

Lifestyle counselling targeting infant's mother during the child's first year and offspring weight development until 4 years of age: a follow-up study of a cluster RCT

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To cite: Mustila T, Raitanen J, Keskinen P, *et al.* Lifestyle counselling targeting infant's mother during the child's first year and offspring weight development until 4 years of age: a follow-up study of a cluster RCT. *BMJ Open* 2012;**2**:e000624. doi:10.1136/bmjopen-2011-000624

► Prepublication history for this paper is available online. To view these files please visit the journal online (<http://dx.doi.org/10.1136/bmjopen-2011-000624>).

Received 16 November 2011
Accepted 22 December 2011

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ABSTRACT

Objective: To investigate the effect of intensified lifestyle counselling targeting infants' mothers on offspring weight development during the first 4 years of life.

Design and setting: Follow-up of a cluster-randomised controlled trial in primary care child health clinics during 2004–2006 in Finland. Participants received a follow-up survey during 2010 concerning weight and height measurements of their offspring. Number of clusters was six and the response rate to the follow-up 71.9% (N=64/89).

Participants: The participants (N=89) were mothers of infants aged 2–10 months.

Intervention: The intervention included individual counselling on diet and physical activity when the infant was 2–10 months of age and an option to attend supervised group exercise sessions.

Primary and secondary outcome measures: The authors analysed the secondary outcome of the intervention study: the weight development of the offspring. The primary outcome was the proportion of women returning to their prepregnancy weight by 10 months post partum, reported earlier.

Results: Multilevel mixed effect non-linear regression models included group, age of the child and interaction between group and age of the child. The increase of BMI z-score between 24 and 48 months was slower among the intervention group offspring (–0.034 to –0.002, $p=0.028$) as compared with control group. Z-scores for weight-for-length/height did not differ between groups when the period 0–48 months was analysed ($p=0.23$) but for the period of 24–48 months, between-group differences were significant ($p=0.012$).

Conclusions: Lifestyle counselling targeting mothers during the child's first year may be effective in slowing offspring weight gain until 4 years of age. However, larger studies are needed to confirm the findings which may have the potential in combatting the obesity epidemic.

Trial registration number: Current Controlled Trials ISRCTN21512277.

ARTICLE SUMMARY

Article focus

- Rapid preschool weight gain is known to increase risk for later obesity.
- There is lack of intervention studies targeting child's first year with follow-up of their weight gain.

Key message

- Results suggested that intensive lifestyle counselling targeting mother during child's first year may slower child's weight gain until 48 months of age.

Strengths and limitations of this study

- A feasible counselling method was used as well as a controlled trial setting and reliable growth data based on repeated measurements by nurses in primary child healthcare. We also utilised the recently updated growth data on Finnish children by using z-scores of weight-for-length/height and BMI-for-age described in that growth data.
- Since the study was a pilot study, number of participants and clusters was low. Also a longer follow-up period could reveal more clear influence of intervention on offspring weight development.

INTRODUCTION

The prevalence of childhood overweight and obesity has increased during the past three decades in the developed world and also in the developing world.^{1–3} However, recent evidence suggests that the increase in childhood obesity prevalence may be abating.⁴ Obesity has detrimental short- and long-term consequences to health, and successful treatment of obesity is difficult even in childhood. Effective preventive means are therefore needed.^{4–6} Because overweight

tends to begin during preschool years, early primary preventive interventions are thought to be the most effective means to combat the obesity epidemic. However, only few randomised, controlled primary prevention lifestyle counselling trials have been reported targeting families during offspring's first year of life, to our knowledge none targeting only mothers and including both diet and physical activity counselling. Some of them have shown slightly positive effects on child's weight development, but evidence of effective preventive means to reduce childhood obesity is still insufficient.⁷⁻⁹

