Childhood infections and common carotid intima media thickness in adolescence

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Abstract

Atherosclerotic changes can be measured as changes in common carotid intima media thickness (CIMT). It is hypothesised that repeated infection-associated inflammatory responses in childhood contribute to the atherosclerotic process. We set out to determine whether the frequency of infectious diseases in childhood is associated with CIMT in adolescence. The study is part of the Prevention and Incidence of Asthma and Mite Allergy (PIAMA) population-based birth cohort. At age 16 years, common CIMT was measured. We collected general practitioner (GP) diagnosed infections and prescribed antibiotics. Parent-reported infections were retrieved from annual questionnaires. Linear regression analysis assessed the association between number of infections during the first 4 years of life and common CIMT. Common CIMT measurement, GP and questionnaire data were available for 221 participants. No association was observed between the infection measures and CIMT. In a subgroup analysis, significant positive associations with CIMT were observed in participants with low parental education for 2–3 or ≥7 GP diagnosed infections (+26.4 µm, 95% CI 0.4–52.4 and +26.8 µm, 95% CI 3.6–49.9, respectively) and ≥3 antibiotic prescriptions (+35.5 µm, 95% CI 15.8–55.3). Overall, early childhood infections were not associated with common CIMT in adolescence. However, a higher number of childhood infections might contribute to the inflammatory process of atherosclerosis in subgroups with low education, this needs to be confirmed in future studies.

Introduction

The inflammatory process underlying atherosclerosis is thought to begin in early childhood. Early atherosclerotic changes can be found in infants as young as 9 months of age [1–4]. The classical risk factors, such as exposure to tobacco smoke, high body mass index (BMI) and high cholesterol, may play an important role in the development of early atherosclerotic changes in childhood by inducing inflammation. Yet, these can only explain part of the variation in atherosclerotic changes observed between individuals [5–8].

In the past decades the role of infectious diseases in the pathogenesis of atherosclerosis has been increasingly recognised [9, 10]. It is hypothesised that infections contribute to the inflammatory process underlying atherosclerosis by stimulating the production of inflammatory cytokines and by changing serum levels of high-density lipoprotein cholesterol (HDLc) and low-density lipoprotein cholesterol [9–11]. As such, the numerous infectious disease episodes a child typically experiences in early life may have a cumulative effect on the atherosclerotic process and this may be most pronounced for infections that elicit a systemic inflammatory response [10].

Short-term effects of acute childhood infectious diseases on vasculature and cholesterol have indeed been documented. Studies have reported reduced endothelium-dependent vasodilatation, increased intima media thickness and decreased HDLc in the weeks or months following the infection [11–13]. It is not known to what extent these effects persist into adolescence and adulthood and to what extent these effects are influenced by other mediating factors of atherosclerosis, including sex and socioeconomic status (SES) [14–18].

Infection induced repeated systemic inflammatory responses are best approximated by quantifying the number of febrile infections [10]. While this can be obtained retrospectively...
from parental questionnaires, such data may not be sufficiently reliable and may be subject to recall bias. Alternatively, infection-related hospitalisation data can be used but this only covers severe infections. Longitudinal general practitioner (GP) medical records may provide a suitable alternative data source, as they include prospectively recorded, doctor diagnosed infectious disease episodes as well as any antibiotic treatment [11, 17].

In the participants of the Prevention and Incidence of Asthma and Mite Allergy (PIAMA) study, we set out to determine the role of repeated febrile infections during early childhood in inducing early atherosclerotic changes, as reflected in common carotid intima media thickness (CIMT) levels in adolescence. We hypothesise that infections that trigger a visit to a GP reflect moderate to severe infections, in particular when associated with antibiotic treatment, and are therefore more likely to be associated with an increase in common CIMT compared to parent-reported infections [10]. This study investigated associations between common CIMT levels in adolescents and the number of (1) GP diagnosed febrile infections, (2) antibiotic prescriptions and (3) parent-reported infections in the first 4 years of life. We restricted the exposure period to the first 4 years of life as the infectious disease incidence is generally highest in these early years of childhood [19, 20]. In addition, it was investigated whether these associations were dependent on sex and SES.

