

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

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

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

Primary study design

Interventional

Study design

Randomised controlled trial

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