Obesity is partly a result of genetic susceptibility, but an obesity epidemic is mainly attributable to societal and environmental changes, with changes in lifestyle.⁴ Pregnant mothers are also more often obese, and prenatally, a child may meet an obesinogenic environment.^{10 11} Mother's prepregnancy BMI and weight gain during pregnancy correlate with the offspring's risk for subsequent overweight and obesity, and mother's glucose intolerance has been shown to increase the offspring's birth weight.¹¹⁻¹³ In some studies, higher birth weight seems to increase the child's risk for overweight and obesity, but the evidence is weak.^{14 15} Children with rapid weight gain during their preschool years and children who reach their BMI rebound earlier are prone to obesity.¹⁶⁻²⁰ Modification of diet in infancy appears to reduce subsequent obesity risk.^{21 22} Since excessive weight gain begins already during preschool years, preventive interventions should start early, before pregnancy, and include pregnancy and infancy.²³ So far, only few such intervention studies have been published.^{4 7-9}

The aim of this study was to investigate whether individual counselling on diet and physical activity targeting first-time mothers with infants aged 2-10 months affects offspring weight gain by the age of 4 years.

METHODS

Study design, participants and methods

A controlled trial was conducted in six maternity and child health clinics in Finland in the cities of Tampere and Hämeenlinna between the years 2004 and 2006. Aim of the trial was to evaluate the feasibility and effects of a lifestyle intervention designed to prevent excessive gestational weight gain and postpartum weight retention. The study protocol was implemented during five visits to maternity or child healthcare clinics (figure 1). The prenatal intervention study will be reported elsewhere. Feasibility of the study protocol and other details have been reported earlier.²⁴⁻²⁷

The intervention study was conducted in six maternity and childcare centers, three of which volunteered to be intervention clinics and the remaining clinics were treated as control clinics. The allocation was performed at clinic level. The clinics were a convenience sample of the clinics in Tampere and Hämeenlinna as they were selected based on the clinics' administrative personnel's suggestion for suitable clinics. The participants consisted

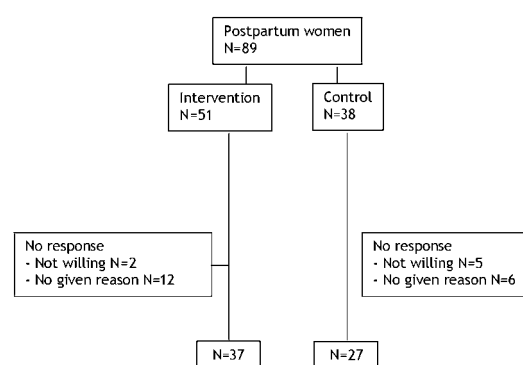


Figure 1 Original sample and the follow-up of the offspring, respondents and non-respondents.

of postpartum primiparous women. The exclusion criteria were age under 18 years, type 1 or type 2 diabetes mellitus (but not gestational diabetes mellitus), twin pregnancy, physical disability preventing exercising, otherwise problematic pregnancy (determined by a physician), substance abuse, treatment or clinical history for any psychiatric illness, inadequate language skills in Finnish and intention to change residence within 3 months. The nurses recruited postpartum women when visiting their home after delivery or at their first visit to the childcare center. The eligibility of all potential participants was assessed, and all eligible women were asked to participate in the study. All participants provided written informed consent for participation. The aim was to recruit at least 40 postpartum participants in the intervention and in the control clinics from August to October 2004.

For the original trial power calculations, we used assumptions from previous literature resulting at 90% power and significance level $\alpha=0.05$, which suggested 82 women per group, in total 164.²⁸ In addition, a conservative estimation of the sample size would be at least 1.5-fold compared with this calculation because cluster randomisation was applied. The estimated dropout rate (25%) was also taken into account in the sample size calculations. With these requirements, at least 300-350 women should be recruited to the original intervention study. However, statistical significance of the results was not a priority in a pilot study and we aimed to recruit at least 60 postpartum women. Of these women, approximately 15 postpartum women were assumed to discontinue the study because of spontaneous abortion, pregnancy complications or for other reasons.

