NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE

Single Technology Appraisal

Lutetium-177 vipivotide tetraxetan in combination for treating PSMA-positive hormone-sensitive metastatic prostate cancer ID6589

Provisional Stakeholder List

Provisional Consultees	Provisional Commentators (no right to submit or appeal)
Company • Novartis Pharmaceuticals UK (Lutetium-177 vipivotide tetraxetan) Patient/carer groups • Black Health Agency for Equality • Bob Champion Cancer Trust • Cancer Black Care • Independent Cancer Patients Voice • Macmillan Cancer Support • Maggie's Centres • Marie Curie • Orchid • PCaSO - Prostate Cancer Support Organisation • Prost8 UK • Prostate Cancer Research • Prostate Cancer UK • Prostate Matters UK • South Asian Health Foundation • Specialised Healthcare Alliance • Tackle Prostate Cancer • Tenovus Cancer Care Healthcare professional groups • Association of Anaesthetists • Association of Cancer Physicians • Association of Surgeons of Great Britain and Ireland • British Association of Urological Nurses • British Geriatrics Society • British Institute of Radiology • British Oncology Pharmacy Association • British Psychosocial Oncology Society	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 - Northern Ireland • Healthcare Improvement Scotland • Medicines and Healthcare products Regulatory Agency • National Association of Primary Care • National Pharmacy Association • NHS Confederation • NHS Wales Joint Commissioning Committee • Scottish Medicines Consortium • Welsh Government Possible comparator companies • Accord (relugolix) • ADVANZ Pharma (cyproterone) • Amarox (abiraterone) • Aspire Pharma (leuprorelin) • AstraZeneca UK (bicalutamide, goserelin) • Astellas Pharma (enzalutamide) • Axunio Pharma GmBh (abiraterone) • Bayer (darolutamide) • Celix Pharma (abiraterone) • Cipla EU (abiraterone) • Genus Pharmaceuticals (abiraterone) • Glenmark Pharmaceuticals Europe (abiraterone) • Hospira (docetaxel)

Provisional Consultees	Provisional Commentators (no right to submit or appeal)
• British Uro-Oncology Group • Cancer Research UK • Prostate Cancer Advisory group • Royal College of Anaesthetists • Royal College of General Practitioners • Royal College of Nursing • Royal College of Pathologists • Royal College of Physicians • Royal College of Radiologists • Royal College of Surgeons • Royal Pharmaceutical Society • Royal Society of Medicine • Society and College of Radiographers • UK Clinical Pharmacy Association • UK Oncology Nursing Society • Urology Foundation <u>Others</u> • Department of Health and Social Care • NHS England	• Ferring Pharmaceuticals (triptorelin) • Ipsen (triptorelin) • Janssen-Cilag (abiraterone, apalutamide) • Krka UK (abiraterone) • Morningside Healthcare (bicalutamide) • Mylan (flutamide) • Neon Healthcare (buserelin) • Ranbaxy UK Limited a Sun Pharmaceutical company (abiraterone, bicalutamide) • Sandoz (abiraterone) • Seacross (docetaxel) • Sovereign Medical (flutamide) • Takeda UK (leuprorelin) • Wockhardt UK (abiraterone) • Zentiva (abiraterone) <u>Relevant research groups</u> • Cochrane Urology • Genomics England • Institute of Cancer Research • MRC Clinical Trials Unit • National Institute for Health Research • Pro Cancer Research Fund <u>Associated Public Health groups</u> • 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:

Consultee or commentator stakeholders are provisional until a signed Confidentiality Agreement & Undertaking form is submitted to NICE at the evaluation stage. Participating stakeholders will be listed on the project information page for the evaluation.

<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.

