## NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE

## **Single Technology Appraisal**

# Belumosudil for treating chronic graft versus host disease after 2 or more therapies ID4021

## **Provisional Stakeholder List**

Consultees	Commentators (no right to submit or appeal)
	appoul
Company	General
Sanofi (belumosudil)	All Wales Inherited Metabolic Disease
	Service Cardiff
Patient/carer groups	All Wales Therapeutics and Toxicology
African-Caribbean Leukaemia Trust	Centre
Anthony Nolan	Allied Health Professionals Federation
Aplastic Anaemia Trust	Board of Community Health Councils in
Blood Cancer UK	Wales
Cianna's Smile	British National Formulary
CML Support Group	Care Quality Commission
• DKMS	Department of Health, Social Services
Immunodeficiency UK	and Public Safety for Northern Ireland
Leukaemia Cancer Society	Healthcare Improvement Scotland
Leukaemia Care	Medicines and Healthcare products
Lymphoma Action	Regulatory Agency
Macmillian Cancer Support	National Association of Primary Care
MDS UK Patient Support Group	<ul> <li>National Pharmacy Association</li> </ul>
MPS Society	National Services Division
Myeloma UK	NHS Confederation
Sickle Cell Society	Scottish Medicines Consortium
Sickle Cell and Young Stroke	Welsh Government
Survivors	Welsh Health Specialised Services
SickleKan	Committee
<ul> <li>South Asian Health Foundation</li> </ul>	
Specialised Healthcare Alliance	Possible comparator companies
The Essenelle Foundation	Accord Healthcare (methotrexate,
UK Thalassaemia Society	mycophenolate mofetil)
	ADVANZ Pharma (methotrexate)  Ciple FIL (methotrexate)
Healthcare professional groups	Cipla EU (methotrexate)
British Blood Transfusion Society	Hospira (methotrexate)
British Committee for Standards in	Medac (methotrexate)     Marningside Healtheare (methotrexate)
Haematology	Morningside Healthcare (methotrexate)     Nordia Pharma (methotrexate)
British Geriatrics Society	Nordic Pharma (methotrexate)
British Society for Genetic Medicine	Novartis Pharmaceuticals UK     (mycenhanalete mefetil)
British Society for Haematology	(mycophenolate mofetil)

Provisional stakeholder list for the evaluation of belumosudil for treating chronic graft versus host disease after 2 or more therapies ID4021

Issue date: September 2022

#### Consultees Commentators (no right to submit or appeal) Orion Pharma UK (methotrexate) British Society for Immunology British Society of Blood and Marrow Pfizer (methotrexate) Transplantation and Cellular Therapy Roche Products (mycophenolate **British Transplantation Society** mofetil) Cancer Research UK Rosemont Pharmaceutical Neonatal and Paediatric Pharmacists (methotrexate) Sandoz Limited (methotrexate, Group mycophenolate mofetil) NHS Blood and Transplant Royal College of General Practitioners Teva Pharma (mycophenolate mofetil) Therakind (methotrexate) Royal College of Nursing Tillomed Laboratories (mycophenolate Royal College of Paediatrics and Child mofetil) Health Royal College of Pathologists Relevant research groups Royal College of Physicians Cochrane Haematological Malignancies Royal Pharmaceutical Society Group Royal Society of Medicine Cochrane UK **UK Clinical Pharmacy Association** Genomics England UK Forum on Haemoglobin Disorders Leukaemia Busters **UK Primary Immunodeficiency** MRC Clinical Trials Unit Network National Institute for Health Research Others Oxford Biomedical Research Centre Department of Health and Social Care Associated Public Health groups • Christie Haematology and Transplant **Public Health Wales** Unit (Manchester) UK Health Security Agency • King's College London - Haematooncology department NHS England St James University Hospital Bone Marrow Transplant Department (Leeds Teaching Hospitals Trust)

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.

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**HSCT Service** 

**University College London Hospitals** 

### **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 statement<sup>1</sup>, 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.

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<sup>&</sup>lt;sup>1</sup> Non-company consultees are invited to submit statements relevant to the group they are representing.