

NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE

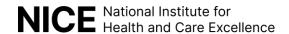
Single Technology Appraisal

Guselkumab for treating moderately to severely active ulcerative colitis ID6237

Provisional Stakeholder List

Consultees	Commentators (no right to submit or appeal)
Company	General
Janssen (guselkumab)	All Wales Therapeutics and Toxicology
 Patient/carer groups Bladder and Bowel Community Bowel Cancer UK Colostomy UK Crohn's and Colitis UK GUTS UK IA: Ileostomy and Internal Pouch Association South Asian Health Foundation Specialised Healthcare Alliance Healthcare professional groups Association of Coloproctology for 	 Centre Allied Health Professionals Federation Board of Community Health Councils in 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
 Great Britain and Ireland British Geriatrics Society British Society of Gastroenterology Primary Care Society for Gastroenterology Royal College of General Practitioners 	 Scottish Medicines Consortium Scottish Society of Gastroenterology Welsh Government Welsh Health Specialised Services Committee
 Royal College of Nursing Royal College of Pathologists Royal College of Physicians Royal Pharmaceutical Society Royal Society of Medicine UK Clinical Pharmacy Association 	 Possible comparator companies AbbVie (adalimumab, risankizumab, upadacitinib) Amgen (adalimumab), Biogen Biosimilars (adalimumab, infliximab), Bristol-Myers Squibb Pharmaceuticals
Others Department of Health and Social Care NHS England	 (ozanimod) Celltrion Healthcare UK (adalimumab, infliximab) Eli Lilly (mirikisumab) Fresenius, Kabi (adalimumab) Galapagos Biotech (filgotinib) Janssen-Cilag (ustekinumab)

Provisional stakeholder list for the evaluation of guselkumab for previously treated moderately to severely active ulcerative colitis ID6237



Consultees	Commentators (no right to submit or appeal)
	 Merck, Sharp and Dohme, UK (golimumab, infliximab) Pfizer (etrasimod, infliximab, tofacitinib) Sandoz (adalimumab, infliximab) Takeda UK (vedolizumab)
	 Relevant research groups Cochrane Inflammatory Bowel Disease and Functional Bowel Disorders Group 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 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 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).

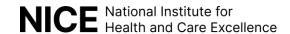
<u>Commentators</u>

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

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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.