NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE

Single Technology Appraisal

Empagliflozin for treating chronic heart failure with preserved ejection fraction ID3945

Provisional Stakeholder List

Consultees	Commentators (no right to submit or appeal)
 Boehringer Ingelheim (empagliflozin) Patient/carer groups Arrhythmia Alliance Atrial Fibrillation Association Blood Pressure UK British Cardiac Patients Association Cardiac Risk in the Young (CRY) Cardiowappathy UK Cardiovascular Care Partnership Network of Sikh Organisations Pumping Marvellous Foundation Somerville Foundation South Asian Health Foundation Specialised Healthcare Alliance Professional groups British and Irish Hypertension Society British Association for Nursing in Cardiovascular Care British Cardiovascular Society British Geriatrics Society 	 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, 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 Alliance NHS Confederation Scottish Medicines Consortium Welsh Health Specialised Services Committee
 British Heart Foundation British Heart Rhythm Society British Nuclear Cardiology Society British Society for Heart Failure British Society of Cardiovascular Imaging National Heart and Lung Institute Primary Care Cardiovascular Society Royal College of Emergency Medicine Royal College of General Practitioners Royal College of Nursing Royal College of Pathologists Royal College of Physicians 	 Possible comparator companies Accord Healthcare (furosemide) Accord UK (furosemide) ADVANZ Pharma (furosemide) Baxter Healthcare (furosemide) Hameln Pharmaceuticals (furosemide) Ipca Laboratories UK (furosemide) M & A Pharmachem (furosemide) Mylan (bumetanide, torasemide) Pinewood Healthcare (furosemide) Rosemont Pharmaceuticals (furosemide)

Provisional stakeholder list for the technology appraisal of empagliflozin for treating chronic heart failure with preserved ejection fraction ID3945. Issue date: January 2022© National Institute for Health and Care Excellence 2022. All rights reserved 1 of 3

Consultees	Commentators (no right to submit or appeal)
 Royal Pharmaceutical Society Royal Society of Medicine Society for Cardiological Science & Technology Society for Vascular Nurses Society for Vascular Technology UK Clinical Pharmacy Association Vascular Society of Great Britain and Ireland Others Department of Health and Social Care NHS England NHS Leeds CCG NHS Telford & Wrekin CCG Welsh Government 	 Thame Laboratories (furosemide) Relevant research groups British Society for Cardiovascular Research Cardiac and Cardiology Research Dept, Barts Circulation Foundation Cochrane Heart Group Cochrane Hypertension Group Cochrane UK Cochrane Vascular European Council for Cardiovascular Research Genomics England Heart Research UK MRC Clinical Trials Unit National Centre for Cardiovascular Preventions and Outcomes National Institute for Health Research Wellcome Trust Associated Public Health groups 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 lists in 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 appraisal; 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.

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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 Appraisal Document (FAD).

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 Appraisal Document (FAD).

Commentators

Organisations that engage in the appraisal process but that are not asked to prepare an evidence submission or statement, are able to respond to consultations and they receive the FAD 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.

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