

Specialiseret stroke service

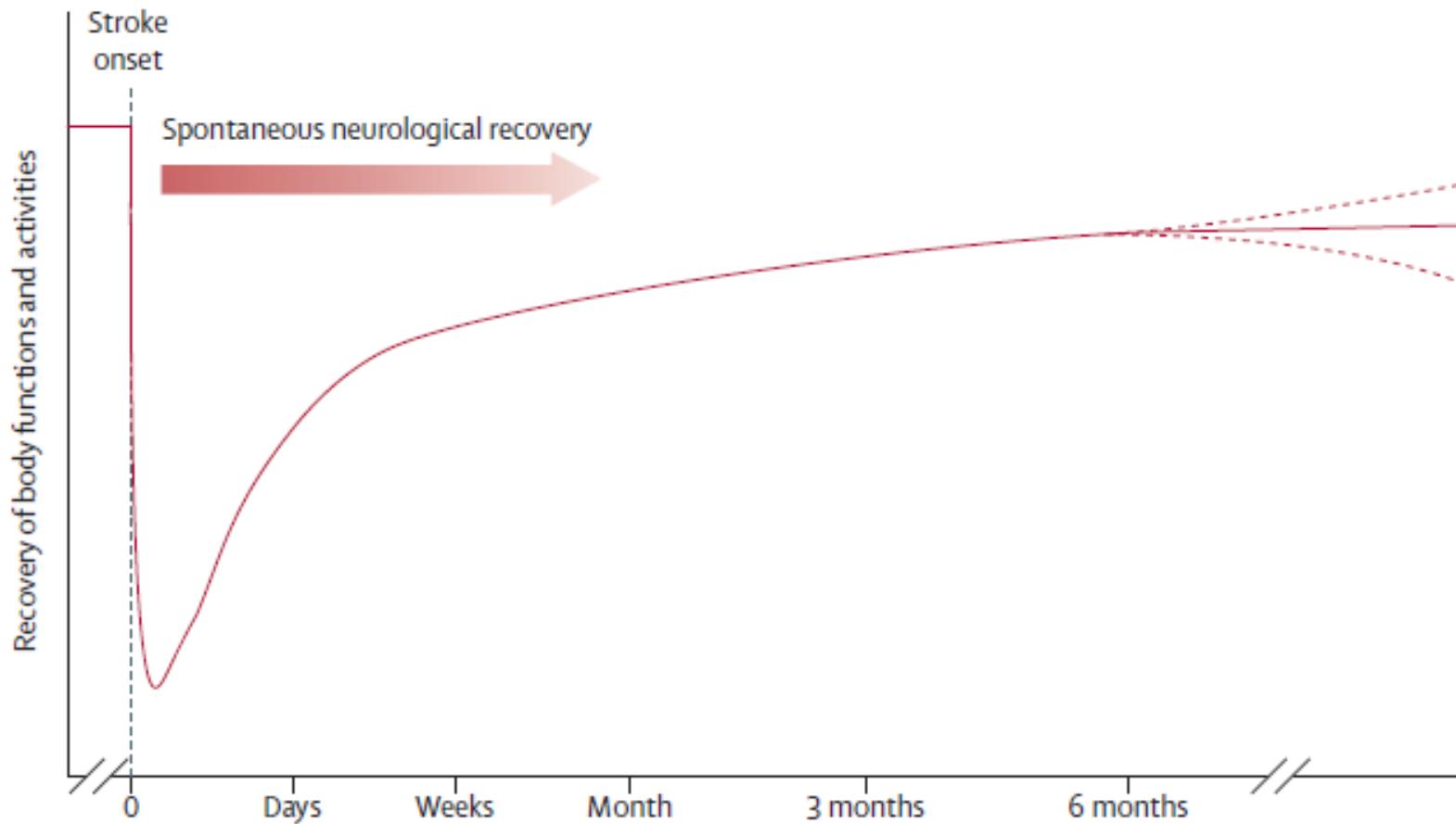
Hvad er der brug for, og hvordan sikrer vi kvaliteten af tilbud?

Har tidlig eller mere intensiv rehabilitering effekt og hvor længe er der brug for rehabilitering.

Hvad ved vi efter AVERT studiet?

