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I VASCULAR COGNITIVE APPROACH

II TREAT IT

THE VASCULAR COGNITIVE IMPAIRMENT APPROACH

- Any cognitive impairment associated with or caused by a vascular factor
- 1. Adopt a common metric system
- 2. Test several provisional criteria simultaneously
- 3. Identify and treat the vascular component

Original Contributions

National Institute of Neurological Disorders and Stroke-Canadian Stroke Network Vascular Cognitive Impairment Harmonization Standards

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Background and Purpose—One in 3 individuals will experience a stroke, dementia or both. Moreover, twice as many individuals will have cognitive impairment short of dementia as either stroke or dementia. The commonly used stroke scales do not measure cognition, while dementia criteria focus on the late stages of cognitive impairment, and are heavily biased toward the diagnosis of Alzheimer disease. No commonly agreed standards exist for identifying and describing individuals with cognitive impairment, particularly in the early stages, and especially with cognitive impairment related to vascular factors, or vascular cognitive impairment.

THE VASCULAR COGNITIVE APPROACH

EVALUATION

- History and examination
- Neuropsychological evaluation
- Investigations
- 4. Identifying the vascular component
- 5. Differential diagnosis

ISCHEMIC SCORE WITH 5 COMPOSITE ITEMS*

Item No.	Item Description	Score if Answer is Yes
1/2	Abrupt onset or stepwise deterioration	Scored electronically
3 / 4	Fluctuating course or nocturnal confusion	Scored electronically
6/8	Depression or emotional incontinence	Scored electronically
9/11	History of hypertension or atherosclerosis	Scored electronically
10/12	History of stroke or focal neurological symptoms	Scored electronically
Total		Scored electronically

^{*} A vascular component of cognitive impairment may be indicated after electronic computation

Hachinski V. et al Arch Neurol 2012:69(2):169-175

STRIVE: **ST**andards for **R**eport**I**ng **V**ascular Changes on **NE**uroimaging, V1, The Lancet Neurology 2013;12:822-38

	Recent small subcortical infarct	White matter hyperintensity	Lacune	Perivascular space	Cerebral microbleeds
Example image					
Schematic	DWI	FLAIR	FLAIR	T2 T1/FLAIR	T2*/SWI
Usual diameter ¹	≤ 20 mm	variable	3-15 mm	≤ 2 mm	≤ 10 mm
Comment	best identified on DWI	located in white matter	usually have hyperintense rim	usually linear without hyperintense rim	detected on GRE seq., round or ovoid, blooming
DWI	↑	\leftrightarrow	↔/(↓)	\leftrightarrow	\leftrightarrow
FLAIR	↑	↑	\	\downarrow	\leftrightarrow
T2	↑	\uparrow	↑	↑	\leftrightarrow
T1	\	\leftrightarrow /(\downarrow)	\	\downarrow	\leftrightarrow
T2* / GRE	\leftrightarrow	↑	↔ (↓ if haemorrhage)	\leftrightarrow	$\downarrow \downarrow$

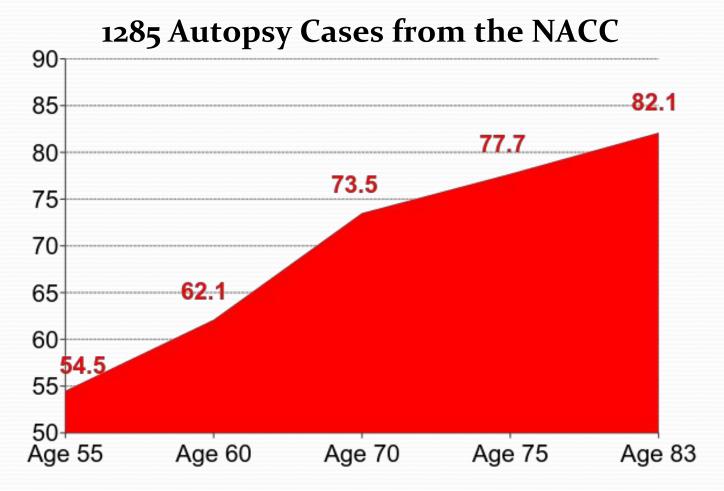
DIFFERENTIAL DIAGNOSIS OF COGNITIVE IMPAIRMENT

1. Drugs

2. Depression

3. Diseases

Prevalence of Vascular Lesions across Ages at Death in Pathology Specimens from Patients with Alzheimer's Disease as Primary Pathological Diagnosis



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THE SPRINT TRIAL

9361 Subjects with BP ≤ 130 mm systolic at increased cardiovascular risk

Randomized to targets of 140mm systolic or

120mm systolic

Trial stopped at 3.26 years

The SPRINT Research Group. NEJM. Nov 9 2015

THE SPRINT TRIAL

120mm systolic target

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\downarrow Mortality HR 0.73 95% CI 0.64 – 0.89
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 \downarrow MI, stroke MR 0.75 95% CI 0.64 – 0.89

heart failure

Serious side effects (hypotension, syncope, electrolyte abnormalities, acute kidney injury or failure)

