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Evaluation of the clinical effectiveness of SLT for children with a primary speech/language disability.

Submission date	Recruitment status	[_] Prospectively registered
23/01/2004	No longer recruiting	[_] Protocol
Registration date	**Overall study status**	[_] Statistical analysis plan
23/01/2004	Completed	[X] Results
Last Edited	**Condition category**	[_] Individual participant data
08/08/2011	Mental and Behavioural Disorders	

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

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Additional identifiers

Protocol serial number
RCT66 Broomfield

Study information

Scientific Title

Study objectives
Is Speech and Language Therapy Effective with Children who have a primary Speech and Language Disability?

Sub-questions are:
1. Is the timing of such intervention significant?
2. Are there early prognostic indicators as compared with outcome?

Ethics approval required
Old ethics approval format

Ethics approval(s)
Not provided at time of registration

Study design
Randomised controlled trial

Primary study design
Interventional

Study type(s)
Not Specified

Health condition(s) or problem(s) studied
Mental and Behavioural Disorders: Speech disorders

Interventions
1. Brief immediate
2. Brief deferred and ongoing intervention

Intervention Type
Other

Phase
Not Specified

Primary outcome(s)
Statistical analyses of variance and regression will compare effects of intervention and prognostic indicators on outcome. The key outcome measure will be the reduction in the severity of the primary speech/language disability.

Key secondary outcome(s))
Not provided at time of registration

Completion date
08/01/2001

Eligibility

Key inclusion criteria
Those children who are referred to Speech and Language Therapy (SLT) who have a primary speech/language disability in the absence of an attributable causal condition e.g. hearing loss, developmental disability or physical disability.

Participant type(s)
Patient

Healthy volunteers allowed
No

Age group
Child

Sex
Not Specified

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
02/01/1999

Date of final enrolment
08/01/2001

Locations

Countries of recruitment
United Kingdom

England

Study participating centre
Tees and North East Yorkshire NHS Trust
Teesside
United Kingdom
TS1 3HF

Sponsor information

Organisation
NHS R&D Regional Programme Register - Department of Health (UK)

Funder(s)

Funder type
Government

Funder Name
NHS Executive Northern and Yorkshire (UK)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary
Not provided at time of registration

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/11/2011		Yes	No