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The primary aim of *World Journal of Gastrointestinal Oncology* (WJGO, *World J Gastrointest Oncol*) is to provide scholars and readers from various fields of gastrointestinal oncology with a platform to publish high-quality basic and clinical research articles and communicate their research findings online.

WJGO mainly publishes articles reporting research results and findings obtained in the field of gastrointestinal oncology and covering a wide range of topics including liver cell adenoma, gastric neoplasms, appendiceal neoplasms, biliary tract neoplasms, hepatocellular carcinoma, pancreatic carcinoma, cecal neoplasms, colonic neoplasms, colorectal neoplasms, duodenal neoplasms, esophageal neoplasms, gallbladder neoplasms, *etc.*

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Clinical and Translational Research

Causal effects of genetic birth weight and gestational age on adult esophageal diseases: Mendelian randomization study

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Abstract

BACKGROUND

Few studies have investigated the association between gestational age, birth weight, and esophageal cancer risk; however, causality remains debated. We aimed to establish causal links between genetic gestational age and birth weight traits and gastroesophageal reflux disease (GERD), Barrett's esophagus (BE), and esophageal adenocarcinoma (EA). Additionally, we explored if known risk factors mediate these links.

AIM

To analyze of the relationship between gestational age, birth weight and GERD, BE, and EA.

METHODS

Genetic data on gestational age and birth weight ($n = 84689$ and 143677) from the Early Growth Genetics Consortium and outcomes for GERD ($n = 467253$), BE ($n = 56429$), and EA ($n = 21271$) from genome-wide association study served as instrumental variables. Mendelian randomization (MR) and mediation analyses were conducted using MR-Egger, weighted median, and inverse variance weighted methods. Robustness was ensured through heterogeneity, pleiotropy tests, and sensitivity analyses.

RESULTS

Birth weight was negatively correlated with GERD and BE risk [odds ratio (OR) = 0.78; 95% confidence interval (CI): 0.69-0.8] and (OR = 0.75; 95%CI: 0.60-0.9), respectively, with no significant association with EA. No causal link was found between gestational age and outcomes. Birth weight was positively correlated with five risk factors: Educational attainment (OR = 1.15; 95%CI: 1.01-1.31), body mass index (OR = 1.06; 95%CI: 1.02-1.1), height (OR = 1.12; 95%CI: 1.06-1.19), weight (OR = 1.13; 95%CI: 1.10-1.1), and alcoholic drinks per week (OR = 1.03; 95%CI: 1.00-1.06). Mediation analysis showed educational attainment and height mediated the birth weight-BE link by 13.99% and 5.46%.

CONCLUSION

Our study supports the protective role of genetically predicted birth weight against GERD, BE, and EA, independent of gestational age and partially mediated by educational attainment and height.

Key Words: Birth weight; Barrett's esophagus; Esophageal adenocarcinoma; Gestational age; Gastroesophageal reflux disease; Preterm birth

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Core Tip: Our study demonstrates that low birth weight, rather than prematurity, is associated with increased risk of gastroesophageal reflux disease (GERD) and Barrett's esophagus (BE) in adulthood, with no association found for esophageal adenocarcinoma. Furthermore, this relationship is influenced by the mediating effects of educational attainment and height. Attention to educational attainment and height during the growth process of low-birth-weight individuals is necessary to reduce the incidence of GERD and BE.

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INTRODUCTION

Esophageal cancer, ranking as the eighth most prevalent cancer globally, exhibits a 5-year survival rate of less than 20%. In 2020, over 604000 new cases were diagnosed worldwide[1,2]. Esophageal adenocarcinoma (EA), as one subtype of esophageal cancer, has witnessed a rapid increase in incidence in Western countries in recent years[3,4]. Barrett's esophagus (BE), characterized by the metaplasia of esophageal squamous epithelium, serves as a precancerous lesion and a high-risk factor for EA. Approximately 3%-5% of BE cases eventually progress to EA[5]. The risk of BE development is associated with recurrent episodes of gastroesophageal reflux disease (GERD), affecting approximately 20% of adults in developed countries[6]. Furthermore, previous research indicates that educational level, socioeconomic status, mental health, obesity, and smoking contribute to varying degrees of EA risk[7-11].

In contemporary research, there is considerable attention on the association between perinatal factors and adult-onset diseases[12]. GERD, a prevalent condition in newborns, exhibits notably elevated risks in preterm infants and those with small-for-gestational-age (SGA)[13]. The debate continues regarding the correlation of preterm birth and SGA with adult-onset GERD, BE, or EA. In a large-scale follow-up study of preterm and SGA individuals, only 8 participants developed esophageal cancer during adulthood, while the standardized incidence of EA increased by more than 7-fold[14]. Conversely, an alternative perspective posits that preterm birth is the primary factor contributing to the elevated risk of EA in adulthood, with SGA having no impact on this risk[15]. In investigations into the risk of developing BE in adulthood, both preterm and SGA individuals demonstrate a significant association with adult BE risk[16]. Notably, in the Swedish BE population, it was observed that SGA individuals, rather than preterm ones, experienced a threefold increase in the risk of developing BE in adulthood[17].

Due to the significant time span from infancy to adulthood, participants' lifestyle habits (smoking, alcohol consumption, obesity, *etc.*) may impact the ultimate development of esophageal diseases through alternative causal pathways. As a result, traditional cohort studies and retrospective research face substantial limitations in this process, potentially leading to misleading interpretations. The principle of Mendelian randomization (MR), involving the random assortment of genes from parents to offspring during gamete formation and fertilization, can significantly mitigate measurement errors and confounding pathways. MR provides an effective means of evaluating the causal nature of modifiable behaviors or environmental exposures[18,19].