**Methods**

**Study population**

This study is part of the PIAMA study, a Dutch population-based birth cohort. The PIAMA study has been described in detail elsewhere [21, 22]. In short, pregnant women were recruited from the general population through antenatal clinics located in the north, center and west of the Netherlands, resulting in a baseline study population of 3963 children born in 1996 and 1997. PIAMA questionnaires were sent to the parents during pregnancy, at age 3 months and thereafter annually until the end of the child until the age of 8 years and included questions on occurrence of infectious diseases. Medical examinations were performed at the ages of 1, 4, 8, 12 and 16 years. At age 16 years, 2159 active participants of two of the three study centres were invited for the medical examination and common CIMT was measured in one of these study centres (Utrecht). At age 18 years, all 3015 active participants in the three study centres (76% of the baseline population) were approached to consent for collection of GP data covering the full 18 years. A total of 1519 (50%) participants gave written informed consent. The medical ethics committees of the participating institutes approved the study protocol.

Eligible for the present study were the participants who were invited for the common CIMT measurement at age 16 years (N=1232), of these participants 60% did not respond or gave no informed consent for the medical examination. In 5.6% of the participants, no IMT measurement was performed due to logistic reasons or measurement errors. The population for analysis included participants with common CIMT measurements and GP data available for at least one of the first 4 years of life (N=221, 18%) (Fig. 1).

**Data collection**

During the medical examination at age 16 years, IMT was measured bilaterally in the distal common carotid artery proximal to the bifurcation at six standard angles (210, 240 and 270 left side and 90, 120, 150 right side) using the automated measurement of the Panasonic CardioHealth® Station (Panasonic Healthcare) by trained research staff. The measurement region was automatically identified by the software and frozen when the accuracy was high; the common CIMT was measured in millimeters over a standard length of 10 mm in the end-diastolic phase. Mean common CIMT was calculated by averaging the common CIMT of the six measurement angles. If <6 measurements were available, mean common CIMT was calculated with the available measurements, the minimum number of measurements was 3. In addition, at the medical examination at age 16 a blood sample was taken and serum total cholesterol (TC) and HDLC were determined enzymatically using Roche automated clinical chemistry analysers (Roche Diagnostics, Indianapolis, IN, USA). The levels of TC and HDLC in mmol/l were used in the analysis to investigate whether cholesterol is a mediator in the association between febrile infections and common CIMT.

GP data were collected by sending letters to the GPs including a questionnaire to obtain participant’s infectious diseases diagnoses and any medication related to infections and allergies. Alternatively, GPs could request on-site data collection by the research team or send an extract of the electronic patient file to the research team. GPs provided a start and end date for the period for which they registered the diagnoses and medication. If a start date was within 6 months after birth and the end date was after 4 years of life, the GP data was considered complete for the first 4 years of life. Complete GP data on at least one of the first 4 years of life was available for 879 (58%) of the 1519 consenting participants and complete information for the first 4 years of life was obtained for 725 (48%).

The number of parent-reported infections as well as indoor smoke exposure in the first 4 years of life were retrieved from
the annual questionnaires. Information regarding education and
allergies of the parents, pre-pregnancy BMI of the mother, birth-
weight of the participant and breastfeeding were retrieved from
the questionnaires completed by the parents during pregnancy
and at age 3 months.

**Definition of exposure potential confounders**

**GP diagnosed infections**

For each participant the number of GP diagnosed infections was
determined by counting the number of GP diagnosed febrile
infections during the first 4 years of life. Febrile infections were
defined as infectious diagnoses according to the International
Classification of Primary Care (ICPC) coding for which a typical
disease course includes one or more days with fever, such as acute
upper or lower respiratory tract infection and urinary tract infec-
tion. A list of ICPC codes and corresponding infectious diagnoses
can be found in Table S1. Similarly, the number of antibiotic
prescriptions in the first 4 years of life was counted including
prophylactic and repeated prescriptions, when two antibiotics were
prescribed on the same day this was counted as one prescription.

