Relationship between allergic manifestations and Toxocara seropositivity: a cross-sectional study among elementary school children

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ABSTRACT: Toxocara (the cause of visceral larva migrans in humans) and allergens may accelerate expression of allergic symptoms at an early age. Toxocara spp., intestinal parasites of dogs and cats, have polyclonal B-cell-activating properties, and can, thereby, induce high serum total immunoglobulin E (IgE) concentrations. Such a polyclonal B-cell stimulation may occur in children who are also sensitized to common allergens, such as house dust mite (Dermatophagoides pteronyssinus) allergen. Frequent exposure to allergens may accelerate expression of allergic symptoms at a young age. Various products of biological origin, such as antigens derived from parasitic worms, have polyclonal B-cell-activating properties, and can, thereby, induce high serum total immunoglobulin E (IgE) concentrations.

Recent investigations suggest an increasing prevalence of childhood asthma [1, 2]. Children at risk are those with an atopic constitution [3, 4]. The age at which allergy becomes manifest depends mainly on the degree of exposure to inhaled allergens. The most obvious being that to house dust mite (Dermatophagoides pteronyssinus) allergen [5]. Frequent exposure to allergens may accelerate expression of allergic symptoms at a young age [6]. Various products of biological origin, such as antigens derived from parasitic worms, have polyclonal B-cell-activating properties, and can, thereby, induce high serum total immunoglobulin E (IgE) concentrations.

**Beta**
In a previous epidemiological study, we demonstrated elevated serum total IgE levels and a more frequent occurrence of inhaled allergen-specific IgE in Toxocara-seropositive as compared to seronegative children. The sample comprised 4–6 yr old elementary schoolchildren from two densely populated Dutch urban areas. This study also suggested an association of Toxocara seroprevalence with physician-confirmed asthma [9]. It was assumed that seropositive children had, or had experienced, an infection with Toxocara.

The endemicity of Toxocara in the Netherlands [9, 16], together with the pathological phenomena described above, suggest that infection with this kind of parasitic roundworm may accelerate expression of asthma-like affections in susceptible children, and possibly aggravates already-existing asthma. Such an association may be important in view of the increasing number of children having such affections.

In the present cross-sectional study, we investigated whether an association between toxocarosis and allergic manifestations could be established in elementary schoolchildren, aged 4–12 yrs, of urban or rural origin. We also investigated whether risk factors for infection with Toxocara could be identified in this age group. The study enabled us to establish whether differences existed between urban and rural areas in the prevalence of Toxocara antibodies and allergic manifestations, and in the relationship between these two conditions. Furthermore, the similar design of the previous and present study allowed comparison of results. The presence of allergic manifestations was to be estimated from questionnaires, from eosinophil numbers, total serum IgE concentration, and from the presence of inhaled allergen-specific IgE.

Materials and methods

Population

The investigation was carried out among 1,379, 4–12 yr old, elementary schoolchildren from urban and rural areas in the provinces of Utrecht and Brabant in the Netherlands. Six hundred and eighty three of the children lived in the province of Utrecht: 335 in Utrecht city (urban), distributed over three schools; and 348 in the countryside (rural), distributed over seven schools. Six hundred and ninety six of the children lived in the province of Brabant: 328 in Eindhoven city (urban), distributed over three schools, and 368 in the countryside (rural), distributed over two schools. The province of Utrecht is located in the centre and that of Brabant in the south of the Netherlands. Both provinces have similar demographic characteristics. Schools were selected in co-operation with the community health services, in order to obtain representative samples of the population.

The surveys were carried out in Utrecht from November 1989 to April 1990, and in Brabant from December 1991 to February 1992. To control for socioeconomic status in the samples from the city populations, schools were selected on the basis of the socioeconomic category (low, middle and high) of the neighbourhoods. National criteria, such as the parents’ education, income and housing, were used to categorize the schools; this information was provided by the Municipal Centers for Research and Statistics. An estimated 85% of the pupils attending a particular school were of the same socioeconomic category as the neighbourhood in general. Division by socioeconomic categories was not meaningful in the countryside. In rural Utrecht, each village had its own small size school attended by all children in the vicinity. In rural Brabant, larger schools served larger areas.

The parents were asked to complete a questionnaire, which, together with a letter of consent for the participation of their child(ren), was returned to the school staff. Per school, 60–85% of the parents returned the letter of consent. Ethical aspects were considered and approved by an independent group of public health authorities of the Public Health Services. No such investigation has been performed in any of the study areas previously.

Blood collection and examination

Sera were collected from children aged 4–12 yrs in Utrecht city (urban) and Brabant province (rural), distributed over seven schools. The sera were collected during the period from November 1989 to April 1990, and from December 1991 to February 1992.

