NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE

Single Technology Appraisal (STA)

Ravulizumab for paroxysmal nocturnal haemoglobinuria [ID1457]

Final stakeholder list of consultees and commentators

Consultees	Commentators (no right to submit or appeal)			
Company				
 Company Alexion Pharma UK (ravulizumab) Patient/carer groups Findacure Genetic Alliance UK PNH Support Specialised Healthcare Alliance Thrombosis UK The Aplastic Anaemia Trust Professional groups 	 General 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 Hospital Information Services – Jehovah's Witnesses Medicines and Healthcare products 			
 British Blood Transfusion Society British Committee for Standards in Haematology British Society for Haematology NHS Blood and Transplant Royal College of Nursing Royal College of Pathologists Royal College of Physicians Royal Pharmaceutical Society Royal Society of Medicine UK Clinical Pharmacy Association 	 Regulatory Agency National Association of Primary Care National Pharmacy Association National Services Division NHS Alliance NHS Confederation Scottish Medicines Consortium Welsh Government Welsh Health Specialised Services Committee 			
Others Department of Health and Social Care NHS England NHS Leeds West CCG NHS Southwark CCG	 Comparator companies Alexion Pharma UK (eculizumab) Relevant research groups Cochrane Haematological Malignancies Group Genomics England MRC Clinical Trials Unit National Institute for Health Research Associated Public Health Groups Public Health England Public Health Wales 			

Final stakeholder list for the technology appraisal of ravulizumab for paroxysmal nocturnal haemoglobinuria [ID1457].

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Appendix C

NICE is committed to promoting equality, eliminating unlawful discrimination and fostering good relations between people who share a protected characteristic and those who do share it. 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.

PTO FOR DEFINITIONS OF CONSULTEES AND COMMENTATORS

Definitions:			
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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.

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 Documentation (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 Documentation (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, 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.