We employed genetic information from a comprehensive alliance of European populations to elucidate the causal associations between fetal prenatal characteristics, birth weight, and the risk of GERD, BE, and EA in adulthood[20-29].

Our methodology incorporated a two-step MR approach, encompassing univariable MR (UVMR) analysis and multivariable MR (MVMR) analysis, mediation analyses, and sensitivity assessments[30]. The primary objective of this study extends beyond elucidating direct causal relationships to actively seeking intermediary factors influencing the development of esophageal diseases. This research endeavors to contribute novel insights for future investigations and clinical practices related to GERD, BE, and EA.

MATERIALS AND METHODS

Study design

Multiple publicly available datasets from genome-wide association study (GWAS) were applied in this study[20-29]. All studies included in the analysis were ethically approved or patient consent, as described in Table 1. The univariate, multivariate, and two-step MR mediating analyses were performed to investigate whether genetically predicted gestational age characteristics, fetal birth weight, and adult susceptibility to GERD, BE, and EA were causally related. An attempt was made to find the mediating risk factors in the above associations and to calculate the corresponding mediating effect size. The route of this study is shown in Figure 1.

GWAS data for exposure

Genetic variation information pertaining to 84689 individuals of European descent, specifically related to the timing of delivery, was sourced from the collaborative efforts of the Early Growth Genetics (EGG) Consortium and the Integrative Psychiatric Research Consortium[25]. This dataset encompasses four fetal traits associated with the fetal genotype. The logistic regression analysis was applied to examine dichotomous outcomes, namely early preterm birth (< 34 wk), preterm birth (< 37 wk), postterm birth (\geq 42 wk). Additionally, gestational duration, subjected to regression on infant sex, had its resulting residuals quantile transformed to a standard normal distribution before undergoing linear regression to assess their association with fetal single nucleotide polymorphisms (SNPs) genotypes. Following gender-specific transformation and adjustments for gestational week and study-specific ancestry covariates, the genetic information pertaining to birth weight was included in the study only for a European population of 143677 individuals from the EGG Consortium[27].

GWAS data for mediator

In previous observational studies, risk factors associated with GERD, BE, and EA have been considered as potential mediators. Utilizing the most recent and largest-scale European GWAS meta-analysis data, we incorporated eight variables, namely, educational attainment (years of education), body mass index (BMI), height, weight, major depressive disorder (MDD), cigarettes smoked per day, alcoholic drinks per week, and *Helicobacter pylori* (*H. pylori*) infection. Genetic information for educational attainment (years of education) was sourced from Loh *et al*[24], employing a mixed-model association approach with a cohort of 461457 European individuals. The genetic characteristics of BMI were derived from the Genetic Investigation of ANthropometric Traits Consortium, encompassing a large-scale meta-analysis of 681275 individuals of European descent[20]. Height's genetic data originated from an exome array association study based on 458235 Europeans from the United Kingdom Biobank[29]. Weight's genetic summary data involved 360116 individuals of European descent[23]. Genetic information for MDD was obtained from the Psychiatric Genomics Consortium through a comprehensive genome-wide association meta-analysis with 59851 cases and 113154 controls[21]. Daily smoking quantity and weekly alcohol consumption data were sourced from the GWAS and Sequencing Consortium of Alcohol and Nicotine use, comprising 337334 and 335394 European individuals, respectively[26]. The genetic characteristics of *H. pylori* infection were derived from GWAS involving 1329 cases and 461604 controls of European ancestry[31,32].

GWAS data for outcome

The summary data for GERD (32957 cases and 434296 controls) was extracted from a large-scale human genetic phenotypic study conducted by Sakaue *et al*[23]. The genetic characteristics of BE (13358 cases and 43071 controls) were amalgamated from a comprehensive data synthesis derived from a large prospective etiological study of complex diseases in middle and older-aged individuals by Sudlow *et al*[22]. As for EA (4112 cases and 17159 controls), the genetic features originated from a large-scale meta-analysis of the entire genome-wide association data. All participants involved in the GERD, BE, and EA studies were of European descent[28].

Statistical analysis

Selection of genetic instrumental variables: Initially, due to stringent criteria ($P < 5E-8$), the genomic significant SNPs acquired for four fetal gestational features (preterm birth, preterm birth, postterm birth, gestational duration) did not yield sufficient power for subsequent MR analyses. Consequently, a more lenient screening threshold ($P < 5E-6$) was incorporated, which has been demonstrated as feasible in previous studies[33]. Similarly, a $P < 5E-5$ threshold was employed in constructing instrumental variables (IVs) for *H. pylori* infection.

Subsequently, SNPs failing to meet the independence criterion were removed through linkage disequilibrium (LD) to obtain independently associated SNPs ($r^2 < 0.001$, kb = 10000, MAF < 0.3). Proxy SNPs ($r^2 > 0.8$) were utilized to address the inability to extract SNPs associated with exposure or mediating IVs in the outcomes of esophageal diseases, achieved through the LD link tool[34]. In this process, SNPs with ambiguous palindromic sequences were excluded.

