

BMJ Open is committed to open peer review. As part of this commitment we make the peer review history of every article we publish publicly available.

When an article is published we post the peer reviewers' comments and the authors' responses online. We also post the versions of the paper that were used during peer review. These are the versions that the peer review comments apply to.

The versions of the paper that follow are the versions that were submitted during the peer review process. They are not the versions of record or the final published versions. They should not be cited or distributed as the published version of this manuscript.

BMJ Open is an open access journal and the full, final, typeset and author-corrected version of record of the manuscript is available on our site with no access controls, subscription charges or pay-per-view fees (http://bmjopen.bmj.com).

If you have any questions on BMJ Open's open peer review process please email info.bmjopen@bmj.com

BMJ Open

Postpartum Hemorrhage Following Vaginal Delivery: A Comprehensive Analysis and Development of Predictive Models for Etiological Subgroups

Journal:	BMJ Open
Manuscript ID	bmjopen-2024-089734
Article Type:	Original research
Date Submitted by the Author:	16-Jun-2024
Complete List of Authors:	Li, Jinke; Shengjing Hospital of China Medical University, Department of Obstetrics and Gynecology Zhang, Dandan; Shengjing Hospital of China Medical University, Department of Obstetrics and Gynecology Lin, Hong; Liaoning Maternal and Child Health Hospital, Department of Obstetrics and Gynecology Shao, Mengyuan; Shenyang Women's and Children's Hospital, Department of Obstetrics and Gynecology Wang, Xiaoxue; Shengjing Hospital of China Medical University, Department of Health Management Chen, Xueting; Shengjing Hospital of China Medical University, Department of Health Management Zhou, Yangzi; Shengjing Hospital of China Medical University, Department of Obstetrics and Gynecology Song, Zixuan; Shengjing Hospital of China Medical University, Department of Obstetrics and Gynecology
Keywords:	Maternal medicine < OBSTETRICS, PERINATOLOGY, Postpartum Women < Postpartum Period, PREVENTIVE MEDICINE

SCHOLARONE™ Manuscripts



I, the Submitting Author has the right to grant and does grant on behalf of all authors of the Work (as defined in the below author licence), an exclusive licence and/or a non-exclusive licence for contributions from authors who are: i) UK Crown employees; ii) where BMJ has agreed a CC-BY licence shall apply, and/or iii) in accordance with the terms applicable for US Federal Government officers or employees acting as part of their official duties; on a worldwide, perpetual, irrevocable, royalty-free basis to BMJ Publishing Group Ltd ("BMJ") its licensees and where the relevant Journal is co-owned by BMJ to the co-owners of the Journal, to publish the Work in this journal and any other BMJ products and to exploit all rights, as set out in our licence.

The Submitting Author accepts and understands that any supply made under these terms is made by BMJ to the Submitting Author unless you are acting as an employee on behalf of your employer or a postgraduate student of an affiliated institution which is paying any applicable article publishing charge ("APC") for Open Access articles. Where the Submitting Author wishes to make the Work available on an Open Access basis (and intends to pay the relevant APC), the terms of reuse of such Open Access shall be governed by a Creative Commons licence – details of these licences and which Creative Commons licence will apply to this Work are set out in our licence referred to above.

Other than as permitted in any relevant BMJ Author's Self Archiving Policies, I confirm this Work has not been accepted for publication elsewhere, is not being considered for publication elsewhere and does not duplicate material already published. I confirm all authors consent to publication of this Work and authorise the granting of this licence.

Postpartum Hemorrhage Following Vaginal Delivery: A Comprehensive Analysis and Development of Predictive Models for Etiological Subgroups

Jinke Li^{1#}, Dandan Zhang^{1#}, Hong Lin², Mengyuan Shao³, Xiaoxue Wang⁴, Xueting Chen⁴, Yangzi Zhou¹ and Zixuan Song^{1*}

- 1 Department of Obstetrics and Gynecology, Shengjing Hospital of China Medical University, Shenyang, China
- 2 Department of Obstetrics and Gynecology, Liaoning Maternal and Child Health Hospital, Shenyang, China
- 3 Department of Obstetrics and Gynecology, Shenyang Women's and Children's Hospital, Shenyang, China
- 4 Department of Health Management, Shengjing Hospital of China Medical University, Shenyang, China
- #These authors contributed equally to this work.

Correspondence:songzixuan666@163.com

Abstract:

Objective: This study aimed to dissect the etiological subgroups of postpartum hemorrhage (PPH) that occur after vaginal delivery. Our goal was to craft and validate predictive models to guide clinical decision-making and optimize resource allocation..

Methods: Conducted across three hospitals from 2016 to 2022, the study enrolled 29,842 women who underwent vaginal delivery. PPH was categorized into uterine atony (UA), placental factors (PF), cervical trauma (CT), and coagulation abnormalities (CA) by etiology. Logistic regression identified risk factors for each PPH subgroup and constructed predictive models. The cohort was segmented into a training dataset (70%) and an internal validation dataset (30%), complemented by an additional cohort for external validation. Model performance was rigorously assessed using R software, with a focus on discrimination, calibration, and clinical utility as benchmarked by Decision Curve Analysis (DCA).

Results: The logistic regression for overall PPH and UA-PPH showcased high discrimination (AUCs of 0.807 and 0.794, respectively), coupled with commendable calibration and DCA-assessed clinical utility, culminating in the development of a nomogram for risk prediction. The PF-PPH model exhibited a modest AUC of 0.739, while the CT-PPH and CA-PPH models demonstrated suboptimal clinical utility and calibration.

Conclusion: The study identified factors associated with PPH and developed models with good performance for overall PPH and UA-PPH. The nomogram offers a valuable tool for risk prediction. However, models for PF-PPH, CT-PPH, and CA-PPH require further refinement. Future research should focus on larger samples and multi-center validation for enhanced model generalizability.

Keywords: Postpartum Hemorrhage, Vaginal Delivery, Etiological Subgroups, Predictive Models, nomogram

Background

Postpartum hemorrhage (PPH) is a widespread and serious medical condition that poses significant risks to women's health around the world. It is particularly devastating in developing

countries, where it is a principal contributor to maternal mortality. ^[1] It is estimated that approximately 1.4 million maternal deaths globally are tied to PPH each year, with the tragic loss of a woman's life to this condition occurring every four minutes. ^[2, 3]. In Australia, the incidence of PPH increased from 6.3% in 2000 to 8.0% in 2009. ^[4] Similarly, in the United States, the rate of PPH rose from 2.7% in 1999 to 3.2% in 2014. ^[5] In China, despite a relatively lower maternal mortality rate of 17.8 per 100,000 in 2019, PPH accounted for one-quarter of these deaths. ^[6]

The World Health Organization (WHO) has conducted an analysis revealing that while PPH is a significant factor in maternal mortality and morbidity, the mortality rates vary considerably across different regions.^[7] In high-income countries, the risk of death due to PPH is significantly lower than in low-income countries.^[8] In high-income nations, the substantial blood loss primarily caused by PPH accounts for 13.4% of overall maternal mortality, while in Africa and Asia, this figure stands at 34% and 30.8%, respectively. ^[7] The international obstetric community is actively engaged in research to better understand the incidence, risk factors, and management strategies for PPH.^[9-11]. Despite the establishment of global clinical guidelines and the identification of various risk factors, further exploration is needed to enhance our understanding and management of PPH.^[12, 13].

PPH can be etiologically classified into uterine atony (UA), placental factors (PF), cervical trauma (CT), and coagulation abnormalities (CA), each requiring distinct clinical management and treatment strategies. ^[14] Clear etiological classification is crucial for developing preventive strategies, formulating management plans, and rational allocation of medical resources. ^[15] While numerous cohort studies have focused on identifying risk factors for PPH, there is a scarcity of studies that quantify and weigh these risk factors for a comprehensive PPH risk assessment. ^[16-18] Given the complexity of PPH and the interplay of multiple risk factors, a holistic approach is necessary to accurately assess the risk of PPH.

Clinical prediction models (CPMs) have been widely applied in clinical settings in recent years. By constructing CPMs, physicians and patients can make better medical decisions, and health departments at all levels can allocate medical resources more rationally. These models play an irreplaceable role in primary prevention (assessing the quantitative risk of future diseases) and secondary prevention (constructing highly sensitive and specific diagnostic schemes, practicing "early detection, early diagnosis, early treatment"), reflecting significant health economic value.

There is a gap in research regarding the development of clinical prediction models for women specifically following vaginal delivery. Many studies are constrained by limited sample sizes, which can affect the robustness of the models^[19]. Other research has focused on PPH prediction models for women undergoing cesarean sections.^[20] Our study aims to address this gap by constructing a clinical prediction model tailored to PPH after vaginal delivery. By analyzing clinical data and risk factors through logistic regression, we can determine the relative impact of each factor on the likelihood of PPH. We further refine our model by performing secondary fitting based on the four etiological subgroups, creating a nomogram that enhances the precision of predicting high-risk populations for PPH. This work provides essential insights for the prevention and management of this critical condition.

Materials and methods

Data Sources and Ethics Statement

This cohort study was conducted at the obstetric wards of Shengjing Hospital of China Medical University, Liaoning Maternal and Child Health Hospital, and Shenyang Women's and Children's Hospital. The study population comprised women who underwent vaginal delivery between January 1, 2016, and December 31, 2022. The outcomes of interest were fetal birth outcomes within the first 24 hours postpartum. Inclusion criteria were women who consented to participate after being informed of the study's scope. Exclusion criteria were defined as follows: age under 18 or over 50 years, delivery occurring at less than 37 weeks or more than 42 weeks of

gestation, multiple births, and instances of induced labor, stillbirth, or fetal death. Comprehensive data encompassing maternal characteristics, obstetric and gynecologic history, pregnancy complications, and details of the delivery process and neonatal conditions were collected (Supplement 1). To protect participant privacy, all data were anonymized. The study protocol was approved by the Ethical Review Committee of Shengjing Hospital of China Medical University (No. 2016PS344K), and written information about the study was provided to all participants.

Sample Size Calculation

The study was designed as an observational analysis to estimate the incidence of PPH at less than 10% and statistical parameters set at a significance level (α) of 0.05 and a power (1- β) of 0.95, the required sample size was determined to be 384 patients to achieve an absolute precision of 5%.

Covariates

A range of covariates were taken into account, including:(1) Age, categorized as <25, 25-29, 30-34, and ≥35 years; (2) Ethnicity, divided into Han, Manchu, and other; (3) Education level, classified as high school or below, bachelor's degree, and postgraduate or above; (4) Occupation, categorized as unemployed, light physical labor, moderate physical labor, and heavy physical labor (based on the International Physical Activity Questionnaire, IPAQ); (5) Monthly household income per capita, divided into <0.5, 0.5-2.0, 2.0-5.0, and >5.0 thousand yuan; (6) Pre-pregnancy BMI, categorized as underweight (<18.5 kg/m²), normal (18.5-23.9 kg/m²), overweight (24-27.9 kg/m²), and obese (≥28 kg/m²); (7) Gravidity, categorized as 1, 2, or ≥ 3 times; (8) Parity, divided into 0, 1, or ≥ 2 times; (9) Gestational age, categorized as <38, 38-40, and >40 weeks; (10) Delivery time, divided into daytime (8-16), evening (17-23), and night (0-7) shifts; (11) Total duration of labor, categorized as normal (≤24 hours) and prolonged (>24 hours); (12) Latent phase of the first stage, categorized as normal (primiparous ≤20 hours, multiparous ≤14 hours) and prolonged (primiparous >20 hours, multiparous >14 hours); (13) Active phase of the first stage, categorized as normal (\le 8 hours) and prolonged (>8 hours); (14) Second stage duration, categorized as normal and prolonged based on specific criteria for primiparous and multiparous women with or without analgesia; (15) Third stage duration, categorized as normal (≤ 30 minutes) and prolonged (≥ 30 minutes).

Etiology Subgroup

Patients were categorized into groups with or without postpartum hemorrhage based on the presence of postpartum bleeding and its underlying etiology, which included UA, PF, CT, and CA.

Model Construction

For the purpose of our investigation, we have categorized the participants from the Shengjing hospital of China Medical University as Cohort I. This cohort was systematically split into a training dataset and an internal validation dataset with a ratio of 7:3. The training dataset was instrumental in developing the predictive model, while the internal validation dataset served to assess the model's predictive accuracy. An additional cohort, comprising participants from two other hospitals, was designated as Cohort II. This external dataset was used to validate the model's general applicability and its efficacy in real-world clinical scenarios.

Within the confines of the datasets, we employed both univariate and multivariate logistic regression analyses to identify potential risk factors across various subgroups. These factors were then subjected to a rigorous selection process for inclusion in the predictive model. The selected factors were further analyzed using multivariate logistic regression in training dataset to discern their discriminative power, thereby establishing them as predictive indicators for the model. The model's features were meticulously optimized through logistic regression, and the performance of these models was corroborated using both the test and validation datasets to ascertain the most accurate predictive model.

Evaluating the Performance of the Models

The area under the receiver operating characteristic curve (AUC) was the primary metric used to evaluate the discrimination of our models. An AUC value above 0.75 suggests excellent model discrimination, while an AUC below 0.6 indicates poor discrimination. Calibration curves were used to assess the models' accuracy, with closer alignment between observed and predicted incidence rates indicating higher model fidelity. Decision Curve Analysis (DCA) was also employed to evaluate the clinical utility of the models, offering a thorough assessment of the models' net benefits across various clinical scenarios.

Nomogram Development

Nomograms for postpartum hemorrhage and its four etiological subgroups were crafted to offer a visual representation of the risk scores derived from the logistic regression analysis. This tool simplifies the interpretation of complex statistical outcomes, providing a more straightforward approach to understanding risk assessments.

Statistical Analysis

All statistical computations, construction of traditional logistic models, and calculations of model discrimination and calibration were carried out using R version 3.6.3 from the R Foundation for Statistical Computing, Vienna, Austria. This software facilitated the development of traditional logistic predictive models and their subsequent evaluation for discriminative power, calibration, and clinical utility. Continuous variables conforming to the normal distribution were expressed as the mean \pm standard deviation (SD), while non-normally distributed continuous variables were presented as medians with interquartile ranges. Categorical data were analyzed using chi-square tests, and continuous variables were analyzed using ANOVA or Mann-Whitney tests, as appropriate. Variables were adjusted as dummy variables, and odds ratios (OR) with corresponding 95% confidence intervals (95% CI) were calculated using univariate and multivariate logistic regression analyses, with significance levels set at P < 0.05 or P < 0.001.

Results

From 2016 to 2022, a total of 27,389 patients underwent vaginal delivery at the Shengjing Hospital of China Medical University. Forty-two patients under 18 years of age or over 50 years old were excluded. Additionally, 2,456 patients with gestational age less than 37 weeks or more than 42 weeks at delivery, 6 patients with multiple births, and 52 patients with induced labor, stillbirth, or fetal death were also excluded. Ultimately, 24,833 patients met the inclusion criteria and were enrolled in the cohort. According to the inclusion and exclusion criteria, a total of 5,099 patients in cohort II were included in the external validation dataset. The general characteristics of all patients are presented in Table 1. All patients were followed up within 24 hours after delivery for neonatal outcomes, with a follow-up rate of 100%. The patient selection criteria flowchart is shown in Figure 1.

Comparison of Basic Characteristics and Risk Analysis for Postpartum Hemorrhage (PPH) and Its Subgroups

Based on the occurrence of postpartum hemorrhage, the parturients in cohort one were divided into two groups: the non-PPH group and the PPH group. Similarly, within the etiological subgroups, they were categorized into UA-PPH and non-UA-PPH groups, PF-PPH and non-PF-PPH groups, CT-PPH and non-CT-PPH groups, and CA-PPH and non-CA-PPH groups. The comparison of basic characteristics and analysis of risk factors for each group are presented in Supplementary Tables 1-5.

In the multivariate analysis of risk factors, apart from age, parity, pre-pregnancy BMI, anemia, premature rupture of membranes, and combined placenta retention/placenta accreta/placental implantation, other specific risk factors were found to be associated with specific etiologies of postpartum hemorrhage. For instance, polyhydramnios was associated with UA-PPH; analgesia during labor, instrumental assistance, and cervical/vaginal/perineal lacerations were associated with the occurrence of CT-PPH (Table 2).

Selection of Predictive Factors for PPH and Its Subgroups in the Training Dataset

Through random sampling of cohort one, 70% of the data (N=17,383) from parturients were used to form the training dataset, with the remaining approximately 30% (N=7,450) forming the internal validation dataset. Multivariate analysis of risk factors for PPH and its subgroups was performed again in the training dataset, with results presented in Supplemental Table 1. After selection, predictive models were constructed for each group using the selected risk factors.

Evaluation of Predictive Model Discrimination

The ROC curves were plotted using R software for the PPH group and its various subgroups across the training dataset, internal and external validation dataset.

The results indicated that the predictive models, namely PH-Logistic, UA-PPH-Logistic, PF-PPH-Logistic, CT-PPH-Logistic, and CA-PPH-Logistic, demonstrated high discriminative power in the training dataset with AUCs of 0.807 (95% CI: 0.792-0.821), 0.794 (95% CI: 0.777-0.811), 0.796 (95% CI: 0.761-0.830), 0.935 (95% CI: 0.901-0.969), and 0.807 (95% CI: 0.792-0.821), respectively. (Figure 2A-E) However, the PF-PPH-Logistic model exhibited only moderate discrimination with an AUC of 0.739 (95% CI: 0.666-0.813) in the internal validation dataset. Furthermore, the CA-PPH-Logistic model showed significantly lower discrimination in the external validation dataset with an AUC of 0.662 (95% CI: 0.450-0.873), which was notably inferior to its performance in the training and test datasets. This discrepancy may be attributed to the lower proportion of patients with coagulation disorders causing PPH in the validation dataset.

Assessment of Predictive Model Calibration

Calibration curves for the PPH and its subgroups were plotted for the Logistic predictive model within the training dataset (Supplemental Figure 1A-E). The performance of the PF-PPH-Logistic, particularly the CT-PPH-Logistic, and CA-PPH-Logistic models was suboptimal in certain aspects, with lower calibration, as observed in the test and external validation datasets (Supplemental Figure 1 F-J, Supplemental Figure 1K-O).

Evaluation of Clinical Utility of Predictive Models

In the evaluation of clinical utility, the PPH-Logistic and UA-PPH-Logistic models demonstrated satisfactory performance across all datasets. However, the clinical utility of the PF-PPH-Logistic, CT-PPH-Logistic, and CA-PPH-Logistic models was found to be relatively poor. (Supplemental Figure 2A-O)

Nomogram Construction

Using R software, we constructed nomograms for PPH and its four subgroups, with the results presented in Figure 3A-E. Physicians can assess the risk probability of PPH occurrence by summing the individual scores on the nomogram. This practical tool aids in a more precise estimation of PPH risk, thereby enhancing clinical decision-making.

Discussion

Maternal mortality has emerged as a pivotal indicator in global maternal and child health, serving as a significant benchmark for assessing the socioeconomic status of nations. Consequently, the effective reduction, prevention, and improvement of conditions leading to maternal deaths have become a focal point for public health initiatives worldwide. Among the various causes of maternal mortality, PPH stands out as a preventable condition that has attracted considerable attention. [17, 18] With the rise in global economic standards and the evolution of medical technologies, there has been an approximate 50% decrease in the worldwide maternal mortality rate between 1990 and 2015. In China, the maternal mortality rate has seen a dramatic reduction of 98.78%^[21] since the establishment of the People's Republic of China. Despite these advancements, a substantial proportion of maternal deaths, estimated between 27% and 40%^[22], remain avoidable due to a range of factors, including inadequate social and medical interventions. PPH is a critical area of focus within this context, and the prediction and prevention of PPH to

 reduce avoidable maternal mortality present a significant challenge on the global stage.

The advent of the big data era has brought new opportunities for the management of PPH. The era is characterized by the digitization and standardization of medical records, along with an increasing volume of data, which has ushered in an era of data-driven management and treatment for maternal care. Leveraging big data analytics for disease risk prediction can contribute to the reduction of avoidable maternal deaths.

A review of the literature reveals over 200 prognostic models in obstetrics, three of which are pertinent to PPH.^[23] However, few models have been applied in routine clinical practice, and the majority of studies have not provided model formulas, hindering independent external validation. The earliest PPH prediction model, dating back to 1994, originated from a case-control study in Zimbabwe^[24], where PPH was defined as blood loss exceeding 600 milliliters following an unassisted vaginal delivery. This study included 150 PPH patients and 299 patients with normal deliveries, with a low positive predictive value of less than 7% and only 35.0% of patients experiencing postpartum bleeding. Since then, approximately ten additional PPH prediction models have been published. These models have varied in focus, with some concentrating on the relationship between placenta previa and PPH, while others have included only vaginal deliveries[24-27] or cesarean sections[20, 28, 29], and some have targeted women with placental implantation disorders^[30] or general obstetric populations^[31]. PPH research has been conducted in hospitals across various countries, including Italy, China, France, the United States, the United Kingdom, South Korea, the Netherlands, Spain, Zimbabwe, Denmark, and Egypt. From the 14 published studies, a total of 124 independent variables were identified as potential predictors (ranging from 5 to 38 per study), and 64 variables were ultimately selected for the final models (an average of 5-15 factors per study). Common predictors included parity, low pre-pregnancy hemoglobin, antenatal bleeding, maternal age over 35, gestational age, high neonatal weight, multiple pregnancies, body mass index (BMI) over 25, previous cesarean section, anterior placenta, and retained placenta. These predictors have also been incorporated into our predictive model.

Once a clinical prediction model is developed, it must undergo validation and evaluation to assess the model's effectiveness, reproducibility, and portability. Published PPH prediction models have reported AUCs ranging from $0.70^{[25]}$ to $0.90^{[27]}$, with external validation AUCs of $0.83^{[20]}$, which are comparable to the results of our study. In addition to discrimination, calibration is essential to evaluate the consistency between the predicted probabilities of clinical outcomes and the observed event probabilities. Only a few studies, such as one by Albright in 2019 on the prediction of PPH following cesarean section, have utilized calibration curves^[32], while most have employed the Hosmer-Lemeshow goodness-of-fit test to compare predicted probabilities with actual event probabilities for significant differences. The Hosmer-Lemeshow test, however, has limited efficacy in small-sample prediction models as it does not quantify model calibration^[33, 34]or provide direction or magnitude of mis calibration^[35].

The Decision Curve Analysis (DCA)^[35] has been used to evaluate the clinical utility of models, focusing on the selection of true positives from positive patients to avoid unnecessary medical resource consumption and reduce harm from overtreatment of false positives. DCA is particularly suited for scenarios where symptoms suggest the possibility of disease but a diagnosis has not yet been confirmed, guiding the decision on whether or what kind of screening method to adopt for disease diagnosis. The DCA's axes represent the threshold probability (P) and net benefit (NB), allowing for the determination of intervention measures based on the predicted probability of adverse events.^[36]

In essence, both ROC and DCA can be used to assess the quality of predictive models, but they differ fundamentally in their theoretical constructs. While ROC combines sensitivity and specificity to compare the accuracy of predictive models through the AUC, the highest AUC does not necessarily represent the optimal model in clinical practice. For instance, in this study, patients in the CA-PPH group, due to coagulation disorders, all underwent cesarean section deliveries to minimize the number of false positives. This requires decision-makers to consider practical issues, as a high ROC does not always indicate the best treatment approach. Furthermore, for some

extreme cases, the accuracy of ROC becomes less critical, and DCA evaluation results are needed for reference.

Statistical analysis of previously published PPH data has shown that factors such as general anesthesia in pregnant women, prolonged use of oxytocin, excessive uterine tension (multiple pregnancies, polyhydramnios), and chorioamnionitis are all associated with uterine atony, potentially increasing the risk of postpartum bleeding. Previous studies have suggested that for pregnant women with high-risk factors, assessing and selecting appropriate treatment options and management based on the type and weight of different risks can reduce the risk of adverse pregnancy outcomes or death^[37]. Early prediction and intervention are key measures in reducing maternal mortality, with studies finding that timely interventions can effectively lower maternal mortality rates by 10%^[38]. Establishing a model that predicts the risk of PPH following vaginal delivery and guides clinical practice is a significant task for maternal and child health.

Limitation

Although some subtypes in this study have shown promising predictive results, the clinical application of the logistic regression clinical prediction models for PF-PPH, CT-PPH, and CA-PPH groups is limited due to the insufficient positive sample size in the subgroups, not fully achieving the initial goal of etiology-based PPH prediction. Compared to previously published prediction models, this study's test set is a single-center study specifically for vaginal delivery PPH, with a limited number of positive samples; the existing external validation datasets are also from hospitals of the same region and level, lacking in diversification and generalizability. Therefore, in addition to future plans for multi-center collaboration and increased sample sizes, we aim to seek higher-quality prediction methods to provide more convincing evidence for clinical prediction.

Conclusion

In conclusion, our study has successfully developed and validated predictive models for PPH following vaginal delivery, offering a novel approach to risk assessment in this critical area of maternal health. The models, particularly for UA-PPH and CT-PPH, demonstrated high discriminative power and clinical utility, with the nomogram providing a user-friendly tool for clinicians. Despite the promising results, limitations exist in the application of the PF-PPH, CT-PPH, and CA-PPH models due to the insufficient positive sample size in these subgroups. The generalizability of our findings may also be limited by the single-center nature of the study and the regional characteristics of the included hospitals. Future research should aim to expand the sample size and include multi-center data to improve the models' applicability and robustness. This study contributes to the growing body of evidence on PPH management and has the potential to influence policy and practice, ultimately enhancing maternal care and outcomes.

