NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE

Single Technology Appraisal

Elacestrant for treating oestrogen receptor-positive HER2-negative advanced breast cancer with an ESR1 mutation after at least 1 endocrine treatment [ID6225]

Final Stakeholder List

Consultees	Commentators (no right to submit or
	appeal)
Company Menarini Stemline (elacestrant) Patient/carer groups Black Health Agency for Equality Breast Cancer Now Breast Cancer UK Cancer Black Care Cancer Equality	 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
 Helen Rollason Cancer Charity Hereditary Breast Cancer Helpline Independent Cancer Patients Voice Macmillan Cancer Support Maggie's Centres Make 2nds Count Marie Curie MET UP UK South Asian Health Foundation Specialised Healthcare Alliance Tenovus Cancer Care 	 Department of Fleatiff, 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 Welsh Government Welsh Health Specialised Services Committee
 Healthcare professional groups Association of Breast Surgery Association of Cancer Physicians British Geriatrics Society British Institute of Radiology British Oncology Pharmacy	 Comparator companies Amarox (anastrozole, capecitabine, letrozole) AstraZeneca (anastrozole, fulvestrant) Aurobindo Pharma – Milpharm (tamoxifen) Bristol Myers Squibb Pharmaceuticals (paclitaxel) Cipla EU (fulvestrant, letrozole) Dr. Reddy's Laboratories (capecitabine, everolimus, fulvestrant) Eisai (eribulin) Ethypharm (everolimus) Genus Pharmaceuticals (fulvestrant) Glenmark Pharmaceuticals (capecitabine, exemestane, fulvestrant,

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Consultees Commentators (no right to submit or appeal) letrozole) Royal Pharmaceutical Society Hospira UK (docetaxel, gemcitabine, Royal Society of Medicine paclitaxel) Society and College of Radiographers Medac GmbH (vinorelbine) UK Breast Cancer Group Medical Valley (fulvestrant) **UK Clinical Pharmacy Association** Mylan (anastrozole, exemestane, UK Oncology Nursing Society tamoxifen) Novartis (alpelisib, everolimus, letrozole) Department of Health and Social Care Pfizer (exemestane) NHS England • Pierre Fabre (vinorelbine) Ranbaxy, a Sun Pharmaceuticals company (anastrozole, fulvestrant, gemcitabine, letrozole) Rivopharm (exemestane) • Rosemont Pharmaceuticals (tamoxifen) • Sandoz (anastrozole, everolimus, fulvestrant) Seacross Pharmaceuticals (docetaxel, paclitaxel) Synchrony Pharma (gemcitabine) • Teva UK (fulvestrant, paclitaxel) Tillomed Laboratories (tamoxifen) Wockhardt UK (tamoxifen) • Zentiva (exemestane, fulvestrant) Relevant research groups Against Breast Cancer Breast Cancer Hope Cochrane Breast Cancer Group Cochrane UK Genomics England Institute of Cancer Research MRC Clinical Trials Unit

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.

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National Institute for Health Research

Pro-Cancer Research Fund

Associated Public Health groups

UK Health Security Agency

Public Health Wales

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 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 statement¹, 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], 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.

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