Infection with Soil-Transmitted Helminths Is Associated with Increased Insulin Sensitivity

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Abstract

Objective
Given that helminth infections have been shown to improve insulin sensitivity in animal studies, which may be explained by beneficial effects on energy balance or by a shift in the immune system to an anti-inflammatory profile, we investigated whether soil-transmitted helminth (STH)-infected subjects are more insulin sensitive than STH-uninfected subjects.

Design
We performed a cross-sectional study on Flores island, Indonesia, an area with high prevalence of STH infections.

Methods
From 646 adults, stool samples were screened for Trichuris trichiura by microscopy and for Ascaris lumbricoides, Necator americanus, Ancylostoma duodenale, and Strongyloides stercoralis by qPCR. No other helminth was found. We collected data on body mass index (BMI, kg/m²), waist-to-hip ratio (WHR), fasting blood glucose (FBG, mmol/L), insulin (pmol/L), high sensitive C-reactive protein (ng/ml) and Immunoglobulin E (IU/ml). The homeostatic model assessment for insulin resistance (HOMAIR) was calculated and regression models were used to assess the association between STH infection status and insulin resistance.
Results

424 (66%) participants had at least one STH infection. STH infected participants had lower BMI (23.2 vs 22.5 kg/m², p value = 0.03) and lower HOMAIR (0.97 vs 0.81, p value = 0.05). In an age-, sex- and BMI-adjusted model a significant association was seen between the number of infections and HOMAIR: for every additional infection with STH species, the HOMAIR decreased by 0.10 (p for linear trend 0.01). This effect was mainly accounted for by a decrease in insulin of 4.9 pmol/L for every infection (p for trend = 0.07).

Conclusion

STH infections are associated with a modest improvement of insulin sensitivity, which is not accounted for by STH effects on BMI alone.

Introduction

The prevalence of type 2 diabetes (T2DM) is rising in low-to-middle income countries (LMIC). The explanations for these trends are complex and multifactorial, but have traditionally been attributed to a shift in infrastructure, technology and food supply that promotes over-nutrition and sedentary lifestyles [1, 2]. There is now accumulating evidence that in addition to a disturbed energy balance, inflammation plays a role in T2DM [3]. Indeed, in T2DM, elevated levels of inflammation-related markers such as interleukin 6 (IL6), IL8, tumor necrosis factor (TNF), and C-reactive protein (CRP) have been reported [3]. The adoption of a Western lifestyle in LMIC is often paralleled by decreased burden of infectious diseases, including helminth infections. Studying these trends in the context of the epidemiological transition theory and the hygiene hypothesis may have important implications for clinical practice, global health policy, and future research within epidemiology [4].

In animal models, helminth infections can enhance glucose tolerance, probably by inducing eosinophilia and preventing obesity [5]. Interestingly, helminth infections have been shown to induce T helper (Th) 2 cells [6–8] and anti-inflammatory immune responses [9–12]. It has been hypothesized that chronic helminth infections decrease systemic inflammation [13] and might be beneficial for the prevention of inflammatory diseases, such as allergy [11,14], inflammatory bowel disease [15], and T2DM [5,10,16]. Based on this information, it could be hypothesized that helminth infections may also have a beneficial influence on glucose metabolism both by preventing obesity and by anti-inflammatory immune responses [16]. In several epidemiological studies, an inverse association between the prevalence of T2DM or metabolic syndrome and helminth infections was found [17–19].

In the present study, we investigated the relationship between helminth infections and insulin resistance, as assessed by homeostatic model assessment for insulin resistance (HOMAIR) [20] in an area endemic for STH on Flores Island, Indonesia. In addition, we studied whether the potential association is explained by changes in body mass index (BMI).

Subjects and Methods

Study objectives

The primary objective of the study was to investigate the relationship between STH infections and HOMAIR in adults. Our hypothesis is that HOMAIR is lower in subjects with STH
infections than in subjects without STH infections. This hypothesis is based on the association between helminth infections and decreased food intake, digestion or nutrient absorption, which will lead to lower BMI [21–23] as well as the proposed effects of helminth infections on systemic inflammation [13]. We studied to what extent the potential relationship between STH infections and HOMAIR is explained by differences BMI.

Study population

The study area is Nangapanda on Flores Island in Indonesia, which is part of East Nusa Tenggara province. The area has a low socioeconomic status according to the recent Indonesian health survey (RISKESDAS 2013). Based on the same survey, prevalence of T2DM in this area was 3.3% in 2013. Previous reports from the area [24,25], have indicated that prevalence of metabolic syndrome according to ATPIII criteria is 11.8% (7.4% male, 14.4% female) [26,27]. Main sources of income are farming, fishing, woving and stone collection with a diet mainly consisting of freshly grown vegetables and fresh fish [28]. The area is highly endemic for STH, whereas no evidence for other helminth species has been found [28,29]. In Nangapanda area, a large investigational project is being conducted on the relationship between STH infections and the immune system (ImmunoSPIN study [28,29]). For the current study, a cross sectional sample was included from all inhabitants aged 18 years and above. Data were collected between May-August 2009.