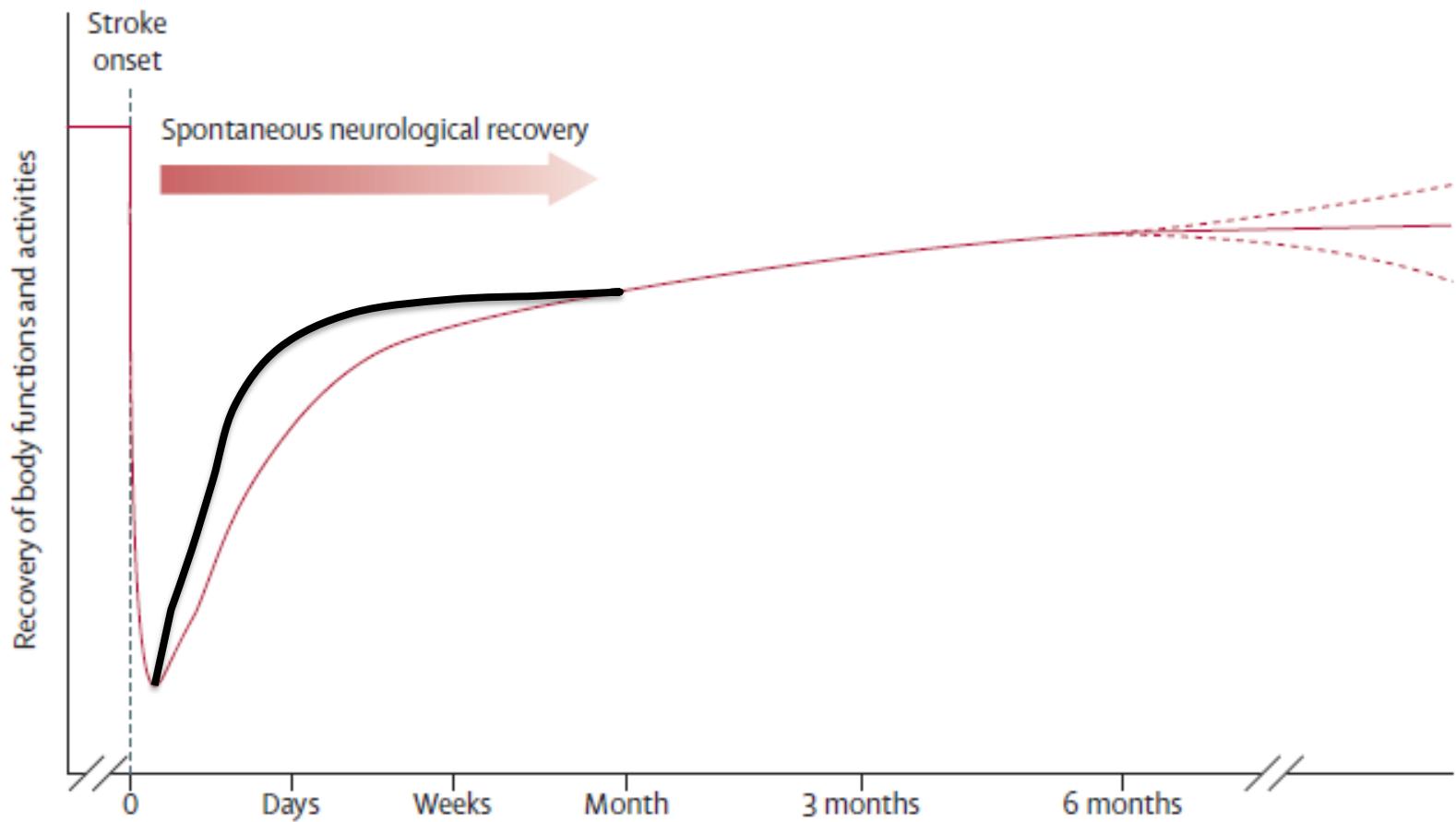
Jørgen Feldbæk Nielsen, professor, overlæge dr.med.