**Parent-reported infections**

For parent-reported infections, we used the number of infections
reported in the annual questionnaires during the first 4 years of
life. The number of parent-reported infections in the first 4 years
of life was defined as the number of infections in the child over
the past 12 months, including any severe respiratory tract infec-
tions (infections of the throat, nose and ears, e.g. influenza, pha-
ryngeitis, otitis media, bronchitis, pneumonia and sinusitis) and
occurrence of chickenpox and physician-diagnosed measles or
whooping cough. From these data, an infection count variable
was created for each year of life.

**Confounder definitions**

Sex, parental education, birth weight, breastfeeding, pre-
pregnancy overweight, allergy of the mother and indoor smoke
exposure up to age 4 were included in all analyses as a priori
potential confounders. The selection of confounders was based
on available literature [12, 14–18]. Figure S1 shows a DAG of these
associations. A binary parental education variable was used as a
measure of SES, defining high parental education as completed
higher vocational or university education by at least one parent.
Breastfeeding was categorised into no breastfeeding, ≤16 weeks of
breastfeeding and >16 weeks of breastfeeding. Pre-pregnancy
overweight was defined as a maternal BMI of ≥25 kg/m² before
pregnancy. Maternal allergy was considered positive if a mother
ever had asthma, pet allergy, house dust mite allergy, or nasal
allergy such as hay fever [23]. Exposure to indoor smoke was con-
sidered present when smoking occurred within the home at least
once a week at ages 3 months, 1, 2, 3 or 4 years.

**Statistical analysis**

The incidence rates per year were calculated for the three infec-
tious disease exposure variables investigated in this study, namely
‘number of GP diagnosed infections’, ‘number of antibiotic pre-
scriptions’, and ‘number of parent-reported infections’. These
variables were categorised into four categories of about equal
size based on quartiles in order to limit the influence of outliers
and to allow for a non-linear association.

The associations between each of these exposure variables and
common CIMT as a continuous outcome variable were investi-
gated using separate linear regression models. All a priori defined
potential confounders were added to the models. Additionally, as
previous evidence suggests that the atherosclerotic effect of infec-
tions is perhaps in part mediated through changes in serum cho-
lesterol, we investigated this by adding TC, HDL, or both to the
adjusted models [9–11]. When this resulted in meaningful
changes of >30%, a threshold selected by the authors, in the par-
parameter estimate for the primary exposure, mediation was consid-
ered present. We investigated whether the association between
number of infections and common CIMT was dependent on
SES or sex by assessing the presence of significant interaction in
the adjusted models as this has been shown by previous studies
[16, 18]. In addition, in explorative post-hoc analyses it was inves-
tigated whether the other potential confounders in our model
showed significant interaction with number of infections. When the
P-value of the interaction term was <0.10, interaction was consid-
ered present.

To prevent bias in the parameter estimates, missing values for
confounders and number of infections were imputed for partici-
pants with GP data available for at least one of the first 4 years of
life and a successful common CIMT measurement (N = 185). We
did not impute data for all 1232 participants invited for the med-
ical examination at age 16, since this may introduce bias due to
the large amount of missing data that would have been imputed.
The imputation model included confounders, outcome and pre-
dictors of childhood infections, namely day care attendance, pres-
ence of older siblings in the household and paternal allergy. We
imputed missing values using Multivariate Imputation by Chained
Equations (MICE). Data were imputed using the Random Forest
method.

Analyses were performed using SPSS version 24.0.0.1 (IBM
Corp., Armonk, New York) and RStudio version 1.0.143 (RStudio,
Boston, Massachusetts). The confidence intervals around the inci-
dence rates were calculated using OpenEpi (Open Source
Epidemiologic Statistics for Public Health, version 3.01) [24].

**Results**

Of 1232 PIAMA participants invited for the medical examination
at age 16 years, 489 (40%) participated and IMT was successfully
measured in 420 (34%). Of these participants, 221 had GP data
available for at least one out of the first 4 years of life and were
included in the current analysis (Fig. 1). GP data were complete
for all 4 years in 185 participants and parent-reported infectious
disease data were available for 207 participants (Table 1).

Table 1 shows the characteristics of the population for analysis;
characteristics of the participants eligible for the current study can
be found in Table S2. The mean common CIMT was 465 μm at a
mean age of 16.3 years. Multiple imputation did not meaningfully
change the distribution of the characteristics of the study popula-
tion as shown in Table 1. The mean incidence of GP diagnosed
infections up to age 4 years was 1.18 per year after imputation
(95% CI 1.12–1.26) and 1.45 (95% CI 1.37–1.53) for parent-
reported infections (Table 2). For further analyses, the imputation
datasets were used.

