NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE

Single Technology Appraisal

Dabrafenib with trametinib for treating BRAF V600E mutation-positive glioma in children and young people aged 1 to 17 ID5104

Provisional Stakeholder List

Consultees	Commentators (no right to submit or
	appeal)
Company	General
Novartis (dabrafenib, trametinib)	All Wales Therapeutics & Toxicology Centre
Patient/carer groups	Allied Health Professionals Federation
Astro Brain Tumour Fund	Board of Community Health Councils in
Black Health Agency for Equality	Wales
Brain and Spine Foundation	British National Formulary
Brain Tumour Action	Care Quality Commission
Brainstrust	Department of Health, Social Services
Cancer 52	and Public Safety for Northern Ireland
Cancer Black Care	Healthcare Improvement Scotland
Cancer Equality	Medicines and Healthcare products
Childhood Cancer Patients Alliance	Regulatory Agency
Children with Cancer	National Association of Primary Care National Pharmacus Association
Headcase Cancer Trust	National Pharmacy Association
Helen Rollason Cancer Charity	NHS Confederation Section Medicines Consertium
Independent Cancer Patients Voice	Scottish Medicines ConsortiumWelsh Government
Macmillan Cancer Support Macmillan Cantrage	Welsh GovernmentWelsh Health Specialised Services
Maggie's Centres Maria Curia	Committee
Marie Curie Neurolagiaal Allianea	Gommittee
Neurological AllianceSouth Asian Health Foundation	Possible comparator companies
Solving Kids Cancer	None
Specialised Healthcare Alliance	
Teenage Cancer Trust	Relevant research groups
Tenovus Cancer Care	Andrew McCartney Trust Fund for Brain
The Brain Charity	Tumour Research
The Brain Tumour Charity	Brain Tumour Research
Young Lives vs Cancer	Brain Tumour Research Campaign
	Cochrane Childhood Cancer Group
Healthcare professional groups	Cochrane UK
Association of British Neurologists	Genomics England
Association of Cancer Physicians	Institute of Cancer Research
Association of Surgeons of Great	Institute of Neurology, UCL
Britain and Ireland	MRC Clinical Trials Unit
British Association of Surgical	National Cancer Research Institute
Oncology	National Institute for Health Research

Provisional stakeholder list for the evaluation of dabrafenib with trametinib for treating BRAF V600E mutation-positive glioma in children and young people aged 1 to 17 ID5104. Issue date: May 2023 © National Institute for Health and Care Excellence 2023. All rights reserved. Page 1 of 3

Consultees	Commentators (no right to submit or appeal)
 British Geriatrics Society British Institute of Radiology British Oncology Pharmacy Association British Psychosocial Oncology Society British Skull Base Society Cancer Research UK Children's Cancer & Leukaemia Group National Neuroscience Advisory Group Neonatal and Paediatric Pharmacists Group 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 Society of British Neurological Surgeons UK Clinical Pharmacy Association UK Oncology Nursing Society 	 Oracle Cancer Trust Associated Public Health groups Public Health Wales UK Health Security Agency
Department of Health and Social CareNHS England	

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 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 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.

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