Study design

From 1841 inhabitants aged 18 years and above in Nangapanda who participated in the ImmunoSPIN project, 646 subjects from whom stool samples were available, were invited to participate in the present cross-sectional study for collection of data on anthropometrics and laboratory measurements. 584 subjects, from whom data on STH infections, BMI, waist-to-hip ratio (WHR) and laboratory measurements were available were included in the present analysis.

The study was approved by the ethical committee of the Faculty of Medicine, University of Indonesia (EC-FMUI), ref: 194/PT02.FK/Etik/2006 with addendum ref: 96/PT02.FK/Etik/2010 and registered as clinical trial ref: ISRCTN83830814 and was filed by the Leiden University Medical Center Committee of Medical Ethics (CME). Because of the high rate of illiteracy amongst elderly participants, either written or verbal informed consent (recorded as signed or with a given thumb-print) was obtained from each participant after explanation of the study and the voluntary nature of the participation.

Clinical and laboratory assessments

Anthropometric measurements of body weight (SECA 761, SECA GMBH & Co. Kg., Hamburg, Germany), height (SECA 206, SECA GMBH & Co. Kg., Hamburg, Germany), waist and hip circumference (SECA 203, SECA GMBH & Co. Kg., Hamburg, Germany) were performed according to the NHLBI practical guidelines (NHLBI web: http://www.nhlbi.nih.gov) by a team of trained researchers. BMI was calculated as weight in kg divided by square of height in meter; WHR was calculated as waist circumference in cm divided by hip circumference in cm.

Participants were instructed to fast after 8 pm the day before blood collection. Fasting blood glucose (FBG) was analyzed using Breeze2 glucose meter (Bayer Health Care LLC, Basel, Switzerland). Insulin was measured using MSD 96-Well MULTI-ARRAY Human insulin assay (Meso Scale Discovery, Gaithersburg, USA). HOMAIR, a well-validated measure of IR, HOMAIR = fasting serum insulin x fasting glucose / 22.5, was calculated to estimate insulin resistance using HOMA2 calculator (https://www.dtu.ox.ac.uk/homacalculator/) [20]. High
sensitive C-reactive protein (HsCRP) level was measured using MSD 96-Well MULTI-ARRAY CRP Assay (Meso Scale Discovery, Gaithersburg, USA).

Whole blood cytokine production after stimulation with *Escherichia coli* lipopolysaccharide (LPS) was performed as described previously [28]. Briefly, heparinized blood was diluted 4x and stimulated within 6 hours after drawing with medium alone or *E. coli* LPS, 1 ng/L Sigma-Aldrich, Zwijndrecht, The Netherlands) and incubated for 24 hours at 37°C and 5% CO₂. The supernatants were frozen at -20°C and TNF and IL10 supernatants were assessed by means of immunobead-based multiplex assays on a Liquichip 200 Workstation (Qiagen, Venlo, The Netherlands) using Liquichip analyzer software (Qiagen, Venlo, The Netherlands). Samples with TNF levels higher than 250 pg/mL in medium stimulation (unstimulated blood) were excluded from further analyses (2 samples) as they are considered unreliable. Immunoglobulin E (IgE) level was measured by an ELISA [30].

Assessment of soil-transmitted helminth infection

Stool samples were collected and preserved in 4% formaldehyde for microscopy examination or frozen (-20°C) unpreserved for PCR detection. The formol-ether acetate concentration method was performed on the formalin preserved stool samples followed by microscopy examination for eggs of STH [28]. For this paper only microscopy results on *Trichuris trichiura* infection were used as no PCR technique yet available. As described in detail before [28], DNA was isolated from approximately 100 mg unpreserved feces and a multiplex real-time PCR for the detection of *Acaris lumbricoides*, *Necator americanus*, *Ancylostoma duodenale*, and *Strongyloides stercoralis* was performed. The real-time PCR output from this system consisted of a cycle-threshold (CT) value, representing the amplification cycle in which the level of fluorescent signal exceeds the background fluorescence, and reflecting the parasite-specific DNA load in the sample tested. Negative and positive control samples were included in each run of the amplification. We have indeed the result of stool cultured (Harada Mori) for detection of hookworm larvae. We have chosen to use PCR results whenever possible because this technique is proven to be very sensitive and specific [31]. We defined a positive case for *T. trichiura* by presence of eggs in stool samples and for *A. lumbricoides*, *N. americanus*, *A. duodenale* and *S. stercoralis* by parasite-specific DNA amplification. Participants were also stratified by number of STH species infections.

