

NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE

Single Technology Appraisal

Glycopyrronium bromide cream for treating severe primary axillary hyperhidrosis ID6487

Final Stakeholder List

Provisional Consultees	Provisional Commentators (no right to submit or appeal)
<p><u>Company</u></p> <ul style="list-style-type: none"> Leith Healthcare (glycopyrronium bromide cream) <p><u>Patient/carers groups</u></p> <ul style="list-style-type: none"> South Asian Health Foundation Specialised Healthcare Alliance <p><u>Healthcare professional groups</u></p> <ul style="list-style-type: none"> British Association of Dermatologists British Dermatological Nursing Group Primary Care Dermatology Society Royal College of General Practitioners Royal College of Nursing Royal College of Pathologists Royal College of Physicians Royal College of Surgeons Royal Pharmaceutical Society Royal Society of Medicine UK Clinical Pharmacy Association <p><u>Others</u></p> <ul style="list-style-type: none"> Department of Health and Social Care NHS England NHS Cornwall and Isles of Scilly ICB NHS North East and North Cumbria ICB 	<p><u>General</u></p> <ul style="list-style-type: none"> 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 - 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 <p><u>Comparator companies</u></p> <ul style="list-style-type: none"> Abbvie (botulinum toxin type A) Aspire Pharma (prolonged-release oxybutynin) Croma Pharma (botulinum toxin type A) Dawa (glycopyrronium bromide) Evolus International (botulinum toxin type A) Galderma (botulinum toxin type A) Grünenthal Meds (propantheline bromide) Merz Aesthetics (botulinum toxin type A) Morningside Healthcare (glycopyrronium bromide) Neon Healthcare (oxybutynin)

Final stakeholder list for the evaluation of glycopyrronium bromide cream for treating severe primary axillary hyperhidrosis ID6487

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Provisional Consultees	Provisional Commentators (no right to submit or appeal)
	<ul style="list-style-type: none"> • Tillomed laboratories (oxybutynin) • Strandhaven Limited t/a Somex Pharma (glycopyrronium bromide) • Strides Pharma (oxybutynin) <p><u>Relevant research groups</u></p> <ul style="list-style-type: none"> • British Skin Foundation • Cochrane Skin Group • Genomics England • MRC Clinical Trials Unit • National Institute for Health Research <p><u>Associated Public Health groups</u></p> <ul style="list-style-type: none"> • 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:

Consultee or commentator stakeholders are provisional until a signed Confidentiality Agreement & Undertaking form is submitted to NICE at the evaluation stage. Participating stakeholders will be listed on the project information page for the evaluation.

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 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.