Intervention

The intervention included individual counselling on physical activity and diet when the child was 2-10 months old and an option to attend supervised group exercise sessions once a week. The content of the intervention is described in greater detail elsewhere.²⁵ The purpose of the intervention was to promote leisure time physical activity and healthy dietary habits, thereby supporting participants' return to their prepregnancy

weight during the study. In the control clinics, the nurses continued their usual counselling practices on physical activity and diet.²⁶

Outcomes

In this study, we analysed the secondary outcome of the intervention study, namely the weight development of the offspring. The primary outcomes of the study have been reported earlier: the proportion of women returning to their prepregnancy weight by 10 months post partum. Dietary outcomes were changes in meal patterns, overall intake of vegetables, fruit and berries, use of high-fibre bread and intake of high-sugar snacks. Physical activity outcomes were MET (metabolic equivalent) minutes.^{25 27}

Follow-up data collection

In 2010, mothers participating in the trial received a postal questionnaire on their offspring's weight development. This questionnaire was chosen for data gathering as direct access to the child health clinic records would have entailed maternal permission, and mothers have the same information on their offspring's growth as the child health clinic records. Finnish children attend child healthcare clinics several times in their first year and once a year thereafter. Children's weight was measured to the nearest 0.1 kg on a standard electronic scale. Children under 2 years were measured in recumbent position and thereafter in standing position to the nearest millimetre with a standard stadiometer. A nurse enters the height and weight measurements in the child's own health booklet for the mothers. The mothers entered these measurements in the postal questionnaire. The mothers were also asked whether their children had any long-term illnesses affecting growth (allergies or other chronic diseases), duration of breast feeding and child's age when starting solid foods.

Statistical methods

Characteristics of the study participants were described using means and SDs or frequencies and proportions. Observed weight trajectories by gender are shown in figure 2. The child's size during follow-up was analysed using weight and length/height converted to BMI (weight (kg)/height (m²))-for-age and weight-for-length/height and their SDs (z-scores) according to the recently updated Finnish growth reference.^{29–31} Exact age of the child was used in all analyses.

Mixed-effects linear regression models were constructed to analyse the association of weight-for-length/height z-score and BMI z-score over time by group (intervention/control). Three-level mixed-effects models consisted of fixed effects (group, child's age in months, non-linear effects AgeInMonths² and AgeInMonths³ and interactions between group and age) and random effects (measurements within child within centre). These models allow for a difference between groups at baseline, linear changes of z-score over time

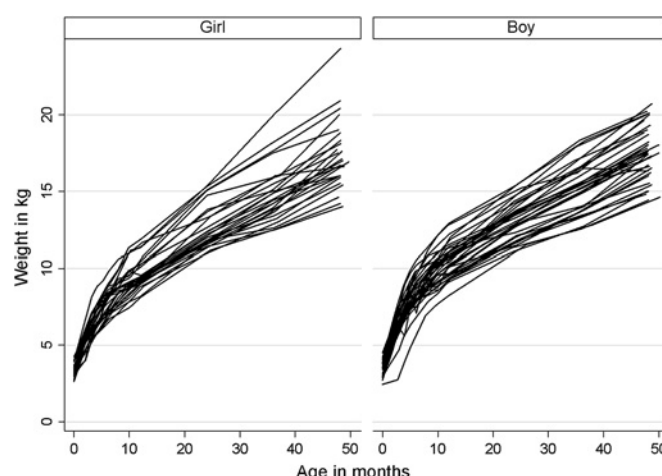


Figure 2 Growth trajectories by gender, exact age of the child and weight from birth to age of 48 months.

and the difference of improvement between groups, which can be viewed as the intervention effect (ie, interaction term). A likelihood ratio test was used for model selection. The parameter estimates were presented with 95% CI and p values. The goodness-of-fit of the models was evaluated visually by normal probability and residual plots and also tested by the normality of the residuals (Kolmogorov–Smirnov test). All analyses were performed using STATA software (V.12.0 for Windows), StataCorp LP.

