NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE

Highly Specialised Technology Evaluation

Ataluren for treating Duchenne muscular dystrophy with a nonsense mutation in the dystrophin gene (review of HST 3)

Provisional matrix of consultees and commentators

Consultees		Commentators (no right to submit or appeal)
Company(ies)		General
	oeutics (ataluren)	All Wales Therapeutics and Toxicology
	,	Centre
Patient/carer groups		Allied Health Professionals Federation
 Action Duch 	nenne	Board of Community Health Councils in
Action for Sick Children		Wales
Alex's Wish		British National Formulary
Arthritis & Musculoskeletal Alliance		Care Quality Commission
Black Health Agency		Department of Health, Social Services
Children's Society		and Public Safety for Northern Ireland
Contact		Healthcare Improvement Scotland
Disability Rights UK		Medicines and Healthcare products
	Children's Trust	Regulatory Agency
Duchenne Family Support Group		National Association of Primary Care
Duchenne N		National Pharmacy Association
Duchenne U		NHS Alliance
Genetic Allia		NHS Commercial Medicines Unit
Harrison's F		NHS Confederation Section Medicines Consertium
Information Advice and Support		Scottish Medicines ConsortiumWelsh Government
Services Network		_
Joining Jack Joining Jack		Welsh Health Specialised Services Committee
	eshire Disability	Johnnie
•	ystrophy UK ildren's Bureau	Possible comparator companies
_	Health Foundation	None
	Healthcare Alliance	···-
•	r Short Lives	Relevant research groups
i ogether for	CHOIL LIVES	British Myology Society
Professional groups		Cochrane Cystic Fibrosis and Genetic
Association of Anaesthetists		Disorders Group
Association of Genetic Nurses &		Cochrane Musculoskeletal Group
Counsellors		Duchenne Research Fund
Association of Respiratory Nurses		John Walton Centre for Muscular
Association	of Surgeons of Great	Dystrophy Research (Newcastle
Britain and I	reland	University)

National Institute for Health and Care Excellence

Provisional matrix for evaluation of ataluren for treating Duchenne muscular dystrophy with a nonsense mutation in the dystrophin gene (review of HST3) [ID1642] Page 1 of 4

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Consultees	Commentators (no right to submit or appeal)
 British Dietetic Association British Institute of Musculoskeletal Medicine British Orthopaedic Association British Paediatric Neurology Association British Paediatric Respiratory Society British Paediatric Respiratory Society British Society for Genetic Medicine British Society of Rehabilitation Medicine British Thoracic Society Chartered Society for Physiotherapy Physiotherapy Pain Association Primary Care Respiratory Society Royal College of Anaesthetists Royal College of General Practitioners Royal College of Paediatrics and Child Health Royal College of Paediatrics and Child Health Royal College of Physicians Royal College of Physicians Royal College of Surgeons Royal Pharmaceutical Society Royal Society of Medicine UK Clinical Pharmacy Association UK Genetic Testing Network Others Department of Health National Commissioning Group for Rare Neuromuscular Disorders NHS England MRC Centre for Neuromuscular Diseases Queen Square Centre for Neuromuscular Diseases Queen Square Centre for Neuromuscular Diseases Queen Square Centre Hospital Cardiff and Vale University Health Board Abertawe Bro Morgannwg University Health Board 	MRC Centre for Neuromuscular Diseases MRC Clinical Trials Unit North Star Clinical Network National Institute for Health Research TREAT-NMD Evidence Review Group National Institute for Health Research Health Technology Assessment Programme Warwick Evidence Associated Guideline Groups National Clinical Guideline Centre Associated Public Health Groups Public Health Wales UK Health security Agency

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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

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Definitions:

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 relevant NHS organisations in England.

The company that markets the technology is invited to make an evidence submission, respond to consultations, nominate clinical specialists and has the right to appeal against recommendations.

All non-company consultees are invited to make an evidence submission or submit a statement¹, respond to consultations, nominate clinical specialists or patient experts and have the right to appeal against the recommendations.

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 final evaluation document for information only, without right of appeal. These organisations are: companies that market 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 HST Evaluation 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.