



## iCAHE JC Critical Appraisal Summary

### Journal Club Details

---

Journal Club location	Flinders Medical Centre
JC Facilitator	Pamela Hewavasam
JC Discipline	Speech Pathology
CAT completed by:	Matt Ransom

### Question

How does the Oxford Cognitive screen compare to the Mini-mental state examination in detecting cognitive symptoms after stroke?

### Review Question/PICO/PACO

- P Patients with cognitive impairments post stroke
- I Oxford Cognitive Screen
- C Mini-mental state examination
- O Ability to detect cognitive symptoms post stroke

### Article/Paper

Mancuso, M., Demeyere, N., Abbruzzese, L., Damora, A., Varalta, V., Pirrotta, F., Antonucci, G., Matano, A., Caputo, M., Caruso, M.G. and Pontiggia, G.T., 2018. Using the Oxford cognitive screen to Detect cognitive impairment in stroke Patients: a comparison with the Mini-Mental state examination. *Frontiers in neurology*, 9, p.101.

*Please note: due to copyright regulations CAHE is unable to supply a copy of the critically appraised paper/article. If you are an employee of the South Australian government you can obtain a copy of articles from the [DOHSA librarian](#).*

**Article Methodology:** Diagnostic

Click [here](#) to access critical appraisal tool

Ques No.	Yes	Can't Tell	No	Comments
1	✓			<p><b>Was there a clear question for the study to address?</b></p> <p>Study aimed to compare the Oxford Cognitive Screen (OCS) with the Mini-mental state examination (MMSE) in detecting cognitive symptoms after stroke, thereby providing further data on the sensitivity and specificity of the OCS in the identification of cognitive deficits in a relatively large sample of first stroke patients.</p>
2	✓			<p><b>Was there a comparison with an appropriate reference standard?</b></p> <p>MMSE, Bamford classification and NIHSS</p> <p><b>Is it worth continuing?</b></p> <p>Yes</p>
3			✓	<p><b>Did all patients get the diagnostic test and reference standard?</b></p> <p>325 first stroke patients were consecutively enrolled</p> <p>For three patients (0.9%) the OCS could not be given at all. In a few cases, some tests could not be administered (see Table 2). The MMSE could not be administered to six patients.</p>
4	✓			<p><b>Could the results of the test have been influenced by the results of the reference standard?</b></p> <p>Yes there is a possibility and the authors have tried to negate this, however, they did not use randomisation - The OCS and MMSE were presented on the same day; order of presentation of the two tests was counterbalanced on an ABAB basis for each rehabilitation centre.</p>

**CONTACTS**

www.unisa.edu.au/cahe  
iCAHE@unisa.edu.au  
Telephone: +61 8 830 22099  
Fax: +61 8 830 22853

University of South Australia  
GPO Box 2471  
Adelaide SA 5001  
Australia

CRICOS Provider Number  
00121B



University of  
South Australia

International Centre for  
Allied Health Evidence

iCAHE

A member of the Sansom Institute

5		✓	<p><b>Is the disease status of the tested population clearly described?</b>                  More detail could have been provided</p> <p><b>TABLE 1   Characteristics of the sample.</b></p> <table border="1"> <thead> <tr> <th></th> <th>Category</th> <th>No. of patients (325)</th> <th>%</th> </tr> </thead> <tbody> <tr> <td rowspan="2">Gender</td> <td>Male</td> <td>178</td> <td>54.7</td> </tr> <tr> <td>Female</td> <td>147</td> <td>45.2</td> </tr> <tr> <td rowspan="2">Etiology</td> <td>Ischemic</td> <td>278</td> <td>85.5</td> </tr> <tr> <td>Hemorrhagic</td> <td>44</td> <td>13.5</td> </tr> <tr> <td rowspan="3">Lesion lateralization</td> <td>Unilateral left hemisphere</td> <td>122</td> <td>37.5</td> </tr> <tr> <td>Unilateral right hemisphere</td> <td>184</td> <td>56.6</td> </tr> <tr> <td>Bilateral/cerebellar</td> <td>19</td> <td>5.8</td> </tr> <tr> <td rowspan="4">Vascular territory affected for ischemic patients: Bamford classification (n = 274)</td> <td>TACI</td> <td>58</td> <td>17.8</td> </tr> <tr> <td>LACI</td> <td>76</td> <td>23.3</td> </tr> <tr> <td>PACI</td> <td>91</td> <td>28</td> </tr> <tr> <td>POCI</td> <td>54</td> <td>16.6</td> </tr> <tr> <td rowspan="4">Stroke severity: NIHSS</td> <td>Minor</td> <td>171</td> <td>52.6</td> </tr> <tr> <td>Moderate</td> <td>136</td> <td>41.8</td> </tr> <tr> <td>Moderate to severe</td> <td>13</td> <td>4</td> </tr> <tr> <td>Severe</td> <td>4</td> <td>1.2</td> </tr> </tbody> </table>		Category	No. of patients (325)	%	Gender	Male	178	54.7	Female	147	45.2	Etiology	Ischemic	278	85.5	Hemorrhagic	44	13.5	Lesion lateralization	Unilateral left hemisphere	122	37.5	Unilateral right hemisphere	184	56.6	Bilateral/cerebellar	19	5.8	Vascular territory affected for ischemic patients: Bamford classification (n = 274)	TACI	58	17.8	LACI	76	23.3	PACI	91	28	POCI	54	16.6	Stroke severity: NIHSS	Minor	171	52.6	Moderate	136	41.8	Moderate to severe	13	4	Severe	4	1.2
	Category	No. of patients (325)	%																																																						
Gender	Male	178	54.7																																																						
	Female	147	45.2																																																						
Etiology	Ischemic	278	85.5																																																						
	Hemorrhagic	44	13.5																																																						
Lesion lateralization	Unilateral left hemisphere	122	37.5																																																						
	Unilateral right hemisphere	184	56.6																																																						
	Bilateral/cerebellar	19	5.8																																																						
Vascular territory affected for ischemic patients: Bamford classification (n = 274)	TACI	58	17.8																																																						
	LACI	76	23.3																																																						
	PACI	91	28																																																						
	POCI	54	16.6																																																						
Stroke severity: NIHSS	Minor	171	52.6																																																						
	Moderate	136	41.8																																																						
	Moderate to severe	13	4																																																						
	Severe	4	1.2																																																						
6	✓		<p><b>Were the methods for performing the test described in detail?</b>                  See Cognitive screening tests under materials and methods</p> <p>The OCS and MMSE were presented on the same day; order of presentation of the two tests was counterbalanced on an ABAB basis for each rehabilitation centre. Tests were administered in a quiet and comfortable setting.</p> <p>More detailed in paper</p>																																																						