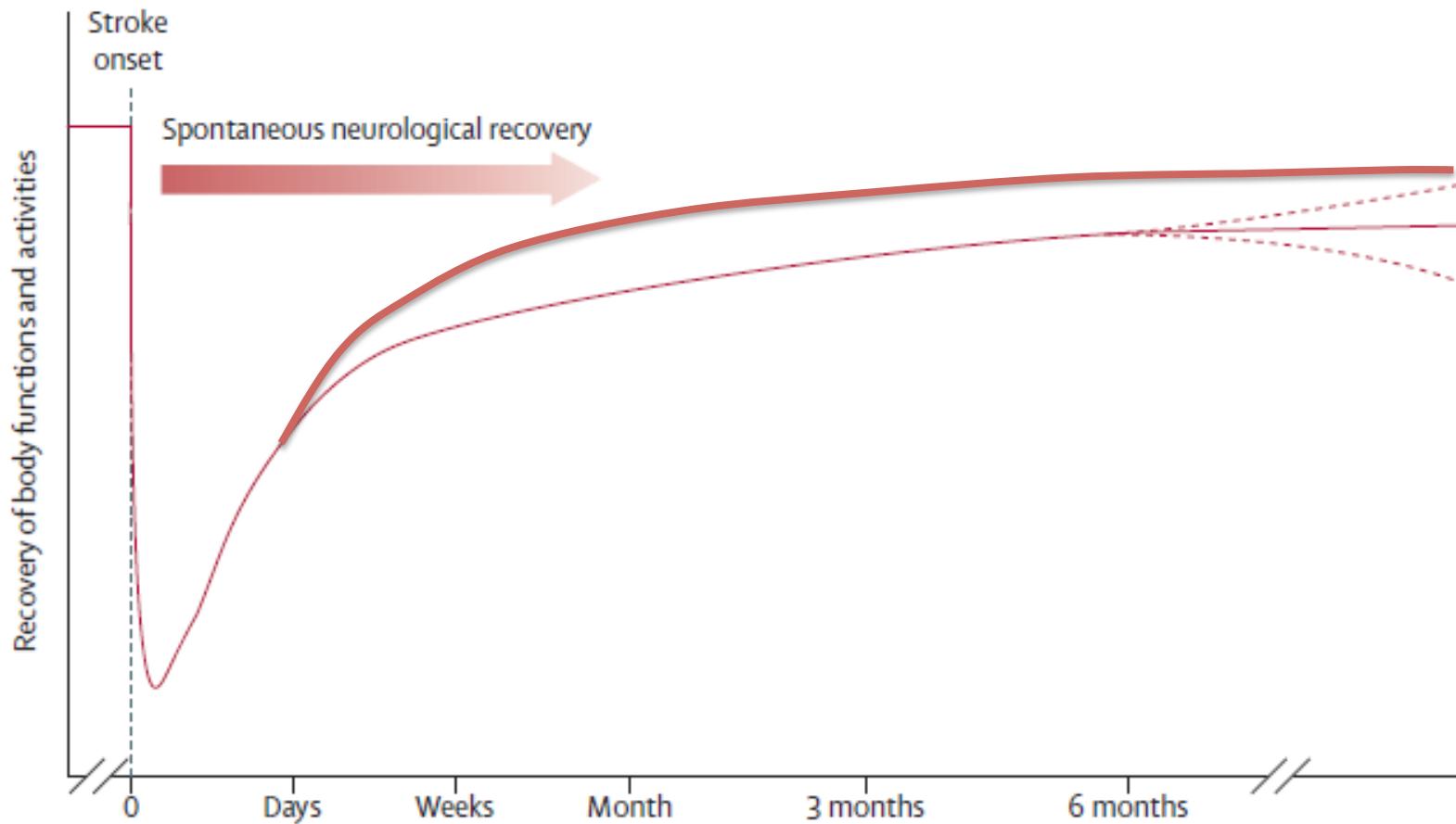
Lancet 2011; 377: 1693-702





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Efficacy and safety of very early mobilisation within 24 h of stroke onset (AVERT): a randomised controlled trial



The AVERT Trial Collaboration group*

Lancet 2015; 386: 46–55

Published Online
April 17, 2015

[http://dx.doi.org/10.1016/
S0140-6736\(15\)60690-0](http://dx.doi.org/10.1016/S0140-6736(15)60690-0)

[See Comment](#)

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Summary

Background Early mobilisation after stroke is thought to contribute to the effects of stroke-unit care; however, the intervention is poorly defined and not underpinned by strong evidence. We aimed to compare the effectiveness of frequent, higher dose, very early mobilisation with usual care after stroke.

Prespecified dose-response analysis for A Very Early Rehabilitation Trial (AVERT)

OPEN



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ABSTRACT

Objective: Our prespecified dose-response analyses of A Very Early Rehabilitation Trial (AVERT) aim to provide practical guidance for clinicians on the timing, frequency, and amount of mobilization following acute stroke.

Methods: Eligible patients were aged ≥ 18 years, had confirmed first (or recurrent) stroke, and were admitted to a stroke unit within 24 hours of stroke onset. Patients were randomized to receive very





Efficacy and safety of very early mobilisation within 24 h of stroke onset (AVERT): a randomised controlled trial



