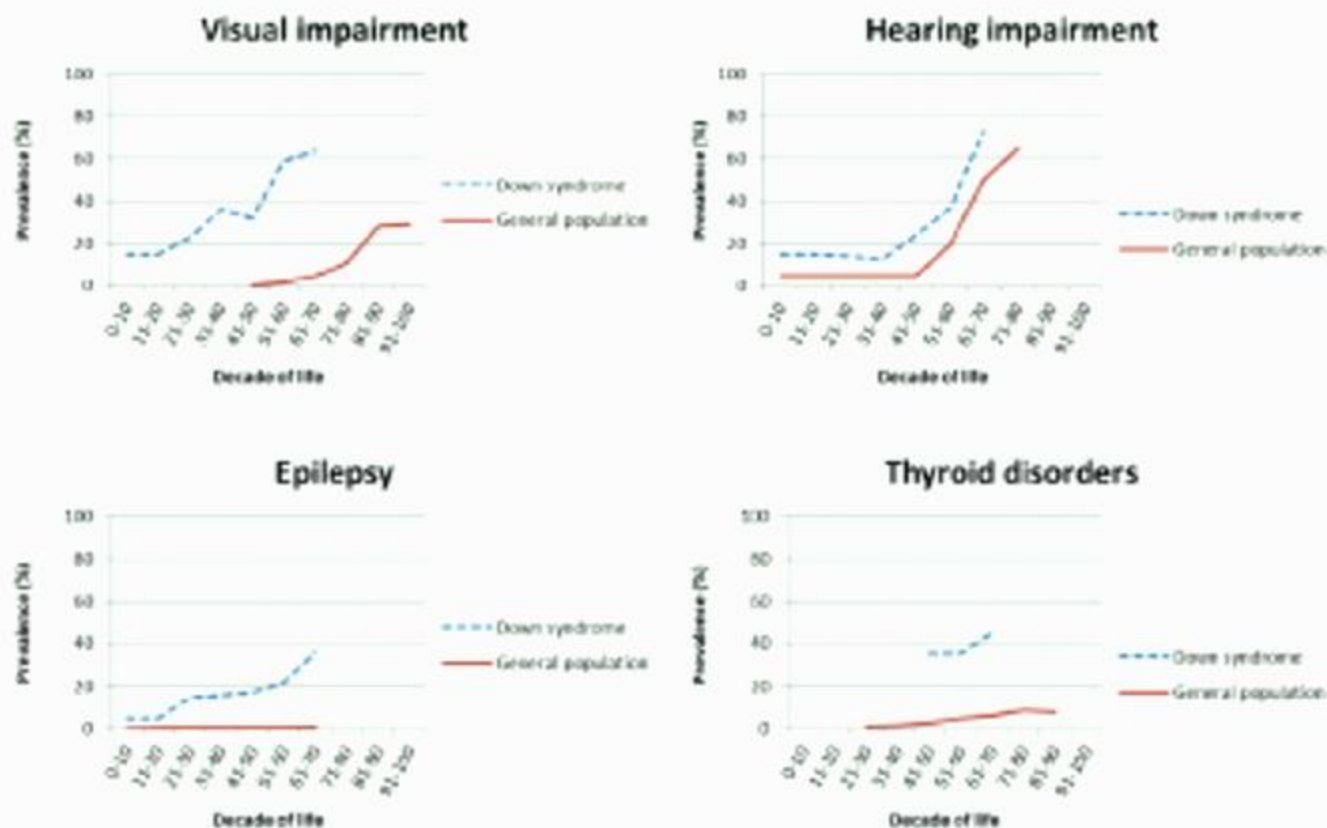
The most common initial symptom was **general deterioration in functioning**, followed by **behavioural or emotional change**.

Deterioration in memory or other cognitive functions was rarely noticed to be prominent in the early stages of the disorder. Other early symptoms included **episodes of confusion**.

Symptoms that significantly discriminated between those with and without dementia in those who screened positive were deterioration in **self-care ability**, deterioration in **IADL**, change in **memory**, development of **muddled thinking**, development of problems with **thinking ahead and planning**, and newly developed **perseveration**.

Declino cognitivo e comorbidità organica e/o psichiatrica

(e) *uncertain with complications*: criteria for definite dementia had been met, but symptoms might be caused by some other substantial concern, usually a medical condition unrelated to a dementing disorder (e.g., loss of vision, poorly resolved hip fracture, loss of social support network due to relocation)



Glasson EJ et al. *J Intellect Disabil Res* 2014;58:393-8.