Table 1 Genome-wide association study data sources and information included in the current study

Phenotype	Unit	Sample size	Ancestry	GWAS ID/data source	PMID
Educational attainment (years of education)	SD increase in years of education	461457	European	ebi-a-GCST90029013	29892013
Body mass index	SD increase in body mass index	681275	European	ieu-b-40	30124842
Height	SD increase in cm of height	458235	European	ebi-a-GCST90025949	34226706
Weight	SD increase in kg of birth adult weight	360116	European	ebi-a-GCST90018949	34594039
Major depressive disorder	Binary categorical variable	173005	European	ieu-a-1188	29700475
Cigarettes smoked per day	SD increase in cigarettes smoked per day	337334	European	ieu-b-25	30643251
Alcoholic drinks per week	SD increase in alcoholic drinks per week	335394	European	ieu-b-73	30643251
Birth weight	SD increase in kg of birth weight	143677	European	ieu-a-1083	27680694
Gestational duration	SD increase in days of gestational duration	84689	European	EKG Consortium	31477735
Early preterm birth	Binary categorical variable	38995	European	EKG Consortium	31477735
Preterm birth	Binary categorical variable	57153	European	EKG Consortium	31477735
Post term birth	Binary categorical variable	60695	European	EKG Consortium	31477735
<i>Helicobacter pylori</i>	Binary categorical variable	462933	European	ukb-b-531	/
Esophageal adenocarcinoma	Binary categorical variable	21271	European	ebi-a-GCST003739	27527254
Barrett's esophagus	Binary categorical variable	56429	European	ebi-a-GCST90000515	25826379
Gastroesophageal reflux disease	Binary categorical variable	467253	European	ebi-a-GCST90018848	34594039

EKG: Early Growth Genetics; GWAS: Genome-wide association study.

Based on previous evidence from observational studies, educational attainment, BMI, smoking, alcohol consumption, depression, or anxiety are considered confounding variables that may violate the assumption of causality[8-10,35]. To mitigate potential horizontal pleiotropy and violations of the correlation assumption in the study of exposure on outcomes, the PhenoScanner network tool was employed to exclude confounding SNPs that might represent alternative pathways, aligning with the exclusivity assumption in the core principles of MR[36]. The MR-PRESSO test was utilized to detect and remove outliers[37], resulting in the final set of IVs for MR analysis. To ensure the adherence to the core assumptions of MR, specifically the exclusion restriction assumption, identified IVs underwent scrutiny with criteria including *F*-statistic values > 10 and statistical power > 0.8, significantly minimizing potential bias introduced by weak IVs[38-40].

UVMR analysis: We conducted multiple rounds of UVMR analyses. Initially, causal estimates were performed for gestational duration-related genetic characteristics, birth weight, and their associations with GERD, BE, and EA. Subsequently, significant fetal traits identified in the initial analysis were included in subsequent UVMR analyses involving eight risk factors. The causal effects between these eight risk factors and esophageal diseases were also estimated using UVMR. Each UVMR analysis underwent tests for heterogeneity and pleiotropy. If the MR-Egger intercept test indicated horizontal pleiotropy ($P < 0.05$)[41], the MR-Egger regression method was employed to mitigate bias introduced by pleiotropy[42]; alternatively, in the absence of such indication, the initial consideration was given to the inverse-variance weighted (IVW) method[43]. If Cochran's *Q* statistic suggested the presence of heterogeneity ($P < 0.05$)[30], the IVW fixed-effects model, rather than the IVW random-effects model, was utilized to minimize bias.

MVMR analysis: To accurately assess the effects of eight risk factors on three esophageal diseases, MVMR analysis method was employed[19]. Adjustments were made for significant neonatal characteristics identified in UVMR analysis. All relevant SNPs were included under a threshold of $P < 5E-8$, and a reevaluation of LD, alignment of effect alleles, removal of palindromic sequences, and multivariate tests for heterogeneity and pleiotropy were conducted.

Mediation analysis: As depicted in the intermediate analysis steps illustrated in Figure 1, UVMR initially computed the total effect of exposure on the outcome (β_c), the direct effect of exposure on the mediator (β_a), and the causal effect of the mediator on the outcome. The subsequent MVMR revealed the results of analyzing the direct effect of exposure on the mediator (β_b) after adjusting for the exposure[19]. The mediation effect was considered established when both the total effect and the mediator effect ($\beta_a \times \beta_b$) acted in the same direction and remained statistically significant[44]. Subsequently, we calculated the proportion of mediation by dividing the mediator effect by the total effect. The delta method was employed to estimate confidence intervals for independent samples[44,45]. Through the integration of mediation analysis with MVMR, concurrently adjusting for multiple intermediary risk factors, we derived an estimation of the comprehensive mediation effect.

