NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE

Single Technology Appraisal

Fezolinetant for treating moderate to severe vasomotor symptoms associated with the menopause ID5071

Provisional consultees Provisional commentators (no right to submit or appeal) Company General Astellas Pharma (fezolinetant) All Wales Therapeutics and • **Toxicology Centre** Patient/carer groups Allied Health Professionals • **Daisy Network** Federation Hysterectomy Association **Board of Community Health Councils** • The Menopause Charity in Wales **British National Formulary** Menopause Matters • • **Care Quality Commission** Menopause UK • Department of Health, Social Services Menopause and Cancer • and Public Safety for Northern Ireland Queer Menopause Collective 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 • **NHS** Confederation Healthcare professional groups • British Geriatrics Society Scottish Medicines Consortium • British Menopause Society Welsh Government • Faculty of Sexual & Reproductive Welsh Health Specialised Services Healthcare Committee Primary Care Women's Health Forum • Possible comparator companies Royal College of General Practitioners HRT Royal College of Nursing ADVANZ Pharma (tibolone) Royal College of Obstetricians & Artisto Pharma (tibolone) • Gynaecologists AtBayer (oestrogen) • Royal College of Pathologists Besins Healthcare (UK) (combined • Royal College of Physicians oestrogen/progesterone) Royal Pharmaceutical Society • Gedeon Richter UK (oestrogen) ٠ Royal Society of Medicine Merus Labs Luxco II S.à R.L. • Society for Endocrinology (oestrogen) **UK Clinical Pharmacy Association** • Mylan (oestrogen, combined • oestrogen and progesterone) Others Novartis (oestrogen) • Department of Health and Social Care Novo Nordisk (combined oestrogen NHS England and progesterone)

Provisional Stakeholder List

Provisional stakeholder list for the evaluation of fezolinetant for treating vasomotor symptoms associated with the menopause ID5071

Issue date: June 2024

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| Provisional consultees | Provisional commentators (no right to submit or appeal) |
|------------------------|--|
| | Orion (oestrogen, combined oestrogen and progesterone) Organon Phama UK (tibolone) Pfizer (oestrogen, combined oestrogen and progesterone) Strides Pharma UK (tibolone) Theramex UK (oestrogen, progesterone, combined oestrogen and progesterone) |
| | Anti-depressants (SSRIs, SNRIs) A. Menarini Farmaceutica Internazionale SRL (dapoxetine) Accord Healthcare (citalopram, escitalopram, fluoxetine, paroxetine, sertraline) Accord UK (citalopram, fluoxetine, paroxetine) ADVANZ Pharma (citalopram, fluoxetine, venlafaxine) Amarox (citalopram, duloxetine) Aspire Pharma (duloxetine, venlafaxine) Aurobindo Pharma-Milpharm (citalopram, duloxetine, escitalopram, fluoxetine, paroxetine, sertraline, venlafaxine) Bristol Laboratories (venlafaxine) Bristol Laboratories (venlafaxine) Brown & Burk UK (escitalopram, fluoxetine) Cipla EU (fluoxetine) Dexcel Pharma (venlafaxine) Dr Reddy's Laboratory (duloxetine, fluoxetine, sertraline) Eli Lilly and Company (duloxetine, fluoxetine) Eli Lilly and Company (duloxetine, fluoxetine) Flamingo Pharma (fluoxetine) Flamingo Pharma (fluoxetine) GlaxoSmithKline (paroxetine) Glenmark Pharmaceuticals Europe (escitalopram) Krka UK (venlafaxine) Lundbeck (citalopram, escitalopram, vortioxetine) Lupin Healthcare (sertraline) |

| Provisional consultees | Provisional commentators (no right to submit or appeal) |
|------------------------|---|
| | Martindale Pharma, an Ethypharm Group Company (venlafaxine) |
| | Morningside Healthcare (paroxetine, venlafaxine) |
| | Mylan (dapoxetine, fluvoxamine, fluoxetine, sertraline) |
| | Neuraxpharm UK (duloxetine) Pinewoord Healthcare (fluoxetine) |
| | Ranbaxy Ltd (sertraline, venlafaxine) |
| | Rivopharm UK (citalopram)Rosemont Pharmaceuticals |
| | (fluoxetine, venlafaxine)Sandoz (citalopram, paroxetine) |
| | Sovereign Medical (citalopram, venlafaxine) |
| | Thame Laboratories (citalopram) |
| | Tillomed Laboratories (venlafaxine)Upjohn UK (sertraline, venlafaxine) |
| | Zentiva (citalopram, sertraline)Zentiva (duloxetine) |
| | Clonidine |
| | SandozSyri |
| | Anti-convulsants |
| | ADVANZ Pharma (pregabalin)Amarox (gabapentin, pregabalin) |
| | Aristo Pharma (pregabalin)Aspire Pharma (pregabalin) |
| | Aurobindo Pharma - Milpharm (gabapentin, pregabalin) |
| | Brown & Burk UK (gabapentin, |
| | pregabalin)Colonis Pharma (gabapentin) |
| | Dr Reddy's Laboratories (UK) (pregabalin) |
| | Glenmark Pharmaceuticals Europe (gabapentin) |
| | MSN Laboratories Europe |
| | (pregabalin)Mylan (pregabalin) |
| | Neuraxpharm UK (pregabalin)Northumbria Pharma (pregabalin) |
| | Ranbaxy (UK) Limited a Sun Pharmaceutical Company |

| Provisional consultees | Provisional commentators (no right to submit or appeal) |
|------------------------|---|
| | (pregabalin) Rivopharm UK (gabapentin) Rosemont Pharmaceuticals Limited (gabapentin, pregabalin) Sandoz Limited (pregabalin) Strikes Pharma UK (gabapentin) Thame Laboratories (gabapentin) Tillomed Laboratories (gabapentin, pregabalin) Upjohn UK (gabapentin, pregabalin) Wockhardt UK (pregabalin) Zentiva (pregabalin) |
| | <u>Relevant research groups</u> Cochrane Menstrual Disorders and Subfertility Group Cochrane UK Genomics England MRC Clinical Trials Unit National Institute for Health Research |
| | <u>Associated Public Health groups</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.

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.

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