

# The effectiveness of adenotonsillectomy in children.

Gepubliceerd: 26-03-2007 Laatste bijgewerkt: 18-08-2022

Adenotonsillectomy in children with mild to moderate symptoms of throat infections or adenotonsillar hypertrophy prevents upper airway infections and fever episodes.

<b>Ethische beoordeling</b>	Positief advies
<b>Status</b>	Werving gestopt
<b>Type aandoening</b>	-
<b>Onderzoekstype</b>	Interventie onderzoek

## Samenvatting

### ID

NL-OMON26888

### Bron

Nationaal Trial Register

### Verkorte titel

NATAN

### Aandoening

1. Throat infections;
2. adenotonsillar hypertrophy;
3. adenotonsillectomy.

## Ondersteuning

**Primaire sponsor:** University Medical Center Utrecht

PO box 85500

3508 GA Utrecht

The Netherlands

**Overige ondersteuning:** CVZ programma Ontwikkelingsgeneeskunde projectnr OG 99-060

## Onderzoeksproduct en/of interventie

## **Uitkomstmaten**

### **Primaire uitkomstmaten**

Incidence of fever episodes defined as a body-temperature of 38.0 C or higher for at least one day.

## **Toelichting onderzoek**

### **Achtergrond van het onderzoek**

Objective:

In the Netherlands children with recurrent mild to moderate symptoms of throat infections or adenotonsillar hypertrophy may ultimately undergo adenotonsillectomy. Evidence regarding the balance between costs and effects of this specific 'low threshold' practice is lacking. A randomized design was used to evaluate the cost-effectiveness of adenotonsillectomy compared to watchful waiting.

Design:

Economic evaluation alongside an open randomised controlled trial.

Setting:

Multi-center: 21 general and 3 university hospitals in the Netherlands.

Participants:

300 children, aged 2 to 8 years selected for adenotonsillectomy according to routine medical practice. Excluded were children with very frequent throat infections and those with suspected obstructive sleep apnoea.

Intervention:

Adenotonsillectomy versus watchful waiting.

Main outcome measures: Incremental cost-effectiveness in terms of costs per fever episode avoided, per throat infection avoided and per upper respiratory tract infection avoided.

Results:

Annual costs incurred in the adenotonsillectomy group amounted to €803 and €551 in the watchful waiting group (46% increase). During a median follow-up period of 22 months, surgery as compared to watchful waiting reduced the number of fever episodes and throat infections by 0.21 per person year (95% confidence intervals -0.12 to 0.54 and 0.06 to 0.36, respectively), and upper respiratory tract infections by 0.53 (95% confidence interval 0.08 to 0.97) episodes. The incremental cost per episode avoided were €1,136, €1,187 and €465 respectively.

Conclusion: For children undergoing adenotonsillectomy for mild to moderate symptoms of throat infections or adenotonsillar hypertrophy the operation resulted in a significant increase in costs without realising relevant clinical benefit. Subgroups of children for whom surgery would be cost-effective may be identified in further research.

## **Doel van het onderzoek**

Adenotonsillectomy in children with mild to moderate symptoms of throat infections or adenotonsillar hypertrophy prevents upper airway infections and fever episodes.

## **Onderzoeksopzet**

N/A

## **Onderzoeksproduct en/of interventie**

Adenotonsillectomy within 6 weeks versus watchful waiting. During the study, the child's temperature was measured daily with a validated infrared tympanic membrane thermometer with an electronic device built in that stored the date and first temperature measurement of each day. Thermometer data were collected by the study physician during scheduled follow-up visits at 3, 6, 12, 18 and 24 months.

During the study, parents kept a diary of complaints of upper respiratory infections in their child; i.e. sore throat, pain/difficulty at swallowing, cough, rhinorrhea, earache and otorrhea. They also noted absence from day-care or school due to upper respiratory infections, and resource use such as prescription and over the counter medication, out-patient visits, additional surgical interventions and out-of-pocket expenses such as babysitters and travel expenses. Diary data were collected by the study physician during scheduled follow-up visits at 3, 6, 12, 18 and 24 months. On the basis of these data incidences of throat infections, sore

throat, upper respiratory infections, absence from day-care or school due to upper respiratory infections and costs were calculated.

