NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE

Single Technology Appraisal

Linzagolix for treating moderate to severe symptoms of uterine fibroids [ID6190]

Stakeholder List

Consultees	Commentators (no right to submit or
Consuitees	appeal)
Company	General
Theramex (linzagolix)	All Wales Therapeutics and Toxicology
,	Centre
Patient/carer groups	Allied Health Professionals Federation
Bladder and Bowel Community	Board of Community Health Councils in
Bladder and Bowel UK	Wales
British Fibroid Trust	 British National Formulary
FEmISA	 Care Quality Commission
Fibroid Forum UK	 Department of Health - Northern Ireland
Fibroid Network	Healthcare Improvement Scotland
South Asian Health Foundation	Medicines and Healthcare products
Specialised Healthcare Alliance	Regulatory Agency
Wellbeing of Women	National Association of Primary Care
Women's Health Concern	National Pharmacy Association
Professional groups	NHS Confederation Section Medicines Consections
Professional groupsRoyal College of General Practitioners	Scottish Medicines ConsortiumWelsh Government
 Royal College of General Fractitioners Royal College of Nursing 	Welsh GovernmentWelsh Health Specialised Services
Royal College of Obstetricians and	Committee
Gynaecologists	Committee
Royal College of Pathologists	Comparator companies
Royal College of Physicians	AstraZeneca (goserelin)
Royal Pharmaceutical Society	Ferring Pharmaceuticals (triptorelin)
Royal Society of Medicine	Gedeon Richter (relugolix-estradiol-
UK Clinical Pharmacy Association	norethisterone acetate)
	 Ipsen (triptorelin)
<u>Others</u>	Takeda UK (leuprorelin)
Department of Health and Social Care	 Typharm (leuprorelin)
NHS England	Delevert was a such and
	Relevant research groups
	Cochrane Gynaecology and Fertility Group
	Group Genomics England
	MRC Clinical Trials Unit
	National Institute for Health Research
	- Radonal module for Health Research
	Associated Public Health groups
	Public Health Wales

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Issue date: June 2023

Appendix C

Consultees	Commentators (no right to submit or appeal)
	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:

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).

<u>Commentators</u>

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.

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