Toxocara serology

Antibodies to Toxocara were measured, as described by Van Knapen et al. [17]. Peroxidase-labelled rabbit-anti-human immunoglobulin G (IgG) (Institut Pasteur, Marnes la Coquette, France) was used as a conjugate. Each microtitre plate contained 7–9 control sera, i.e. 6–8 known negative samples and one standard serum with known Toxocara antibody titre. Since contact of the children with Toxocara (and not disease) was subject to study, the sera were screened in a 1:20 dilution, the lowest dilution at which specific antibodies were detected. Results were considered positive when the extinction value was higher than the mean extinction value of 6–8 negative control sera raised with three times the standard deviation. Results were expressed as either negative or positive. Antibody measurement was performed according to good laboratory practice procedures, including the required controls for approval of new batches of reagents.

Estimations of serum total IgE concentration and of IgE specific for inhaled allergens

The PharmaciaCAP System IgE radioimmunoassay (RIA) and the PharmaciaCAP System radioallergosorbent test (RAST) were used for the estimation of serum
total IgE and of inhaled allergen-specific IgE, respectively. The CAP System included equipment, test kits and all required control and standard sera. The tests are based on the ImmunoCAP technology. Briefly, the antigens, anti-IgE or allergens, are covalently coupled to cyanogen bromide-activated sepharose beads (ImmunoCAP). IgE or inhaled allergen-specific IgE, when present in the serum, reacts with the ImmunoCAP. Radioactively labelled antibodies against IgE are then added, which bind to the complex. The radioactivity of the complex formed is measured in a gamma counter. The amount of bound radioactivity is a measure of the quantity of total IgE or inhaled allergen-specific IgE. The detection limit for total IgE is 0.8 kU·L⁻¹ and for inhaled allergen-specific IgE is <0.35 kU·L⁻¹.

The allergens tested were Dermatophagoides pteronyssinus, Canis familiaris and Felis domesticus. Because, the majority of the sera containing IgE specific for F. domesticus and/or C. familiaris also contained D. pteronyssinus-specific IgE, the samples were categorized as either nonreacting or reacting with at least one allergen. Results are expressed as kU IgE·L⁻¹ for serum total IgE, and as either negative or positive for inhaled allergen-specific IgE. A concentration of >0.35 kU of inhaled allergen-specific IgE·L⁻¹ was considered positive.

Estimations of Toxocara-specific IgE

Toxocara-specific IgE was estimated by J-F. Magnaval (Toulouse, France), in six Toxocara-seropositive and seronegative samples, according to the method described by Magnaval et al. [18]. The Toxocara-positive sera were selected on the basis of varying total IgE concentrations. The Toxocara-negative samples (controls) were selected on the basis of high total IgE concentrations.

Questionnaires

Prior to visiting the school, the school staff, the parents and the general practitioners were informed in writing about the aim of the investigation, the life cycle of the parasite, and the route of infection. Questionnaires were distributed to the parents requesting: 1) name, date of birth, and sex of the child; 2) information concerning putative risk factors for infection, i.e. the presence of dogs and cats in the home, contact with animals outside the home, and use of public playgrounds; 3) information concerning allergic complaints, i.e. "Does the child suffer from asthma/recurrent bronchitis?"; "Does the child react with allergic symptoms during or after contact with animals?"; "Does the child suffer from eczema?"; and 4) whether the child had ever been hospitalized with asthma. Questions about allergic symptoms were followed by: "If the answer is yes, has it been confirmed by a physician?". Asthma was diagnosed using national criteria, which are concordant with those of the American Thoracic Society. Recurrent bronchitis was mentioned considering that 4 and 5 yr olds are suspected of an asthmatic condition when they have recurrent bronchial ailments at that age. Answers were accepted only when allergic disorders observed by the parents were subsequently confirmed by a physician.

Risk factors

Putative risk factors were also investigated by questionnaire. The parents were asked: 1) whether or not the family owned dogs and/or cats, now or ever; 2) if their children played at public playgrounds; 3) and if the children had contact with animals which did not belong to the household.

Statistical analysis

Logistic regression analysis was used to study the associations between Toxocara seroprevalence and allergic manifestations, inhaled allergen-specific IgE, the risk factors, and the background variables: district (Utrecht versus Eindhoven), environment (urban vs rural), sex, school and age. These analyses were made using the EGRET package [19]. Firstly, the relationships between Toxocara seroprevalence and the five background variables were studied, and subsequently those between Toxocara seroprevalence and allergic manifestations, between Toxocara seroprevalence and inhaled allergen-specific IgE, and between Toxocara seroprevalence and the risk factors. The latter three associations were corrected for confounding effects of the background variables. School was entered in the models as a random effect, to account for the fact that children from the same school were more alike than children from different schools. Not adjusting for school in this way may cause an underestimation of confidence intervals of the odds ratios, which in its turn may lead to spuriously significant results [19]. For each model, all first order interactions between fixed effects were examined, and statistically significant ones were retained.