Figure legend

Figure 1. Patient Selection Criteria Flowchart.

Figure 2. AUC Curve for Postpartum Hemorrhage (PPH) Group and four subgroups. A. PPH Group; B. Uterine Atony PPH Group; C. Placental Factors PPH Group; D. Cervical Trauma PPH Group; E. Coagulation Abnormalities PPH Group. The blue line signifies the training dataset, which is employed to evaluate the model's predictive capabilities following the training phases. The green line corresponds to the internal validation dataset, pivotal for refining model parameters and for conducting initial assessments of the model's accuracy. The purple line denotes the external validation dataset, which is utilized to ascertain the model's generalizability and to verify its performance in an independent dataset.

Figure 3. Nomograms for Postpartum Hemorrhage (PPH) and Uterine Atony PPH Group. A. PPH Group; B. Uterine Atony PPH Group.

Supplemental Figure 1. Calibration curves for Postpartum Hemorrhage (PPH) Group and four subgroups. A\F\K. PPH Group; B\G\L. Uterine Atony PPH Group; C\H\M. Placental Factors PPH Group; D\I\N. Cervical Trauma PPH Group; E\J\O. Coagulation Abnormalities PPH Group. The blue line signifies the training dataset; The green line signifies the internal validation dataset; The purple line signifies the external validation dataset.

Supplemental Figure 2. Decision Curve Analysis for Postpartum Hemorrhage (PPH) Group and four subgroups. A. PPH Group; B. Uterine Atony PPH Group; C. Placental Factors PPH Group; D. Cervical Trauma PPH Group; E. Coagulation Abnormalities PPH Group. The blue line signifies the training dataset; The green line signifies the internal validation dataset; The purple line signifies the external validation dataset.

List of abbreviations

RP: Refined Peyton;

TTM: Traditional Teaching-mode

DOPS: direct observation of procedural skills

OB-GYN: Obstetrics and Gynecology CBVI: computer-based video instruction

PEARLS: Promoting Excellence and Reflective Learning in Simulation

OSCE: objective structured clinical assessment

Declarations

Ethics approval and consent to participate

Ethics approval and consent to participate: The study was approved by the Ethics Committee of Shengjing Hospital of China Medical University (No. 2016PS344K, Date.17/12/2016). All participants provided informed consent.

Consent for publication

Not Applicable

Availability of data and material

The data that support the findings of this study are available from the corresponding author upon reasonable request.

Competing interests

No potential conflict of interest was reported by the author(s).

Funding

This study was supported in part by grants from 345 Talent Project of Shengjing Hospital of China Medical University (No. M0946), and Medical Education Research Project of Liaoning Province (No. 2022-N005-03).

Authors' contributions

JL, DZ and KZ designed the study and drafted the manuscript. HL, MS, and XW done the data collection. YZ and XC designed the statistical analysis plan. DZ has participated the training and reviewed and co-authored the manuscript with ZS.

Acknowledgements

We would like to express our gratitude to all those who helped us during the writing of this manuscript. Thanks to all the peer reviewers for their opinions and suggestions. We would like to acknowledge that Jinke Li and Dandan Zhang have contributed equally to this work.

References:

- [1] World Health Organization. Trends in maternal mortality 2000 to 2017: estimates by WHO, UNICEF, UNFPA, World Bank Group and the United Nations Population Division [Z].
- [2] WHO U J G, DEPARTMENT OF REPRODUCTIVE HEALTH, RESEARCH W. UNFPA: Maternal Mortality in 2000: Estimates developed by WHO, UNICEF and UNFPA [J]. 2004.
- [3] ABOUZAHR C. Global burden of maternal death and disability [J]. British Medical Bulletin, 2003, 67(1): 1-11.
- [4] MEHRABADI A, HUTCHEON J A, LEE L, et al. Trends in postpartum hemorrhage from 2000 to 2009: a population-based study [J]. BMC Pregnancy Childbirth, 2012, 12: 108.
- [5] REALE S C, EASTER S R, XU X, et al. Trends in Postpartum Hemorrhage in the United States From 2010 to 2014 [J]. Anesth Analg, 2020, 130(5): e119-e22.
- [6] YOU J H S, LEUNG T Y. Cost-effectiveness analysis of carbetocin for prevention of postpartum hemorrhage in a low-burden high-resource city of China [J]. PloS one, 2022, 17(12): e0279130.
- [7] ABOUZAHR C J B M B. Global burden of maternal death and disability [J]. 2003, 67(1): 1-11.
- [8] MOUSA H A, WALKINSHAW S J C O I O, GYNECOLOGY. Major postpartum haemorrhage [J]. 2001, 13(6): 595-603.
- [9] AKTER S, FORBES G, VAZQUEZ CORONA M, et al. Perceptions and experiences of the prevention, detection, and management of postpartum haemorrhage: a qualitative evidence synthesis [J]. The Cochrane database of systematic reviews, 2023, 11(11): Cd013795.
- [10] ZHANG R, CAO X, FENG H, et al. Review of clinical practice guidelines for postpartum hemorrhage according to AGREE II [J]. Midwifery, 2023, 121: 103659.
- [11] ZDANOWICZ J A, SCHNEIDER S, MARTIGNONI C, et al. A Retrospective before and after Assessment of Multidisciplinary Management for Postpartum Hemorrhage [J]. Journal of clinical medicine, 2023, 12(23).

- [12] GIOULEKA S, TSAKIRIDIS I, KALOGIANNIDIS I, et al. Postpartum Hemorrhage: A Comprehensive Review of Guidelines [J]. Obstetrical & gynecological survey, 2022, 77(11): 665-82.
- [13] DE VRIES P L M, DENEUX-THARAUX C, BAUD D, et al. Postpartum haemorrhage in high-resource settings: Variations in clinical management and future research directions based on a comparative study of national guidelines [J]. BJOG: an international journal of obstetrics and gynaecology, 2023, 130(13): 1639-52.
- [14] KLUFIO C A, AMOA A B, KARIWIGA G. Primary postpartum haemorrhage: causes, aetiological risk factors, prevention and management [J]. Papua and New Guinea medical journal, 1995, 38(2): 133-49.
- [15] GYAMFI-BANNERMAN C, SRINIVAS S K, WRIGHT J D, et al. Postpartum hemorrhage outcomes and race [J]. American Journal of Obstetrics and Gynecology, 2018, 219(2): 185.e1-.e10.
- [16] !!! INVALID CITATION !!! [50, 51].
- [17] DESALE M, THINKHAMROP J, LUMBIGANON P, et al. Ending preventable maternal and newborn deaths due to infection [J]. Best practice & research Clinical obstetrics & gynaecology, 2016, 36: 116-30.
- [18] SOTUNSA J O, ADENIYI A A, IMARALU J O, et al. Maternal near-miss and death among women with postpartum haemorrhage: a secondary analysis of the Nigeria Near-miss and Maternal Death Survey [J]. BJOG: an international journal of obstetrics and gynaecology, 2019, 126 Suppl 3: 19-25.
- [19] GOAD L, ROCKHILL K, SCHWARZ J, et al. Development and validation of a prediction model for postpartum hemorrhage at a single safety net tertiary care center [J]. American journal of obstetrics & gynecology MFM, 2021, 3(5): 100404.
- [20] ALBRIGHT C M, SPILLANE T E, HUGHES B L, et al. A Regression Model for Prediction of Cesarean-Associated Blood Transfusion [J]. Am J Perinatol, 2019, 36(9): 879-85.
- [21] LIANG J, LI X, KANG C, et al. Maternal mortality ratios in 2852 Chinese counties, 1996-2015, and achievement of Millennium Development Goal 5 in China: a subnational analysis of the Global Burden of Disease Study 2016 [J]. Lancet, 2019, 393(10168): 241-52.
- [22] GAO Y, ZHOU H, SINGH N S, et al. Progress and challenges in maternal health in western China: a Countdown to 2015 national case study [J]. Lancet Glob Health, 2017, 5(5): e523-e36.
- [23] KLEINROUWELER C E, CHEONG-SEE F M, COLLINS G S, et al. Prognostic models in obstetrics: available, but far from applicable [J]. Am J Obstet Gynecol, 2016, 214(1): 79-90.e36.
- [24] TSU V D. Antenatal screening: its use in assessing obstetric risk factors in Zimbabwe [J]. J Epidemiol Community Health, 1994, 48(3): 297-305.
- [25] BIGUZZI E, FRANCHI F, AMBROGI F, et al. Risk factors for postpartum hemorrhage in a cohort of 6011 Italian women [J]. Thromb Res, 2012, 129(4): e1-7.
- [26] PRATA N, HAMZA S, BELL S, et al. Inability to predict postpartum hemorrhage: insights from Egyptian intervention data [J]. BMC Pregnancy Childbirth, 2011, 11: 97.
- [27] RUBIO-ÁLVAREZ A, MOLINA-ALARCÓN M, ARIAS-ARIAS Á, et al. Development and validation of a predictive model for excessive postpartum blood loss: A retrospective, cohort study [J]. Int J Nurs Stud, 2018, 79: 114-21.
- [28] AHMADZIA H K, PHILLIPS J M, JAMES A H, et al. Predicting peripartum blood transfusion in women undergoing cesarean delivery: A risk prediction model [J]. PloS one, 2018, 13(12):

e0208417.

- [29] DUNKERTON S E, JEVE Y B, WALKINSHAW N, et al. Predicting Postpartum Hemorrhage (PPH) during Cesarean Delivery Using the Leicester PPH Predict Tool: A Retrospective Cohort Study [J]. Am J Perinatol, 2018, 35(2): 163-9.
- [30] YOON S Y, YOU J Y, CHOI S J, et al. A combined ultrasound and clinical scoring model for the prediction of peripartum complications in pregnancies complicated by placenta previa [J]. Eur J Obstet Gynecol Reprod Biol, 2014, 180: 111-5.
- [31] CHI Z, ZHANG S, WANG Y, et al. Research of the assessable method of postpartum hemorrhage [J]. Technol Health Care, 2016, 24 Suppl 2: S465-9.
- [32] !!! INVALID CITATION !!! [73].
- [33] MOHER D, LIBERATI A, TETZLAFF J, et al. Preferred reporting items for systematic reviews and meta-analyses: the PRISMA statement [J]. PLoS Med, 2009, 6(7): e1000097.
- [34] STEYERBERG E W, VERGOUWE Y. Towards better clinical prediction models: seven steps for development and an ABCD for validation [J]. Eur Heart J, 2014, 35(29): 1925-31.
- [35] VICKERS A J, ELKIN E B. Decision curve analysis: a novel method for evaluating prediction models [J]. Med Decis Making, 2006, 26(6): 565-74.
- [36] HIJAZI Z, OLDGREN J, LINDBäCK J, et al. The novel biomarker-based ABC (age, biomarkers, clinical history)-bleeding risk score for patients with atrial fibrillation: a derivation and validation study [J]. Lancet, 2016, 387(10035): 2302-11.
- [37] ZUCKERWISE L C, LIPKIND H S. Maternal early warning systems-Towards reducing preventable maternal mortality and severe maternal morbidity through improved clinical surveillance and responsiveness [J]. Semin Perinatol, 2017, 41(3): 161-5.
- [38] AOYAMA K, D'SOUZA R, PINTO R, et al. Risk prediction models for maternal mortality: A systematic review and meta-analysis [J]. PloS one, 2018, 13(12): e0208563.

Table 1 The general view of the maternal.

Table 1 The general view of the mat	ernal.		
	Cohort population	Validation population	
Characteristics	(N=24,833)	(N=5009)	P
Age (years) *, N (%)			0.393
<25	1,309 (5.27%)	266 (5.31%)	
25-29	11,736 (47.26%)	2340 (46.71%)	
30-34	9,445 (38.03%)	1959 (39.11%)	
≥35	2,343 (9.44%)	444 (8.86%)	
Ethnicity, N (%)			0.983
Han	22,222 (89.49%)	4,475 (89%)	
Manchu	1,872 (7.54%)	389 (7.8%)	
Other ethnic groups	739 (2.98%)	145 (2.9%)	
Educational Attainment, N (%)			0.115
High school or below	8,635 (34.77%)	1,742 (35%)	
Bachelor's degree	13,639 (54.92%)	2,703 (54%)	
Postgraduate or higher	2,559 (10.30%)	564 (11%)	
Occupation, N (%)			0.777
Unemployed	11,373 (45.80%)	2,266 (45%)	
Light physical labor	2,825 (11.38%)	569 (11%)	
Moderate physical labor	10,011 (40.31%)	2,038 (41%)	
Heavy physical labor	624 (2.51%)	136 (2.7%)	
Family Per Capita Monthly			
Income (10,000 yuan), N (%)			0.9862
<0.5	10,325 (41.58%)	2,080 (42%)	
0.5-2.0	9,534 (38.39%)	1,922 (38%)	
2.0-5.0	3,584 (14.43%)	720 (14%)	
>5.0	1,390 (5.60%)	287 (5.7%)	
Pre-pregnancy BMI (Kg/m2)			
*, N (%)			< 0.001
<18.5 (Underweight)	7,294 (29.37%)	1475 (29.45%)	
18.5-23.9 (Normal)	15,005 (60.42%)	2963 (59.16%)	
24.0-27.9 (Overweight)	1,700 (6.85%)	307 (6.13%)	
>28.0 (Obesity)	834 (3.36%)	264 (5.29%)	
Pregnancy History, N (%)			0.565
1	14,985 (60.34%)	3,005 (60%)	
2	6,513 (26.23%)	1,303 (26%)	
≥3	3,335 (13.43%)	701 (14%)	
Parity (number of deliveries), N			
(%)			0.775
0	20,550 (82.75%)	4,127 (82%)	
1	4,152 (16.72%)	853 (17%)	
≥2	131 (0.53%)	29 (0.6%)	
Gestational Age at Delivery			
Sesentional rige at 2011, or 3			

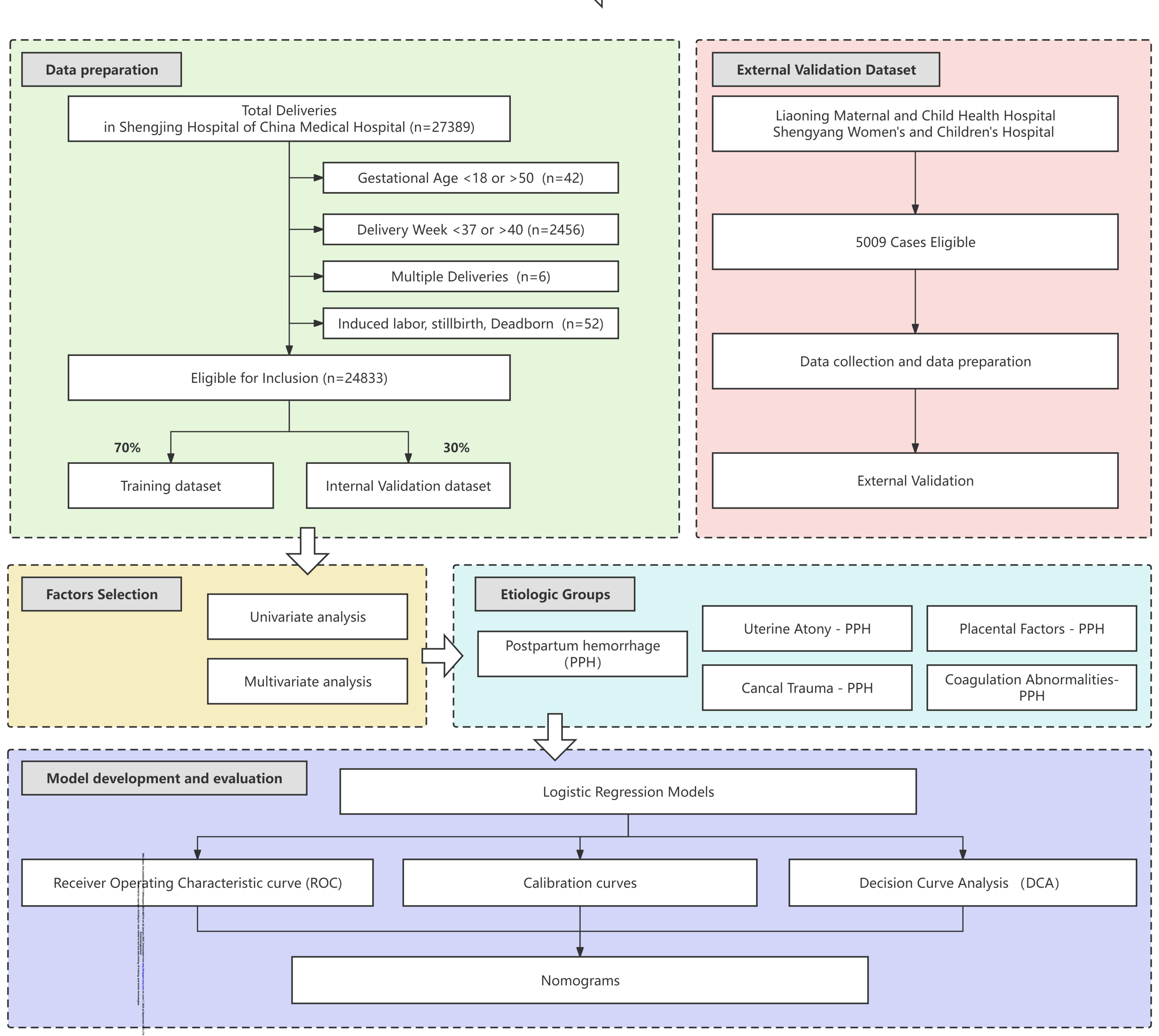
<38	1,507 (6.07%)	296 (5.91%)	
38-40	13,023 (52.44%)	2589 (51.59%)	
>40	10,303 (41.49%)	2129 (42.50%)	
Blood Loss (ml)	393.54±92.53	413.48±124.65	0.081
Postpartum Hemorrhage, N (%)	1,623 (6.54%)	286 (5.71%)	0.032
Due to uterine atony, N (%)	1,225 (4.93%)	266 (5.31%)	0.279
Due to placental factors, N (%)	242 (0.97%)	43 (0.86%)	0.489
Due to Cancal Trauma, N (%)	139 (0.56%)	31 (0.62%)	0.686
Due to coagulation disorders, N			
(%)	76 (0.31%)	17 (0.34%)	0.705

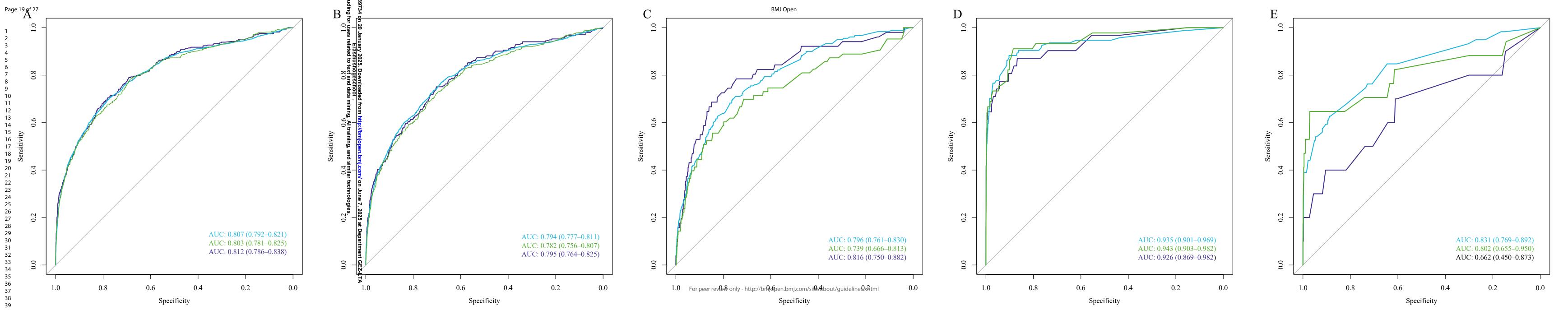
		BMJ Open		86/bmjopen-2024-089734 ป by copyright, including	
Table 2: Multivariate Risk Factor Analysis Characteristics	for Postpartum Hen PPH	orrhage (PPH) within UA-PPH	Subgroups PF-PPH	on	СА-РРН
Characteristics	rrn	General view	rr-rrn	Se CT-PPH	СА-ГГП
Age (years)	*	*	_	* ' * ' ' ' January 2025. Downlos Erasmushogesch	_
Ethnicity	**	**	**	* / 20 #Sm ed t	_
Educational Attainment		_	_	25. ush o te	_
Occupation	**	**	**	Xt a	*
Family Per Capita Monthly Income				vnlo	
(10,000 yuan)	100	-	-	oadeo hool	-
Pre-pregnancy BMI (Kg/m2)	**	**	**	omii **	*
Smoking	_		_	ning -	*
Alcohol Consumption	_	' /	_	a, http	_
Aconor Consumption	Obe	tetric and Gynecologic	- History	i.//b	_
Pregnancy History	-	-	-		_
Parity (number of deliveries)	**	**	**	α	*
History of miscarriage				nd:	
_	-	-		sim.	-
Spontaneous abortion	-	-	-0/-	ilar om/	-
Induced abortion or medication	-	-	-	ogeschool . At and data mining, Al training, and similar technologies	-
abortion				Jun	
Induced labor	-	-	-	7, <u>نون</u>	-
Assisted reproductive technology	-	-	-	 2025	-
Gestational Age at Delivery (weeks)	**	**	*	න - න	-
Diabetes	**	**	**	* * * at Departm	-
Hypertension	**	**	**	i) par	-
Anemia	**	**	**	**	*

