NATIONAL INSTITUTE FOR HEALTH AND CLINICAL EXCELLENCE

Single Technology Appraisal

Lenalidomide in combination with dexamethasone for previously untreated multiple myeloma [ID474]

Matrix of consultees and commentators

Consultees	Commentators (no right to submit or appeal)
Company	General
Celgene (lenalidomide)	All Wales Therapeutics and Toxicology Centre
Patient/carer group	Allied Health Professionals Federation
Black Health Agency	Board of Community Health Councils in
Bloodwise	Wales
Cancer Black Care	British National Formulary
Cancer Equality	Care Quality Commission
Cancer52	Department of Health, Social Services
• DKMS	and Public Safety for Northern Ireland
HAWC	Healthcare Improvement Scotland
Helen Rollason Cancer Charity	 Medicines and Healthcare products
 Independent Cancer Patients Voice 	Regulatory Agency
Leukaemia Cancer Society	National Association for Primary Care
Leukaemia CARE	National Pharmacy Association
Lymphoma Association	NHS Alliance
Macmillan Cancer Support	NHS Commercial Medicines Unit
Maggie's Centres	NHS Confederation
Marie Curie	Scottish Medicines Consortium
Muslim Council of Britain	Welsh Health Specialised Services
Myeloma UK	Committee
South Asian Health Foundation	Possible comparator manufacturers
Specialised Healthcare Alliance	Actavis UK (prednisolone)
Tenovus Cancer Care	Alliance Pharmaceuticals (prednisolone)
Professional groups	Amdipharm (prednisolone)
Professional groupsAssociation of Cancer Physicians	Aspen Europe (melphalan)
British Committee for Standards in	Auden McKenzie (Pharma Division)
Haematology (BCSH)	(dexamethasone)
 British Geriatrics Society 	Baxter Healthcare (cyclophosphamide)
British Psychosocial Oncology Society	Cardinal Health Martindale Products
British Society for Haematology	(dexamethasone)
Cancer Research UK	Celgene (thalidomide)
Royal College of General Practitioners	Concordia International (prednisolone)

National Institute for Health and Clinical Excellence

Matrix for the Single Technology Appraisal lenalidomide in combination with dexamethasone for previously untreated multiple myeloma

Issue date: August 2017

Consultees	Commentators (no right to submit or appeal)
 Royal College of Nursing Royal College of Pathologists Royal College of Physicians Royal Pharmaceutical Society Royal Society of Medicine UK Clinical Pharmacy Association UK Health Forum UK Myeloma Forum UK Oncology Nursing Society Others Department of Health NHS England NHS Havering CCG NHS Sandwell and West Birmingham CCG Welsh Government 	 Hospira UK (dexamethasone) Janssen (bortezomib) Intrapharm Laboratories (prednisolone) Rosemont Pharmaceuticals (dexamethasone) Sandoz Limited (cyclophosphamide) Wockhardt UK (dexamethasone) Zentiva (prednisolone) Relevant research groups Cochrane Haematological Malignancies Group Institute of Cancer Research Leuka 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 not. Please let us know if we have missed any important organisations from the lists in the matrix, and which organisations we should include that have a particular focus on relevant equality issues.

PTO FOR DEFINITIONS OF CONSULTEES AND COMMENTATORS

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 the Welsh Government and relevant NHS organisations in England.

National Institute for Health and Clinical Excellence Matrix for the Single Technology Appraisal lenalidomide in combination with dexamethasone for previously untreated multiple myeloma Issue date: August 2017 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 Appraisal Determination (FAD).

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 Appraisal Determination (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; the relevant National Collaborating Centre (a group commissioned by the Institute to develop clinical guidelines); 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 NHS Commercial Medicines Unit, and the *British National Formulary*.

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

Evidence Review Group (ERG)

An independent academic group commissioned by the National Institute for Health Research (NIHR) Health Technology Assessment Programme (HTA Programme) to assist the Appraisal Committee in reviewing the company evidence submission to the Institute.

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