Risks of major congenital malformations in relation to maternal overweight and obesity severity: cohort study of 1.2 million singletons

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ABSTRACT

OBJECTIVE
To estimate the risks of major congenital malformations in the offspring of mothers who are underweight (body mass index (BMI) <18.5), overweight (BMI 25 to <30), or in obesity classes I (BMI 30 to <35), II (35 to <40), or III (≥40) compared with offspring of normal weight mothers (BMI 18.5 to <25) in early pregnancy.

DESIGN
Population based cohort study.

SETTING
Nationwide Swedish registries.

PARTICIPANTS
1243 957 liveborn singleton infants from 2001 to 2014 in Sweden. Data on maternal and pregnancy characteristics were obtained by individual record linkages.

EXPOSURE
Maternal BMI at the first prenatal visit.

MAIN OUTCOME MEASURES
Offspring with any major congenital malformation, and subgroups of organ specific malformations diagnosed during the first year of life. Risk ratios were estimated using generalised linear models adjusted for maternal factors, sex of offspring, and birth year.

RESULTS
A total of 43 550 (3.5%) offspring had any major congenital malformation, and the most common subgroup was for congenital heart defects (n=20 074; 1.6%). Compared with offspring of normal weight mothers (risk of malformations 3.4%), the proportions and adjusted risk ratios of any major congenital malformation among the offspring of mothers with higher BMI were: overweight, 3.5% and 1.05 (95% confidence interval 1.02 to 1.07); obesity class I, 3.8% and 1.12 (1.08 to 1.15), obesity class II, 4.2% and 1.23 (1.17 to 1.30), and obesity class III, 4.7% and 1.37 (1.26 to 1.49). The risks of congenital heart defects, malformations of the nervous system, and limb defects also progressively increased with BMI from overweight to obesity class III. The largest organ specific relative risks related to maternal overweight and increasing obesity were observed for malformations of the nervous system. Malformations of the genital and digestive systems were also increased in offspring of obese mothers.

CONCLUSIONS
Risks of any major congenital malformation and several subgroups of organ specific malformations progressively increased with maternal overweight and increasing severity of obesity. For women who are planning pregnancy, efforts should be encouraged to reduce adiposity in those with a BMI above the normal range.

Introduction
Obesity has reached epidemic proportions globally and is now a major health concern in pregnancy, in both high and low income countries. In particular the prevalence of severe obesity, commonly defined as obesity classes II and III (body mass index (BMI) ≥35) is rapidly increasing.2-4 In the US, approximately half of the women are overweight or obese at the first antenatal visit,5 and the high prevalence of obesity class III (BMI ≥40; 10-12%) in women of reproductive age is of concern.6 In Sweden, the prevalence of early pregnancy obesity (BMI ≥30) increased from 6.0% to 12.9% from 1992 to 2014.7 Globally, it was recently reported that the number of women aged 18 years and older with a BMI ≥35 doubled from approximately 50 million to 100 million between 2000 and 2010.8

Obesity in pregnancy adversely influences both fetal and neonatal outcomes,9,15 including increased risks of major congenital malformations, which are a common cause of stillbirth16 and a major cause of infant mortality and long term morbidity.14,17 A meta-analysis reported that the offspring of obese mothers are at increased risk of a wide range of congenital malformations, including neural tube defects, cardiovascular anomalies, cleft lip and palate, anorectal atresia, and limb reduction anomalies.18 However, associations between increasing severity of obesity and risks of malformations were not analysed and it is not clear if risks are also increased in the offspring of overweight mothers. Several of the included studies also used non-standard definitions of normal weight, overweight, and obesity.19-28 Since the prevalence of severe obesity is increasing among women of reproductive age, it is of
interest to investigate whether risks of major congenital malformations increase with severity of obesity in mothers.

In this study we included information on 1243 957 live singleton births in Sweden recorded in the medical birth register between 2001 and 2014. We investigated associations between maternal BMI in early pregnancy and risks of any major congenital malformation as well as risks of the most prevalent subgroups of organ specific malformations.

**Methods**

**Setting**
The study was performed in Sweden, where prenatal and delivery care is publicly funded, and participation in the standardised prenatal care programme is almost 100%. The Swedish medical birth register includes information on close to 100% of all births in Sweden since 1973. Using standardised prenatal, obstetrical, and neonatal records, information is prospectively collected during pregnancy, delivery, and the neonatal period. Individual information on maternal height and weight has been included since 1992.