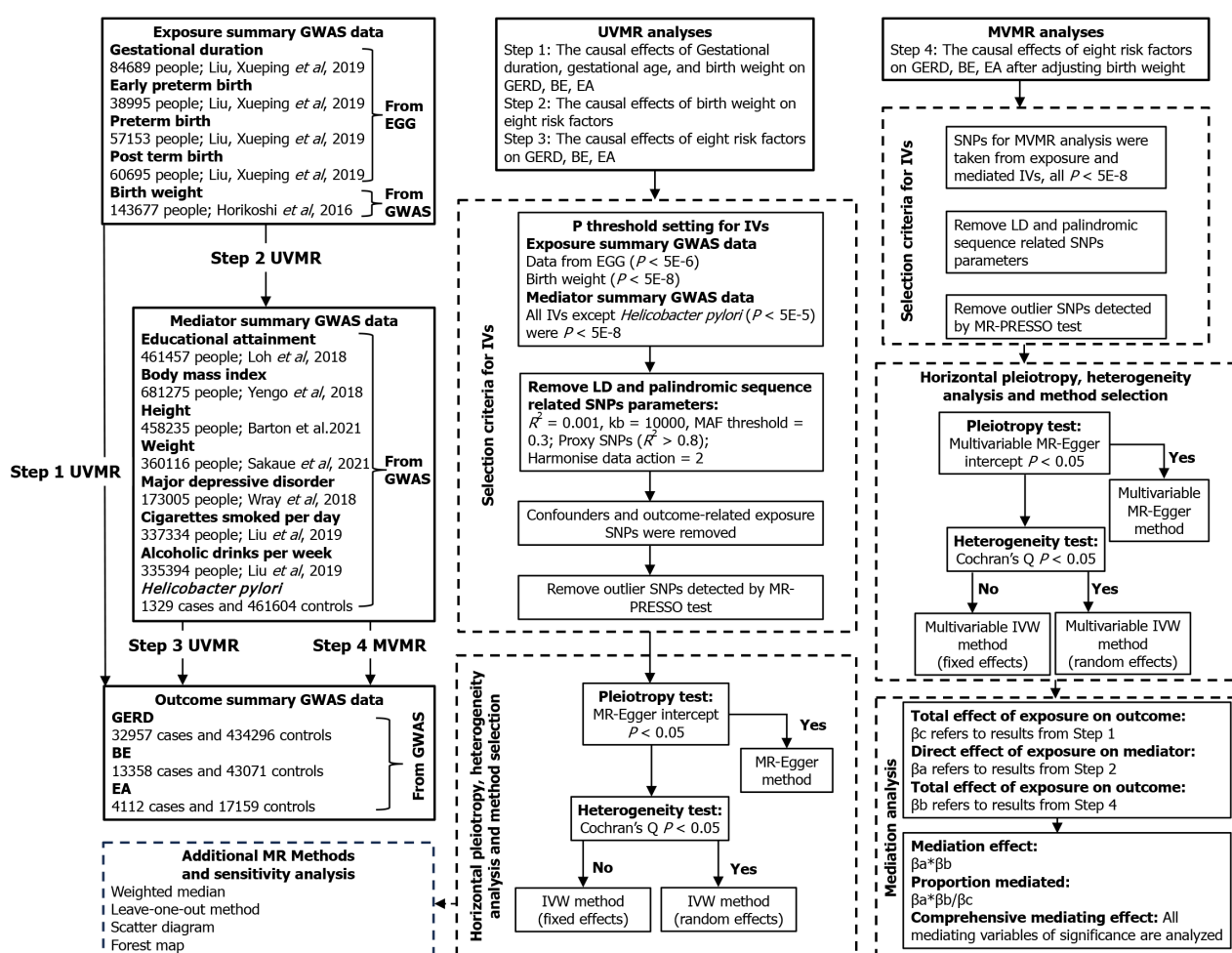


Figure 1 Flowchart of Mendelian randomization analyses in this study. Step 1: The causal effects of gestational duration, gestational age, and birth weight on gastroesophageal reflux disease (GERD), Barrett's esophagus (BE), esophageal adenocarcinoma (EA); Step 2: The causal effects of birth weight on eight risk factors; Step 3: The causal effects of eight risk factors on GERD, BE, EA; Step 4: The causal effects of eight risk factors on GERD, BE, EA after adjusting birth weight. BE: Barrett's esophagus; EA: Esophageal adenocarcinoma; EGG: Early Growth Genetics; GWAS: Genome-wide association study; IVs: Instrumental variables; IVW: Inverse variance weighted; MR-PRESSO: MR-Pleiotropy RESidual sum and outlier; UVMR: Univariable Mendelian randomization; SNPs: Single nucleotide polymorphisms; GERD: Gastroesophageal reflux disease; MVMR: Multivariable Mendelian randomization; LD: Linkage disequilibrium.

Sensitivity analysis: Sensitivity analysis was employed to assess the robustness of MR results. In addition to IVW method (both fixed-effects and random-effects models) and the MR-Egger method, the weighted median method was also utilized as a reference[41]. Ensuring consistency in the direction of effects among different MR methods, with statistical significance met, enhances the credibility of the results. Supplementary scatter plots, forest plots, and Leave-One-Out (LOO) analysis were incorporated to ensure the validity of causal inferences in this study[46]. In the LOO analysis results, SNPs that were statistically significant and led to a reversal in the direction of effects were excluded. Statistical analyses were carried out using R version 4.2.1, and the R packages TwoSampleMR (version 0.5.7), Mendelian-Randomization (version 0.3.0), and forestploter (version 1.1.1) were employed for these analyses.

RESULTS

Causal links between four gestational traits and birth weight on GERD, BE, and EA

After rigorously excluding outlier SNPs, the final IVs associated with four gestational phenotypes (gestational duration, early preterm birth, preterm birth, postterm birth) was 2.38%, with F -statistics ranging from 26.99 to 199.63. For the IVs related to birth weight, the explained variance was 2.23%, with F -statistics between 607.81 and 740.38 (Supplementary Tables 1 and 2). Figure 2A visually presents the results of UVMR analysis. An increase in birth weight demonstrated a significant effect in reducing the risk of GERD and BE, while showing no association with EA risk. GERD (OR = 0.78, 95%CI: 0.69-0.8) and BE (OR = 0.75, 95%CI: 0.60-0.9). However, no significant associations were detected for gestational duration, early preterm birth, preterm birth, or postterm birth in the outcomes of the three esophageal disorders. Additional results from other MR methods are reported in Supplementary Table 3. No horizontal pleiotropy was observed in all analyses ($P_{\text{MR-Egger Intercept}} > 0.05$), and random-effects models were applied to correct for heterogeneity.

