NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE

Single Technology Appraisal

Vedolizumab for treating moderately to severely active Crohn's disease after prior therapy

Provisional matrix of consultees and commentators

Consultees	Commentators (no right to submit or appeal)
Manufacturers/sponsors	General
Takeda UK (vedolizumab)	Allied Health Professionals Federation
	 Board of Community Health Councils
Patient/carer groups	in Wales
Afiya Trust	British National Formulary
Black Health Agency	Care Quality Commission
Bladder and Bowel Foundation	 Department of Health, Social Services
Bowel Cancer UK	and Public Safety for Northern Ireland
Colostomy Association	Healthcare Improvement Scotland
 Crohn's and Colitis UK (NACC) 	 Medicines and Healthcare products
Equalities National Council	Regulatory Agency
 IA: Ileostomy and Internal Pouch 	 National Association of Primary Care
Support Group	National Pharmacy Association
Muslim Council of Britain	NHS Alliance
Muslim Health Network	NHS Commercial Medicines Unit
Ostomy Lifestyle Centre	NHS Confederation
South Asian Health Foundation	 Scottish Medicines Consortium
Specialised Healthcare Alliance	
	Possible comparator manufacturer(s)
Professional groups	AbbVie (adalimumab)
 Association of Coloproctology of 	• Accord Healthcare UK (methotrexate)
Great Britain and Ireland	Actavis UK (prednisolone,
British Association for Services to	azathioprine)
the Elderly	Alliance Pharmaceuticals
British Geriatrics Society	(prednisolone)
British Institute of Radiology	Almirall (balsalazide)
British Society of Gastroenterology	Amdipharm Mercury Company
British Society of Paediatric	(hydrocortisone)
Gastroenterology, Hepatology and	 Aspen Pharma (azathioprine,
Nutrition	mercaptopurine)
Royal College of General	 AstraZeneca UK (budesonide)
Practitioners	Beacon Pharmaceuticals
Royal College of Nursing	(methylprednisolone)

National Institute for Health and Care Excellence

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Consultees	Commentators (no right to submit or appeal)
 Royal College of Pathologists Royal College of Physicians Royal College of Radiologists Royal Pharmaceutical Society Royal Society of Medicine Society and College of Radiographers United Kingdom Clinical Pharmacy Association Others Department of Health NHS England NHS Somerset CCG NHS Southampton City CCG Welsh Government 	 Dr Falk Pharma UK (budesonide, mesalazine) Ferring Pharmaceuticals (mesalazine) Hospira UK (methotrexate) Merck Sharp & Dohme (infliximab) Meda Pharmaceuticals UK (hydrocortisone) Medac GmbH (methotrexate) Napp Pharmaceuticals (prednisolone) Nova Laboratories (mercaptopurine) Orion Pharma UK (methotrexate) Pfizer (methylprednisolone, methotrexate, sulfasalazine) Sandoz (azathioprine, mesalazine, methotrexate, sulfasalazine) Shire Pharmaceuticals UK (mesalazine) Shire Pharma (usalazine) Shire Pharma (nesalazine) UCB Pharma (losalazine) UCB Pharma (losalazine) Viropharma (hydrocortisone) Warner Chilcott UK (mesalazine) Wockhardt UK (prednisolone) Zentiva UK (prednisolone) Zentiva UK (prednisolone) Cochrane Inflammatory Bowel Disease and Functional Bowel Disorders Group CORE - The Digestive Disorders Foundation Health Research Authority MRC Clinical Trials Unit National Institute for Health Research Research Institute for Health Research Health Technology Assessment Programme Warwick Evidence

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Consultees	Commentators (no right to submit or appeal)
	 Associated Guideline Groups National Clinical Guideline Centre
	 <u>Associated Public Health Groups</u> Public Health England Public Health Wales NHS 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 lists in the matrix, and which organisations we should include that have a particular focus on relevant equality issues.

PTO FOR DEFINITIONS OF CONSULTEES AND COMMENTATORS

Definitions:

Consultees

Organisations that accept an invitation to participate in the appraisal; the manufacturer(s) or sponsor(s) of the technology; national professional organisations; national patient organisations; the Department of Health and the Welsh Government and relevant NHS organisations in England.

The manufacturer/sponsor of the technology is invited to make an evidence submission, respond to consultations, nominate clinical specialists and has the right to appeal against the Final Appraisal Determination (FAD).

All non-manufacturer/sponsor consultees are invited to submit a statement¹, respond to consultations, nominate clinical specialists or patient experts and have the right to appeal against the Final Appraisal Determination (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: manufacturers of comparator technologies; Healthcare Improvement Scotland; the relevant National Collaborating Centre (a group commissioned by the Institute to develop clinical guidelines); other 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 NHS Commercial Medicines Unit, and the *British National Formulary*.

All non-manufacturers/sponsors commentators are invited to nominate clinical specialists or patient experts.

Evidence Review Group (ERG)

An independent academic group commissioned by the National Institute for Health Research (NIHR) Health Technology Assessment Programme (HTA Programme) to assist the Appraisal Committee in reviewing the manufacturer/sponsor evidence submission to the Institute.

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