NATIONAL INSTITUTE FOR HEALTH AND CLINICAL EXCELLENCE

Provisional Single Technology Appraisal (STA)

Lenalidomide for the treatment of deletion 5q in myelodysplastic syndrome

Final matrix of consultees and commentators

Consultees	Commentators (no right to submit or
	appeal)
Manufacturers/sponsors Celgene (lenalidomide) Patient/carer groups Afiya Trust African Caribbean Leukaemia Trust	 General Allied Health Professionals Federation Board of Community Health Councils in Wales British National Formulary Care Quality Commission
 Anthony Nolan Bone Marrow Trust Aplastic Anaemia Trust Black Health Agency Cancer Black Care Cancer Equality Cancer 52 Chronic Myeloid Leukaemia Support Group Counsel and Care Equalities National Council Helen Rollason Heal Cancer Charity Leukaemia CARE Leukaemia Cancer Society Macmillan Cancer Support Maggie's Centres Marie Curie Cancer Care 	 Commissioning Support Appraisals Service Department of Health, Social Services and Public Safety for Northern Ireland Healthcare Improvement Scotland Medicines and Healthcare products Regulatory Agency National Association of Primary Care National Pharmacy Association NHS Alliance NHS Commercial Medicines Unit NHS Confederation Public Health Wales NHS Trust Scottish Medicines Consortium
 MDS UK Patient Support Group Muslim Council of Britain Muslim Health Network Rarer Cancers Foundation South Asian Health Foundation Specialised Healthcare Alliance Sue Ryder Care Tenovus Professional groups Association of Cancer Physicians British Association for Services to the Elderly British Committee for Standardisation 	 Celgene (azacitadine) Relevant research groups Cochrane Haematology Malignancies Group Elimination of Leukaemia Fund Institute of Cancer Research Leukaemia & Lymphoma Research Leukaemia Busters MRC Clinical Trials Unit National Cancer Research Institute National Cancer Research Network National Institute for Health Research Policy Research Institute on Ageing and Ethnicity

National Institute for Health and Clinical Excellence

Matrix for the technology appraisal of lenalidomide for the treatment of deletion 5q in myelodysplastic syndrome

Issue date: September 2012

Consultees	Commentators (no right to submit or appeal)
 in Haematology British Geriatrics Society British Psychosocial Oncology Society British Society for Haematology Cancer Networks Pharmacists Forum Cancer Research UK Royal College of General Practitioners Royal College of Nursing Royal College of Pathologists Royal College of Physicians Royal Pharmaceutical Society Royal Society of Medicine United Kingdom Clinical Pharmacy Association United Kingdom Oncology Nursing Society 	 Research Institute for the Care of Older People Evidence Review Group Kleijnen Systematic Reviews Ltd National Institute for Health Research Health Technology Assessment Programme Associated Guideline Groups National Collaborating Centre for Cancer Associated Public Health Groups None
 Others Department of Health Inner North East London (PCT Cluster) South East London (PCT Cluster) Welsh Assembly Government 	

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 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

National Institute for Health and Clinical Excellence Matrix for the technology appraisal of lenalidomide for the treatment of deletion 5q in myelodysplastic syndrome

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Definitions:

Consultees

Organisations that accept an invitation to participate in the appraisal; the manufacturer(s) or sponsor(s) of the technology; national professional organisations; national patient organisations; the Department of Health and the Welsh Assembly Government and relevant NHS organisations in England.

The manufacturer/sponsor of the technology is invited to make an evidence submission, respond to consultations and has the right to appeal against the Final Appraisal Determination (FAD).

All non-manufacturer/sponsor 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 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: manufacturers of comparator technologies; NHS Quality 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-manufacturers/sponsors commentators are invited to nominate clinical specialists 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 manufacturer/sponsor evidence submission to the Institute.

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