NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE

Single Technology Appraisal (STA)

Dabrafenib in combination with trametinib for treating advanced, metastatic BRAF V600E mutation-positive non-small-cell lung cancer [ID3851]

Consultees	Commentators (no right to submit or appeal)
 Company Novartis pharmaceuticals (dabrafenib and trametinib) Patient/carer groups Asthma and Lung UK Black Health Agency British Lung Foundation Cancer Black Care Cancer Equality Helen Rollason Cancer Charity Independent Cancer Patients Voice Macmillan Cancer Support Maggie's Centres Marie Curie Roy Castle Lung Cancer Foundation Ruth Strauss Foundation Specialised Healthcare Alliance Tenovus Cancer Care UK Lung Cancer Coalition 	 <u>General commentators</u> All Wales Therapeutics 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
 <u>Professional groups</u> Association of Anaesthetists Association of Cancer Physicians Association of Respiratory Nurse Specialists Association of Surgeons of Great Britain and Ireland Association for Cancer Surgery British Geriatrics Society British Institute of Radiology British Oncology Pharmacy Association 	 <u>Comparator companies</u> Accord Healthcare (pemetrexed, docetaxel, carboplatin, cisplatin, gemcitabine, paclitaxel, vinorelbine) Aspire pharma (pemetrexed) Boehringer Ingelheim (nintedanib) Bristol Myers Squibb Pharmaceuticals (paclitaxel, nivolumab) Eli Lilly and company (pemetrexed) Genus pharmaceuticals (pemetrexed) Hospira UK (carboplatin, docetaxel, gemcitabine, paclitaxel, cisplatin)

Final Stakeholder List

Final stakeholder list for the single technology appraisal of dabrafenib in combination with trametinib for treating advanced, metastatic BRAF V600E mutation-positive non-small-cell lung cancer [ID3851] Issue date: June 2022

 Cancer Research UK Lung Cancer Nursing UK National Heart and Lung Institute National Lung Cancer Forum for	 (pembrolizumab) Mylan (pemetrexed) Pfizer (pemetrexed, bevacizumab) Pierre Fabre (vinorelbine) Roche (atezolizumab, bevacizumab) Sandoz (pemetrexed, docetaxel) Seacross pharmaceuticals
Nurses Primary Care Respiratory Society Royal College of Anesthetists Royal College of General Practitioners Royal College of Pathologists Royal College of Physicians Royal College of Radiologists Royal College of Surgeons Royal Society of Medicine Society and College of Radiographers UK Clinical Pharmacy Association UK Oncology Nursing Society	(pemetrexed, docetaxel, paclitaxel) Sun Pharmaceuticals (pemetrexed, gemcitabine) Teva UK (paclitaxel) Zentiva (pemetrexed, bevacizumab) Relevant research groups British Association of Lung Research Cochrane Lung Cancer Group Cochrane UK Genomics England Institute of Cancer Research MRC Clinical Trials Unit National Cancer Research Network National Institute for Health & Care Research Associated Public Health Groups UK Health Security Agency
	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.

Final stakeholder list for the single technology appraisal of dabrafenib in combination with trametinib for treating advanced, metastatic BRAF V600E mutation-positive non-small-cell lung cancer [ID3851] Issue date: June 2022

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 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 experts and has the right to appeal against the Final Draft Guidance (FDG).

All non-company consultees are invited to submit a statement¹, respond to consultations, nominate clinical or patient experts and have the right to appeal against the Final Draft Guidance (FDG).

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], 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 or patient experts.

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

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