At inclusion and the scheduled follow-up visits at 3, 6, 12, 18 and 24 months disease-specific and health-related quality of life questionnaires (TAPQoL, TACQoL, and CHQpf50) were filled out. An ear, nose and throat examination was performed including tympanometry and length and weight were measured. These data were used to establish the effect of adenotonsillectomy on middle ear status, sleeping and eating pattern, length and weight and health-related quality of life.

Serum samples were collected at baseline and at 1 year follow-up to evaluate changes in serum immunoglobulin levels in relation to surgery and occurrence of URIs.

Oropharyngeal swabs were taken at baseline and at 3 and 12 months follow-up to study the effect of adenotonsillectomy on carriage of potential pathogenic bacteria in the oropharynx at 3 and 12 months follow-up and the association between carriage of these potential pathogens and the number of throat infections during the 12 months follow-up.

## Contactpersonen

### Publiek

University Medical Center Utrecht <br>  
Wilhelmina Children's Hospital <br>  
Department of Otorhinolaryngology <br>  
PO Box 85090  
Anne G.M. Schilder  
Utrecht 3508 AB  
The Netherlands  
+31 (0)30 2504004

### Wetenschappelijk

University Medical Center Utrecht <br>  
Wilhelmina Children's Hospital <br>  
Department of Otorhinolaryngology <br>  
PO Box 85090  
Anne G.M. Schilder  
Utrecht 3508 AB  
The Netherlands  
+31 (0)30 2504004

## Deelname eisen

## Belangrijkste voorwaarden om deel te mogen nemen (Inclusiecriteria)

Children aged 2 to 8 years indicated for adenotonsillectomy according to current medical practice. These included children with recurrent throat infections (3 or more episodes per year) or other indications such as obstructive complaints or recurrent upper respiratory infections.

## Belangrijkste redenen om niet deel te kunnen nemen (Exclusiecriteria)

Children with:

1. A history of 7 or more throat infections in the preceding year, or 5 or more in each of the two preceding years, or 3 or more in each of the 3 preceding years (Paradise criteria);
2. High suspicion of obstructive sleep apnoea, i.e. Brouillette's OSA-score of more than 3.5;
3. Down's syndrome;
4. Craniofacial malformation, such as cleft palate;
5. Documented immunodeficiency, other than IgA or IgG2 deficiencies.

## Onderzoeksopzet

### Opzet

Type:	Interventie onderzoek
Onderzoeksmodel:	Parallel
Toewijzing:	Gerandomiseerd
Blinding:	Open / niet geblindeerd
Controle:	Actieve controle groep

### Deelname

Nederland	
Status:	Werving gestopt
(Verwachte) startdatum:	01-03-2000

Aantal proefpersonen: 300  
Type: Werkelijke startdatum

## Ethische beoordeling

Positief advies  
Datum: 26-03-2007  
Soort: Eerste indiening

## Registraties

### Opgevolgd door onderstaande (mogelijk meer actuele) registratie

Geen registraties gevonden.

### Andere (mogelijk minder actuele) registraties in dit register

Geen registraties gevonden.

### In overige registers

Register	ID
NTR-new	NL916
NTR-old	NTR940
Ander register	:
ISRCTN	ISRCTN04973569

## Resultaten

### Samenvatting resultaten

1. van Staaïj BK, van den Akker EH, Rovers MM, Hordijk GJ, Hoes AW, Schilder AG. Effectiveness of adenotonsillectomy in children with mild symptoms of throat infections or adenotonsillar hypertrophy: open, randomised controlled trial. BMJ. 2004 Sep 18;329(7467):651. <br>
2. Arch Otolaryngol Head Neck Surg. 2007 Nov;133(11):1083-8.<br>

