

In Memoriam

From Bench to Bedside in Neuropsychology

Professor Glyn W. Humphreys' Legacy

NELE DEMEYERE

It has now been slightly more than a year since we suddenly and tragically lost one of the great neuropsychologists of our time. Professor Glyn Humphreys passed away unexpectedly on January 14, 2016, while in Hong Kong as a distinguished visiting professor. A year has passed, but those who worked close to him and whose lives were touched by his understated leadership, sharp mind, clear insights, and humanity will always miss his gentle guidance. He leaves a tremendous legacy, in many research domains within psychology and neuroscience, including neuropsychology, cognitive psychology, computational modeling, social neuroscience, neurorehabilitation, and cognitive screening and diagnostics. His enormous influence and standing in the field were recognized internationally by many awards and accolades, such as The Spearman Medal (1986), the British Psychological Society's Cognitive Psychology section prize (1999 and 2012) and President's Award (1999), a Humboldt Research Fellowship (1998), the Royal Society Wolfson merit award (2007), the Donald Broadbrent Prize from the European Society for Cognitive Psychology (2013), and the Leibniz Professorship and Special Professorship of the Chinese Academy of Sciences, as well as the British Psychological Society Life Time Achievement Award (2015). He was elected a member of the Royal Society for Medicine (2008) and a fellow of the British Academy (2009).

These esteemed awards complement the daily leadership tasks he took on, as editor of the *Quarterly Journal of Experimental Psychology, Visual Cognition* (as founding Editor), and the *Journal of Experimental Psychology: Human Perception and Performance*. He was head of the School of Psychology at the University of Birmingham for 15 years, and head of the Department of Experimental Psychology at the University of Oxford from 2011, where he established the Cognitive Neuropsychology Centre.

Listing these markers of esteem, however, cannot do justice to the kind of person Glyn Humphreys was. He was much more than a star academic; he really cared deeply about his colleagues, his students, and the many patients who contributed to the research. Throughout his career he and Professor Jane Riddoch set about improving neuropsychological assessments and neurorehabilitation for patients after brain injury. It is this legacy in neuropsychological applications and the journey from laboratory-based experiments to applied and widely used cognitive screening and assessments that this tribute will focus on.

Professor Humphreys' early investigations in visual cognition and visual agnosia directly led to the development of the Birmingham Object Recognition Battery [BORB]¹). It started with a theoretical interest in Marr's posthumously published work on the derivation of canonical axes from an image.² In particular,

the importance of the axes of elongation for object recognition, also highlighted by Biederman's work in his "recognition by components" theory³ struck a chord. When reviewing some of the classic work on impairments to visual object recognition following right hemisphere lesions, particularly on an unusual views object naming task,⁴ he was driven to investigate the components underlying these unusual views. He did so by differentiating between views that foreshorten the main axis and views that maintained the main elongation axis, but minimized the saliency of identifying features.

In their seminal paper, Humphreys and Riddoch⁵ found a double dissociation, with four patients with acquired brain damage, demonstrating a specific impairment in matching objects from foreshortened views to a target canonical view, with no problems identifying the minimal feature view, and a single patient who showed impaired matching only when the saliency of the target's distinctive features was reduced. This one patient with visual agnosia was of particular interest and became the subject of many articles to follow with the famous initials H.J.A., most notably described in the book To See But Not To See: A Case Study of Visual Agnosia.6 H.J.A. demonstrated intact low level vision and binding features into edges, intact shape discriminations, and intact copying, but severely impaired figureground segmentation. This provided evidence for a further fractionation of Lissauer's⁷ two stage framework for understanding disorders of object processing: apperception and association, because, within apperceptive agnosia, H.J.A.'s impairment in object recognition was specific to an inability to integrate form information. Unlike patients with form agnosia, such as the patients reported by Efron⁸ and Benson and Greenberg, H.J.A. was able to copy and to discriminate between shapes. Humphreys and Riddoch coined the concept and term "integrative agnosia." The tests used in these initial patient studies were structured under four headers: Low Level Vision, Mid Level Vision, Semantics, and Object Recognition, and were published in the first standardized neuropsychological clinical test from the Humphreys' laboratory: the BORB. 11