Factors associated with depression and anxiety in older adults with intellectual disabilities: results of the healthy ageing and intellectual disabilities study

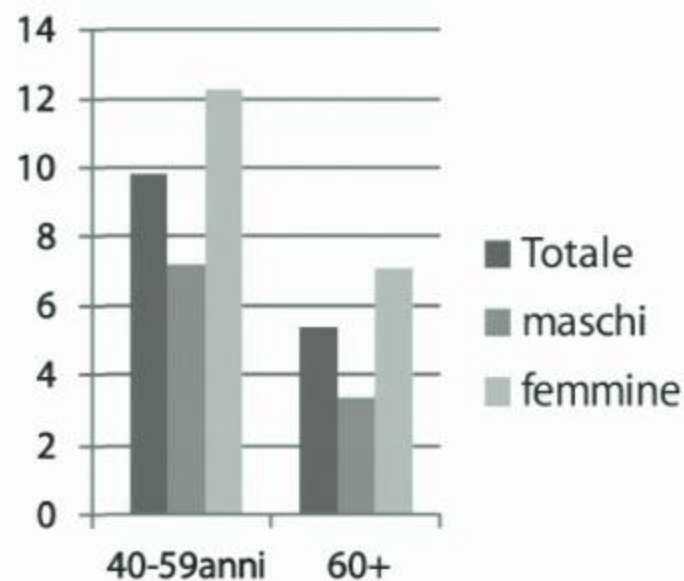
Heidi Hermans^{1,2} and Heleen M. Evenhuis¹

HA-ID study: 980 soggetti con DI, di cui 861 con SD; età media 61,5 anni (50-93)

Depression and anxiety (%)		
Increased depressive symptoms	166 (16.8)	138 (47.6)
Increased anxiety symptoms	161 (16.3)	128 (44.1)
Major depression	³	42 (14.5)
Anxiety disorders	³	23 (7.9)

³ Studied in 290 participants only

Increased depressive symptoms were positively associated with increased anxiety symptoms, number of life events during the past year and chronic diseases (heart failure, stroke, chronic obstructive pulmonary disease, coronary artery disease, diabetes mellitus and malignancy in the previous 5 years)