Response rate to the follow-up questionnaire was 71.9% (N=64/89). We also performed an analysis to compare mothers lost to follow-up and respondents. According to the loss-of-follow-up analysis, mothers participating in the intervention and responding to the follow-up questionnaire reported significantly lower weight before pregnancy than non-responders (61.1 vs 66.3 kg, p=0.04). There were no differences in age, employment status or smoking before pregnancy, but the responding mothers tended to be more highly educated than non-respondents (highest education group 58.7% vs 33.3%, p=0.07).

We also estimated the power for the future studies using the current sample. With a multi-level structure, the ordinary sample size estimates need to be inflated by the design effect $(1+(n-1)\rho)$, where n is the average cluster size and ρ is the estimated intracluster correlation coefficient. When we have repeated measurements on the same child, the child is considered as the cluster. We applied the design effect after calculating a sample size (STATA, *sampclus*). New power estimates are shown in the Results section.

The study was approved by the Ethics Committee of the Pirkanmaa Hospital District.

RESULTS

Mothers in the intervention group who responded to the questionnaire were slightly older than the control mothers (mean age 29.6 vs 28.4 years, p=0.195). There

Table 1 Baseline characteristics of the trial groups (mean $\pm$ SD or frequency and %, difference between the groups and 95% CI)

	Intervention	Control	Difference (95% CI)	p Value	Missing
N	37	27			
Age of the mother at delivery	29.6 $\pm$ 3.6	28.4 $\pm$ 4.0	1.26 (−0.69 to 3.20)	0.195*	1, 0
Prepregnancy weight (kg)	61.8 $\pm$ 11.1	60.1 $\pm$ 8.1	1.64 (−3.19 to 6.47)	0.519*	1, 0
Prepregnancy BMI (kg/m ²)	22.4 $\pm$ 3.7	21.8 $\pm$ 2.4	0.68 (−0.85 to 2.21)	0.402*	1, 0
Range (kg)	18.1–35.4	17.3–27.9			
Prepregnancy BMI (kg/m ²)				0.593†	
<25	30 (83.3%)	25 (92.6%)	−9.3% (−24.9% to 6.4%)		1, 0
25–29.9	4 (11.1%)	2 (7.4%)	3.7% (−10.5% to 18.0%)		
30+	2 (5.6%)	—	5.6% (−1.9% to 13.0%)		
Gestational weight gain (kg)	15.8 $\pm$ 5.5	16.0 $\pm$ 5.0	−0.19 (−2.86 to 2.48)	0.888*	1, 0
Weight gain recommendations during pregnancy				0.965‡	
Lower	9 (25.0%)	6 (22.2%)	2.8% (−18.3% to 23.9%)		1, 0
At the range of the recommendations	10 (27.8%)	8 (29.6%)	−1.9% (−24.5% to 20.7%)		
Higher	17 (47.2%)	13 (48.1%)	−0.9% (−25.8% to 24.0%)		
Education				0.603‡	
Low	8 (22.2%)	8 (29.6%)	−7.4% (−29.3% to 14.5%)		1, 0
Medium	7 (19.4%)	3 (11.1%)	8.3% (−9.2% to 25.9%)		
High	21 (58.3%)	16 (59.3%)	−0.9% (−25.5% to 23.6%)		
Employed	32 (88.9%)	24 (88.9%)	0.0% (−15.7% to 15.7%)	1.000‡	1, 0
Ever-smokers	18 (50.0%)	17 (63.0%)	−13.0% (−37.4% to 11.5%)	0.306‡	1, 0
Smoking during pregnancy	4 (11.1%)	6 (22.2%)	−11.1% (−29.9% to 7.6%)	0.232‡	3, 1
Sex of the child—boy	21 (56.8%)	16 (59.3%)	−2.5% (−27.0% to 22.0%)	0.841‡	—
Proportion of children with SGA	7 (19.4%)	4 (15.4%)	4.1% (−14.9% to 23.0%)	0.680‡	1, 1
Proportion of children with LGA	1 (2.8%)	1 (3.8%)	−1.1% (−10.2% to 8.1%)	1.000‡	1, 1
Macrosomia, birth weight >4000 g	5 (13.5%)	3 (11.5%)	2.0% (−14.5% to 18.5%)	0.817‡	0, 1
Breast feeding (no other nutrition) (months)	4.0 $\pm$ 1.8	3.5 $\pm$ 2.4	0.45 (−0.64 to 1.54)	0.391*	—
Partial breast feeding (months)	6.0 $\pm$ 4.3	5.9 $\pm$ 5.8	0.12 (−2.52 to 2.76)	0.657§	—
Age of the child receiving solid foods (months)	4.8 $\pm$ 1.0	4.8 $\pm$ 1.3	−0.03 (−0.64 to 0.58)	0.870§	1, 0