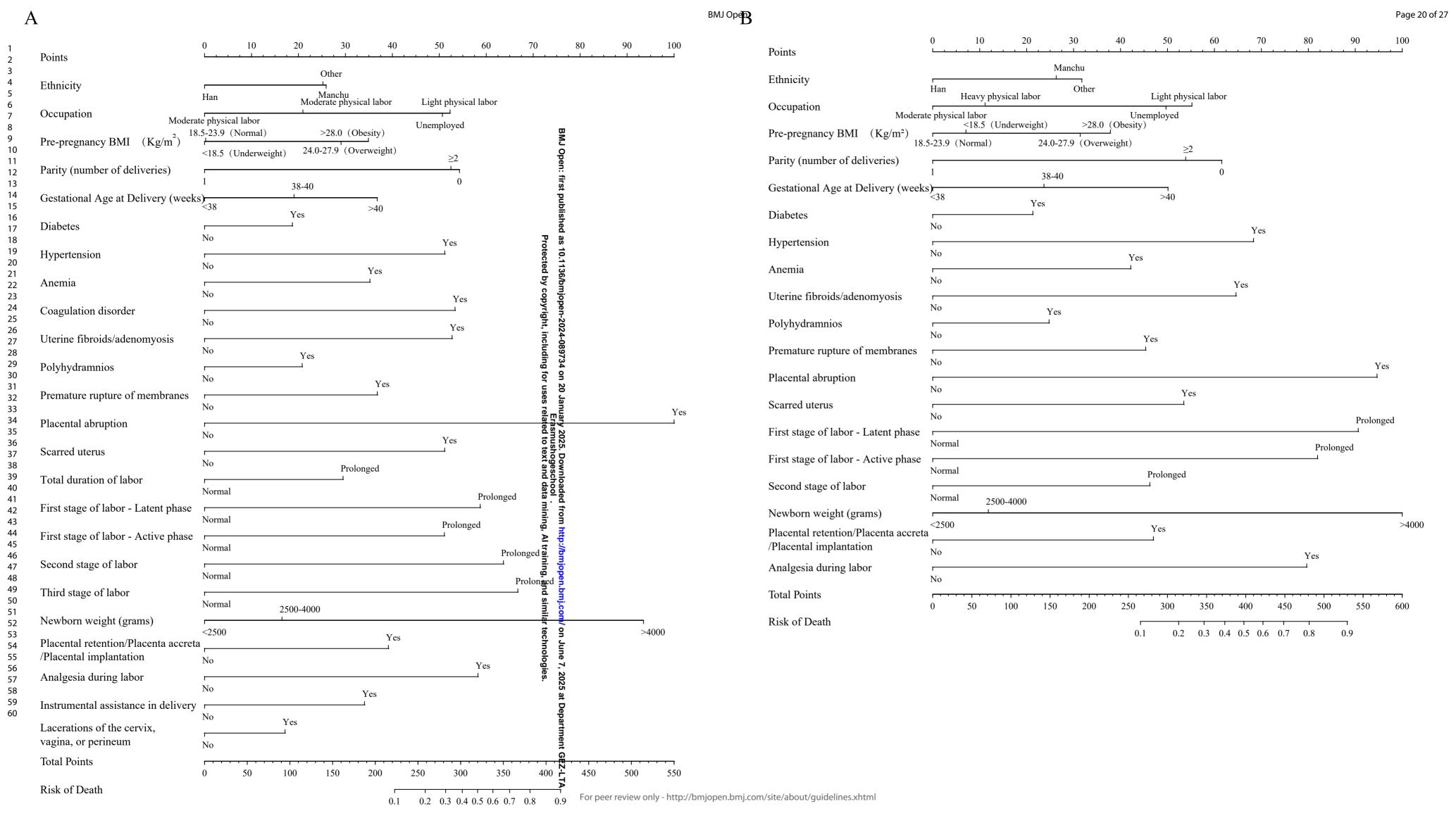
36/bmjopen-2024 d by copyright, ir

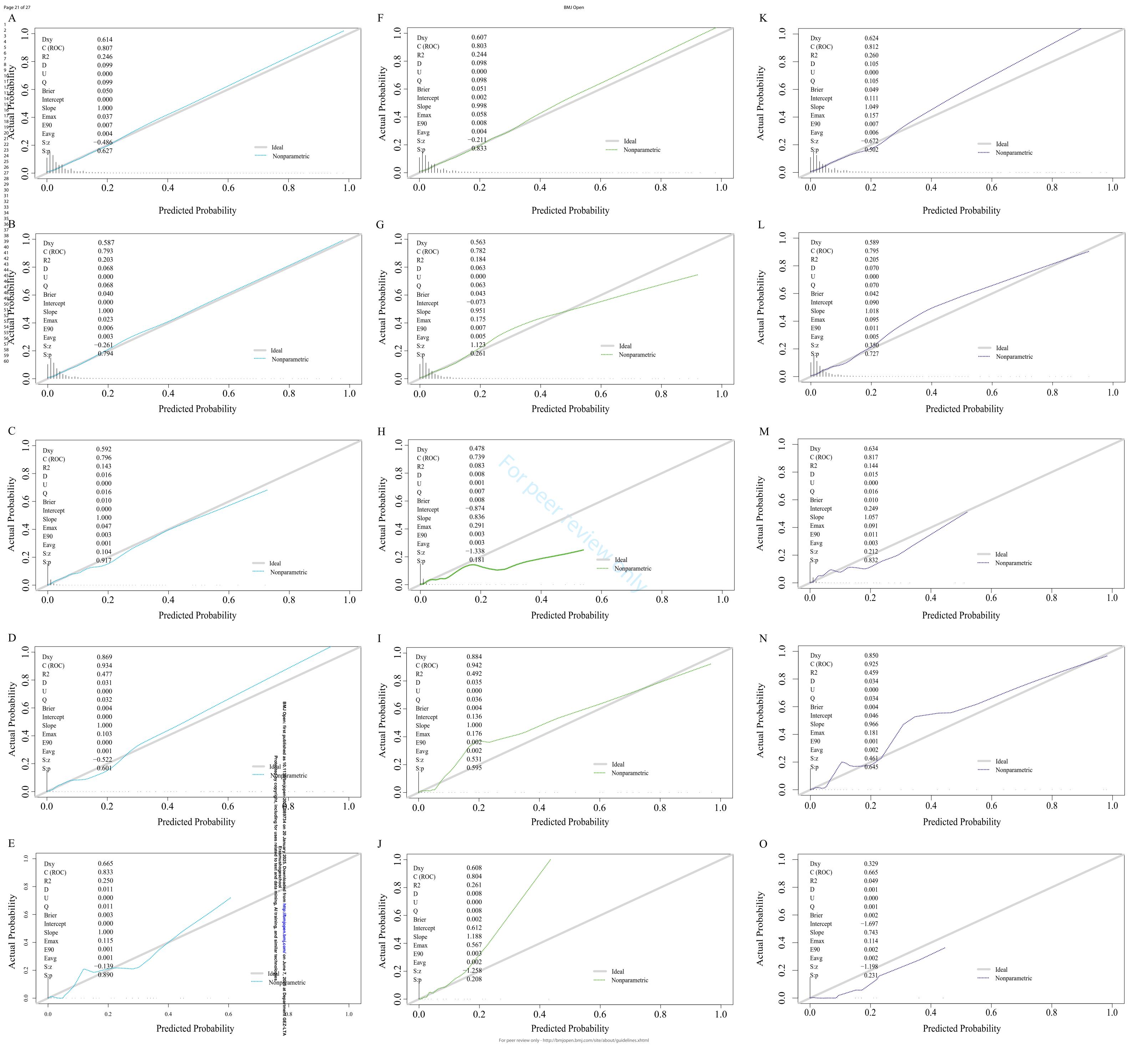
				4-089; includ		
Coagulation disorder	**	-	-	9734 Iding	-	*
Uterine fibroids/adenomyosis	**	**	**	on	*	-
Polyhydramnios	**	**	*	20 J use	-	-
Umbilical cord entanglement	_	-	_	Janı es r	-	-
Premature rupture of membranes	**	**	**	uary Era elate	*	**
Placental abruption	**	**	_	202 Smu ed to	_	-
Vaginal bleeding during pregnancy	-	-	-	ısho o te:	-	-
Scarred uterus	**	**	-	Xt age	-	*
	Deliver	y Process and Neonatal	Conditions	nd c		
Time of delivery		<u> </u>	-	adec ool lata	-	-
Total duration of labor	**	*	*	d from minin	*	**
First stage of labor - Latent phase	**	**	-	(0	-	-
First stage of labor - Active phase	**	**	-	http://bmjopen.bmj ,, Al training, and s	-	-
Second stage of labor	**	**	*	ttp://bmjope Al training,	**	-
Third stage of labor	**	-	**	ning	*	-
Placental retention/Placenta	**	**	4.4	en.br I, and	**	**
accreta/Placental implantation	<i>ሉ</i> ሉ	<i>*</i> *	**	od si	ጥ ጥ	<i>*</i> *
Analgesia during labor	**	**	- 0	nj.com/ similar	*	**
Instrumental assistance in delivery	**	-	-		**	-
Lacerations of the cervix, vagina, or	**			n Ju		
perineum	<i>ሉ</i> ሉ	-	-	June		-
Newborn weight (grams)	**	**	-	on June 7, 20 technologies	-	-
Newborn length (centimeters)	-	-	-	2025 a	-	-
*: P<0.05; **: P<0.001				at De		
				<u></u>		

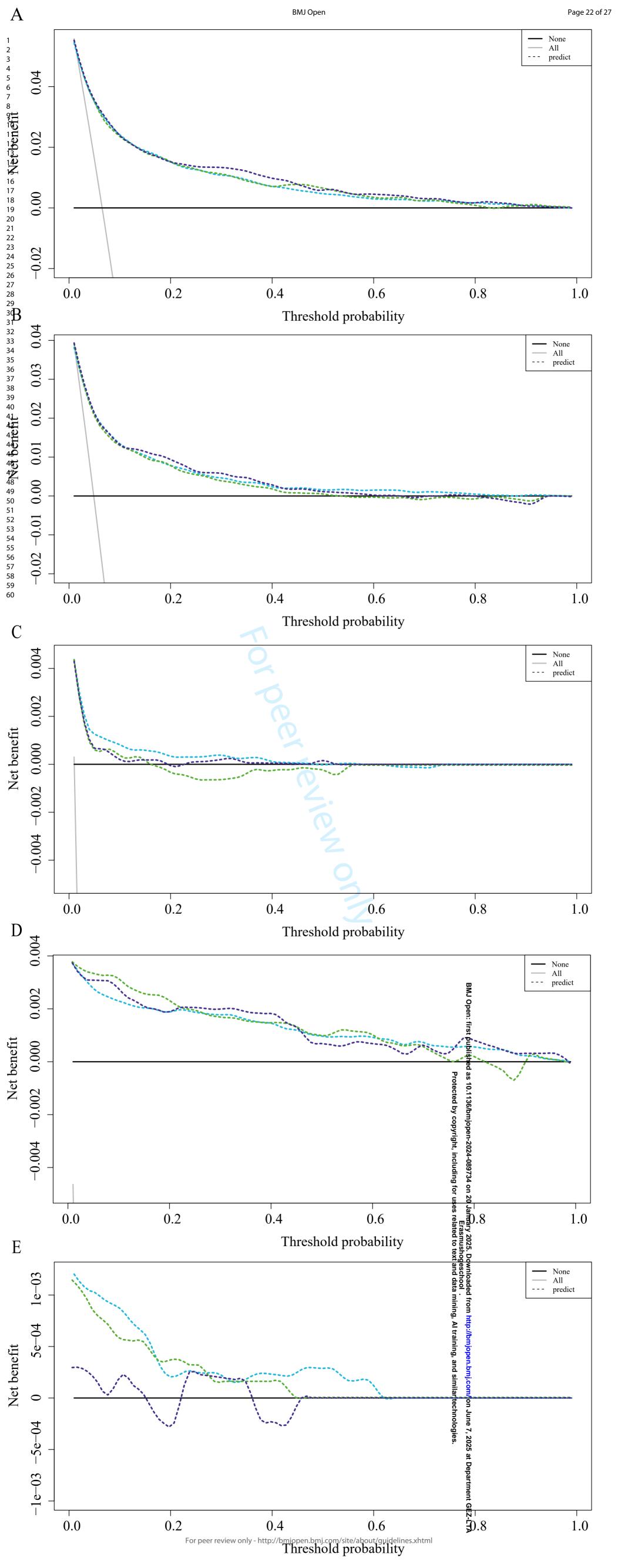












						В	МЈ Оре	en			36/bmjop d by copy				
Supplemental Table 1 Re Subgroups	esults of	f Predict Fac	tors Selection	on for t	he Multivari	ate Logistic	Regress	sion Model ir	ı the Traini	ng Data	ven-2024-089734 on persons on persons on the person of the	oartum Hei	norrhag	e (PPH) and I	Its
		PPH			UA-PPH			PF-PPH			`N			СА-РРН	
特征	OR	95% CI	P	OR	95% CI	P	OR	95% CI	P	OR	es selated t	P	OR	95% CI	P
						Ger	eral vie	ew			Eras Bate				
Age (years)											smushoge ed to text				
<25	Ref.			Ref.							5. Do Ishog Text				
25-29	0.94	0.67, 1.35	0.746	1.04	0.70, 1.59	0.852					Downloaded from ogeschool .ext and data mining				
30-34	1.28	0.91, 1.84	0.166	1.41	0.95, 2.16	0.099					nloa scho nd d:				
≥35	1.22	0.81, 1.85	0.349	1.25	0.78, 2.03	0.369					padec 1001 data				
thnicity											from h mining,				
Han	Ref.			Ref.			Ref.			Ref.	ing.				
Manchu	1.82	1.46, 2.24	<0.001**	1.59	1.24, 2.02	<0.001**	2.02	1.27, 3.08	0.002*	1.53	9 0.63 2 3.3‡	0.313			
Other ethnic groups	1.77	1.25, 2.45	<0.001**	1.75	1.20, 2.50	0.003*	1.47	0.57, 3.11	0.361	3.01	1.10 7.1	0.020*			
Educational											1.10 7.1 mjope				
Attainment											i, an				
High school or below											id si				
Bachelor's degree															
Postgraduate or higher											n√o ar te				
Occupation											//hamjopen.bmj.com/ on June 7, 20 tra/ning, and similar technologies.				
Unemployed	Ref.			Ref.			Ref.				nolo		Ref.		
Light physical labor	1.05	0.86, 1.27	0.647	1.12	0.91, 1.38	0.283	0.72	0.44, 1.13	0.171		7, 2 ogies		1.48	0.68, 3.01	0.2
Moderate physical labor			<0.001**	0.53	0.45, 0.64	<0.001**	0.43	0.29, 0.61	<0.001**		25		0.66	0.34, 1.25	0.2
Heavy physical labor		0.17, 0.53			0.17, 0.61		0.29	0.05, 0.94	0.090				0.44	0.02, 2.47	0.4
amily Per Capita		-			•			,			Оера			•	
Ionthly Income											artn				
•											at Department GEZ-LTA				
											GE				
											Σ̈-L				

BMJ Open

2 3 4

5 6

7

8

9 10

11

12

13 14

15

16 17

18 19 20

21

22 23

24

25

26 27

28

29

30

31 32

33

34

35 36

37

44 45 46 Page 24 of 27

BMJ Open

Page 25 of 27

2 3 4

5 6

8

9 10

11

12

13

14

15

16

17

18 19

20

21

22 23

24

25

26

27

28

29

30

31 32

33

34

35

36

						В	МЈ Оре	en			86/bmjopen-2024-089734 on 20 January 2025. Downloaded from http://bmjopen.bmj.com/ on June 7, 2025 at Departmen ERasmushogeschool . I by copyright, including for uses related to text and data mining, Al training, ਕ੍ਰੀ ਕ੍ਰਿੰਡimilar technologies.				
											2024-089 ht, inclu				
Premature rupture of											734 ding				
membranes											for				
No	Ref.			Ref.			Ref.			Ref.	20 J use		Ref.		
											anu s re				<0.001*
Yes	2.32	2.01, 2.68	<0.001**	2.18	1.85, 2.55	<0.001**	1.82	1.31, 2.51	<0.001**	1.33	0.77 6 7 6 7	0.294	2.63	1.50, 4.59	ŵ
Placental abruption											202 d to				
No	Ref.			Ref.							5. D sho tex				
		4.22,		11.0							owr ges t an				
Yes	9.98	22.51	<0.001**	0	4.66, 24.6	<0.001**					าloa cho d da				
Scarred uterus											ded ata i				
No	Ref.			Ref.							nini		Ref.		
Yes	3.01	1.65, 5.18	<0.001**	2.96	1.55, 5.29	<0.001**					ing,		6.11	0.87, 24.5	0.027*
					Deliver	ry Process a	nd Neo	natal Condit	ions		≱ t				
Total duration of labor											/bm rain				
normal	Ref.			Ref.			Ref.			Ref.	ing,		Ref.		
											291,				
prolonged	1.94	1.29, 2.88	0.001*	1.50	0.97, 2.28	0.064	2.42	1.03, 5.01	0.027*	5.16	1263	<0.001**	3.78	0.95, 11.2	0.032*
First stage of labor -											nila				
Latent phase											/ on				
normal	Ref.			Ref.							를 다				
prolonged	3.83	2.65, 5.48	<0.001**	4.42	3.02, 6.38	<0.001**					ne 7 oloç				
First stage of labor -											', 20 jies				
Active phase											. 125				
normal	Ref.			Ref.							at D				
prolonged	3.17	2.30, 4.32	<0.001**	3.81	2.74, 5.24	<0.001**					ера				
Second stage of labor											rtmen				

BMJ Open

Page 27 of 27

2 3

5

6

7

8

9 10

11

12

13

14 15

16

17

18 19

20

21

22 23

24

25

26 27

28

29

30

31

32

33

34

35 36

37

36/bmjopen-20

							36.1	024-08973 t, includin
Yes	1.47	1.19, 1.79	<0.001**				0	6.2014 On <0.001**
Newborn weight								20 J
(grams)								anu s re
<2500	Ref.			Ref.	1.02,			ary 202 Erasmu lated to
2500-4000	1.50	0.69, 4.00	0.358	3.30 15.1	20.31 4.57,	0.100		5. Dow ishoges
>4000	8.50	3.79, 23.0	<0.001**	0	93.62	<0.001**		nloa schc
*: P<0.05; **: P<	0.001; OR	: Odds Rati	o; CI: Conf	fidence I	nterval;		evien on	000 0000 0000 0000 0000 0000 0000 0000 0000

^{*:} P<0.05; **: P<0.001; OR: Odds Ratio; CI: Confidence Interval;

BMJ Open

Postpartum Hemorrhage Following Vaginal Delivery: A Comprehensive Analysis and Development of Predictive Models for Etiological Subgroups in Chinese Women

Journal:	BMJ Open						
Manuscript ID	bmjopen-2024-089734.R1						
Article Type:	Original research						
Date Submitted by the Author:	08-Nov-2024						
Complete List of Authors:	Li, Jinke; Shengjing Hospital of China Medical University, Department of Obstetrics and Gynecology Zhang, Dandan; Shengjing Hospital of China Medical University, Department of Obstetrics and Gynecology Lin, Hong; Liaoning Maternal and Child Health Hospital, Department of Obstetrics and Gynecology Shao, Mengyuan; Shenyang Women's and Children's Hospital, Department of Obstetrics and Gynecology Wang, Xiaoxue; Shengjing Hospital of China Medical University, Department of Health Management Chen, Xueting; Shengjing Hospital of China Medical University, Department of Health Management Zhou, Yangzi; Shengjing Hospital of China Medical University, Department of Obstetrics and Gynecology Song, Zixuan; Shengjing Hospital of China Medical University, Department of Obstetrics and Gynecology						
Primary Subject Heading :	Obstetrics and gynaecology						
Secondary Subject Heading:	Obstetrics and gynaecology						
Keywords:	Maternal medicine < OBSTETRICS, PERINATOLOGY, Postpartum Women < Postpartum Period, PREVENTIVE MEDICINE						

SCHOLARONE™ Manuscripts



I, the Submitting Author has the right to grant and does grant on behalf of all authors of the Work (as defined in the below author licence), an exclusive licence and/or a non-exclusive licence for contributions from authors who are: i) UK Crown employees; ii) where BMJ has agreed a CC-BY licence shall apply, and/or iii) in accordance with the terms applicable for US Federal Government officers or employees acting as part of their official duties; on a worldwide, perpetual, irrevocable, royalty-free basis to BMJ Publishing Group Ltd ("BMJ") its licensees and where the relevant Journal is co-owned by BMJ to the co-owners of the Journal, to publish the Work in this journal and any other BMJ products and to exploit all rights, as set out in our licence.

The Submitting Author accepts and understands that any supply made under these terms is made by BMJ to the Submitting Author unless you are acting as an employee on behalf of your employer or a postgraduate student of an affiliated institution which is paying any applicable article publishing charge ("APC") for Open Access articles. Where the Submitting Author wishes to make the Work available on an Open Access basis (and intends to pay the relevant APC), the terms of reuse of such Open Access shall be governed by a Creative Commons licence – details of these licences and which Creative Commons licence will apply to this Work are set out in our licence referred to above.

Other than as permitted in any relevant BMJ Author's Self Archiving Policies, I confirm this Work has not been accepted for publication elsewhere, is not being considered for publication elsewhere and does not duplicate material already published. I confirm all authors consent to publication of this Work and authorise the granting of this licence.

Jinke Li^{1#}, Dandan Zhang^{1#}, Hong Lin², Mengyuan Shao³, Xiaoxue Wang⁴, Xueting Chen⁴, Yangzi Zhou¹ and Zixuan Song^{1*}

- 1 Department of Obstetrics and Gynecology, Shengjing Hospital of China Medical University, Shenyang, China
- 2 Department of Obstetrics and Gynecology, Liaoning Maternal and Child Health Hospital, Shenyang, China
- 3 Department of Obstetrics and Gynecology, Shenyang Women's and Children's Hospital, Shenyang, China
- 4 Department of Health Management, Shengjing Hospital of China Medical University, Shenyang, China
- #These authors contributed equally to this work.
- Correspondence:songzixuan666@163.com

Abstract:

Objectives: This study aimed to dissect the etiological subgroups of postpartum hemorrhage (PPH) that occur after vaginal delivery in women with full-term singleton pregnancies. Our goal was to craft and validate predictive models to guide clinical decision-making and optimize resource allocation.

Design: A retrospective cohort study.

Setting: Shengjing Hospital of China Medical University, Liaoning Maternal and Child Health Hospital, and Shenyang Women's and Children's Hospital.

Participants: 29,842 women who underwent vaginal delivery were enrolled in the study across three hospitals from 2016 to 2022.

Primary outcome measures: PPH, categorized into uterine atony (UA), placental factors (PF), cervical trauma (CT), and coagulation abnormalities (CA) by etiology.

Results: The logistic regression for overall PPH and UA-PPH showcased high discrimination (AUCs of 0.807 and 0.794, respectively), coupled with commendable calibration and DCA-assessed clinical utility, culminating in the development of a nomogram for risk prediction. The PF-PPH model exhibited a modest AUC of 0.739, while the CT-PPH and CA-PPH models demonstrated suboptimal clinical utility and calibration.

Conclusions: The study identified factors associated with PPH and developed models with good performance for overall PPH and UA-PPH. The nomogram offers a valuable tool for risk prediction. However, models for PF-PPH, CT-PPH, and CA-PPH require further refinement. Future research should focus on larger samples and multi-center validation for enhanced model generalizability.

Keywords: Postpartum Hemorrhage, Vaginal Delivery, Etiological Subgroups, Predictive Models, nomogram

Strengths and limitations of this study

- This study included data from large multicenter cohorts in China, comprising 29,842 women to enhance the statistical power of the analysis.
- Predictive models were developed for postpartum hemorrhage following vaginal delivery and for different etiological subgroups.

- This study's test set is a single-center study specifically focused on vaginal delivery PPH, with a limited number of positive samples.
- The existing external validation datasets are from hospitals of the same region and level, resulting in a lack of diversification and generalizability.

Background

Postpartum hemorrhage (PPH) is defined as blood loss exceeding 500 milliliters within 24 hours following vaginal delivery or exceeding 1,000 milliliters within 24 hours following cesarean delivery.¹ PPH is a widespread and serious medical condition that poses significant risks to women's health around the world. It is particularly devastating in developing countries, where it is a principal contributor to maternal mortality.² It is estimated that approximately 1.4 million maternal deaths globally are tied to PPH each year, with the tragic loss of a woman's life to this condition occurring every four minutes.³⁴. In Australia, the incidence of PPH increased from 6.3% in 2000 to 8.0% in 2009. ⁵ Similarly, in the United States, the rate of PPH rose from 2.7% in 1999 to 3.2% in 2014. ⁶ In China, despite a relatively lower maternal mortality rate of 17.8 per 100,000 in 2019, PPH accounted for one-quarter of these deaths.⁷

The World Health Organization (WHO) has conducted an analysis revealing that while PPH is a significant factor in maternal mortality and morbidity, the mortality rates vary considerably across different regions.⁸ In high-income countries, the risk of death due to PPH is significantly lower than in low-income countries.⁹ In high-income nations, the substantial blood loss primarily caused by PPH accounts for 13.4% of overall maternal mortality, while in Africa and Asia, this figure stands at 34% and 30.8%, respectively. ⁸ The international obstetric community is actively engaged in research to better understand the incidence, risk factors, and management strategies for PPH.¹⁰⁻¹². Despite the establishment of global clinical guidelines and the identification of various risk factors, further exploration is needed to enhance our understanding and management of PPH.^{13 14}.

PPH can be etiologically classified into uterine atony (UA), placental factors (PF), cervical trauma (CT), and coagulation abnormalities (CA), each requiring distinct clinical management and treatment strategies. ¹⁵ Clear etiological classification is crucial for developing preventive strategies, formulating management plans, and rational allocation of medical resources. ¹⁶ While numerous cohort studies have focused on identifying risk factors for PPH, there is a scarcity of studies that quantify and weigh these risk factors for a comprehensive PPH risk assessment. ¹⁷ ¹⁸ Given the complexity of PPH and the interplay of multiple risk factors, a holistic approach is necessary to accurately assess the risk of PPH.

Clinical prediction models (CPMs) have been widely applied in clinical settings in recent years. By constructing CPMs, physicians and patients can make better medical decisions, and health departments at all levels can allocate medical resources more rationally. These models play an irreplaceable role in primary prevention (assessing the quantitative risk of future diseases) and secondary prevention (constructing highly sensitive and specific diagnostic schemes, practicing "early detection, early diagnosis, early treatment"), reflecting significant health economic value.

There is a gap in research regarding the development of clinical prediction models for women specifically following vaginal delivery. Many studies are constrained by limited sample sizes, which can affect the robustness of the models 19. Other research has focused on PPH prediction models for women undergoing cesarean sections. 20 Our study aims to address this gap by constructing a clinical prediction model tailored to PPH after vaginal delivery. By analyzing clinical data and risk factors through logistic regression, we can determine the relative impact of each factor on the likelihood of PPH. We further refine our model by performing secondary fitting based on the four etiological subgroups, creating a nomogram that enhances the precision of predicting high-risk populations for PPH. This work provides essential insights for the prevention and management of this critical condition.

Materials and methods

Data Sources and Ethics Statement

This cohort study was conducted at the obstetric wards of Shengjing Hospital of China Medical University, Liaoning Maternal and Child Health Hospital, and Shenyang Women's and Children's Hospital. The study population comprised women who underwent vaginal delivery between January 1, 2016, and December 31, 2022. The outcomes of interest were fetal birth outcomes within the first 24 hours postpartum. Inclusion criteria were women who consented to participate after being informed of the study's scope. Exclusion criteria were defined as follows: age under 18 or over 50 years, delivery occurring at less than 37 weeks or more than 42 weeks of gestation, multiple births, and instances of induced labor, stillbirth, or fetal death. Comprehensive data encompassing maternal characteristics, obstetric and gynecologic history, pregnancy complications, and details of the delivery process and neonatal conditions were collected (Supplement 1). To protect participant privacy, all data were anonymized. The study protocol was approved by the Ethical Review Committee of Shengjing Hospital of China Medical University (No. 2016PS344K), and written information about the study was provided to all participants.

Sample Size Calculation

According to the obstetric big data from Shengjing Hospital of China Medical University, the incidence of PPH is approximately 6%-7%. Based on the sample size estimation using the psamplesize package in R software, the minimum sample size required for constructing clinical prediction models is estimated to be between 1,048 and 1,536. The sample size included in this study far exceeds the minimum requirement.