**Data sources**

By using the unique personal identification number assigned to each Swedish resident,26 we linked data from the medical birth register to the national patient register,27 cause of death register, education register, and total population register.28 The national patient register includes diagnoses and dates on visits for hospital based inpatient and outpatient care. Diagnoses are coded according to the Swedish version of the ICD-10 (international classification of diseases, 10th revision). In Sweden, all pregnant women are offered an ultrasonic examination at around 18 weeks of gestation. Abortion after 18 until 22 completed weeks must be approved by the National Board of Health and Welfare. Severe birth defects as indication for late abortion are generally approved. In 2015, the rate of abortions in Sweden was 20.9/1000 women (aged 15-44 years; 93% of abortions were performed before 12 weeks of gestation and 83% before nine weeks of gestation).

**Study population**

From 1 January 2001 to 31 December 2014, we retrieved information on 1 480 892 deliveries recorded in the Swedish medical birth register. We excluded 42 638 (2.9%) multiple births (since they differ from single births for malformation outcomes30) and 4598 (0.3%) stillbirths (where diagnosis and registration of malformations is poor or missing). Of the remaining 1 433 656 live singleton births, 19 863 (1.4%) were excluded owing to lack of valid personal identification numbers (18 929 for the mother and 934 for the infant), as these births could not be linked to other registries. Because we assumed that adiposity is unlikely to cause malformations for which another cause is known, we also excluded 7514 (0.5%) infants with chromosomal aberrations, genetic syndromes, malformation syndromes with known causes, and viral infections having a possible association with malformations (see supplementary eMethods).30 Finally, seven infants did not have data on sex and were therefore excluded. After these exclusions, 1 406 272 singletons remained, of whom 1 243 957 (88.5%) had complete data on all covariates.

Women missing data on BMI were more likely to have missing data on smoking status and family situation (see supplementary eTable 1). Maternal age, level of education, parity, and proportion of women of Nordic origin were similar in women with missing BMI data compared with normal weight women. However, women missing BMI data were more likely to have high education and to be primiparous compared with obese women.

**Patient involvement**

No patients were involved in setting the research question or the outcome measures, nor were they involved in developing plans for design, or implementation of the study. No patients were asked to advise on interpretation or writing up of results. There are no plans to disseminate the results of the research to study participants or the relevant patient community.

**Main exposure**

We calculated early maternal pregnancy BMI from measured weight and self reported height at the first prenatal visit, which takes place during the first trimester (first 14 weeks of gestation) for 90% of all women.32 We calculated the median height of each mother using information from all available pregnancies.31 Data on maternal height and/or weight were missing in 9.6% of live singleton births (see supplementary eTable 1). Based on BMI, we categorised the women as underweight (BMI <18.5), of normal weight (18.5 to <25), overweight (25 to <30), or in obesity class I (30 to <35), obesity class II (35 to <40), or obesity class III (≥40).32

**Covariates**

Data on parity, whether the mother cohabited with a partner or not, and self reported smoking were registered at the first prenatal visit. Information on maternal age was obtained at delivery, highest attained education was retrieved from the education register, and mother’s country of birth was retrieved from the total population register.

**Outcomes**

The main outcome was presence of any major congenital malformations in liveborn infants (from 22 completed gestational weeks), as recorded in either the medical birth register, the national patient register within one year of birth (including inpatient and hospital based outpatient care), or the cause of death register. Major congenital malformations were coded according to the ICD-10 classification and were defined according to the European Surveillance of Congenital
Anomalies classification (EUROCAT; www.eurocat-network.eu; see supplementary eTables 2 and 3 for ICD-10 codes).

We analysed the most common subgroups of major congenital malformations with a prevalence of ≥0.1% as additional outcomes. These included major congenital malformations of the heart, limbs, genital organs, urinary system, digestive system, orofacial clefts, eye, nervous system, and other malformations. In keeping with the EUROCAT definition, patent ductus arteriosus in preterm infants and persistent pulmonary stenosis were not considered as major congenital malformations. Other minor malformations were also excluded in accordance with the EUROCAT classification.