*Independent samples t test.

†Fisher's exact test.

‡ χ^2 Test.

§Mann–Whitney U test.

LGA, large for gestational age; SGA, small for gestational age.

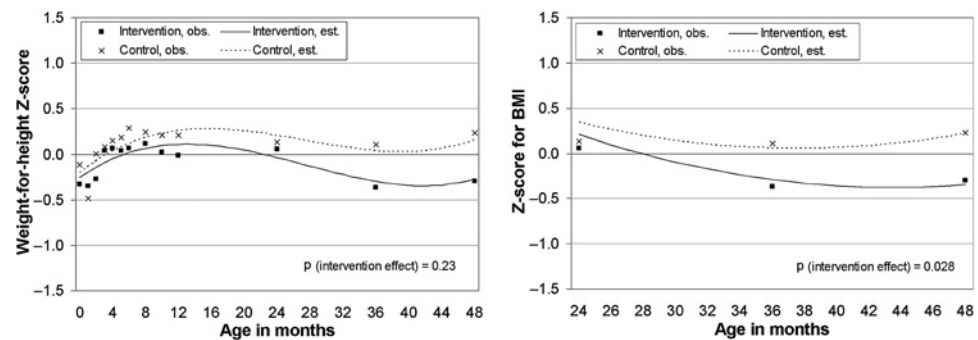
were no obese mothers (BMI $\geq$ 30 kg/m²) in the control group, whereas in the intervention group, there were two mothers who were obese before pregnancy, but mean prepregnancy BMI did not differ between groups ($p=0.40$) (table 1). Smoking during pregnancy or duration of breast feeding did not differ significantly between groups (table 1). Number of mothers reporting children's allergies (two in intervention group, one in control group) or any chronic diseases (four children in intervention and five in control group) was low and did not result to exclusion of these children. Proportion of missing mother–child dyads due to non-response was similar among intervention (N=14) and control (N=11) groups (figure 1).

Observed weight trajectories were slightly wider among girls than among boys until age of 48 months (figure 2). The weight gain from birth to 48 months of child's age measured as weight-for-length/height was no significantly different between the groups (figure 3).

Multilevel mixed effect non-linear regression models included group, age of the child and interaction between group and age of the child. The increase of BMI z-score between 24 and 48 months was slower among the intervention group offspring (−0.034 to −0.002, $p=0.028$) as compared with control group. Z-scores for weight-for-length/height did not differ between groups when the period 0–48 months was analysed ($p=0.23$), but for the period of 24–48 months between-group differences were significant ($p=0.012$) (table 2 and figure 3).

Based on the current data, we also estimated the sample size of the study which should be needed to achieve sufficient power for the study. From the current sample, we assume group means (SDs) of −0.3 (1.1) vs 0.2 (1.2) and intracluster correlation coefficient of 0.63 for z-score on weight for height. For future studies, 730 observations are needed, divided by the number of observations per child, 730/13=56.2, 57 children per group.

Figure 3 Weight-for-height from 0 to 48 months and BMI z-scores from 24 to 48 months. p Values denote for the significance of intervention effects (interaction between group and child's age at months). Non-linear model including age of the child and interaction between group $\times$ age. Obs., observed; est., estimated.