Statistical analysis

Normally distributed continuous data were reported as mean and standard deviation. Normal distribution was assessed by verifying data distribution in a histogram graph relative to a normal distribution line. Non-normally distributed continuous data were expressed as median and interquartile range (insulin, HOMAIR, hsCRP, cytokines, IgE). Categorical data were expressed as proportions. Non-normally distributed data were log-transformed for analyses.

Differences in study parameters between subjects with and without any STH infection were analysed by linear regression. In addition, we also analysed the potential association between the number of helminth species per subject and HOMAIR. In all analyses, we adjusted for age and gender. To assess whether potential differences or associations are mediated through an effect of STH on BMI, we also adjusted for BMI in a separate analysis. Differences in immune parameters and HOMAIR between infected and uninfected participants were reported as mean differences with 95% confidence intervals (95% CI). P values <0.05 were considered to be statistically significant. In a supplementary analysis (S1 Table), we investigated the relationship between BMI categorized according to reference values for Asian subjects [32] and FBG,
insulin and HOMAIR. Normality test (histogram and Shapiro–Wilk test) and statistical analyses were performed with SPSS 17.0.2 (SPSS Inc., Chicago, Illinois, The USA).

### Results

#### Characteristics of study participants

A total of 424 participants who were infected with at least one species of STH were compared to 222 uninfected participants (Table 1). Gender and age distribution were comparable between the 2 groups. The most prevalent STH species among the study subjects were *N. americanus* (51.7%), *A. lumbricoides* (21.8%) and *T. trichiura* (19.7%). The proportion of participants infected with *A. duodenale* (3.7%) and with *S. stercoralis* (0.6%) was clearly lower. 261 participants were infected with one STH species only, 124 with two STH species and 39 with 3 or more STH species. Intestinal protozoa found in the participants were *Blatocystis hominis* 15 (2.3%), *Entamoeba hystolitica* 6 (0.9%), *Entamoeba coli* 37 (5.7%), *Giardia lamblia*

| Table 1. Characteristics of the study population. |
|--------------------------------------------------|
| Whole study population | No infection with soil transmitted helminths | Infection with soil-transmitted helminths | P-value for difference infected vs non-infected |
|------------------------|---------------------------------------------|------------------------------------------|------------------------------------------------|
| (n = 646)              | (n = 222)                                   | (n = 424)                                |                                                 |
| Age (year) (mean, SD)  | 44.9 (13.9)                                | 44.4 (13.2)                              | 45.2 (14.2)                                   | 0.48                                          |
| Female, N (%)          | 410 (63.5)                                 | 147 (66.2)                               | 263 (62.0)                                   | 0.29                                          |
| *Trichuris trichiura* N (%) | 127 (19.7)                                | 127 (30.0)                               |                                                 |                                               |
| *Ascaris lumbricoides* N (%) | 141 (21.8)                                | 141 (33.3)                               |                                                 |                                               |
| *Necator americanus* N (%) | 334 (51.7)                                | 334 (78.8)                               |                                                 |                                               |
| *Ancylostoma duodenale* N (%) | 24 (3.7)                                  | 24 (5.7)                                 |                                                 |                                               |
| *Strongyloides stercoralis* N (%) | 4 (0.6)                                   | 4 (0.9)                                  |                                                 |                                               |
| Single STH species infection N (%) | 261 (40.4) | 261 (61.6)                               |                                                 |                                               |
| Two STH species infecton N (%) | 124 (19.2) | 124 (29.2)                               |                                                 |                                               |
| Three or more STH species infection N (%) | 39 (6.0) | 39 (9.2)                                  |                                                 |                                               |
| BMI (Kg/m²) (mean, SD) | 22.7 (3.8)                                 | 23.2 (3.7)                               | 22.5 (3.8)                                   | 0.03                                          |
| WHR (mean, SD)         | 0.88 (0.07)                                | 0.89 (0.07)                              | 0.88 (0.06)                                   | 0.08                                          |
| FBG (mmol/L) (mean, SD) | 5.90 (1.6)                                | 5.92 (1.5)                               | 5.88 (1.6)                                   | 0.76                                          |
| Insulin (pmol/L) (mean, SD) | 46.5 (55.3) | 49.5 (43.7)                               | 45.0 (60.1)                                   | 0.40                                          |
| HOMAIR (Mean, SD)      | 0.86 (0.86)                                | 0.97 (0.84)                              | 0.81 (0.86)                                   | 0.05                                          |
| HsCRP (ng/ml) (Median, interquartile range) | 469 (185–1354) | 437 (159–1435)                           | 488 (202–1344)                               | 0.88*                                         |
| TNF (pg/ml) (Median, interquartile range) | 304 (146–547) | 279 (138–510)                            | 326 (152–592)                                 | 0.07*                                         |
| IL10 (pg/ml) (Median, interquartile range) | 133 (74–231) | 145 (82–239)                             | 131 (74–230)                                 | 0.75*                                         |
| IgE (IU/ml) (Median, interquartile range) | 964 (514–2073) | 755 (481–1625)                  | 1105 (540–2248)                              | 0.002*                                         |

