ISRCTN64609553 https://doi.org/10.1186/ISRCTN64609553

Predicting the outcome of conservative (non-surgical) voice therapy for adults with laryngeal pathologies associated with hyperfunctional voice use

Submission date 10/10/2002	**Recruitment status** No longer recruiting	[_] Prospectively registered [_] Protocol
Registration date 10/10/2002	**Overall study status** Completed	[_] Statistical analysis plan [_] Results
Last Edited 01/07/2009	**Condition category** Respiratory	[_] Individual participant data [_] Record updated in last year

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
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-
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-

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
821007

Study information

Scientific Title

Study objectives
Not provided at time of registration

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

Secondary study design
Randomised controlled trial

Study setting(s)
Hospital

Study type(s)
Treatment

Participant information sheet

Health condition(s) or problem(s) studied
Voice disorders

Interventions
Patients will be randomly assigned to either the treatment or non-treatment group:
1. Treatment group: each subject in the treatment group will receive a structured voice therapy programme individually. The program consists of 10 weekly sessions involving vocal hygiene education and vocal exercises
2. Control group: the non-treatment group will receive no therapy for 10 weeks before being given voice therapy

Intervention Type
Other

Phase
Not Specified

Primary outcome measure

Not provided at time of registration

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/01/1998

Completion date
01/01/2001

Eligibility

Key inclusion criteria
1. Subjects with voice disorders should have one of the following pathologies:
1.1. Vocal nodule
1.2. Vocal polyp
1.3. Cysts
1.4. Chronic laryngitis
1.5. Odema
2. Must have normal hearing
3. Aged between 21 to 50 years old
4. Must be non-smokers
5. Consume less than one standard alcohol drink a day
6. Had not received any voice therapy
7. Not professional voice users (e.g., singers, actors, etc)

Participant type(s)
Patient

Age group
Adult

Sex
Both

Target number of participants
Not provided at time of registration

Key exclusion criteria
Does not meet inclusion criteria

Date of first enrolment
01/01/1998

Date of final enrolment
01/01/2001

Locations

Countries of recruitment
Hong Kong

Study participating centre
Department of Speech & Hearing Sciences
-
Hong Kong
-

Sponsor information

Organisation
Hong Kong Health Services Research Fund (Hong Kong)

Sponsor details
Health Welfare and Food Bureau
Government Secretariat, HKSAR
20th floor Murray Building
Garden Road
-
Hong Kong
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hsrf@hwfb.gov.hk

Sponsor type
Government

Website
http://www.fhb.gov.hk/grants/english/funds/funds_hhsrf/funds_hhsrf_abt/funds_hhsrf_abt.html

ROR
https://ror.org/03qh32912

Funder(s)

Funder type
Government

Funder Name
Hong Kong Health Services Research Fund (Hong Kong)

Results and Publications

Publication and dissemination plan
Not provided at time of registration

Intention to publish date

Individual participant data (IPD) sharing plan

IPD sharing plan summary
Not provided at time of registration