In the total study population no significant associations were
found between any of the cumulative incidence measures of child-
hood infections and common CIMT (Table 3). The adjusted
models included all a priori defined potential confounders and
one of the infection measures, adding TC and/or HDLC did
not result in meaningful changes of the parameter estimate for any of the infection measures (Table S3).

The number of antibiotic prescriptions and GP diagnosed infections showed statistically significant interaction with parental education for the association with common CIMT, therefore, an exploratory analysis was performed stratifying the results according to education level (Fig. 2). The number of GP diagnosed infections and antibiotic prescriptions during the first 4 years were positively associated with common CIMT in adolescence in participants with a low level, but not in those with a high level of parental education. Low parental education participants with 2–3 or ≥7 GP diagnosed infections had a +26.4 µm and +26.8 µm higher common CIMT, respectively, than those with 0 or 1 GP diagnosed infections. Also, participants with ≥3 prescriptions had a +35.5 µm higher common CIMT than those without antibiotic prescriptions. No significant association between 4 and 6 GP diagnosed infections and common CIMT was observed for low parental education participants. None of the infectious disease exposures showed statistically significant interaction with sex. P-values of the interaction terms for the other confounders can be found in Table S4.

**Table 1.** Characteristics of study population with common carotid intima media thickness (IMT) measurement and general practitioner data on at least one of the first 4 years of life

<table>
<thead>
<tr>
<th>Study population before MI</th>
<th>Study population after MI, N = 221</th>
</tr>
</thead>
<tbody>
<tr>
<td>Complete parent-reported data on the first 4 years of life</td>
<td>93.7 % 207</td>
</tr>
<tr>
<td>Complete GP data on the first 4 years of life</td>
<td>83.7 % 185</td>
</tr>
<tr>
<td>Sex</td>
<td>221</td>
</tr>
<tr>
<td>Male</td>
<td>47.1 %</td>
</tr>
<tr>
<td>Female</td>
<td>52.9 %</td>
</tr>
<tr>
<td>Overweight mother before pregnancy</td>
<td>212</td>
</tr>
<tr>
<td>No</td>
<td>84.9 % 83.3 %</td>
</tr>
<tr>
<td>Yes</td>
<td>15.1 % 16.7 %</td>
</tr>
<tr>
<td>Parental education</td>
<td>220</td>
</tr>
<tr>
<td>Low</td>
<td>34.1 % 34.3 %</td>
</tr>
<tr>
<td>High</td>
<td>65.9 % 65.7 %</td>
</tr>
<tr>
<td>Allergic mother</td>
<td>221</td>
</tr>
<tr>
<td>No</td>
<td>68.3 % 68.3 %</td>
</tr>
<tr>
<td>Yes</td>
<td>31.7 % 31.7 %</td>
</tr>
<tr>
<td>Exposure to any smoking indoors first 4 years of life</td>
<td>216</td>
</tr>
<tr>
<td>No</td>
<td>75.0 % 75.2 %</td>
</tr>
<tr>
<td>Yes</td>
<td>25.0 % 24.8 %</td>
</tr>
<tr>
<td>Breastfeeding</td>
<td>221</td>
</tr>
<tr>
<td>No breastfeeding</td>
<td>10.9 % 10.9 %</td>
</tr>
<tr>
<td>&lt;16 weeks of breastfeeding</td>
<td>43.9 % 43.9 %</td>
</tr>
<tr>
<td>&gt;16 weeks of breastfeeding</td>
<td>45.2 % 45.2 %</td>
</tr>
<tr>
<td>Mean</td>
<td>s.e. (N)</td>
</tr>
<tr>
<td>Birth weight gram</td>
<td>3548 37.3 (220)</td>
</tr>
<tr>
<td>IMT in µm age 16 years*</td>
<td>465 2.6 (221)</td>
</tr>
<tr>
<td>Total cholesterol mmol/l age 16 years*</td>
<td>3.84 0.05 (209)</td>
</tr>
<tr>
<td>HDL cholesterol mmol/l age 16 years*</td>
<td>1.34 0.02 (209)</td>
</tr>
<tr>
<td>Systolic blood pressure mm Hg age 16 years*</td>
<td>116 0.64 (220)</td>
</tr>
<tr>
<td>Body mass index kg/m² age 16 years*</td>
<td>20.75 0.18 (221)</td>
</tr>
</tbody>
</table>

GP, general practitioner; IMT, intima media thickness of common carotid artery; MI, multiple imputation; s.e., standard error.