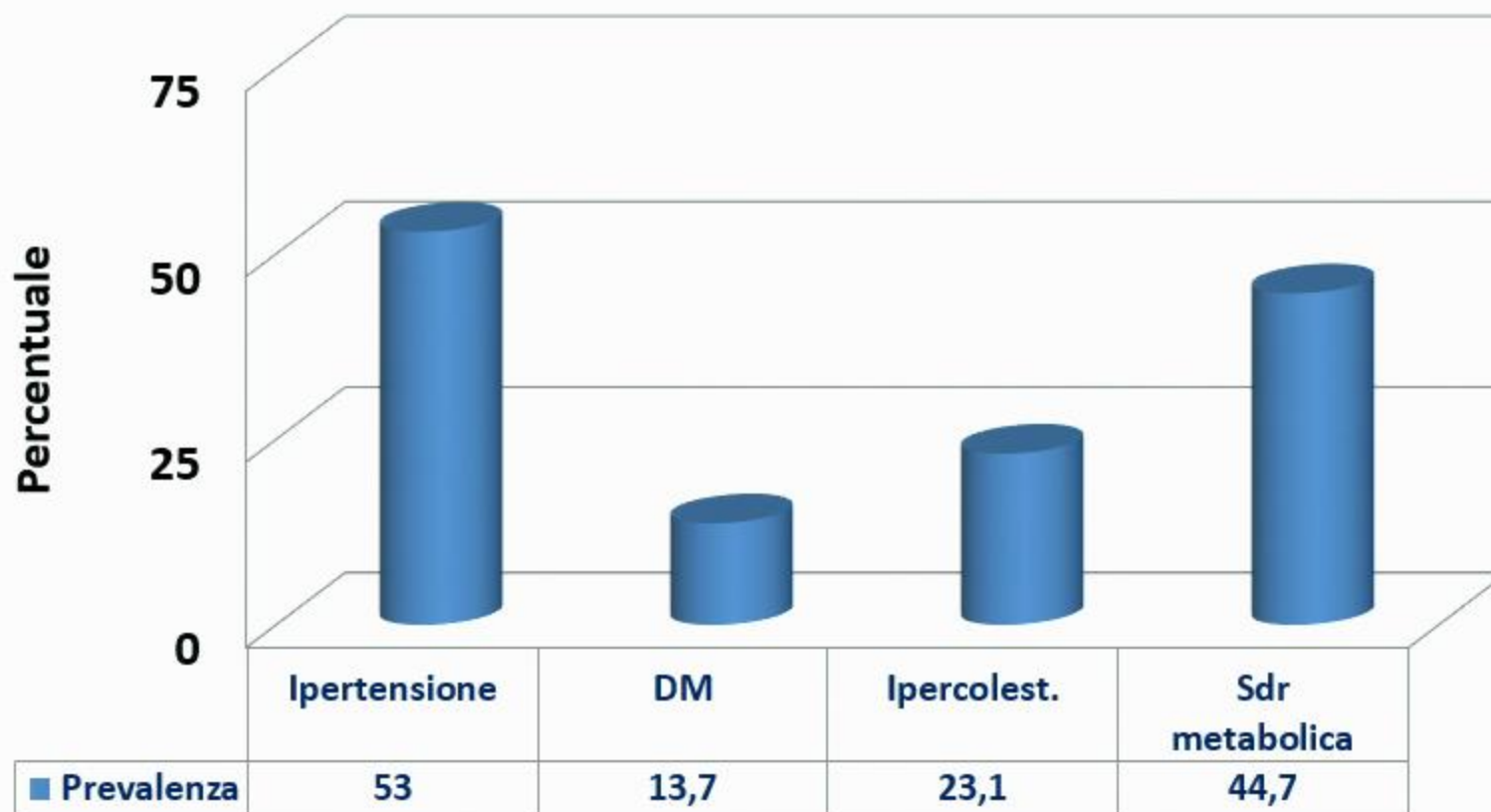




Cardiovascular risk factors (diabetes, hypertension, hypercholesterolemia and metabolic syndrome) in older people with intellectual disability: Results of the HA-ID study

C.F. de Winter^{a,b,*}, L.P. Bastiaanse^{a,c}, T.I.M. Hilgenkamp^{a,d}, H.M. Evenhuis^a, M.A. Echteld^a

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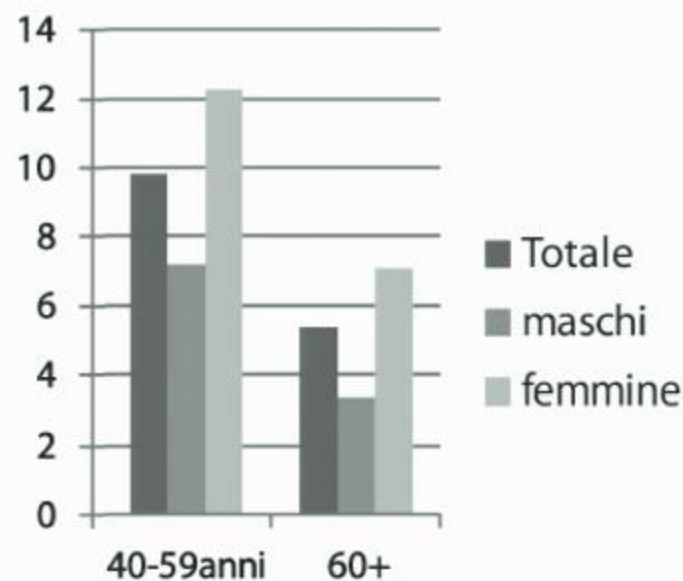
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II. Physical and Mental Health

