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LETTER TO THE EDITOR

Good recovery after stroke may hide widespread cognitive deficits

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Sir,

I was very interested to read the recently published paper by Planton et al. [1] not least because we are also studying neuropsychological functioning in those with a ‘good recovery’ after stroke: in our CompAS (Complaints After Stroke) study. Planton et al.’s paper rightly deserved its own editorial by van Dijk and de Leeuw (same issue). It is wonderful to see neuropsychological outcome after stroke finally finding the place it deserves in the neuropsychological literature, so much so that it even has a new name: ‘post-stroke cognitive impairment’ [2]. As the authors rightly point out, many stroke patients are sent home deemed to have ‘good recovery’ only to discover that their cognitive functioning is not working as well as it might and that these problems have a massive impact on their daily lives.

They found that survivors tested at 3 months post-stroke performed more poorly than controls on every cognitive domain tested. The main goal of their study was thus fulfilled – to assess neuropsychological outcome (in the short term) in good recoverers with a first-ever stroke. Their second goal – to identify both the profile of possible neuropsychological impairment and its relationship with brain lesions – was harder for the authors to fulfil. They had unequal subgroups that were also relatively small; no significant links were found between lesion location and cognitive impairment. This may simply be an issue of low statistical power or not choosing the correct statistical analyses (the focus is on the group rather than on the individual level). Unfortunately, the authors also provide no data on the vascular risk factors in the control group nor did they measure MRIs in the controls making the (significant) link between increased white matter lesions and more cognitive deficits premature: they cannot rule out the possibility of such lesions in the control group caused for example by silent strokes.

The authors also found that whilst patients were more apathetic than controls, they were not more depressed. Depression is common after stroke, but perhaps the patients were not home long enough to be fully aware of their cognitive disabilities or, as the authors themselves suggested, depression after stroke is more common in those who are more disabled than their patient population. This too was reflected in the fact that the levels of subjective cognitive complaints in both patients and controls were low (as assessed by the CCQ – a French questionnaire, maximum score 10 and perhaps not sensitive enough to pick up subtle cognitive complaints?) Cognitive complaints are what we are focusing on in our CompAS study, and as such, we see these as very different to actual cognitive functioning; they are after all what patients themselves think about their cognition and do not always correlate with actual cognitive test performance [in stroke: see 3].

There are a number of issues I would like to ask the authors about: (i) Why did they use spouses or relatives of the patients as controls? (close relatives, especially spouses, do not make for good controls) (ii) Why did the authors not use the Oxfordshire Community Stroke Project classification (OCSP) [4,5] to assess stroke type/severity? And, (iii) why do they refer to cognitive complaint, quality of life and depression after stroke as not indicative of cognitive impairment? They also underlined the importance of taking confounding factors (e.g. previous cognitive decline, clinical variables) into account in future prospective studies of neuropsychological functioning after stroke, something that many studies have regretfully not considered. Highlighting the need to follow-up patients who have been deemed ‘recovered’ is clearly necessary if we are to develop better ways to treat them. What matters to stroke patients and how they function needs to be further explored not just in the short term but also in the longer term, something that we hope to address in the CompAS study.

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