NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE

Single Technology Appraisal

Andexanet alfa for reversing anticoagulation from apixaban or rivaroxaban in people with intracranial haemorrhage ID6335 (part review of TA697)

Final Stakeholder List

Consultees	Commentators (no right to submit or appeal)
 Company AstraZeneca (andexanet alfa) Patient/carer groups Arrhythmia Alliance Atrial Fibrillation Association Black Health Agency British Cardiac Patients Association Cardiovascular Care Partnership (UK) Circulation Foundation Different Strokes HEART UK Muslim Council of Britain Pumping Marvellous Foundation Somerville Foundation South Asian Health Foundation Specialised Healthcare Alliance Stroke Association Thrombosis UK Professional groups Association of Anaesthetists Association of Surgeons of Great Britain and Ireland 	 General Allied Health Professionals Federation All Wales Therapeutics and Toxicology Centre 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 Hospital Information Services - Jehovah's Witnesses Medicines and Healthcare products Regulatory Agency National Association of Primary Care National Pharmacy Association NHS Alliance NHS Commercial Medicines Unit NHS Confederation Scottish Medicines Consortium Welsh Health Specialised Services Committee
 British Association for Accident and Emergency Medicine British Association for Immediate Care British Association of Stroke Physicians British Cardiovascular Society British Geriatrics Society British Trauma Society British Orthopaedic Association British Society of Interventional 	 Possible comparator companies ADVANZ Pharma (tranexamic acid) AAH Pharmaceuticals (tranexamic acid) Accord Healthcare (tranexamic acid) AMCo (tranexamic acid) Aurobindo Pharma – Milpharm (tranexamic acid) Behring UK (prothrombin complex) Bowmed Ibisqus (tranexamic acid)

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Consultees Commentators (no right to submit or appeal) Radiology Mylan (tranexamic acid) **British Society of Neuroradiologists** Octapharma (prothrombin complex British Society for Haematology concentrate) British Society for Haemostasis and • Pfizer (tranexamic acid) **Thrombosis** Rivopharm UK (tranexamic acid) British Institute of Radiology Sovereign Medical (tranexamic acid) **British Thoracic Society** Takeda (prothrombin complex) British Vein Institute Tillomed Laboratories (tranexamic Clinical Leaders of Thrombosis acid) College of Paramedics Other relevant companies Royal College of Anaesthetists Royal College of Emergency Medicine Axunio Pharma (apixaban) Royal College of General Practitioners • Bayer (rivaroxaban) Royal College of Nursing Bristol-Myers Squibb (apixaban) Royal College of Pathologists Daiichi Sankyo (edoxaban) Royal Pharmaceutical Society Mylan (apixaban) Royal College of Physicians Pfizer (apixaban) Royal College of Surgeons Sandoz (apixaban) Royal Society of Medicine Teva (apixaban) Society of British Neurological Surgeons Relevant research groups Society of Cardiothoracic Surgeons MRC Clinical Trials Unit Society for Vascular Technology National Centre for Cardiovascular Society of Vascular Nurses **Preventions and Outcomes** St Johns Ambulance National Institute for Health Research The Intensive Care Society Thrombosis Research Institute **UK Clinical Pharmacy Association** Trauma Audit Research Network Vascular Anaesthesia Society of Wellcome Trust Great Britain and Ireland Vascular Society of Great Britain and Associated Public Health Groups Ireland • Public Health England Public Health Wales Others Department of Health NHS England NHS Hounslow CCG NHS Sutton CCG Welsh Government

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

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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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