NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE

Single Technology Appraisal (STA)

Dabigatran etexilate for the treatment and secondary prevention of deep vein thrombosis and/or pulmonary embolism

Matrix of consultees and commentators

Consultees	Commentators (no right to submit or appeal)
Manufacturers/sponsors	General
 Boehringer Ingelheim (dabigatran etexilate) 	 Allied Health Professionals Federation Board of Community Health Councils in Wales
Patient/carer groups	British Cardiovascular Industry
Afiya Trust	Association
Anticoagulation Europe	British National Formulary
Black Health Agency	Care Quality Commission
British Lung Foundation	Commissioning Support Appraisals
Equalities National Council	Service
Lifeblood: The Thrombosis Charity	Department of Health, Social Services
Muslim Council of Britain	and Public Safety for Northern Ireland
Muslim Health Network	Healthcare Improvement Scotland
 South Asian Health Foundation 	 Medicines and Healthcare products
Specialised Healthcare Alliance	Regulatory Agency
	National Association of Primary Care
Professional groups	 National Pharmacy Association
Anticoagulation Specialist Association	NHS Alliance
British Cardiovascular Society	NHS Commercial Medicines Unit
British Geriatrics Society	NHS Confederation
British Orthopaedic Association	Scottish Medicines Consortium
British Society for Haematology	
British Society for Haemostasis and	Comparator manufacturers
Thrombosis	Alliance Pharma (acenocoumarol)
British Thoracic Society	Amdipharm Mercury Pharmaceuticals (abaginational superformation)
Clinical Leaders of Thrombosis	(phenindione, warfarin)
(CLOT)	Bayer (rivaroxaban) Briatal Laboratoriaa (warfaria)
Royal College of General Practitioners	Bristol Laboratories (warfarin)
Royal College of Nursing	Crescent Pharma (warfarin)
Royal College of Pathologists	GlaxoSmithKline (fondaparinux)
Royal College of Physicians	LEO Pharma (tinzaparin)
Royal Pharmaceutical Society Devial Society of Medicine	Pfizer (dalteparin) Sandaz (warfarin)
Royal Society of Medicine	 Sandoz (warfarin) Sanofi (anovanarin)
Society of Vascular Nurses	Sanofi (enoxaparin) Tara Dharmasautiasia LIK (warfarin)
 United Kingdom Clinical Pharmacy 	 Taro Pharmaceuticals UK (warfarin)

National Institute for Health and Care Excellence

Matrix for the technology appraisal of dabigatran etexilate for the treatment and secondary prevention of deep vein thrombosis and/or pulmonary embolism Issue date: December 2013

Consultees	Commentators (no right to submit or appeal)
Association Others • Department of Health • NHS England • NHS Newark & Sherwood CCG • NHS West Leicestershire CCG • Welsh Government	 Teva UK (warfarin) Zentiva UK (warfarin) Zentiva UK (warfarin) <u>Relevant research groups</u> Antithrombotic Trialists' (ATT) Collaboration Cochrane Peripheral Vascular Disease Group MRC Clinical Trials Unit National Institute for Health Research Research Institute for the Care of Older People Thrombosis Research Institute <u>Evidence Review Group</u> BMJ - TAG National Institute for Health Research Health Technology Assessment Programme <u>Associated Guideline Groups</u> National Clinical Guidelines Centre <u>Associated Public Health Groups</u> Public Health England Public Health Wales NHS Trust

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 matrix, and which organisations we should include that have a particular focus on relevant equality issues.

PTO FOR DEFINITIONS OF CONSULTEES AND COMMENTATORS

Definitions:

Consultees

Organisations that accept an invitation to participate in the appraisal; the manufacturer(s) or sponsor(s) of the technology; national professional organisations; national patient organisations; the Department of Health and the Welsh Government and relevant NHS organisations in England.

The manufacturer/sponsor of the technology is invited to make an evidence submission, respond to consultations, nominate clinical specialists and has the right to appeal against the Final Appraisal Determination (FAD).

All non-manufacturer/sponsor consultees are invited to submit a statement¹, respond to consultations, nominate clinical specialists or patient experts and have the right to appeal against the Final Appraisal Determination (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: manufacturers of comparator technologies; Healthcare Improvement Scotland; the relevant National Collaborating Centre (a group commissioned by the Institute to develop clinical guidelines); other 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 NHS Commercial Medicines Unit, and the *British National Formulary*.

All non-manufacturers/sponsors commentators are invited to nominate clinical specialists or patient experts.

Evidence Review Group (ERG)

An independent academic group commissioned by the National Institute for Health Research (NIHR) Health Technology Assessment Programme (HTA Programme) to assist the Appraisal Committee in reviewing the manufacturer/sponsor evidence submission to the Institute.

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