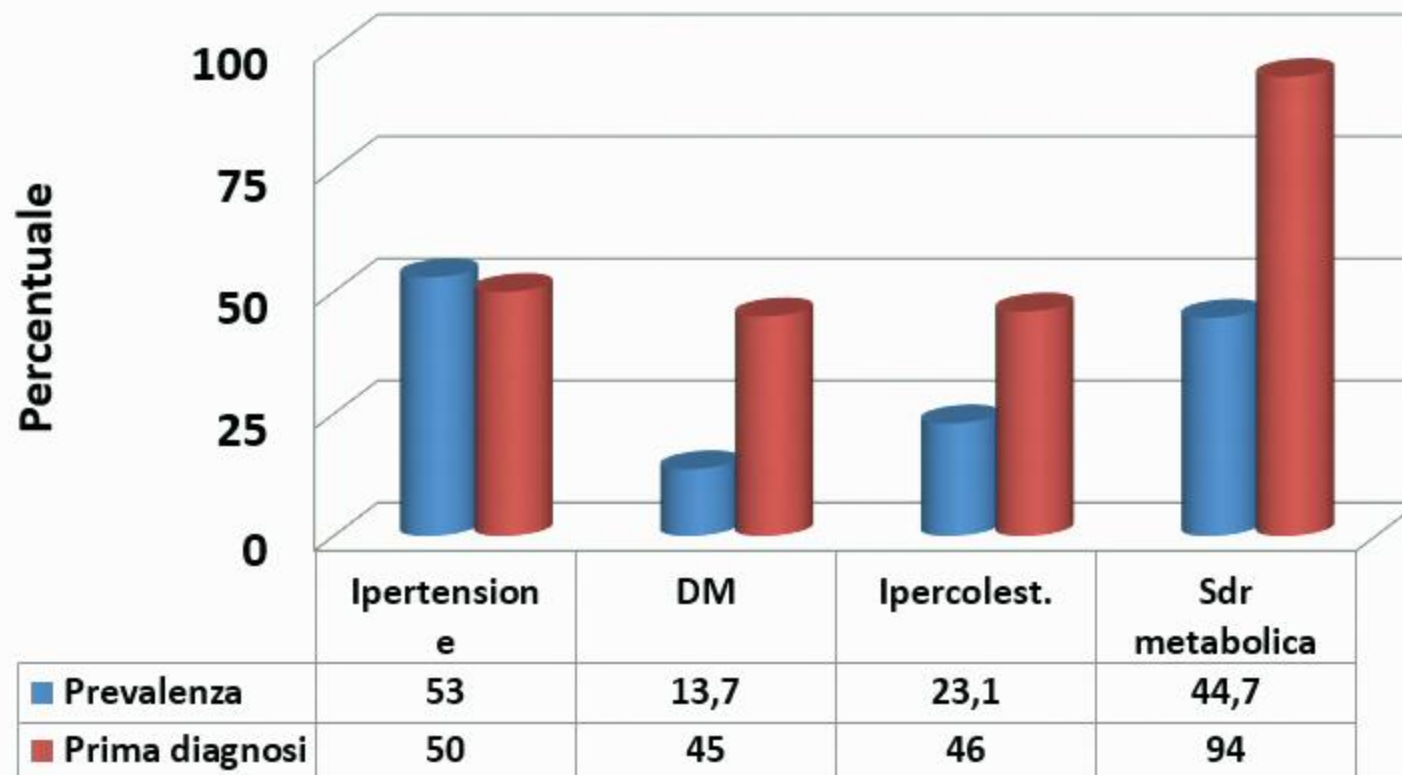
Many health problems remain unrecognised

- Communication/insight difficulties
- Physical limitations/reluctance of family members
- Behaviour problems
- Diagnostic overshadowing
- Lack of specialist care in ID



Cardiovascular risk factors (diabetes, hypertension, hypercholesterolemia and metabolic syndrome) in older people with intellectual disability: Results of the HA-ID study

C.F. de Winter^{a,b,*}, L.P. Bastiaanse^{a,c}, T.I.M. Hilgenkamp^{a,d}, H.M. Evenhuis^a, M.A. Echteld^a



RESEARCH ARTICLE

Open Access

A systematic review of hospital experiences of people with intellectual disability

Teresa Iacono^{1*}, Christine Bigby², Carolyn Unsworth³, Jacinta Douglas⁴ and Petya Fitzpatrick²

A number of reviews have explored underlying contributors to poor hospital experiences by people with intellectual disability [13,28-30]. These reviews have similarly revealed that some people with intellectual disability are fearful of hospital encounters, there is reliance on carers during the entirety of stays, and problem attitudes and limited knowledge of hospital staff, sometimes with dire outcomes.

Delirium nelle persone con DI: un gran sconosciuto

«I do not know of any screening tool or algorithm in older ID persons with or without dementia.

I do not know of any major research study in this area.

I think the field of ID is still trying to detect dementia *per se* rather than focus on subgroups.

Maybe in the future.»

Brief report

The triple challenges associated with age-related comorbidities in Down syndromeE. J. Glasson,¹ D. E. Dye² & A. H. Bittles^{3,4}

with their ill-effects exacerbated by **long-term medication, polypharmacy**, immobility and general inactivity.

REVIEW ARTICLE

Pharmacological Management of Behavioral and Psychiatric Symptoms in Older Adults with Intellectual Disability

Nicole Eady · Ken Courtenay · André Strydom

Abstract Given medical and social advances, the life expectancy of individuals with intellectual disability (ID) has increased dramatically, leading to a generation of older individuals with such disabilities. This review focuses on the pharmacological treatment of behavioral and psychiatric symptoms and disorders in older adults with ID. Older adults with ID often present with medical co-morbidities and mental health issues. Medication management of behavioral and psychiatric problems is complicated by a higher risk for adverse events, lack of decision-making capacity, and complex care networks. Some studies have shown that individuals with ID and co-morbid mental disorders are undertreated in comparison with those with similar disorders in the general population, resulting in poorer outcomes. However, older adults with ID are also at risk of polypharmacy, and older age is a risk factor for development of side effects. A general principle is that medication treatment for psychiatric disorders in older individuals with ID should be started at low dosages and

