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THE MS-SMART TRIAL IN SECONDARY PROGRESSIVE MS— CURRENT UPDATE

Jeremy Chataway,¹ Siddharthan Chandran,¹ David Miller,² Peter Connick,² Gavin Giovannoni,³ Sue Pavitt,⁴ Nigel Stallard,⁵ Clive Hawkins,⁶ Basil Sharrack,⁷ Domenico Plantone^{1,2,3,4,5,6,7,8}. ¹ University College London; ² University of Edinburgh; ³ Queen Mary University of London; ⁴ University of Leeds; ⁵ University of Warwick; ⁶ University Hospital of North Staffordshire; ⁷ University of Sheffield; ⁸ on behalf of the MS-SMART trialists

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MS-SMART is an ongoing multicentre, multi-arm, double blind, placebo-controlled phase IIIb randomised controlled trial to establish whether putative neuroprotective drugs (fluoxetine, riluzole, amiloride or placebo) can slow down the progression of brain volume loss in secondary progressive multiple sclerosis (SPMS) over 96 weeks using MRI-derived Percentage Brain Volume Change (PBVC) as the primary outcome. 360 patients have been screened so far, 328 (92%) consented and 272 randomized (65% of the total UK cohort—440). Patients will have outcome-data collected after 0, 24, 48 and 96 weeks. The trial is using a range of conventional and novel imaging techniques to better understand neuroprotection in SPMS. Core data includes: new and enlarging T2 lesions count, persistent new T1 hypointense lesion, whole brain atrophy, grey matter volume, and advanced imaging in 2 centres [cervical cord area change, MR spectroscopy and Magnetic Transfer Ratio (MTR)]. Additional measures include: CSF neurofilament levels and optical coherence tomography (OCT) to measure Retinal Nerve Fibre Layer (RNFL) thickness. This abstract describes the ongoing status of the trial. This independent research is awarded by the Efficacy and Mechanism Evaluation Programme (EME) and funded by the Medical Research Council (MRC) and the Multiple Sclerosis Society (MS Society) and managed by the National Institute for Health Research (NIHR) on behalf of the MRC-NIHR partnership.