NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE

Single Technology Appraisal

Lisocabtagene maraleucel for treating relapsed or refractory aggressive B-cell non-Hodgkin lymphoma ID3887

Provisional Stakeholder List

Consultees	Commentators (no right to submit or
	appeal)
 Company Bristol-Myers Squibb (Lisocabtagene maraleucel) Patient/carer groups African Caribbean Leukaemia Trust Anthony Nolan Black Health Agency for Equality Blood Cancer UK Cancer Black Care Cancer Equality Cancer52 DKMS Helen Rollason Cancer Charity Independent Cancer Patients Voice Kevin Kararwa Leukaemia Trust Leukaemia Cancer Society Leukaemia CARE Lymphoma Action 	General All Wales Therapeutics & 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 Healthcare Improvement Scotland Medicines and Healthcare products Regulatory Agency National Association of Primary Care National Pharmacy Association NHS Confederation Scottish Medicines Consortium Welsh Government
 Lympnoma Action Macmillan Cancer Support Maggie's Centres Marie Curie South Asian Health Foundation Specialised Healthcare Alliance Tenovus Cancer Care WMUK 	 Welsh Health Specialised Services Committee Possible comparator companies Accord Healthcare (carboplatin, cisplatin, cytarabine, epirubicin, etoposide, gemcitabine) Advanz Pharma (dexamethasone)
 Healthcare professional groups 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 	 Aspen (dexamethasone) Baxter Healthcare (ifosfamide) Cheplapharm Arzneimittel GmbH (etoposide) Glenmark Pharmaceuticals (dexamethasone) Hameln Pharma (dexamethasone) Hospira UK (carboplatin, cisplatin,

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Issue date: January 2024

Consultees	Commentators (no right to submit or appeal)
 British Society for Haematology Cancer Research UK Royal College of General Practitioners Royal College of Nursing Royal College of Pathologists Royal College of Physicians Royal College of Radiologists Royal Pharmaceutical Society Royal Society of Medicine Society and College of Radiographers UK Clinical Pharmacy Association UK Cutaneous Lymphoma Group UK Oncology Nursing Society Others Department of Health and Social Care NHS England 	cytarabine, dexamethasone, gemcitabine) Jazz Pharmaceuticals UK (cytarabine) Martindale Pharma (dexamethasone) medac GmbH (epirubicin, etoposide) Napp Pharmaceuticals (rituximab) Panpharma UK (dexamethasone) Pfizer (cytarabine, epirubicin, methylprednisolone, rituximab) Roche Products (rituximab) Rosemont Pharmaceuticals (dexamethasone) Sun Pharmaceutical (gemcitabine) Synchrony Pharma (dexamethasone) Sandoz (cisplatin, rituximab) Thame Laboratories (dexamethasone) Wockhardt UK (dexamethasone)
	Relevant research groups Cochrane Haematological Group Cochrane UK Genomics England Institute of Cancer Research Leukaemia Busters Leukaemia UK Lymphoma Research Trust MRC Clinical Trials Unit National Cancer Research Institute High Grade Lymphoma Subgroup 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:

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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.