NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE

Single Technology Appraisal

Pembrolizumab for treating relapsed or refractory classical Hodgkin lymphoma after multi-agent chemotherapy

Provisional stakeholder list of consultees and commentators

Consultees	Commentators (no right to submit or appeal)
Company	General
Merck Sharp & Dohme	Allied Health Professionals Federation
(pembrolizumab)	All Wales Therapeutics and Toxicology Centre
Patient/carer groups	Board of Community Health Councils in
African Caribbean Leukaemia Trust	Wales
Anthony Nolan	British National Formulary
Black Health Agency	Care Quality Commission
Bloodwise	Department of Health, Social Services
Cancer Black Care	and Public Safety for Northern Ireland
Cancer Equality	Healthcare Improvement Scotland
Cancer 52	Medicines and Healthcare products
DKMS	Regulatory Agency
Equalities National Council	National Association of Primary Care
HAWC	National Pharmacy Association
Helen Rollason Cancer Charity	NHS Alliance
Independent Cancer Patients Voice	NHS Confederation
Leukaemia Cancer Society	Scottish Medicines Consortium
Leukaemia CARE	Welsh Health Specialised Services
Lymphoma Action	Committee
Macmillan Cancer Support	
Maggie's Centres	Possible comparator companies
Marie Curie	Takeda (brentuximab vedotin)
Muslim Council of Britain	
 South Asian Health Foundation 	Relevant research groups
Specialised Healthcare Alliance	Cochrane Haematological Malignancies Croup
Tenovus Cancer Care	Group
WMUK	Genomics England Institute of Consor Besserah
	Institute of Cancer ResearchLeuka
Professional groups	Leuka Leukaemia Busters
Association of Cancer Physicians	
British Committee for Standards in	Lymphoma Research TrustMRC Clinical Trials Unit
Haematology	
British Geriatrics Society	National Cancer Research Institute National Cancer Research Nativers
 British Institute of Radiology 	National Cancer Research Network

Provisional stakeholder list for the technology appraisal of pembrolizumab for treating relapsed or refractory classical Hodgkin lymphoma after multi-agent chemotherapy Issue date: June 2019

Consultees	Commentators (no right to submit or appeal)
 British Psychosocial Oncology Society British Society for Haematology Cancer Research UK 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 Clinical Pharmacy Association UK Health Forum UK Oncology Nursing Society 	 National Institute for Health Research Associated Public Health Groups Public Health England Public Health Wales
Others Department of Health and Social Care NHS Bromley CCG NHS England NHS North Durham CCG Welsh 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 not. Please let us know if we have missed any important organisations from the lists in the stakeholder list, and which organisations we should include that have a particular focus on relevant equality issues.

PTO FOR DEFINITIONS OF CONSULTEES AND COMMENTATORS

Provisional stakeholder list for the technology appraisal of pembrolizumab for treating relapsed or refractory classical Hodgkin lymphoma after multi-agent chemotherapy Issue date: June 2019

Definitions:

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.

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