

(Dis-) advantages of the Wand

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|--|---|---|
| Submission date 27/01/2006 | Recruitment status No longer recruiting | <input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol |
| Registration date 27/01/2006 | Overall study status Completed | <input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results |
| Last Edited 26/08/2009 | Condition category Surgery | <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

Protocol serial number

NTR475

Study information

Scientific Title

Computerized anaesthesia delivery system (Wand) versus traditional syringe: comparing pain and pain-related behaviour in children

Acronym

WAND

Study objectives

Does the use of the Wand system reduce pain related behavior during the local anaesthesia injection in children?

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Randomised open label active controlled parallel group trial

Primary study design

Interventional

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Anesthesia injection, local

Interventions

Local anesthesia injection with a computerised anaesthesia delivery system (Wand) or the traditional syringe.

Intervention Type

Procedure/Surgery

Phase

Not Specified

Primary outcome(s)

Occurrence of 5 pain related behaviors observed in 15 s intervals of the injection time.

Key secondary outcome(s)

Venham's Distress score (0-5) in 15 s intervals of the injection time and self-reported pain (0-10)

Completion date

01/06/2006

Eligibility**Key inclusion criteria**

1. Age 4-11 years
2. Needing two treatment sessions
3. Dutch speaking

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Child

Lower age limit

4 years

Upper age limit

11 years

Sex

All

Key exclusion criteria

Special education

Date of first enrolment

01/10/2005

Date of final enrolment

01/06/2006

Locations**Countries of recruitment**

Netherlands

Study participating centre

Academisch Centrum Tandheelkunde Amsterdam (ACTA)

Amsterdam

Netherlands

1066 EA

Sponsor information**Organisation**

Academic Centre Dentistry Amsterdam (ACTA) (Netherlands)

ROR

<https://ror.org/04x5wnb75>

Funder(s)

Funder type

University/education

Funder Name

Interuniversity Dentistry Research School (IOT) (Netherlands)

Funder Name

Academic Centre Dentistry Amsterdam (ACTA) (Netherlands)

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 | 12/07/2008 | | Yes | No |