NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE

Single Technology Appraisal

Ibrutinib with venetoclax for untreated chronic lymphocytic leukaemia [ID3860]

Final stakeholder list

Consultees	Commentators (no right to submit or appeal)
 Company Janssen-Cilag (ibrutinib) Patient/carer groups African Caribbean Leukaemia Trust Anthony Nolan Black Health Agency for Equality Blood Cancer UK Cancer Black Care Cancer Equality Cancer52 Chronic Lymphocytic Leukaemia Support Association DKMS Helen Rollason Cancer Charity Independent Cancer Patients Voice Leukaemia Care Lymphoma Action Macmillan Cancer Support Maggie's Centres Marie Curie South Asian Health Foundation Specialised Healthcare Alliance Tenovus Cancer Care WMUK Professional groups 	 General 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 Hospital Information Services – Jehovah's Witnesses Medicines and Healthcare products Regulatory Agency National Association of Primary Care National Pharmacy Association NHS Alliance NHS Blood and Transplant NHS Confederation Scottish Medicines Consortium Welsh Health Specialised Services Committee Comparator companies AbbVie (venetoclax) Accord Healthcare (bendamustine,
 Association of Cancer Physicians British Blood Transfusion Society British Committee for Standards in Haematology British Geriatrics Society British Institute of Radiology British Psychosocial Oncology Society British Society for Haematology 	fludarabine) Aspen (chlorambucil) AstraZeneca UK (acalabrutinib) Baxter Healthcare (cyclophosphamide) Dr Reddy's Laboratories (bendamustine) Gilead Sciences (idelalisib)

Final stakeholder list for the single technology appraisal of ibrutinib with venetoclax for untreated chronic lymphocytic leukaemia [ID3860]

Issue date: April 2022

Consultees Commentators (no right to submit or appeal) Janssen-Cilag (ibrutinib) Cancer Research UK Royal College of General Practitioners Napp Pharmaceuticals (rituximab) Royal College of Nursing Pfizer (rituximab) Royal College of Pathologists Roche Products (obinutuzumab, Royal College of Physicians rituximab) Sandoz (cyclophosphamide, Royal College of Radiologists rituximab) Royal Pharmaceutical Society Sanofi (fludarabine) Royal Society of Medicine Seacross Pharmaceuticals Society and College of Radiographers (bendamustine) **UK Clinical Pharmacy Association** Zentiva (bendamustine) **UK CLL Forum UK Oncology Nursing Society** Relevant research groups Cochrane Haematological Malignancies Group Department of Health and Social Care Cochrane UK NHS England Genomics England NHS Leeds CCG Institute of Cancer Research NHS Telford & Wrekin CCG Leukaemia Busters Welsh Government Leukaemia UK Lymphoma Research Trust MRC Clinical Trials Unit National Cancer Research Institute National Cancer Research Network 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 appraisal; the company that markets the technology; national professional organisations; national patient

Final stakeholder list for the single technology appraisal of ibrutinib with venetoclax for untreated chronic lymphocytic leukaemia [ID3860]

Issue date: April 2022

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 Appraisal Document (FAD).

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 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; 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.

Final stakeholder list for the single technology appraisal of ibrutinib with venetoclax for untreated chronic lymphocytic leukaemia [ID3860]

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