NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE

Single Technology Appraisal

Ivosidenib for treating advanced cholangiocarcinoma with an IDH1 mutation after at least 1 therapy ID6164

Provisional Stakeholder List

Consultees	Commentators (no right to submit or appeal)
 <u>Company</u> Servier Laboratories (ivosidenib) <u>Patient/carer groups</u> AMMF – The Cholangiocarcinoma Charity Black Health Agency for Equality British Liver Trust Cancer Black Care Cancer Equality Cancer52 GUTS UK Helen Rollason Cancer Charity Independent Cancer Patients Voice Liver4Life Macmillan Cancer Support Maggie's Centres Marie Curie South Asian Health Foundation Specialised Healthcare Alliance Tenovus Cancer Care 	 <u>General</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 Confederation Scottish Medicines Consortium Scottish Society of Gastroenterology Welsh Government Welsh Health Specialised Services Committee
 Healthcare professional groups Association of Anaesthetists Association of Cancer Physicians Association of Surgeons of Great Britain and Ireland Association of Upper Gastrointestinal Surgeons of Great Britain and Ireland British Association of Surgical Oncology British Geriatric Society British Institute of Radiology British Psychosocial Oncology Society British Society of Gastroenterology British Society of Interventional 	 <u>Possible comparator companies</u> AAH Pharmaceuticals (calcium folinate) Accord Healthcare (fluorouracil, oxaliplatin) Alliance Healthcare (calcium folinate) Consilient Health (oxaliplatin) DE Pharmaceuticals (calcium folinate) Fresenius Kabi (oxaliplatin) Medac Healthcare (fluorouracil, oxaliplatin) Medihealth (calcium folinate) Pfizer (calcium folinate, fluorouracil, oxaliplatin) Seacross Pharmaceuticals (oxaliplatin)

Provisional stakeholder list for evaluation of ivosidenib for treating advanced cholangiocarcinoma with an IDH1 mutation after at least 1 therapy ID6164 Issue date: November 2022

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Consultees	Commentators (no right to submit or appeal)
 Radiology Cancer Research UK Cholangiocarcinoma-UK Primary Care Society for Gastroenterology 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 Medicine Society and College of Radiographers UK Clinical Pharmacy Association UK Oncology Nursing Society Others Department of Health and Social Care NHS England 	 Sigma Pharmaceuticals (calcium folinate) Sun Pharmaceutical Industries (oxaliplatin) Teva (calcium folinate) Relevant research groups Cochrane UK Cochrane Upper Gastrointestinal and Pancreatic Diseases Group Genomics England Institute of Cancer Research MRC Clinical Trials Unit National Cancer Research Institute National Institute for Health Research Associated Public Health groups Public Health Wales UK Health Security Agency

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.

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.

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