Over the years, broader interests came with the arrival of many different patients. Professor Humphreys and his group published countless articles on a variety of neuropsychological syndromes and symptoms, furthering our theoretical understanding of cognition, contributing to knowledge about underlying neuroanatomical and functional processes, and demonstrating effective methods of intervention. For the purpose of this tribute, it is impossible to be exhaustive, but some examples include studies on neglect, 12,13,14 extinction, 15,16,17 aphasia, 18,19 alexia, 20,21 letter by letter reading,²² prosopagnosia,²³ numerical cognition, 24,25 apraxia, 26,27,28 short-term memory, ^{29,30} simultanagnosia, 31,32,33 and even deficits in theory of mind. 34,35,36

With this varied experience and expertise also came an awareness of a gap in cognitive neuropsychological assessments: the need for an initial cognitive screen for patients with acquired brain injuries (e.g., stroke), that would be both "broad and shallow." Many neuropsychological domain-specific batteries existed, but to use all of them for every new patient would take hours to complete. To complete only one specific domain test would ignore the many often co-occurring cognitive difficulties in attention, memory, and numerical cognition. There was a need for a neuropsychological screening approach that would cover a range of cognitive domains, and that could be delivered efficiently. In addition, it was to be designed to maximize inclusion by being aphasia and neglect friendly. This led to the development of the Birmingham Cognitive Screen (BCoS).³⁷ The test philosophy was made explicit: to use short high-frequency words throughout, use vertical layouts and multimodal presentations, and to maximize time efficiency by designing tests that would incorporate several measures. The tests were made to be sensitive (so that they would detect a problem if one were present) and indicative of domain deficits, although not detailed. For example, the language tests would pick up a problem in object naming, but would not investigate the nature of the exact naming problem. For the first time, neuropsychological expertise in the form of a user-friendly first line cognitive screen was within the reach of interested allied health professionals, who were not necessarily neuropsychologists. BCoS can be delivered within 1 hour, at subacute stages of stroke, and gives a relatively detailed breakdown of cognitive function in five domains: Attention and Executive Function, Language, Memory, Numerical Ability, and Praxis. A recent article summarized the ability of BCoS to identify differential cognitive profiles, informing rehabilitation and contributing to the prediction of longer-term outcomes.³⁸

Although BCoS filled a gap in cognitive profiling, where there was time to do so (where patients and staff could support an hour of assessment), the timepressured environments of acute stroke units required a shorter screen. At the time, no stroke-specific cognitive screen existed, and short dementia screens, such as the Mini Mental State Examination³⁹ or Montreal Cognitive Assessment⁴⁰ were routinely used. However, these were developed to screen for dementia, and not necessarily suited to screen for common post- stroke cognitive impairments: the cognitive profile of a stroke survivor with a focal infarct or bleed is

very different from a neurodegenerative global impairment profile. In particular, the language demands for these screens are high, and the presence of neglect may contaminate task performance.⁴¹ The Oxford Cognitive Screen (OCS)⁴² provided a tailored stroke-specific solution, one that was not relying on spoken responses, and not impacted by neglect. The OCS focussed on briefly screening for impairments in the same five cognitive domains as set out in the BCoS, delivering a cognitive profile to inform the multidisciplinary team about specific cognitive impairments, rather than an overall pass/fail judgement. This shorter screen, which can be completed in under 20 minutes, has gained great traction in clinical practice, and has at the point of writing already been adopted in over 250 National Health Service (NHS) units in the United Kingdom and several translations have been completed^{43,44} or are underway (www.ocs-test.org).

The latest developments in Professor Humphreys' efforts in improving neuropsychological screening were through the development of further tablet-based assessments of the domain-specific OCS as well as OCS-Plus: aimed to briefly and sensitively assess broader domaingeneral functions, using the mobile technology to automatically derive not only outcome, but also measurements of process and strategies. This work is being continued by our group at the Cognitive Neuropsychology Centre, and a first study using this latest approach to demonstrate effects of ageing and cognitive reserve in a large low-literacy cohort in South Africa was published in his name posthumously. 45

Professor Glyn Humphreys' legacy is immense, and although his specific impact on improving real-life clinical practice is but one strand of his lasting influence, this tribute specifically set out to celebrate this aspect of his career, which served as a bridge joining bench

and bedside. His portfolio of assessments and screening, built on neuropsychological theory expertise and translated to clinical practice improvements will, no doubt, continue to inspire future generations of researchers to walk in the footsteps of this giant in neuropsychology.

Notes

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In Memoriam

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