

# Evaluation of initial fitting of young children using intraoperative neural response telemetry (NRT)

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		<input type="checkbox"/> Protocol
<b>Registration date</b> 30/09/2004	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
<b>Last Edited</b> 07/09/2015	<b>Condition category</b> Ear, Nose and Throat	<input type="checkbox"/> Individual participant data
		<input type="checkbox"/> Record updated in last year

**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**  
N0045122960

## Study information

**Scientific Title**

# Evaluation of initial fitting of young children using intraoperative neural response telemetry (NRT)

## **Study objectives**

How do the two existing methods of mapping cochlear implants, intraoperative neural response telemetry measures and behavioural assessment compare?

## **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)**

Treatment

## **Health condition(s) or problem(s) studied**

Cochlear implants

## **Interventions**

Patients are going to be randomised to either intraoperative neural response telemetry (NRT) or initial fitting based on behavioural assessment.

## **Intervention Type**

Other

## **Phase**

Not Specified

## **Primary outcome(s)**

1. Performance on Infant Toddler Meaningful Auditory Integrated Scale (IT-MAIS)
2. Performance on phonemes differentiation test (APE)
3. Soundfield Threshold Assessment
4. Time spent performing clinical activities involved

## **Key secondary outcome(s)**

Not provided at time of registration

## **Completion date**

01/09/2004

## **Eligibility**

**Key inclusion criteria**

Children aged <4 years of age with congenital deafness and no cochlear malformations such as dysplasia. Minimum of 10 children.

**Participant type(s)**

Patient

**Healthy volunteers allowed**

No

**Age group**

Child

**Upper age limit**

4 years

**Sex**

All

**Key exclusion criteria**

Does not match inclusion criteria

**Date of first enrolment**

22/10/2002

**Date of final enrolment**

01/09/2004

**Locations****Countries of recruitment**

United Kingdom

England

**Study participating centre**

**Aston University**

Birmingham

United Kingdom

B4 7ET

**Sponsor information****Organisation**

Department of Health

# Funder(s)

## Funder type

Hospital/treatment centre

## Funder Name

Birmingham Children's Hospital NHS Trust (UK)

# Results and Publications

## Individual participant data (IPD) sharing plan

### IPD sharing plan summary

Not provided at time of registration