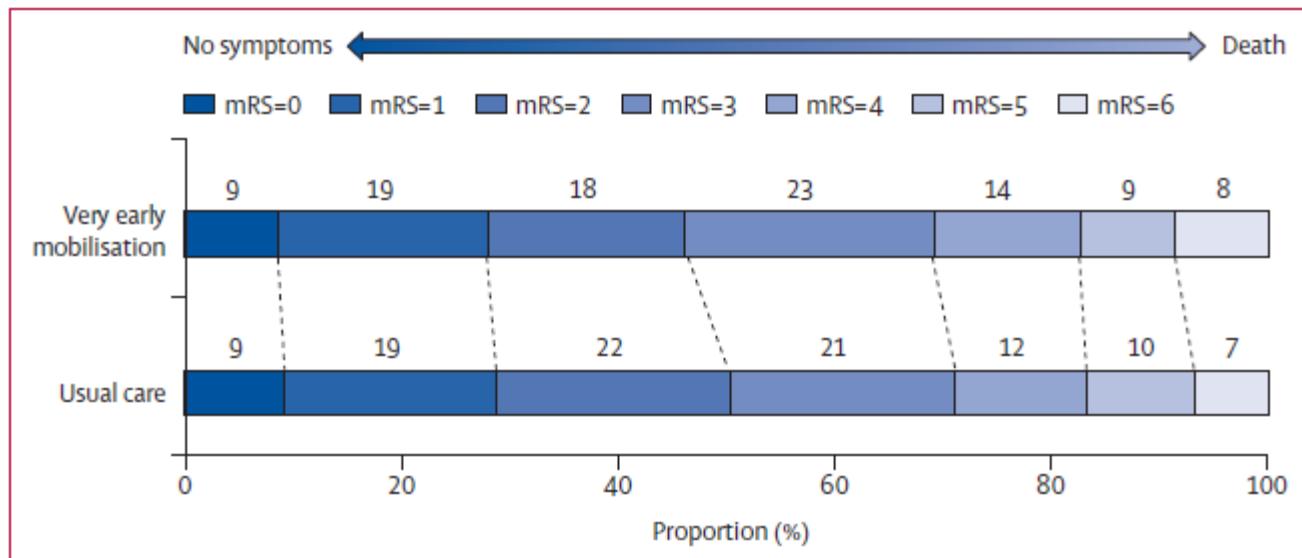
The AVERT Trial Collaboration group*

	Very early mobilisation (n=1054)	Usual care (n=1050)	Very early mobilisation (n=1054)	Usual care (n=1050)
(Continued from previous column)				
Recruitment region				
Australia and New Zealand	617 (59%)	626 (60%)	Lacunar infarct	255 (24%)
Asia	126 (12%)	125 (12%)	Intracerebral haemorrhage	142 (14%)
UK	311 (29%)	299 (28%)	rtPA treatment	268 (26%)
Age (years)	72·3 (62·3–80·3)	72·7 (63·4–80·4)	Yes	116 (11%)
<65	331 (31%)	298 (28%)	Baseline walking (Mobility Scale for Acute Stroke walking score)	247 (23%)
65–80	448 (43%)	481 (46%)	Independent	260 (25%)
>80	275 (26%)	271 (26%)	Supervised or assisted	439 (42%)
			Unable to walk	522 (49%)
			Unknown	96 (9%)
				0 (0%)

	Very early mobilisation (n=1054)	Usual care (n=1050)	p value	Median shift (95% CI)
Time to first mobilisation (h)	18·5 (12·8–22·3; n=1042*)	22·4 (16·5–29·3; n=1036*)	<0·0001	4·8 (4·1–5·7)
Frequency per person†	6·5 (4·0–9·5)	3 (2·0–4·5)	<0·0001	3 (3·3–5)
Daily amount per person (min)‡	31 (16·5–50·5)	10 (0–18)	<0·0001	21·0 (20–22·5)
Total amount per person (min)§	201·5 (108–340)	70 (32–130)	<0·0001	117 (107–128)



Very early mobilisation (n=1038*)	Usual care (n=1045*)	Adjusted analysis		Unadjusted analysis		
		OR, generalised OR, or HR† (95% CI)	p value	OR generalised OR, or HR† (95% CI)	p value	
Primary						
Favourable outcome‡	480 (46%)	525 (50%)	0.73 (0.59-0.90)	0.004	0.85 (0.72-1.0)	0.068



mRS = 0 Helt symptomfri

mRS = 1 Ikke betydende handikap på trods af lette symptomer; klarer alle sædvanlige opgaver og aktiviteter

mRS = 2 Lettere handikap; klarer ikke samtlige tidligere udførte aktiviteter, men er i stand til at klare sig selv uden hjælp til personlige gøremål fra andre personer (må godt have praktisk hjælp til f.eks. rengøring og indkøb)

mRS = 3 Moderat handikap; har brug for nogen hjælp til personlige gøremål (f.eks. påklædning eller hjælp til at klare en bankforretning), men er i stand til at gå uden hjælp fra andre personer (må godt bruge stok eller rollator)

mRS = 4 Moderat til svært handikap; ude af stand til at gå uden hjælp fra andre personer, og ude af stand til at klare personlige behov uden hjælp fra andre personer

mRS = 5 Svært handikap; sengeliggende, inkontinent, konstant pleje- og opmærksomhedskrævende
Død



Prespecified dose-response analysis for A Very Early Rehabilitation Trial (AVERT)

