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W J C C World Journal of Clinical Cases

Contents

Thrice Monthly Volume 11 Number 30 October 26, 2023

MINIREVIEWS

| 7261 | Lower limb amputation rehabilitation status in India: A review | | | | | | |
|---------------|--|--|--|--|--|--|--|
| | Swarnakar R, Yadav SL, Surendran D | | | | | | |
| 53 (0) | | | | | | | |

Magnetic resonance imaging for acute pancreatitis in type 2 diabetes patients 7268 Ni YH, Song LJ, Xiao B

ORIGINAL ARTICLE

Retrospective Study

7277 Efficacy of lidocaine wet compress combined with red-light irradiation for chronic wounds Bao MZ, Zhou LB, Zhao L, Zhang H, Li Y, Yang L, Tai AT

- 7284 Clinical implications of forkhead box M1, cyclooxygenase-2, and glucose-regulated protein 78 in breast invasive ductal carcinoma Bai J, Li Y, Cai L
- 7294 Six-year analysis of key monitoring for bacterial strain distribution and antibiotic sensitivity in a hospital Li ZY, Yang D, Hao CH
- 7302 Clinical pharmacists' involvement in carbapenem antibiotics management at Wenzhou Integrated Hospital Xu XM, Pan CY, Zeng DL

Observational Study

High risk for obstructive sleep apnea and risk of hypertension in military personnel: The CHIEF sleep 7309 study

Liu WN, Lin KH, Tsai KZ, Chu CC, Chang YC, Kwon Y, Lin GM

EVIDENCE-BASED MEDICINE

7318 Causal relationship association of cheese intake with gestational hypertension and diabetes result from a Mendelian randomization study

Zhong T, Huang YQ, Wang GM

META-ANALYSIS

7329 Left lateral decubitus sleeping position is associated with improved gastroesophageal reflux disease symptoms: A systematic review and meta-analysis

Simadibrata DM, Lesmana E, Amangku BR, Wardoyo MP, Simadibrata M

7337 Efficacy and safety of anti-vascular endothelial growth factor agents on corneal neovascularization: A meta-analysis

Lai SC, Loh EW, Chiou DI, Hong CT



| Conter | |
|--------|--|
| | Thrice Monthly Volume 11 Number 30 October 26, 202 |
| 7350 | Efficacy and safety of different anti-osteoporotic drugs for the spinal fusion surgery: A network meta analysis |
| | He XY, Chen HX, Zhao ZR |
| | SCIENTOMETRICS |
| 7363 | Construction of clinical research nurse training program based on position competence |
| | Sun J, Shan WC, Liu JM, Zhang QQ, Ye Y, Huang ST, Zhong K |
| | CASE REPORT |
| 7372 | Fatal hemophagocytic lymphohistiocytosis-induced multiorgan dysfunction secondary to <i>Burkholderi</i> pseudomallei sepsis: A case report |
| | Sui MZ, Wan KC, Chen YL, Li HL, Wang SS, Chen ZF |
| 7380 | Interpeduncular cistern intrathecal targeted drug delivery for intractable postherpetic neuralgia: A cas report |
| | Fu F, Jiang XF, Wang JJ, Gong L, Yun C, Sun HT, Tang FW |
| 7386 | Using shape-memory alloy staples to treat comminuted manubrium sterni fractures: A case report |
| | Zhang M, Jiang W, Wang ZX, Zhou ZM |
| 7393 | Lead helix winding tricuspid chordae tendineae: A case report |
| | Liu TF, Ding CH |
| 7398 | Fournier gangrene in an infant, complicated with severe sepsis and liver dysfunction: A case report |
| | Bakalli I, Heta S, Kola E, Celaj E |
| 7403 | Prenatal ultrasound diagnosis of congenital infantile fibrosarcoma and congenital hemangioma: Three cas reports |
| | Liang RN, Jiang J, Zhang J, Liu X, Ma MY, Liu QL, Ma L, Zhou L, Wang Y, Wang J, Zhou Q, Yu SS |
| 7413 | Iatrogenic bladder neck rupture due to traumatic urethral catheterization: A case report |
| | Ekici O, Keskin E, Kocoglu F, Bozkurt AS |
| 7418 | Near obstructing painful anorectal mass and facial rash in a man with monkeypox: A case report |
| | Akpoigbe K, Yannick J, Culpepper-Morgan J |
| 7424 | Traditional Chinese medicine for foot pain in a patient with complex regional pain syndrome: A cas report |
| | Shin WC, Kim H, Chung WS |
| 7432 | Diffuse large B-cell lymphoma successfully treated with amplified natural killer therapy alone: A cas report |
| | Nagai K, Nagai S, Okubo Y, Teshigawara K |
| 7440 | Pharmacogenomics-based individualized treatment of hypertension in preterm infants: A case report an review of the literature |
| | Tang LF, Xu A, Liu K |



