NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE

Single Technology Appraisal

Ruxolitinib for treating non-segmental vitiligo in people 12 years and over ID3998

Provisional Stakeholder List

Consultees	Commentators (no right to submit or appeal)
Company	General
Incyte (ruxolitinib)	All Wales Therapeutics and Toxicology Centre
Patient/carer groups	Allied Health Professionals Federation
Changing Faces	Board of Community Health Councils in
Contact	Wales
Gene People	British National Formulary
Genetic Alliance UK	Care Quality Commission
Let's Face It	Department of Health, Social Services
South Asian Health Foundation	and Public Safety for Northern Ireland
Specialised Healthcare Alliance	Healthcare Improvement Scotland
Vitiligo Society	 Medicines and Healthcare products Regulatory Agency
Healthcare professional groups	National Association of Primary Care
British Association of Dermatologists	 National Pharmacy Association
British Dermatological Nursing Group	NHS Confederation
British Geriatrics Society	Scottish Medicines Consortium
Immunodeficiency UK	Welsh Government
Neonatal and Paediatric Pharmacists	Welsh Health Specialised Services
Group	Committee
Primary Care Dermatology Society Payol College of Capacal Prostitionare	Possible comparator companies
Royal College of General PractitionersRoyal College of Nursing	None
 Royal College of Nursing Royal College of Paediatrics & Child 	- Mone
Health	Relevant research groups
Royal College of Pathologists	British Skin Foundation
Royal College of Physicians	Cochrane Skin Group
Royal Pharmaceutical Society	Cochrane UK
Royal Society of Medicine	Genomics England
UK Clinical Pharmacy Association	MRC Clinical Trials Unit
	National Institute for Health Research
<u>Others</u>	St John's Institute of Dermatology
Department of Health and Social Care	Associated Dublic Har III
NHS England	Associated Public Health groups
	Public Health WalesUK Health Security Agency

Provisional stakeholder list for the evaluation of ruxolitinib for treating non-segmental vitiligo in people 12 years and over ID3998

Issue date: March 2023

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:

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 statement¹, 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.

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