NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE

Single Technology Appraisal (STA)

Lorlatinib for treating ALK-positive advanced non-small-cell lung cancer

Provisional matrix of consultees and commentators

Consultees	Commentators (no right to submit or appeal)
Company Pfizer (Iorlatinib) Patient/carer groups Black Health Agency British Lung Foundation Cancer Black Care Cancer Equality HAWC Helen Rollason Cancer Charity Independent Cancer Patients Voice Macmillan Cancer Support Maggie's Centres Marie Curie Muslim Council of Britain Roy Castle Lung Cancer Foundation South Asian Health Foundation Specialised Healthcare Alliance Tenovus Cancer Care UK Lung Cancer Coalition	 General All Wales Therapeutic and Toxicology Centre Allied Health Professionals Federation Board of Community Health Councils in Wales British National Formulary Care Quality Commission 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 Confederation Scottish Medicines Consortium Welsh Health Specialised Services Committee
 Professional groups Association of Cancer Physicians Association of Respiratory Nurse Specialists British Geriatrics Society British Institute of Radiology British Psychosocial Oncology Society British Thoracic Oncology Group British Thoracic Society Cancer Research UK National Lung Cancer Forum for Nurses Primary Care Respiratory Society UK Royal College of General Practitioners Royal College of Pathologists Royal College of Physicians 	 Possible comparator companies Accord Healthcare (carboplatin, cisplatin, docetaxel, gemcitabine, paclitaxel, vinorelbine) Boehringer Ingelheim (nintedanib) Bristol-Myers Squibb Pharmaceuticals (nivolumab) Celgene (paclitaxel) Eli Lilly (gemcitabine) Hospira UK (carboplatin, cisplatin, docetaxel, gemcitabine, paclitaxel) Medac GmbH (docetaxel, paclitaxel, vinorelbine) Merck Sharp & Dohme (pembrolizumab) Novartis (ceritinib)

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Consultees Commentators (no right to submit or appeal) Royal College of Radiologists Pfizer (crizotinib) Royal Pharmaceutical Society Pierre Fabre (vinorelbine) Roche Products (alectinib, Royal Society of Medicine Society and College of Radiographers atezolizumab) **UK Clinical Pharmacy Association** Sandoz (cisplatin) Sanofi (docetaxel) **UK Health Forum Seacross Pharmaceuticals** UK Oncology Nursing Society (docetaxel) Sun Pharmaceuticals (gemcitabine) Others Takeda (brigatinib) Department of Health and Social Care NHS England Relevant research groups NHS Havering CCG Cochrane Lung Cancer Group NHS West Suffolk CCG Institute of Cancer Research Welsh Government 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 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 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 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.