NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE

Proposed Single Technology Appraisal (STA)

Durvalumab with tremelimumab for untreated metastatic urothelial cancer ID1335

Provisional matrix of consultees and commentators

| Consultees | Commentators (no right to submit or |
|---|--|
| | appeal) |
| Company AstraZeneca (durvalumab with tremelimumab) Patient/carer groups Action Bladder Cancer UK Black Health Agency Cancer 52 Cancer Black Care Cancer Equality Fight Bladder Cancer HAWC Helen Rollason Cancer Charity Independent Cancer Patients Voice Macmillan Cancer Support Maggie's Centres Marie Curie Muslim Council of Britain | 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 NHS Confederation Scottish Medicines Consortium |
| Pelican Cancer Foundation South Asian Health Foundation Specialised Healthcare Alliance | Welsh Health Specialised Services Committee |
| Tenovus Cancer Care | Possible comparator companies Accord Healthcare (carboplatin, |
| Professional groupsAssociation of Anaesthetists | cisplatin, doxorubicin, gemcitabine, methotrexate Tenovus Cancer Care |
| Association of Cancer PhysiciansAssociation of Surgeons of Great Britain and Ireland | Allergan (gemcitabine) Amgen (pegfilgrastim) Chugai Pharma (lenograstim) |
| 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 | Concordia International (methotrexate) Hospira UK (carboplatin, cisplatin, gemcitabine, methotrexate, vinblastine) Janssen-Cilag (doxorubicin) Lilly UK (gemcitabine) Medac GmbH (doxorubicin, gemcitabine) Nordic Pharma (methotrexate) |

Consultees Commentators (no right to submit or appeal) British Society of Urogenital Radiology Orion Pharma (methotrexate) British Society of Urogynaecology Pfizer (doxorubicin, methotrexate, British Uro-Oncology Group vinblastine) Rosemont Pharmaceuticals Cancer Research UK Royal College of Anaesthetists (methotrexate) Sandoz (cisplatin, methotrexate) Royal College of General Practitioners Seacross Pharmaceuticals Royal College of Nursing (doxorubicin) Royal College of Pathologists Sun Pharma (carboplatin, gemcitabine) Royal College of Physicians Teva UK (doxorubicin, lipegfilgrastim, Royal College of Radiologists methotrexate) Royal College of Surgeons Therakind (methotrexate) Royal Pharmaceutical Society Royal Society of Medicine Relevant research groups Society and College of Radiographers Cochrane Prostate Diseases and **UK Clinical Pharmacy Association Urologic Cancers Group UK Health Forum** Cochrane Urology **UK Oncology Nursing Society** Genomics England **Urology Foundation** Institute of Cancer Research MRC Clinical Trials Unit **Others** National Cancer Research Institute Department of Health and Social Care National Cancer Research Network NHS England National Institute for Health Research NHS North Kirklees CCG Urothelial Cancers Research Group, NHS Oldham CCG Leeds Institute of Cancer & Pathology Welsh Government Associated Public Health Groups Public Health England **Public Health Wales**

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 lists in the matrix, and which organisations we should include that have a particular focus on relevant equality issues.

PTO FOR DEFINITIONS OF CONSULTEES AND COMMENTATORS

Definitions:

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).

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.

Provisional matrix for the proposed technology appraisal of durvalumab with tremelimumab for untreated metastatic urothelial cancer ID1335. Issue date: October 2018

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