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CORRIGENDUM

I. Zerr, K. Kallenberg, D. M. Summers, C. Romero, A. Taratuto, U. Heinemann, M. Breithaupt, D. Varges, B. Meissner, A. Ladogana, M. Schuur, S. Haik, S. J. Collins, Gerard H. Jansen, G. B. Stokin, J. Pimentel, E. Hewer, D. Collie, P. Smith, H. Roberts, J. P. Brandel, C. van Duijn, M. Pocchiari, C. Begue, P. Cras, R. G. Will and P. Sanchez-Juan. **Updated clinical diagnostic criteria for sporadic Creutzfeldt-Jakob disease.** *Brain* 2009; **132**: 2659–2668; doi:10.1093/brain/awp191.

The authors would like to apologize for an oversight in Figure 1. The corrected version of the criteria is given below:
Figure 1 Amended clinical criteria for sporadic CJD.
Rapidly progressive dementia (obligatory) plus
I. clinical signs

1. myoclonus
2. cerebellar or visual
3. pyramidal or extrapyramidal
4. akinetic mutism

II. tests

1. PSWCs in EEG
2. 14-3-3 detection in CSF
3. high signal abnormalities in caudate nucleus and striatum or at least two cortical regions (temporal-parietal-occipital) either in DWI or FLAIR

Probable CJD
rapidly progressive dementia and two out of I and at least one out of II

Possible CJD
rapidly progressive dementia and two out of I and duration less than 2 years

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