

Curriculum Vitae Professor Dr Alastair Compston



Name: David Alastair Standish Compston

Date of birth: 23 January 1948

Research Priorities: Multiple sclerosis, pathogenesis and novel therapies, genetics

Alastair Compston is a British neurologist. Now retired, he has been researching the aetiology, disease mechanisms and therapeutics of multiple sclerosis (MS) since the mid-1970s. He is acknowledged as a leader in this research field over several decades. With his many collaborators in Cambridge and internationally, he has been associated with the identification of genetic risk factors for MS, he developed a novel therapy approach with drug development through to licencing of Lemtrada©, and he has advanced concepts on the complex pathogenesis and evolution of disease progression.

Academic and Professional Career

2004	Founder and Chairman, Department of Clinical Neurosciences, University of Cambridge Clinical School, University of Cambridge, Cambridge, UK
2001	Founder and Co-Chairman, Cambridge Neuroscience, Cambridge, UK
1994 - 2001	Consultant Advisor in Neurology to the Chief Medical Officer, UK
1992	Founder and Chairman, Medical Research Council Cambridge Centre for Brain Repair, Cambridge, UK
1989 - 2015	Professor of Neurology, University of Cambridge, Cambridge, UK
1987 - 1988	Professor of Neurology, University of Wales, Cardiff, UK
1982 - 1986	Consultant Neurologist, University of Wales, Cardiff, UK
1978	PhD in Medicine, University of London, London, UK
1975 - 1982	Training in Neurology, National Hospital for Neurology and Neurosurgery, London, UK

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Functions in Scientific Societies and Committees

2019 - 2022	Member and Chairman, Sectional Committee 10 (Health and Human Science), Royal Society of London
2009 - 2010	President, Association of British Neurologists, UK
2004 - 2013	Editor, Brain: A Journal of Neurology
2002 - 2003	President, European Neurological Society
2001 - 2004	Chairman, Panel "Neuroscience and Mental Health", Wellcome Trust, London, UK

Honours and Awarded Memberships

2018	Koetser Award, Betty & David Koetser Foundation for Brain Research, Zurich, Switzerland
2018	Jean Hunter Prize, Royal College of Physicians, London, UK
2016	ABN Medal, Association of British Neurologists, UK
since 2016	Fellow, Royal Society, UK
since 2016	Commander, Order of the British Empire (CBE), UK
2016	Galen Medal, Society of Apothecaries, London, UK
2015	Honorary Fellow, European Academy of Neurology
2015	Hughlings Jackson Medal, London, UK
2015	John Dystel Prize for Multiple Sclerosis Research, National Multiple Sclerosis Society, UK and American Academy of Neurology (AAN), USA
2015	Richard and Mary Cave Award, National Multiple Sclerosis Society, UK
2013	Medal for Scientific Achievement in Neurology, World Federation of Neurology
since 2012	Foreign Member, National Academy of Medicine, USA
2011	Ian McDonald Award, National Multiple Sclerosis Society, UK
2010	Gertrud-Reemtsma Foundation Prize, Max Planck Society, Munich, Germany
since 2008	Member, German National Academy of Sciences Leopoldina, Germany
2007	Charcot Award, Multiple Sclerosis International Federation
2002	Sobek Research Prize, Roman, Marga und Mareille Sobek Stiftung, Stuttgart, Germany

since 2000	Foreign Member, Royal Physiographic Society of Lund, Sweden
since 2000	Fellow, Royal Society of Biology, UK
since 1998	Fellow, Academy of Medical Sciences, UK
since 1997	Fellow, Royal Society of Arts, UK
since 1986	Fellow, Royal College of Physicians, London, UK

Research Priorities

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Multiple sclerosis is an inflammatory disease of the central nervous system. It is established that the body's own immunological defence system intermittently attacks the protective membranes (myelin sheath) of nerve fibres, and the underlying axons, in the brain and spinal cord. This results in episodes of neurological dysfunction manifesting, for example, as impaired vision, loss of sensation and weakness.

Starting with his co-demonstration of the link to HLA-DR15 (1976), Alastair Compston and his many colleagues have systematically shown that multiple sclerosis develops against the background of genetic susceptibility in which many risk variants, each individually exerting a small effect, underly the predisposition to disease. In related work, he has studied the cellular neurobiology of glia, especially myelin-forming cells in the central nervous system, as the basis for understanding the possibilities for enhancing endogenous and exogenous repair.

With others, Alastair Compston developed a novel therapy approach in which the attacking immune cells are depleted using biological medicines (monoclonal antibodies) leaving a sufficient number to deflect coincidental infections but too few to maintain disease activity in the nervous system. Later the repertoire is reconstituted providing significant longer term disease protection but not without adverse effects, especially secondary autoimmunity affecting other organs. The antibody Alemtuzumab (Campath-1H) is now marketed as Lemtrada©. Alastair Compston has shown that it is possible to slow down activity of the disease in the early stages of multiple sclerosis in most patients, and thereby to delay or even prevent subsequent progression. Alemtuzumab is now an established means of treatment amongst a growing number of other therapies for multiple sclerosis, each varying in their risk-benefit ratios.