Covariates

A range of covariates were taken into account, including:(1) Age, categorized as <25, 25-29, 30-34, and ≥35 years; (2) Ethnicity, divided into Han, Manchu, and other; (3) Education level, classified as high school or below, bachelor's degree, and postgraduate or above; (4) Occupation, categorized as unemployed, light physical labor, moderate physical labor, and heavy physical labor (based on the International Physical Activity Questionnaire, IPAQ); (5) Monthly household income per capita, divided into <0.5, 0.5-2.0, 2.0-5.0, and >5.0 thousand yuan; (6) Pre-pregnancy BMI, categorized as underweight (<18.5 kg/m²), normal (18.5-23.9 kg/m²), overweight (24-27.9 kg/m²), and obese (\geq 28 kg/m²); (7) Smoking and alcohol consumption history; (8) Gravidity, categorized as 1, 2, or \geq 3 times; (9) Parity, divided into 0, 1, or ≥ 2 times; (10) History of miscarriage and induced labor; (11) Assisted reproductive technology; (11) Gestational age, categorized as <38, 38-40, and >40 weeks; (12) Pregnancy complications: diabetes, hypertension, anemia, coagulation disorder, uterine fibroids/adenomyosis, polyhydramnios, umbilical cord entanglement, premature rupture of membranes, placental abruption, vaginal bleeding during pregnancy, and scarred uterus; (13) Delivery time, divided into daytime (8-16), evening (17-23), and night (0-7) shifts; (14) Total duration of labor, categorized as normal (≤24 hours) and prolonged (>24 hours); (15) Latent phase of the first stage, categorized as normal (primiparous ≤20 hours, multiparous ≤14 hours) and prolonged (primiparous >20 hours, multiparous >14 hours); (16) Active phase of the first stage, categorized as normal (≤8 hours) and prolonged (>8 hours); (17) Second stage duration, categorized as normal and prolonged based on specific criteria for primiparous and multiparous women with or without analgesia; (18) Third stage duration, categorized as normal (≤30 minutes) and prolonged (>30 minutes); (19) Placental retention/placenta accreta/placental implantation; (20) Analgesia during labor; (21) Instrumental assistance in delivery; (22) Lacerations of the cervix, vagina, or perineum; (23) Newborn weight and length.

Etiology Subgroups

Patients were categorized into groups with or without postpartum hemorrhage based on the presence of postpartum bleeding and its underlying etiology, which included UA, PF, CT, and CA.

Model Construction

For the purpose of our investigation, we have categorized the participants from the Shengjing hospital of China Medical University as Cohort I. This cohort was systematically split into a training dataset and an internal validation dataset with a ratio of 7:3. The training dataset was instrumental in developing the predictive model, while the internal validation dataset served to assess the model's predictive accuracy. An additional cohort, comprising participants from two other hospitals, was designated as Cohort II. This external dataset was used to validate the model's general applicability and its efficacy in real-world clinical scenarios.

Within the confines of the datasets, we employed both univariate and multivariate logistic regression analyses to identify potential risk factors across various subgroups. These factors were then subjected to a rigorous selection process for inclusion in the predictive model. The selected factors were further analyzed using multivariate logistic regression (Bidirectional elimination) in training dataset to discern their discriminative power, thereby establishing them as predictive indicators for the model. The women in the training set were divided into PPH and non-PPH groups based on PPH as the outcome variable. Subsequently, they were further categorized by etiology into UA-PPH and non-UA-PPH groups, PF-PPH and non-PF-PPH groups, CT-PPH and non-CT-PPH groups, and CA-PPH and non-CA-PPH groups. Five predictive models were constructed sequentially, and the performance of these models was corroborated using both the test and validation datasets to ascertain the most accurate predictive model.

Evaluating the Performance of the Models

The area under the receiver operating characteristic curve (AUC) was the primary metric used to evaluate the discrimination of our models. An AUC value above 0.75 suggests excellent model discrimination, while an AUC below 0.6 indicates poor discrimination. Calibration curves were used to assess the models' accuracy, with closer alignment between observed and predicted incidence rates indicating higher model fidelity. Decision Curve Analysis (DCA) was also employed to evaluate the clinical utility of the models, offering a thorough assessment of the models' net benefits across various clinical scenarios.

Nomogram Development

Nomograms for postpartum hemorrhage and its four etiological subgroups were crafted to offer a visual representation of the risk scores derived from the logistic regression analysis. This tool simplifies the interpretation of complex statistical outcomes, providing a more straightforward approach to understanding risk assessments.

Statistical Analysis

All statistical computations, construction of traditional logistic models, and calculations of model discrimination and calibration were carried out using R version 3.6.3 from the R Foundation for Statistical Computing, Vienna, Austria. This software facilitated the development of traditional logistic predictive models and their subsequent evaluation for discriminative power, calibration, and clinical utility. Continuous variables conforming to the normal distribution were expressed as the mean \pm standard deviation (SD), while non-normally distributed continuous variables were presented as medians with interquartile ranges. Categorical data were analyzed using chi-square tests, and continuous variables were analyzed using ANOVA or Mann-Whitney tests, as appropriate. Variables were adjusted as dummy variables, and odds ratios (OR) with corresponding 95% confidence intervals (95% CI) were calculated using univariate and multivariate logistic regression analyses, with significance level set at P < 0.05.

Patient and public involvement

Patients and/or the public were not involved in the design, conduct, reporting or dissemination plans of this research.

From 2016 to 2022, a total of 27,389 patients underwent vaginal delivery at the Shengjing Hospital of China Medical University. Forty-two patients under 18 years of age or over 50 years old were excluded. Additionally, 2,456 patients with gestational age less than 37 weeks or more than 42 weeks at delivery, 6 patients with multiple births, and 52 patients with induced labor, stillbirth, or fetal death were also excluded. Ultimately, 24,833 patients met the inclusion criteria and were enrolled in the cohort. According to the inclusion and exclusion criteria, a total of 5,099 patients in cohort II were included in the external validation dataset. The general characteristics of all patients are presented in Table 1. All patients were followed up within 24 hours after delivery for neonatal outcomes, with a follow-up rate of 100%. The patient selection criteria flowchart is shown in Figure 1.

Comparison of Basic Characteristics and Risk Analysis for Postpartum Hemorrhage (PPH) and Its Subgroups

Based on the occurrence of postpartum hemorrhage, the parturients in cohort one were divided into two groups: the non-PPH group and the PPH group. Similarly, within the etiological subgroups, they were categorized into UA-PPH and non-UA-PPH groups, PF-PPH and non-PF-PPH groups, CT-PPH and non-CT-PPH groups, and CA-PPH and non-CA-PPH groups. The comparison of basic characteristics and analysis of risk factors for each group are presented in Supplementary Tables 1-5.

In the multivariate analysis of risk factors, apart from age, parity, pre-pregnancy BMI, anemia, premature rupture of membranes, and combined placenta retention/placenta accreta/placental implantation, other specific risk factors were found to be associated with specific etiologies of postpartum hemorrhage. For instance, polyhydramnios was associated with UA-PPH; analgesia during labor, instrumental assistance, and cervical/vaginal/perineal lacerations were associated with the occurrence of CT-PPH (Table 2).

Selection of Predictive Factors for PPH and Its Subgroups in the Training Dataset

Through random sampling of cohort one, 70% of the data (N=17,383) from parturients were used to form the training dataset, with the remaining approximately 30% (N=7,450) forming the internal validation dataset. Multivariate analysis of risk factors for PPH and its subgroups was performed again in the training dataset, with results presented in Supplemental Table 1. After selection, predictive models were constructed for each group using the selected risk factors.

Evaluation of Predictive Model Discrimination

The ROC curves were plotted using R software for the PPH group and its various subgroups across the training dataset, internal and external validation dataset.

The results indicated that the predictive models, namely PPH-Logistic, UA-PPH-Logistic, PF-PPH-Logistic, CT-PPH-Logistic, and CA-PPH-Logistic, demonstrated high discriminative power in the training dataset with AUCs of 0.807 (95% CI: 0.792-0.821), 0.794 (95% CI: 0.777-0.811), 0.796 (95% CI: 0.761-0.830), 0.935 (95% CI: 0.901-0.969), and 0.802 (95% CI: 0.769-0.892), respectively.(Figure 2A-E) However, the PF-PPH-Logistic model exhibited only moderate discrimination with an AUC of 0.739 (95% CI: 0.666-0.813) in the internal validation dataset. Furthermore, the CA-PPH-Logistic model showed significantly lower discrimination in the external validation dataset with an AUC of 0.662 (95% CI: 0.450-0.873), which was notably inferior to its performance in the training and test datasets. This discrepancy may be attributed to the lower proportion of patients with coagulation disorders causing PPH in the validation dataset.

Assessment of Predictive Model Calibration

Calibration curves for the PPH and its subgroups were plotted for the Logistic predictive model within the training dataset (Supplemental Figure 1A-E). The performance of the PF-PPH-Logistic, particularly the CT-PPH-Logistic, and CA-PPH-Logistic models was suboptimal in certain aspects, with lower calibration, as observed in the test and external validation datasets (Supplemental Figure 1 F-J, Supplemental Figure 1K-O).

Evaluation of Clinical Utility of Predictive Models

In the evaluation of clinical utility, the PPH-Logistic and UA-PPH-Logistic models demonstrated satisfactory performance across all datasets. However, the clinical utility of the PF-PPH-Logistic, CT-PPH-Logistic, and CA-PPH-Logistic models was found to be relatively poor. (Supplemental Figure 2A-O)

Nomogram Construction

Using R software, we constructed nomograms for PPH and UA-PPH, with the results presented in Figure 3A-B. Physicians can assess the risk probability of PPH occurrence by summing the individual scores on the nomogram. This practical tool aids in a more precise estimation of PPH risk, thereby enhancing clinical decision-making.

Discussion

Maternal mortality has emerged as a pivotal indicator in global maternal and child health, serving as a significant benchmark for assessing the socioeconomic status of nations. Consequently, the effective reduction, prevention, and improvement of conditions leading to maternal deaths have become a focal point for public health initiatives worldwide. Among the various causes of maternal mortality, PPH stands out as a preventable condition that has attracted considerable attention. ¹⁷ ¹⁸ With the rise in global economic standards and the evolution of medical technologies, there has been an approximate 50% decrease in the worldwide maternal mortality rate between 1990 and 2015. In China, the maternal mortality rate has seen a dramatic reduction of 98.78%²¹ since the establishment of the People's Republic of China. Despite these advancements, a substantial proportion of maternal deaths, estimated between 27% and 40%²², remain avoidable due to a range of factors, including inadequate social and medical interventions. PPH is a critical area of focus within this context, and the prediction and prevention of PPH to reduce avoidable maternal mortality present a significant challenge on the global stage.

Early prediction or identification of PPH and timely preventive or intervention measures are extremely valuable, necessitating a clear understanding of the etiologies of PPH for targeted management. In this study, the predictive models developed for PPH and UA-PPH demonstrated excellent performance in effectively identifying high-risk populations. For women at high risk for UA-PPH, early cord clamping should be considered. After the placenta is delivered, uterine fundal massage can be performed, and oxytocin should be administered immediately to promote uterine contraction and reduce bleeding. If necessary, mechanical compression or uterine artery ligation can be employed for hemostasis. Overall, high-risk women for PPH should be closely monitored, and proactive interventions should be implemented, such as promoting uterine contractions, advising patients to avoid excessive straining that could cause lacerations, and timely correction of coagulopathy.

The advent of the big data era has brought new opportunities for the management of PPH. The era is characterized by the digitization and standardization of medical records, along with an increasing volume of data, which has ushered in an era of data-driven management and treatment for maternal care. Leveraging big data analytics for disease risk prediction can contribute to the reduction of avoidable maternal deaths.

A review of the literature reveals over 200 prognostic models in obstetrics, three of which are pertinent to PPH.²³ However, few models have been applied in routine clinical practice, and the majority of studies have not provided model formulas, hindering independent external validation. The earliest PPH prediction model, dating back to 1994, originated from a case-control study in Zimbabwe²⁴, where PPH was defined as blood loss exceeding 600 milliliters following an unassisted vaginal delivery. This study included 150 PPH patients and 299 patients with normal deliveries, with a low positive predictive value of less than 7% and only 35.0% of patients experiencing postpartum bleeding. Since then, approximately ten additional PPH prediction models have been published. These models have varied in focus, with some concentrating on the relationship between placenta previa and PPH, while others have included only vaginal

Once a clinical prediction model is developed, it must undergo validation and evaluation to assess the model's effectiveness, reproducibility, and portability. Published PPH prediction models have reported AUCs ranging from 0.70^{25} to 0.90^{27} , with external validation AUCs of 0.83^{20} , which are comparable to the results of our study. In addition to discrimination, calibration is essential to evaluate the consistency between the predicted probabilities of clinical outcomes and the observed event probabilities. Only a few studies, such as one by Albright in 2019 on the prediction of PPH following cesarean section, have utilized calibration curves³², while most have employed the Hosmer-Lemeshow goodness-of-fit test to compare predicted probabilities with actual event probabilities for significant differences. The Hosmer-Lemeshow test, however, has limited efficacy in small-sample prediction models as it does not quantify model calibration^{33 34}or provide direction or magnitude of mis calibration³⁵.

The Decision Curve Analysis (DCA)³⁵ has been used to evaluate the clinical utility of models, focusing on the selection of true positives from positive patients to avoid unnecessary medical resource consumption and reduce harm from overtreatment of false positives. DCA is particularly suited for scenarios where symptoms suggest the possibility of disease but a diagnosis has not yet been confirmed, guiding the decision on whether or what kind of screening method to adopt for disease diagnosis. The DCA's axes represent the threshold probability (P) and net benefit (NB), allowing for the determination of intervention measures based on the predicted probability of adverse events.³⁶

In essence, both ROC and DCA can be used to assess the quality of predictive models, but they differ fundamentally in their theoretical constructs. While ROC combines sensitivity and specificity to compare the accuracy of predictive models through the AUC, the highest AUC does not necessarily represent the optimal model in clinical practice. For instance, in this study, patients in the CA-PPH group, due to coagulation disorders, all underwent cesarean section deliveries to minimize the number of false positives. This requires decision-makers to consider practical issues, as a high ROC does not always indicate the best treatment approach. Furthermore, for some extreme cases, the accuracy of ROC becomes less critical, and DCA evaluation results are needed for reference.

Statistical analysis of previously published PPH data has shown that factors such as general anesthesia in pregnant women, prolonged use of oxytocin, excessive uterine tension (multiple pregnancies, polyhydramnios), and chorioamnionitis are all associated with uterine atony, potentially increasing the risk of postpartum bleeding. Previous studies have suggested that for pregnant women with high-risk factors, assessing and selecting appropriate treatment options and management based on the type and weight of different risks can reduce the risk of adverse pregnancy outcomes or death³⁷. Early prediction and intervention are key measures in reducing maternal mortality, with studies finding that timely interventions can effectively lower maternal mortality rates by 10%³⁸. Establishing a model that predicts the risk of PPH following vaginal delivery and guides clinical practice is a significant task for maternal and child health.

Conclusion

In conclusion, our study has successfully developed and validated predictive models for PPH following vaginal delivery, offering a novel approach to risk assessment in this critical area of maternal health. The models, particularly for overall PPH and UA-PPH, demonstrated high

discriminative power and clinical utility, with the nomogram providing a user-friendly tool for clinicians. Despite the promising results, limitations exist in the application of the PF-PPH, CT-PPH, and CA-PPH models due to the insufficient positive sample size in these subgroups. The generalizability of our findings may also be limited by the single-center nature of the study and the regional characteristics of the included hospitals. Future research should aim to expand the sample size and include multi-center data to improve the models' applicability and robustness. This study contributes to the growing body of evidence on PPH management and has the potential to influence policy and practice, ultimately enhancing maternal care and outcomes.



List of abbreviations

RP: Refined Peyton;

TTM: Traditional Teaching-mode

DOPS: direct observation of procedural skills

OB-GYN: Obstetrics and Gynecology CBVI: computer-based video instruction

PEARLS: Promoting Excellence and Reflective Learning in Simulation

OSCE: objective structured clinical assessment

Declarations

Ethics approval and consent to participate

Ethics approval and consent to participate: The study was approved by the Ethics Committee of Shengjing Hospital of China Medical University (No. 2016PS344K, Date.17/12/2016). All participants provided informed consent.

Consent for publication

Not Applicable

Availability of data and material

The data that support the findings of this study are available from the corresponding author upon reasonable request.

Competing interests

No potential conflict of interest was reported by the author(s).

Funding

This study was supported in part by grants from 345 Talent Project of Shengjing Hospital of China Medical University (No. M0946), and Medical Education Research Project of Liaoning Province (No. 2022-N005-03).

Authors' contributions

Contributors JL and DZ contributed equally to this study. JL, DZ and KZ designed the study and drafted the manuscript. HL, MS, and XW done the data collection. YZ and XC designed the statistical analysis plan. DZ has participated the training and reviewed and co-authored the manuscript with ZS. All authors have read and approved this manuscript. ZS is responsible for the overall content as guarantor.

Acknowledgements

We would like to express our gratitude to all those who helped us during the writing of this manuscript. Thanks to all the peer reviewers for their opinions and suggestions. We would like to acknowledge that Jinke Li and Dandan Zhang have contributed equally to this work.

References:

1. Practice Bulletin No. 183 Summary: Postpartum Hemorrhage. Obstetrics and gynecology

- 2. World Health Organization. Trends in maternal mortality 2000 to 2017: estimates by WHO, UNICEF, UNFPA, World Bank Group and the United Nations Population Division [Available from: https://www.who.int/publications/i/item/9789240012202. accessed 2020 12 22.
- 3. Who UJG, Department of Reproductive Health, Research W. UNFPA: Maternal Mortality in 2000: Estimates developed by WHO, UNICEF and UNFPA. 2004
- 4. AbouZahr C. Global burden of maternal death and disability. *British Medical Bulletin* 2003;67(1):1-11. doi: 10.1093/bmb/ldg015 %J British Medical Bulletin
- Mehrabadi A, Hutcheon JA, Lee L, et al. Trends in postpartum hemorrhage from 2000 to 2009: a population-based study. BMC Pregnancy Childbirth 2012;12:108. doi: 10.1186/1471-2393-12-108 [published Online First: 2012/10/13]
- 6. Reale SC, Easter SR, Xu X, et al. Trends in Postpartum Hemorrhage in the United States From 2010 to 2014. *Anesth Analg* 2020;130(5):e119-e22. doi: 10.1213/ane.0000000000004424 [published Online First: 2019/10/01]
- You JHS, Leung TY. Cost-effectiveness analysis of carbetocin for prevention of postpartum hemorrhage in a low-burden high-resource city of China. *PloS one* 2022;17(12):e0279130. doi: 10.1371/journal.pone.0279130 [published Online First: 2022/12/16]
- 8. AbouZahr CJBmb. Global burden of maternal death and disability. 2003;67(1):1-11.
- 9. Mousa HA, Walkinshaw SJCoiO, Gynecology. Major postpartum haemorrhage. 2001;13(6):595-603.
- 10. Akter S, Forbes G, Vazquez Corona M, et al. Perceptions and experiences of the prevention, detection, and management of postpartum haemorrhage: a qualitative evidence synthesis. The Cochrane database of systematic reviews 2023;11(11):Cd013795. doi: 10.1002/14651858.CD013795.pub2 [published Online First: 2023/11/27]
- 11. Zhang R, Cao X, Feng H, et al. Review of clinical practice guidelines for postpartum hemorrhage according to AGREE II. *Midwifery* 2023;121:103659. doi: 10.1016/j.midw.2023.103659 [published Online First: 2023/03/30]
- 12. Zdanowicz JA, Schneider S, Martignoni C, et al. A Retrospective before and after Assessment of Multidisciplinary Management for Postpartum Hemorrhage. *Journal of clinical medicine* 2023;12(23) doi: 10.3390/jcm12237471 [published Online First: 2023/12/09]
- 13. Giouleka S, Tsakiridis I, Kalogiannidis I, et al. Postpartum Hemorrhage: A Comprehensive Review of Guidelines. *Obstetrical & gynecological survey* 2022;77(11):665-82. doi: 10.1097/ogx.00000000001061 [published Online First: 2022/11/09]
- 14. de Vries PLM, Deneux-Tharaux C, Baud D, et al. Postpartum haemorrhage in high-resource settings: Variations in clinical management and future research directions based on a comparative study of national guidelines. *BJOG : an international journal of obstetrics and gynaecology* 2023;130(13):1639-52. doi: 10.1111/1471-0528.17551 [published Online First: 2023/06/01]
- 15. Klufio CA, Amoa AB, Kariwiga G. Primary postpartum haemorrhage: causes, aetiological risk factors, prevention and management. *Papua and New Guinea medical journal* 1995;38(2):133-49. [published Online First: 1995/06/01]
- 16. Gyamfi-Bannerman C, Srinivas SK, Wright JD, et al. Postpartum hemorrhage outcomes and race.

 **American Journal of Obstetrics and Gynecology 2018;219(2):185.e1-85.e10. doi: https://doi.org/10.1016/j.ajog.2018.04.052
- 17. Desale M, Thinkhamrop J, Lumbiganon P, et al. Ending preventable maternal and newborn deaths

- 18. Sotunsa JO, Adeniyi AA, Imaralu JO, et al. Maternal near-miss and death among women with postpartum haemorrhage: a secondary analysis of the Nigeria Near-miss and Maternal Death Survey. *BJOG*: an international journal of obstetrics and gynaecology 2019;126 Suppl 3:19-25. doi: 10.1111/1471-0528.15624 [published Online First: 2019/03/22]
- Goad L, Rockhill K, Schwarz J, et al. Development and validation of a prediction model for postpartum hemorrhage at a single safety net tertiary care center. *American journal of obstetrics & gynecology MFM* 2021;3(5):100404. doi: 10.1016/j.ajogmf.2021.100404 [published Online First: 2021/05/29]
- 20. Albright CM, Spillane TE, Hughes BL, et al. A Regression Model for Prediction of Cesarean-Associated Blood Transfusion. Am J Perinatol 2019;36(9):879-85. doi: 10.1055/s-0039-1678604 [published Online First: 2019/02/12]
- 21. Liang J, Li X, Kang C, et al. Maternal mortality ratios in 2852 Chinese counties, 1996-2015, and achievement of Millennium Development Goal 5 in China: a subnational analysis of the Global Burden of Disease Study 2016. *Lancet* 2019;393(10168):241-52. doi: 10.1016/s0140-6736(18)31712-4 [published Online First: 2018/12/18]
- 22. Gao Y, Zhou H, Singh NS, et al. Progress and challenges in maternal health in western China: a Countdown to 2015 national case study. *Lancet Glob Health* 2017;5(5):e523-e36. doi: 10.1016/s2214-109x(17)30100-6 [published Online First: 2017/03/28]
- 23. Kleinrouweler CE, Cheong-See FM, Collins GS, et al. Prognostic models in obstetrics: available, but far from applicable. *Am J Obstet Gynecol* 2016;214(1):79-90.e36. doi: 10.1016/j.ajog.2015.06.013 [published Online First: 2015/06/14]
- 24. Tsu VD. Antenatal screening: its use in assessing obstetric risk factors in Zimbabwe. *J Epidemiol Community Health* 1994;48(3):297-305. doi: 10.1136/jech.48.3.297 [published Online First: 1994/06/01]
- 25. Biguzzi E, Franchi F, Ambrogi F, et al. Risk factors for postpartum hemorrhage in a cohort of 6011 Italian women. *Thromb Res* 2012;129(4):e1-7. doi: 10.1016/j.thromres.2011.09.010 [published Online First: 2011/10/25]
- 26. Prata N, Hamza S, Bell S, et al. Inability to predict postpartum hemorrhage: insights from Egyptian intervention data. BMC Pregnancy Childbirth 2011;11:97. doi: 10.1186/1471-2393-11-97 [published Online First: 2011/11/30]
- Rubio-Álvarez A, Molina-Alarcón M, Arias-Arias Á, et al. Development and validation of a predictive model for excessive postpartum blood loss: A retrospective, cohort study. *Int J Nurs Stud* 2018;79:114-21. doi: 10.1016/j.ijnurstu.2017.11.009 [published Online First: 2017/12/11]
- 28. Ahmadzia HK, Phillips JM, James AH, et al. Predicting peripartum blood transfusion in women undergoing cesarean delivery: A risk prediction model. *PloS one* 2018;13(12):e0208417. doi: 10.1371/journal.pone.0208417 [published Online First: 2018/12/15]
- 29. Dunkerton SE, Jeve YB, Walkinshaw N, et al. Predicting Postpartum Hemorrhage (PPH) during Cesarean Delivery Using the Leicester PPH Predict Tool: A Retrospective Cohort Study. *Am J Perinatol* 2018;35(2):163-69. doi: 10.1055/s-0037-1606332 [published Online First: 2017/08/29]
- 30. Yoon SY, You JY, Choi SJ, et al. A combined ultrasound and clinical scoring model for the prediction

- of peripartum complications in pregnancies complicated by placenta previa. *Eur J Obstet Gynecol Reprod Biol* 2014;180:111-5. doi: 10.1016/j.ejogrb.2014.06.030 [published Online First: 2014/08/01]
- 31. Chi Z, Zhang S, Wang Y, et al. Research of the assessable method of postpartum hemorrhage. *Technol Health Care* 2016;24 Suppl 2:S465-9. doi: 10.3233/thc-161169 [published Online First: 2016/05/11]
- 32. !!! INVALID CITATION !!! [73]
- 33. Moher D, Liberati A, Tetzlaff J, et al. Preferred reporting items for systematic reviews and meta-analyses: the PRISMA statement. *PLoS Med* 2009;6(7):e1000097. doi: 10.1371/journal.pmed.1000097 [published Online First: 2009/07/22]
- 34. Steyerberg EW, Vergouwe Y. Towards better clinical prediction models: seven steps for development and an ABCD for validation. *Eur Heart J* 2014;35(29):1925-31. doi: 10.1093/eurheartj/ehu207 [published Online First: 2014/06/06]
- 35. Vickers AJ, Elkin EB. Decision curve analysis: a novel method for evaluating prediction models.