OPEN ▲

Neurology® 2016;86:1-8

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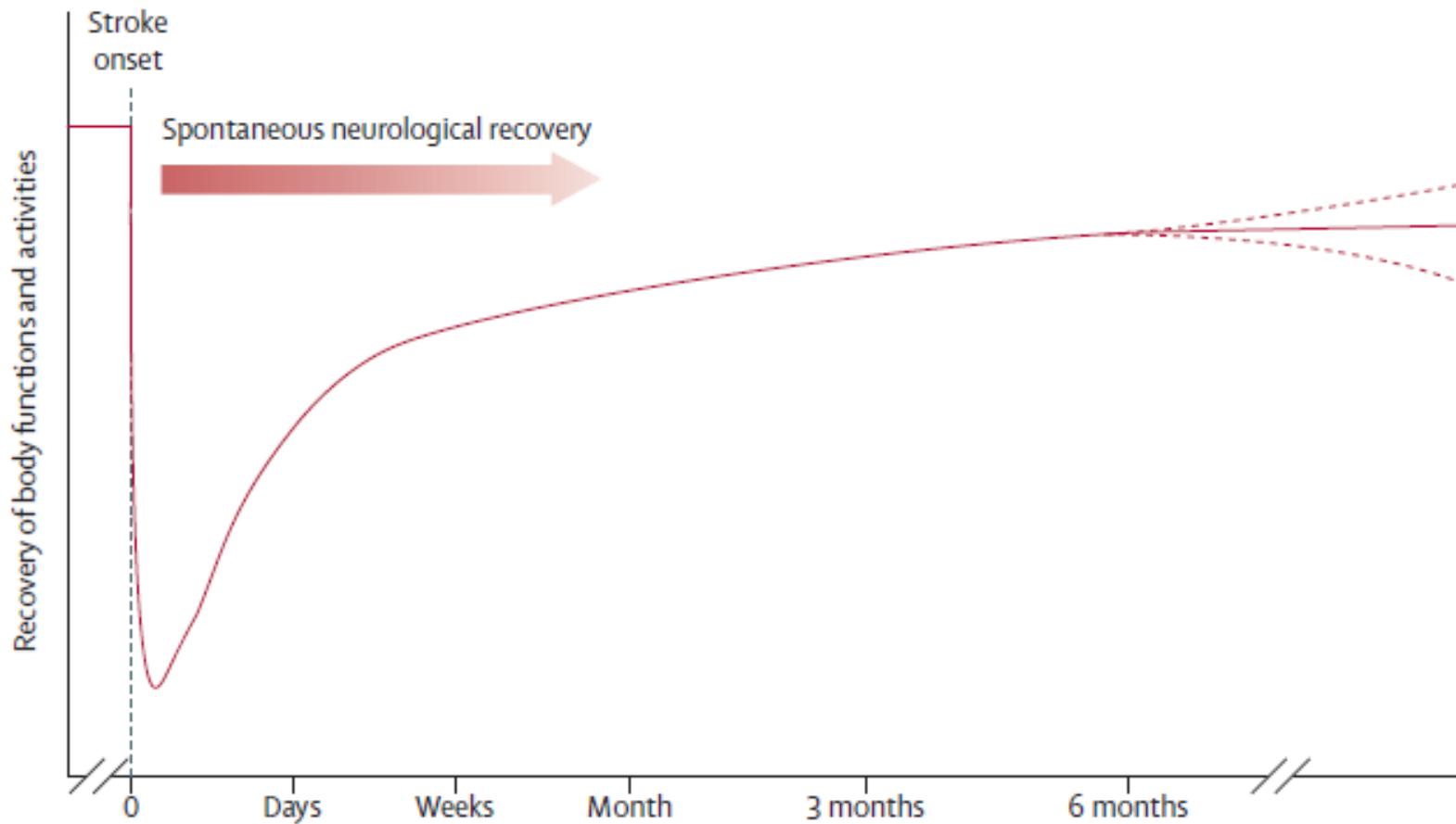
ABSTRACT

Objective: Our prespecified dose-response analyses of A Very Early Rehabilitation Trial (AVERT) aim to provide practical guidance for clinicians on the timing, frequency, and amount of mobilization following acute stroke.

Table 3 Effect of intervention characteristics on favorable outcome (mRS 0-2) and unassisted walking

Efficacy	Favorable outcome (mRS 0-2)		Walking unassisted 50 meters		
	OR (95% CI)	p Value	Binary OR (95% CI)	p Value	Cox hazard ratio (95% CI)
Model 1					
Time to first mobilization (per extra hour)	0.99 (0.98-1.0)	0.036	1.0 (0.99-1.0)	0.40	0.99 (0.98-0.99)
Frequency, median daily sessions ^a (per one extra session)	1.13 (1.09-1.18)	<0.001	1.66 (1.53-1.80)	<0.001	<0.001
Daily amount, median (per extra 5 minutes)	0.94 (0.91-0.97)	<0.001	0.85 (0.81-0.89)	<0.001	0.96 (0.94-0.97)