DISCUSSION

The main finding of our study was that the offspring of the mothers receiving intensified lifestyle counselling during the period from 2 to 10 months of infant's age may have slower weight gain measured as BMI z-scores between 24 and 48 months than the children in the control group. The STRIP Study showed that children overweight at 13 years had a steeper weight gain starting at 2 or 3 years.¹⁸ Thus, our result suggests that the lifestyle intervention might reduce the risk for obesity.

Since the unfavourable health consequences of obesity already begin during childhood and the treatment of childhood obesity tends not to lead to permanent results, early preventive measures are needed.^{4–6 23} One of the early determinants for obesity, type 2 diabetes and cardiovascular disease is rapid growth in early childhood.^{16 18 19 32 33} Most of the evidence published so far on rapid early growth and subsequent increased risk for obesity has concerned infants, but there is also similar evidence regarding later preschool years.^{18 19 34 35} To the best of our knowledge, no previous controlled intervention trials have targeted only mothers during child's first year and included both diet and physical activity counselling.^{8 9} Our study was follow-up of a cluster-randomised trial conducted in child health clinics. The

participants were first-time mothers without specifically sought risk determinants for having overweight offspring. There were no statistically significant differences between the groups regarding mother's age before pregnancy, prepregnancy BMI, gestational weight gain, education, smoking during pregnancy or duration of breast feeding. In the intervention clinics, the mothers received individual counselling on diet and physical activity and the option to attend supervised group exercise sessions once a week during the first 10 months of infant's life. The control group received conventional healthcare counselling.

The strengths of our study include a feasible counselling method, controlled trial setting and reliable growth data based on repeated measurements by nurses in primary child healthcare.^{25 26} We also utilised the recently updated growth data on Finnish children by using z-scores of weight-for-length/height and BMI-for-age described in that growth data.²⁹ Our sample included healthy first-time mothers thereby constituting a more homogeneous group than mothers with earlier deliveries. We were also able to take account of confounding factors on childhood growth, such as mothers' smoking and prepregnancy BMI. We have shown earlier that intensified counselling both during pregnancy and postpartum results in changes in mother's dietary and physical activity behaviour.^{24–27 36} Therefore, the beneficial sequelae in offspring weight gain found in this study are more probable than without an effective counselling method.

The weaknesses of our study include the relatively small number of participants and clusters. In spite of this, clinic level was taken into account in the models. The respondents were also more often highly educated and had lower prepregnancy weight than the non-respondents. Therefore, selective response may have influenced the result, and the abovementioned selection may have diminished the intervention effect. The possibility of Hawthorne effect cannot be denied either. Another reason for the small observed differences between groups may be that the participant mothers as a group had no special risk characteristics of having overweight children, such as obesity or low social class.⁴ The proportion of overweight children tends to increase with age, and longer follow-up time might have revealed increasing differences between the groups.^{4 37 38}

Table 2 Estimates and 95% CIs for z-scores for weight-for-length/height and body mass index

	Coefficient	95% CI	p Value
Weight-for-length/height z-score from 0 to 48 months of age			
Group	−0.056	−0.487 to 0.375	0.80
Age	0.071	0.044 to 0.098	<0.001
Age ²	−0.003	−0.005 to −0.002	<0.001
Age ³	0.000	0.000 to 0.000	<0.001
Group $\times$ age	−0.008	−0.021 to 0.005	0.23
BMI z-score from 24 to 48 months of age			
Group	0.308	−0.480 to 1.095	0.44
Age	−0.115	−0.174 to −0.057	<0.001
Age ²	0.002	0.001 to 0.002	<0.001
Group $\times$ age	−0.018	−0.034 to −0.002	0.028

Results from separate multilevel mixed-effects non-linear regression models including group (intervention/control), age and interaction between age of the child and group.