 Med Decis Making 2006;26(6):565-74. doi: 10.1177/0272989x06295361 [published Online First: 2006/11/14]
- 36. Hijazi Z, Oldgren J, Lindbäck J, et al. The novel biomarker-based ABC (age, biomarkers, clinical history)-bleeding risk score for patients with atrial fibrillation: a derivation and validation study. *Lancet* 2016;387(10035):2302-11. doi: 10.1016/s0140-6736(16)00741-8 [published Online First: 2016/04/09]
- 37. Zuckerwise LC, Lipkind HS. Maternal early warning systems-Towards reducing preventable maternal mortality and severe maternal morbidity through improved clinical surveillance and responsiveness. *Semin Perinatol* 2017;41(3):161-65. doi: 10.1053/j.semperi.2017.03.005 [published Online First: 2017/04/19]
- 38. Aoyama K, D'Souza R, Pinto R, et al. Risk prediction models for maternal mortality: A systematic review and meta-analysis. *PloS* one 2018;13(12):e0208563. doi: 10.1371/journal.pone.0208563 [published Online First: 2018/12/05]

Table 1 The general view of the ma			
	Cohort population	Validation population	
Characteristics	(N=24,833)	(N=5009)	P
Age (years) *, N (%)			0.393
<25	1,309 (5.27%)	266 (5.31%)	
25-29	11,736 (47.26%)	2340 (46.71%)	
30-34	9,445 (38.03%)	1959 (39.11%)	
≥35	2,343 (9.44%)	444 (8.86%)	
Ethnicity, N (%)			0.983
Han	22,222 (89.49%)	4,475 (89%)	
Manchu	1,872 (7.54%)	389 (7.8%)	
Other ethnic groups	739 (2.98%)	145 (2.9%)	
Educational Attainment, N (%)			0.115
High school or below	8,635 (34.77%)	1,742 (35%)	
Bachelor's degree	13,639 (54.92%)	2,703 (54%)	
Postgraduate or higher	2,559 (10.30%)	564 (11%)	
Occupation, N (%)			0.777
Unemployed	11,373 (45.80%)	2,266 (45%)	
Light physical labor	2,825 (11.38%)	569 (11%)	
Moderate physical labor	10,011 (40.31%)	2,038 (41%)	
Heavy physical labor	624 (2.51%)	136 (2.7%)	
Family Per Capita Monthly			
Income (10,000 yuan), N (%)			0.9862
<0.5	10,325 (41.58%)	2,080 (42%)	
0.5-2.0	9,534 (38.39%)	1,922 (38%)	
2.0-5.0	3,584 (14.43%)	720 (14%)	
>5.0	1,390 (5.60%)	287 (5.7%)	
Pre-pregnancy BMI (Kg/m2)			
*, N (%)			< 0.001
<18.5 (Underweight)	7,294 (29.37%)	1475 (29.45%)	
18.5-23.9 (Normal)	15,005 (60.42%)	2963 (59.16%)	
24.0-27.9 (Overweight)	1,700 (6.85%)	307 (6.13%)	
>28.0 (Obesity)	834 (3.36%)	264 (5.29%)	
Pregnancy History, N (%)			0.565
1	14,985 (60.34%)	3,005 (60%)	
2	6,513 (26.23%)	1,303 (26%)	
≥3	3,335 (13.43%)	701 (14%)	
Parity (number of deliveries), N			
(%)			0.775
0	20,550 (82.75%)	4,127 (82%)	
1	4,152 (16.72%)	853 (17%)	
≥2	131 (0.53%)	29 (0.6%)	
Gestational Age at Delivery			
(weeks), N (%)			0.434

<38	1,507 (6.07%)	296 (5.91%)	
38-40	13,023 (52.44%)	2589 (51.59%)	
>40	10,303 (41.49%)	2129 (42.50%)	
Blood Loss (ml)	393.54±92.53	413.48±124.65	0.081
Postpartum Hemorrhage, N (%)	1,623 (6.54%)	286 (5.71%)	0.032
Due to uterine atony, N (%)	1,225 (4.93%)	266 (5.31%)	0.279
Due to placental factors, N (%)	242 (0.97%)	43 (0.86%)	0.489
Due to Cancal Trauma, N (%)	139 (0.56%)	31 (0.62%)	0.686
Due to coagulation disorders, N			
(%)	76 (0.31%)	17 (0.34%)	0.705



_		
0		
1		
2 3		
4 5		
6		
7		
8		
9		
0		
1		
2 3		
4		
5		
6 7		
8		
9		
0		
1		
2		
3		
4 5		
5 6		
0		

		BMJ Open		by copyright, including		
Table 2: Multivariate Risk Factor Analysis				for) 5	CA DRW
Characteristics	PPH	UA-PPH	PF-PPH	uses	T-PPH	СА-РРН
A ()	*	General view *		rela		
Age (years)		·	- -	asn ted	₹ - 2	-
Ethnicity	**	**	**	Erasmushogeschoo s related to text and dat	ラ ^本 3	-
Educational Attainment		-	-	ext ext	3 ⁻	-
Occupation	**	**	**	esc	<u> </u>	*
Family Per Capita Monthly Income	<i>-</i> () ₋	<u>-</u>	-	ogeschool . ext and data r	_	-
10,000 yuan)				a - e	3	
Pre-pregnancy BMI (Kg/m2)	**	**	**	mining,	₹ ** }	*
Smoking	-		-		-	*
Alcohol Consumption	-	(0)	-	nining, Al training, and similar technologies.	-	-
	Ob	stetric and Gynecologic	History	aini		
Pregnancy History	-	-	(Q), -	ng,	-	-
Parity (number of deliveries)	**	**	**	and	*	*
History of miscarriage	-	-	-	sin	<u>.</u> -	-
Spontaneous abortion	-	-	-UA	nila	-	-
Induced abortion or medication	_	_	_	tec	2 -	_
abortion				Hing Su	Ę	
Induced labor	-	-	_	plog	3 -	-
Assisted reproductive technology	-	-	-	ies.	<u> </u>	-
Gestational Age at Delivery (weeks)	**	**	*	5	ว ก - ม	-
Diabetes	**	**	**	ָ ק	* *	-
Hypertension	**	**	**	2	*	-
Anemia	**	**	**		**	*
				at Debalulient GEZ-ETA		
				[]	ń N	
				_	<u>-</u> -1	

		·		cop njo	
				njopen: copyriç	
				-202 yht,	
				-2024-089 ght, incluc	
Coagulation disorder	**			njopen-2024-089734 copyright, including	*
Uterine fibroids/adenomyosis	**	**	**	g for	_
Polyhydramnios	**	**	*	20 Ja r uses	_
Umbilical cord entanglement	_	_	_	-	_
Premature rupture of membranes	**	**	**	uar Er	**
Placental abruption	**	**	_	y 20 asm	_
Vaginal bleeding during pregnancy		_	_	ush o te	_
Scarred uterus	**	**	_	Dov Dov	*
scarred attrus	Deliver	y Process and Neonatal (- Conditions	vnlc sch	
Fime of delivery	_ Denver	-	-	ade lool data	_
Time of delivery Total duration of labor	**	*	*	a . id *	**
First stage of labor - Latent phase	**	**	_	* - from h	_
First stage of labor - Active phase	**	**	_	y, http.	_
Second stage of labor	**	**	*	tra **	_
Third stage of labor	**	_ //	**	· * * * * * http://bmjopen.bmj.com/ , Al training, and similar	_
Placental retention/Placenta				g <mark>pe</mark> n	
accreta/Placental implantation	**	**	**	and s	**
Analgesia during labor	**	**	-0.	simi. *	**
Instrumental assistance in delivery	**	_		lar t	_
Lacerations of the cervix, vagina, or				on J	
perineum	**	-	-	June	-
Newborn weight (grams)	**	**	-	ر بق	-
Newborn length (centimeters)	-	-	-	2025 ; es.	-
*: P<0.05; **: P<0.001				THE DO	
				y par	
				partment GEZ-LTA	
				nt C	
				ξEZ	
				ή	

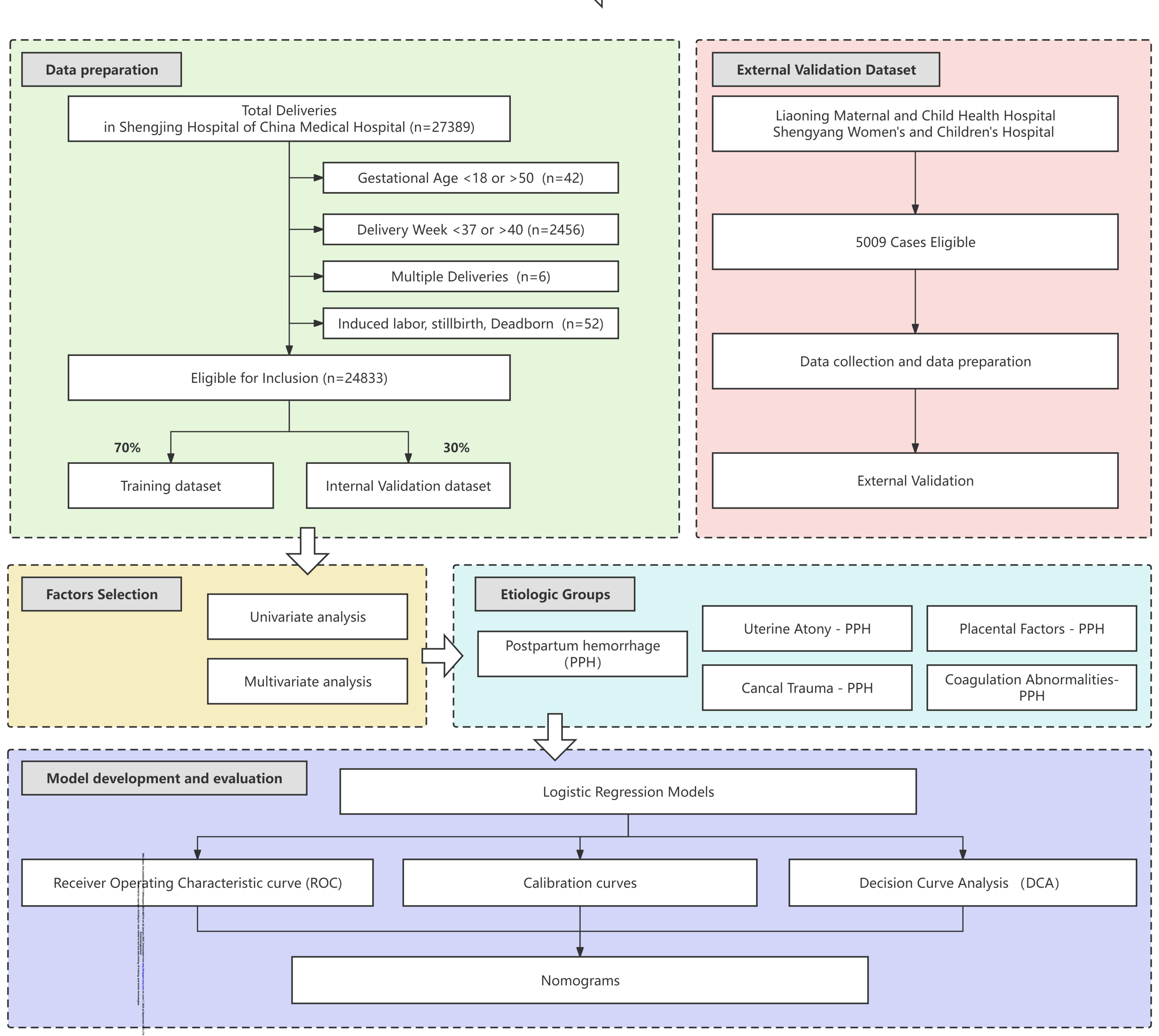
Page 17 of 68

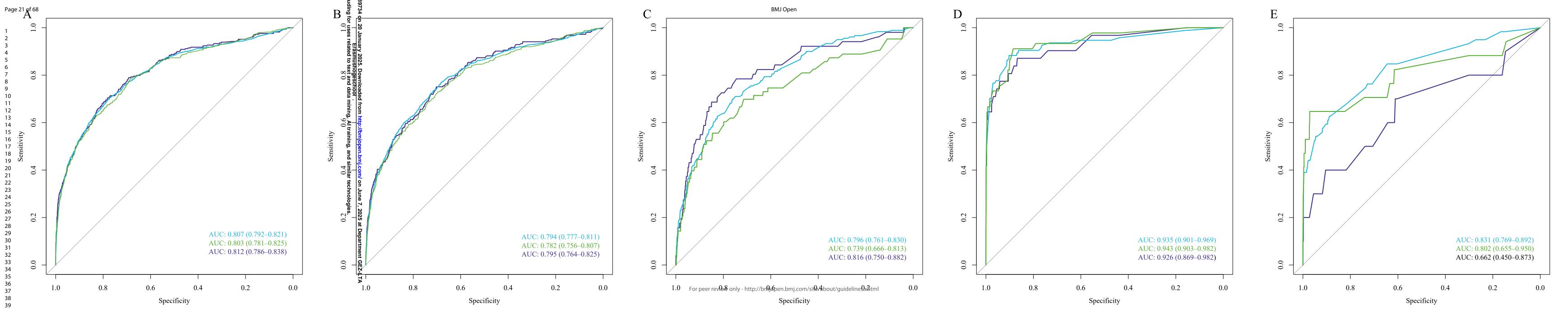
d by copyright, including for 36/bmjopen-2024-089734 on

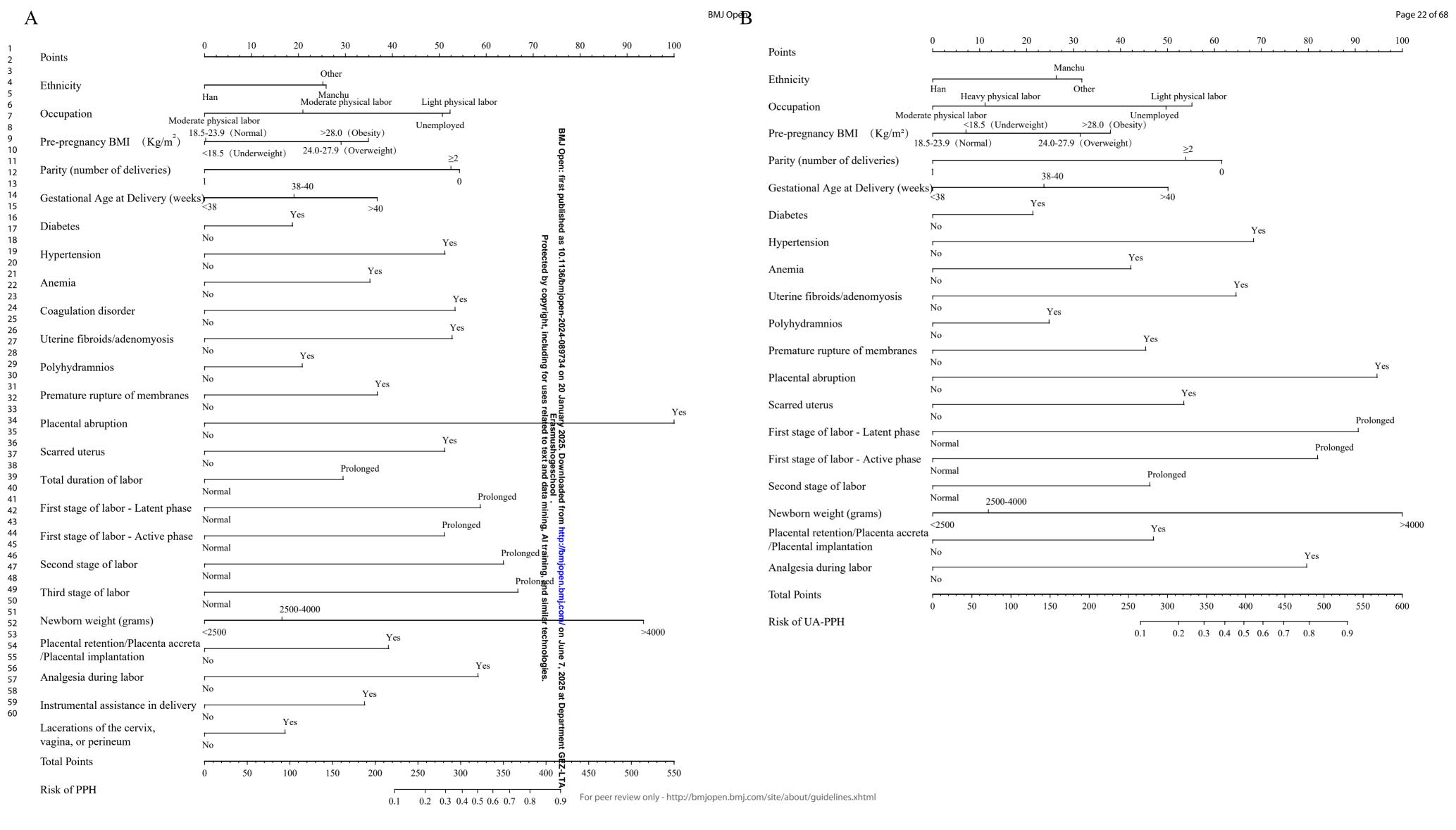
Figure legend

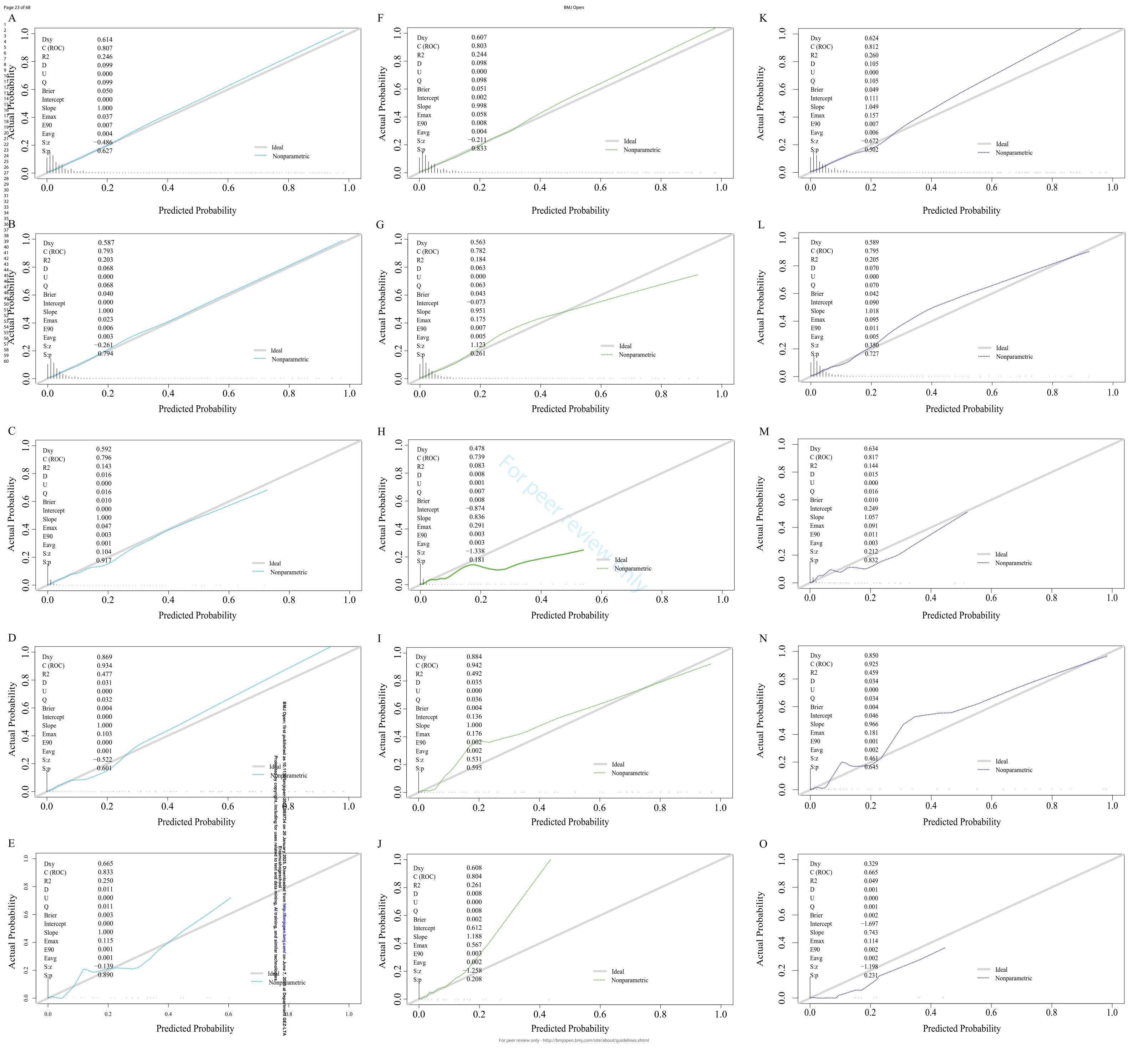
- Figure 1. Patient Selection Criteria Flowchart.
- Figure 2. AUC Curve for Postpartum Hemorrhage (PPH) Group and four subgroups. A. PPH Group; B. Uterine Atopy PPH Group; C. Placental Factors PPH Group; D. Cervical Trauma PPH Group; E. Coagulation Abnormalities PPH Group. The blue line signifies the training dataset, which is employed to evaluate the model's predictive capabilities following the training phases. The green line corresponds to the internal validation data votal for refining model parameters and for conducting initial assessments of the model's accuracy. The purple line denotes the external validation datase which is utilized to ascertain the model's generalizability and to verify its performance in an independent dataset.
- Figure 3. Nomograms for Postpartum Hemorrhage (PPH) and Uterine Atony PPH Group. A. PPH Group; B. Uterang Atony PPH Group.
- Supplemental Figure 1. Calibration curves for Postpartum Hemorrhage (PPH) Group and four subgroups. A PH Group; B\G\L. Uterine Atony PPH dataset; The green line signifies the internal validation dataset; The purple line signifies the external validation dataset.
- Supplemental Figure 2. Decision Curve Analysis for Postpartum Hemorrhage (PPH) Group and four subgroups. E PH Group; B. Uterine Atony PPH Group; C. Placental Factors PPH Group; D. Cervical Trauma PPH Group; E. Coagulation Abnormalities PPH Group. The blu in signifies the training dataset; The green training, and similar technologies line signifies the internal validation dataset; The purple line signifies the external validation dataset. /bmjopen.bmj.com/ on June 7, 2025 at Department GEZ-LTA