Analysis of covariance (ANCOVA) was used to study the relationships between Toxocara seroprevalence and serum immunoglobulin E concentration, and between Toxocara seroprevalence and eosinophil number. PROC GLM of the Statistical Analysis System (SAS) package [20] was used to perform these analyses. IgE and eosinophil values were log-transformed in order to achieve approximate normality. Toxocara seroprevalence and the background variables, district, environment and sex, were entered as fixed factors in the models. School was entered as a fixed blocking factor, nested under district and environment. Age was included as covariate. This design contained 2 × 2 × 15 = 60 cells. One of the cells was empty in the IgE data and two cells were empty in the eosinophil data. To avoid the complication of analysing designs having some cells empty, one artificial observation was placed in each of these cells. These observations had the average IgE or eosinophil value for the particular combination of levels of the factors, Toxocara seroprevalence, district, environment and sex they were in, and also the average value for age. These observations were not included in the numbers of observations shown in the tables.

Initially, models containing all possible interactions of all orders between Toxocara seroprevalence, district, environment and sex were fitted. Interactions with school
Results

Sample characteristics

The number of children observed in each of the 15 schools, as well as the seroprevalence of *Toxocara*, asthma/recurrent bronchitis and inhaled allergen-specific IgE are presented in table 1. The numbers of subjects varied considerably among schools. Relatively small numbers were observed in the seven schools in the rural vicinity of Utrecht. However, the numbers of children in each of the four regions were very similar, namely approximately 340. Also, the number of boys and girls, 688 and 691 respectively, were almost equal.

We first studied whether living in the district of Utrecht or in the district of Eindhoven (district), living in an urban or in a rural environment (environment), sex, age or the school attended were associated with *Toxocara* seroprevalence. The overall seroprevalence of *Toxocara* was 8%. We did not find a relationship with district (odds ratio (OR) 1.18; 95% confidence interval (95% CI) 0.66–2.11) or with environment (OR 0.83; 95% CI 0.46–1.49). Significantly less girls (7%) than boys (10%) had *Toxocara* antibodies. Using boys as the baseline category, the OR for sex was 0.64 (95% CI 0.44–0.95). However, this OR varied between the four subpopulations formed by combining the levels of district and environment in the sample studied. This was analysed by studying the association between *Toxocara* seropositivity and sex separately in each of the four strata. This revealed that the odds for *Toxocara* seropositivity were always smaller in girls than in boys, except in the city of Eindhoven, where girls had the larger odds. The OR for sex was significant in the city of Utrecht (OR 0.35; 95% CI 0.13–0.93) and in rural Utrecht (OR 0.43; 95% CI 0.18–1.05), but not in Eindhoven city (OR 1.49; 95% CI 0.72–3.13) or in rural Eindhoven (OR 0.51; 95% CI 0.19–1.30).

*Toxocara* seroprevalence varied widely between the schools from a minimum of 2% to a maximum of 24% (table 1), which resulted in a highly significant random effect of school (p<0.001, likelihood ratio test) in the logistic regression model. The median age in the total sample was 8 yrs. Age did not show a relationship with *Toxocara* seroprevalence. The overall seroprevalence of *Toxocara* was 8%. We did not find a relationship with district (odds ratio (OR) 1.18; 95% confidence interval (95% CI) 0.66–2.11) or with environment (OR 0.83; 95% CI 0.46–1.49). Significantly less girls (7%) than boys (10%) had *Toxocara* antibodies. Using boys as the baseline category, the OR for sex was 0.64 (95% CI 0.44–0.95). However, this OR varied between the four subpopulations formed by combining the levels of district and environment in the sample studied. This was analysed by studying the association between *Toxocara* seropositivity and sex separately in each of the four strata. This revealed that the odds for *Toxocara* seropositivity were always smaller in girls than in boys, except in the city of Eindhoven, where girls had the larger odds. The OR for sex was significant in the city of Utrecht (OR 0.35; 95% CI 0.13–0.93) and in rural Utrecht (OR 0.43; 95% CI 0.18–1.05), but not in Eindhoven city (OR 1.49; 95% CI 0.72–3.13) or in rural Eindhoven (OR 0.51; 95% CI 0.19–1.30).