\*Variable was not imputed.

**Discussion**

Overall, we observed no association between number of infections or antibiotic prescriptions during the first 4 years of life and common CIMT in adolescence. Analysis in strata of low and high parental education showed that a higher frequency of GP diagnosed
childhood infections and antibiotic prescriptions were associated with an increased common CIMT at age 16 years only for participants with a low parental education level.

We hypothesised that repeated febrile infections severe enough to warrant a GP consultation, and in particular when requiring antibiotic treatment, would be associated with increased common CIMT in adolescence due to the repeated inflammatory responses elicited by the infections [10, 15–17]. This hypothesis was not confirmed in the total study population of the current study and findings of previous studies have been inconsistent [11, 15–17]. This hypothesis was not confirmed in the total study population of the current study and findings of previous studies have been inconsistent [11, 15–17]. This hypothesis was not confirmed in the total study population of the current study and findings of previous studies have been inconsistent [11, 15–17]. This hypothesis was not confirmed in the total study population of the current study and findings of previous studies have been inconsistent [11, 15–17]. 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between persons with low and high SES could potentially explain the inconsistent findings of previous studies if they did not assess such interactions.

Similarly, in participants with lower educated parents we found that the number of infections treated with antibiotics in the first 4 years of life was positively associated with common CIMT at age 16 years. An antibiotic prescription could be a marker of more severe infection, which might explain the larger effect estimate for antibiotic prescriptions compared with GP diagnosed infections. This is in line with the study of Evelein et al., which reported an association between antibiotic prescriptions and an increase in common CIMT [17].

The current study has some limitations, for instance parent-reported infections were mostly limited to respiratory tract infections. However, as respiratory tract infections are the most common infections in childhood, we consider it unlikely that this limitation has influenced the results. The study population of the current study consisted mostly of Dutch participants. The development of atherosclerosis is suggested to be influenced by ethnicity, but whether the effects of infections, if any, are different for different ethnic groups is unknown [36, 37]. The current study investigated a subsample of the PIAMA study population due to data availability limitations, however when comparing Table 1 with Table S2, the differences in participant characteristics are limited. Therefore, we consider it unlikely that our results would have been different when we would have been able to analyse the entire eligible population. The percentage of participants with a high SES, as defined by educational level of the parents, was higher than the percentage with low SES in this study. As a low SES is associated with a higher prevalence of childhood infections and an increased susceptibility to infections, this might have influenced the strength of the associations reported in this study and could explain the absence of a significant association for the 4–6 GP diagnosed infections category in participants with low parental education [18, 34, 35, 38]. Due to sample size restrictions the subgroup analyses have limited statistical power and should be interpreted with caution. There is a chance for false positive findings in small samples, however the chance of not finding any interaction would presumably be higher. In addition, statistically significant interaction was observed for multiple infectious disease exposures, therefore we consider it unlikely that the reported interaction is a chance finding, however, the results of the current study need to be confirmed in future studies.

**Conclusion**

The study suggests that at population level the number of early childhood infections is not associated with the development of early atherosclerotic changes in adolescence. In participants with lower parental education, GP diagnosed and antibiotic treated infectious diseases, probably indicative of more severe infections were associated with increased common CIMT at age 16 years. Although it is based on relatively small number, this suggests that infections might contribute to other (inflammatory) processes, ultimately resulting in increased common CIMT and atherosclerosis, only in subgroups of the population. As the development of atherosclerosis can start in early childhood, identifying determinants of this early development is important for the recognition of potential risk populations as well as potential targets for prevention.

**Supplementary material.** The supplementary material for this article can be found at [https://doi.org/10.1017/S095026881800287X](https://doi.org/10.1017/S095026881800287X)

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