NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE

Single Technology Appraisal

Amivantamab with carboplatin and pemetrexed for untreated EGFR exon 20 insertion mutation-positive advanced non-small-cell lung cancer ID5110

Provisional Stakeholder List

Provisional Consultees	Provisional Commentators (no right to
	submit or appeal)
Company	<u>General</u>
 Janssen-Cilag (amivantamab) 	All Wales Therapeutics and Toxicology
	Centre
Patient/carer groups	Allied Health Professionals Federation
Asthma and Lung UK	Board of Community Health Councils in
Black Health Agency for Equality	Wales
Cancer Black Care	British National Formulary
Cancer Equality	Care Quality Commission
Cancer52	Department of Health - Northern Ireland
EGFR Positive UK	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	NHS Confederation
 Oncogene-Driven Lung Cancer 	Scottish Medicines Consortium
Patient Alliance UK	Welsh Government
 Roy Castle Lung Cancer Foundation 	Welsh Health Specialised Services
Ruth Strauss Foundation	Committee
South Asian Health Foundation	
Specialised Healthcare Alliance	Possible comparator companies
Tenovus Cancer Care	Amarox (erlotinib)
UK Lung Cancer Coalition	Aspire pharma (pemetrexed)
	AstraZeneca (gefitinib, osimertinib)
Healthcare professional groups	Boehringer Ingelheim (afatinib)
Association of Anaesthetists	Bristol Myers Squibb (paclitaxel)
Association of Cancer Physicians	Cipla EU (gefitinib)
Association of Respiratory Nurse	Eli Lilly and company (pemetrexed)
Specialists	Genus pharmaceuticals (pemetrexed,
Association of Surgeons of Great Britain and Instant.	gefitinib)
Britain and Ireland	Glenmark Pharmaceutical (erlotinib)
British Geriatrics Society	Hospira UK (carboplatin, cisplatin, depotation of the provided in the pr
British Institute of Radiology	docetaxel, gemcitabine, paclitaxel)
British Oncology Pharmacy Association	Medac GmbH (vinorelbine) Marak Chara & Dahras
Association	Merck, Sharp & Dohme (nambra ligumah)
British Psychosocial Oncology Society	(pembrolizumab)

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Provisional Consultees	Provisional Commentators (no right to submit or appeal)
 British Society of Interventional Radiology British Thoracic Oncology Group British Thoracic Society British Transplantation Society Cancer Research UK Lung Cancer and Mesothelioma Clinical Expert Group Lung Cancer Nursing UK National Heart and Lung Institute NHS Blood and Transplant Primary Care Respiratory Society Royal College of Anaesthetists Royal College of General Practitioners Royal College of Pathologists Royal College of Physicians Royal College of Radiologists Royal College of Surgeons Royal College of Surgeons Royal Pharmaceutical Society Royal Society of Medicine Society and College of Radiographers UK Clinical Pharmacy Association UK Oncology Nursing Society Others Department of Health and Social Care NHS England 	 Mylan (erlotinib) Pfizer (pemetrexed, dacomitinib) Pierre Fabre (vinorelbine) Ranbaxy UK (gemcitabine) Roche (atezolizumab, erlotinib) Sandoz (cisplatin, docetaxel, pemetrexed, erlotinib, gefitinib) Seacross pharmaceuticals (docetaxel, paclitaxel, pemetrexed) Sun Pharmaceuticals (gemcitabine, pemetrexed) Synchrony Pharma (gemcitabine) Teva UK (paclitaxel) Zentiva (pemetrexed, erlotinib) Relevant research groups British Association of Lung Research Cochrane Lung Cancer Group Genomics England Institute of Cancer Research MRC Clinical Trials Unit National Institute for Health Research Associated Public Health groups Public Health Wales UK Health Security Agency

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.

Definitions:

Consultees

Organisations that accept an invitation to participate in the evaluation; the company that markets the technology; national professional organisations; national patient organisations; the Department of Health and Social Care and relevant NHS organisations in England.

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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 Draft Guidance (FDG).

All non-company consultees are invited to submit a statement relevant to the group they are representing, respond to consultations, nominate clinical or patient experts and have the right to appeal against the Final Draft Guidance (FDG).

Consultee or commentator stakeholders are provisional until a CA&U form is signed at appraisal stage.

Commentators

Organisations that engage in the evaluation process but that are not asked to prepare an evidence submission or statement, are able to respond to consultations and they receive the FDG for information only, without right of appeal. These organisations are: companies that market comparator technologies; Healthcare Improvement Scotland; related research groups where appropriate (for example, the Medical Research Council [MRC]); other groups (for example, the NHS Confederation and the British National Formulary).

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

Consultee or commentator stakeholders are provisional until a CA&U form is signed at appraisal stage.