| World Journal of Clinical | | | | | | |
|---------------------------|--|--|--|--|--|--|
| Conte | nts Thrice Monthly Volume 11 Number 30 October 26, 2023 | | | | | |
| 7450 | Warthin-like papillary renal cell carcinoma: A case report | | | | | |
| | Li XF, Wang ZJ, Zhang HM, Yang MQ | | | | | |
| 7457 | Bladder stone due to late clip migration after prostatic urethral lift procedure: A case report | | | | | |
| | Bozkurt AS, Ekici O, Keskin E, Kocoglu F | | | | | |
| 7463 | Acute-on-chronic liver failure induced by antiviral therapy for chronic hepatitis C: A case report | | | | | |
| | Zhong JL, Zhao LW, Chen YH, Luo YW | | | | | |
| 7469 | Hemodynamic instability following intravenous dexmedetomidine infusion for sedation under brachial plexus block: Two case reports | | | | | |
| | Kim YS, Lee C, Oh J, Nam S, Doo AR | | | | | |
| 7475 | Neonatal methicillin-resistant <i>Staphylococcus aureus</i> pneumonia-related recurrent fatal pyopneumothorax: A case report and review of literature | | | | | |
| | Li XC, Sun L, Li T | | | | | |
| 7485 | Infrequent organ involvement in immunoglobulin G4-related prostate disease: A case report | | | | | |
| | Yu Y, Wang QQ, Jian L, Yang DC | | | | | |
| 7492 | Gouty tenosynovitis with compartment syndrome in the hand: A case report | | | | | |
| | Lee DY, Eo S, Lim S, Yoon JS | | | | | |
| 7497 | Acute myocardial infarction after initially diagnosed with unprovoked venous thromboembolism: A case report | | | | | |
| | Seo J, Lee J, Shin YH, Jang AY, Suh SY | | | | | |
| 7502 | Distal clavicle fractures treated by anteroinferior plating with a single screw: Two case reports | | | | | |
| | Zhao XL, Liu YQ, Wang JG, Liu YC, Zhou JX, Wang BY, Zhang YJ | | | | | |
| | | | | | | |
| | | | | | | |
| | | | | | | |
| | | | | | | |



Contents

Thrice Monthly Volume 11 Number 30 October 26, 2023

ABOUT COVER

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WJCC mainly publishes articles reporting research results and findings obtained in the field of clinical medicine and covering a wide range of topics, including case control studies, retrospective cohort studies, retrospective studies, clinical trials studies, observational studies, prospective studies, randomized controlled trials, randomized clinical trials, systematic reviews, meta-analysis, and case reports.

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EVIDENCE-BASED MEDICINE

Causal relationship association of cheese intake with gestational hypertension and diabetes result from a Mendelian randomization study

Tao Zhong, Yu-Qing Huang, Gui-Ming Wang

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Abstract

BACKGROUND

The evidence from observational studies has been inconclusive on the causal relationship between cheese intake and gestational hypertension or diabetes.

AIM

To determine whether cheese consumption was causally related to hypertension and diabetes during pregnancy.

METHODS

This was a two-sample Mendelian randomized (MR) study. Summary-level genetic data for cheese intake was exposure and corresponding outcome data for gestational hypertension and gestational diabetes were extracted from the IEU OpenGWAS database. MR analysis was conducted using inverse variance weighting. For sensitivity analyses, MR-Egger regression, weighted median, weighted mode, and leave-one-out methods were conducted. A fixed-effect model was used to meta-analyze two sample MR estimates. The traits of gestational hypertension were pregnancy hypertension (123579 individuals) and oedema, proteinuria and hypertensive disorders in pregnancy, childbirth and the puerperium (123579 individuals), and traits of gestational diabetes were gestational diabetes (123579 individuals) and diabetes mellitus in pregnancy (116363 individuals), respectively.

RESULTS

Cheese intake per standard deviation increase has causally reduced the risks of gestational hypertension [odds ratio (OR) = 0.60, 95% confidence interval (CI): 0.47-0.76, *P* < 0.001] and gestational diabetes (OR = 0.41, 95%CI: 0.30-0.55, *P* <



(0.001) in inverse variance weighted analysis. Sensitivity analysis showed no heterogeneity (all P > 0.05) nor horizontal pleiotropy (all P > 0.05) in the relationship between cheese intake and gestational hypertension, but heterogeneity presented (all P < 0.05) in relation to gestational diabetes in the two-sample MR analysis.

CONCLUSION

Cheese intake was inversely associated with gestational hypertension and gestational diabetes in MR analysis, suggesting that cheese consumption may be beneficial in preventing hypertension and diabetes during pregnancy.

Key Words: Cheese intake; Gestational hypertension; Gestational diabetes; Mendelian randomization

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Core Tip: Gestational hypertension and gestational diabetes were associated with an increased risk of complications for both the mother and fetus during pregnancy. We found that cheese intake was inversely associated with gestational hypertension and gestational diabetes in Mendelian randomization analysis, suggesting that cheese consumption may be beneficial in preventing hypertension and diabetes during pregnancy. These findings suggested that dietary interventions, especially increasing cheese intake, may be effective in the prevention gestational hypertension and gestational diabetes, and should be promoted in more regions.

