NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE

Single Technology Appraisal

Mirikizumab for treating moderately to severely active ulcerative colitis ID3973

Provisional Stakeholder List

Consultees	Commentators (no right to submit or appeal)
<u>Company</u>	General
Eli Lilly (mirikizumab)	All Wales Therapeutics and Toxicology
	Centre
Patient/carer groups	Allied Health Professionals Federation
Bladder and Bowel Community	Board of Community Health Councils in
Bowel Cancer UK	Wales
CICRA	British National Formulary
Colostomy UK Crabba's and Califia LIK	Care Quality Commission
 Crohn's and Colitis UK GUTS UK 	Department of Health, Social Services and Public Safety for Northern Iroland
	and Public Safety for Northern IrelandHealthcare Improvement Scotland
	 Medicines and Healthcare Products
Specialised Healthcare Alliance	Regulatory Agency
Healthcare professional groups	 National Association of Primary Care
 Association of Coloproctology for 	 National Pharmacy Association
Great Britain and Ireland	NHS Alliance
British Geriatrics Society	NHS Confederation
British Society of Gastroenterology	Scottish Medicines Consortium
Primary Care Society for	 Scottish Society of Gastroenterology
Gastroenterology	Welsh Health Specialised Services
Royal College of General Practitioners	Committee
Royal College of Nursing	
Royal College of Pathologists	 Possible comparator companies
Royal College of Physicians	AbbVie (adalimumab, upadacitinib)
Royal Society of Medicine	Amgen (adalimumab)
Royal Pharmaceutical Society	BioGen Biosimilars (infliximab,
UK Clinical Pharmacy Association	adalimumab)
Othoro	Bristol Myers-Squibb (ozanimod)
Others	 Celltrion Healthcare (infliximab, adalimumab)
Department of Health and Social CareNHS England	adalimumab) • Fresenius Kabi (adalimumab)
 NHS England NHS Leeds CCG 	 Fresenius Kabi (adalimumab) Galapagos BioTech (filgotinib)
 NHS Leeds CCG NHS Telford & Wrekin CCG 	 Janssen-Cilag (ustekinumab)
 Welsh Government 	 Merck Sharp & Dohme (infliximab,
	golimumab)
	 Pfizer (infliximab, tofacitinib)
	 Sandoz (infliximab, adalimumab)

Provisional stakeholder list for the evaluation of mirikizumab for treating moderately to severely active ulcerative colitis ID3973 Issue date: June 2022

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Consultees	Commentators (no right to submit or appeal)
	Takeda (vedolizumab)
	 <u>Relevant research groups</u> Cochrane Inflammatory Bowel Disease and Functional Bowel Disorders Group Cochrane UK Genomics 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

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¹ Non company consultees are invited to submit statements relevant to the group they are representing.

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.