# NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE

## Single Technology Appraisal

# Glofitamab with gemcitabine and oxaliplatin for treating relapsed or refractory diffuse large B-cell lymphoma ID6202

## **Provisional Stakeholder List**

Commentators (no right to submit or
appeal)
General
All Wales Therapeutics and Toxicology Centre
<ul> <li>Allied Health Professionals Federation</li> <li>Board of Community Health Councils in Wales</li> <li>British National Formulary</li> <li>Care Quality Commission</li> <li>Department of Health, Social Services and Public Safety for Northern Ireland</li> <li>Healthcare Improvement Scotland</li> <li>Medicines and Healthcare products Regulatory Agency</li> <li>National Association of Primary Care</li> <li>National Pharmacy Association</li> <li>NHS Confederation</li> <li>Scottish Medicines Consortium</li> <li>Welsh Government</li> <li>Welsh Health Specialised Services Committee</li> </ul> Possible comparator companies <ul> <li>AbbVie (epcoritamab)</li> <li>Accord Healthcare (bendamustine, etoposide, gemcitabine, prednisolone, oxaliplatin)</li> <li>Actavis (bendamustine, etoposide, gemcitabine, prednisolone)</li> <li>Advanz Pharma (prednisolone)</li> <li>Aspen (chlorambucil)</li> </ul>
• Baxter Healthcare (cyclophosphamide,
ifosfamide)
Bristol-Myers Squibb Pharmaceuticals

Provisional stakeholder list for evaluation of epcoritamab for glofitamab with gemcitabine and oxaliplatin for treating relapsed or refractory diffuse large B-cell lymphoma ID6202 Issue date: March 2024

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Consultees	Commentators (no right to submit or
<ul> <li>British Society for Haematology</li> <li>British Society of Interventional Radiology</li> <li>British Transplantation Society</li> <li>Cancer Research UK</li> <li>NHS Blood and Transplant</li> <li>Royal College of General Practitioners</li> <li>Royal College of Pathologists</li> <li>Royal College of Physicians</li> <li>Royal College of Radiologists</li> <li>Royal College of Radiologists</li> <li>Royal College of Radiologists</li> <li>Royal College of Medicine</li> <li>Society and College of Radiographers</li> <li>UK Clinical Pharmacy Association</li> <li>UK Cutaneous Lymphoma Group</li> <li>UK Oncology Nursing Society</li> </ul> Others <ul> <li>Department of Health and Social Care</li> <li>NHS England</li> </ul>	<ul> <li>appeal) <ul> <li>(etoposide)</li> <li>Consilient Health (dexamethasone)</li> <li>Dr Reddy's Laboratories UK (bendamustine)</li> <li>Eli Lilly and Company (gemcitabine)</li> <li>Hospira UK (gemcitabine, oxaliplatin)</li> <li>Kite, a Gilead company (axicabtagene ciloleucel)</li> <li>Medac UK (bendamustine, etoposide, lomustine, oxaliplatin)</li> <li>Napp Pharmaceuticals (bendamustine, , rituximab)</li> <li>Pfizer (gemcitabine, vincristine)</li> <li>Ranbaxy (UK), a Sun Pharmaceutical Company (gemcitabine)</li> <li>Roche (glofitamab, polatuzumab vedotin, rituximab)</li> <li>Sandoz (cyclophosphamide, rituximab)</li> <li>Seacross pharmaceuticals (bendamustine)</li> <li>Servier Laboratories (pixantrone)</li> <li>Swedish Orphan Biovitrum (loncastuximab tesirine)</li> <li>Teva Nederland B.V. (bleomycin)</li> <li>Zentiva (bendamustine, prednisolone)</li> </ul> Relevant research groups <ul> <li>Cochrane Haematology Group</li> <li>Cochrane UK</li> <li>Genomics England</li> <li>Institute of Cancer Research</li> <li>Leukaemia Busters</li> <li>Leukaemia UK</li> <li>Lymphoma Research Trust</li> <li>MRC Clinical Trials Unit</li> <li>National Cancer Research Institute High Grade Lymphoma Subgroup</li> <li>National Institute for Health Research</li> <li>Public Health Wales</li> <li>UK Health Security Agency</li> </ul></li></ul>

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:

#### <u>Consultees</u>

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

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.