

NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE

Single Technology Appraisal

Ribociclib with an aromatase inhibitor for adjuvant treatment of hormone receptor-positive, HER2-negative early breast cancer [ID6153]

Final Stakeholder List

Provisional consultees	Provisional commentators (no right to submit or appeal)
<p><u>Company</u></p> <ul style="list-style-type: none"> • Novartis (ribociclib) <p><u>Patient/carer groups</u></p> <ul style="list-style-type: none"> • Black Health Agency for Equality • Breast Cancer Now • Breast Cancer UK • Cancer Black Care • Cancer Equality • Helen Rollason Cancer Charity • Hereditary Breast Cancer Helpline • Independent Cancer Patients Voice • Inflammatory Breast Cancer Network UK • Lobular Breast Cancer UK • Macmillan Cancer Support • Maggie’s Centres • Marie Curie • MET UP UK • South Asian Health Foundation • Specialised Healthcare Alliance • Tenovus Cancer Care • UK Charity for Triple Negative Breast Cancer <p><u>Healthcare professional groups</u></p> <ul style="list-style-type: none"> • Association of Breast Surgery • Association of Cancer Physicians • British Geriatrics Society • British Institute of Radiology • British Oncology Pharmacy Association • British Psychosocial Oncology Society • Cancer Research UK • Royal College of General Practitioners • Royal College of Nursing • Royal College of Pathologists 	<p><u>General</u></p> <ul style="list-style-type: none"> • 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 Alliance • NHS Confederation • Scottish Medicines Consortium • Welsh Government • Welsh Health Specialised Services Committee <p><u>Comparator companies</u></p> <ul style="list-style-type: none"> • Eli Lilly (abemaciclib) • AstraZeneca (olaparib) <p><u>Relevant research groups</u></p> <ul style="list-style-type: none"> • Against Breast Cancer • Breast Cancer Hope • Cochrane Breast Cancer Group • Cochrane UK • Genomics England • Institute of Cancer Research • MRC Clinical Trials Unit • National Institute for Health Research • Pro-Cancer Research Fund

Final stakeholder list for evaluation of ribociclib with an aromatase inhibitor for adjuvant treatment of hormone receptor-positive, HER2-negative early breast cancer [ID6153]

Issue date: August 2024

© National Institute for Health and Care Excellence 2024. All rights reserved.

Page 1 of 3

Provisional consultees	Provisional commentators (no right to submit or appeal)
<ul style="list-style-type: none"> • Royal College of Physicians • Royal College of Radiologists • Royal Pharmaceutical Society • Royal Society of Medicine • Society and College of Radiographers • UK Breast Cancer Group • UK Clinical Pharmacy Association • UK Oncology Nursing Society <p><u>Others</u></p> <ul style="list-style-type: none"> • Department of Health and Social Care • NHS England 	<p><u>Associated Public Health groups</u></p> <ul style="list-style-type: none"> • 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).

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.