NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE

Single Technology Appraisal

Melphalan flufenamide with dexamethasone for treating relapsed or refractory multiple myeloma ID3862

Provisional stakeholder list

Consultees	Commentators (no right to submit or appeal)
Company	General
Oncopeptides AB (melphalan	All Wales Therapeutics and Toxicology
flufenamide)	Centre
Patient/carer group African Caribbean Leukaemia Trust	 Allied Health Professionals Federation Board of Community Health Councils in Wales
 Anthony Nolan 	
 Black Health Agency 	British National FormularyCare Quality Commission
 Blood Cancer UK 	 Department of Health, Social Services
Cancer Black Care	and Public Safety for Northern Ireland
 Cancer Equality 	 Healthcare Improvement Scotland
Cancer52	 Medicines and Healthcare Products
DKMS	Regulatory Agency
 Helen Rollason Cancer Charity 	 National Association of Primary Care
Independent Cancer Patients Voice	National Pharmacy Association
Leukaemia Cancer Society	NHS Alliance
Leukaemia CARE	NHS Confederation
Leukaemia UK	Scottish Medicines Consortium
Lymphoma Action	Welsh Health Specialised Services
Macmillan Cancer Support	Committee
Maggie's Centres	
Marie Curie	Possible comparator companies
Myeloma UK	 Actavis UK Ltd (bortezomib)
South Asian Health Foundation	 Advanz pharma (dexamethasone)
Specialised Healthcare Alliance	Amgen (carfilzomib)
Tenovus Cancer Care	Aspen (dexamethasone)
	Aspire pharma (bortezomib,
Professional groups	dexamethasone)
Association of Cancer Physicians	Bristol-Myers Squibb Pharmaceuticals
British Blood Transfusion Society	limited (lenalidomide)
British Committee for Standards in	Celgene (pomalidomide) Consiliant health (devemathesene)
Haematology	 Consilient health (dexamethasone) Dr Boddy's laboratories (bortozomib)
British Geriatrics Society	Dr Reddy's laboratories (bortezomib)
British Institute of Radiology	 Glenmark pharmaceuricals (dexamethasone)
British Psychosocial Oncology Society	
British Society for Haematology	S ()
Cancer Research UK	Martindale pharma (dexamethasone)

Provisional stakeholder list for the technology appraisal of melphalan flufenamide with dexamethasone for treating relapsed or refractory multiple myeloma ID3862. Issue date: February 2021 © National Institute for Health and Care Excellence 2021. All rights reserved. Page 1 of 3

Consultees	Commentators (no right to submit or appeal)
 NHS Blood and Transplant Royal College of General Practitioners Royal College of Nursing Royal College of Pathologists Royal College of Physicians Royal College of Radiologists Royal Pharmaceutical Society Royal Society of Medicine Society and College of Radiographers UK CLL Forum UK CLL Forum UK Clinical Pharmacy Association UK Myeloma Forum UK Oncology Nursing Society Others Department of Health and Social Care NHS England NHS Southport and Formby CCG NHS Durham Dales, Easington and Sedgefield CCG Welsh Government 	 Mylan (bortezomib) Novartis pharmaceuticals (panobinostat) Sanofi Genzyme (Isatuximab) Synchrony Pharma Ltd (dexamethasone) Teva (dexamethasone) Thame Laboratories (dexamethasone) Thornton & Ross (bortezomib) Rosemont pharmaceuticals limited (dexamethasone) Thornton & Ross (bortezomib) Rosemont pharmaceuticals limited (dexamethasone) Zentiva (bortezomib) Relevant research groups Cochrane Haematological Malignancies Group Genomics England Institute of Cancer Research Leukaemia UK Leukaemia Busters MRC Clinical Trials Unit National Cancer Research Institute National Cancer Research Network National Institute for Health Research Associated Public Health Groups Public Health England Public Health Wales

NICE is committed to promoting equality, eliminating unlawful discrimination and fostering good relations between people who share a protected characteristic and those who do share it. 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 appraisal; 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 specialists and has the right to appeal against the Final Appraisal Document (FAD).

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

Commentators

Organisations that engage in the appraisal process but that are not asked to prepare an evidence submission or statement, are able to respond to consultations and they receive the FAD for information only, without right of appeal. These organisations are: companies that market comparator technologies; Healthcare Improvement Scotland; other 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 specialists or patient experts.

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