Power of the study was insufficient, but the primary aim of the original trial was to evaluate the feasibility of the counselling protocol. According to our estimates based on the current sample, at least 57 children are needed per group for future intervention studies concerning childhood obesity prevention.

The positive intervention effect on offspring weight gain is probably mediated by the healthier diet and increased physical activity adopted by the intervention mothers. The role of parents is vital in facilitating sustainable lifestyle behaviour in their offspring, and early childhood is a critical period in the acquisition of food preferences and physical activity habits.^{23 39} The impact of the intervention via mother and her breast milk on infant's early nutrition could partly explain the effect of the lifestyle intervention on offspring weight gain: infants have been shown to acquire a flavour bridge through breast milk, which is influenced by mother's diet, making it easier for a child to accept these flavours in her diet.⁴⁰

CONCLUSIONS

In our study, the intensified lifestyle intervention targeting mothers during child's first year may reduce weight gain in the offspring until 4 years of age. By slowing the weight gain, such an intervention targeting this crucial growth period could be one means of combating the obesity epidemic. To break this inter-generational circle of obesity and its complications, initiating early prevention programmes targeting mothers before, during and after pregnancy is essential, likewise community-based preventive actions.^{4 5 23} Larger randomised controlled trials are needed to gather more evidence for selecting the most effective preventive programmes.

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Acknowledgements We thank Tarja I Kinnunen and Minna Aittasalo for planning the physical activity and diet counselling of the trial, we thank Ms Päivi Viitanen of the UKK Institute for Health Promotion who participated in data collection and coding; Kirsi Mansikkamäki, MSc; Mrs Ulla Hakala; Mrs Ulla Honkanen; Mrs Taru Helenius and Mrs Sirke Rasinperä of the UKK Institute laboratory participated in data collection. We gratefully acknowledge the participating parents and children and nurses in maternity and child healthcare in the six clinics in Tampere and Hämeenlinna.

Funding This project was funded by the Competitive Research Funding of the Tampere University Hospital (Dr Riitta Luoto, grant 9G042, Dr Päivi Keskinen), Medical Research Fund of Vaasa Hospital District (Dr Taina Mustila), this project was funded by the UKK Institute for Health Promotion (Tampere, Finland), the Ministry of Health and Welfare, the Academy of Finland and the Foundation of Pediatric Research (Finland).

Competing interests None.

Ethics approval The study was approved by the Ethics Committee of the Pirkanmaa Hospital District.

Contributors RL is the guarantor of the study. TM, PK and RL planned the follow-up questionnaire to the mothers. TM coded the data together with a research assistant. AS produced the BMI-for-age statistics and participated in the interpretation of the BMI-for-age results. JR performed the statistical analyses. All contributors participated in drafting the manuscript and approved the final manuscript. All authors had full access to all data (including statistical reports and tables) in the study and take responsibility for the integrity of the data and the accuracy of the data analysis.

Provenance and peer review Not commissioned; externally peer reviewed.

Data sharing statement Technical appendix, statistical codes and data set available from the authors at email riitta.luoto@uta.fi. Consent for data sharing was obtained from the participants.

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STROBE 2007 (v4) checklist of items to be included in reports of observational studies in epidemiology*

Checklist for cohort, case-control, and cross-sectional studies (combined)

