NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE

Single Technology Appraisal

Natalizumab for treating highly active relapsing-remitting multiple sclerosis ID4020

Provisional Stakeholder List

Consultees	Commentators (no right to submit or appeal)
<u>Company</u>	General
Biogen (natalizumab)	All Wales Therapeutics and Toxicology Control
Patient/carer groups	CentreAllied Health Professionals Federation
Brain and Spine Foundation	Board of Community Health Councils in
Brain CharityMS-UK	Wales ■ British National Formulary
Multiple Sclerosis Society	Care Quality Commission
Multiple Sclerosis Trust	Department of Health, Social Services
Neurological AllianceShift.ms	and Public Safety for Northern IrelandHealthcare Improvement Scotland
South Asian Health Foundation	Medicines and Healthcare products
Specialised Healthcare Alliance	Regulatory Agency
Sue Ryder	Multiple Sclerosis Society WalesNational Association of Primary Care
Healthcare professional groups	National Pharmacy Association
Association of British Neurologists	Neurological Alliance of Scotland
British Association of Neuroscience Nurses	NHS AllianceNHS Confederation
British Geriatrics Society	Scottish Medicines Consortium
British Neuropathological Society	Wales Neurological Alliance
 British Society for Blood and Marrow Transplantation 	 Welsh Health Specialised Services Committee
British Society of Rehabilitation	Committee
Medicine	Possible comparator companies
Chartered Society of PhysiotherapyInstitute of Neurology	Glenmark Pharmaceuticals (fingolimod)Merck Serono (cladribine)
London MS-AHSCT Collaborative	 Novartis Pharmaceuticals (fingolimod)
Group	Roche Products (ocrelizumab)
 National Neurosciences Advisory Group 	Sanofi Genzyme (alemtuzumab)Tillomed Laboratories (fingolimod)
Primary Care and Community	Tillottied Laboratories (illigoliifiod)
Neurology Society	Research groups
Royal College of General PractitionersRoyal College of Nursing	Brain Research UK British Neurological Research Trust
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Provisional stakeholder list for the evaluation of natalizumab for treating highly active relapsing-remitting multiple sclerosis ID4020

Issue date: May 2022

Appendix C

Consultees	Commentators (no right to submit or appeal)
 Royal College of Occupational Therapists Royal College of Pathologists Royal College of Physicians Royal Pharmaceutical Society Royal Society of Medicine Therapists in MS UK Clinical Pharmacy Association UK Multiple Sclerosis Specialist Nurse Association Others Department of Health and Social Care NHS Castle Point and Rochford CCG NHS England NHS North Central London CCG Welsh Government 	 Cochrane Multiple Sclerosis and Rare Diseases of the Central Nervous System Group Cochrane UK Genomics England MRC Clinical Trials Unit National Institute for Health Research Associated Public Health groups Public Health Wales UK Health Security Agency

NICE is committed to promoting equality, eliminating unlawful discrimination and fostering good relations between people who share a protected characteristic and those who do not. Please let us know if we have missed any important organisations from the stakeholder list, and which organisations we should include that have a particular focus on relevant equality issues.

Definitions:

Consultees

Organisations that accept an invitation to participate in the evaluation; the company that markets the technology; national professional organisations; national patient organisations; the Department of Health and Social Care and the Welsh Government and relevant NHS organisations in England.

The company that markets the technology is invited to make an evidence submission, respond to consultations, nominate clinical experts and has the right to appeal against the Final Draft Guidance (FDG).

All non-company consultees are invited to submit a statement¹, respond to consultations, nominate clinical or patient experts, and have the right to appeal against the Final Draft Guidance (FDG).

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¹ Non-company consultees are invited to submit statements relevant to the group they are representing.

Commentators

Organisations that engage in the evaluation process but that are not asked to prepare an evidence submission or statement, are able to respond to consultations and they receive the FDG for information only, without right of appeal. These organisations are: companies that market comparator technologies; Healthcare Improvement Scotland; related research groups where appropriate (for example, the Medical Research Council [MRC], National Cancer Research Institute); other groups (for example, the NHS Confederation, NHS Alliance, and the British National Formulary).

All non-company commentators are invited to nominate clinical or patient experts.