NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE

Single Technology Appraisal

Olaparib for maintenance treatment of BRCA-mutated ovarian, fallopian tube and peritoneal cancer after response to first-line platinum-based chemotherapy ID6191 (Review of TA598)

Provisional Stakeholder List

Consultees	Commentators (no right to submit or appeal)
Company	General
AstraZeneca (olaparib)	Allied Health Professionals Federation
	 All Wales Therapeutics and Toxicology
Patient/carer groups	Centre
Black Health Agency	 Board of Community Health Councils in
Cancer Black Care	Wales
Cancer Equality	 British National Formulary
Cancer 52	 Care Quality Commission
Eve Appeal	 Department of Health, Social Services
Go Girls	and Public Safety for Northern Ireland
Helen Rollason Cancer Charity	Healthcare Improvement Scotland
Independent Cancer Patients Voice	Medicines and Healthcare products
Macmillan Cancer Support	Regulatory Agency
Maggie's Centres	National Association of Primary Care
Marie Curie	National Pharmacy Association
Ovacome	NHS Alliance
Ovarian Cancer Action	NHS Confederation
Robin Cancer Trust	Scottish Medicines Consortium
South Asian Health Foundation	Welsh Health Specialised Services
Specialised Healthcare Alliance	Committee
Target Ovarian Cancer	D 11.
Tenovus Cancer Care	Possible comparator companies
Wellbeing of Women	Celltrion Healthcare UK (bevacizumab)
Women's Health Concern	Organon Pharma UK (bevacizumab)
	Pfizer (bevacizumab)
Professional groups	Roche (bevacizumab)
Association of Cancer Physicians	Thornton & Ross (bevacizumab)
British Geriatrics Society	 Zentiva (bevacizumab)
British Gynaecological Cancer Society	Delevent need and
British Institute of Radiology	Relevant research groups
British Psychosocial Oncology Society	Cochrane Gynaecological Cancer Craver
British Society of Urogynaecology	Group
Cancer Research UK	Cochrane UK

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Provisional stakeholder list for the evaluation of olaparib for maintenance treatment of BRCA-mutated ovarian, fallopian tube and peritoneal cancer after response to first-line platinum-based chemotherapy ID6191 (Review of TA598) Issue date: May 2023

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Consultees	Commentators (no right to submit or appeal)
 National Forum of Gynaecological Oncology Nurses Royal College of General Practitioners Royal College of Nursing Royal College of Obstetricians & Gynaecologists Royal College of Pathologists Royal College of Physicians Royal College of Radiologists Royal Pharmaceutical Society Royal Society of Medicine Society and College of Radiographers UK Clinical Pharmacy Association UK Oncology Nursing Society 	 Genomics England Institute of Cancer Research MRC Clinical Trials Unit National Cancer Research Institute National Institute for Health Research Ovarian & Prostate Cancer Research Trust Associated Public Health Groups Public Health Wales UK Health Security Agency
 Department of Health and Social Care NHS England 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 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

Provisional stakeholder list for the evaluation of olaparib for maintenance treatment of BRCA-mutated ovarian, fallopian tube and peritoneal cancer after response to first-line platinum-based chemotherapy ID6191 (Review of TA598) Issue date: May 2023

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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.

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