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INTRODUCTION

Gestational hypertension and gestational diabetes were associated with an increased risk of complications for both the mother and fetus during pregnancy, and also been associated with an increased risk of several long-term health outcomes in pregnant women and intermediate outcomes in their children[1-5]. Although gestational hypertension and gestational diabetes posed a huge public problem to society, effective prevention has remained a major challenge in most countries, and measures were very limited. In recent years, some studies have found that dietary modification including cheesecontaining dietary pattern might play an important role in the prevention and alleviate of hypertension and diabetes during pregnancy, and this method could be implemented in maternity care[6-11]. Although several previous studies have found a link between cheese consumption and reduced risk of pregnancy-related complications, most findings were based on observational findings. However, observational studies lacked of randomization, made it difficult to effectively control for confounding, making causal relationship difficult to be established.

In recent years, the introduction of Mendelian randomized (MR) has provided an effective way to make causal inferences in observational studies, and the public release of a large amount of genome-wide association studies (GWAS) summary data has contributed to the flourishing of two-sample MR[12]. MR studies are performed by using instrumental variable (IV) analysis of genetic variation using randomization during meiosis and conception, providing unbiased and unconfounded estimates^[12]. Briefly, MR is a technique that can detect and estimate phenotypic causal effects unbiasedly [13]. With the popularity of GWAS and GWAS meta-analysis, MR has become an effective and feasible method to study causal relationships^[14]. Two-sample MR is a method for estimating the causal effect of exposure on prognosis using only summary statistics from the GWAS, in which genetic variant-exposure factor association data and genetic variant-disease prognosis association data from two independent samples with similar distribution characteristics were used[15,16]. In this study, we have performed a two-sample MR study based on public GWAS data to analyse whether cheese intake was associated with gestational hypertension or gestational diabetes.

MATERIALS AND METHODS

Overall study design

In this study, we conducted a two-sample MR analysis to explore the causal association of cheese intake with gestational hypertension and gestational diabetes in the summary level GWAS dataset, as well as the risk of gestational hypertension and gestational diabetes with single nucleotide polymorphisms (SNPs) defined as IVs (Figure 1). While all the pooled data in the present study was obtained from publicly available datasets that obtained relevant ethical approval and participant consent, the design and analysis of this study was also approved by the Ethics Committee of Guangdong Provincial People's Hospital (KY-Q-2021-244-01).



Zhong T et al. Cheese, gestational hypertension and diabetes

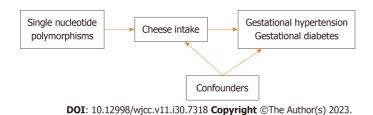


Figure 1 Flow chart of the design of the present study. The selected genetic variant (single nucleotide polymorphisms) was strongly associated with cheese intake. The selected genetic variants were strongly associated with confounding factors, but not associated with gestational hypertension gestational diabetes

Study exposure and outcomes

Summary-level genetic data for cheese intake as exposure and corresponding outcome data for gestational hypertension and gestational diabetes was extracted from the IEU OpenGWAS database (https://gwas.mrcieu.ac.uk/). The traits of gestational hypertension were characterized by gestational hypertension (123579 cases, 7686 cases/control 115893 cases) and gestational edema, proteinuria and hypertensive disorders (123579 cases, 8844 cases/control 114735 cases); gestational diabetes mellitus was characterized by gestational diabetes (123579 cases, 5687 cases/control 117892 cases) and gestational diabetes mellitus (116363 cases, 6033 cases/control 110330 cases), respectively. All the above SNPs associated with cheese intake were found in UK Biobank, and with gestational hypertension and gestational diabetes were both found in FinnGen Biobank. The detailed descriptions of included traits were summarized in Table 1. All data was downloaded and analyzed on May 20, 2023.

Mendelian randomization

The analytical strategy for this study was based on the strengthening the reporting of observational studies in epidemiology using MR statement^[17]. For all SNPs instrumental selection, genetic variants related to exposure factors with a genome-wide significant level ($P < 5 \times 10^{\circ}$), not in linkage disequilibrium (LD) at a threshold of $R^2 < 0.001$. If a particular exposure SNP was not present in an outcome dataset, proxy SNPs should be used instead through LD tagging with minimum LD R^2 value > 0.8. The causality was tested primarily by using the inverse variance weighting (IVW) method. To test for robustness, we performed sensitivity analyses using the MR-Egger regression, weighted median, and weighted mode methods. Weighted mode methods were conducted by leave one-out sensitivity analysis, in which the effect of each single instrument on the overall effect was shown in the plot of the leave-one-out analysis. Heterogeneity tests were performed using IVW and MR-Egger methods. To test for heterogeneity, we used the Cochrane Q statistic. The MR-Egger regression method could detect horizontal pleiotropy. The MR-Egger intercept was non-zero with statistical significance (P < 0.05) if possible horizontal pleiotropy of IVs existed. Scatter plots, forest plots, and funnel plots were used to present the results, And the effect of each SNP was presented in a forest plot. The MR estimates for exposure and outcomes were presented as odds ratios (OR) with corresponding 95% confidence intervals (CI). In addition, as outcomes were from different datasets, the MR outcomes were pooled meta-analysis by using a mixed-effects model to obtain an overall causal estimate, assuming no between-method heterogeneity. MR analysis was performed by using the R software (version 4.0.3, http://www.r-project.org) and the R package "TwoSampleMR" version 0.5.5 (https://mrcieu.github.io/ TwoSampleMR/).