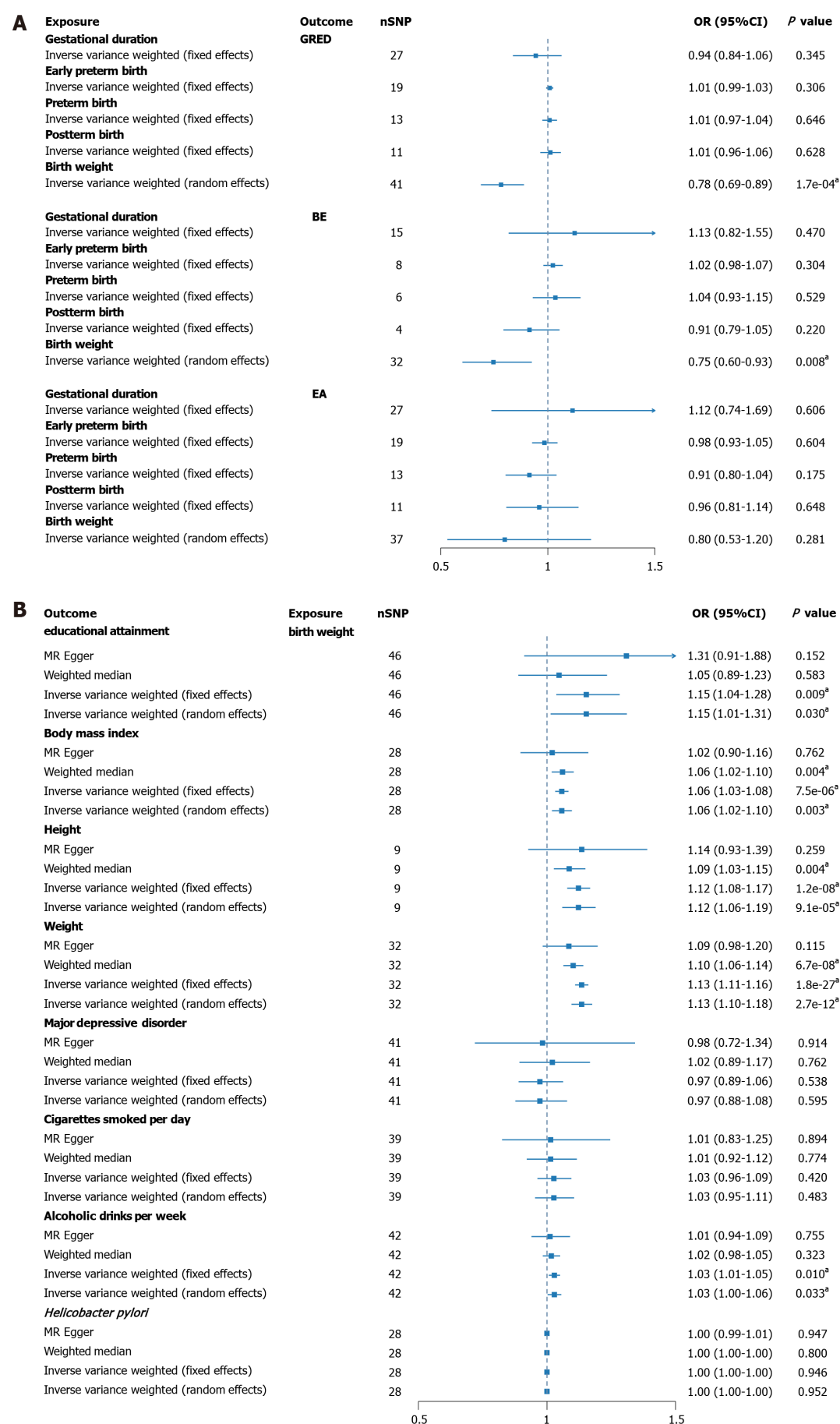


Figure 2 Univariable Mendelian randomization analysis. A: Univariable Mendelian randomization analysis of gestational age-related genetic characteristics, birth weight and causality of esophageal disease; B: Univariable Mendelian randomization analysis of causal effects of genetic birth weight on risk

factors. ^a $P < 0.05$. BE: Barrett's esophagus; CI: Confidence interval; EA: Esophageal adenocarcinoma; GERD: Gastroesophageal reflux disease; nSNPs: Number of single nucleotide polymorphisms; OR: Odds ratio.

Sensitivity analysis indicated robust results, with no significant outliers identified in the LOO analysis (Supplementary Figures 1-15).

Causal links between birth weight on multiple risk factors

Due to statistical significance observed only between birth weight and the outcome in UVMR, subsequent analyses were focused solely on birth weight. The explanatory power of birth weight among various risk factors was 3.64%, with F -statistics ranging from 160.61 to 833.99 (Supplementary Tables 4 and 5). Figure 2B illustrates a significant positive correlation between birth weight and five risk factors, while showing no association with others. Educational attainment (OR = 1.15, 95%CI: 1.01-1.31), BMI (OR = 1.06, 95%CI: 1.02-1.1), height (OR = 1.12, 95%CI: 1.06-1.19), weight (OR = 1.13, 95%CI: 1.10-1.1), and alcoholic drinks per week (OR = 1.03, 95%CI: 1.00-1.06) all exhibited significant associations (Supplementary Table 6). No evidence of pleiotropy was detected, and all MR methods showed consistent effect directions. Sensitivity analysis also revealed no significant outliers, ensuring the reliability of the results (Supplementary Figures 16-23).

Causal links of risk factors on GERD, BE, and EA considering adjusted and unadjusted birth weight

