NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE

Single Technology Appraisal

Polatuzumab vedotin in combination for untreated diffuse large B-cell lymphoma [ID3901]

Final Stakeholder List

Consultees	Commentators (no right to submit or appeal)
 Company Roche (Polatuzumab vedotin) 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 Leukaemia Cancer Society Leukaemia CARE Lymphoma Action Macmillan Cancer Support Maggie's Centres Marie Curie South Asian Health Foundation 	 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 Medicines and Healthcare Products Regulatory Agency National Association of Primary Care National Pharmacy Association NHS Alliance NHS Confederation Scottish Medicines Consortium Welsh Health Specialised Services Committee
 Specialised Healthcare Alliance Tenovus Cancer Care WMUK 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 British Society for Haematology Cancer Research UK 	 Comparator companies Accord Healthcare Limited (doxorubicin) Accord UK Limited (prednisolone) ADVANZ Pharma (prednisolone) Allergan Ltd (prednisolone) Bausch & Lomb UK Ltd (prednisolone) Baxter Healthcare (cyclophosphamide, doxorubicin) Chemidex Pharma Ltd (prednisolone) Hospira UK Ltd (vincristine) Karo Pharma AB (prednisolone)

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Consultees Commentators (no right to submit or appeal) Royal College of General Practitioners Logixx Pharma Ltd (prednisolone) Royal College of Nursing Medac GmbH (doxorubicin) Royal College of Pathologists Napp Pharmaceuticals Limited (rituximab) Royal College of Physicians Royal College of Radiologists Pfizer Limited (doxorubicin, rituximab) Phoenix Labs (prednisolone) Royal Pharmaceutical Society Recipharm Pharmaceuticals AB Royal Society of Medicine Society of Radiographers (prednisolone) Roche Products Limited (rituximab) **UK Clinical Pharmacy Association** Sandoz Limited (cyclophosphamide, UK Oncology Nursing Society rituximab) UK Cutaneous Lymphoma Group **Seacross Pharmaceuticals** (doxorubicin) Others Wockhardt UK Ltd (prednisolone) Department of Health and Social Care Zentiva (prednisolone) NHS England NHS Lambeth CCG Relevant research groups NHS Wolverhampton CCG Cochrane Haematological Welsh Government Malignancies Group Cochrane UK Genomics England Institute of Cancer Research Leukaemia Busters Leukaemia UK Lymphoma Research Trust MRC Clinical Trials Unit 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

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Organisations that accept an invitation to participate in the appraisal; 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 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.

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