RESULTS

MR estimates

We selected two subtype traits from FinnGen Biobank database with a genotype of gestational diabetes (finn-b-GEST_DIABETES and finn-b-O15_PREG_DM), gestational hypertension (finn-b-O15_HYPTENSPREG and finn-b-O15_OEDEM_PROTUR_HYPERT) and one group with a trait of cheese intake from UK Biobank database (ukb-b-1489). All participants were of European ethnicity. The numbers of SNPs related to cheese intake were 9851867, and those for gestational hypertension and gestational diabetes were both 16379784. The detailed descriptions of included traits were summarized in Table 1.

The two-sample MR estimates for the association of cheese intake with gestational hypertension and gestational diabetes were summarized in Table 2. The MR analyses on the result of the IVW method showed an inversely causal effect of genetically predicted cheese intake with pregnancy hypertension (OR = 0.57, 95%CI: 0.40-0.80; P = 1.377e-3), oedema, proteinuria and hypertensive disorders in pregnancy, childbirth and the puerperium (OR = 0.63, 95% CI: 0.46-0.88; P = 0.006), gestational diabetes (OR = 0.39, 95% CI: 0.26-0.60; P = 1.822e-5) and diabetes mellitus in pregnancy (OR = 0.43, 95% CI: 0.28-0.67; P = 1.718e-4) by using 62 SNPs as the instruments, respectively. The weighted median estimator showed consistent result when the study outcomes were pregnancy hypertension (OR = 0.55, 95% CI: 0.34-0.90; P = 0.017), gestational diabetes (OR = 0.40, 95%CI: 0.23-0.71; P = 1.507e-3) and diabetes mellitus in pregnancy (OR = 0.35, 95%CI: 0.20-0.62; P = 2.998e-4), but not consistent of oedema, proteinuria and hypertensive disorders in pregnancy, childbirth and the puerperium (OR = 0.69, 95% CI: 0.45-1.05; P = 0.083). In addition, although this negative correlation could also be observed in other methods including MR Egger and Weighted mode, it was not statistically significant (all P > 0.05)



| Table 1 Description of included traits | | | | | | | | | |
|--|--|------------------------------------|-------------------|------------|----------------|--------------|--------------------|------------|------|
| Exposure | Trait | ID | Number of SNPs | Population | Sample size | Case/control | Database | Consortium | Year |
| Cheese intake | Cheese intake | ukb-b-1489 | 9851867 | European | 451486 | NA | UK Biobank | MRC-IEU | 2018 |
| Outcomes | | | | | | | | | |
| Gestational hypertension | Pregnancy hypertension | finn-b-O15_HYPTENSPREG | 16379784 | European | 123579 | 7686/115893 | FinnGen Biobank | NA | 2021 |
| | Oedema, proteinuria and hypertensive disorders in pregnancy, childbirth and the puerperium | finn-b- O15_OEDEM_PROTUR_HYPERT | 16379784 | European | 123579 | 8844/114735 | FinnGen Biobank | NA | 2021 |
| Gestational diabetes | Gestational diabetes (for exclusion) | finn-b-GEST_DIABETES | 16379784 | European | 123579 | 5687/117892 | FinnGen Biobank | NA | 2021 |
| | Diabetes mellitus in pregnancy | finn-b-O15_PREG_DM | 16379684 | European | 116363 | 6033/110330 | FinnGen Biobank | NA | 2021 |

SNP: Single nucleotide polymorphism; NA: Not available; MRC-IEU: MRC integrative epidemiology unit.

(Table 2). A meta-analysis using fixed-effect IVW models showed that genetically predicted cheese intake was inversely associated with the risk of gestational hypertension (OR = 0.60, 95% CI: 0.47-0.76; P < 0.0001) and gestational diabetes (OR = 0.41, 95% CI: 0.30-0.55; P < 0.0001) (Figure 2). The SNP effects on the outcome were plotted against SNP effects on the exposure were displayed in Figure 3.

Sensitivity analyses

MR-Egger intercept tests were used to test for pleiotropy, indicating that there was no horizontal pleiotropy existed on the association between cheese intake and gestational hypertension or gestational diabetes (all P > 0.05). Heterogeneity tests demonstrated that there was no heterogeneity between cheese intake and gestational hypertension (all P > 0.05), but heterogeneity was observed in the Q test analysis on the association between cheese intake and gestational diabetes (all P < 0.05) (Table 2). Funnel plots can show the directional horizontal pleiotropy of IVs by drawing a single Wald ratio for each SNP. Sensitivity analyses using the leave-one-out analysis demonstrated that there was significant association of cheese intake with gestational hypertension or gestational diabetes (Figure 4). The overall estimates of MR effect size, calculated by IVW showed significant associations of cheese intake with gestational hypertension or gestational diabetes, but the MR-Egger method was not statistically significant despite finding also a negative correlation (Figure 5).