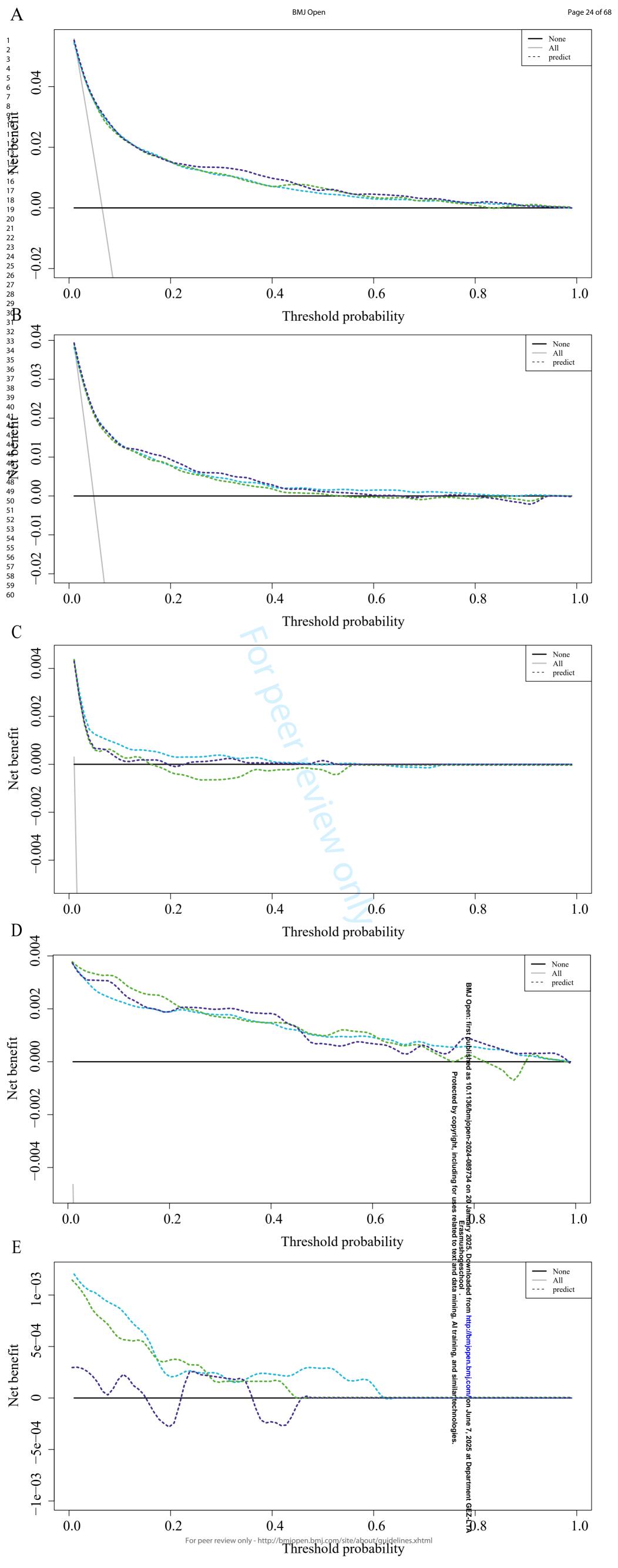












							BMJ Open	ı		√bmjo _l		
Supplementary	table 1	Compar	rison of l	Basic Cha	racterist	ics and Ris	k Factor Scr	eening between	Non-PPH an	56/bmjopen-2024-089734 By copyright, includia		
		-PPH		PH						on for		
Characteristi	(N=2	3,210)	(N=	1,623)			Univa	riate Logistic R	egression	uses Multiv	ariate Logistic l	Regression
Characteristi			Sam		χ^2	P				_		
cs	Sampl		ple							nuary 2 Erasr related		
	e size	n%	size	n%			OR	95% CI	P	ed to 15	95% CI	P
General view										<u>8</u> . → .		
						<0.001*				Downloaded logeschool . Ref.		
Age (years)					77.07	*				hoader Ref.		
<25	1,251	5.39%	58	3.57%			Ref.			Ref.		
25-29	11,047	47.60%	689	42.45%			1.35	1.03, 1.79	0.034*	ning, 1.45	0.85, 1.49	0.430
30-34	8,719	37.57%	726	44.73%			1.80	1.38, 2.39	<0.001**	ر ق 1. 4 5	1.11, 1.94	0.009*
≥35	2,193	9.45%	150	9.24%			1.48	1.09, 2.03	0.014*	1.25	0.92, 1.72	0.169
						<0.001*				omj ainii		
Ethnicity					86.05	*				ng,		
Han	20,848	89.82%	1,374	84.66%			Ref.		•	Ref.		
Manchu	1,693	7.29%	179	11.03%			1.60	1.36, 1.88	<0.001**	<u>s</u> . 1. 5 9	1.34, 1.87	<0.001**
Other ethnic		• 0001	= ^	4.0107			1.50	1.00 - 22	0.00544	om/	101-0-	.0.00
groups	669	2.88%	70	4.31%		.0.00**	1.59	1.22, 2.03	<0.001**	fcomontation (bmjopen.brg).com/on June 7, 20	1.24, 2.07	<0.001**
Educational					20.46	<0.001*	D. C			Jun _c		
Attainment					39.46		Ref.			Ref.		
High school	0 1 4 2	25.000/	402	20.200/			1.20	1 00 1 25	0.001*	202	0.04 1.07	0.202
or below Bachelor's	8,142	35.08%	493	30.38%			1.20	1.08, 1.35	0.001*	,2025 at Department Gi	0.84, 1.07	0.392
degree	12,713	54.77%	926	57.05%			1.43	1.21, 1.69	<0.001**	D ee 1 16 6	0.88, 1.27	0.520
Postgraduate	2,355	10.15%	204	12.57%			1.43	1.41, 1.09	~0.001	. 	0.00, 1.27	0.320

or higher										'34 ing		
						<0.001*				on		
Occupation					359.52	*				e R R)89734 on 20 January 2625. Downloaded from http://bmjopen.bmj.com/ on June 7, 20 Erasmushogeschoo⊅. luding for uses related to text and data mining, Al training, and similar technologies.		
Unemployed	10,441	44.98%	932	57.42%			Ref.			Ref.		
Light physical										ary Eras		
labor	2,573	11.09%	252	15.53%			1.10	0.95, 1.27	0.212	od to	0.92, 1.24	0.363
Moderate										5. D Isho		
physical labor	9,591	41.32%	420	25.88%			0.49	0.44, 0.55	<0.001**	t ge. an	0.44, 0.56	<0.001**
Heavy										nloa icho id di		
physical labor	605	2.61%	19	1.17%			0.35	0.21, 0.54	<0.001**	ata ⊖.663	0.20, 0.51	<0.001**
Family Per Ca	pita Mon	thly								min		
Income (10,000) yuan)				2.98	0.887				ing,		
< 0.5	9,657	41.61%	668	41.16%				-	-	<u> ≱</u>	-	-
0.5-2.0	8,898	38.34%	636	39.19%					-	.rain	-	-
2.0-5.0	3,362	14.49%	222	13.68%			_		-	ing -	-	-
>5.0	1,293	5.57%	97	5.98%			-		-	en.t	-	-
Pre-pregnancy	BMI					<0.001*				d si.		
(Kg/m^2)					299.11	*				con mila		
<18.5										ar te		
(Underweig										chn L		
ht)	6,869	29.60%	425	26.19%			Ref.			Ref.		
18.5-23.9										7, 20 gies		
(Normal)	14,116	60.82%	889	54.78%			1.02	0.90, 1.15	0.771	1 A> >	0.90, 1.15	0.803
24.0-27.9										2. Thent		
Overweight										ера		
)	1,500	6.46%	200	12.32%			2.15	1.80, 2.57	<0.001**	2. ₫ 6	1.81, 2.58	<0.001**

Page 27 of 68

2 3 4

5 6

7

8

9 10

11

12

13

14

15

16

17

18 19

20

21

22 23

24

25

26 27

28

29

30

31 32

33

34

35 36

37

Yes 2, induced abortion or medication abortion	20,874 2,336	89.94% 10.06%	1,465 158	90.26% 9.74%						6/bmjopen-2024-089734 on 20 January 202: Erasmu by copyright, including for uses related to			_
No 20 Yes 2, induced abortion or medication abortion							_			34 ng			
Yes 2, induced abortion or medication abortion							_			on .			
induced abortion or medication abortion No 18	2,336	10.06%	158	9.74%			-	-	-	n 20 or u:	-	-	
abortion or medication abortion No 18							-	-	-	20 January 2025. Erasmus	-	-	
medication abortion No 18										າuar Er rela			
abortion No 18										y 20 asm ted t			
No 18					1.02	0.797				· S 5.			
	8,688	80.52%	1,295	79.79%	/	0.171	-	_	-	Downloaded from logeschool .	_	_	
168 4,	4,522	19.48%	328	20.21%			-	-	-	· · · wnloa Jescho and d	-	_	
induced						<0.001*				adec pol lata			
labor					26.61	*				y fro			
No 22	22,717	97.88%	1,609	99.14%			Ref.						
Yes 4	493	2.12%	14	0.86%			0.40	0.22, 0.66	<0.001**	≥ 0. ₹ 2	0.44, 1.40	0.489	
assisted										rain //bm			
reproductive										jope ing,			
technology					3.67	0.299				http://bmjopen.bmj.com/ on June 7, 2025 g, Al training, and similar technologies.			
	22,595	97.35%	1,589	97.91%			-	-	-	sin -	-	-	
	615	2.65%	34	2.09%			-	-	UA	om/	-	-	
Gestational										tec			
Age at						-0 001*				Jun			
Delivery (weeks)					241.48	<0.001* *				e 7,			
	1,441	6.21%	66	4.07%	241.40		Ref.			2025 Ref			
		53.21%		41.53%			1.19	0.93, 1.56	0.184		1.17, 1.99	0.002*	
		40.59%		54.41%			2.05	1.60, 2.67	<0.001**	2.62	2.26, 3.84	<0.001**	
Diabetes	·,	• • • • • • • • • • • • • • • • • • • •	~ -		126.81	<0.001*		2000,	•••	ak-Desiartment GEZ-LTA	_·_·, - · ·	****	

entanglement										jopen-2024-089734 . opyright, including		
No	16,254	70.03%	1,143	70.43%			_	-	_	on for	_	_
Yes	6,956	29.97%	480	29.57%			-	-	-	on 20 January Era for uses relate	-	_
Premature	•									Janu Ss re		
rupture of						<0.001*				uary Era:		
membranes					336.49	*				2025, smus		
No	18,247	78.62%	1,051	64.76%			Ref.			ma ⊐ Ret		
Yes	4,963	21.38%	572	35.24%			2.00	1.80, 2.22	<0.001**	Downloaded	1.92, 2.41	<0.001**
Placental						<0.001*				nloa icho id di		
abruption					150.97	*				ided ool .		
No	23,180	99.87%	1,605	98.89%			Ref.			mining, 8.1		
Yes	30	0.13%	18	1.11%			8.67	4.73, 15.4	<0.001**	4.28, 15.30	<0.001**
Vaginal										≽i t		
bleeding										/bm		
during										rttp://bmjopen.bmj.com/ on June 7,50 Al training, and similar technologies.		
pregnancy					0.73	0.867				en.b		
No	22,030	94.92%	1,546	95.26%			-	-	-	d sir	-	-
Yes	1,180	5.08%	77	4.74%			-	-	OA	nila -	-	-
Scarred						<0.001*				r te		
uterus					21.53	*				chn Ju		
No	22,980	99.01%	1,593	98.15%			Ref.					
Yes	230	0.99%	30	1.85%			1.88	1.26, 2.72	<0.001*	ie 3.05	1.98, 4.53	<0.001**
Delivery Proce	ess and N	eonatal C	onditior	ıs						7.53025 at Departm 3.79025 at Departm		
Time of										∓ De		
delivery					0.13	0.989				y par		
Day shift	9,722	41.89%	685	42.21%			-	-	-	tment GEZ-LTA	-	-

 Page 30 of 68

Page 31 of 68

2 3 4

5 6

7

8

9 10

11

12

13

14

15

16

17

18 19

20

21

22 23

24

25

26 27

28

29

30

31 32

33

34

35

36

37

2 3 4

5 6

8

9 10

11

12

13

14 15

16

17

18 19

20

21

22 23

24

25

26

27

28

29

30

31 32

33

34

35 36

37

38

44 45 46 Page 32 of 68

							BMJ Oper	า		36/bmjope		
										e R 86/bmjopen-2024-0897∰ on 20 January 2025∰Oo∰nloaded from http://bmjopen.bmj.com/ on June 7, 202 Erasmushogeschool . ⊖ 9 by copyright, including for uses related to text and data mining, Al training, and similar technologies.		
Yes	2,071	8.92%	225	13.86%			1.64	1.41, 1.90	<0.001**	7.23 ng 1.23	1.26, 1.74	<0.001**
Newborn										n 20 or u		
weight					1510.0	<0.001*) Ja ses		
(grams)					2	*				nua rela		
<2500	221	0.95%	7	0.43%			Ref.			ited Ref.		
2500-40		:							0.44-	025 nus to 1		0.40-
00	21,944	94.55%	1,281	78.93%			1.84	0.94, 4.34	0.112	ext of	0.84, 3.94	0.187
>4000	1,045	4.50%	335	20.64%			10.1	5.09, 24.0	<0.001**	Masc and	4.68, 22.40	<0.001**
Newborn										oad hoo I da		
length						<0.001*				ia m		
(centimeters)	22.555	05.000	1 7 15	05.0107	52.57		F 0			inin		
≤ 55	22,566	97.23%	1,542	95.01%			Ref.		0.004**	Kef.	0.62.4.00	
>55	644	2.77%	81	4.99%			1.84	1.44, 2.32	<0.001**	∠ 0. 5 3	0.63, 1.08	0.172
										ain j		
										jopen.bmj.com/ on June 7, 2025 at Department GEZ-LTA ing, and similar technologies.		
										nd s		
										j.co simi		
										m/ c		
										on J		
										une		
										, 7, <i>1</i>		
										2025 s.		
										at		
										Dep		
										oartr		
										nen		
										t <u>G</u> E		
										:Z-L		
				For peer	review or	nly - http://l	omjopen.bmj	.com/site/about/	guidelines.xh	tml		

							BMJ Open	eening between		:6/bmjopen-202 I by copyright,		
Supplementary	Non-U	A-PPH	UA	-PPH	racterist	ics and Ris				익 ㅋ		
Characteristi	(N=2	3,608)	Sam	1,225)	χ^2	P	Univa	riate Logistic R	egression	ses r	ariate Logistic	Regression
cs	Sampl e size	n%	ple size	n%	λ.	•	OR	95% CI	P	ulu R M OR R Erasmushogeschool . 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	95% CI	P
General view										5. D		
Age (years)					67.01	<0.001* *				ownloa gescho t and da		
<25	1,269	5.38%	40	3.27%			Ref.			at of the Ref.		
25-29	11,212	47.49%	524	42.78%			1.48	1.08, 2.09	0.018*	<u> 1.</u> 2	0.88, 1.71	0.266
30-34	8,891	37.66%	554	45.22%			1.98	1.45, 2.78	<0.001**	g 1.35	1.13, 2.19	0.009*
≥35	2,236	9.47%	107	8.73%		<0.001*	1.52	1.06, 2.22	0.027*	f. Ref. Ref. l.	0.86, 1.82	0.259
Ethnicity					39.45	*				ing.		
Han	21,172	89.68%	1,050	85.71%			Ref.			Ref.		
Manchu Other ethnic	1,749	7.41%	123	10.04%			1.42	1.16, 1.71	<0.001**	und 11.4	1.15, 1.70	<0.001**
groups	687	2.91%	52	4.24%			1.53	1.13, 2.02	0.004*	1.55	1.15, 2.05	0.003*
Educational						<0.001*				chn Ju		
Attainment					54.87	*				ne 7 olog		
High school										', 20 jies.		
or below	8,278	35.06%	357	29.14%			Ref.			Ref.		
Bachelor's										at De		
degree	12,939	54.81%	700	57.14%			1.25	1.10, 1.43	<0.001**	0. 29	0.86, 1.13	0.831
Postgraduate	2,391	10.13%	168	13.71%			1.63	1.35, 1.96	<0.001**	1. 23 0	0.98, 1.47	0.075

Page 35 of 68

2 3 4

5 6

7

8

9 10

11

12

13

14

15

16

17

18 19

20

21

22 23

24

25

26 27

28

29

30

31 32

33

34

35 36

37

38

										open-2024-089734 on 20 Jai		
										1-0897 nclud		
(Normal)										734 c		
24.0-27.9										on 20 for us		
Overweight										0 Ja		
)	1,556	6.59%	144	11.76%			1.86	1.52, 2.28	<0.001**) Ja&uary ⊡Eras ses relate	1.51, 2.27	<0.001**
>28.0										©uary 2 ⊣ Eras⊩ related		
(Obesity)	764	3.24%	70	5.71%			1.85	1.40, 2.40	<0.001**	2025. Smisis	1.39, 2.39	<0.001**
Smoking					2.68	0.444				w → .		
No	23,522	99.64%	1,218	99.43%			-	-	-	- bwnl gesc	-	-
Yes	86	0.36%	7	0.57%			-	-	-	Downloaded	-	-
Alcohol										ded ol . ita n		
Consumption					0.66	0.883				from http://bmjopen.bmj.		
No	23,550	99.75%	1,223	99.84%			<u></u>	-	-	ng, <mark>m</mark> -	-	-
Yes	58	0.25%	2	0.16%			10.	-	-	Al t	-	-
Obstetric and	Gynecolo	gic Histor	ry							/bm rain		
Pregnancy						<0.001*				ing,		
History					33.29	*				en.b		
1	14,179	60.06%	806	65.80%			Ref.			Ref.		
2	6,244	26.45%	269	21.96%			0.76	0.66, 0.87	<0.001**	<u>≅</u> . 0. § 8	0.84, 1.14	0.830
≥3	3,185	13.49%	150	12.24%			0.83	0.69, 0.99	0.039*	imilar technolo	0.95, 1.42	0.131
Parity										chn		
(number of						<0.001*				f. Ref I from http://bmjopen.bmj.com O mining, Al training, and similar technologies		
deliveries)					138.11	*				7, 20 ogies		
0	19,431	82.31%	1,119	91.35%			Ref.			Ref.		
1	4,053	17.17%	99	8.08%			0.42	0.34, 0.52	<0.001**	0. 월	0.32, 0.53	<0.001**
≥2	124	0.53%	7	0.57%			0.98	0.41, 1.95	0.959	0. 2 6	0.31, 1.61	0.513
<i>></i> 2					0.08	0.994				artment GEZ-LTA		

 Page 36 of 68

Page 37 of 68

2 3 4

5 6

8

9 10

11

12

13

14 15

16

17

18 19

20

21

22 23

24

25

26 27

28

29

30

31

32

33

34

35 36

37

										-089 734 Ref.		
<38	1,458	6.18%	49	4.00%			Ref.			Ref.		
38-40	12,518	53.02%	505	41.22%			1.20	0.90, 1.64	0.230	학 1. 월 1	1.12, 2.06	0.008*
>40	9,632	40.80%	671	54.78%			2.07	1.56, 2.82	<0.001**	us 2.90	2.16, 3.97	<0.001**
						<0.001*				anu s re		
Diabetes					96.97	*				f. Re Re 26-January 2025-Downloade 2 Erasmus h ogeschool uses related to text and data		
No	20,527	86.95%	980	80.00%			Ref.			To a Ref.		
Yes	3,081	13.05%	245	20.00%			1.67	1.44, 1.92	<0.001**	tex 50	1.38, 1.86	<0.001**
						<0.001*				owi ges		
hypertension					256.89	*				nloa ichc		
No	21,998	93.18%	1,036	84.57%			Ref.			ef. Ref. 26 January 2025 Downloaded 2 Erasmushogeschool . uses related to text and data r		
Yes	1,610	6.82%	189	15.43%			2.49	2.11, 2.93	<0.001**	m: 2. ₹ 7	2.33, 3.28	<0.001**
						<0.001*				m h		
anemia					150.71	*				Al 1		
No	19,667	83.31%	903	73.71%			Ref.			ef. Ref. Ref. Paded Fom http://bmj@pen.bmj.com/ on June 7, 20 1001 . 2. data mining, Al training, and similar technologies		
Yes	3,941	16.69%	322	26.29%			1.78	1.56, 2.03	<0.001**	ing 1.89	1.64, 2.16	<0.001**
coagulation										en.t		
disorder					0.24	0.972				d si		
No	23,454	99.35%	1,218	99.43%			-	-		milia -	-	-
Yes	154	0.65%	7	0.57%			-	-	- /-)/	ar te	-	-
uterine										1 June		
fibroids/aden						<0.001*				ne do		
omyosis					205.10	*				7, 20 ogies		
No	22,966	97.28%	1,130	92.24%			Ref.			. % Ref.		
Yes	642	2.72%	95	7.76%			3.01	2.39, 3.74	<0.001**	3. 24	2.40, 3.82	<0.001**
polyhydramn						<0.001*				a∰Departmo		
ios					50.52	*				ir th		

5

6

8

9 10

11

12

13 14

15

16

17

18 19

20

21

22 23

24

25

26 27

28

29

30

31

32

33

34

35 36

37

38

2 3 4

5 6

8

9 10

11

12

13

14

15

16

17

18 19

20

21

22 23

24

25

26

27

28

29

30

31 32

33

34

35 36

37

44 45 46 Page 40 of 68

Page 41 of 68

2 3 4

5 6

8

9 10

11

12

13

14

15

16

17

18 19

20

21

22 23

24

25

26 27

28

29

30

31 32

33

34

35 36

37

2 3 4

5 6

7

8

9 10

11

12

13

14

15

16

17

18 19

20

21

22 23

24

25

44 45 46 Page 42 of 68

							BMJ Open	ening between		6/bmjopen-2024		
	Non-P	Compar F-PPH 4,591)	PF	Basic Cha -PPH =242)	racterist	ics and Risk		ening between		익 크	roups ariate Logistic R	egression
Characteristi cs	Sampl	,	Sam ple	_	χ^2	P		J	n	January 2 Eras ss relatec	C	J
General view	e size	n%	size	n%			OR	95% CI	P	to OR to Sh	95% CI	P
Age (years)				11.46	0.120					Dow oges ar		
<25	1,297	5.27%	12	4.96%	A		-	-	_	nlos scho	-	_
25-29	11,639	47.33%	97	40.08%			-	-	-	ided ool ata	-	-
30-34	9,340	37.98%	105	43.39%			_	-	-	min -	-	-
≥35	2,315	9.41%	28	11.57%			<u> </u>	-	-	ing, -	-	-
						<0.001*				≱ <mark>tt</mark>		
Ethnicity					29.18	*				/bm		
Han	22,023	89.56%	199	82.23%			Ref.			Ref.		
Manchu	1,839	7.48%	33	13.64%			1.99	1.35, 2.84	<0.001**	an	1.32, 2.78	<0.001*
Other ethnic groups	729	2.96%	10	4.13%			1.52	0.75, 2.73	0.201	d simila 1.55	0.76, 2.78	0.181
Educational					3.44	0.632				r tec		
Attainment					J. 44	0.032				- Jur		
High school or below	8,559	34.81%	76	31.40%			-	-	<u>-</u>	ne 7, 20 blogies	-	-
Bachelor's degree	13,496	54.88%	143	59.09%			-	-	-	125 at D	-	-
Postgraduate or higher	2,536	10.31%	23	9.50%			-	_	-	at Departm	_	_

					77.00	<0.001*			(9734 on 20 Ja		
Occupation					77.00	*				for On :		
Unemployed	11,218	45.62%	155	64.05%			Ref.			Ref.		
Light physical labor	2,796	11.37%	29	11.98%			0.75	0.49, 1.10	0.159	Ref. Ref. 0.74 Ref. 0.23 Erasmushogeschool or uses related to text and data	0.49, 1.08	0.137
Moderate physical labor	9,955	40.48%	56	23.14%			0.41	0.30, 0.55	<0.001**	2025. C smushc	0.30, 0.55	<0.001*
Heavy physical labor	622	2.53%	2	0.83%			0.23	0.04, 0.73	0.041*	2 Downloaded from http://bmjopen.bmj.com/ on June logeschool .	0.04, 0.71	0.038*
Family Per										adec ool ata		
Capita										min of the		
Monthly					2.01	0.959			(ing T		
Income									·	≥ 		
(10,000 yuan)										maii traii		
< 0.5	10,222	41.57%	103	42.56%			-	/O:	- ,	ning -	-	-
0.5-2.0	9,440	38.39%	94	38.84%			-		-	, an	-	-
2.0-5.0	3,549	14.43%	35	14.46%			-	-	_	d si	-	-
>5.0	1,380	5.61%	10	4.13%			-	-			-	-
Pre-pregnan cy BMI					64.09	<0.001*				oaded from http://bmjopen.bmj.com/ on June 7, 20 hool . data mining, Al training, and similar technologies		
(Kg/m2)					04.09	*						
<18.5										ğ 7		
(Underweig	7,235	29.42%	59	24.38%			Ref.			es. 2025 Ref.		
ht)	1,233	∠J.¬∠/0	3)	27. 30/0			ICI.			<u>a</u>		
18.5-23.9	14,873	60.48%	132	54.55%			1.09	0.80, 1.49	0.590	at Departmen	0.81, 1.50	0.578
(Normal)										<u>f</u>		

Page 45 of 68

2 3 4

5

6

7 8

9

10

11

12

13

14 15

16

17

18 19

20

21

22 23

24

25

26

27

28

29

30

31 32

33

34

35 36

37

							BMJ Open			, , , 6/bmjopen-2024-089734 on 20 Ja by copyright, including for uses			Page 46 of
										4-0897; ncludii			_
No	18,087	73.55%	180	74.38%			-	-	-	- 34 o ng f	-	-	
Yes	6,504	26.45%	62	25.62%			-	-	-	on 20	-	-	
spontaneous										D Ja			
abortion		20.060/		00.5=01	0.04	0.998				nua rela			
No	22,122	89.96%	217	89.67%			-	-	-	nuary 2029 Erasmu related to	-	-	
Yes	2,469	10.04%	25	10.33%			-	-	-	2025, Smus	-	-	
induced										5. Do shog text			
abortion or										ry 2025. Downloaded rasmushogeschool . ated to text and data r			
medication						2265				oad hoo I dat			
abortion	10.706	00.460/	107	01 400/	0.27	0.965				ta m			
No	19,786	80.46%	197	81.40%			-	-	-	inin	-	-	
Yes	4,805	19.54%	45	18.60%			h-	-	-	n htt	-	-	
induced					C 40	2 200				7 tr			
labor	24.005	27.040/	0.41	00.700/	6.48	0.090				ainii			
No	24,085	97.94%	241	99.59%			- 1	10 ;	-	ng,	-	-	
Yes	506	2.06%	1	0.41%			-	W/	-	n.br	-	-	
assisted										sim .c			
reproductive					2.24	0.504				om/ nilar			
technology	22.051	0 7 400/	222	26.2007	2.34	0.504				wnloaded from http://bmjopen.bmj.com/ on June 7, 20 eschool . and data mining, Al training, and similar technologies.			
No	23,951	97.40%	233	96.28%			-	-	- 4	Jun -	-	-	
Yes	640	2.60%	9	3.72%			-	-	-	le 7,	-	-	
Gestational										2025 ies.			
Age at						-0.001*				:5 a			
Delivery					26.60	<0.001*				5 at Department GEZ-L1			
(weeks)	1 400	C 100/	0	2 210/	26.69		D 0			par			
<38	1,499	6.10%	8	3.31%			Ref.			🗲 Ref.			

