NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE

Single Technology Appraisal

Brodalumab for treating moderate to severe plaque psoriasis after systemic therapy ID878

Final matrix of consultees and commentators

Consultees	Commentators (no right to submit or appeal)
Company	General
Leo Pharma (brodalumab)	All Wales Therapeutics and Toxicology Centre
Patient/carer groups	Allied Health Professionals Federation
Action Against Allergy	Board of Community Health Councils in
Allergy UK	Wales
British Skin Foundation	 British National Formulary
Changing Faces	Care Quality Commission
Muslim Council of Britain	 Department of Health, Social Services
Psoriasis Association	and Public Safety for Northern Ireland
 Psoriasis and Psoriatic Arthritis 	 Healthcare Improvement Scotland
Alliance	 Medicines and Healthcare products
Psoriasis Help Organisation	Regulatory Agency
South Asian Health Foundation	 National Association of Primary Care
Specialised Healthcare Alliance	 National Pharmacy Association
	NHS Alliance
Professional groups	NHS Commercial Medicines Unit
British Association of Dermatologists	NHS Confederation
British Dermatological Nursing Group	Scottish Medicines Consortium
British Geriatrics Society British Geriatrics Society Allegenerations Allegeneratio	Welsh Health Specialised Services
British Society for Cutaneous Allergy British Society for Dhawrood lawy	Committee
British Society for Rheumatology Drive and Care Parametel and Care in the Care Parametel and Care in the	Possible comparator companies
Primary Care Dermatology Society Payal Callage of Capacal Practitionare	AbbVie Limited (adalimumab)
Royal College of General Practitioners Doyal College of Nursing	 Accord Healthcare (methotrexate)
Royal College of Nursing Royal College of Pathologists	Accord Fleatificare (methotrexate)Allergan (acitretin)
Royal College of PathologistsRoyal College of Physicians	Astellas (tacrolimus)
Royal College of PhysiciansRoyal Pharmaceutical Society	 B&S Colorama Pharmaceuticals
 Royal Frialmaceutical Society Royal Society of Medicine 	(ciclosporin)
UK Clinical Pharmacy Association	Cubic Pharmaceuticals (ciclosporin)
- Ort Ollinour Harmady / toodiation	 Dexcel-Pharma (ciclosporin, tacrolimus)
<u>Others</u>	Eli Lilly (ixekizumab)
Department of Health	Genus Pharmaceuticals (acitretin)
NHS England	Hameln Pharmaceuticals
NHS Tower Hamlets CCG	(methotrexate)

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Consultees	Commentators (no right to submit or appeal)
NHS Trafford CCG Welsh Government	 Hospira UK (infliximab, methotrexate) Janssen (ustekinumab) Medac (methotrexate) Merck Sharp & Dohme (infliximab) Mylan UK (ciclosporin, tacrolimus) Napp (infliximab) Novartis Pharmaceuticals (secukinumab, ciclosporin) Orion Pharma UK (methotrexate) Pfizer (etanercept, methotrexate) Sandoz (methotrexate, tacrolimus) Teva UK (methotrexate, tacrolimus) Wockhardt UK (methotrexate) Relevant research groups British Epidermo-Epidemiology Society Centre of Evidence-based Dermatology, University of Nottingham Cochrane Skin Group MRC Clinical Trials Unit National Institute for Health Research Skin Research Centre Skin Treatment & Research Trust
	 Associated Public Health groups Public Health England Public Health Wales

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

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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 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 specialists and has the right to appeal against the Final Appraisal Determination (FAD).

All non-company 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: companies that market comparator technologies; Healthcare Improvement Scotland; 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-company commentators are invited to nominate clinical specialists or patient experts.

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