DISCUSSION

In this two-sample MR study, based on pooled statistics from a large GWAS, we clarified that cheese intake could reduce the risk of hypertension and diabetes during pregnancy. A pooled meta-analysis based on the results of the IVW method, using a fixed effects model to calculate the overall causal effect, showed that cheese intake was significantly and inversely associated with gestational hypertension and gestational diabetes. Heterogeneity and horizontal polymorphism in gestational hypertension were not found in the two-sample MR analysis, but heterogeneity in gestational diabetes was present.

Our results were similar to those of some previous observational or small sample randomised controlled trials (RCTs) that cheese intake might prevent gestational hypertension and gestational diabetes during pregnancy. Chinese intake in Canadian and Iranian women prior to pregnancy have been shown to be an important intervention target to reduce the likelihood of pregnancy-related complications, such as gestational hypertension[11,18]. According to a study of pregnant United States women, consuming more cheese before and during pregnancy could alleviate or reduce gestational diabetes risk[6,7]. Taking cheese during pregnancy might lead to modest dietary improvements in pregnant women at high risk of gestational diabetes, according to the Finnish gestational diabetes prevention study [8,9,19]. A small sample RCT study suggested that an increased intake of low-fat but not regular-fat cheese between pre-pregnancy and early pregnancy was associated with a lower risk of gestational diabetes in high-risk women[20]. However, despite an association between cheese consumption and gestational diabetes, a study in Singapore did not find a significant association[21]. Previous results differed from the present results mainly because some potential confounding variables

Table 2 Mendelian randomization for the association of cheese intake with gestational hypertension and gestational diabetes

| | Trait | Number of instruments | Method C | | 95%CI | P value | Heterogeneity | | Pleiotropy | |
|--------------------------|---|-----------------------|---------------------------------|------|---------------|--------------|----------------|--------------|------------|------------|
| Outcomes | | | | OR | | | Q statistic | P value | Intercept | P value |
| Gestational hypertension | Pregnancy hypertension | 62 | MR Egger | 0.40 | 0.09- 1.75 | 0.229 | 73.04 | 0.120 | 0.006 | 0.634 |
| | | 62 | Weighted median | 0.55 | 0.34- 0.90 | 0.017 | | | | |
| | | 62 | Inverse variance weighted | 0.57 | 0.40- 0.80 | 1.377e- 3 | 73.31 | 0.134 | | |
| | | 62 | Weighted mode | 0.49 | 0.20- 1.19 | 0.121 | | | | |
| | Oedema, proteinuria and hypertensive disorders in pregnancy, childbirth and the puerperium | 62 | MR Egger | 0.62 | 0.15- 2.52 | 0.505 | 76.44 | 0.075 | 4.3e-4 | 0.971 |
| | | 62 | Weighted median | 0.69 | 0.45- 1.05 | 0.083 | | | | |
| | | 62 | Inverse variance weighted | 0.63 | 0.46- 0.88 | 0.006 | 76.44 | 0.088 | | |
| | | 62 | Weighted mode | 0.69 | 0.28- 1.68 | 0.416 | | | | |
| Gestational diabetes | Gestational diabetes (for exclusion) | 62 | MR Egger | 0.23 | 0.04- 1.38 | 0.113 | 83.73 | 0.023 | 9.5 e-3 | 0.537 |
| | | 62 | Weighted median | 0.40 | 0.23- 0.71 | 1.507e- 3 | | | | |
| | | 62 | Inverse variance weighted | 0.39 | 0.26- 0.60 | 1.822e- 5 | 84.26 | 0.025 | | |
| | | 62 | Weighted mode | 0.32 | 0.10- 1.03 | 0.062 | | | | |
| | Diabetes mellitus in pregnancy | 62 | MR Egger | 0.46 | 0.07- 3.00 | 0.418 | 91.39 | 5.586e- 3 | -0.001 | 0.948 |
| | | 62 | Weighted median | 0.35 | 0.20- 0.62 | 2.998e- 4 | | | | |
| | | 62 | Inverse variance weighted | 0.43 | 0.28- 0.67 | 1.718 e-4 | 91.4 | 7.089e- 3 | | |
| | | 62 | Weighted mode | 0.21 | 0.04- 1.20 | 0.085 | | | | |

CI: Confidence interval; OR: Odds ratio; MR: Mendelian randomization.

were not adequately adjusted. However, in this study, the influence of confounding variables on the results was overcome through the two-sample MR study. In addition, although this study found heterogeneity in SNPs among gestational diabetes, this may be due to part of gestational diabetes itself existed gene mutation including pregnancy in women with monogenic diabetes or maternal genetic characteristics. Previous studies suggested that monogenic diabetes was an underdiagnosed type of diabetes mellitus, which could be harmful in pregnancy and was easily misdiagnosed as gestational diabetes[22-24]. However, the present study does not distinguish whether pregnant women have monogenic diabetes, missing data on monogenic diabetes was also an important shortcoming of this study.