Lancet 2011; 377: 1693-702



Fysisk rehabilitering



Fysisk rehabilitering - principper

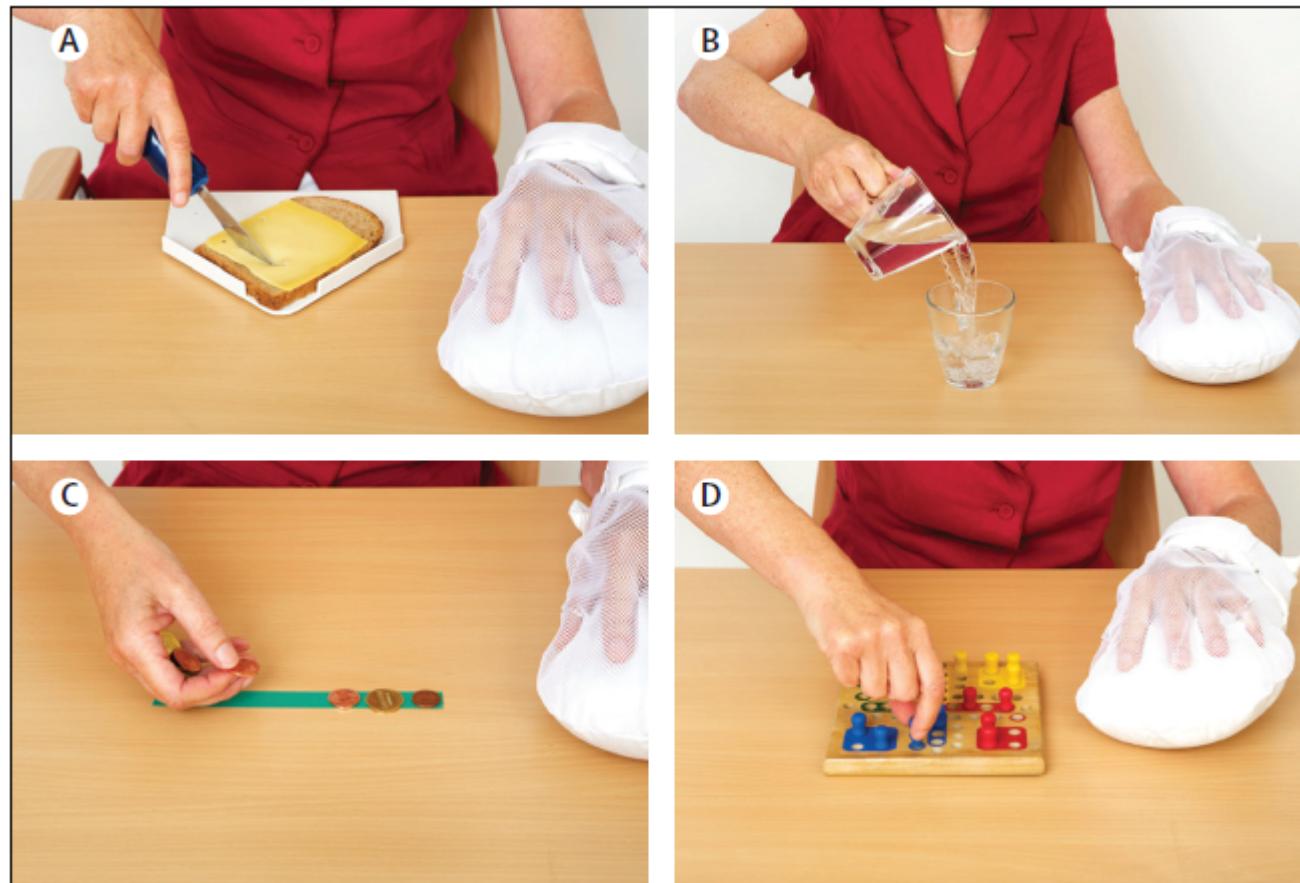
- ▶ Opgave specifik – man bliver god til det man træner
- ▶ Kontekst-afhængig
- ▶ Høj intensitet – mange repetitioner
- ▶ Motivation
- ▶ Begynd tidlig