Through UVMR, the causal effects of risk factors on GERD, BE, and EA were estimated without adjusting for birth weight (Supplementary Tables 7-9), with corresponding sensitivity analyses presented in Supplementary Figures 24-47. The application of MVMR yielded the direct effects of various risk factors on GERD, BE, and EA after adjusting for birth weight (Figure 3, Supplementary Table 10). Only weight and partially adjusted BMI showed evidence of horizontal pleiotropy, and MR-Egger was employed to report the results. Educational attainment significantly reduced the risk of GERD, BE, both before adjustment (OR = 0.89, 95%CI: 0.87-0.91; OR = 0.76, 95%CI: 0.73-0.81), and after adjustment (OR = 0.89, 95%CI: 0.87-0.91; OR = 0.75, 95%CI: 0.71-0.78), and EA (OR = 0.90, 95%CI: 0.84-0.96). BMI increased the susceptibility to all three esophageal diseases, both before adjustment (OR = 1.29, 95%CI: 1.21-1.39; OR = 2.64, 95%CI: 2.32-3.00; OR = 1.50, 95%CI: 1.22-1.86), and after adjustment (OR = 1.28, 95%CI: 1.20-1.36; OR = 2.78, 95%CI: 2.46-3.15; OR = 1.60, 95%CI: 1.32-1.94). Height demonstrated a protective effect in BE (OR = 0.95, 95%CI: 0.92-0.99) and EA (OR = 0.86, 95%CI: 0.79-0.93), with a significant protective effect observed only in BE after adjustment (OR = 0.87, 95%CI: 0.79-0.96). Conversely, weight significantly reduced the likelihood of GERD (OR = 0.73, 95%CI: 0.55-0.97) and BE before adjustment (OR = 0.27, 95%CI: 0.14-0.54). However, after adjustment, weight exhibited an opposite effect, with significant differences observed for GERD (OR = 1.21, 95%CI: 1.11-1.31), BE (OR = 1.86, 95%CI: 1.55-2.24), and EA (OR = 1.63, 95%CI: 1.28-2.06). A similar contrast phenomenon was observed in MDD, where MDD, before adjustment, acted as a factor increasing the risk of GERD (OR = 1.37, 95%CI: 0.94-1.98) and BE (OR = 2.70, 95%CI: 1.91-3.83). However, after adjustment, MDD exhibited a protective effect (OR = 0.79, 95%CI: 0.72-0.88; OR = 0.76, 95%CI: 0.61-0.95). Daily smoking quantity, before adjustment, did not show significance in any outcome, but after adjustment, smoking increased the risk of BE (OR = 1.23, 95%CI: 1.04-1.46). Before adjustment, a substantial increase in weekly alcohol consumption raised the risk of EA (OR = 3.88, 95%CI: 1.53-9.89), but in the adjusted results, no significant effect was observed. *H. pylori* infection was not reported to be significant before adjustment, and in the analysis adjusting for birth weight, due to sample size limitations, independent SNPs were not obtained for MVMR analysis at the threshold of $P < 5E-5$.

Mediation effect analysis

For factors demonstrating consistent and statistically significant total and mediation effects, we conducted confidence interval calculations and estimated mediation proportions (Supplementary Table 11). Despite the consistent negative association of educational attainment with GERD observed in both UVMR and MVMR, the mediation effect did not reach statistical significance ($\beta = -0.017$, 95%CI: -0.052 to 0.019). Conversely, education level (proportion mediated = 13.99%, $\beta = -0.041$, 95%CI: -0.080, to -0.003) and height (proportion mediated = 5.46%, $\beta = -0.016$; 95%CI: -0.029 to -0.002) mediated the impact of birth weight on BE, both exhibiting negative correlations. Subsequently, Figure 4 and Supplementary Tables 12 and 13 present the comprehensive mediation effects of education level and height on BE. For educational attainment (proportion mediated = 15.36%, $\beta = -0.045$; 95%CI: -0.086 to -0.003) and height (proportion mediated = 4.78%, $\beta = -0.014$, 95%CI: -0.026 to -0.002), the results indicate significant and consistent mediation effects. No evidence of horizontal pleiotropy was found, with both multivariable IVW and MVMR-Egger results demonstrating significance and consistent direction, supporting the robustness of our findings.

DISCUSSION

In this study, we employed a large-scale genetic dataset and conducted UVMR, MVMR, and mediation analyses to comprehensively assess the causal relationships between fetal genetic characteristics during pregnancy, birth weight, and the onset risk of GERD, BE, and EA in adulthood. Additionally, we analyzed potential mediation effects of known risk factors. Our research confirms that gestational age does not impact the risk of GERD, BE, and EA in adulthood. This conclusion remains consistent whether analyzed as a continuous variable based on gestational weeks or as a binary variable distinguishing early preterm, preterm, and post-term births. Lower birth weight is causally associated with an

