NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE

Health Technology Appraisal

Tofacitinib for treating juvenile idiopathic arthritis [ID2718]

Final stakeholder list of consultees and commentators

Consultees	Commentators (no right to submit or appeal)
Company Pfizer (tofacitinib) Patient/carer groups Action on Pain Arthritis Action Arthritis and Musculoskeletal Alliance Children's Chronic Arthritis Association National Rheumatoid Arthritis Society Pain Concern Pain Relief Foundation Pain UK Psoriasis and Psoriatic Arthritis Alliance Psoriasis Help Organisation	 <u>General</u> All Wales Therapeutics and Toxicology 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 Alliance NHS Confederation
 South Asian Health Foundation Specialised Healthcare Alliance Versus Arthritis 	 Scottish Medicines Consortium Welsh Health Specialised Services Committee
 <u>Professional groups</u> British Association of Dermatologists British Dermatological Nursing Group British Institute of Musculoskeletal Medicine British Orthopaedic Association British Pain Society British Skin Foundation British Society for Rheumatology British Society for Paediatric and Adolescent Rheumatology British Society of Rehabilitation Medicine Chartered Society of Physiotherapy Physiotherapy Pain Association Primary Care Dermatology Society 	 Possible comparator companies AbbVie (adalimumab) Accord Healthcare UK (methotrexate) Advanz Pharma (methotrexate) Amgen UK (adalimumab) Biogen (adalimumab, etanercept, infliximab) Bristol-Myers Squibb Pharmaceuticals (abatacept) Celltrion Healthcare UK (infliximab) Fresenius Kabi (adalimumab) Hospira UK (methotrexate) Medac GmbH (methotrexate) Merck Sharpe & Dohme (infliximab) Mylan UK (adalimumab) Napp Pharmaceuticals (rituximab)

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Consultees	Commentators (no right to submit or appeal)
 Primary Care Rheumatology and Musculoskeletal Medicine Society Rheumatoid Arthritis Surgical Society Royal College of General Practitioners Royal College of Nursing Royal College of Paediatrics and Child Health Royal College of Pathologists Royal College of Physicians Royal College of Physicians Royal Society of Medicine Society and College of Radiographers UK Clinical Pharmacy Association Others Department of Health and Social Care NHS England NHS Herts Valley CCG NHS Telford & Wrekin CCG Welsh Government 	 Nordic Pharma (methotrexate) Orion Pharma UK (methotrexate) Pfizer (etanercept, infliximab, methotrexate) Roche Products (rituximab, tocilizumab) Rosemont Pharmaceuticals (methotrexate) Sandoz (adalimumab, etanercept, infliximab, methotrexate, rituximab) Swedish Orphan Biovitrum (anakinra) Therakind (methotrexate) Relevant research groups Bone Research Society British Psoriatic Arthritis Consortium Centre of Evidence-based Dermatology, University of Nottingham Chronic Pain Policy Coalition Cochrane Musculoskeletal Group Genomics England MRC Clinical Trials Unit National Institute for Health Research Orthopaedic Research UK Skin Treatment and Research Trust Society of Back Pain Research 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 stakeholder list, and which organisations we should include that have a particular focus on relevant equality issues.

Consultees

Organisations that accept an invitation to participate in the appraisal; the company that markets the technology; national professional organisations; national patient

Final stakeholder list for the Technology Appraisal of tofacitinib for treating juvenile idiopathic arthritis [ID2718] Issue date: December 2020 © National Institute for Health and Care Excellence 2020. All rights reserved 2 of 3 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 specialists 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 specialists 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; 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 the British National Formulary.

All non-company commentators are invited to nominate clinical specialists or patient experts.

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

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