

NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE

Single Technology Appraisal

Belantamab mafodotin with pomalidomide and dexamethasone for treating relapsed or refractory multiple myeloma after 1 or more treatments ID6211

Provisional Stakeholder List

| Consultees | Commentators (no right to submit or appeal) |
|---|--|
| <p><u>Company</u></p> <ul style="list-style-type: none"> GlaxoSmithKline (belantamab mafodotin) <p><u>Patient/carer groups</u></p> <ul style="list-style-type: none"> 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 Karawa Leukaemia Trust Leukaemia Cancer Society Leukaemia Care Macmillan Cancer Support Maggie's Centres Marie Curie Myeloma UK South Asian Health Foundation Specialised Healthcare Alliance Tenovus Cancer Care <p><u>Healthcare professional groups</u></p> <ul style="list-style-type: none"> Association of Cancer Physicians British Blood Transfusion Society British Committee for Standards in Haematology British Geriatrics Society British Institute of Radiology British Oncology Pharmacy Association | <p><u>General</u></p> <ul style="list-style-type: none"> 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 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 <p><u>Possible comparator companies</u></p> <ul style="list-style-type: none"> AbbVie (dexamethasone) ADVANZ Pharma (dexamethasone, lenalidomide) Amarox (lenalidomide) Amgen (carfilzomib) AS Kalceks (dexamethasone) Aspen (dexamethasone) Aspire Pharma (bortezomib, dexamethasone) Aurobindo Pharma (bortezomib) Bausch & Lomb UK (dexamethasone) Biocon Pharma UK (lenalidomide) Bristol Myers Squibb Pharmaceuticals |

Provisional stakeholder list for evaluation of belantamab mafodotin with pomalidomide and dexamethasone for treating relapsed or refractory multiple myeloma after 1 or more treatments ID6211

Issue date: March 2024

| Consultees | Commentators (no right to submit or appeal) |
|--|---|
| <ul style="list-style-type: none"> • 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 • 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 Myeloma Society • UK Oncology Nursing Society <p><u>Others</u></p> <ul style="list-style-type: none"> • Department of Health and Social Care • NHS England | <ul style="list-style-type: none"> (lenalidomide, pomalidomide) • Dr. Reddy's Laboratories UK (bortezomib) • Glenmark Pharmaceuticals Europe (dexamethasone, lenalidomide) • Hameln pharma (dexamethasone) • Hospira UK (dexamethasone) • Janssen-Cilag (bortezomib, daratumumab) • Krka UK (dexamethasone) • Martindale Pharma, an Ethypharm Group Company (dexamethasone) • Medac GmbH (bortezomib) • Menarini Stemline UK (selinexor) • MSN Laboratories Europe (bortezomib) • Mylan (bortezomib, lenalidomide) • Novartis Pharmaceuticals UK (dexamethasone) • Panpharma UK (dexamethasone) • ParaPharm Development (dexamethasone) • Pfizer (bortezomib, elranatamab) • Piramal Critical Care (lenalidomide) • Ranbaxy UK Limited a Sun Pharmaceutical Company (bortezomib, lenalidomide) • Rayner Pharmaceuticals (dexamethasone) • Rosemont Pharmaceuticals (dexamethasone) • Sandoz (bortezomib, lenalidomide) • Sanofi (dexamethasone, isatuximab) • Santen UK (dexamethasone) • Synchrony Pharma (dexamethasone) • Takeda UK (ixazomib) • Teva UK (dexamethasone, lenalidomide) • Thame Laboratories (dexamethasone) • Thea Pharmaceuticals (dexamethasone) • Thornton & Ross (bortezomib, lenalidomide) • Tillomed Laboratories (bortezomib) • Wockhardt UK (dexamethasone) |

Provisional stakeholder list for evaluation of belantamab mafodotin with pomalidomide and dexamethasone for treating relapsed or refractory multiple myeloma after 1 or more treatments ID6211

Issue date: March 2024

| Consultees | Commentators (no right to submit or appeal) |
|------------|--|
| | <ul style="list-style-type: none"> • Zentiva (lenalidomide) • ZR Pharma & GmbH (panobinostat) <p><u>Relevant research groups</u></p> <ul style="list-style-type: none"> • Cochrane Haematological Malignancies Group • Cochrane UK • Genomics England • Institute of Cancer Research • Leukaemia Busters • Leukaemia UK • MRC Clinical Trials Unit • National Institute for Health Research <p><u>Associated Public Health groups</u></p> <ul style="list-style-type: none"> • 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

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 Provisional stakeholder list for evaluation of belantamab mafodotin with pomalidomide and dexamethasone for treating relapsed or refractory multiple myeloma after 1 or more treatments ID6211

Issue date: March 2024

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.