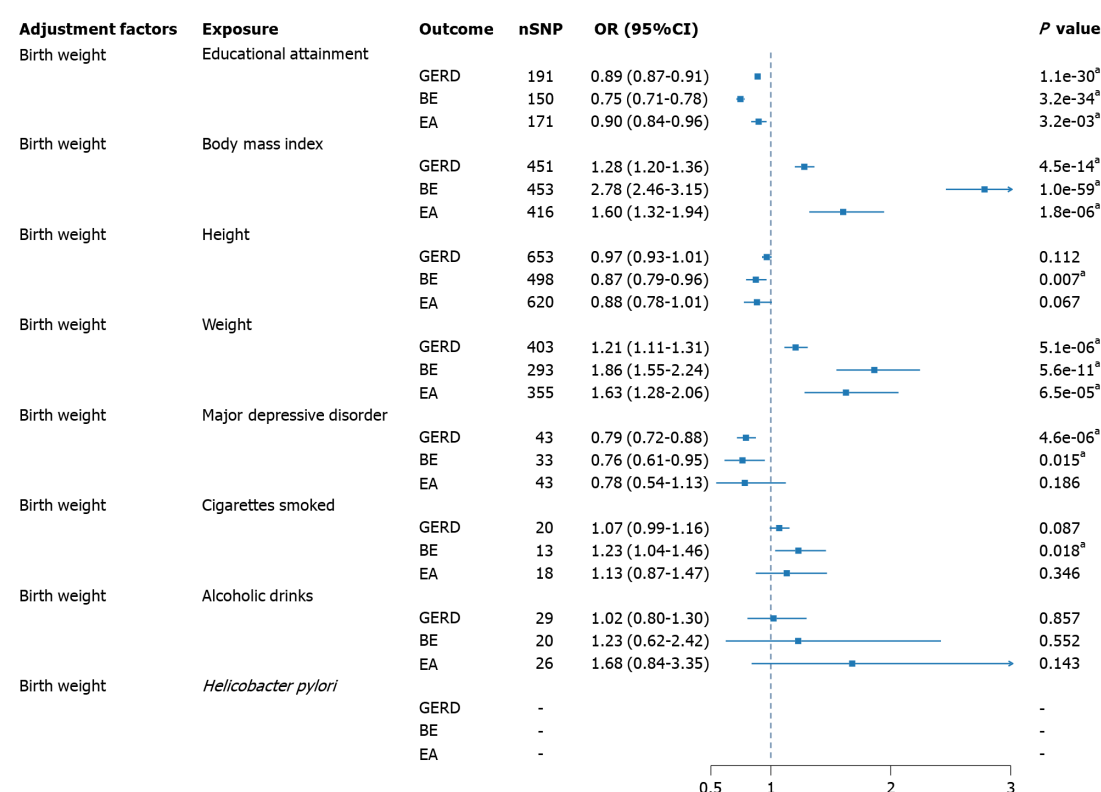


Figure 3 Multivariable Mendelian randomization analysis of causal effects of risk factors on esophageal disease after adjusting for birth weight. ^a $P < 0.05$. *Helicobacter pylori* phenotypes do not acquire sufficient single nucleotide polymorphisms in multivariable Mendelian randomization. GERD: Gastroesophageal reflux disease; BE: Barrett's esophagus; EA: Esophageal adenocarcinoma; OR: Odds ratio; CI: Confidence interval; nSNPs: Number of single nucleotide polymorphisms.

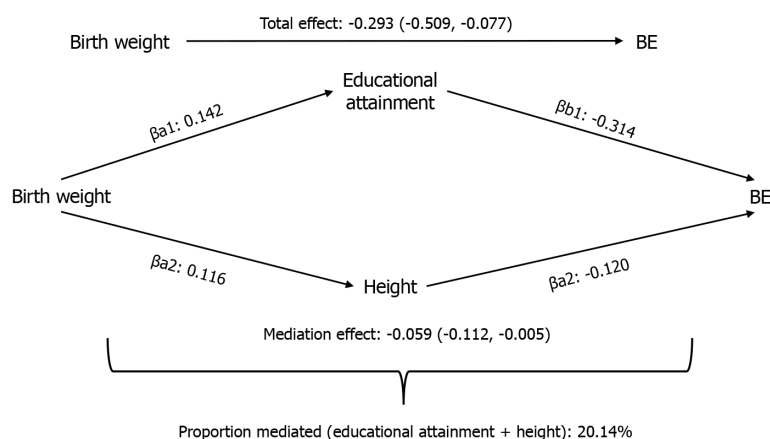


Figure 4 Combined effects of multiple mediators on esophageal diseases. The combined proportions mediated by educational attainment and height for the effect of birth weight on Barrett's esophagus. β_a : The effect of birth weight on the risk of modifiable risk factors; β_b : The effect of risk factors on the risk of Barrett's esophagus after combination of mediating factors; Total effect: The effect of birth weight on the risk of Barrett's esophagus; BE: Barrett's esophagus.

increased risk of GERD and BE in adults. Conversely, higher birth weight shows causal relationships with higher education levels, obesity (high BMI), increased height and weight, and a higher weekly alcohol intake. Furthermore, within these factors, education level and height mediate 15.36% and 4.78%, respectively, of the total effect of birth weight on BE. All analysis results withstand sensitivity and multicollinearity tests, ensuring the reliability of our findings.

Considering the strong concealment of EA, it remains a challenging subtype in the early diagnosis and prevention of esophageal cancer worldwide, with an increasing incidence in many Western countries[4,47]. Most EA patients face a poor prognosis, with the overall 5-year survival rate for esophageal cancer still below 20%[2]. Meanwhile, BE, as the only precancerous lesion of EA, has an annual incidence rate of approximately 0.2%-0.5%[5]. In developed countries, adult GERD accounts for about 20%[6], showing an association with increased risks of BE and EA. Therefore, understanding the high-risk populations and risk factors for the onset of GERD, BE, and EA is crucial for early screening and prevention decisions regarding esophageal cancer.

In recent years, limited research has focused on the potential connections between neonatal characteristics and the risk of GERD, BE, and EA in adulthood. However, these studies have yielded inconsistent viewpoints. Although GERD is common in preterm or SGA infants[13], there is no reported association between GERD risk in adults and a history of preterm birth in cohort studies. Shiota *et al*[16] proposed that preterm birth increases the risk of adult BE, with this risk decreasing in infants larger than the gestational age, and this phenomenon was not mediated by known high-risk factors. In contrast, Forssell *et al*[17] suggested a different perspective, indicating that the increased risk of BE in adulthood is due to the impact of SGA and is unrelated to preterm birth. This aligns with our study's findings, where individuals with higher birth weight have a 25% lower risk of developing BE, with no evidence supporting the influence of preterm birth on BE. In a long-term follow-up involving 3364 preterm/SGA infants, eight individuals developed esophageal cancer in adulthood compared to only one in the control group. This resulted in an 11.5-fold increase in EA risk in SGA, but concerns were raised about potential false associations due to the small sample size[14]. Some argue that the increased risk of adult EA is unrelated to birth weight but associated with gestational weeks[15], while prospective studies, including ours, find no correlation between gestational weeks, birth weight, and EA risk[48]. These phenomena may be due to the adaptive response of SGA in the early stage of the disease, which leads to the lagging growth and development of tissue and organ function through metabolic or hormonal changes, resulting in an increased risk of disease. It is also possible that perinatal promoter methylation affects gene expression in pathways associated with a range of physiological processes and continues into adulthood[49].