120mmHg group. 4.7% vs 2.5% 140 mmHg group

HR 1.88 p < 0.001

Orthostatic hypotension significantly less common in 120mmHg group

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3 STEPS IN PREVENTION

RISK MOTIVATION

Personality

Decision

stage Harnessing technologies

and social media

NABLEMENT

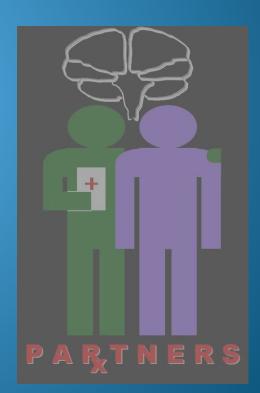
School/Work

Communities

Environment

PARTNERS

A Canadian Multi-Center, Randomized, Controlled, Open-Labeled, Blinded Adjudication Clinical Trial







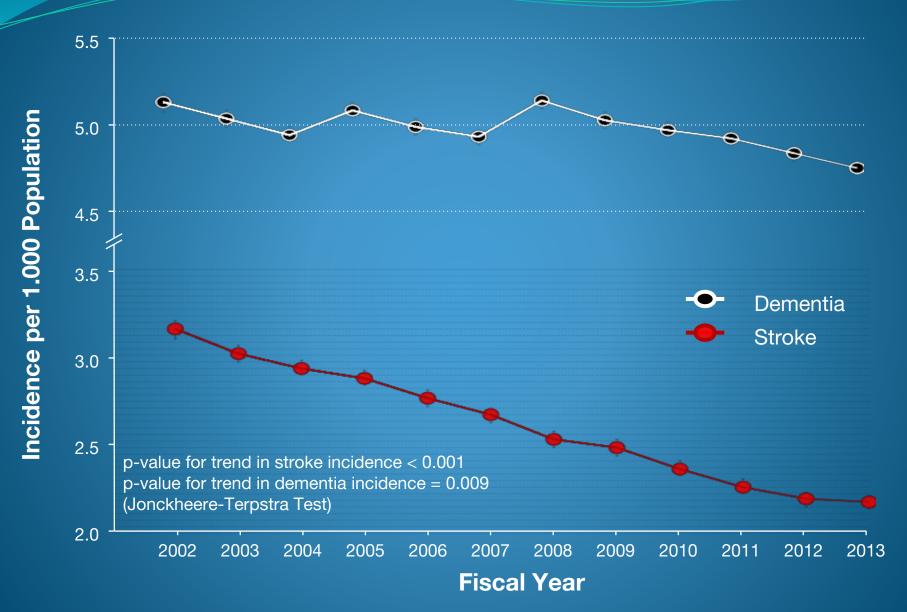






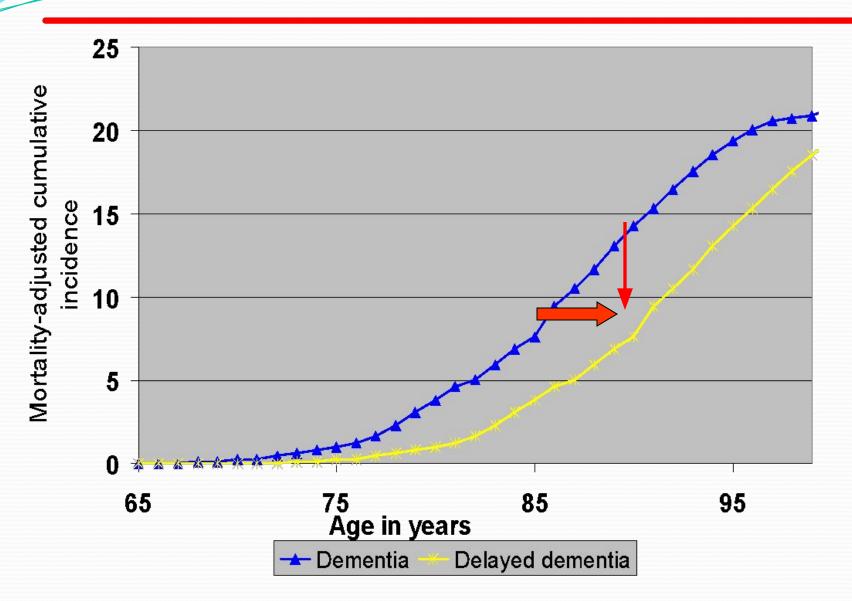
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Sposato LA.....Hachinski V. JAMA Neurol. 2015;72:1529-1531

Lifetime Risk of Dementia: Women - 65 yrs



Seshadri et al. Lifetime Risk of Stroke: The Framingham Study. Stroke 2006; 37: 344-349.