	N	%
Use as antiepileptic	33	12.0
Use as mood stabiliser	20	7.2
Use not specified	8	2.9
No ATC N03 drug use	215	77.9

22,1%

61 subjects (22.1% of the study sample) are under ATC N03 drug therapy

	N	%
FGA	40	14.5
SGA	14	5.1
FGA/SGA not specified	19	6.9
FGA e SGA in combined use	5	1.8
No ATC N05A use	198	71.7

28,3%

FGA: first generation antipsychotics, SGA: second generation antipsychotics; (28.3% of the study sample) are under ATC N05A drug therapy.

	N	%
No ATCN06 use	256	92.8
Antidepressant therapy	20	7.2

14 subjects in SSRI therapy, 4 subjects treated with Trazodone, one with NaSSA, and one subject with SNRI

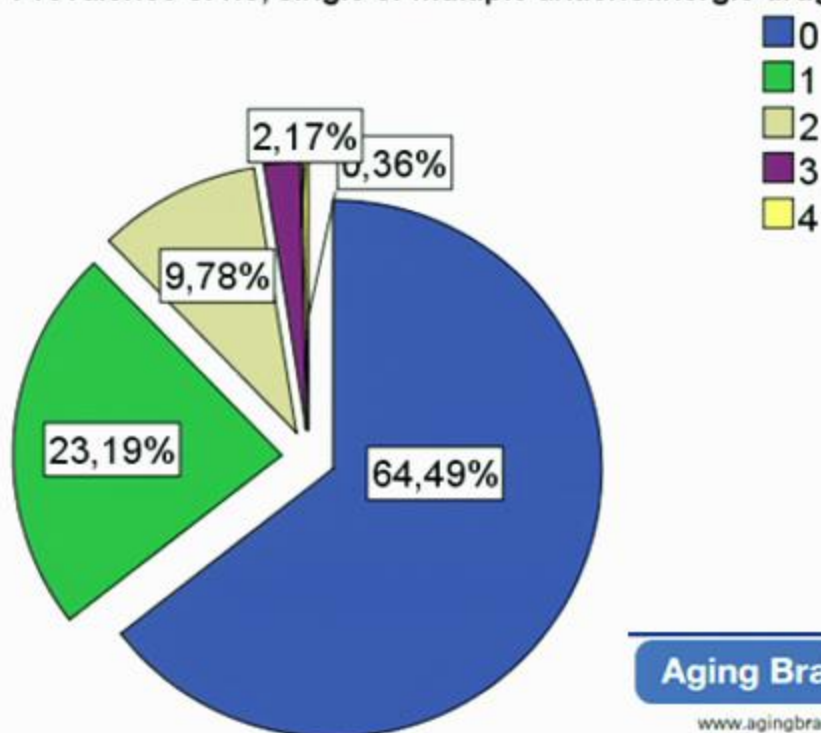
	N	%
No ATCN05B use	239	86.6
Anxiolytic use	32	11.6
Combined use (n=2)	5	1.8

13,4%

	N	%
No ATC N0C use	259	93.8
Hypnotic and sedative use	11	4.0
Combined use (n=2)	6	2.2

6,2%

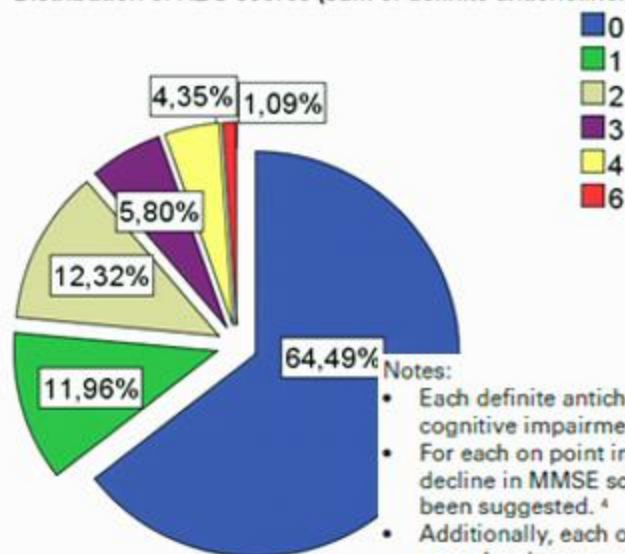
Prevalence of no, single or multiple anticholinergic drug use



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Distribution of ABC scores (sum of definite anticholinergics)



Notes:

- Each definite anticholinergic may increase the risk of cognitive impairment by 46% over 6 years.³
- For each one point increase in the ABC total score, a decline in MMSE score of 0.33 points over 2 years has been suggested.⁴
- Additionally, each one point increase in the ABC total score has been correlated with a 26% increase in the risk of death.⁴

Criteria diagnostici

Table 1 Criteria for dementia in the classification systems

Impaired domain/ symptoms	DSM-IV	ICD-10	DC-LD
Memory			
Short- and/or long-term	+	+	+
Higher cortical functions²			
Executive function	○		
Thinking		○	○
Judgement		○	○
Other cognitive skills			○
Information processing		○	
Aphasia/ language skills	○		
Apraxia	○		
Agnosia	○		
Behavioural and emotional function²			
Emotional lability		○	○
Irritability		○	○
Apathy		○	○
Social behaviour		○	○
Other criteria			
Change from premorbid state/decline in level of functioning ¹	+		+
Duration of at least 6 months		+	+
Exclusions			
Not caused by delirium	+	+	+
Not caused by mental illness	+		+

+, required for diagnosis.

1. The ICD-10 classification requires a decline in memory and other cognitive function, but does not have a separate criterion for change or deterioration in function.

2. At least one of the circled is required.

Diagnostic criteria for Psychiatric Disorders for Use with Adults with Learning Disabilities/Mental Retardation

Validity of Criteria for Dementia in Older People With Intellectual Disability

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poor outcomes at T2. Conclusion: Dementia diagnostic criteria show substantial reliability and satisfactory validity in ID. The diagnoses were, however, less stable than in the general population and some caution is advisable in those with more severe ID or additional sensory disability. MCI definitions require further consideration in the ID population. (Am J Geriatr Psychiatry 2013; 21:279–288)

Intellectual Disability, Mild Cognitive Impairment, and Risk for Dementia

Wayne P. Silverman¹, Warren B. Zigman², Sharon J. Krinsky-McHale², Robert Ryan², and Nicole Schupf³

The concept of mild cognitive impairment (MCI) has gained currency in the general population as a 'preclinical' stage of dementia in which the cognitive and functional losses are greater than would be expected as part of normal ageing, but are not sufficient to warrant formal diagnosis.

The validity of this term has been questioned, even within the normal population, and **there are practical difficulties of applying it to people with intellectual disability**

Trattamento farmacologico

Risultati negativi

- **Donepezil**

- De la Torre R. et al. *Prog Brain Res* 2012;197:1-14.

- **Memantina**

- Hanney M. et al. *Lancet* 2012;379:528-36.

- **Rivastigmina**

- Prasher V. et al. *Int J Geriatr Psychiatry* 2013;28:219-20.

Trattamento non farmacologico

The projected effect of risk factor reduction on Alzheimer's disease prevalence



Deborah E Barnes, Kristine Yaffe

At present, about 33.9 million people worldwide have Alzheimer's disease (AD), and prevalence is expected to triple over the next 40 years. The aim of this Review was to summarise the evidence regarding seven potentially modifiable risk factors for AD: diabetes, midlife hypertension, midlife obesity, smoking, depression, cognitive inactivity or low educational attainment, and physical inactivity. Additionally, we projected the effect of risk factor reduction on AD prevalence by calculating population attributable risks (the percent of cases attributable to a given factor) and the number of AD cases that might be prevented by risk factor reductions of 10% and 25% worldwide and in the USA. Together, up to half of AD cases worldwide (17.2 million) and in the USA (2.9 million) are potentially attributable to these factors. A 10–25% reduction in all seven risk factors could potentially prevent as many as 1.1–3.0 million AD cases worldwide and 184 000–492 000 cases in the USA.

Lancet Neurol 2011; 10: 819–28

- Diabete
- Ipertensione
- Fumo
- Obesità
- Depressione
- Inattività cognitiva
- Bassa scolarizzazione
- Inattività fisica

Guidelines for Structuring Community Care and Supports for People With Intellectual Disabilities Affected by Dementia

Nancy Jokinen*, Matthew P. Janicki[†], Seth M. Keller[‡], Philip McCallion[§], Lawrence T. Force[¶], and the National Task Group on Intellectual Disabilities and Dementia Practices

*University of Northern British Columbia, Prince George, British Columbia, Canada; [†]University of Illinois at Chicago, IL; [‡]American Academy of Developmental Medicine and Dentistry, Lumberton, NJ; [§]University at Albany, Albany, NY; and [¶]Mt. St. Mary College, Newburgh, NY, USA

The guidelines also provide information on nonpharmacological options for providing community care for persons affected by dementia as well as commentary on abuse, financial, managing choice and liability, medication, and nutritional issues.