Page 47 of 68

2 3 4

5

6

7

8

9 10

11

12

13

14 15

16

17

18 19

20

21

22 23

24

25

26 27

28

29

30

31 32

33

34

35 36

37

38

2 3 4

5 6

7

8

9 10

11

12

13

14 15

16

17

18 19

20

21

22 23

24

25

26 27

28

29

30

31

32

33

34

35 36

37

44 45 46 Page 48 of 68

Page 49 of 68

2 3 4

5 6

7

8

9 10

11

12

13

14 15

16

17

18 19

20

21

22 23

24

25

26 27

28

29

30

31 32

33

34

35 36

37

38

2 3 4

5 6

7

8

9 10

11

12

13

14 15

16

17

18 19

20

21

22 23

24

25

26 27

28

29

30

31 32

33

34

35 36

37

44 45 46 Page 50 of 68

≤55

>55

23,875

716

97.09%

2.91%

233

9

96.28%

3.72%

16

17

18 19

20

21

22 23

24

44 45 46 0.27, 5.39

1.21, 24.90

0.860

0.059

BMJ Open

							BMJ Open	eening between		6/bmjopen-2024-(by copyright, inc		
Supplementary	Non-C	Compar T-PPH 4,694)	CI	Basic Cha T-PPH =139)	racterist	ics and Ris		eening between		우 ㅋ	PH groups	Regression
Characteristi cs	Sampl e size	n%	Sam ple size	n%	χ^2	P	OR	95% CI	P P	M OR Ref Ref Properties of the Ref Ref Ref Ref Ref Ref Ref Ref Ref Re	_	P.
General view										usho tex		
Age (years)				<i>\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\</i>	11.23	0.129				æges ges an		
<25	1,303	5.28%	6	4.32%			-	-	-	nloa icho id di	-	-
25-29	11,682	47.31%	54	38.85%			-	-	-	ided ol . ata -	-	-
30-34	9,379	37.98%	66	47.48%			-	-	-	mini	-	-
≥35	2,330	9.44%	13	9.35%			<u> </u>	-	-	ing, -	-	-
						<0.001*				Al t		
Ethnicity					24.67	*				/bm rain		
Han	22,109	89.53%	113	81.29%			Ref.			Ref	•	
Manchu	1,856	7.52%	16	11.51%			1.69	0.96, 2.77	0.051	a 1.67	0.95, 2.75	0.057
Other ethnic										y sir		
groups	729	2.95%	10	7.19%			2.68	1.31, 4.89	0.003*	<u>a</u> 2. § 1	1.32, 4.95	0.003*
Educational										· tec		
Attainment					4.25	0.515				ef Per paded from http://bmjopeஞ்bmj.cn/ on June 7, 20 hool . data mining, Al training, and similar technologies.		
High school										,7 gi		
or below	8,585	34.77%	50	35.97%			-	-	-	2025 ies.	-	-
Bachelor's	10.566	54050/	5 0	50.050/								
degree	13,569	54.95%	70	50.36%			-	-	-	: De	-	-
Postgraduate	2.540	10.2007	10	12 (70)						at Departm		
or higher	2,540	10.29%	19	13.67%			-	-	-	:ment GE	-	-

Page 53 of 68

2 3 4

5 6

7

8

9 10

11

12

13

14

15

16

17

18 19

20

21

22 23

24

25

26 27

28

29

30

31 32

33

34

35 36

37

Overweight										jopen-2024-089734 on : opyright, including for		
)										on 20 for us		
>28.0										20 Ja		
(Obesity)	810	3.28%	24	17.27%			10.80	5.93, 19.82	<0.001**	a 10 2 /0	5.91, 19.71	<0.001*
Smoking					1.05	0.789				£ 8j ~		
				100.00						2025 smus		
No	24,601	99.62%	139	%			-	-	-	hog text	-	-
Yes	93	0.38%	0	0.00%			-	-	-	- wnl Jesc and	-	-
Alcohol					0.50	0.050				oad hoo dat		
Consumption				100.00	0.68	0.879				a m ed f		
N	24.624	00.760/	120	100.00						d from http://bmjopen.bmj.com/ on June mining, Al training, and similar technolo		
No	24,634	99.76%	139	%			/ _	-	-	g, A	-	-
Yes	60 Cymanala	0.24%	0	0.00%			(5)	-	-	p://k	-	-
Obstetric and	Gynecolo	gic Histor	y									
Pregnancy History					8.22	0.145				ng, a		
History 1	14,892	60.31%	93	66.91%	0.22	0.143				ind by		
2	6,487	26.27%	26	18.71%			-	-		nj.cc	-	_
≥ ≥3	3,315	13.42%	20	14.39%			_	-		om/	-	_
Parity	3,313	13.42/0	20	14.57/0			_	_	~ 7//	on J	_	_
(number of										lune		
deliveries)					18.31	0.003*				7,		
0	20,422	82.70%	128	92.09%	10.51	0.000	Ref.			2025 Ref.		
1	4,142	16.77%	10	7.19%			0.39	0.19, 0.70	0.004*	0.40	0.20, 0.73	0.006*
≥2				0.72%			1.23		0.839	1. 2 5	0.07, 5.70	
history of					0.38	0.945		,		artment GEZ-LTA	,	

 Page 54 of 68

Page 55 of 68

2 3 4

5 6

8

9 10

11

12

13

14 15

16

17

18 19

20

21

22 23

24

25

26 27

28

29

30

31

32

33

34

35 36

37

2 3 4

5 6

7

8

9 10

11

12

13

14 15

16

17

18 19

20

21

22 23

24

25

26 27

28

29

30

31 32

33

34

35 36

37

38

44 45 46 Page 56 of 68

Page 57 of 68

2 3 4

5 6

7

8

9 10

11

12

13

14 15

16

17

18 19

20

21

22 23

24

25

26

27

28

29

30

31 32

33

34

35 36

37

2 3 4

5 6

8

9 10

11

12

13

14

15

16

17

18 19

20

21

22 23

24

25

26

27

28

29

30

31 32

33

34

35 36

37

38

44 45 46 Page 58 of 68

Page 59 of 68

2 3 4

5 6

7

8

9 10

11

12

13

14 15

16

17

18 19

20

21

22 23

24

25

26

27

28

29

30

31 32

33

34

35 36

37

										089734 cluding		
perineum										Ref. Ref. 734 on 20 January 2 Erasr ling for uses related		
No	22,506	91.14%	31	22.30%			Ref.			f. Ref. 24 on 20-25. Dow 3 Erasmushoges		
Yes	2,188	8.86%	108	77.70%			35.82	24.3, 54.4	<0.001**	15 37 2 42	23.80, 61.01	<0.001**
Newborn										anu: s re		
weight						<0.001*				ary Eras		
(grams)					669.48	*				202: mu: d to		
<2500	227	0.92%	1	0.72%			Ref.			ref.		
2500-40										Downloaded		
00	23,144	93.72%	81	58.27%			0.79	0.18, 14.0	0.820	d 8 5 5 5	0.12, 12.3	0.689
>4000	1,323	5.36%	57	41.01%			9.78	2.14, 173	0.024*		1.15, 120	0.089
Newborn										nini nini		
length						<0.001*				from http://bmjäpen. nining, Al training, ar		
(centimeters)					18.02	*				Al t		
≤55	23,979	97.10%	129	92.81%			Ref.			Ref.		
>55	715	2.90%	10	7.19%			2.60	1.27, 4.72	0.004*	jj 1 <mark>2</mark> 4	0.47, 2.50	0.762
										ાં જે www.loaded from http://bmjäpen.bmj.com/ on June 7, 2025 at Department GEZ-LTA jeschoob . and data mining, Al training, and similar technologies.		
				For peer	review or	nly - http://l	omjopen.bmj.	.com/site/about/	′guidelines.xh			

							BMJ Open			6/bmjop		
Supplementary	Non-C	CA-PPH	CA	A-PPH	racteristi	ics and Risk		ening between No		윽 ㅋ		
Characteristi	(N=2	4,757)	Sam	N=76)	χ^2	P	Univar	iate Logistic Reg	ression	iti III R O Downloaded from http://bmjopen.bmj.com/ on June 7, 20 20 January 2025. Downloaded from http://bmjopen.bmj.com/ on June 7, 20 Erasmushogeschool . uses related to text and data mining, Al training, and similar technologies.	riate Logistic Re	gression
cs	Sampl		ple		λ	1				uary Era elate		
	e size	n%	size	n%			OR	95% CI	P	S OR	95% CI	P
General view										5. D		
Age (years)					6.09	0.530				own ges		
<25	1,308	5.28%	1	1.32%			-	-	-	lload chood da	-	-
25-29	11,702	47.27%	34	44.74%			-	-	-	ded -	-	-
30-34	9,412	38.02%	33	43.42%			-	-	-	fror -	-	-
≥35	2,335	9.43%	8	10.53%			/ →-	-	-	ng,	-	-
Ethnicity					6.92	0.227				Al t		
Han	22,158	89.50%	64	84.21%				-	-	/bm rain	-	-
Manchu	1,862	7.52%	10	13.16%			-	10:	-	ing,	-	-
Other ethnic										n.b anc		
groups	737	2.98%	2	2.63%			-	-	-	isir =	-	-
Educational										om,		
Attainment					1.65	0.895				on tec		
High school										Jun		
or below	8,611	34.78%	24	31.58%			-	-	_	ne 7,	-	-
Bachelor's										, 2025 ies.		
degree	13,597	54.92%	42	55.26%			-	-	-	25 - a	-	-
Postgraduate										at Department		
or higher	2,549	10.30%	10	13.16%			-	-	-	· par	-	-
Occupation					21.85	0.003*				\$		

Unemployed	11,330	45.76%	43	56.58%			Ref.			njopen-2024-089734 Ref.		
Light physical	11,550	43.7070	43	30.3070			Ref.			g for		
labor	2,811	11.35%	14	18.42%			1.31	0.69, 2.34	0.378		0.68, 2.30	0.411
Moderate	_,							,		Janu es ru	,	
physical labor	9,993	40.36%	18	23.68%			0.47	0.27, 0.81	0.008*	26 Janua 1 Erai uses relate	0.27, 0.81	0.008*
Heavy	,							,		202 Smu	,	
physical labor	623	2.52%	1	1.32%			0.42	0.02, 1.94	0.395	2025⊖Do smustbog d to text	0.02, 1.93	0.393
Family Per										oge ar ar		
Capita										nlos scho		
Monthly										oaded าool . data r		
Income										វ from mining		
(10,000 yuan)					6.42	0.492				ing,		
< 0.5	10,297	41.59%	28	36.84%				-	-	http://bmjopen.bmj.com/ on June y, Al training, and similar technolo	-	-
0.5-2.0	9,503	38.39%	31	40.79%				, • -	-	/bm	-	-
2.0-5.0	3,569	14.42%	15	19.74%			<u>-</u>		-	ing -	-	-
>5.0	1,388	5.61%	2	2.63%			-	1	-	en.k	-	-
Pre-pregnan										d <u>si</u>		
cy BMI										con mila		
(Kg/m2)					15.86	0.026*				Ir te		
<18.5										chn Ju		
(Underweig										ne 7 olog		
ht)	7,282	29.41%	12	15.79%			Ref.			f. Ref. I from http://bmjopen.bmj.com/ on June 7, 2025		
18.5-23.9										. 125		
(Normal)	14,953	60.40%	52	68.42%			2.11	1.17, 4.15	0.020*	2.	1.16, 4.11	0.021*
24.0-27.9										epar		
Overweight	1,692	6.83%	8	10.53%			2.87	1.12, 6.95	0.021*	parænent GEZ-LTA	1.12, 6.97	0.021*

 Page 62 of 68

Page 63 of 68

7

8

9 10

11

12

13

14 15

16

17

18 19

20

21

22 23

24

25

26 27

28

29

30

31 32

33

34

35 36

37

							BMJ Open			o homjopen-2024-089734 on 20 Jaby copyright, including for uses			Page 64
										024-089 t, incluc			
No	18,210	73.55%	57	75.00%			-	-	-	734 #ing	-	-	_
Yes	6,547	26.45%	19	25.00%			-	-	-	on 2	-	-	
spontaneous										20 J			
abortion					0.04	0.998				January Eras Ses relate			
No	22,271	89.96%	68	89.47%			-	-	-	nuary 2029 Erasmu related to	-	-	
Yes	2,486	10.04%	8	10.53%			-	-	-	ry 2025. rasmusited to t	-	-	
induced													
abortion or										t an			
medication										. Downloaded from http://bmjopen.bmj.com/ on June 7, 20 shogeschool . text and data mining, AI training, and similar technologies.			
abortion					0.57	0.903				ded ol . ata i			
No	19,920	80.46%	63	82.89%			-	-	-	froi	-	-	
Yes	4,837	19.54%	13	17.11%			<u>_</u>	-	-	r r r r r r r r r r r r r r r r r r r	-	-	
induced										A t Ett			
labor					3.18	0.365				/bm			
				100.00						ing,			
No	24,250	97.95%	76	%			-		-	and b	-	-	
Yes	507	2.05%	0	0.00%			-	- 1	-	d sir	-	-	
assisted										nila on			
reproductive										r te			
technology					0.02	0.999				chn Ju			
No	24,110	97.39%	74	97.37%			-	-	- =	ne 7	-	-	
Yes	647	2.61%	2	2.63%			-	-	-	, 7, 2025 ogies.	-	-	
Gestational										25			
Age at										D D			
Delivery										at Department GEZ			
(weeks)					14.46	0.013*				a			

Page 65 of 68

2 3 4

5

6

7

8

9 10

11

12

13

14 15

16

17

18 19

20

21

22 23

24

25

26

27

28

29

30

31 32

33

34

35 36

37

Yes	2,458	9.93%	9	11.84%			-	-	-	ijopen-2024-089734 on i copyright, including for	-	-
umbilical										on 2		
cord										20 Ja		
entanglement					0.39	0.943				re _nu		
No	17,342	70.05%	55	72.37%			-	-	-	ary : Eras	-	-
Yes	7,415	29.95%	21	27.63%			-	-	-	202: d to	-	-
premature										5. D shootex		
rupture of						<0.001*				own ges		
membranes					39.31	*				Nownloaded Ref. ogeschool .		
No	19,255	77.78%	43	56.58%			Ref.			Ref.		
Yes	5,502	22.22%	33	43.42%			2.69	1.69, 4.22	<0.001**	ai: 2. 3	1.72, 4.51	<0.001**
placental										n ht		
abruption					9.96	0.019*	10,			tp://		
No	24,710	99.81%	75	98.68%			Ref.			Ref.		
Yes	47	0.19%	1	1.32%			7.01	0.39, 32.7	0.056	ng,	-	-
vaginal										n.br and		
bleeding										nj.c sim		
during					0.20	0.042				om/ nilar		
pregnancy	22.502	04.020/	72	06.0504	0.39	0.942				fe. Refer on http://bmjopen.bmj.com/ on June		
No	23,503	94.93%	73	96.05%			-	-	-	Jun	-	-
Yes	1,254	5.07%	3	3.95%			-	-	-		-	-
scarred					12.38	0.006*				7, 2025 at Depa		
uterus No	24,500	98.96%	73	96.05%	12.36	0.000	Ref.			a Ref.		
Yes	257	1.04%		3.95%			3.92	0.96, 10.6	0.021*	6 8 9	1.48, 21.10	0.004*
Delivery Proce							3.74	0.70, 10.0	0.021	o. y art	1.70, 21.10	v.vv4

 Page 66 of 68

Page 67 of 68

2 3 4

5 6

7

8

9 10

11

12

13

14 15

16

17

18 19

20

21

22 23

24

25

26 27

28

29

30

31 32

33

34

35 36

37

38

prolonge									Ć	opvright including for uses		
d	192	0.78%	3	3.95%			5.26	1.28, 14.3	0.005*	g 3.13	0.71, 9.65	0.078
Third stage										o Ja		
of labo					0.45	0.930				ren E E		
normal prolonge	24,553	99.18%	75	98.68%			-	-	-	20 January 2025.	-	-
d	204	0.82%	1	1.32%			-	-	-	v ⊇	-	-
Placental										Downloaded		
retention/Pla										nloa cho		
centa										aded from http://bmjopen.bmj.com/ on June 7, 20 ool .		
accreta/Place										from http://bm/open.bm/.com/ on Ref. 3.54		
ntal						<0.001*			Ç	ng <mark>H</mark>		
implantation					40.92	*				Al i t		
No	20,721	83.70%	49	64.47%			Ref.			Ref.		
Yes	4,036	16.30%	27	35.53%			2.83	1.74, 4.49	<0.001**	2 .97	1.83, 4.73	<0.001*
Analgesia						<0.001*				מ <mark>א</mark>		
during labor					27.45	*						
No	23,831	96.26%	67	88.16%			Ref.		Uh.	Ref.		
Yes	926	3.74%	9	11.84%			3.46	1.60, 6.59	<0.001**	3.54	1.64, 6.79	<0.001*
Instrumental										June		
assistance in										ne 7		
delivery					0.81	0.848				, 20		
No	24,153	97.56%	75	98.68%			-	-	-	125 -	-	-
Yes	604	2.44%	1	1.32%			-	-	-	7, 2025 at De	-	-
Lacerations										ера		
of the cervix,					0.01	0.999				partm		

 Page 68 of 68

Postpartum Hemorrhage Following Vaginal Delivery: A Comprehensive Analysis and Development of Predictive Models for Etiological Subgroups in Chinese Women

Journal:	BMJ Open
Manuscript ID	bmjopen-2024-089734.R2
Article Type:	Original research
Date Submitted by the Author:	02-Dec-2024
Complete List of Authors:	Li, Jinke; Shengjing Hospital of China Medical University, Department of Obstetrics and Gynecology Zhang, Dandan; Shengjing Hospital of China Medical University, Department of Obstetrics and Gynecology Lin, Hong; Liaoning Maternal and Child Health Hospital, Department of Obstetrics and Gynecology Shao, Mengyuan; Shenyang Women's and Children's Hospital, Department of Obstetrics and Gynecology Wang, Xiaoxue; Shengjing Hospital of China Medical University, Department of Health Management Chen, Xueting; Shengjing Hospital of China Medical University, Department of Health Management Zhou, Yangzi; Shengjing Hospital of China Medical University, Department of Obstetrics and Gynecology Song, Zixuan; Shengjing Hospital of China Medical University, Department of Obstetrics and Gynecology
Primary Subject Heading :	Obstetrics and gynaecology
Secondary Subject Heading:	Obstetrics and gynaecology
Keywords:	Maternal medicine < OBSTETRICS, PERINATOLOGY, Postpartum Women < Postpartum Period, PREVENTIVE MEDICINE

SCHOLARONE™ Manuscripts



I, the Submitting Author has the right to grant and does grant on behalf of all authors of the Work (as defined in the below author licence), an exclusive licence and/or a non-exclusive licence for contributions from authors who are: i) UK Crown employees; ii) where BMJ has agreed a CC-BY licence shall apply, and/or iii) in accordance with the terms applicable for US Federal Government officers or employees acting as part of their official duties; on a worldwide, perpetual, irrevocable, royalty-free basis to BMJ Publishing Group Ltd ("BMJ") its licensees and where the relevant Journal is co-owned by BMJ to the co-owners of the Journal, to publish the Work in this journal and any other BMJ products and to exploit all rights, as set out in our licence.

The Submitting Author accepts and understands that any supply made under these terms is made by BMJ to the Submitting Author unless you are acting as an employee on behalf of your employer or a postgraduate student of an affiliated institution which is paying any applicable article publishing charge ("APC") for Open Access articles. Where the Submitting Author wishes to make the Work available on an Open Access basis (and intends to pay the relevant APC), the terms of reuse of such Open Access shall be governed by a Creative Commons licence – details of these licences and which Creative Commons licence will apply to this Work are set out in our licence referred to above.

Other than as permitted in any relevant BMJ Author's Self Archiving Policies, I confirm this Work has not been accepted for publication elsewhere, is not being considered for publication elsewhere and does not duplicate material already published. I confirm all authors consent to publication of this Work and authorise the granting of this licence.

Postpartum Hemorrhage Following Vaginal Delivery: A Comprehensive Analysis and Development of Predictive Models for Etiological Subgroups in Chinese Women

Jinke Li^{1#}, Dandan Zhang^{1#}, Hong Lin², Mengyuan Shao³, Xiaoxue Wang⁴, Xueting Chen⁴, Yangzi Zhou¹ and Zixuan Song^{1*}

- 1 Department of Obstetrics and Gynecology, Shengjing Hospital of China Medical University, Shenyang, China
- 2 Department of Obstetrics and Gynecology, Liaoning Maternal and Child Health Hospital, Shenyang, China
- 3 Department of Obstetrics and Gynecology, Shenyang Women's and Children's Hospital, Shenyang, China
- 4 Department of Health Management, Shengjing Hospital of China Medical University, Shenyang, China
- #These authors contributed equally to this work.
- Correspondence:songzixuan666@163.com

Abstract:

Objectives: This study aimed to dissect the etiological subgroups of postpartum hemorrhage (PPH) that occur after vaginal delivery in women with full-term singleton pregnancies. Our goal was to craft and validate predictive models to guide clinical decision-making and optimize resource allocation.

Design: A retrospective cohort study.

Setting: Shengjing Hospital of China Medical University, Liaoning Maternal and Child Health Hospital, and Shenyang Women's and Children's Hospital.

Participants: 29,842 women who underwent vaginal delivery were enrolled in the study across three hospitals from 2016 to 2022.

Primary outcome measures: PPH, categorized into uterine atony (UA), placental factors (PF), cervical trauma (CT), and coagulation abnormalities (CA) by etiology.

Results: The logistic regression for overall PPH and UA-PPH showcased high discrimination (AUCs of 0.807 and 0.794, respectively), coupled with commendable calibration and DCA-assessed clinical utility, culminating in the development of a nomogram for risk prediction. The PF-PPH model exhibited a modest AUC of 0.739, while the CT-PPH and CA-PPH models demonstrated suboptimal clinical utility and calibration.

Conclusions: The study identified factors associated with PPH and developed models with good performance for overall PPH and UA-PPH. The nomogram offers a valuable tool for risk prediction. However, models for PF-PPH, CT-PPH, and CA-PPH require further refinement. Future research should focus on larger samples and multi-center validation for enhanced model generalizability.

Keywords: Postpartum Hemorrhage, Vaginal Delivery, Etiological Subgroups, Predictive Models, nomogram

Strengths and limitations of this study

- This study included data from large multicenter cohorts in China, comprising 29,842 women to enhance the statistical power of the analysis.
- Predictive models were developed for postpartum hemorrhage following vaginal delivery and for different etiological subgroups.

- This study's test set is a single-center study specifically focused on vaginal delivery PPH, with a limited number of positive samples.
- The existing external validation datasets are from hospitals of the same region and level, resulting in a lack of diversification and generalizability.

Background

Postpartum hemorrhage (PPH) is defined as blood loss exceeding 500 milliliters within 24 hours following vaginal delivery or exceeding 1,000 milliliters within 24 hours following cesarean delivery.¹ PPH is a widespread and serious medical condition that poses significant risks to women's health around the world. It is particularly devastating in developing countries, where it is a principal contributor to maternal mortality.² It is estimated that approximately 1.4 million maternal deaths globally are tied to PPH each year, with the tragic loss of a woman's life to this condition occurring every four minutes.³⁴. In Australia, the incidence of PPH increased from 6.3% in 2000 to 8.0% in 2009. ⁵ Similarly, in the United States, the rate of PPH rose from 2.7% in 1999 to 3.2% in 2014. ⁶ In China, despite a relatively lower maternal mortality rate of 17.8 per 100,000 in 2019, PPH accounted for one-quarter of these deaths.⁷

The World Health Organization (WHO) has conducted an analysis revealing that while PPH is a significant factor in maternal mortality and morbidity, the mortality rates vary considerably across different regions.⁸ In high-income countries, the risk of death due to PPH is significantly lower than in low-income countries.⁹ In high-income nations, the substantial blood loss primarily caused by PPH accounts for 13.4% of overall maternal mortality, while in Africa and Asia, this figure stands at 34% and 30.8%, respectively. ⁸ The international obstetric community is actively engaged in research to better understand the incidence, risk factors, and management strategies for PPH.¹⁰⁻¹². Despite the establishment of global clinical guidelines and the identification of various risk factors, further exploration is needed to enhance our understanding and management of PPH.¹³ ¹⁴.

PPH can be etiologically classified into uterine atony (UA), placental factors (PF), cervical trauma (CT), and coagulation abnormalities (CA), each requiring distinct clinical management and treatment strategies. ¹⁵ Clear etiological classification is crucial for developing preventive strategies, formulating management plans, and rational allocation of medical resources. ¹⁶ While numerous cohort studies have focused on identifying risk factors for PPH, there is a scarcity of studies that quantify and weigh these risk factors for a comprehensive PPH risk assessment. ¹⁷ ¹⁸ Given the complexity of PPH and the interplay of multiple risk factors, a holistic approach is necessary to accurately assess the risk of PPH.

Clinical prediction models (CPMs) have been widely applied in clinical settings in recent years. By constructing CPMs, physicians and patients can make better medical decisions, and health departments at all levels can allocate medical resources more rationally. These models play an irreplaceable role in primary prevention (assessing the quantitative risk of future diseases) and secondary prevention (constructing highly sensitive and specific diagnostic schemes, practicing "early detection, early diagnosis, early treatment"), reflecting significant health economic value.

There is a gap in research regarding the development of clinical prediction models for women specifically following vaginal delivery. Many studies are constrained by limited sample sizes, which can affect the robustness of the models 19. Other research has focused on PPH prediction models for women undergoing cesarean sections. 20 Our study aims to address this gap by constructing a clinical prediction model tailored to PPH after vaginal delivery. By analyzing clinical data and risk factors through logistic regression, we can determine the relative impact of each factor on the likelihood of PPH. We further refine our model by performing secondary fitting based on the four etiological subgroups, creating a nomogram that enhances the precision of predicting high-risk populations for PPH. This work provides essential insights for the prevention and management of this critical condition.