Abbreviations: BMI = body mass index, WHR = waist to hip ratio, FBG = fasting blood glucose, HOMAIR = Homeostasis model assessment for insulin resistance, HsCRP = High sensitive C reactive protein, TNF = tumor necrosis factor, IL10 = interleukin 10, IgE = Immunoglobulin E.

*after logarithmic transformation

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These proportions were considered too low to be used for further statistical analyses. BMI distribution and the relationship with FBG and HOMAIR is provided in a S1 Table.

Association between STH infection and glucose metabolism (Table 2)

BMI and WHR were lower in infected (22.5 kg/m² and 0.88 respectively) than in uninfected participants (23.2 kg/m² and 0.89), which were independent of age and sex. No differences in FBG glucose concentration were present between subjects with and without STH infection. In comparison with uninfected participants, participants with any STH infection had a trend towards lower HOMAIR but this difference was not statistically significant (mean difference 0.15, p = 0.06).

We found an association between number of STH species per subject and BMI, with a decrease of 0.3 kg/m² for every increase in the number of species (p for linear trend = 0.04, Table 3); a similar association was found for WHR (p for linear trend = 0.01). In an age-, sex- and BMI-adjusted model an association was found between the number of STH species per

Table 2. Parameters of glucose metabolism parameters in soil-transmitted helminth uninfected and infected participants.

|                  | Mean difference adjusted for age and sex (95% confidence interval) | Mean difference adjusted for age, sex and BMI (95% confidence interval) | Trend analysis # adjusted for age, sex and BMI (95% confidence interval) |
|------------------|---------------------------------------------------------------------|-----------------------------------------------------------------------|-----------------------------------------------------------------------|
| BMI (Kg/m2)      | -0.6 (-1.2, -0.02), p = 0.04                                        | -0.3 (-0.7, -0.02), p = 0.04                                          |                                                                       |
| WHR*             | -0.01 (-0.02, -0.001), p = 0.02                                      | -0.007 (-0.012, -0.002), p = 0.01                                     |                                                                       |
| FBG (mmol/L)     | -0.05 (-0.3, 0.2), p = 0.7                                           | 0.01 (-0.2, 0.3), p = 0.9                                             |                                                                       |
| Insulin (pmol/L) | -4.2 (-14.7, 6.2), p = 0.4                                           | -1.5 (-11.5, 8.5), p = 0.8                                            |                                                                       |
| HOMAIR (index**) | -0.15 (-0.32, 0.01), p = 0.06                                        | -0.10 (-0.25, 0.05), p = 0.2                                          |                                                                       |

Abbreviations: BMI = body mass index, WHR = waist to hip ratio, FBG = fasting blood glucose, HOMAIR = Homeostasis model assessment for insulin resistance.

* WHR is calculated by waist circumference (cm) / hip circumference (cm)

** HOMAIR index is calculated with HOMAIR formula = fasting serum insulin x fasting glucose / 22.5, using HOMA2 calculator (https://www.dtu.ox.ac.uk/homacalculator/)

# The difference is expressed as increase or decrease in the parameter per increasing number of helminth species per patient (maximum = 3). Insulin and HOMAIR were log-transformed.

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Table 3. Immune parameters in soil-transmitted helminth uninfected and infected participants.

|                  | Mean difference adjusted for age and sex (95% confidence interval) | Mean difference adjusted for age, sex and BMI (95% confidence interval) | Trend analysis # adjusted for age, sex and BMI (95% confidence interval) |
|------------------|---------------------------------------------------------------------|-----------------------------------------------------------------------|-----------------------------------------------------------------------|
| HsCRP (ng/ml)    | -0.00 (-0.13, 0.12), p = 0.96                                       | 0.10 (-0.11, 0.13), p = 0.88                                          | -0.02 (-0.08, 0.05), p = 0.60                                         |
| (n = 496)        |                                                                     |                                                                       |                                                                       |
| TNF (pg/ml)      | 0.10 (-0.00, 0.21), p = 0.06                                        | 0.10 (-0.01, 0.21), p = 0.07                                          | 0.06 (0.00, 0.12), p = 0.04                                          |
| (n = 346)        |                                                                     |                                                                       |                                                                       |
| IL10 (pg/ml)     | -0.01 (-0.10, 0.08), p = 0.82                                        | -0.02 (-0.11, 0.08), p = 0.74                                         | -0.01 (-0.06, 0.04), p = 0.65                                        |
| (n = 346)        |                                                                     |                                                                       |                                                                       |
| IgE (IU/ml)      | 0.15 (0.06, 0.24), p = 0.002                                        | 0.15 (0.05, 0.24), p = 0.002                                          | 0.01 (0.05, 0.15), p<0.0001                                          |
| (n = 510)        |                                                                     |                                                                       |                                                                       |