Statistical analysis
Using generalised linear models with a robust sandwich estimator, we estimated risk ratios and 95% confidence intervals for the outcomes in offspring of underweight mothers (BMI <18.5), overweight mothers (25 to <30), and mothers in obesity classes I (30 to <35), II (35 to <40), and III (≥40) compared with the offspring of normal weight mothers (18.5 to <25). To adjust for the possible dependence in outcome introduced by repeated births in the same mother, we constructed models with mother’s identification number as a cluster. The outcome was assumed to follow a Poisson distribution. We made adjustments for maternal age, height, parity, early pregnancy smoking status (0, 1-9, or ≥10 cigarettes daily), educational level, mother’s country of birth, family situation (living or not living with a partner), and sex of offspring.

Subgroup analyses—sex specific subgroup analyses were performed for any congenital malformations as well as for subgroups of major congenital malformations.

Sensitivity analyses—diabetes is a known teratogen33-35 and may also be in the causal pathway between obesity and major congenital malformations. In sensitivity analyses, we excluded women with pregestational diabetes to investigate whether the associations between BMI and major congenital malformations were affected. Although gestational diabetes generally develops after the developmentally critical period for malformations, we also performed an analysis where we excluded both women with pregestational diabetes and women with gestational diabetes.

An additional analysis was performed for the association between BMI and the outcome of any chromosomal aberration, genetic syndrome, malformation syndrome with known causes, and viral infection having a possible association with malformations (infants with these conditions were excluded from the study population of the main analyses; n=7514, of whom 6495 had complete data on all variables).

Data were analysed using SAS (version 9.4). We considered two sided P values <0.05 to be statistically significant. No adjustment was made for multiple comparisons.

Results
Major congenital malformations
During the study period a total of 1243957 liveborn singleton infants were included in the cohort, 43550 (3.5%) of whom had any major congenital malformation. Congenital heart defects were the most common malformation subtype (1.6%) followed by malformations of the genital organs (0.5%), limbs (0.4%), urinary system (0.3%), other (0.2%), eye (0.2%), digestive system (0.2%), orofacial clefts (0.1%), and nervous system (0.1%; fig 1).

BMI and major congenital malformations
The proportions of offspring with any major congenital malformation were 3.4% for underweight mothers, 3.4% for normal weight mothers, 3.5% for overweight mothers, and 3.8% for mothers in obesity class I, 4.2% in obesity class II, and 4.7% in obesity class III (table 1). In unadjusted analyses, risk ratios of any malformation were increased in offspring of overweight mothers, and increased with severity of obesity. A weak U-shaped relation was found between maternal age and risk of any major congenital malformation. Risks of any major congenital malformations were higher in boys than in girls and in offspring of smokers, primiparous women, mothers of low stature, and mothers who were not cohabiting with their partner (table 1). Supplementary eTable 1 provides maternal characteristics by BMI categories.

The adjusted risk ratios of any major congenital malformation increased with maternal overweight and severity of obesity (fig 2). Compared with offspring of normal weight mothers, the adjusted risk ratios for any major congenital malformation increased with maternal BMI: 1.05 (95% confidence interval 1.02 to 1.07) in overweight mothers, 1.12 (1.08 to 1.15) in mothers in obesity class I, 1.23 (1.17 to 1.30) in mothers in obesity class II, and 1.37 (1.26 to 1.49) in mothers in obesity class III (fig 2).

The overall risk of any major congenital malformation was higher in boys (4.1%) than in girls (2.8%). In
analyses stratified by sex of offspring, the risk of any malformation in boys of underweight mothers was 4.0%, normal weight mothers was 4.0%, and overweight mothers was 4.3% and in offspring of mothers in obesity classes I, II, and III was 4.5%, 4.9%, and 5.3%, respectively (fig 2). The corresponding risks in girls were 2.8%, 2.8%, 2.8%, 3.0%, 3.4%, and 4.0%. The risk ratios of any malformation increased with maternal severity of obesity in both boys and girls (fig 2).

BMI and specific types of major congenital malformations

The adjusted risk ratios of malformations of the nervous system, heart, digestive system, genital organs, limbs, and other malformations all increased with increasing maternal BMI (P for trend ≤0.03 in all analyses; figs 3 and 4).

