NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE

Health Technology Appraisal

Cemiplimab for untreated PD-L1-positive advanced or metastatic non-smallcell lung cancer ID3839

Provisional stakeholder list

Consultees	Commentators (no right to submit or appeal)
Company	<u>General</u>
Sanofi (cemiplimab)	 All Wales Therapeutic and Toxicology Centre
Patient/carer groups	Allied Health Professionals Federation
Black Health Agency	Board of Community Health Councils in
British Lung FoundationCancer Black Care	Wales
Cancer Black CareCancer Equality	British National FormularyCare Quality Commission
 Helen Rollason Cancer Charity 	 Department of Health, Social Services
 Independent Cancer Patients Voice 	and Public Safety for Northern Ireland
Macmillan Cancer Support	Healthcare Improvement Scotland
Maggie's Centres	Medicines and Healthcare Products
Marie Curie	Regulatory Agency
Roy Castle Lung Cancer Foundation	National Association of Primary Care
 South Asian Health Foundation 	 National Pharmacy Association
Specialised Healthcare Alliance	NHS Alliance
Tenovus Cancer Care	NHS Confederation
UK Lung Cancer Coalition	Scottish Medicines Consortium
Drefessional groups	Welsh Health Specialised Services Committee
Professional groups	Committee
Association of Cancer PhysiciansAssociation of Respiratory Nurse Specialists	Possible comparator companies
 British Geriatrics Society 	Accord Healthcare (cisplatin, carboplatin,
 British Institute of Radiology 	docetaxel, gemcitabine, paclitaxel,
 British Psychosocial Oncology Society 	vinorelbine)
British Thoracic Oncology Group	Celgene (paclitaxel)
British Thoracic Society	Consilient Health (carboplatin,
Cancer Research UK	gemcitabine, vinorelbine)
Lung Cancer Nursing UK	• Eli Lilly (pemetrexed)
 National Heart and Lung Institute 	Hospira UK (carboplatin, cisplatin,
Primary Care Respiratory Society UK	docetaxel, gemcitabine, paclitaxel)
Royal College of General Practitioners	Medac GmbH (vinorelbine) Morek, Sharp & Dohmo (boyacizumab)
Royal College of Nursing	 Merck, Sharp & Dohme (bevacizumab, pembrolizumab)
Royal College of Pathologists	 Pfizer (bevacizumab)
Royal College of Physicians	 Pierre Fabre (vinorelbine)
Royal College of Radiologists	 Ranbaxy (gemcitabine)
Royal Pharmaceutical Society	Roche (atezolizumab, bevacizumab)
Royal Society of Medicine Society and College of Pediagraphere	• Sandoz (cisplatin)
Society and College of Radiographers	Seacross Pharmaceuticals (docetaxel,
UK Clinical Pharmacy Association	pemetrexed, paclitaxel)

Provisional stakeholder list for the health technology appraisal of cemiplimab for untreated PD-L² positive advanced or metastatic non-small-cell lung cancer ID3839 Issue date: January 2021

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Consultees	Commentators (no right to submit or appeal)
 UK Oncology Nursing Society <u>Others</u> Department of Health and Social care NHS England NHS Oxfordshire CCG NHS Swindon CCG Welsh Government 	Relevant research groups• Cochrane Lung Cancer Group• Genomics England• Institute of Cancer Research• MRC Clinical Trials Unit• National Cancer Research Institute• National Cancer Research Network• National Institute for Health Research
	 <u>Associated Public Health Groups</u> 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 stakeholder list, and which organisations we should include that have a particular focus on relevant equality issues.

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.

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

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