NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE

Proposed Single Technology Appraisal (STA)

Alirocumab for treating primary hypercholesterolemia and mixed dyslipidaemia [ID779]

Provisional matrix of consultees and commentators

Consultees	Commentators (no right to submit or appeal)
Companies	General
Regeneron	Allied Health Professionals Federation
Sanofi	Board of Community Health Councils in
	Wales
Patient/carer groups	British Cardiovascular Industry
Afiya Trust Displict Location Agreement	Association
Black Health AgencyBlood Pressure UK	British National FormularyCare Quality Commission
British Cardiac Patients Association	Care Quality CommissionDepartment of Health, Social Services
Cardiovascular Care Partnership	and Public Safety for Northern Ireland
Coronary Prevention Group	Healthcare Improvement Scotland
Equalities National Council	Medicines and Healthcare products
HEART UK	Regulatory Agency
Muslim Council of Great Britain	National Association of Primary Care
Muslim Health Network	 National Pharmacy Association
 Network of Sikh Organisations 	NHS Alliance
South Asian Health Foundation	NHS Commercial Medicines Unit
Specialised Healthcare Alliance	NHS Confederation
Dueta asian al guarra	Scottish Medicines Consortium
Professional groups	Possible comparator companies
British Association for Nursing in Cardiovascular Care	Accord Healthcare (fluvastatin,
British Cardiovascular Intervention	pravastatin, simvastatin)
Society	Actavis (atorvastatin, fluvastatin,
British Cardiovascular Society	pravastatin, simvastatin)
British Dietetic Association	Arrow Generics (atorvastatin,
British Geriatrics Society	pravastatin, simvastatin)
British Heart Foundation	Aspire Pharma (atorvastatin, fluvastatin)
British Inherited Metabolic Disease	AstraZeneca (rosuvastatin) Association Discrete (size of action)
Group	Aurobindo Pharma (simvastatin) Bristal Laboratoriae (simvastatin)
British Nuclear Cardiology Society British Cardiology Society	Bristol Laboratories (simvastatin) Bristol Myora Squibb (proyectatin)
British Society of Cardiovascular Imaging	Bristol-Myers Squibb (pravastatin)Chanelle Medical (simvastatin)
Imaging British Society for Genetic Medicine	Dexcel-Pharma (atorvastatin,
 British Society for Genetic Medicine 	• Dexcel-Pharma (atorvastatin,

National Institute for Health and Care Excellence

Provisional matrix for the appraisal of alirocumab for treating primary hypercholesterolemia and mixed dyslipidaemia [ID779]

Consultees Commentators (no right to submit or appeal) Primary Care Cardiovascular Society simvastatin) Royal College of General Practitioners **Discovery Pharmaceuticals** (simvastatin) Royal College of Nursing • Dr Reddy's Laboratories (atorvastatin) Royal College of Pathologists Kent (pravastatin) Royal College of Physicians McNeil Products (simvastatin) Royal Pharmaceutical Society Medreich (pravastatin, simvastatin) Royal Society of Medicine Society for Cardiological Science & Merck, Sharp & Dohme (ezetimibe, simvastatin, simvastatin with ezetimibe) Technology • Mylan (fluvastatin, pravastatin, Society for Endocrinology simvastatin) Society for Vascular Technology Novartis Pharmaceuticals (fluvastatin) Society of Vascular Nurses Pfizer (atorvastatin) **UK Clinical Pharmacy Association** Ranbaxy (atorvastatin, pravastatin, UK Health Forum simvastatin) Vascular Society of Great Britain & Rosemont Pharma (simvastatin) Ireland Sandoz (fluvastatin, pravastatin, simvastatin) Others Sanofi (simvastatin) Somex (simvastatin) Department of Health NHS Birmingham South and Central Teva UK (atorvastatin, fluvastatin) pravastatin, simvastatin) Wockhardt (atorvastatin, simvastatin) NHS East Leicestershire and Rutland Zentiva (atorvastatin, fluvastatin, CCG NHS England pravastatin, simvastatin) Welsh Government Relevant research groups British Society for Cardiovascular Research Central Cardiac Audit Database Cochrane Heart Group CORDA MRC Clinical Trials Unit National Heart Research Fund National Institute for Health Research Wellcome Trust - Cardiovascular Research Initiative **Evidence Review Group** Evidence Review Group tbc National Institute for Health Research Health Technology Assessment Programme

National Institute for Health and Care Excellence Provisional matrix for the appraisal of alirocumab for treating primary hypercholesterolemia and mixed dyslipidaemia [ID779]

Appendix C

Consultees	Commentators (no right to submit or appeal)
	Associated Guideline Groups National Clinical Guidelines Centre
	 Associated Public Health Groups Public Health England Public Health Wales

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 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 company that manufactures the technology; national professional organisations; national patient organisations; the Department of Health and the Welsh Government and relevant NHS organisations in England.

The company that manufactures 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-company 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-company 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 company evidence submission to the Institute.

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