In addition, this study found no horizontal pleiotropy in SNPs, but some of the sensitivity analysis results were not statistically significant, which may be due to the presence of other pleiotropy bias in some SNPs. However, a metaanalysis of fixed effects models showed that cheese intake was inversely associated with gestational diabetes and gestational hypertension with no heterogeneity. The fixed-effect meta-analysis model used an inverse-variance weight (variance of the observed effect size) and assumed that all studies share a single common effect and, as a result, all of the variance in observed effect sizes was attributable to sampling error[25]. These analytical methods, such as MR analysis, leave-one-out sensitivity analysis and pooled analysis of fixed effects models ensured the reliability of the results in the present MR study, indicating that cheese consumption could help preventing gestational diabetes and hypertension, but warranted further research to understand underlying pathophysiology of gestational diabetes and hypertension

| Outcomes | Data source | | OR (95%CI) | P value |
|--------------------------------|-----------------|--------------------------|-----------------------|-------------|
| Gestational hypertension | | 1 | | |
| Pregnancy hypertension | FinnGen Biobank | _ | 0.57 (0.40 to 0.80) | 1.4E-3 |
| Hypertensive in pregnancy | FinnGen Biobank | _ - | 0.63 (0.46 to 0.88) | 6.5E-3 |
| Meta-analysis | | _ | 0.60 (0.47 to 0.76) | < 0.0001 |
| Gestational diabetes | | | | |
| Gestational diabetes | FinnGen Biobank | | 0.39 (0.26 to 0.60) | < 0.0001 |
| Diabetes mellitus in pregnancy | FinnGen Biobank | | 0.43 (0.28 to 0.67) | < 0.0001 |
| Meta-analysis | | | 0.41 (0.30 to 0.55) | < 0.0001 |
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Figure 2 Forest plot showing the main Mendelian randomization estimates for the association of cheese intake with gestational hypertension and gestational diabetes by a meta-analysis of fixed-effect model. OR: Odds ratio; CI: Confidence interval.

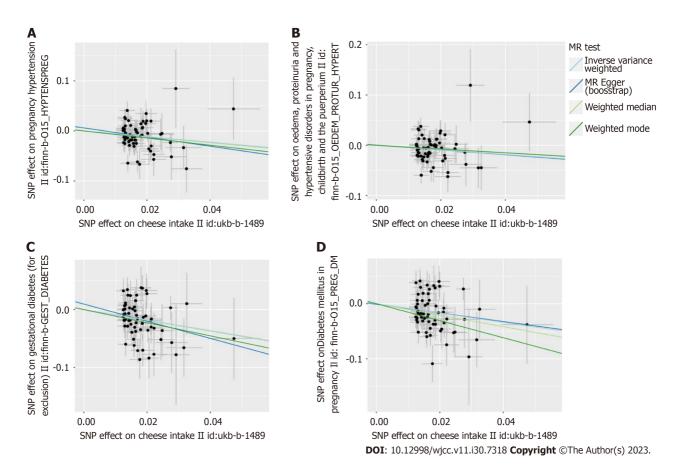


Figure 3 Scatter plot to visualize causal effect of cheese intake on gestational hypertension and gestational diabetes risk using four Mendelian randomization methods. The slope of the straight line indicates the magnitude of the causal association. The plot presents the effect sizes of the single nucleotide polymorphisms (SNP)-cheese intake association (X-axis, standard deviation units) and the SNP-gestational hypertension and gestational diabetes association [Y-axis, log (odds ratio)] with 95% confidence intervals. The regression slopes of the lines correspond to causal estimates using the four Mendelian randomization methods. A: The Scatter plot visualize causal effect of cheese intake on gestational hypertension risk in the FinnGen Biobank database; B: The Scatter plot visualize causal effect of cheese intake on gestational diabetes risk in the FinnGen Biobank database; D: The Scatter plot visualize causal effect of cheese intake on gestational diabetes risk in the FinnGen Biobank database. SNPs: Single nucleotide polymorphisms; MR: Mendelian randomization.

associated with dietary behaviors during pregnancy, such as increased cheese intake.

The possible mechanisms linking cheese intake to a reduced risk of hypertension and diabetes in pregnancy were unclear and we speculated that there were several possible reasons for this. Firstly, cheese was rich in calcium, and previous meta-analysis studies have found that calcium supplementation in pregnancy has been associated with a reduced risk of pregnancy-induced hypertension and diabetes, but further high-quality evidence was needed[26,27]. Secondly, cheese contained lots of lactic acid bacteria, which were healthy for the body and could maintain intestinal flora balance[28]. It was important to note that intestinal flora imbalance was an important cause of gestational hypertension and gestational diabetes during pregnancy[29-31]. Thirdly, the benefits of eating cheese could counteract the negative effects of its high saturated fat content. This was because cheese not only contained nutrients such as protein, calcium,