Impact of Dementia-derived Nonpharmacological Intervention Procedures on Cognition and Behavior in Older Adults With Intellectual Disabilities: A 3-year Follow-up Study

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		SCU n = 14				DC n = 22				NH n = 24			
Sex	Male	7 (50%)				11 (50%)				12 (50%)			
DS Present	DS	10 (71.4%)				15 (66.2%)				5 (16.7%)			
Age	Mean	53.2				55.2				51.9			
	SD	(6.9)				(7.5)				(5.5)			
IQ values	Mean	29.9				32.7				27.3			
	SD	(11.6)				(14.8)				(13.3)			
Level of IQ N/%	Category	Pr	Se	Mo	Mi	Pr	Se	Mo	Mi	Pr	Se	Mo	Mi
		6 43%	4 28.5%	4 28.5%	0 0	8 36%	3 14%	8 36%	3 14%	13 54.2%	5 20.8%	4 16.7%	2 8.3%
DMR/SCS	Mean	25.4				13.6				15.1			
	SD	(10.9)				(14.3)				(10.9)			
DMR/SOS	Mean	19.3				16.9				13.9			
	SD	(10.6)				(12.4)				(8.7)			

Pr = Profound (≤ 25); Se = Severe (26–35); Mo = Moderate (36–49); Mi = Mild (≥ 50); SCU = special care unit; DC = day center; NH = nursing home

Gentle Care
Person-centred Care
Stimolazione multisensoriale
Attività a mediazione di un cane
Musicoterapia
Ambiente →

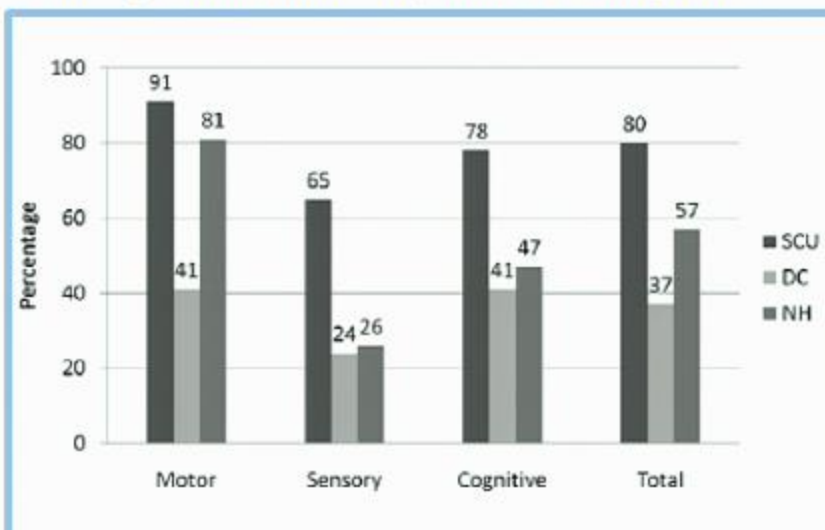


FIGURE 1

Appropriateness¹ of the environment in the three settings.

¹Based on the questionnaire (Chiogna & Dalprà, 2009).