These long-term cohort studies lack sufficient exploration of other risk factors' potential effects on GERD, BE, and EA, and are burdened by significant limitations. It remains unclear if preterm/SGA individuals exhibit growth tendencies or engage in high-risk behaviors like smoking, drinking, or obesity[17,50]. Additionally, the increased risk of adult depression linked to low birth weight complicates inferences[51], as depression or anxiety further heightens gastrointestinal disease risk[15,52]. Scarce research on preterm/SGA individuals regarding esophageal diseases and a lack of reports on intermediate factors prompted our study. Leveraging large-scale genetic data from the European population (84689 for gestational age, 143677 for birth weight)[25,27] and utilizing MR analysis, we overcame confounding factors, facilitating precise causal inferences[18].

Past studies on risk factors for esophageal diseases did not consider the potential impact of neonatal characteristics. This research offers new insights into the role of adjusted risk factors after birth weight modification in esophageal diseases. Firstly, the increase in education level, as found by Jansson *et al*[9] and Zhang *et al*[11], is associated with a decreased risk of GERD, BE, and esophageal cancer. Our results align with this, showing that education level remains a protective factor after adjusting for birth weight. Obesity (high BMI/overweight) has been widely recognized for its detrimental effects on gastrointestinal diseases. In our study, adjusting for birth weight increased the risks of GERD, BE, and EA by 28%, more than double, and 60%, respectively. The association between these factors may be attributed to visceral obesity, resulting in increased abdominal fat thickness and heightened intra-abdominal pressure. Mechanical damage to the esophagus is more common in obese individuals, increasing the chances of acid exposure and thereby elevating the risk of GERD. Additionally, elevated levels of serum adipocyte-derived cytokines, intensified by inflammatory responses mediated through adiponectin or leptin, contribute to an increased risk of BE[53]. Thrift *et al*[54] found that increased height contributes to a lower probability of BE and EA in males, with no significant effect in females. In our study, adjusted height was only significantly associated with BE, possibly due to the genetic information related to height explaining the overall effects in both male and female populations, minimizing the impact of gender subgroups. Regarding the impact of MDD on esophageal diseases, some argue that MDD is unrelated to BE and EA but is associated with an increased risk of GERD[35]. In our study, after adjusting for birth weight, MDD had statistical significance for both GERD and BE, but caution is needed in interpreting this due to inconsistent directions in effects before and after adjustment and the limited number of SNPs obtained. Smoking is associated with a 23% increased risk of BE in this study, aligning with findings by Yuan and Larsson[55]. Smoking has been shown to induce damage to esophageal epithelial cells, exacerbating metaplasia. Infection with *H. pylori* (adjusted before) and alcohol consumption (adjusted after) is unrelated to the occurrence of GERD, BE, and EA, aligning with some research conclusions[56,57] but differing from others[1,6,58]. Perhaps this is due to the impact of low socioeconomic status and insufficient education levels in SGA individuals during adulthood, leading to increased risks of smoking, drinking, *H. pylori* infection, *etc*[59]. Neglecting birth weight in observational studies masks the true effects of relevant factors, potentially resulting in incorrect inferences.

This study provides support for education level being an intermediate factor in the increased risk of BE associated with high birth weight. In several large birth cohort studies, early improvements in birth weight have been shown to effectively increase education levels, cognitive abilities, and social status, subsequently reducing the risk of chronic diseases in adulthood[60,61]. Higher education levels can further reduce the harms brought by adverse habits[24,62]. The protective intermediate effect of height may be due to growth restriction in the womb for SGA individuals, resulting in a relatively small stature during growth, increasing the risk of gastrointestinal diseases such as GERD due to increased intra-abdominal pressure[54,63]. This process may eventually evolve into BE through repeated stimulation.

The findings of this study address previous research limitations by proposing the impact of birth weight-adjusted relevant risk factors on esophageal diseases and highlighting the intermediate roles of education level and height. New perspectives on the effects of pathogenic factors like smoking and drinking have been introduced. This study is expected to enhance our understanding of the complex mechanisms of genetic factors in esophageal diseases. Early intervention measures and personalized treatment should be developed for SGA individuals, and early screening and prevention of esophageal diseases should be conducted in adulthood for these high-risk populations. However, the mediating mechanism of height between SGA children and precancerous lesions of esophageal cancer remains unclear, requiring further research to explain this effect.

Limitations

This study focused solely on the European population, and the generalization of findings to other ethnic groups remains inconclusive. In the MVMR analysis after adjusting for birth weight, *H. pylori* infection lacked a sufficient number of SNPs for subsequent analysis. Despite incorporating multiple known risk factors into our analysis, we cannot eliminate the potential effects of unknown factors.

CONCLUSION

Our research supports the association between low birth weight and an increased risk of GERD and BE, but not EA. Gestational weeks are unrelated to all three esophageal diseases. The causal effect of birth weight on BE is jointly mediated by education level and height, accounting for approximately 20%. Furthermore, our study provides a comprehensive effect of known risk factors on esophageal diseases after adjusting for birth weight.

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FOOTNOTES

Author contributions: Ruan LC and Zhang Y contributed equally to this work. Ruan LC conceived the project and wrote the paper; Zhang Y and Su L collaborated in designing the analysis plans; Zhu LX and Wan BG collected relevant gene phenotype data; Wang SL, Quo Q, Qiu SY and Hu S provided technical support throughout the research process; Wei YP and Zheng QL contributed equally to this study; Zheng QL and Wei YP both rigorously reviewed and proofread the content of the article, and put forward reasonable improvement plans; and all authors have read and approved the final manuscript.

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