NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE

Single Technology Appraisal

Bimekizumab for treating moderate to severe hidradenitis suppurativa ID6134

Final Stakeholder List

Consultees	Commentators (no right to submit or appeal)
Company	General
UCB Pharma (bimekizumab) Patient/carer groups Action on Pain	 All Wales Therapeutics and Toxicology Centre Allied Health Professionals Federation Board of Community Health Councils in
 African Health Policy Network Black Health Agency for Equality Changing Faces Crohn's and Colitis UK Hidradenitis Suppurativa Trust HS Support Network Pain Concern Pain Relief Foundation Pain UK South Asian Health Foundation Specialised Healthcare Alliance The Hidradenitis Suppurativa Trust Healthcare professional groups 	 Wales British National Formulary Care Quality Commission Department of Health, Social Services and Public Safety for Northern Ireland Healthcare Improvement Scotland Medicines and Healthcare products Regulatory Agency National Association of Primary Care National Pharmacy Association NHS Confederation Scottish Medicines Consortium Welsh Government Welsh Health Specialised Services
 British Association of Dermatologists British Dermatological Nursing Group 	Committee
British Geriatrics Society	Possible comparator companies
Primary Care Dermatology Society	AbbVie (adalimumab)
 Royal College of General Practitioners 	Amgen (adalimumab)
Royal College of Nursing	Biogen (adalimumab)
 Royal College of Pathologists Royal College of Physicians Royal Pharmaceutical Society Royal Society of Medicine 	 Celltrion Healthcare UK (adalimumab) Fresenius Kabi (adalimumab) Novartis (secukinumab) Sandoz (adalimumab)
St John's Institute of DermatologyUK Clinical Pharmacy Association	Relevant research groups British Skin Foundation
Others Department of Health and Social Care NHS England	 Centre of Evidence-based Dermatology, University of Nottingham Cochrane Skin Group Cochrane UK
	Dermatrust

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Issue date: February 2024

Consultees	Commentators (no right to submit or appeal)
	 Genomics England MRC Clinical Trials Unit National Institute for Health Research
	 Associated Public Health groups 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.

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