NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE

Single Technology Appraisal

Encorafenib with binimetinib and cetuximab for previously treated BRAF V600E mutation-positive metastatic colorectal cancer ID1598

Stakeholder list of consultees and commentators

Consultees	Commentators (no right to submit or appeal)
<u>Company</u>	General
Pierre Fabre (binimetinib, encorafenib)	 All Wales Therapeutics and Toxicology Centre
 Patient/carer groups Beating Bowel Cancer Black Health Agency Bladder and Bowel Community Bowel Cancer Information Bowel Cancer UK Cancer Black Care Cancer Equality Cancer52 Colostomy UK Helen Rollason Cancer Charity IA: Ileostomy and Internal Pouch Support Group Independent Cancer Support 	 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
 Macmillan Cancer Support Maggie's Centres Marie Curie Muslim Council of Britain Pelican Cancer Foundation 	 Scottish Medicines Consortium Scottish Society of Gastroenterology Welsh Health Specialised Services Committee
 South Asian Health Foundation Specialised Healthcare Alliance Tenovus Cancer Care 	 <u>Possible comparator companies</u> Accord Healthcare (capecitabine, fluorouracil, irinotecan, oxaliplatin) Consilient Healthcare (fluorouracil)
 Professional groups Association of Cancer Physicians Association of Coloproctology of Great Britain and Ireland British Geriatrics Society British Institute for Radiology British Psychosocial Oncology Society British Society of Gastroenterology Cancer Research UK Pelican Cancer Foundation Primary Care Society of 	 Dr Reddy's Laboratories (capecitabine) Medac (capecitabine, fluorouracil, folinic acid, irinotecan, oxaliplatin) Merck (cetuximab) Mylan (capecitabine) Pfizer (fluorouracil, irinotecan, oxaliplatin, raltitrexed) Roche (capecitabine) Sandoz (oxaliplatin) Sanofi (oxaliplatin) Seacross (irinotecan)

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Consultees	Commentators (no right to submit or
	appeal)
Gastroenterology	• Servier Laboratories (trifluridine-tipitacil)
Royal College of General Practitioners	 Sun Pharma (oxaliplatin)
Royal College of Nursing	
Royal College of Pathologists	Relevant research groups
Royal College of Physicians	Bowel & Cancer Research
Royal College of Radiologists	Cochrane Colorectal Cancer Group
Royal Pharmaceutical Society	Guts UK
Royal Society of Medicine	Genomics England
• Society and College of Radiographers	Institute of Cancer Research
UK Clinical Pharmacy Association	MRC Clinical Trials Unit
UK Health Forum	National Cancer Research Institute
UK Oncology Nursing Society	National Cancer Research Network
	National Institute for Health Research
<u>Others</u>	
• Department of Health and Social Care	Associated Public Health Groups
NHS Bradford City CCG	Public Health England
NHS Eastern Cheshire CCG	Public Health Wales
NHS England	
Welsh Government	

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.

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 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.