NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE

Single Technology Appraisal

Bimekizumab for treating active psoriatic arthritis ID4009

Provisional Stakeholder List

Consultees	**Commentators (no right to submit or appeal)**
Company • UCB Pharma (bimekizumab) Patient/carer groups • Action on Pain • Arthritis Action • Arthritis and Musculoskeletal Alliance • Pain Concern • Pain UK • Psoriasis and Psoriatic Arthritis Alliance • Psoriasis Association • Psoriasis Help Organisation • South Asian Health Foundation • Specialised Healthcare Alliance • Versus Arthritis Healthcare professional groups • British Association of Dermatologists • British Dermatological Nursing Group • British Geriatrics Society • British Institute of Musculoskeletal Medicine • British Orthopaedic Association • British Pain Society • British Society for Rheumatology • British Society of Rehabilitation Medicine • Chartered Society of Physiotherapy • Physiotherapy Pain Association • Primary Care Dermatology Society • Primary Care Rheumatology and Musculoskeletal Medicine Society Society • Rheumatoid Arthritis Surgical Society	General • 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 Confederation • Scottish Medicines Consortium • Welsh Government • Welsh Health Specialised Services Committee Possible comparator companies • AbbVie (upadacitinib, risankizumab, adalimumab) • Accord Healthcare (methotrexate) • ADVANZ Pharma (methotrexate) • Amgen (adalimumab, apremilast) • Aspire Pharma (leflunomide) • Biogen Biosimilars (infliximab, adalimumab, etanercept) • Celltrion Healthcare (infliximab, adalimumab) • Cipla EU (methotrexate) • Eli Lilly (ixekizumab) • Fresenius Kabi (adalimumab)

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Consultees	Commentators (no right to submit or appeal)
• Royal College of General Practitioners • Royal College of Nursing • Royal College of Pathologists • Royal College of Physicians • Royal Pharmaceutical Society • Royal Society of Medicine • Society and College of Radiographers • UK Clinical Pharmacy Association <u>Others</u> • Department of Health and Social Care • NHS England	• Hospira UK (methotrexate) • Janssen-Cilag (guselkumab, ustekinumab) • Medac GmbH (methotrexate, leflunomide) • Merck Sharp & Dohme (golimumab, infliximab) • Morningside Healthcare (methotrexate) • Mylan (leflunomide) • Nordic Pharma (methotrexate) • Novartis (secukinumab) • Orion Pharma (methotrexate) • Pfizer (methotrexate, sulfasalazine, tofacitinib, infliximab, etanercept) • Rivopharm UK (leflunomide) • Rosemont Pharmaceuticals (methotrexate, sulfasalazine) • Sandoz (methotrexate, leflunomide, infliximab, adalimumab, etanercept) • Sanofi (leflunomide) • Therakind (methotrexate) • Tillomed Laboratories (leflunomide) <u>Relevant research groups</u> • British Psoriatic Arthritis Consortium • British Skin Foundation • Centre of Evidence-based Dermatology, University of Nottingham • Chronic Pain Policy Coalition • Cochrane Musculoskeletal Group • Cochrane Skin Group • Cochrane UK • Genomics England • MRC Clinical Trials Unit • National Institute for Health Research • Orthopaedic Research UK • Pain Relief Foundation <u>Associated Public Health groups</u> • Public Health Wales • UK Health Security Agency

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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 evaluation; 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 Draft Guidance (FDG).

All non-company consultees are invited to submit a statement[1], respond to consultations, nominate clinical or patient experts and have the right to appeal against the Final Draft Guidance (FDG).

Commentators

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

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

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