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Alzheimer's کئ Dementia

Alzheimer's & Dementia (2017) 1-13 Featured Article Progress toward standardized diagnosis of vascular cognitive impairment: Guidelines from the Vascular Impairment of Cognition Classification Consensus Study Olivia A. Skrobot^a, Sandra E. Black^b, Christopher Chen^c, Charles DeCarli^d, Timo Erkinjuntti^e, Gary A. Ford^f, Rajesh N. Kalaria^g, John O'Brien^h, Leonardo Pantoniⁱ, Florence Pasquier^j, Gustavo C. Roman^k, Anders Wallin¹, Perminder Sachdev^{m,n}, Ingmar Skoog^o, VICCCS group¹, Yoav Ben-Shlomo^p, Anthony P. Passmore^q, Seth Love^a, Patrick G. Kehoe^{a,*} ^aDementia Research Group, School of Clinical Sciences, Faculty of Health Sciences, University of Bristol, Level 1, Learning & Research, Southmead Hospital, Bristol, UK ^bSunnybrook Research Institute, University of Toronto, Canada ^cMemory Aging & Cognition Centre, Department of Pharmacology, National University of Singapore, Singapore ^dAlzheimer's Disease Center and Imaging of Dementia and Aging (IDeA) Laboratory, Department of Neurology and Center for Neuroscience, University of California at Davis, 4860 Y Street, Suite 3700, Sacramento, CA 95817, USA ^eClinical Neurosciences, Neurology, University of Helsinki and Helsinki University Hospital, Finland, POB 300, FIN-00290, HUS, Finland ^fDivison of Medical Sciences, Oxford University, Magdalen Centre North, Oxford Science Park, OX4 4GA, UK ⁸Institute of Neuroscience, Campus for Ageing & Vitality Newcastle upon Tyne, NIHR Biomedical Research Building, NE4 5PL, UK ^hDepartment of Psychiatry, University of Cambridge School of Clinical Medicine, Cambridge Biomedical Campus Box 189, Level E4, Cambridge, CB2 0SP UK ⁱNEUROFARBA Department, University of Florence, Florence, Italy ^jUniversity of Lille, Inserm U1171, Degenerative and vascular disorders, CHU, Distalz, F-59000, Lille, France ^kMethodist Neurological Institute, 6560 Fannin Street, Suite 802, Houston, TX 77030, USA ¹Institute of Neuroscience and Physiology at Sahlgrenska Academy, University of Gothenburg, Memory Clinic at Department of Neuropsychiatry, Sahlgrenska University Hospital, Wallinsgatan 6, SE-431 41 Mölndal, Sweden ^mSchool of Psychiatry, University of New South Wales, Sydney, Australia ⁿCHeBA (Centre for Healthy Brain Ageing), Neuropsychiatric Institute, Prince of Wales Hospital, Randwick NSW 2031, Australia ^oCenter for Health and Ageing (AGECAP), Institute of Neuroscience and Physiology, Sahlgrenska Academy, University of Gothenburg, Sweden ^PSchool of Social and Community Medicine, University of Bristol, Canynge Hall, 39 Whatley Road, Bristol, BS8 2PS, UK 34^{Q2} ⁴Institute of Clinical Sciences, Block B, Queens University Belfast, Royal Victoria Hospital, Belfast, BT12 6BA Abstract Introduction: Progress in understanding and management of vascular cognitive impairment (VCI) has been hampered by lack of consensus on diagnosis, reflecting the use of multiple different assessment protocols. A large multinational group of clinicians and researchers partici-pated in a two-phase Vascular Impairment of Cognition Classification Consensus Study (VICCCS) to agree on principles (VICCCS-1) and protocols (VICCCS-2) for diagnosis of VCI. We present VICCCS-2. Methods: We used VICCCS-1 principles and published diagnostic guidelines as points of reference for an online Delphi survey aimed at achieving consensus on clinical diagnosis of VCI. Results: Six survey rounds comprising 65-79 participants agreed guidelines for diagnosis of VICCCS-revised mild and major forms of VCI and endorsed the National Institute of Neurological Disorders-Canadian Stroke Network neuropsychological assessment protocols and recommenda-tions for imaging. ¹Members are listed at the end of the article. E-mail address: patrick.kehoe@bristol.ac.uk 54^{Q3} *Corresponding author. Tel.: 0117 414 7821.

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Discussion: The VICCCS-2 suggests standardized use of the National Institute of Neurological Disorders-Canadian Stroke Network recommendations on neuropsychological and imaging assessment for diagnosis of VCI so as to promote research collaboration. © 2017 the Alzheimer's Association. Published by Elsevier Inc. All rights reserved.

