

NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE

Single Technology Appraisal

Epcoritamab for treating relapsed or refractory follicular lymphoma ID6338

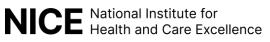
Provisional Stakeholder List

Consultees	Commentators (no right to submit or appeal)
Company	
 <u>Company</u> AbbVie (epcoritamab) 	 <u>General</u> All Wales Therapeutics and Toxicology Centre
 Patient/carer groups African Caribbean Leukaemia Trust Anthony Nolan Blood Cancer UK Cancer Black Care Follicular Lymphoma Foundation Helen Rollason Cancer Charity Independent Cancer Patients Voice Kevin Kararwa Leukaemia Trust Leukaemia Cancer Society Leukaemia UK Lymphoma Action Macmillan Cancer Support Maggie's Centres Marie Curie South Asian Health Foundation 	 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 Welsh Government Welsh Health Specialised Services Committee
 Specialised Healthcare Alliance Tenovus Cancer Care <u>Healthcare professional groups</u> Association of Cancer Physicians British Geriatrics Society British Institute of Radiology British Oncology Pharmacy Association British Psychosocial Oncology Society British Society for Haematology British Society of Interventional Radiology British Transplantation Society Cancer Research UK NHS Blood and Transplant Royal College of General Practitioners 	 Possible comparator companies Advanz Pharma (lenalidomide) Amarox (lenalidomide) Biocon Pharma (lenalidomide) Bristol Myers Squibb (lenalidomide, lisocabtagene maraleucel) Celltrion Healthcare (rituximab) Dr Reddy's Laboratories (bendamustine) Grindeks Kalceks UK (lenalidomide) Mylan (lenalidomide) Pfizer (rituximab) Piramal Critical Care (lenalidomide) Ranbaxy (lenalidomide) Roche (obinutuzumab, rituximab) Sandoz (lenalidomide, rituximab)

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Issue date: November 2024

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Consultees	Commentators (no right to submit or appeal)
Royal College of Nursing	Seacross Pharmaceuticals
Royal College of Pathologists	(bendamustine)
Royal College of Physicians	 Teva UK (lenalidomide)
Royal College of Radiologists	 Thornton & Ross (lenalidomide)
Royal Pharmaceutical Society	 Zentiva (bendamustine)
Royal Society of Medicine	
• Society and College of Radiographers	Relevant research groups
UK Clinical Pharmacy Association	 Cochrane Haematology Group
UK Cutaneous Lymphoma Group	 Genomics England
UK Oncology Nursing Society	 Institute of Cancer Research
	Leukaemia Busters
<u>Others</u>	 Lymphoma Research Trust
Department of Health and Social Care	MRC Clinical Trials Unit
NHS England	National Cancer Research Institute High
Ŭ	Grade Lymphoma Subgroup
	National Institute for Health Research
	Associated Public Health group
	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

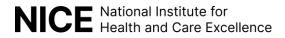
Organisations that accept an invitation to participate in the evaluation; the company that markets the technology; national professional organisations; national patient organisations; the Department of Health and Social Care and relevant NHS organisations in England.

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 statement relevant to the group they are representing, respond to consultations, nominate clinical or patient experts and have the right to appeal against the Final Draft Guidance (FDG).

Commentators

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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]); other groups (for example, the NHS Confederation and the British National Formulary).

All non-company commentators are invited to nominate clinical or patient experts.