

A randomised controlled trial to compare three therapeutic interventions in patients with self-declared illnesses due to dental amalgam fillings

Submission date 10/02/2005	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 29/07/2005	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 10/09/2009	Condition category Oral Health	<input type="checkbox"/> 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

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N/A

Study information

Scientific Title

Acronym

GAT (German Amalgam Trial)

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)

Not specified

Study type(s)

Treatment

Participant information sheet

Health condition(s) or problem(s) studied

Symptoms attributed to amalgam fillings

Interventions

Group 1: Removal of amalgam fillings and restoration with composites and other dental materials

Group 2: Removal of amalgam fillings and restoration with composites and other dental materials + mobilisation of mercury with complementary drug treatment

Group 3: No removal of amalgam fillings; psychosocial group intervention

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

01/11/1998

Completion date

31/07/2004

Eligibility

Key inclusion criteria

90 patients with symptoms attributed to amalgam fillings

Participant type(s)

Patient

Age group

Adult

Sex

Both

Target number of participants

90

Key exclusion criteria

Patients with confirmed mental disorders or organic diseases were excluded

Date of first enrolment

01/11/1998

Date of final enrolment

31/07/2004

Locations

Countries of recruitment

Germany

Study participating centre

Centre for Complementary Medicine Research
Munich
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80801

Sponsor information

Organisation

Munich Technical University - Centre for Complementary Medicine Research (Germany)

Sponsor details

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Sponsor type

University/education

ROR

<https://ror.org/02kkvpp62>

Funder(s)

Funder type

Research organisation

Funder Name

German Association of Foundations for Science (Stifterverband der Deutschen Wissenschaft)
(Germany)

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

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/04/2008		Yes	No