NATIONAL INSTITUTE FOR HEALTH AND CLINICAL EXCELLENCE

Single Technology Appraisal (STA)

Abatacept for the treatment of rheumatoid arthritis after the failure of conventional disease-modifying anti-rheumatic drugs

Matrix of consultees and commentators

Consultees	Commentators (no right to submit or appeal)
Manufacturers/sponsors	General
Bristol-Myers Squibb (abatacept)	Board of Community Health Councils in Wales
Patient/carer groups	 British National Formulary
Action on Pain	Care Quality Commission
Afiya Trust	 Commissioning Support Appraisals
Arthritic Association	Service
Arthritis & Musculoskeletal Alliance	 Department of Health, Social Services
(ARMA)	and Public Safety for Northern Ireland
Arthritis Care	 Medicines and Healthcare products
Black Health Agency	Regulatory Agency
 Chinese National Healthy Living 	 National Association of Primary Care
Centre	 NHS Alliance
Counsel and Care	 NHS Commercial Medicines Unit
Equalities National Council	 NHS Confederation
 Leonard Cheshire Disability 	 NHS Quality Improvement Scotland
Muslim Council of Britain	 Public Health Wales NHS Trust
Muslim Health Network	 Scottish Medicines Consortium
National Rheumatoid Arthritis Society	
Pain Concern	Possible comparator manufacturer(s)
Pain Relief Foundation	Abbott Laboratories (adalimumab)
Royal Association for Disability &	Actavis (azathioprine, sulfasalazine,
Rehabilitation (RADAR)	penicillamine)
South Asian Health Foundation	Alliance Pharmaceuticals (nonicillamina)
Specialised Healthcare Alliance	(penicillamine)
Drafaccional groups	Almus Pharmaceuticals (sulfasalazine) Arrow Congrigo (azathianzina)
Professional groups	Arrow Generics (azathioprine) Actallag (auranafin)
Association of Surgeons of Great Pritoin Output Description Output Description Descri	Astellas (auranofin) AstraZanaea (ablaraguina)
British Association for Sorvices to the	AstraZeneca (chloroquine) Focus Pharmacouticals (azathioprine)
 British Association for Services to the Elderly 	Focus Pharmaceuticals (azathioprine) Glave Smith Kling (azathioprine)
 British Geriatrics Society 	GlaxoSmithKline (azathioprine)IVAX Pharmaceuticals (azathioprine,
 British Health Professionals in 	IVAX Pnarmaceuticals (azatnioprine, sulfasalazine)
Rheumatology	 Kent Pharmaceuticals (azathioprine,
British Institute of Musculoskeletal	sulfasalazine, penicillamine)

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Consultees Commentators (no right to submit or appeal) Medicine Mylan (azathioprine, sulfasalazine, British Institute of Radiology penicillamine) **British Orthopaedic Association** Novartis (ciclosporin) **British Pain Society** Pfizer (sulfasalazine) British Society for Rheumatology Sandoz (azathioprine) British Society of Rehabilitation Sanofi Aventis (hydroxychloroquine, leflunomide, sodium aurothiomalate) Medicine Physiotherapy Pain Association Schering-Plough (infliximab, Primary Care Rheumatology Society golimumab) Teva UK (azathioprine, sulfasalazine, Royal College of Anaesthetists penicillamine) Royal College of General Practitioners UCB Pharma (certolizumab pegol) Royal College of Nursing Waymade Healthcare (sulfasalazine) Royal College of Pathologists Wyeth Pharmaceuticals (etanercept) Royal College of Physicians Royal College of Radiologists Relevant research groups Royal College of Surgeons Arthritis Research Campaign Royal Pharmaceutical Society Chronic Pain Policy Coalition Royal Society of Medicine -MRC Clinical Trials Unit Intellectual Disabilities Forum National Institute for Health Research Society and College of Radiographers Policy Research Institute on Ageing and United Kingdom Clinical Pharmacy Ethnicity Association Research Institute for the Care of Older People **Others** Department of Health **Evidence Review Group** Heart of Birmingham Teaching PCT School of Health & Related Research NHS Telford and Wrekin Sheffield (ScHARR) Welsh Assembly Government National Institute for Health Research Health Technology Assessment Programme Associated Guideline Groups National Clinical Guideline Centre Associated Public Health Groups tbc

NICE is committed to promoting equality and eliminating unlawful discrimination. Please let us know if we have missed any important organisations from the lists

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Appendix B

contained within the matrix and which organisations we should include who 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 Assembly Government and relevant NHS organisations in England.

The manufacturer/sponsor of the technology is invited to make an evidence submission, respond to consultations 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; NHS Quality 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 Information Authority and NHS Purchasing and Supplies Agency, 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.