The adjusted risk ratios for congenital heart defects by maternal BMI were 1.05 (95% confidence interval 1.01 to 1.08) for overweight mothers, 1.15 (1.09 to 1.20) for mothers in obesity class I, 1.26 (1.16 to 1.37) for mothers in obesity class II, and 1.44 (1.27 to 1.63) for mothers in obesity class III. The largest organ specific increases in risk ratios related to overweight and severity of obesity were observed for malformations of the nervous system. Compared with offspring of normal weight mothers, the adjusted risk ratios for malformations in the nervous system were 1.15 (95% confidence interval 1.00 to 1.31) for overweight mothers, 1.44 (1.20 to 1.73) for mothers in obesity class I, 1.65 (1.23 to 2.21) for mothers in obesity class II, and 1.88 (1.20 to 2.94) for mothers in obesity class III (fig 3).

When comparing subgroups of malformations in boys and girls, the largest sex difference (with higher proportions for boys) was noted for genital malformations (boys 0.9% and girls 0.04%) and malformations in the urinary system (boys 0.5% and girls 0.2%; see supplementary eFigure 2). In the sex stratified analyses of malformation subtypes, comparisons were hampered by reduced statistical power in the different BMI categories, especially in girls. The risks of congenital heart defects increased with maternal overweight and severity of obesity for boys, whereas only female offspring of mothers in obesity classes I to III were at increased risks. Maternal BMI was associated with statistically significant increased risks of malformations in the digestive system in boys but not in girls (see supplementary eFigure 2).

Overall, 0.5% of infants who had diagnoses associated with increased risk of malformations were excluded from the main analyses, with a range from 0.5% in the offspring of normal weight mothers to 0.8% in the offspring of mothers in obesity class III. Risks of these malformations also generally increased with maternal overweight and severity of obesity (see supplementary eTable 4).

Sensitivity analysis: exclusion of women with diabetes—after exclusion of women with pregestational and gestational diabetes, the associations between BMI and overall major congenital malformations were materially unchanged (see supplementary eTable 5).

Discussion

This large population based study found that overall risks of major congenital malformations and risks of several organ specific groups of malformations progressively increase with maternal overweight and severity of obesity. By finding a dose-response relation throughout the spectrum of an above normal body mass index (BMI) and associations between maternal overweight and a range of malformation subgroups, this study substantially expands on previous data, which largely rest on a meta-analysis of pooled studies with varying definitions of obesity and no information on severity of obesity.38
Fig 2 | Major congenital malformations in liveborn singletons by maternal body mass index (BMI) in underweight (BMI <18.5; n=29 864), normal weight (BMI 18.5 to <25; n=756 432), and overweight (BMI 25 to <30; n=311 339) women, and in women in obesity classes I (BMI 30 to <35; n=103 085), II (BMI 35 to <40; n=31 883), and III (BMI ≥40; n=11 354). Adjustment was made for maternal age (13-24, 25-29, 30-34, ≥35 years), height (130-154, 155-159, 160-164, 165-169, 170-174, 175-200 cm), parity (primiparous, multiparous), early pregnancy smoking status (non-smoker, 1-9, ≥10 cigarettes daily), educational level (<10, 10-12, >12 years), maternal country of birth (Nordic (Sweden, Denmark, Finland, Iceland, and Norway), non-Nordic), family situation (living with partner, not living with partner), and sex of offspring.

Congenital heart defects were the most prevalent subtype of organ specific malformation, and risks increased with maternal overweight and increasing obesity in a dose-response pattern. This is consistent with findings from a meta-analysis focused on congenital heart defects, reporting increasing risks with maternal overweight, and with women in obesity class I (BMI 30 to <35) and obesity classes II and III (BMI ≥35).16

The largest organ specific risk ratios related to increasing maternal obesity were observed for malformations of the nervous system. Compared with offspring of normal weight mothers, offspring of mothers in obesity class III had an almost doubled risk of major congenital malformations of the nervous system. In line with the present finding, a meta-analysis on maternal BMI and risk of malformations reported a close two-fold increased risk of neural tube defects in offspring of obese mothers.18 However, the obesity related risks of malformations in the nervous system must be interpreted with caution as antenatal detection of these malformations might be more difficult in obese women compared with normal weight women.