Toxocara seroprevalence varied widely between the schools from a minimum of 2% to a maximum of 24% (table 1), which resulted in a highly significant random effect of school (p<0.001, likelihood ratio test) in the logistic regression model. The median age in the total sample was 8 yrs. Age did not show a relationship with *Toxocara* seroprevalence. Furthermore, no significant interactions were found between age and the other variables investigated.

Association between *Toxocara* seroprevalence and allergic manifestations

ORs for the associations between *Toxocara* seroprevalence and allergic manifestations, presented in table 2, were adjusted for the background variables, district, environment, sex and school. The allergic manifestations were not related to age. Age was, therefore, omitted from the models. Asthma/recurrent bronchitis and allergic reaction on animal contact were significantly associated with *Toxocara* seroprevalence, whereas eczema and hospitalization due to asthma/recurrent bronchitis were not. No statistically significant interactions between allergic manifestations and the background variables were found.

Association of inhalant allergen-specific IgE with *Toxocara* seropositivity and sex

Inhaled allergen-specific IgE was found significantly more often in the *Toxocara*-seropositive group as compared to the group in which no antibodies were
demonstrated (table 3). Furthermore, inhaled allergen-specific IgE was found significantly more often in children from urban than from rural backgrounds, and more often in boys than in girls. Also, its occurrence increased with age. No difference was observed between the districts of Utrecht and Eindhoven.

The considerable variation among schools in percentage of children presenting allergen-specific IgE, ranging 5–27% (table 1), turned out to be nonsignificant in this analysis after adjustment for the other variables in the model (table 3).

Association of eosinophil numbers and serum IgE levels with Toxocara seroprevalence

Overall, the number of eosinophils was higher in the Toxocara-seropositive group (221 cells·µL⁻¹; 95% CI 188–259 cells·µL⁻¹) as compared to the seronegative group (175 cells·µL⁻¹; 95% CI 167–184 cells·µL⁻¹), as indicated by a statistically significant main effect of Toxocara seroprevalence (p=0.008). However, the simplest ANCOVA model that adequately described the relationships between eosinophil number and Toxocara, district, environment, sex, school and age included, in addition to the main effects of all factors, a significant second order interaction between Toxocara seroprevalence, district and environment (p=0.043). We have interpreted this as the effect of Toxocara seroprevalence on eosinophil numbers being different in each of the four subgroups formed by combining the levels of district and environment. Results for these groups are presented in table 4. Although the direction of the difference between Toxocara-seronegative and seropositive groups was the same in all groups, the magnitudes of the differences varied considerably. The largest and only significant difference was found in the rural vicinity of ASSOCIATION OF TOXOCARA WITH ALLERGY 1471

Table 2. – Logistic regressions of allergic manifestations on Toxocara seroprevalence: Dutch elementary schoolchildren 1989–1992

| Outcome                  | Cases         | Total* (%) | Crude† OR  | Adjusted‡ OR  | 95% CI‡ |
|--------------------------|---------------|------------|------------|---------------|---------|
| Asthma/RB -ve            | 77/1257       | 6          | Baseline   | Baseline      | 1.41–4.39 |
| +ve                      | 17/114        | 15         | 2.69       | 2.49          |         |
| Allergic reaction -ve    | 41/1251       | 3          | Baseline   | Baseline      | 1.65–6.42 |
| on animal contact +ve    | 12/113        | 11         | 3.51       | 3.25          |         |
| Eczema -ve               | 61/1252       | 5          | Baseline   | Baseline      | 0.34–2.19 |
| +ve                      | 5/112         | 4          | 1.18       | 0.86          |         |
| Hospitalization -ve      | 31/1258       | 2          | Baseline   | Baseline      | 0.59–4.22 |
| due to asthma/RB +ve     | 5/114         | 4          | 1.82       | 1.58          |         |

*: number of cases of the outcome variable versus the total number of children in that Toxocara category; †: odds ratios (OR) versus baseline category; ‡: ORs adjusted for district, urban/rural environment, sex and school; †: confidence intervals (CI) of the adjusted OR. -ve: negative; +ve: positive; RB: recurrent bronchitis.

