NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE

Single Technology Appraisal

Relugolix for treating hormone-sensitive prostate cancer ID6187

Provisional Stakeholder List

Consultees	Commentators (no right to submit or appeal)
Company Accord Healthcare Limited and Myovant Sciences (relugolix) Patient/carer groups • Black Health Agency for Equality • Bob Champion Cancer Trust • Cancer Black Care • Cancer Equality • Helen Rollason Cancer Charity • Independent Cancer Patients Voice • Macmillan Cancer Support • Maggie's Centres • Marie Curie • Orchid • PCaSO - Prostate Cancer Support Organisation • Pelican Cancer Foundation • Prost8 UK • Prostate Cancer 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 Association of Urological Surgeons • British Geriatrics Society • British Institute of Radiology • British Oncology Pharmacy Association	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 Confederation • Scottish Medicines Consortium • Welsh Government • Welsh Health Specialised Services Committee Possible comparator companies • Accord (bicalutamide, docetaxel) • Astellas Pharma (enzalutamide) • AstraZeneca (bicalutamide, goserelin) • Bayer (cyproterone acetate, darolutamide) • Ferring Pharmaceuticals (degarelix, triptorelin) • Hospira UK (docetaxel) • Ipsen (triptorelin) • Janssen-Cilag (apalutamide) • Kent Pharma (cyproterone acetate) • Mylan (flutamide) • Neon Healthcare (buserelin) • Seacross (docetaxel) • Sovereign Medical (flutamide) • Sun Pharma (bicalutamide)

Consultees	**Commentators (no right to submit or appeal)**
• British Psychosocial Oncology Society • 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	• Takeda (leuprorelin) • Typharm (leuprorelin) <u>Relevant research groups</u> • Cochrane UK • Cochrane Urology • Genomics England • Institute of Cancer Research • MRC Clinical Trials Unit • National Institute for Health Research • Pro Cancer Research Fund • Prostate Cancer Research Centre <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.

<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 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 Draft Guidance (FDG).

All non-company consultees are invited to submit a statement[1], respond to consultations, nominate clinical or patient experts and have the right to appeal against the Final Draft Guidance (FDG).

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

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], 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.