Strengths and limitations of this study
Strengths of the present study include the population based study design, with more than one million singleton births. The large sample size enabled us to investigate the effects of overweight and severity of obesity on risks of major congenital malformations and several specific malformation subgroups. Data on exposures and outcomes were prospectively collected within the universally accessible Swedish healthcare system. Maternal BMI was calculated based on measured weight, which limits recall bias, but height was self reported. We used standard BMI categories as defined by WHO.32 Major congenital malformations were classified according to the EUROCAT categorisation. Because we used data from several nationwide registries, we had an opportunity to identify the majority of infants with a diagnosis of a major congenital malformation within the first year of life. Furthermore, we were able to adjust risk estimates for important confounders.

Our study was restricted to live births. Malformations are more common in pregnancies ending in miscarriage or stillbirths, and some prenatally diagnosed malformations may also lead to induced abortions. In particular, most pregnancies complicated by neural tube defects are terminated by induced abortion.7 The Swedish national registries with patient level data do not include individual data on malformations in pregnancies with miscarriages, stillbirths, and induced abortions. However, aggregated data for Sweden reported on the EUROCAT website for the period 2007-13 show that there were 551 cases of neural tube defects, of which 396 (72%) were classified as termination of pregnancy for fetal anomaly.32

Antenatal detection of congenital malformations may be more difficult in obese than in normal weight women.38 If some malformations in offspring of obese women (notably neural tube defects) were less likely to be diagnosed prenatally and the women underwent an induced abortion, we may have overestimated the risks of malformations in offspring of obese mothers. However, risks may be underestimated if obesity is associated with malformations leading to spontaneous abortion. This hypothesis is supported by findings from studies including information about pregnancy terminations and reporting a doubled risk of neural tube defects and a 50% increased risk of cardiovascular anomalies in pregnancies with obesity.18

In this study we used BMI as a proxy for adiposity. This is a reasonable assumption given the strong correlation between BMI and fat mass in early pregnancy.39 We did not have information on the fat distribution, which may be of interest to further explore the association between overweight and severity of obesity and risks of malformations. In addition, we cannot rule out the possibility of residual confounding by unknown or unmeasured factors, such as alcohol use. Power was limited for analyses of less prevalent malformation subgroups by increasing severity of maternal obesity.

Potential mechanisms
The pathophysiology of malformations is multifactorial, with interactions between genetic and environmental
Nervous system

<table>
<thead>
<tr>
<th>BMI (kg/m²)</th>
<th>Events (%)</th>
<th>Adjusted risk ratio (95% CI)</th>
</tr>
</thead>
<tbody>
<tr>
<td>&lt;18.5</td>
<td>1272 (0.10)</td>
<td>1.28 (0.91 to 1.79)</td>
</tr>
<tr>
<td>18.5 to &lt;25</td>
<td>38 (0.13)</td>
<td>1.00 (ref)</td>
</tr>
<tr>
<td>25 to &lt;30</td>
<td>695 (0.09)</td>
<td>1.15 (1.00 to 1.31)</td>
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<tr>
<td>30 to &lt;35</td>
<td>331 (0.11)</td>
<td>1.44 (1.20 to 1.73)</td>
</tr>
<tr>
<td>35 to &lt;40</td>
<td>139 (0.13)</td>
<td>1.65 (1.23 to 2.21)</td>
</tr>
<tr>
<td>≥40</td>
<td>49 (0.15)</td>
<td>1.88 (1.20 to 2.94)</td>
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</tbody>
</table>

Congenital heart defects

<table>
<thead>
<tr>
<th>BMI (kg/m²)</th>
<th>Events (%)</th>
<th>Adjusted risk ratio (95% CI)</th>
</tr>
</thead>
<tbody>
<tr>
<td>&lt;18.5</td>
<td>20  (0.11)</td>
<td>0.93 (0.71 to 1.23)</td>
</tr>
<tr>
<td>18.5 to &lt;25</td>
<td>2407 (0.19)</td>
<td>1.00 (ref)</td>
</tr>
<tr>
<td>25 to &lt;30</td>
<td>54 (0.18)</td>
<td>1.00 (0.91 to 1.10)</td>
</tr>
<tr>
<td>30 to &lt;35</td>
<td>1466 (0.19)</td>
<td>1.00 (0.86 to 1.16)</td>
</tr>
<tr>
<td>35 to &lt;40</td>
<td>603 (0.19)</td>
<td>1.02 (0.79 to 1.32)</td>
</tr>
<tr>
<td>≥40</td>
<td>198 (0.19)</td>
<td>1.03 (0.68 to 1.57)</td>
</tr>
</tbody>
</table>

