NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE

Single Technology Appraisal (STA)

Pembrolizumab for treating advanced or recurrent PD-L1 positive non-small-cell lung cancer after progression with platinum-based chemotherapy [ID840]

Final matrix of consultees and commentators

Consultees	Commentators (no right to submit or appeal)
Company	General
Merck Sharp & Dohme	Allied Health Professionals
(pembrolizumab)	Federation
	Board of Community Health Councils
Patient/carer groups	in Wales
Black Health Agency	 British National Formulary
British Lung Foundation	Care Quality Commission
Cancer Black Care	Department of Health, Social Services
Cancer Equality	and Public Safety for Northern Ireland
HAWC	Healthcare Improvement Scotland
Helen Rollason Cancer Charity	Medicines and Healthcare Products
Independent Cancer Patients Voice	Regulatory Agency
Macmillan Cancer Support	National Association of Primary Care
Maggie's Centres	National Pharmacy Association
Marie Curie Cancer Care	NHS Alliance
Muslim Council of Britain	NHS Commercial Medicines Unit
Roy Castle Lung Cancer Foundation	NHS Confederation
South Asian Health Foundation	Scottish Medicines Consortium
Specialised Healthcare Alliance	2
Tenovus Cancer Care	Comparator companies
UK Lung Cancer Coalition	Accord Healthcare (docetaxel)
	Actavis UK (docetaxel)
Professional groups	Boehringer Ingelheim (nintedanib,
Association of Cancer Physicians	afatinib)
Association of Respiratory Nurse Section	Dr. Reddy's Laboratories (docetaxel)
Specialists	Hospira UK (docetaxel) Lilly UK (remusirumah)
British Geriatrics Society British Institute of Redialague	Lilly UK (ramucirumab) Modes CmbH (desetavel)
British Institute of Radiology British Bayeshasasial Charles and Casiatry	Medac GmbH (docetaxel) Decha Bradueta (arletinib)
British Psychosocial Oncology Society British Thomasia Oncology Crown	Roche Products (erlotinib) Sanafi (decetavel)
British Thoracic Oncology Group British Thoracic Society	Sanofi (docetaxel) Pfizor (crizotinib)
British Thoracic Society	Pfizer (crizotinib) Pristal Myore Squibb (piyolumab)
Cancer Research UK Netional Lynn Conser Forum for Nurses	Bristol-Myers Squibb (nivolumab)
National Lung Cancer Forum for Nurses Daylot Callege of Cancer Propriities are	Relevant research groups
Royal College of General Practitioners Payer College of Nivering	British Association for Lung Research
Royal College of Nursing	Dillish Association for Lung Research

National Institute for Health and Care Excellence

Matrix for the single technology appraisal of pembrolizumab for treating advanced or recurrent PD-L1 positive non-small-cell lung cancer after progression with platinum-based chemotherapy [ID840] Issue date: January 2016 Page 1 of 3

Consultees	Commentators (no right to submit or appeal)
 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 Others Department of Health NHS Basildon and Brentwood CCG NHS Cambridgeshire and Peterborough CCG NHS England Welsh Government 	 Cochrane Airways Group Cochrane Lung Cancer Group Institute of Cancer Research 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.

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 Determination (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 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; 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 specialists or patient experts.

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