Vascular cognitive impairment; Vascular dementia; Guidelines; Criteria; consensus; Delphi

1. Introduction

Keywords:

Since Hachinski et al [1] proposed the term multi-infarct dementia to describe dementia complicating ischemic vascular disease, numerous other descriptors have been used to encompass the heterogeneous clinical and etiological spectrum of cognitive impairment due to cerebrovascular disease (CVD). These include vascular dementia (VaD), vascular cognitive impairment (VCI), subcortical (ischemic) vascular dementia, and vascular cognitive disorders (VCDs), variably diagnosed according to multiple different guidelines or protocols [2-14], some agreed by national institutions or research networks, for example, Alzheimer's Disease Diagnostic and Treatment Centers [11], International Statistical Classification of Diseases, 10th revision 135 [15], the National Institute of Neurological Disorders and 136 Stroke (NINDS)-Association Internationale pour la Re-137 cherche et l'Enseignement en Neurosciences [16], and Diag-138 nostic and Statistical Manual of Mental Disorders, fourth 139 and fifth editions (DSM-4 and DSM-5; [17,18]). 140

141 Studies comparing some of these protocols have shown 142 they are not readily interchangeable [19-21]. After the 143 commencement of the Vascular Impairment of Cognition 144 Classification Consensus Study phase 1 (VICCCS-1), the 145 American Heart Association/American Stroke Association 146 published a statement on vascular (AHA/ASA) 147 contributions to cognitive impairment and dementia [22]. 148 This supported the use of assessment protocols previously 149 published by NINDS-Canadian Stroke Network (CSN) 150 [13]. There have been other recent contributions to this field 151 from the International Society of Vascular Behavioral and 152 Cognitive Disorders (VASCOG) [23] and the DSM-5 [18]. 153 154 The level of take up of these recent guidelines is still unclear. 155 Only those published during VICCCS-1, before commence-156 ment of VICCCS-2, could be included for consideration in 157 the present study [22,24]. 158

The aim of VICCCS was to achieve broad international 159 consensus on diagnosis of VCI, through participation of a 160 large pool of international researchers and clinicians in an 161 iterative survey using the Delphi approach. After two initial 162 survey rounds, the study was separated into two phases: 163 VICCCS-1, addressing key concepts in our understanding 164 165 and terminology of cognitive impairment resulting from 166 CVD [25], and VICCCS-2, focusing on the formulation of 167 practical guidelines for diagnosis.

168 The VICCCS-1 achieved broad consensus on concepts of 169 VCI. It supported the use of "mild" and "major" subdivisions 170

of the severity of impairment, aligning with the revised terminology in the DSM-5. VICCCS-1 participants concluded that attempts to separate mild VCI into further subtypes according to affected cognitive domains were at present premature but agreed that this should be an area of future research. VICCCS-1 agreed (Fig. 1, reproduced from [25]) that the major forms of VCI (VaD) should be classified into four main subtypes: (i) post-stroke dementia (PSD); (ii) subcortical ischemic vascular dementia (SIVaD); (iii) multi-infarct (cortical) dementia (MID); and (iv) mixed dementias (further subdivided according to additional neurodegenerative pathologies). Framed by these concepts, VICCCS-2 used the same Delphi methodology to agree diagnostic guidelines on determination of severity of VCI and discrimination of subtypes.

2. Methods

Participants in VICCCS-1 [25] were invited to participate in VICCCS-2 (Supplementary Figure 1). Although 149 initially agreed to participate, only approximately half were active and committed respondents in three or more rounds, with low attrition and little variation in participation throughout the six rounds (65-79 participants in each round, a mean of 72). Of the active participants, 63%–75% of participants (mean 68%) were clinicians with direct involvement in clinical assessment or health service patient care. The remainder were nonclinical (i.e., supporting clinical work technically or otherwise, but not involved in clinical decision-making, or predominantly involved in research). Individual round representation is provided in Supplementary Table 1.

2.1. Data collection

We used the Delphi method, an iterative, multistaged series of structured questionnaires with feedback of anonymized responses and progressive refinement of questions to reach consensus [26]. The process was co-ordinated by a nonparticipating researcher (O.A.S). Anonymization of responses facilitated free expression of opinion throughout the study. Feedback of summary responses after each round informed subsequent questions and allowed unbiased evolution of group judgment. A threshold of two-thirds agreement was chosen to signify consensus [27] for issues refined iteratively through multiple rounds, as in VICCCS-1 [25]. For issues where this threshold was not reached, we present 171 172

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