Orofacial clefts

<table>
<thead>
<tr>
<th>BMI (kg/m²)</th>
<th>Events (%)</th>
<th>Adjusted risk ratio (95% CI)</th>
</tr>
</thead>
<tbody>
<tr>
<td>&lt;18.5</td>
<td>2074 (1.61)</td>
<td>0.99 (0.90 to 1.09)</td>
</tr>
<tr>
<td>18.5 to &lt;25</td>
<td>458 (1.53)</td>
<td>1.00 (ref)</td>
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<tr>
<td>25 to &lt;30</td>
<td>11807 (1.56)</td>
<td>1.05 (1.01 to 1.08)</td>
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<tr>
<td>30 to &lt;35</td>
<td>5082 (1.63)</td>
<td>1.15 (1.09 to 1.20)</td>
</tr>
<tr>
<td>35 to &lt;40</td>
<td>1840 (1.78)</td>
<td>1.26 (1.16 to 1.37)</td>
</tr>
<tr>
<td>≥40</td>
<td>630 (1.98)</td>
<td>1.44 (1.27 to 1.63)</td>
</tr>
</tbody>
</table>

Adjusted risk ratios were calculated after adjusting for abnormal glucose tolerance, overweight, smoking status, age of offspring, sex of offspring, maternal smoking, alcohol use, socioeconomic factors, antiepileptic drugs, deficiency of folic acid and other vitamins, and diabetes. The possible teratogenic role of other metabolic derangements associated with obesity, such as insulin resistance, hyperlipidemia, and inflammation, is unclear. The adipose tissue is an active metabolic and endocrine organ, distributed subcutaneously and in the visceral compartment. Visceral obesity in pregnancy is associated with a state of inflammation, vascular dysfunction, and abnormal placental metabolism, which may adversely influence organogenesis and fetal development.

