NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE

Single Technology Appraisal

Niraparib for maintenance treatment of advanced ovarian, fallopian tube and peritoneal cancer after response to first-line platinum-based chemotherapy [Review of TA673] (ID6403)

Final stakeholder List

Consultees	Commentators (no right to submit or appeal)
Company	General
GlaxoSmithKline (niraparib)	All Wales Therapeutics and Toxicology Centre
Patient/carer groups	Allied Health Professionals Federation
Black Health Agency for Equality	Board of Community Health Councils in
Cancer 52	Wales
Cancer Black Care	British National Formulary
Eve Appeal	Care Quality Commission
Go Girls	Department of Health, Social Services
Helen Rollason Cancer Charity	and Public Safety for Northern Ireland
Independent Cancer Patients Voice	Healthcare Improvement Scotland
Macmillan Cancer Support	Medicines and Healthcare products
Maggie's Centres	Regulatory Agency
Marie Curie	National Association of Primary Care
Ovacome	National Pharmacy Association
Ovarian Cancer Action	NHS Confederation
Robin Cancer Trust	Scottish Medicines Consortium
South Asian Health Foundation	Welsh Government
Specialised Healthcare Alliance	Welsh Health Specialised Services
Target Ovarian Cancer	Committee
Tenovus Cancer Care	
Wellbeing of Women	Possible comparator companies
Women's Health Concern	AstraZeneca (olaparib)
	Celltrion Healthcare (bevacizumab)
Healthcare professional groups	Clovis Oncology (rucaparib)
Association of Cancer Physicians	Dr. Reddy's Laboratories (bases since a b.)
British Geriatrics Society	(bevacizumab)
British Gynaecological Cancer Society	Organon Pharma (bevacizumab)
British Institute of Radiology	Pfizer (bevacizumab) Pecha (bevacizumab)
British Psychosocial Oncology Society	Roche (bevacizumab) Thernton & Roce (bevacizumab)
British Society of Urogynaecology	Thornton & Ross (bevacizumab) Zontive (bevacizumab)
Cancer Research UK	Zentiva (bevacizumab) Dharmand CmbH (rusanarib)
National Forum of Gynaecological	Pharmaand GmbH (rucaparib)

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Issue date: May 2025

Consultees	Commentators (no right to submit or appeal)
 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 Others Department of Health and Social Care NHS England 	Relevant research groups Cochrane Gynaecological Cancer Group Cochrane UK Genomics England Institute of Cancer Research MRC Clinical Trials Unit National Cancer Research Institute National Institute for Health Research Associated Public Health groups 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.

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 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).

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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.