

Autologous myoblasts and fibroblasts versus collagen for treatment of stress urinary incontinence in women: a randomised trial

Submission date 19/06/2007	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 16/07/2007	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 03/10/2008	Condition category Urological and Genital Diseases	<input type="checkbox"/> Statistical analysis plan
		<input checked="" type="checkbox"/> Results
		<input type="checkbox"/> Individual participant data

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

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Additional identifiers

Protocol serial number

2002/42-21

Study information

Scientific Title

Study objectives

Incontinence can be better treated with ultrasound-guided application of autologous cells than with standard endoscopic collagen injectons.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Austrian Ministry of Health. Date of approval: 27th August 2002 (ref: GZ 2.481.159/1-VI/A/4/02)

Study design

Prospective randomized controlled trial.

Primary study design

Interventional

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Urinary incontinence

Interventions

Transurethral ultrasound guided injection of autologus myo- and fibroblasts versus transurethral endoscopic injection of collagen.

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

The following were assessed pre-operatively and 3 and 12 months post-operatively:

1. Incontinence score
2. Ultrasound parameters

Key secondary outcome(s))

The following were assessed pre-operatively and 3 and 12 months post-operatively:

1. Quality of life
2. Urodynamic parameters

Completion date

12/05/2005

Eligibility**Key inclusion criteria**

1. Women who were admitted to Department of Urology, Medical University of Innsbruck, Austria with stress incontinence.
2. Aged between 35 and 85 years

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

Female

Key exclusion criteria

1. Marked hypermobility
2. Urge incontinence

Date of first enrolment

01/09/2002

Date of final enrolment

12/05/2005

Locations**Countries of recruitment**

Austria

Study participating centre**Department of Urology**

Innsbruck

Austria

6020

Sponsor information**Organisation**

Medical University of Innsbruck, Department of Urology (Austria)

ROR

<https://ror.org/054pv6659>

Funder(s)

Funder type

Industry

Funder Name

Innovacell Biotechnologie (Austria)

Funder Name

FWF Austrian Science Fund (Fonds zur Foerderung der wissenschaftlichen Forschung) (ref: P-12828)

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	30/06/2007		Yes	No