Table 3. – Logistic regressions of inhalant allergen-specific immunoglobulin E on Toxocara seroprevalence, environment, district, sex, age and school: Dutch elementary schoolchildren 1989–1992

| Variable                  | Cases         | Total* (%) | Crude† OR  | Adjusted‡ OR  | 95% CI‡ |
|---------------------------|---------------|------------|------------|---------------|---------|
| Toxocara                  |               |            |            |               |         |
| Negative                  | 193/1238 (16) | BL         | BL         |               |         |
| Positive                  | 28/112 (25)   | 1.81       | 1.66       | 1.04–2.65     |         |
| Environment               |               |            |            |               |         |
| Urban                     | 126/647 (20)  | BL         | BL         |               |         |
| Rural                     | 95/703 (14)   | 0.65       | 0.61       | 0.43–0.87     |         |
| District                  |               |            |            |               |         |
| Utrecht                  | 97/647 (15)   | BL         | BL         | 0.86–1.74     |         |
| Eindhoven                 | 124/684 (18)  | 1.30       | 1.23       |               |         |
| Sex                       |               |            |            |               |         |
| Boys                      | 133/671 (20)  | BL         | BL         |               |         |
| Girls                     | 88/679 (13)   | 0.60       | 0.61       | 0.45–0.82     |         |
| Age (yrs) continuous      |               | 1.10       | 1.10       | 1.04–1.17     |         |
| School (random)           |               | p-value=0.186 |         |               |         |

*: number of children with inhalant allergen-specific immunoglobulin E versus the total number of children in that category; †: odds ratio (OR) adjusted for all other variables in the table; ‡: 95% confidence intervals (CI) of the adjusted ORs; BL: baseline.

Table 4. – Analysis of covariance of eosinophil numbers in relation to Toxocara seroprevalence, district, environment, sex, age and school: Dutch elementary schoolchildren 1989–1992

| Eosinophils cells·µL⁻¹ | n     | L-S Mean† | 95% CI† | p-value |
|------------------------|-------|-----------|---------|---------|
| Utrecht: urban         |       |           |         |         |
| Toxocara -ve           | 303   | 198       | 180–219 | 0.1404† |
| +ve                    | 24    | 260       | 183–367 |         |
| Utrecht: rural         |       |           |         |         |
| Toxocara -ve           | 304   | 178       | 161–197 | 0.6050† |
| +ve                    | 26    | 195       | 141–269 |         |
| Eindhoven: urban       |       |           |         |         |
| Toxocara -ve           | 287   | 190       | 172–210 | 0.8767† |
| +ve                    | 38    | 195       | 148–255 |         |
| Eindhoven: rural       |       |           |         |         |
| Toxocara -ve           | 340   | 141       | 129–154 | 0.0030† |
| +ve                    | 24    | 241       | 171–339 |         |
| Sex                    |       |           |         |         |
| Boys                   | 672   | 205       | 187–226 | 0.0669† |
| Girls                  | 674   | 189       | 171–208 | 0.0576† |
| Age (yrs)              | -0.0179; 0.00944 | 0.0576† |
| School                 | 0.3259 |           |         |         |

n: number of children in that category; -ve: negative; +ve: positive. †: least squares (L-S) means and their 95% confidence intervals (CI), transformed back to the original scale; †: p-value belonging to the tests of the contrast between Toxocara-negative and seropositive groups within each of the four combinations of the levels of the factors, district and environment; ‡: p-value of the difference in eosinophil numbers between the sexes; §: coefficient and standard error of the covariable age indicating the change in log eosinophil numbers by a change of 1 yr in age.
Eindhoven, whilst in Eindhoven city a very small difference was found. The two samples from Utrecht were intermediate. There was a considerable variation among the seropositive groups, and less so among the seronegative ones.

The number of eosinophils was higher in boys than in girls, a difference that was borderline significant (table 4). Furthermore, a small and almost significant negative trend with age was found, corresponding with a decrease of almost 2% in the number of eosinophils for a 1 yr increase in age. No significant differences between schools were found. Least squares means transformed back to the original scale ranged 166–247 cells·μL⁻¹ (detailed results not shown).

For the IgE values, the same model was found to be adequate as that used for the eosinophil numbers. Therefore, for this variable also, a second order interaction between district, environment and Toxocara (p=0.0402) was included in the model. Overall, a significantly higher serum total IgE concentration was found in the seropositive group (126.9 IU·mL⁻¹; 95% CI 94–172 IU·mL⁻¹) than in the seronegative group (66.3 IU·mL⁻¹; 95% CI 61–73 IU·mL⁻¹; p=0.0001). In this case, also, examination of this difference in the four subpopulations separately revealed a substantial variation (table 5). Large and statistically significant differences were found in the city of Utrecht and in rural Eindhoven. A much smaller difference was observed in the city of Eindhoven, and a very small difference in rural vicinity of Utrecht. Again, the direction of the difference was the same in all subgroups. Serum IgE levels were higher in children