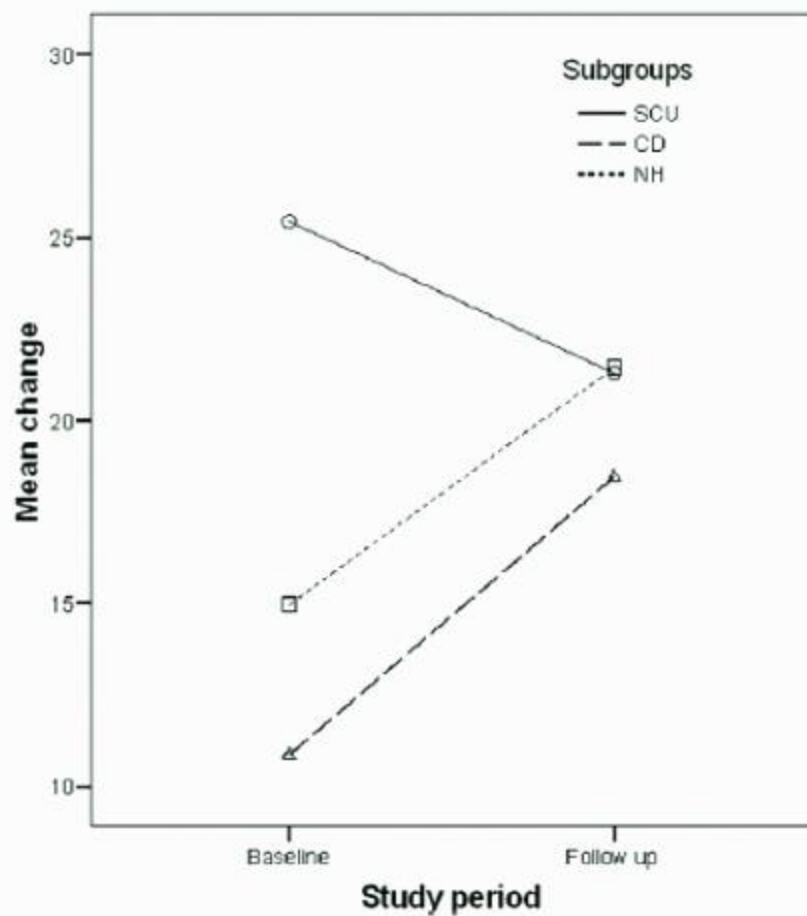


FIGURE 3
Mean DMR/SCS obtained by the three study groups across the 3-year period.

measures have not been included in this study. There were however, some indirect indicators of cost-effectiveness of the DAD Project, such as the hospitalization rate and number of traumatic falls. SCU residents were admitted to the hospital for 17 days against an average of 46 days in the control settings. There were only three falls among the SCU residents during the whole 3-year period, whereas the incidence of falls in the control settings was three falls per year.

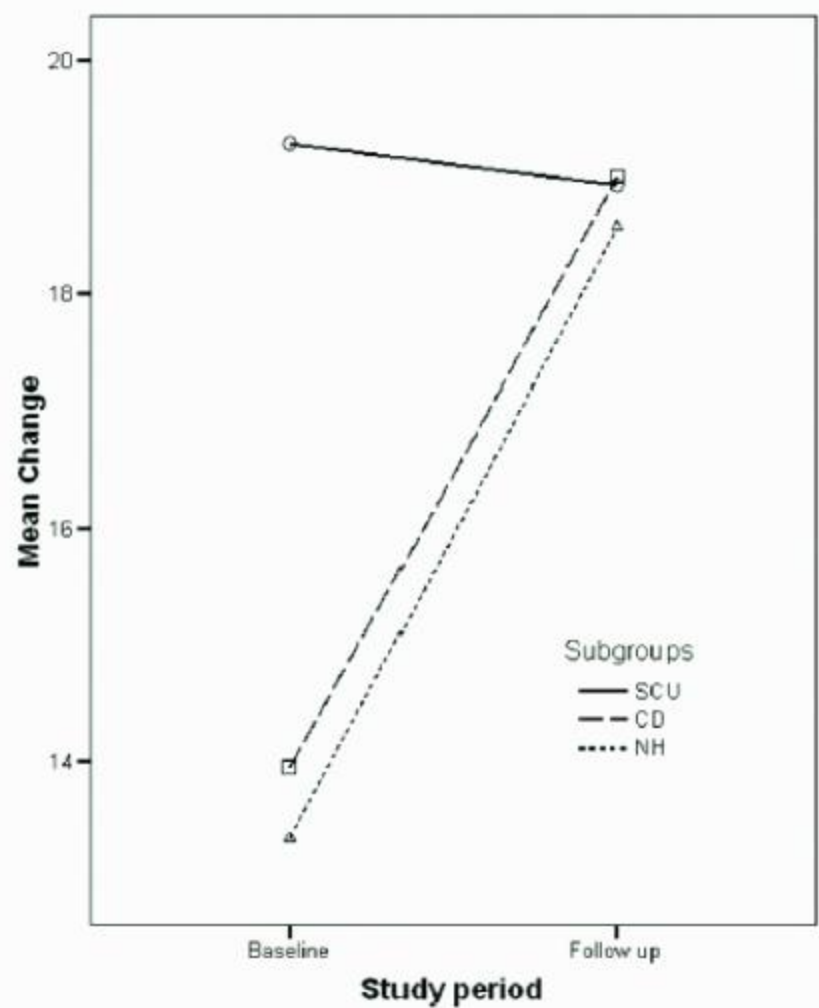


FIGURE 4
Mean DMR/SOS obtained by the three study groups across the 3-year period.

To take home...

- Adulti/anziani con DI sono ad elevato rischio per la demenza
- La valutazione e la diagnosi di demenza non è agevole ma è possibile
- La frequente multimorbidità impone interventi di prevenzione primaria e secondaria
- Sono urgenti studi sul *delirium* e sugli interventi psicosociali per queste persone «le più fragili tra le fragili»