Abbreviations: HsCRP = High sensitive C reactive protein, TNF = tumor necrosis factor, IL10 = interleukin 10, IgE = Immunoglobulin E. HsCRP, TNF, IL10 and IgE are log-transformed.

# The difference is expressed as increase or decrease in the parameter per increasing number of infections per patient (maximum = 3).

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subject and HOMAIR: for every additional STH species, HOMAIR decreased by 0.10 (p for linear trend = 0.01) (Table 3, Fig 1). No clear associations were found between STH infections and hsCRP, TNF or IL10 after whole blood stimulation with LPS (TNF-LPS and IL10-LPS). We found a small but positive association between TNF-LPS with increasing number of STH species per subject with the highest TNF-LPS-levels in participants with 3 or more infections (Table 3). This association was not changed after adjustment for age, sex and BMI. No clear association was found between IgE levels and HOMAIR (p = 0.99, crude, p = 0.87, adjusted for age and sex, p = 0.60, adjusted for age, sex and BMI).

We also investigated whether there were differences in the association between the species of STH infections and HOMAIR. We found no individual associations between N. americanus, A. lumbricoides, or T. trichiura (prevalence of S. stercoralis and A. duodenale were too low to be considered), with HOMAIR, although these analyses might lack sufficient statistical power.

**Discussion**

In this study we investigated the relationship between infection with STH and insulin resistance in a population residing in an area highly endemic for STH. The hypothesis being tested is that STH infections may have a beneficial effect on glucose metabolism, by influencing BMI or their ability to skew immune responses to Th2 and an anti-inflammatory profile. The beneficial influence of STH infections on glucose metabolism has been shown in animal models of T2DM [5]. Mice on high fat diet that were subsequently infected with helminths, became less obese and less insulin resistant which seemed to be in conjunction with maintenance of alternative activated macrophages in adipose tissues [5,16].

In our study, we observed a trend towards lower HOMAIR in subjects with STH infection as compared with uninfected subjects. Furthermore, we found that infection with an increasing number of STH species incrementally enhanced insulin sensitivity. This effect may be partly...
explained by the lower BMI in STH infected subjects. It should be noted that BMI reference
values in Asia are different from Western countries [32]. It is well known that a relationship ex-
ists between helminths and energy metabolism. This relationship may be multifactorial and in-
clude changes in digestion and decreased absorption of nutrients [21,33]. However, after
adjustment for BMI, the negative association between STH infections and HOMAIR persisted,
indicating that this association cannot be explained by effects of STH infections on BMI alone.
In the analysis using BMI reference values in Asia, we found a significant positive association
between the BMI classification and FBG, insulin level and HOMAIR, as expected.

It is tempting to speculate that STH infections in our study may have influenced systemic
inflammation which would then lead to improved glucose tolerance. However we found no in-
dication for differences in systemic inflammation between subjects with and without
STH infections.

The fact that we did not find statistically significant differences in IL10 levels between in-
fected and non-infected subjects points to the complexity of IL10 regulation, IL10 for instance
can be inhibited by insulin, which can have implications for obesity [34].

It should be noted that the magnitude of the effect of STH on HOMAIR in our study is
modest. The reduction of 13% in HOMAIR by presence of STH, might reflect the fact that the
population is lean and insulin sensitive. Indeed, the low HOMAIR value in our study popula-
tion as well as the lower BMI and WHR represent the rural character of the area with accompa-
nying healthier lifestyle [35,36].

Alternatively, the difference between currently STH infected and uninfected subjects might
be small due to the possibility that the energy balance and immune profiles could still be modu-
lated by previous infections, recent deworming treatment, or sub-microscopic infection. We
chose to analyse *A. lumbricoides*, Hookworm (*N. americanus* and *A. duodenale*) and *Strongylo-
ides stercoralis* using qPCR, a method that is more sensitive and reliable than microscopy
[31]. However, as we do not have an optimized qPCR method to detect *T. trichiura*, while the
species is also endemic in the area, we measured *T. trichiura* infection using the microscopy
method. The fact that we were not able to verify these factors is a limitation of the study.