Zhong T et al. Cheese, gestational hypertension and diabetes

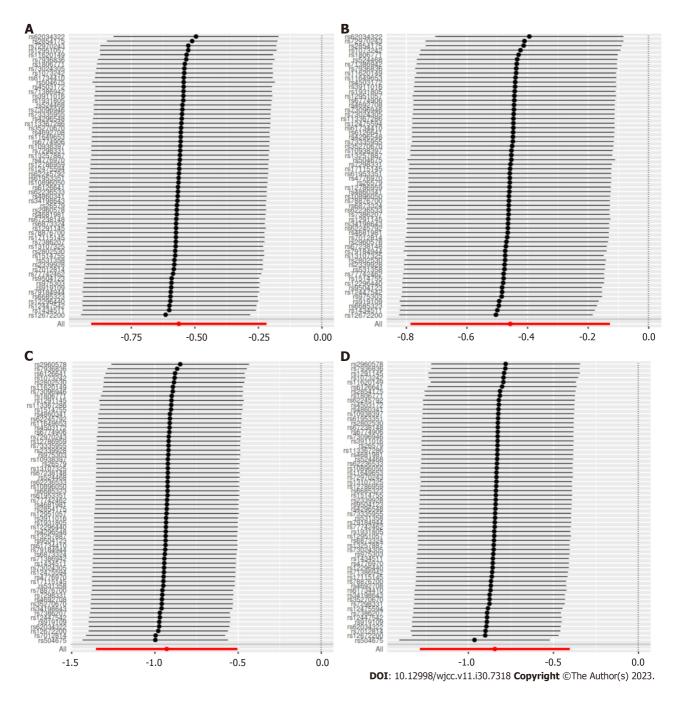


Figure 4 Leave-one-out of each single nucleotide polymorphisms associated of cheese intake with gestational hypertension and gestational diabetes risk. Each black point represents result of the inverse variance weighted (IVW) Mendelian randomization method applied to estimate the causal effect of cheese intake on gestational hypertension and gestational diabetes excluding particular single nucleotide polymorphism (SNP) from the analysis. Each red point depicts the IVW estimate using all SNPs. A: Mendelian randomization (MR) leave-one-out sensitivity analysis of gestational hypertension risk in FinnGen Biobank database; B: MR leave-one-out sensitivity analysis of gestational hypertension in FinnGen Biobank database; C: MR leave-one-out sensitivity analysis of gestational diabetes in FinnGen Biobank database; D: MR leave-one-out sensitivity analysis of gestational diabetes in FinnGen Biobank database.

iron, zinc, vitamins A, B1, B2 and folic acid, but also probiotics, which would reduce the state of inflammation and oxidative stress in the body [32,33]. Inflammation and oxidative stress were both closely related to gestational hypertension and gestational diabetes[34,35]. In addition, cheese also contained conjugated linoleic acid, an unsaturated fatty acid that could raise the level of high density lipoprotein cholesterol and lower the level of low density lipoprotein cholesterol^[36]. Previous studies have shown that specific lipid biomarkers in early pregnancy may be associated with gestational hypertension and gestational diabetes [37,38]. Finally, animal experiments have shown the consumption of goat milk cheese could prevent obesity, insulin resistance, inflammation, and hepatic steatosis while on a high-fat diet induced obesity in mice[39].

There were several strengths to the current study. On the one hand, it was the first study to investigate the causal relationship between cheese consumption and gestational diabetes and hypertension through a MR analysis. On the other hand, a key strength of MR analysis was the use of randomly allocated genetic variants to help overcome environmental confounding, which was analogous to randomization of treatment allocation in clinical trials. In addition, a two-sample

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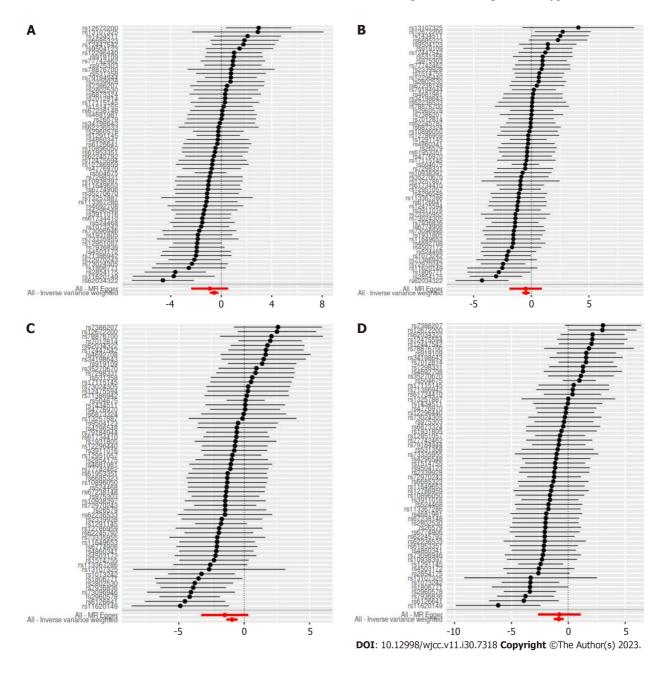