**CONTACTS**

www.unisa.edu.au/cahe  
 iCAHE@unisa.edu.au  
 Telephone: +61 8 830 22099  
 Fax: +61 8 830 22853

University of South Australia  
 GPO Box 2471  
 Adelaide SA 5001  
 Australia

CRICOS Provider Number  
 00121B



University of  
 South Australia

International Centre for  
 Allied Health Evidence  
 iCAHE

A member of the Sansom Institute

7		<p><b>What are the results?</b></p> <p>About a third of patients (35.3%) had a performance lower than the cutoff (&lt;22) on the MMSE, whereas 91.6% were impaired in at least one OCS domain, indicating higher incidences of impairment for the OCS. More than 80% of patients showed an impairment in two or more cognitive domains of the OCS. Using the MMSE as a standard of clinical practice, the comparative sensitivity of OCS was 100%. Out of the 208 patients with normal MMSE performance 180 showed impaired performance in at least one domain of the OCS. The discrepancy between OCS and MMSE was particularly strong for patients with milder strokes. As for subtypes of cerebral infarction, fewer patients demonstrated widespread impairments in the OCS in the Posterior Circulation Infarcts category than in the other categories.</p> <p>Overall, the results showed a much higher incidence of cognitive impairment with the OCS than with the MMSE and demonstrated no false negatives for OCS vs MMSE. The authors concluded that OCS is a sensitive screen tool for cognitive deficits after stroke. In particular, the OCS detects high incidences of stroke-specific cognitive impairments, not detected by the MMSE, demonstrating the importance of cognitive profiling.</p>
8	Journal Club to discuss	<p><b>How sure are we about the results? (consequences and cost of alternatives performed?)</b></p> <p>Could discuss how results may be affected by inclusion/exclusion criteria, may limit generalizability to those patients with comorbidities and difficulties with concentrating.</p> <p>- Exclusion criteria stated: Patients with the presence of premorbid psychiatric or neurological disease; patients unable to give informed consent; patients without ability to concentrate for &lt;20 min (as judged by the care team).</p>
9		<p><b>Do you believe the results?</b></p>

**CONTACTS**

www.unisa.edu.au/cahe  
iCAHE@unisa.edu.au  
Telephone: +61 8 830 22099  
Fax: +61 8 830 22853

University of South Australia  
GPO Box 2471  
Adelaide SA 5001  
Australia

CRICOS Provider Number  
00121B



University of  
South Australia

International Centre for  
Allied Health Evidence

iCAHE

A member of the Sansom Institute

**CONTACTS**

www.unisa.edu.au/cahe  
 iCAHE@unisa.edu.au  
 Telephone: +61 8 830 22099  
 Fax: +61 8 830 22853

University of South Australia  
 GPO Box 2471  
 Adelaide SA 5001  
 Australia

CRICOS Provider Number  
 00121B



University of  
 South Australia

International Centre for  
 Allied Health Evidence

iCAHE

A member of the Sansom Institute

10		<p><b>Can the results be applied to the local population? Choose relevant context issues. The following are only suggestions to prompt discussion.</b></p> <p><b>CONTEXT ASSESSMENT</b></p> <ul style="list-style-type: none"> <li>- Infrastructure</li> <li>- Available workforce (? Need for substitute workforce?)</li> <li>- Patient characteristics</li> <li>- Training and upskilling, accreditation, recognition</li> <li>- Ready access to information sources</li> <li>- Legislative, financial &amp; systems support</li> <li>- Health service system, referral processes and decision-makers</li> <li>- Communication</li> <li>- Best ways of presenting information to different end-users</li> <li>- Availability of relevant equipment</li> <li>- Cultural acceptability of recommendations</li> <li>- Others</li> </ul>
11		<p><b>Were all outcomes important to the individual or population considered?</b></p> <p><b>What would be the impact of using this test on your patients/population?</b></p>
12		<p><b>Are the benefits worth the harms and costs?</b></p>
13		<p><b>What do the study findings mean to practice (i.e. clinical practice, systems or processes)?</b></p>
14		<p><b>What are your next steps?</b></p> <p><b>ADOPT, CONTEXTUALISE, ADAPT</b></p> <p><b>And then (e.g. evaluate clinical practice against evidence-based recommendations; organise the next four journal club meetings around this topic to build the evidence base; organize training for staff, etc.)</b></p>
15		<p><b>What is required to implement these next steps?</b></p>