

# Randomised controlled trial (RCT): Conservative voice treatment (CVT) versus electrostimulation therapy (EST) for unilateral vocal cord paresis (UVCP)

<b>Submission date</b>	<b>Recruitment status</b>	<input type="checkbox"/> Prospectively registered
13/09/2004	No longer recruiting	<input type="checkbox"/> Protocol
<b>Registration date</b>	<b>Overall study status</b>	<input type="checkbox"/> Statistical analysis plan
11/10/2004	Completed	<input checked="" type="checkbox"/> Results
<b>Last Edited</b>	<b>Condition category</b>	<input type="checkbox"/> Individual participant data
10/12/2008	Ear, Nose and Throat	

## Plain English summary of protocol

Not provided at time of registration

## Contact information

### Type(s)

Scientific

### Contact name

Prof Martin Ptok

### Contact details

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## Additional identifiers

### Protocol serial number

N/A

## Study information

## Scientific Title

### Acronym

CVT vs EST in UVCP: RCT

### Study objectives

Added as of 10/12/2008:

Electrical stimulation therapy is as effective as voice therapy in patients with unilateral recurrent laryngeal nerve paresis.

### Ethics approval required

Old ethics approval format

### Ethics approval(s)

The study was approved by the Ethics Committee of the Hannover Medical School.

### Study design

Randomised controlled trial

### Primary study design

Interventional

### Study type(s)

Treatment

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

Vocal cord paresis

### Interventions

Conservative voice treatment versus electrostimulation therapy

### Intervention Type

Other

### Phase

Not Applicable

### Primary outcome(s)

Not provided at time of registration

### Key secondary outcome(s)

Not provided at time of registration

### Completion date

31/12/2003

## Eligibility

### Key inclusion criteria

**Patients with unilateral vocal cord paresis**

**Participant type(s)**

Patient

**Healthy volunteers allowed**

No

**Age group**

Not Specified

**Sex**

All

**Key exclusion criteria**

Added as of 10/12/2008:

1. Voice therapy or any other therapy prior to T0
2. Paresis onset less than 2 weeks or more than 6 months prior to therapy
3. Age under 18 years
4. Paresis following resection of the recurrent nerve
5. Pareses due to muscle or joint injuries
6. Hearing loss 40 dB in the 5003000 Hz range
7. Presence of other conditions possibly interfering with the therapy, such as cognitive deficits or other serious diseases

**Date of first enrolment**

01/01/2003

**Date of final enrolment**

31/12/2003

## Locations

**Countries of recruitment**

Germany

**Study participating centre**

Klinik für Phoniatrie und Pädaudiologie

Hannover

Germany

30625

## Sponsor information

**Organisation**

Phoniatriy and Paedaudiology Clinic, Hannover Medical School (Germany)

**ROR**

<https://ror.org/00f2yqf98>

## Funder(s)

### Funder type

Other

### Funder Name

Added as of 10/12/2008:

### Funder Name

Hannover Medical School, Clinical Department of Communication Disorders (Germany)

## 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?
<a href="#">Results article</a>	final results	01/08/2008		Yes	No