As mentioned above, STH infection can modulate the host’s immune system but this was
not clearly shown in our study participants. An interesting additional explanation may be that
that STH affect the gut microbiome which on its turn will also affect the host immune system
[37–41].

We acknowledge the limitations of this cross-sectional study, which prevent conclusions on
causal relationships between STH infections and T2DM. The population studied was not at
risk of T2DM, so potential effects of STH on insulin sensitivity might have been difficult to de-
tect. In addition, no consistent effects of STH on the immune system were found. Furthermore,
no data on the history and intensity of helminth infections nor history of deworming are pres-
ent, which prevents strong conclusions on the relationship between STH infections, the im-
mune system and insulin sensitivity. Furthermore, the fact that only participants were included
who provided stool samples may also have led to selection bias.

Using HOMAIR, which is measured in fasting state, may fail to detect early disturbances in
insulin sensitivity, which is typically a post-prandial disturbance [42]. An oral glucose tolerance
test may therefore have been worthwhile to perform.

It is clear that further and specifically designed investigations are needed to clarify the rela-
tionship between innate and adaptive immune responses, inflammation, STH infections and
insulin sensitivity in humans.

We acknowledge that the pathogenesis of insulin resistance and diabetes is multifactorial
and extremely complex. However, we believe that our study provides interesting data that add
a layer of complexity that needs to be taken into account in the relationship between rural-urban transition in LMIC and the development of T2DM.

Supporting Information
S1 Table. Body mass index of study participants in relation to fasting blood glucose, insulin and HOMAIR.

(DOCX)

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Author Contributions
Conceived and designed the experiments: AEW FP ES TS MY JWAS. Performed the experiments: AEW FH LJW MAP. Analyzed the data: AEW LM OMD MY JWAS. Contributed reagents/materials/analysis tools: MMMK JJV. Wrote the paper: AEW JWAS. Reviewed the paper: ES OMD BG MY JWAS.

References
1. Chan JCN, Malik V, Jia W, Kadowaki T, Yajnik CS, Yoon K-H, et al. Diabetes in Asia: epidemiology, risk factors, and pathophysiology. JAMA. 2009; 301: 2129–2140. doi:10.1001/jama.2009.726 PMID: 19470990
2. Ramachandran A, Ma RCW, Snehalatha C. Diabetes in Asia. Lancet. 2010; 375: 408–418. doi: 10.1016/S0140-6736(09)60937-5 PMID: 19875164
3. Shoelston SE, Lee J, Goldfine AB. Inflammation and insulin resistance. J Clin Invest. 2006; 116: 1793–1801. PMID: 16823477
4. Zuckerman MK, Harper KN, Barrett R, Armelagos GJ. The evolution of disease: anthropological perspectives on epidemiologic transitions. Glob Health Action. 2014; 7: 23303. doi: 10.3402/gha.v7.23303 PMID: 24848652
5. Wu D, Molofsky AB, Liang H-E, Ricardo-Gonzalez RR, Jouihan HA, Bando JK, et al. Eosinophils sustain adipose alternatively activated macrophages associated with glucose homeostasis. Science. 2011; 332: 243–247. doi: 10.1126/science.1201475 PMID: 21436399
6. Chen F, Liu Z, Wu W, Rozo C, Bowdridge S, Millman A, et al. An essential role for Th2-type responses in limiting acute tissue damage during experimental helminth infection. Nat Med. 2012; 18: 260–266. doi:10.1038/nm.2628 PMID: 22245779
7. Everts B, Perona-Wright G, Smits HH, Hokke CH, van der Ham AJ, Fitzsimmons CM, et al. Omega-1, a glycoprotein secreted by Schistosoma mansoni eggs, drives Th2 responses. J Exp Med. 2009; 206: 1673–1680. doi: 10.1084/jem.20082460 PMID: 19635864
8. Jackson JA, Turner JD, Rentoul L, Faulkner H, Behnke JM, Hoyle M, et al. T helper cell type 2 responsiveness predicts future susceptibility to gastrointestinal nematodes in humans. J Infect Dis. 2004; 190: 1804–1811. PMID: 15499537
9. Sawant DV, Gravano DM, Vogel P, Giacomin P, Artis D, Vignali DAA. Regulatory T cells limit induction of protective immunity and promote immune pathology following intestinal helminth infection. J Immunol. 2014; 192: 2904–2912. doi: 10.4049/jimmunol.1202502 PMID: 24532574
10. Wiria AE, Djuardi Y, Supali T, Sartono E, Yazdanbaksh M. Helminth infection in populations undergoing epidemiological transition: A friend or foe? Seminars in Immunopathology. 2012; 34: 889–901. doi:10.1007/s00281-012-0358-0 PMID: 23129304
11. Van den Biggelaar AH, van Ree R, Rodrigues LC, Leli B, Deelder AM, Kremers-R PG, et al. Decreased atopy in children infected with Schistosoma haematobium: a role for parasite-induced interleukin-10. Lancet. 2000; 356: 1723–1727. PMID: 11095260
12. D’Elia R, Behnke JM, Bradley JE, Else KJ. Regulatory T cells: a role in the control of helminth-driven intestinal pathology and worm survival. J Immunol. 2009; 182: 2340–2348. doi:10.4049/jimmunol.0802767 PMID: 19201888