Table 5. – Analysis of covariance of serum total immunoglobulin E (IgE) levels in relation to Toxocara seroprevalence, district, environment, sex, age and school: Dutch elementary schoolchildren 1989–1992

|                | n     | L-S Mean | 95% CI† | p-value |
|----------------|-------|----------|---------|---------|
| Utrecht:urban  |       |          |         |         |
| Toxocara -ve   | 306   | 85.1     | 70.7–102.5 | 0.0078‡ |
| +ve            | 24    | 212.4    | 110.6–407.8 |         |
| Utrecht:rural  |       |          |         |         |
| Toxocara -ve   | 315   | 45.7     | 38.1–54.9 | 0.0801‡ |
| +ve            | 27    | 54.6     | 29.8–100.0 | 0.5824‡ |
| Eindhoven:urban|       |          |         |         |
| Toxocara -ve   | 286   | 76.0     | 63.0–91.7 | 0.0998‡ |
| +ve            | 38    | 120.2    | 72.1–200.2 |         |
| Eindhoven:rural|       |          |         |         |
| Toxocara -ve   | 340   | 65.4     | 55.2–77.6 | 0.0021‡ |
| +ve            | 24    | 186.0    | 97.8–353.6 |         |
| Sex            |       |          |         |         |
| Boys           | 678   | 101.2    | 84.9–120.6 |         |
| Girls          | 674   | 83.2     | 69.1–100.0 | 0.0249‡ |
| Age (yrs)      | (0.0844; 0.0176) | 0.0001 |
| School         |       | 0.0001   |         |         |

n: number of children in that category; -ve: negative; +ve: positive. †: least squares (L-S) means and their 95% confidence intervals (CI), transformed back to the original scale; ‡: p-value belonging to the tests of the contrast between Toxocara-negative and positive groups within each of the four combinations of the levels of the factors, district and environment; §: p-value of the difference in IgE concentrations between the sexes; †: coefficient and standard error of the covariable age indicating the change in ln IgE level by a change of 1 yr in age.

Table 6. – Serum total immunoglobulin E (IgE) concentration versus Toxocara-specific IgE in Toxocara-seropositive and seronegative samples

| No. | Total IgE† | Tox.-specific IgE‡ |
|-----|------------|--------------------|
| 1   | 727        | 40.0               |
| 2   | >2000      | 2.5                |
| 3   | >2000      | <1.0               |
| 4   | 673        | 1000.0             |
| 5   | 481        | <1.0               |
| 6   | 866        | 3.0                |
| 7   | 1261       | 4.0                |
| 8   | 1669       | <1.0               |
| 9   | >2000      | <1.0               |
| 10  | 1243       | <1.0               |
| 11  | 1759       | <1.0               |
| 12  | >2000      | <1.0               |

†: serum total IgE concentration expressed in international kilo units per litre of serum; ‡: Toxocara-specific IgE expressed in Toxocara units per litre of serum; Tox.: Toxocara.

from an urban (113.4 IU·mL⁻¹) than from a rural environment (74.2 IU·mL⁻¹, p=0.0055), higher in males than in females, and increased almost 9% with a 1 yr increase in age (table 5). IgE levels varied widely among schools, ranging 29.4–237.8 IU·mL⁻¹, a variation that was highly significant.

Toxocara-specific IgE

Serum samples, selected from the Toxocara-seropositive and seronegative groups on the basis of total serum IgE concentrations, were analysed for Toxocara-specific IgE. No relationship was found between levels of total IgE and Toxocara-specific IgE concentrations in the six seropositive samples, two of which even demonstrated absence of specific IgE. Five of the six seronegative samples were negative for specific IgE. One sample, however, demonstrated a low positive reaction (table 6).

Risk factors

Relationships between Toxocara seroprevalence and the putative risk factors were adjusted for confounding effects of district, environment, school, sex and age. Having kept a dog as a pet at sometime in the life of a child resulted in higher odds for having Toxocara antibodies (OR 1.52; 95% CI 1.01–2.29). Currently keeping a dog, raised the odds for having Toxocara antibodies only slightly, and not significantly (OR 1.22; 95% CI 0.80–1.86). Similarly, having a dog in the past but not at present raised the odds, but not significantly (OR 1.47; 95% CI 0.98–2.22). Having kept a cat in the past, or having a cat in the past but not at present, also yielded (not significantly) higher odds (OR 1.53; 95% CI 1.01–2.33, respectively). Currently having a cat was not related to Toxocara seroprevalence at all (OR 0.99; 95% CI 0.59–1.67). The dog
ownership variables were included in the models describing the associations between Toxocara seroprevalence and allergic manifestations to check for possible confounding effects. This did not appreciably lower the odds ratios for Toxocara, and thus did not change the findings.

