NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE

Single Technology Appraisal

Upadacitinib for treating moderately to severely active ulcerative colitis ID3953

Provisional Stakeholder List

Consultees	Commentators (no right to submit or appeal)
Company	General
AbbVie (upadacitinib)	All Wales Therapeutics and Toxicology
Royal Society of Medicine	Amgen (adalimumab) Picture Richard (adalimumab)
 Royal Pharmaceutical Society Neonatal & Paediatric Pharmacist Group UK Clinical Pharmacy Association 	 Biogen Biosimilars (adalimumab, infliximab) Celltrion Healthcare UK (adalimumab, infliximab) Fresenius Kabi (adalimumab)
Others	Gilead Sciences (filgotinib) Gilead Sciences (filgotinib)
Department of Health and Social Care NUS England	Janssen-Cilag (ustekinumab) Morek Sharp & Dohmo (golimumab)
NHS EnglandNHS Northern, Eastern and Western	 Merck Sharp & Dohme (golimumab, infliximab)
Devon CCG	Pfizer (infliximab, tofacitinib)
NHS Windsor, Ascot and Maidenhead	Sandoz (adalimumab, infliximab)
CCG	Takeda UK (vedolizumab)
Welsh Government	,
	Relevant research groups Cochrane Inflammatory Bowel Disease and

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Issue date: November 2021

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Consultees	Commentators (no right to submit or appeal)
	Functional Bowel Disorders Group Cochrane UK 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 appraisal; 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 Appraisal Document (FAD).

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 Appraisal Document (FAD).

Commentators

Organisations that engage in the appraisal process but that are not asked to prepare an evidence submission or statement, are able to respond to consultations and they receive the FAD 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.

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