Fig 3 | Major congenital malformations in nervous system, eye, heart, and oral clefts in liveborn singletons by maternal body mass index (BMI) in underweight (BMI <18.5; n=29 864), normal weight (BMI 18.5 to <25; n=756 432), and overweight (BMI 25 to <30; n=311 339) women, and in women in obesity classes I (BMI 30 to <35; n=103 085), II (BMI 35 to <40; n=318 833), and III (BMI ≥40; n=11 354). Adjustment was made for maternal age (13-24, 25-29, 30-34, ≥35 years), height (130-154, 155-159, 160-164, 165-169, 170-174, 175-200 cm), parity (primiparous, multiparous), early pregnancy smoking status (non-smoker, 1-9, ≥10 cigarettes daily), educational level (<10, 10-12, >12 years), maternal country of birth (Nordic (Sweden, Denmark, Finland, Iceland, and Norway), non-Nordic), family situation (living with partner, not living with partner), and sex of offspring.
BMI (kg/m²)  | Events (%) | Adjusted risk ratio (95% CI) | Adjusted risk ratio (95% CI)
--- | --- | --- | ---
Digestive system | | | |
<18.5 | 43 (0.14) | 0.98 (0.72 to 1.33) | 1.00 (ref)
18.5 to <25 | 1134 (0.15) | 0.97 (0.87 to 1.09) | 1.01 (0.94 to 1.09)
25 to <30 | 452 (0.15) | 1.41 (1.21 to 1.63) | 0.98 (0.88 to 1.09)
30 to <35 | 215 (0.21) | 1.33 (1.03 to 1.72) | 0.91 (0.74 to 1.11)
35 to <40 | 63 (0.20) | 1.54 (1.05 to 2.28) | 1.19 (0.88 to 1.60)
≥40 | 26 (0.23) | | |
Urinary system | | | |
<18.5 | 115 (0.39) | 1.18 (0.98 to 1.43) | 1.03 (0.87 to 1.22)
18.5 to <25 | 2544 (0.34) | 1.00 (ref) | 1.00 (ref)
25 to <30 | 1068 (0.34) | 1.01 (0.94 to 1.09) | 1.01 (0.94 to 1.09)
30 to <35 | 342 (0.33) | 0.98 (0.88 to 1.10) | 0.98 (0.88 to 1.10)
35 to <40 | 97 (0.30) | 0.91 (0.74 to 1.11) | 0.91 (0.74 to 1.11)
≥40 | 45 (0.40) | 1.19 (0.88 to 1.60) | 1.19 (0.88 to 1.60)
Genital organs | | | |
<18.5 | 5921 (0.48) | 1.02 (0.83 to 1.24) | 1.02 (0.83 to 1.24)
18.5 to <25 | 145 (0.49) | 1.00 (ref) | 1.00 (ref)
25 to <30 | 1496 (0.48) | 1.04 (0.98 to 1.11) | 1.04 (0.98 to 1.11)
30 to <35 | 529 (0.51) | 1.12 (1.02 to 1.23) | 1.12 (1.02 to 1.23)
35 to <40 | 182 (0.57) | 1.27 (1.09 to 1.47) | 1.27 (1.09 to 1.47)
≥40 | 72 (0.63) | 1.43 (1.13 to 1.80) | 1.43 (1.13 to 1.80)
Limb | | | |
<18.5 | 103 (0.34) | 1.01 (0.83 to 1.24) | 1.10 (1.01 to 1.18)
18.5 to <25 | 2616 (0.35) | 1.13 (1.02 to 1.25) | 1.13 (1.02 to 1.25)
25 to <30 | 1182 (0.38) | 1.12 (1.02 to 1.34) | 1.12 (1.02 to 1.34)
30 to <35 | 402 (0.39) | 1.19 (0.98 to 1.40) | 1.19 (0.98 to 1.40)
35 to <40 | 124 (0.39) | | |
≥40 | 51 (0.45) | | |
Other | | | |
<18.5 | 2672 (0.21) | 0.78 (0.59 to 1.06) | 0.78 (0.59 to 1.06)
18.5 to <25 | 49 (0.16) | 1.00 (ref) | 1.00 (ref)
25 to <30 | 1618 (0.21) | 0.96 (0.88 to 1.06) | 0.96 (0.88 to 1.06)
30 to <35 | 644 (0.21) | 1.00 (0.87 to 1.15) | 1.00 (0.87 to 1.15)
35 to <40 | 221 (0.21) | 1.55 (1.28 to 1.89) | 1.55 (1.28 to 1.89)
≥40 | 106 (0.33) | 1.39 (0.99 to 1.95) | 1.39 (0.99 to 1.95)

Fig 4 | Major congenital malformations in digestive, urinary, and genital systems; limbs; and other malformations anliveborn singletons by maternal body mass index (BMI) in overweight (BMI <18.5; n=29 864), normal weight (BMI 18.5 to <25; n=756 432), and overweight (BMI 25 to <30; n=31 339) women, and in women in obesity classes I (BMI 30 to <35; n=103 085), II (BMI 35 to <40; n=31 883), and III (BMI ≥40; n=11 354). Adjustment was made for maternal age (13-24, 25-29, 30-34, ≥35 years), height (130-154, 155-159, 160-164, 165-169, 170-174, 175-200 cm), parity (primiparous, multiparous), early pregnancy smoking status (non-smoker, 1-9, ≥10 cigarettes daily), educational level (<10, 10-12, >12 years), maternal country of birth (Nordic (Sweden, Denmark, Finland, Iceland, and Norway), non-Nordic), family situation (living with partner, not living with partner), and sex of offspring.

Contributors: MN and JS had full access to all of the data in the study and take full responsibility for the integrity of the data and the accuracy of the data analysis. SC, MN, BP, MP, OS, and EV conceived and designed the study. All authors acquired, analysed, and interpreted the data and critically revised the manuscript for important intellectual content: SC, MN, and MP drafted the manuscript. MN and JS carried out the statistical analysis. SC, MN, and OS obtained funding and provided administrative, technical, or material support. MN and JS are the guarantors.

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Data sharing: No additional data available.

Transparency: The lead author (MP) affirms that the manuscript is an honest, accurate, and transparent account of the study being reported; that no important aspects of the study have been omitted; and that any discrepancies from the study as planned (and, if relevant, registered) have been explained.

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**Supplementary web appendix:** eTables 1-5 and eFigures 1 and 2.