Use of public playgrounds and the amount of contact with animals outside the home were measured on a three-point scale, having the categories: no, infrequent, and frequent use or contact. Of the children reporting no contact with animals outside the home, 14.5% were Toxocara seropositive. In the groups reporting infrequent and frequent contact these percentages were 6 and 10.5%, respectively. Frequent and infrequent contact, therefore, resulted in lower odds for Toxocara seroprevalence when compared to the group that reported no contact. The OR for infrequent contact was 0.42 (95% CI 0.24–0.74), which was significant, and for frequent contact 0.77 (95% CI 0.42–1.41), which was not significant.

Of the children who never played in public playgrounds, 9.3% were Toxocara seropositive; 6.9% were seropositive in the group who played only infrequently in these playgrounds; and 10.4% in those who did so frequently. Thus, when using the group who never played in public playgrounds as the baseline category, the OR for frequent use was 0.73 (95% CI 0.42–1.29) and for frequent use 1.11 (95% CI 0.62–2.00), neither of them being significant.

### Discussion

The main aim of the present study was to investigate whether or not an association exists between allergy-related phenomena and Toxocara seroprevalence. Such an association was hypothesized on the basis of characteristics that toxocarosis and allergy have in common, i.e., excessive IgE production, eosinophilia, and respiratory complaints [1, 3, 13]. Excessive IgE production after contact with IgE-stimulating agents is an inherited phenomenon [21]. Young children with an atopic condition risk development of allergic asthma during growth [22]. Since accumulated exposure to inhaled allergens of 4–12 yr old children may not yet have initiated allergic manifestations, this association could be best studied in this age group.

We found that asthma/recurrent bronchitis and allergic reaction on animal contact were diagnosed significantly more often in the Toxocara-seropositive group than in the seronegative group. Moreover, inhaled allergen-specific IgE was observed significantly more often among Toxocara-seropositive than among seronegative individuals. Furthermore, overall eosinophil numbers and total IgE levels were significantly increased in the seropositive group as compared to the seronegative group. However, an analysis of the four subsamples revealed that, although the direction of the differences was the same in all four subgroups, a significant increase in eosinophil numbers existed in rural Eindhoven, and significantly raised serum total IgE concentrations in urban Utrecht and rural Eindhoven.

The observed associations between Toxocara seroprevalence and allergy-related variables, including allergen-specific IgE, strongly suggest that Toxocara stimulates the production of allergen-specific IgE. Since various parasitic infections induce polyclonal B-cell activation [14, 15], it is conceivable that infection with IgE-inducing parasites, including Toxocara, results in nonspecific stimulation of dormant allergic manifestations in children prone to atopy. The causative mechanism may be that both conditions stimulate the immune response in a similar fashion. Allergens and parasite-derived antigens stimulate type 0 T-helper cells to develop into type 2 T-helper cells (Th2), which produce the cytokines interleukin (IL)-4 and IL-5 [23, 24]. IL-4 stimulates IgE production by inducing the B-cell switch from µ to ε expression [24]. IL-5 stimulates eosinophil proliferation and maturation [23]. Under normal conditions, type 1 T-helper cells (Th1), which produce interferon-gamma (IFN-γ), are also activated. IFN-γ is responsible for the downregulation of Th2 cell activity [25]. The balance between the two T-helper cell subpopulations in allergy susceptible individuals may be disturbed in such a way that no downregulation of the Th2 cells occurs.

The ability of Toxocara larvae to survive in their hosts for many months may stimulate Th2 cells and, consequently, IgE production for a longer period. We demonstrated in Toxocara-infected mice that elevated serum total IgE levels lasted for at least 3 months postinfection [8]. Recently, van Ommen et al. [26] demonstrated in a trinitrophenol (TNP)-keyhole limpet haemocyanin (KLH) immunization model, that longstanding elevated IgE levels induced a decrease in antigen-specific IgE, but an increase in serum total IgE. It was demonstrated that parasites, such as Toxocara, induced high serum total IgE levels, of which only a fraction was parasite-specific [14, 15]. Thus, it may be the case that allergen-primed resting B-cell, induced previously to Toxocara infection, and newly Toxocara-primed B-cells are stimulated, simultaneously, to produce allergen-specific IgE and Toxocara-specific IgE, respectively. This supposition was supported by the results of Toxocara-specific IgE estimated in a few selected serum samples from infected and noninfected children participating in the present study.

