NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE

Single Technology Appraisal

Effornithine for treating high-risk neuroblastoma with complete or partial response after immunotherapy [ID4060]

Final Stakeholder List

Consultees	Commentators (no right to submit or appeal)
 Company Norgine (eflornithine) Patient/carer groups Black Health Agency for Equality 	 General All Wales Therapeutics and Toxicology Centre Allied Health Professionals Federation
 Cancer Black Care Cancer Equality Cancer52 Childhood Cancer Parents Alliance Children with Cancer UK Children's Cancer and Leukaemia 	 Board of Community Health Councils in Wales British National Formulary Care Quality Commission Department of Health - Northern Ireland
 Group Contact Gene People Genetic Alliance UK Helen Rollason Cancer Charity Independent Cancer Patients Voice Macmillan Cancer Support Maggie's Centres Marie Curie Mitchell's Miracles Neuroblastoma UK Solving Kids' Cancer UK South Asian Health Foundation 	 Healthcare Improvement Scotland Hospital Information Services – Jehovah's Witnesses Medicines and Healthcare products Regulatory Agency National Association for Primary Care National Pharmacy Association NHS Alliance NHS Confederation Scottish Medicines Consortium Welsh Government Welsh Health Specialised Services Committee
 Specialised Healthcare Alliance Teenage Cancer Trust Tenovus Cancer Care Together for Shorter Lives 	Possible comparator companies None
 Young Lives vs Cancer Healthcare professional groups Association of Cancer Physicians British Institute of Radiology British Paediatric Neurology Association British Psychosocial Oncology Society 	 Relevant research groups Cochrane Childhood Cancer Group Cochrane UK Genomics England Institute of Cancer Research MRC Clinical Trials Unit National Institute for Health Research

Provisional stakeholder list for the single technology appraisal of eflornithine for treating highrisk neuroblastoma with complete or partial response after immunotherapy [ID4060] Issue date: September 2024

Consultees Commentators (no right to submit or appeal) Cancer Research UK Associated Public Health Groups Children's Cancer and Leukaemia Public Health Wales Group UK Health Security Agency Neonatal & Paediatric Pharmacists Group Royal College of General Practitioners Royal College of Nursing Royal College of Paediatrics and Child Health 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** Others Department of Health and Social Care **NHS** England

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 relevant NHS organisations in England.

Provisional stakeholder list for the single technology appraisal of eflornithine for treating highrisk neuroblastoma with complete or partial response after immunotherapy [ID4060] Issue date: September 2024 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.