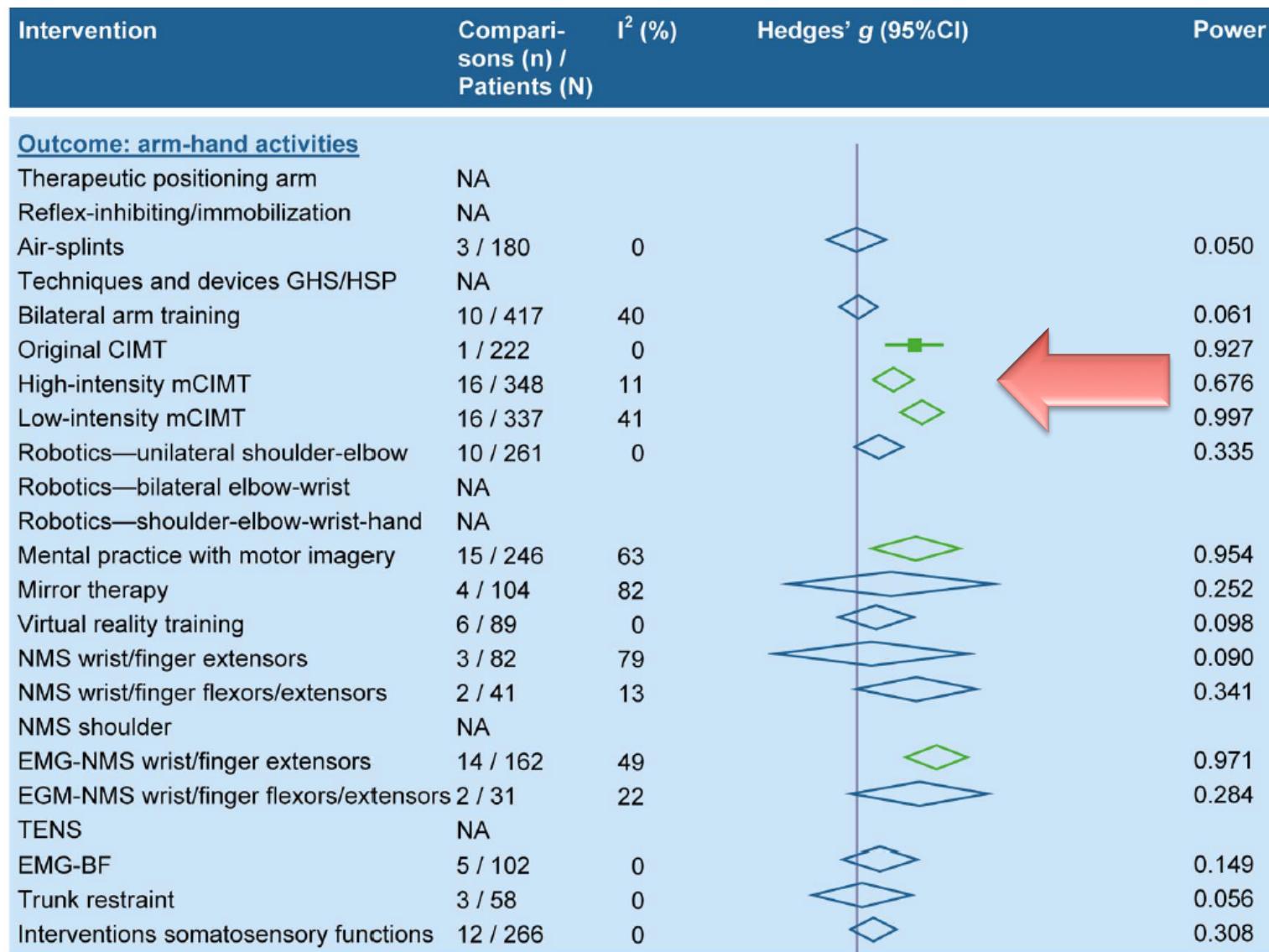




Constraint-induced movement therapy after stroke

Gert Kwakkel, Janne M Veerbeek, Erwin E H van Wegen, Steven L Wolf





Citation: Veerbeek JM, van Wegen E, van Peppen R, van der Wees PJ, Hendriks E, et al. (2014) What Is the Evidence for Physical Therapy Poststroke? A Systematic Review and Meta-Analysis. PLoS ONE 9(2): e87987. doi:10.1371/journal.pone.0087987



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Cochrane Database of Systematic Reviews

Physical rehabilitation approaches for the recovery of function and mobility following stroke (Review)

Pollock A, Baer G, Campbell P, Choo PL, Forster A, Morris J, Pomeroy VM, Langhorne P



AARHUS UNIVERSITY

Hammel Neurorehabilitation Centre and University Research Clinic

17.06.2016

Hvad begrænser os?

- ▶ træthed
 - ▶ depression
 - ▶ kognitive følger
 - ▶ co-morbiditet
 - ▶ mangel på viden
 - ▶ implementering er svær
 - ▶ manglende sammenhæng over sektorgrænser
 - ▶ dristighed til at gøre op med gamle vaner
- 
- patienten/borgeren
- sundhedsvæsenet

....men vi har et problem med de kognitive følger...



ORIGINAL ARTICLE

Post-stroke cognitive impairment is common even after successful clinical recovery

H. Jokinen, S. Melkas, R. Ylikoski, T. Pohjasvaara, M. Kaste, T. Erkinjuntti and M. Hietanen

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Keywords:
cognition, cognitive
impairment,
neuropsychology, stroke

Received 20 January 2015

Accepted 6 April 2015

*European Journal of
Neurology* 2015, **22**:
1288–1294

doi:10.1111/ene.12743

Background and purpose: Cognitive impairment is common after stroke, but the prevalence and long-term significance of the diverse neuropsychological deficits on functional outcome are still not well known. The frequency and prognostic value of domain-specific cognitive impairments were investigated in a large cohort of ischaemic stroke patients.

Methods: Consecutive patients ($n = 409$), aged 55–85 years, from the acute stroke unit of the Helsinki University Hospital, Finland, were evaluated with extensive clinical and neuropsychological assessments 3 months post-stroke. Impairments within nine cognitive domains were determined according to age-appropriate normative data from a random healthy population. Functional disability was evaluated with the modified Rankin scale (mRS) 3 and 15 months post-stroke.