13. Mishra PK, Palma M, Bleich D, Loke P, Gause WC. Systemic impact of intestinal helminth infections. Mucosal Immunol. 2014; 7: 753–762. doi:10.1038/mi.2014.23 PMID: 24736234

14. Endara P, Vaca M, Chico ME, Erazo S, Oviedo G, Quinzo I, et al. Long-term periodic anthelmintic treatments are associated with increased allergen skin reactivity. Clin Exp Allergy. 2010; 40: 1669–1677. doi:10.1111/j.1365-2222.2010.03559.x PMID: 21039971

15. Summers RW, Elliott DE, Thompson RA, Weinstock JV. Trichuris suis therapy for active ulcerative colitis: a randomized controlled trial. Gastroenterology. 2005; 128: 825–832. PMID: 15825065

16. Wiria AE, Sartono E, Supali T, Yazdanbakhsh M. Helminth infections, type-2 immune response, and metabolic syndrome. PLoS Pathog. 2014; 10: e1004140. doi: 10.1371/journal.ppat.1004140 PMID: 24992724

17. Aravindhan V, Mohan V, Surendar J, Muralidhara Rao M, Pavankumar N, Deepa M, et al. Decreased prevalence of lymphatic filariasis among diabetic subjects associated with a diminished pro-inflammatory cytokine response (CURES 83). PLoS Negl Trop Dis. 2010; 4: e707. doi:10.1371/journal.pntd.0000707 PMID: 20559443

18. Nazligul Y, Sabuncu T, Ozbilge H. Is there a predisposition to intestinal parasitosis in diabetic patients? Diabetes Care. 2001; 24: 1503–1504. PMID: 11473100

19. Chen Y, Lu J, Huang Y, Wang T, Xu Y, Xu M, et al. Association of previous schistosome infection with diabetes and metabolic syndrome: a cross-sectional study in rural China. J Clin Endocrinol Metab, 2013; 98: E283–E287. doi:10.1210/jc.2012-2517 PMID: 23725524

20. Wallace TM, Levy JC, Matthews DR. Use and abuse of HOMA modeling. Diabetes Care. 2004; 27: 1487–1495. PMID: 15161807

21. Stephenson LS, Latham MC, Ottesen EA. Malnutrition and parasitic helminth infections. Parasitology. 2000; 121: S23–S38. PMID: 11386688

22. Saldívar SR, Silveira AS, Filippi ST, Torres DM, Mangini AC, Dias RM, et al. Ascaris-Trichuris association and malnutrition in Brazilian children. Paediatr Perinat Epidemiol. 1999; 13: 89–98. PMID: 9987788

23. Oberhelman RA, Guerrero ES, Fernandez ML, Silio M, Mercado D, Comiskey N, et al. Correlations between intestinal parasitosis, physical growth, and psychomotor development among infants and children from rural Nicaragua. Am J Trop Med Hyg. 1998; 58: 470–475. PMID: 9574794

24. Wiria AE, Wammes LJ, Hamid F, Dekkers OM, Prasetyani MA, May L, et al. Relationship between Carotid Intima Media Thickness and Helminth Infections on Flores Island, Indonesia. PLoS One. 2013; 8: e54855. doi: 10.1371/journal.pone.0054855 PMID: 23365679

25. Pasha SM, Wiria AE, Wammes LJ, Smit JWA, Partono F, Supali T, et al. Blood pressure class and carotid artery intima-media thickness in a population at the secondary epidemiological transition. J Hypertens. 2011; 29: 2194–2200. doi:10.1097/HJH.0b013e32834bbba8 PMID: 21941206

26. Alberti KGMM, Zimmel P, Shaw J. The metabolic syndrome—a new worldwide definition. Lancet. 2005; 366: 1059–1062. PMID: 16182882

27. Kahn R, Buse J, Ferrannini E, Stern M. The metabolic syndrome: time for a critical appraisal; joint statement from the American Diabetes Association and the European Association for the Study of Diabetes. Diabetes Care. 2005; 28: 2289–2304. PMID: 16123508

