NATIONAL INSTITUTE FOR HEALTH AND CLINICAL EXCELLENCE

Proposed Single Technology Appraisal (STA)

Afatinib for the treatment of epidermal growth factor receptor mutation positive locally advanced or metastatic non-small cell lung cancer

Provisional matrix of consultees and commentators

	Commentators (no right to submit or appeal)
 Boehringer Ingelheim (afatinib) <u>Patient/carer groups</u> Afiya Trust Black Health Agency British Lung Foundation Cancer Black Care Cancer Equality Counsel and Care Equalities National Council Helen Rollason Heal Cancer Charity Macmillan Cancer Support Maggie's Centres Marie Curie Cancer Care Muslim Council of Britain Muslim Health Network Roy Castle Lung Cancer Foundation Specialised Healthcare Alliance Tenovus UK Lung Cancer Coalition 	 General Allied Health Professionals Federation Board of Community Health Councils in Wales British National Formulary Care Quality Commission Commissioning Support Appraisals Service 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 Commercial Medicines Unit NHS Confederation Public Health Wales NHS Trust Scottish Medicines Consortium Possible comparator manufacturers Accord Healthcare (carboplatin, cisplatin, gemcitabine, paclitaxel) Actavis UK (docetaxel, gemcitabine, paclitaxel, vinorelbine) AstraZeneca (gefitinib) Bristol-Myers Squibb (carboplatin, cisplatin, paclitaxel) Hospira UK (carboplatin, cisplatin docetaxel, gemcitabine, paclitaxel, vinorelbine) Lilly UK (gemcitabine, pemetrexed)

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Consultees	Commentators (no right to submit or appeal)
 Cancer Research UK National Lung Cancer Forum for Nurses Primary Care Respiratory Society UK Royal College of General Practitioners Royal College of Nursing Royal College of Pathologists Royal College of Physicians Royal College of Radiologists Royal College of Radiologists Royal Pharmaceutical Society Royal Society of Medicine Society and College of Radiographers United Kingdom Clinical Pharmacy Association United Kingdom Oncology Nursing Society 	 Medac UK (docetaxel, gemcitabine, paclitaxel, vinorelbine) Mylan UK (gemcitabine) Pfizer (cisplatin) Pierre Fabre (vinorelbine) Roche Products (erlotinib) Sandoz (carboplatin, cisplatin, docetaxel, gemcitabine, paclitaxel, vinorelbine) Sanofi (docetaxel) Sun Pharmaceuticals (carboplatin, gemcitabine) Teva UK (carboplatin, cisplatin, gemcitabine, paclitaxel) Wockhardt UK (carboplatin, cisplatin, paclitaxel, vinorelbine)
Others Department of Health North West London PCT Cluster South East London PCT Cluster Welsh Government 	Relevant research groups• British Thoracic Oncology 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• Research Institute for the Care of Older PeopleEvidence Review Group• Evidence Review Group tbc• National Institute for Health Research Health Technology Assessment ProgrammeAssociated Guideline Groups• National Collaborating Centre for CancerAssociated Public Health Groups• tbc

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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 manufacturer(s) or sponsor(s) of the technology; national professional organisations; national patient organisations; the Department of Health and the Welsh Government and relevant NHS organisations in England.

The manufacturer/sponsor of 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-manufacturer/sponsor 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: manufacturers of comparator technologies; Healthcare Improvement Scotland ; the relevant National Collaborating Centre (a group commissioned by the Institute to develop clinical guidelines); 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-manufacturers/sponsors commentators are invited to nominate clinical specialists or patient experts.

Evidence Review Group (ERG)

An independent academic group commissioned by the National Institute for Health Research (NIHR) Health Technology Assessment Programme (HTA Programme) to assist the Appraisal Committee in reviewing the manufacturer/sponsor evidence submission to the Institute.

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

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