NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE

Single Technology Appraisal

Exagamglogene autotemcel for treating sickle cell disease [ID4016]

Final Stakeholder List

Consultees	Commentators (no right to submit or appeal)
 <u>Company</u> Vertex (Exagamglogene autotemcel) <u>Patient/carer groups</u> Anthony Nolan AOFAC Foundation Black Health Agency for Equality Caribbean and African Health Network Cianna's Smile Contact Diamond Blackfan Anaemia UK Support Group (BDA) Gene People Genetic Alliance UK Pain UK Sickle Cell and Young Stroke Survivors 	 appeal) <u>General</u> All Wales Therapeutics and Toxicology Centre Allied Health Professionals Federation Board of Community Health Councils in Wales British National Formulary Care Quality Commission Cell and Gene Therapy Catapult Department of Health, Social Services and Public Safety for Northern Ireland Healthcare Improvement Scotland Hospital Information Services – Jehovah's Witnesses Medicines and Healthcare products Regulatory Agency National Association of Primary Care
 Sickle Cell Society SickleKan South Asian Health Foundation Specialised Healthcare Alliance The Essenelle Foundation United Kingdom Thalassaemia	 National Pharmacy Association NHS Confederation Scottish Medicines Consortium Welsh Government Welsh Health Specialised Services
Society	Committee
 Healthcare professional groups Anaemia Nurse Specialist Association Association of Genetic Nurses &	 Masters Specialty Pharma
Counsellors British Blood Transfusion Society British Committee for Standards in	(hydroxycarbamide) Medac GmbH (hydroxycarbamide) Neon Laboratories (hydroxycarbamide) Nova Laboratories (hydroxycarbamide) Nova Laboratories (hydroxycarbamide) Relevant research groups Cochrane Cystic Fibrosis & Genetic
Haematology British Geriatrics Society British Society for Gene and Cell	Disorders Group Cochrane Haematological Malignancies
Therapy British Society for Genetic Medicine British Society for Haematology	Group Cochrane UK Genomics England

Final stakeholder list for the evaluation of exagamglogene autotemcel for treating sickle cell disease ID4016 Issue date: June 2023

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Consultees	Commentators (no right to submit or appeal)
 British Society of Bone Marrow Transplantation and Cellular Therapy Immunodeficiency UK Neonatal and Paediatric Pharmacists Group NHS Blood and Transplant NHS Genomic Medicine Service Alliance Neonatal Paediatric Pharmacist Group Royal College of General Practitioners Royal College of Paediatrics and Child Health Royal College of Pathologists Royal College of Physicians Royal College of Physicians Royal College of Physicians Royal College of Medicine Sickle cell and thalassaemia nurses, midwives and allied health care professionals (STANMAP) UK Clinical Pharmacy Association UK Forum on Haemoglobin Disorders UK National Haemoglobinopathy Panel UK National Screening Committee Others Department of Health and Social Care NHS England 	 MRC Clinical Trials Unit National Institute for Health Research <u>Associated Public Health groups</u> 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:

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.