28. Wiria AE, Prasetyani MA, Hamid F, Wammes LJ, Leel B, Ariawan I, et al. Does treatment of intestinal helminths infection influence malaria? Background and methodology of a longitudinal study of clinical, parasitological and immunological parameters in Nangapanda, Flores, Indonesia (ImmuNoSPIN Study). BMC Infect Dis. 2010; 10: 77. doi:10.1186/1471-2334-10-77 PMID: 20338054

29. Hamid F, Wiria AE, Wammes LJ, Kaisar MM, Leel B, Ariawan I, et al. A longitudinal study of allergy and intestinal helminth infections in semi urban and rural areas of Flores, Indonesia (ImmuNoSPIN Study). BMC Infect Dis. 2011; 11: 83. doi:10.1186/1471-2334-11-83 PMID: 21457559

30. Terhell AJ, Stolk WA, Haarbink M, Mangali A, Van Oortmarssen GJ, Yazdanbakhsh M. Regulation of anti-filarial IgE by infection pressure. Parasitology. 2002; 124: 509–519. PMID: 12049413

31. Verweij JJ, Brienien EAT, Ziem J, Yelifiari L, Polderman AM, Van Lieshout L. Simultaneous detection and quantification of Ancylostoma duodenale, Necator americanus, and Oesophagostomum bifurcum in fecal samples using multiplex real-time PCR. Am J Trop Med Hyg. 2007; 77: 685–690. PMID: 17978072

32. WHO Expert Consultation. Appropriate body-mass index for Asian populations and its implications for policy and intervention strategies. Lancet. 2004; 363: 157–163. PMID: 14726171
33. Bethony J, Brooker S, Albonico M, Geiger SM, Loukas A, Diemert D, et al. Soil-transmitted helminth infections: ascariasis, trichuriasis, and hookworm. Lancet. 2006; 367: 1521–1532. PMID: 16679166

34. Han JM, Patterson SJ, Speck M, Ehses JA, Levings MK. Insulin inhibits IL-10-mediated regulatory T cell function: implications for obesity. J Immunol. 2014; 192: 623–629. doi: 10.4049/jimmunol.1302181 PMID: 24323581

35. Khoo CM, Sairazi S, Taslim S, Gardner D, Wu Y, Lee J, et al. Ethnicity modifies the relationships of insulin resistance, inflammation, and adiponectin with obesity in a multiethnic Asian population. Diabetes Care. 2011; 34: 1120–1126. doi: 10.2337/dc10-2097 PMID: 21931814

36. Celis-Morales CA, Perez-Bravo F, Ibañes L, Sanzana R, Hormazabal E, Ulloa N, et al. Insulin resistance in Chileans of European and indigenous descent: evidence for an ethnicity x environment interaction. PLoS One. 2011; 6: e24690. doi: 10.1371/journal.pone.0024690 PMID: 21931814

37. Walk ST, Blum AM, Ewing SA-S, Weinstock JV, Young V. Alteration of the murine gut microbiota during infection with the parasitic helminth Heligmosomoides polygyrus. Inflamm Bowel Dis. 2010; 16: 1841–1849. doi: 10.1002/ibd.21299 PMID: 20848461

38. Kabat AM, Srinivasan N, Maloy KJ. Modulation of immune development and function by intestinal microbiota. Trends Immunol. 2014; 35: 507–517. doi: 10.1016/j.it.2014.07.010 PMID: 25172617

39. Reynolds LA, Smith KA, Filbey KJ, Harcus Y, Hewitson JP, Redpath SA, et al. Commensal-pathogen interactions in the intestinal tract: lactobacilli promote infection with, and are promoted by, helminth parasites. Gut Microbes. 2014; 5: 522–532. doi: 10.4161/gmic.32155 PMID: 25144609

40. Osborne LC, Monticelli LA, Nice TJ, Sutherland TE, Siracusa MC, Hepworth MR, et al. Virus-helminth coinfection reveals a microbiota-independent mechanism of immunomodulation. Science. 2014; 345: 578–582. doi: 10.1126/science.1256942 PMID: 25082704

41. Cho I, Blaser MJ. The human microbiome: at the interface of health and disease. Nat Rev Genet. 2012; 13: 260–270. doi: 10.1038/nrg3182 PMID: 22411464

42. Martinez-Hervas S, Argente C, Garcia-Jodar J, Priego A, Real JT, Carratala A, et al. Misclassification of subjects with insulin resistance and associated cardiovascular risk factors by homeostasis model assessment index. Utility of a postprandial method based on oral glucose tolerance test. Metabolism. 2011; 60: 740–746. doi: 10.1016/j.metabol.2010.07.024 PMID: 20850158