Section/Topic	Item #	Recommendation	Reported on page #
Title and abstract	1	(a) Indicate the study’s design with a commonly used term in the title or the abstract	1,2
		(b) Provide in the abstract an informative and balanced summary of what was done and what was found	1,2
Introduction			
Background/rationale	2	Explain the scientific background and rationale for the investigation being reported	4,6
Objectives	3	State specific objectives, including any pre-specified hypotheses	6
Methods			
Study design	4	Present key elements of study design early in the paper	6,7
Setting	5	Describe the setting, locations, and relevant dates, including periods of recruitment, exposure, follow-up, and data collection	6,7
Participants	6	(a) Cohort study—Give the eligibility criteria, and the sources and methods of selection of participants. Describe methods of follow-up Case-control study—Give the eligibility criteria, and the sources and methods of case ascertainment and control selection. Give the rationale for the choice of cases and controls Cross-sectional study—Give the eligibility criteria, and the sources and methods of selection of participants	6,7
		(b) Cohort study—For matched studies, give matching criteria and number of exposed and unexposed Case-control study—For matched studies, give matching criteria and the number of controls per case	6,7
Variables	7	Clearly define all outcomes, exposures, predictors, potential confounders, and effect modifiers. Give diagnostic criteria, if applicable	7,8
Data sources/ measurement	8*	For each variable of interest, give sources of data and details of methods of assessment (measurement). Describe comparability of assessment methods if there is more than one group	8
Bias	9	Describe any efforts to address potential sources of bias	5,6,7,8
Study size	10	Explain how the study size was arrived at	7
Quantitative variables	11	Explain how quantitative variables were handled in the analyses. If applicable, describe which groupings were chosen and why	8,9
Statistical methods	12	(a) Describe all statistical methods, including those used to control for confounding	8,9
		(b) Describe any methods used to examine subgroups and interactions	8,9
		(c) Explain how missing data were addressed	8,9

		(d) <i>Cohort study</i> —If applicable, explain how loss to follow-up was addressed <i>Case-control study</i> —If applicable, explain how matching of cases and controls was addressed <i>Cross-sectional study</i> —If applicable, describe analytical methods taking account of sampling strategy (e) Describe any sensitivity analyses	6,7,8,9
Results			
Participants	13*	(a) Report numbers of individuals at each stage of study—eg numbers potentially eligible, examined for eligibility, confirmed eligible, included in the study, completing follow-up, and analysed (b) Give reasons for non-participation at each stage (c) Consider use of a flow diagram	figure 1, 9,10 figure 1 figure 1
Descriptive data	14*	(a) Give characteristics of study participants (eg demographic, clinical, social) and information on exposures and potential confounders (b) Indicate number of participants with missing data for each variable of interest (c) <i>Cohort study</i> —Summarise follow-up time (eg, average and total amount)	9,10, table 1 9,10 8
Outcome data	15*	<i>Cohort study</i> —Report numbers of outcome events or summary measures over time <i>Case-control study</i> —Report numbers in each exposure category, or summary measures of exposure <i>Cross-sectional study</i> —Report numbers of outcome events or summary measures	6,7 - -
Main results	16	(a) Give unadjusted estimates and, if applicable, confounder-adjusted estimates and their precision (eg, 95% confidence interval). Make clear which confounders were adjusted for and why they were included (b) Report category boundaries when continuous variables were categorized (c) If relevant, consider translating estimates of relative risk into absolute risk for a meaningful time period	9,10,11 9,10,11
Other analyses	17	Report other analyses done—eg analyses of subgroups and interactions, and sensitivity analyses	9,10,11
Discussion			
Key results	18	Summarise key results with reference to study objectives	11
Limitations	19	Discuss limitations of the study, taking into account sources of potential bias or imprecision. Discuss both direction and magnitude of any potential bias	11,12,13
Interpretation	20	Give a cautious overall interpretation of results considering objectives, limitations, multiplicity of analyses, results from similar studies, and other relevant evidence	11,12,13
Generalisability	21	Discuss the generalisability (external validity) of the study results	11,12,13
Other information			
Funding	22	Give the source of funding and the role of the funders for the present study and, if applicable, for the original study on which the present article is based	14

*Give information separately for cases and controls in case-control studies and, if applicable, for exposed and unexposed groups in cohort and cross-sectional studies.

Note: An Explanation and Elaboration article discusses each checklist item and gives methodological background and published examples of transparent reporting. The STROBE checklist is best used in conjunction with this article (freely available on the Web sites of PLoS Medicine at <http://www.plosmedicine.org/>, Annals of Internal Medicine at <http://www.annals.org/>, and Epidemiology at <http://www.epidem.com/>). Information on the STROBE Initiative is available at www.strobe-statement.org.