NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE

Single Technology Appraisal

Enfortumab vedotin for treating locally advanced or metastatic urothelial cancer after platinum-containing chemotherapy and a PD-1 or PD-L1 inhibitor ID3845

Final Stakeholder List

Consultees	Commentators (no right to submit or appeal)
 Company Astellas Pharma Ltd (enfortumab vedotin) Patient/carer groups Action Bladder Cancer UK Black Health Agency Bladder and Bowel UK Cancer 52 Cancer Black Care Cancer Equality Fight Bladder Cancer Helen Rollason Cancer Charity Independent Cancer Patients Voice Macmillan Cancer Support Maggie's Centres 	 General 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 Alliance
 Marie Curie Pelican Cancer Foundation South Asian Health Foundation Specialised Healthcare Alliance Tenovus Cancer Care 	 NHS Confederation Scottish Medicines Consortium Welsh Health Specialised Services Committee
 Professional groups Association of Cancer Physicians British Association of Urological Nurses British Association of Urological Surgeons British Geriatrics Society British Gynaecological Cancer Society British Institute of Radiology British Psychosocial Oncology Society British Society of Urogenital Radiology British Society of Urogynaecology British Uro-Oncology Group 	 Possible comparator companies Accord Healthcare (docetaxel, paclitaxel) Celgene (paclitaxel) Hospira UK (docetaxel, paclitaxel) Seacross Pharmaceuticals (docetaxel, paclitaxel) Teva UK (paclitaxel) Relevant research groups Cochrane Urology Genomics England Heart Research UK Institute of Cancer Research

Final stakeholder list for the technology appraisal of enfortumab vedotin for treating locally advanced or metastatic urothelial cancer after platinum-containing chemotherapy and a PD-1 or PD-L1 inhibitor ID3845 Issue date: April 2021

Consultees Commentators (no right to submit or appeal) MRC Clinical Trials Unit Cancer Research UK Royal College of General Practitioners National Cancer Research Institute Royal College of Nursing National Cancer Research Network Royal College of Pathologists National Institute for Health Research Royal College of Physicians Urothelial Cancers Research Group. Leeds Institute of Cancer & Pathology Royal College of Radiologists Royal Pharmaceutical Society Associated Public Health groups Royal Society of Medicine Public Health England Society and College of Radiographers **Public Health Wales UK Clinical Pharmacy Association UK Oncology Nursing Society Urology Foundation Others** Department of Health and Social Care NHS England NHS Sheffield CCG NHS Wokingham CCG Welsh Government

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.

PTO FOR DEFINITIONS OF CONSULTEES AND COMMENTATORS

Consultees

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 specialists and has the right to appeal against the Final Appraisal Document (FAD).

Final stakeholder list for the technology appraisal of enfortumab vedotin for treating locally advanced or metastatic urothelial cancer after platinum-containing chemotherapy and a PD-1 or PD-L1 inhibitor ID3845 Issue date: April 2021

All non-company consultees are invited to submit a statement¹, respond to consultations, nominate clinical specialists 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; other 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 specialists or patient experts.

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

Final stakeholder list for the technology appraisal of enfortumab vedotin for treating locally advanced or metastatic urothelial cancer after platinum-containing chemotherapy and a PD-1 or PD-L1 inhibitor ID3845 Issue date: April 2021

[©] National Institute for Health and Care Excellence 2021. All rights reserved