Figure 5 Forest plot showing the association of each single nucleotide polymorphisms with gestational hypertension and gestational diabetes. Each black point represents result of the inverse variance weighted (IVW) Mendelian randomization (MR) method applied to estimate the causal effect of each single nucleotide polymorphism (SNP) on gestational hypertension and gestational diabetes. Each red point depicts the IVW and MR egger estimate using all SNPs. A: Forest plot of gestational hypertension in FinnGen Biobank database; B: Forest plot of gestational hypertension in FinnGen Biobank database; C: Forest plot of gestational diabetes in FinnGen Biobank database; D: Forest plot of gestational diabetes in FinnGen Biobank database.

MR design and high-quality GWAS summary statistics based on a large sample size was also an advantage of this study. Finally, to ensure the stability of the results, MR-Egger regression tests were performed, and no pleiotropy were found. Of course, several limitations of the present MR study should also be considered. Firstly, it should be noted that the study population was all of European ancestry, so the results cannot be generalized to other races. Secondly, self-reported information contributed to some of history, so recalled bias cannot be eliminated. Thirdly, although IVW, MR-Egger regression, weighted median and weighted mode methods found an inverse association of cheese intake with gestational diabetes and gestational hypertension in MR analysis, the results of some methods were not statistically significant. Fourthly, even though we found some heterogeneity in gestational diabetes, our fixed-effect meta-analysis found that cheese intake was significantly associated with gestational diabetes according to IVW results. Fifthly, genetic polymorphisms were difficult to validate, and even using the MR-Egger method, misclassification cannot be completely ruled out. In addition, we were currently unable to obtain the type of cheese and the specific amount of the daily intake, as well as neither cheese intake nor gestational hypertension or gestational diabetes could be examined for their potential doseeffect relationships. Finally, the individual-level data on gestational diabetes and gestational hypertension were unavailable, therefore stratified subgroup analyses could not be conducted. As a result, more MR studies were needed to investigate the causal association of cheese intake with gestational diabetes and hypertension in the future.

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CONCLUSION

The present study indicated an inversely genetic causal association of cheese intake prior to pregnancy with pregnancy complications such as gestational hypertension and gestational diabetes during pregnancy, which suggesting that the consumption of cheese during pregnancy may contribute to the prevention of gestational hypertension and gestational diabetes. These findings suggested that dietary interventions, especially increasing cheese intake, may be effective in the prevention gestational hypertension and gestational diabetes, and should be promoted in more regions.

ARTICLE HIGHLIGHTS

Research background

Evidence from observational studies has not been able to establish a causal association of cheese intake with hypertension or diabetes during pregnancy.

Research motivation

Eating cheese during pregnancy may help prevent hypertension and diabetes.

Research objectives

The objective was to determine whether cheese consumption was causally related to hypertension and diabetes during pregnancy.

Research methods

A Mendelian randomised (MR) study with two samples was conducted. The IEU OpenGWAS database's corresponding outcome data for gestational diabetes and gestational hypertension were taken out, and summary-level genetic information for cheese consumption was exposed. MR analysis was performed by using inverse variance weighting as the main method. Methods used for sensitivity studies included MR-Egger regression, weighted median, weighted mode, and leave-one-out techniques. The meta-analysis of two sample MR estimations was performed using a fixed-effect model. The characteristics of gestational diabetes were gestational diabetes (123579 individuals) and diabetes mellitus in pregnancy (116363 individuals), whereas the characteristics of gestational hypertension were pregnancy hypertension (123579 individuals) and oedema, proteinuria, and hypertensive disorders in pregnancy, childbirth, and the puerperium (123579 individuals).

Research results

Inverse variance weighted analysis has shown a causal relationship between cheese consumption per standard deviation increase and the risks of gestational hypertension and gestational diabetes (odds ratio = 0.41, 95% confidence interval: 0.30-0.55, P < 0.001). The two-sample MR analysis of the relationship between cheese intake and gestational hypertension revealed no heterogeneity (all P > 0.05) or horizontal pleiotropy, but there was heterogeneity (all P > 0.05) in relation to gestational diabetes.

Research conclusions

In this MR analysis, cheese consumption was found to be inversely related to gestational hypertension and gestational diabetes, implying that cheese intake may be beneficial in preventing gestational hypertension and gestational diabetes.

Research perspectives

These findings indicated that dietary interventions, particularly increasing cheese consumption, could be effective in preventing hypertension and diabetes during pregnancy, and should be promoted in more areas.

FOOTNOTES

Author contributions: All authors contributed to the design; Huang YQ contributed to analysis of the study; all authors prepared the written manuscript; and all authors critically reviewed and edited the manuscript.

Conflict-of-interest statement: All the authors report no relevant conflicts of interest for this article.

PRISMA 2009 Checklist statement: The authors have read the PRISMA 2009 Checklist, and the manuscript was prepared and revised according to the PRISMA 2009 Checklist.

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