NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE

Single Technology Appraisal

Dupilumab for treating moderate to severe chronic obstructive pulmonary disease ID6235

Provisional Stakeholder List

Provisional Consultees	Provisional Commentators (no right to submit or appeal)
Company • Sanofi (dupilumab) Patient/carer groups • Action on Smoking and Health (ASH) • Asthma + Lung UK • Breathing Matters • NARA – The Breathing Charity • South Asian Health Foundation • Specialised Healthcare Alliance Healthcare professional groups • Association of Respiratory Nurse Specialists • British Geriatrics Society • British Thoracic Society • ILD-IN: Interstitial Lung Diseases Interdisciplinary Network • National Heart and Lung Institute • Primary Care Respiratory Society UK • Royal College of General Practitioners • Royal College of Nursing • Royal College of Pathologists • Royal College of Physicians • Royal Pharmaceutical Society • Royal Society of Medicine • UK Clinical Pharmacy Association Others • Department of Health and Social Care • NHS England	General • All Wales Therapeutics and Toxicology Centre • Allied Health Professionals Federation • Board of Community Health Councils in Wales • British National Formulary • Care Quality Commission • Chest, Heart and Stroke Scotland • 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 • Northern Ireland Chest, Heart and Stroke Association • Scottish Medicines Consortium • Welsh Government • Welsh Health Specialised Services Committee Possible comparator companies • ADVANZ Pharma (aminophylline) • AstraZeneca UK Limited (formoterol, symbicort, roflumilast) • Aspire Pharma Ltd (salmeterol) • Aurobindo Pharma – Milpharm Ltd. (roflumilast) • Boehringer Ingelheim Limited (tiotropium) • Chiesi Limited (formoterol, fostair) • Cipla EU Ltd (formoterol, salmeterol)

Provisional Consultees	Provisional Commentators (no right to submit or appeal)
	• Ennogen Healthcare Ltd (aminophylline) • Genus Pharmaceuticals (salmeterol) • GlaxoSmithKline UK (seretide, salmeterol) • Glenmark Pharmaceuticals Europe Ltd (tiotropium, salmeterol) • Hameln Pharma Ltd (aminophylline) • Mylan (tiotropium, salmeterol) • Novartis Pharmaceuticals UK Ltd (formoterol) • Orion Pharma (UK) Limited (formoterol, salmeterol) • Sandoz Limited (salmeterol) • Teva Pharma B.V. (formoterol) • Teva UK Limited (tiotropium, salmeterol) • Wockhardt UK Ltd (formoterol) <u>Relevant research groups</u> • British Association for Lung Research • Cochrane Airways Group • Genomics England • MRC Clinical Trials Unit • National Institute for Health Research <u>Associated Public Health group</u> • Public Health Wales • UK Health Security Agency

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 stakeholder list, and which organisations we should include that have a particular focus on relevant equality issues.

Definitions:

<u>Consultees</u>

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 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 relevant to the group they are representing, 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]); other groups (for example, the NHS Confederation and the British National Formulary).

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

Consultee or commentator stakeholders are provisional until a Confidentiality Agreement & Undertakings form is signed at appraisal stage.