Results: In all, 83% patients showed impairment in at least one cognitive domain, whereas 50% patients were impaired in multiple (≥ 3) domains. In

Table 2. Neuropsychological test battery of the Helsinki Stroke Aging Memory Study

Executive functions and attention
Trail making test, parts A and B
Stroop test, colour naming and interference
Wisconsin card sorting test
Verbal fluency, phonemic and semantic
Digit span, forward and backward
Memory
WMS-R logical memory, story A, immediate and delayed recall
WMS-R visual reproduction, immediate and delayed recall
Fuld object memory evaluation, 10-item learning test, 5 trials and delayed recall
Visuoconstructional and spatial functions
WAIS-R block design
Figure copying test
Clock drawing test
Abstract thinking
WAIS-R similarities
Aphasia
Token test
Boston naming test
Boston diagnostic aphasia examination, overall speech evaluation
Reading and writing
Samples
Arithmetic
Arithmetic operations
Neglect
Bells test
Agnosia
Poppelreuter



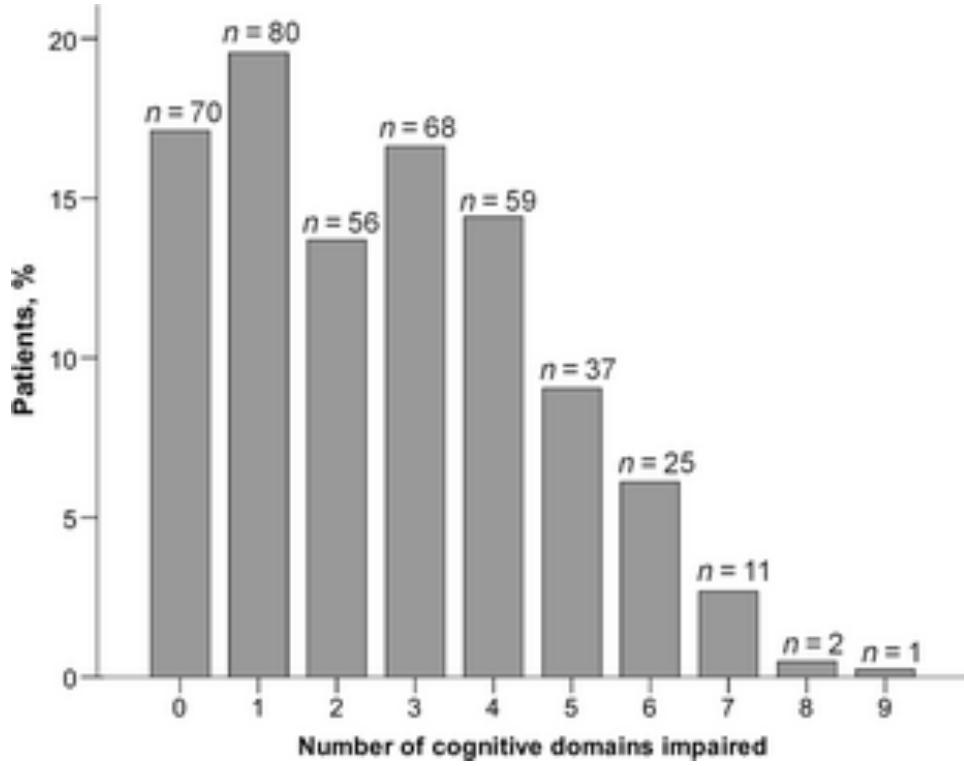


Table 3. Frequencies of impairment in each cognitive domain in the whole cohort and in a subgroup of patients with excellent clinical recovery (mRS = 0–1) 3 months after ischaemic stroke in the Helsinki Stroke Aging Memory Study

Cognitive domain	<i>n (%)</i>	
	All patients <i>n</i> = 409	Patients with mRS = 0–1 <i>n</i> = 152
Memory functions	227 (60)	77 (52)
Visuoconstructional and spatial functions	216 (55)	54 (36)
Executive functions and attention	181 (49)	52 (34)
Aphasia	114 (29)	23 (15)
Reading and writing	112 (30)	16 (11)
Abstract reasoning	106 (29)	26 (17)
Arithmetic	71 (20)	12 (8)
Neglect	29 (8)	2 (1)
Agnosia	13 (3)	1 (1)

Table 4. Cognitive impairments 3 months post-stroke as predictors of functional dependence (modified Rankin Scale >2) at 15-month follow-up in the Helsinki Stroke Aging Memory Study

Impairment	Logistic regression	
	OR (P value)	95% CI for OR
Memory functions	2.2 (0.008)	1.2–3.9
Visuoconstructional and spatial functions	5.1 (<0.001)	2.7–9.1
Executive functions and attention	3.2 (<0.001)	1.8–5.7
Aphasia	2.1 (0.017)	1.1–3.9
Reading and writing	2.3 (0.011)	1.2–4.3
Abstract reasoning	2.3 (0.006)	1.3–4.2
Arithmetic	1.9 (0.063)	1.0–3.9

Statistical analyses are adjusted for age, sex, years of education and stroke severity (National Institutes of Health Stroke Scale score). OR, odds ratio; CI, confidence interval.