No relationship was found between the level of total IgE concentration and that of Toxocara-specific IgE in the seropositive samples. MAGNAVAL [27] reported similar observations from patients with toxocarosis in France. We performed a study among children selected on the basis that they (and their relatives) did not have or had not had established allergic asthma. Both groups demonstrated increased IgE levels in the Toxocara-seropositive groups. IgE levels, although increased as compared to the seronegative children, remained within normal limits in the asthma-free group, whereas in the asthma group the already highly increased IgE level in seronegative children was doubled in the seropositive children [28]. It is plausible that Toxocara infection initially induces an IgE increase in a majority of infected individuals, and that only those with unbalanced T-helper cell function demonstrate excessive IgE production.

Our results are in agreement with the observations of Hagel et al. [29] from a study performed in Venezuela. They observed that rural children with low socioeconomic background had a high and diverse parasite burden,
and that urban children from medium to high socioeconomic backgrounds had a low and less diverse parasite burden. Both had highly increased serum total IgE levels, but the rural children presented significantly less often with positive skin tests on allergen challenge than urban children. They proposed that, in the case of high parasite burden, mast cell receptors were saturated with IgE molecules of great diversity. This prevented the bridging of two identical IgE molecules upon allergen challenge and the subsequent histamine release. In the case of low and less diverse parasite burden, the IgE molecules on mast cell receptors would be more homogeneous, allowing the bridging of IgE molecules upon allergen challenge followed by histamine release. Differences between socioeconomic classes in the Netherlands are much smaller than in Venezuela, and all classes are probably comparable to Venezuela's highest socioeconomic class. Children harbouring different helminthic parasites have become rare in the Netherlands [28].

A relationship between eosinophil numbers and Toxocara seroprevalence was demonstrated in the sample, although the increase was only significant in the subpopulation of rural Eindhoven. Eosinophils are tissue cells and migrate to inflammatory sites. In a mouse study, we found that infection caused a rapid increase in blood eosinophils, returning to near normal values after 4 weeks. Eosinophils were still present at inflammatory sites in the tissues (lungs) 3 months after infection [8]. Taylor et al. [7] investigated Toxocara seroprevalence among children reporting with allergic complaints at an outpatient department. Although blood eosinophilia was observed, it was not a consistent characteristic. It is plausible that, during the acute phase, eosinophils migrate from bone marrow to inflamed tissues via the peripheral circulation. When the inflammation has become chronic, the chemotactic stimulus and, thus, the stimulus for eosinophil migration decreases.

In addition to the association between allergy-related phenomena and Toxocara infection, we studied differences in Toxocara seroprevalence among schools, between girls and boys, between urban and rural environments, and between the districts of Utrecht and Eindhoven, and we tried to identify risk factors for Toxocara infection.

In the sample as a whole, significantly more boys than girls had Toxocara antibodies. This difference varied in the four subpopulations, and was even reversed in the city of Eindhoven, though not significantly so. There were no differences between the districts of Utrecht and Eindhoven, nor between urban and rural environments. However, we did find a very large variation among the schools, which is hard to explain. Variations in seropositivity within countries have been reported by other investigators [30–32]. This observation was attributed mainly to factors such as social class [33] and contact with animals. In the present study, no relationship was found between seroprevalence and socioeconomic class, which was investigated in the cities only (results not shown). Rural schools are attended by children irrespective of socioeconomic background, living both in villages and on farms in the neighbourhood. Analysis of apparent risk factors for Toxocara infection yielded no clear relationships. Similar results (including the variability among schools) were obtained in a previous study carried out at 15 schools in two Dutch cities [9]. Although contamination of the outside environment by pets is the main source of infection, environment-related risks analysed in a cross-sectional study did not clarify how children became infected, nor did it explain the large variation among schools. A longitudinal study and investigation into factors such as children’s behaviour when playing, and hygiene of the inside environment at home and at school may provide a better understanding of infection dynamics and merit investigation.

Our study showed that children from urban areas more often than those from rural areas had allergen-specific IgE. One explanation may be that the quality both of indoor and outdoor environment in cities is different from that in rural areas, in such a way that the immune response in susceptible individuals is triggered to produce IgE at a younger age.

In conclusion, an association was demonstrated between Toxocara seroprevalence and various manifestations of allergic disease, including allergic asthma. It was suggested that the mechanisms by which Toxocara stimulates these manifestations were based on a combination of a hereditary tendency for skewing to type 2 T-helper cell function and the longevity of the parasite within its host. Longstanding elevated levels of interleukin-4 may have induced excessive production of polyclonal immunoglobulin E which was directed partly against inhaled allergens. As excessive immunoglobulin E production, which has been shown to be hereditary [21], appears to play a role, we speculate that only children with an atopic predisposition demonstrate an association between Toxocara infection and allergic manifestations.

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