Materials and methods

Data Sources and Ethics Statement

This cohort study was conducted at the obstetric wards of Shengjing Hospital of China Medical University, Liaoning Maternal and Child Health Hospital, and Shenyang Women's and Children's Hospital. The study population comprised women who underwent vaginal delivery between January 1, 2016, and December 31, 2022. The outcomes of interest were fetal birth outcomes within the first 24 hours postpartum. Inclusion criteria were women who consented to participate after being informed of the study's scope. Exclusion criteria were defined as follows: age under 18 or over 50 years, delivery occurring at less than 37 weeks or more than 42 weeks of gestation, multiple births, and instances of induced labor, stillbirth, or fetal death. These factors were excluded to focus on women with full-term, singleton pregnancies undergoing vaginal delivery. Preterm and postterm pregnancies, multiple pregnancies, and induced labor are associated with different physiological characteristics and obstetric risks, which could confound the analysis of PPH risk in this cohort. Additionally, stillbirth and fetal death involve other pathological processes and significant complications that are outside the scope of this study's focus on live births and PPH risk.

Comprehensive data encompassing maternal characteristics, obstetric and gynecologic history, pregnancy complications, and details of the delivery process and neonatal conditions were collected (Supplement 1). To protect participant privacy, all data were anonymized. The study protocol was approved by the Ethical Review Committee of Shengjing Hospital of China Medical University (No. 2016PS344K), and written information about the study was provided to all participants.

Sample Size Calculation

According to the obstetric big data from Shengjing Hospital of China Medical University, the incidence of PPH is approximately 6%-7%. Based on the sample size estimation using the psamplesize package in R software, the minimum sample size required for constructing clinical prediction models is estimated to be between 1,048 and 1,536. The sample size included in this study far exceeds the minimum requirement.

Covariates

A range of covariates were taken into account, including:(1) Age, categorized as <25, 25-29, 30-34, and ≥35 years; (2) Ethnicity, divided into Han, Manchu, and other; (3) Education level, classified as high school or below, bachelor's degree, and postgraduate or above; (4) Occupation, categorized as unemployed, light physical labor, moderate physical labor, and heavy physical labor (based on the International Physical Activity Questionnaire, IPAQ); (5) Monthly household income per capita, divided into <0.5, 0.5-2.0, 2.0-5.0, and >5.0 thousand yuan; (6) Pre-pregnancy BMI, categorized as underweight (<18.5 kg/m²), normal (18.5-23.9 kg/m²), overweight (24-27.9 kg/m²), and obese (≥28 kg/m²); (7) Smoking and alcohol consumption history; (8) Gravidity, categorized as 1, 2, or \geq 3 times; (9) Parity, divided into 0, 1, or ≥ 2 times; (10) History of miscarriage and induced labor; (11) Assisted reproductive technology; (11) Gestational age, categorized as <38, 38-40, and >40 weeks; (12) Pregnancy complications: diabetes, hypertension, anemia, coagulation disorder, uterine fibroids/adenomyosis, polyhydramnios, umbilical cord entanglement, premature rupture of membranes, placental abruption, vaginal bleeding during pregnancy, and scarred uterus; (13) Delivery time, divided into daytime (8-16), evening (17-23), and night (0-7) shifts; (14) Total duration of labor, categorized as normal (≤24 hours) and prolonged (>24 hours); (15) Latent phase of the first stage, categorized as normal (primiparous ≤20 hours, multiparous ≤ 14 hours) and prolonged (primiparous ≥ 20 hours, multiparous ≥ 14 hours); (16) Active phase of the first stage, categorized as normal (≤ 8 hours) and prolonged (> 8 hours); (17) Second stage duration, categorized as normal and prolonged based on specific criteria for primiparous and multiparous women with or without analgesia; (18) Third stage duration, categorized as normal (≤30 minutes) and prolonged (>30 minutes); (19) Placental retention/placenta accreta/placental implantation; (20) Analgesia during labor; (21)

Instrumental assistance in delivery; (22) Lacerations of the cervix, vagina, or perineum; (23) Newborn weight and length.

Etiology Subgroups

In this study, PPH was defined according to WHO standards as blood loss exceeding 500 milliliters following vaginal delivery. Patients were categorized into those with and without PPH based on this definition, and further analysis was conducted on the underlying etiologies, including UA, PF, CT, and CA.

Model Construction

For the purpose of our investigation, we have categorized the participants from the Shengjing hospital of China Medical University as Cohort I. This cohort was systematically split into a training dataset and an internal validation dataset with a ratio of 7:3. The training dataset was instrumental in developing the predictive model, while the internal validation dataset served to assess the model's predictive accuracy. An additional cohort, comprising participants from two other hospitals, was designated as Cohort II. This external dataset was used to validate the model's general applicability and its efficacy in real-world clinical scenarios.

Within the confines of the datasets, we employed both univariate and multivariate logistic regression analyses to identify potential risk factors across various subgroups. These factors were then subjected to a rigorous selection process for inclusion in the predictive model. The selected factors were further analyzed using multivariate logistic regression (Bidirectional elimination) in training dataset to discern their discriminative power, thereby establishing them as predictive indicators for the model. The women in the training set were divided into PPH and non-PPH groups based on PPH as the outcome variable. Subsequently, they were further categorized by etiology into UA-PPH and non-UA-PPH groups, PF-PPH and non-PF-PPH groups, CT-PPH and non-CT-PPH groups, and CA-PPH and non-CA-PPH groups. Five predictive models were constructed sequentially, and the performance of these models was corroborated using both the test and validation datasets to ascertain the most accurate predictive model.

Evaluating the Performance of the Models

The area under the receiver operating characteristic curve (AUC) was the primary metric used to evaluate the discrimination of our models. An AUC value above 0.75 suggests excellent model discrimination, while an AUC below 0.6 indicates poor discrimination. Calibration curves were used to assess the models' accuracy, with closer alignment between observed and predicted incidence rates indicating higher model fidelity. Decision Curve Analysis (DCA) was also employed to evaluate the clinical utility of the models, offering a thorough assessment of the models' net benefits across various clinical scenarios.

Nomogram Development

Nomograms for postpartum hemorrhage and its four etiological subgroups were crafted to offer a visual representation of the risk scores derived from the logistic regression analysis. This tool simplifies the interpretation of complex statistical outcomes, providing a more straightforward approach to understanding risk assessments.

Statistical Analysis

All statistical computations, construction of traditional logistic models, and calculations of model discrimination and calibration were carried out using R version 3.6.3 from the R Foundation for Statistical Computing, Vienna, Austria. This software facilitated the development of traditional logistic predictive models and their subsequent evaluation for discriminative power, calibration, and clinical utility. Continuous variables conforming to the normal distribution were expressed as the mean \pm standard deviation (SD), while non-normally distributed continuous variables were presented as medians with interquartile ranges. Categorical data were analyzed using chi-square tests, and continuous variables were analyzed using ANOVA or Mann-Whitney

tests, as appropriate. Variables were adjusted as dummy variables, and odds ratios (OR) with corresponding 95% confidence intervals (95% CI) were calculated using univariate and multivariate logistic regression analyses, with significance level set at P < 0.05.

Patient and public involvement

Patients and/or the public were not involved in the design, conduct, reporting or dissemination plans of this research.

Results

From 2016 to 2022, a total of 27,389 patients underwent vaginal delivery at the Shengjing Hospital of China Medical University. Forty-two patients under 18 years of age or over 50 years old were excluded. Additionally, 2,456 patients with gestational age less than 37 weeks or more than 42 weeks at delivery, 6 patients with multiple births, and 52 patients with induced labor, stillbirth, or fetal death were also excluded. Ultimately, 24,833 patients met the inclusion criteria and were enrolled in the cohort. According to the inclusion and exclusion criteria, a total of 5,099 patients in cohort II were included in the external validation dataset. The general characteristics of all patients are presented in Table 1. All patients were followed up within 24 hours after delivery for neonatal outcomes, with a follow-up rate of 100%. The patient selection criteria flowchart is shown in Figure 1.

Comparison of Basic Characteristics and Risk Analysis for Postpartum Hemorrhage (PPH) and Its Subgroups

Based on the occurrence of postpartum hemorrhage, the parturients in cohort one were divided into two groups: the non-PPH group and the PPH group. Similarly, within the etiological subgroups, they were categorized into UA-PPH and non-UA-PPH groups, PF-PPH and non-PF-PPH groups, CT-PPH and non-CT-PPH groups, and CA-PPH and non-CA-PPH groups. The comparison of basic characteristics and analysis of risk factors for each group are presented in Supplementary Tables 1-5.

In the multivariate analysis of risk factors, apart from age, parity, pre-pregnancy BMI, anemia, premature rupture of membranes, and combined placenta retention/placenta accreta/placental implantation, other specific risk factors were found to be associated with specific etiologies of postpartum hemorrhage. For instance, polyhydramnios was associated with UA-PPH; analgesia during labor, instrumental assistance, and cervical/vaginal/perineal lacerations were associated with the occurrence of CT-PPH (Table 2).

Selection of Predictive Factors for PPH and Its Subgroups in the Training Dataset

Through random sampling of cohort one, 70% of the data (N=17,383) from parturients were used to form the training dataset, with the remaining approximately 30% (N=7,450) forming the internal validation dataset. Multivariate analysis of risk factors for PPH and its subgroups was performed again in the training dataset, with results presented in Supplemental Table 1. After selection, predictive models were constructed for each group using the selected risk factors.

Evaluation of Predictive Model Discrimination

The ROC curves were plotted using R software for the PPH group and its various subgroups across the training dataset, internal and external validation dataset.

The results indicated that the predictive models, namely PPH-Logistic, UA-PPH-Logistic, PF-PPH-Logistic, CT-PPH-Logistic, and CA-PPH-Logistic, demonstrated high discriminative power in the training dataset with AUCs of 0.807 (95% CI: 0.792-0.821), 0.794 (95% CI: 0.777-0.811), 0.796 (95% CI: 0.761-0.830), 0.935 (95% CI: 0.901-0.969), and 0.802 (95% CI: 0.769-0.892), respectively.(Figure 2A-E) However, the PF-PPH-Logistic model exhibited only moderate discrimination with an AUC of 0.739 (95% CI: 0.666-0.813) in the internal validation dataset. Furthermore, the CA-PPH-Logistic model showed significantly lower discrimination in the external validation dataset with an AUC of 0.662 (95% CI: 0.450-0.873), which was notably inferior to its performance in the training and test datasets. This discrepancy may be attributed to

Assessment of Predictive Model Calibration

Calibration curves for the PPH and its subgroups were plotted for the Logistic predictive model within the training dataset (Supplemental Figure 1A-E). The performance of the PF-PPH-Logistic, particularly the CT-PPH-Logistic, and CA-PPH-Logistic models was suboptimal in certain aspects, with lower calibration, as observed in the test and external validation datasets (Supplemental Figure 1 F-J, Supplemental Figure 1K-O).

Evaluation of Clinical Utility of Predictive Models

In the evaluation of clinical utility, the PPH-Logistic and UA-PPH-Logistic models demonstrated satisfactory performance across all datasets. However, the clinical utility of the PF-PPH-Logistic, CT-PPH-Logistic, and CA-PPH-Logistic models was found to be relatively poor. (Supplemental Figure 2A-O)

Nomogram Construction

Using R software, we constructed nomograms for PPH and UA-PPH, with the results presented in Figure 3A-B. Physicians can assess the risk probability of PPH occurrence by summing the individual scores on the nomogram. This practical tool aids in a more precise estimation of PPH risk, thereby enhancing clinical decision-making.

Discussion

Maternal mortality has emerged as a pivotal indicator in global maternal and child health, serving as a significant benchmark for assessing the socioeconomic status of nations. Consequently, the effective reduction, prevention, and improvement of conditions leading to maternal deaths have become a focal point for public health initiatives worldwide. Among the various causes of maternal mortality, PPH stands out as a preventable condition that has attracted considerable attention. ^{17 18} With the rise in global economic standards and the evolution of medical technologies, there has been an approximate 50% decrease in the worldwide maternal mortality rate between 1990 and 2015. In China, the maternal mortality rate has seen a dramatic reduction of 98.78%²¹ since the establishment of the People's Republic of China. Despite these advancements, a substantial proportion of maternal deaths, estimated between 27% and 40%²², remain avoidable due to a range of factors, including inadequate social and medical interventions. PPH is a critical area of focus within this context, and the prediction and prevention of PPH to reduce avoidable maternal mortality present a significant challenge on the global stage.

Early prediction or identification of PPH and timely preventive or intervention measures are extremely valuable, necessitating a clear understanding of the etiologies of PPH for targeted management. The overall PPH model provides a comprehensive perspective that captures common risk factors for postpartum hemorrhage, offering a baseline risk assessment for women. This broad assessment helps in formulating general preventive measures and policies, ensuring that all women at risk of PPH are monitored. Although the overall PPH model demonstrates higher overall accuracy, it may mask the heterogeneity between different PPH subtypes in clinical practice, limiting its guidance for individualized management. Subtype-specific models, on the other hand, can delve deeper into the specific risk factors of each subtype, helping to enhance our understanding of the potential mechanisms of PPH and meet the diverse needs of clinical practice. The value provided by these models in treatment is more critical, as they can guide clinicians to adopt effective therapeutic strategies tailored to the specific etiology, thereby reducing complications and mortality and promoting the development of precision medicine. In this study, the predictive models developed for PPH and UA-PPH demonstrated excellent performance in effectively identifying high-risk populations. For women at high risk for UA-PPH, early cord clamping should be considered. After the placenta is delivered, uterine fundal massage can be performed, and oxytocin should be administered immediately to promote uterine contraction and reduce bleeding. If necessary, mechanical compression or uterine artery ligation can be employed for hemostasis. In other PPH subtype models, the performance may be suboptimal due to factors

The advent of the big data era has brought new opportunities for the management of PPH. The era is characterized by the digitization and standardization of medical records, along with an increasing volume of data, which has ushered in an era of data-driven management and treatment for maternal care. Leveraging big data analytics for disease risk prediction can contribute to the reduction of avoidable maternal deaths.

A review of the literature reveals over 200 prognostic models in obstetrics, three of which are pertinent to PPH.²³ However, few models have been applied in routine clinical practice, and the majority of studies have not provided model formulas, hindering independent external validation. The earliest PPH prediction model, dating back to 1994, originated from a case-control study in Zimbabwe²⁴, where PPH was defined as blood loss exceeding 600 milliliters following an unassisted vaginal delivery. This study included 150 PPH patients and 299 patients with normal deliveries, with a low positive predictive value of less than 7% and only 35.0% of patients experiencing postpartum bleeding. Since then, approximately ten additional PPH prediction models have been published. These models have varied in focus, with some concentrating on the relationship between placenta previa and PPH, while others have included only vaginal deliveries²⁴⁻²⁷ or cesarean sections²⁰ ²⁸ ²⁹, and some have targeted women with placental implantation disorders³⁰ or general obstetric populations³¹. PPH research has been conducted in hospitals across various countries, including Italy, China, France, the United States, the United Kingdom, South Korea, the Netherlands, Spain, Zimbabwe, Denmark, and Egypt. From the 14 published studies, a total of 124 independent variables were identified as potential predictors (ranging from 5 to 38 per study), and 64 variables were ultimately selected for the final models (an average of 5-15 factors per study). Common predictors included parity, low pre-pregnancy hemoglobin, antenatal bleeding, maternal age over 35, gestational age, high neonatal weight, multiple pregnancies, body mass index (BMI) over 25, previous cesarean section, anterior placenta, and retained placenta. These predictors have also been incorporated into our predictive model

Once a clinical prediction model is developed, it must undergo validation and evaluation to assess the model's effectiveness, reproducibility, and portability. Published PPH prediction models have reported AUCs ranging from 0.70^{25} to 0.90^{27} , with external validation AUCs of 0.83^{20} , which are comparable to the results of our study. In addition to discrimination, calibration is essential to evaluate the consistency between the predicted probabilities of clinical outcomes and the observed event probabilities. Only a few studies, such as one by Albright in 2019 on the prediction of PPH following cesarean section, have utilized calibration curves³², while most have employed the Hosmer-Lemeshow goodness-of-fit test to compare predicted probabilities with actual event probabilities for significant differences. The Hosmer-Lemeshow test, however, has limited efficacy in small-sample prediction models as it does not quantify model calibration^{33 34}or provide direction or magnitude of mis calibration³⁵.

The Decision Curve Analysis (DCA)³⁵ has been used to evaluate the clinical utility of models, focusing on the selection of true positives from positive patients to avoid unnecessary medical resource consumption and reduce harm from overtreatment of false positives. DCA is particularly suited for scenarios where symptoms suggest the possibility of disease but a diagnosis has not yet been confirmed, guiding the decision on whether or what kind of screening method to adopt for disease diagnosis. The DCA's axes represent the threshold probability (P) and net benefit (NB), allowing for the determination of intervention measures based on the predicted probability of adverse events.³⁶

In essence, both ROC and DCA can be used to assess the quality of predictive models, but they differ fundamentally in their theoretical constructs. While ROC combines sensitivity and specificity to compare the accuracy of predictive models through the AUC, the highest AUC does not necessarily represent the optimal model in clinical practice. For instance, in this study, patients

Statistical analysis of previously published PPH data has shown that factors such as general anesthesia in pregnant women, prolonged use of oxytocin, excessive uterine tension (multiple pregnancies, polyhydramnios), and chorioamnionitis are all associated with uterine atony, potentially increasing the risk of postpartum bleeding. Previous studies have suggested that for pregnant women with high-risk factors, assessing and selecting appropriate treatment options and management based on the type and weight of different risks can reduce the risk of adverse pregnancy outcomes or death³⁷. Early prediction and intervention are key measures in reducing maternal mortality, with studies finding that timely interventions can effectively lower maternal mortality rates by 10%³⁸. Establishing a model that predicts the risk of PPH following vaginal delivery and guides clinical practice is a significant task for maternal and child health.

Limitation

Although some subtypes in this study showed promising predictive results, the clinical application of the logistic regression models for the PF-PPH, CT-PPH, and CA-PPH groups is limited by insufficient positive sample sizes in these subgroups, which prevents fully achieving the initial goal of etiology-based PPH prediction. Compared to previously published models, this study's test set is a single-center study focusing specifically on vaginal delivery PPH, with a limited number of positive samples. Additionally, the external validation datasets come from hospitals of the same region and level, lacking diversity and generalizability. Therefore, future plans include multi-center collaboration, increasing sample sizes, and seeking higher-quality prediction methods to provide more robust clinical evidence.

Furthermore, the PPH scoring model developed in this study is not applicable to women with preterm or postterm labor, induced labor, or multiple pregnancies. These populations have unique physiological and pathological factors, along with higher complication rates, that fall outside the scope of this study. Further research is needed to develop individualized risk assessment models for these groups. It is important to note that these models incorporate not only antepartum factors but also postpartum factors such as labor characteristics and newborn weight. This comprehensive approach aids in understanding the mechanisms of PPH and provides a basis for targeted prevention or intervention strategies aimed at risk factors. However, these models are not entirely suitable for antepartum clinical decision-making or real-time prediction.

Conclusion

In conclusion, our study has successfully developed and validated predictive models for PPH following vaginal delivery, offering a novel approach to risk assessment in this critical area of maternal health. The models, particularly for overall PPH and UA-PPH, demonstrated high discriminative power and clinical utility, with the nomogram providing a user-friendly tool for clinicians. Despite the promising results, limitations exist in the application of the PF-PPH, CT-PPH, and CA-PPH models due to the insufficient positive sample size in these subgroups. The generalizability of our findings may also be limited by the single-center nature of the study and the regional characteristics of the included hospitals. Future research should aim to expand the sample size and include multi-center data to improve the models' applicability and robustness. This study contributes to the growing body of evidence on PPH management and has the potential to influence policy and practice, ultimately enhancing maternal care and outcomes.

List of abbreviations

RP: Refined Peyton;

TTM: Traditional Teaching-mode

DOPS: direct observation of procedural skills

OB-GYN: Obstetrics and Gynecology CBVI: computer-based video instruction

PEARLS: Promoting Excellence and Reflective Learning in Simulation

OSCE: objective structured clinical assessment

Declarations

Ethics approval and consent to participate

Ethics approval and consent to participate: The study was approved by the Ethics Committee of Shengjing Hospital of China Medical University (No. 2016PS344K, Date.17/12/2016). All participants provided informed consent.

Consent for publication

Not Applicable

Availability of data and material

The data that support the findings of this study are available from the corresponding author upon reasonable request.

Competing interests

No potential conflict of interest was reported by the author(s).

Funding

This study was supported in part by grants from 345 Talent Project of Shengjing Hospital of China Medical University (No. M0946), and Medical Education Research Project of Liaoning Province (No. 2022-N005-03).

Authors' contributions

Contributors JL and DZ contributed equally to this study. JL, DZ and KZ designed the study and drafted the manuscript. HL, MS, and XW done the data collection. YZ and XC designed the statistical analysis plan. DZ has participated the training and reviewed and co-authored the manuscript with ZS. All authors have read and approved this manuscript. ZS is responsible for the overall content as guarantor.

Acknowledgements

We would like to express our gratitude to all those who helped us during the writing of this manuscript. Thanks to all the peer reviewers for their opinions and suggestions. We would like to acknowledge that Jinke Li and Dandan Zhang have contributed equally to this work.

References:

1. Practice Bulletin No. 183 Summary: Postpartum Hemorrhage. Obstetrics and gynecology

- 2017;130(4):923-25. doi: 10.1097/aog.000000000002346
- 2. World Health Organization. Trends in maternal mortality 2000 to 2017: estimates by WHO, UNICEF, UNFPA, World Bank Group and the United Nations Population Division [Available from: https://www. who. int/ publications/i/item/9789240012202. accessed 2020 12 22.
- 3. Who UJG, Department of Reproductive Health, Research W. UNFPA: Maternal Mortality in 2000: Estimates developed by WHO, UNICEF and UNFPA. 2004
- 4. AbouZahr C. Global burden of maternal death and disability. *British Medical Bulletin* 2003;67(1):1-11. doi: 10.1093/bmb/ldg015 %J British Medical Bulletin
- Mehrabadi A, Hutcheon JA, Lee L, et al. Trends in postpartum hemorrhage from 2000 to 2009: a population-based study. BMC Pregnancy Childbirth 2012;12:108. doi: 10.1186/1471-2393-12-108 [published Online First: 2012/10/13]
- 6. Reale SC, Easter SR, Xu X, et al. Trends in Postpartum Hemorrhage in the United States From 2010 to 2014. *Anesth Analg* 2020;130(5):e119-e22. doi: 10.1213/ane.0000000000004424 [published Online First: 2019/10/01]
- You JHS, Leung TY. Cost-effectiveness analysis of carbetocin for prevention of postpartum hemorrhage in a low-burden high-resource city of China. *PloS one* 2022;17(12):e0279130. doi: 10.1371/journal.pone.0279130 [published Online First: 2022/12/16]
- 8. AbouZahr CJBmb. Global burden of maternal death and disability. 2003;67(1):1-11.
- 9. Mousa HA, Walkinshaw SJCoiO, Gynecology. Major postpartum haemorrhage. 2001;13(6):595-603.
- 10. Akter S, Forbes G, Vazquez Corona M, et al. Perceptions and experiences of the prevention, detection, and management of postpartum haemorrhage: a qualitative evidence synthesis. The Cochrane database of systematic reviews 2023;11(11):Cd013795. doi: 10.1002/14651858.CD013795.pub2 [published Online First: 2023/11/27]
- 11. Zhang R, Cao X, Feng H, et al. Review of clinical practice guidelines for postpartum hemorrhage according to AGREE II. *Midwifery* 2023;121:103659. doi: 10.1016/j.midw.2023.103659 [published Online First: 2023/03/30]
- 12. Zdanowicz JA, Schneider S, Martignoni C, et al. A Retrospective before and after Assessment of Multidisciplinary Management for Postpartum Hemorrhage. *Journal of clinical medicine* 2023;12(23) doi: 10.3390/jcm12237471 [published Online First: 2023/12/09]
- 13. Giouleka S, Tsakiridis I, Kalogiannidis I, et al. Postpartum Hemorrhage: A Comprehensive Review of Guidelines. *Obstetrical & gynecological survey* 2022;77(11):665-82. doi: 10.1097/ogx.00000000001061 [published Online First: 2022/11/09]
- 14. de Vries PLM, Deneux-Tharaux C, Baud D, et al. Postpartum haemorrhage in high-resource settings: Variations in clinical management and future research directions based on a comparative study of national guidelines. *BJOG : an international journal of obstetrics and gynaecology* 2023;130(13):1639-52. doi: 10.1111/1471-0528.17551 [published Online First: 2023/06/01]
- 15. Klufio CA, Amoa AB, Kariwiga G. Primary postpartum haemorrhage: causes, aetiological risk factors, prevention and management. *Papua and New Guinea medical journal* 1995;38(2):133-49. [published Online First: 1995/06/01]
- 16. Gyamfi-Bannerman C, Srinivas SK, Wright JD, et al. Postpartum hemorrhage outcomes and race.

 **American Journal of Obstetrics and Gynecology 2018;219(2):185.e1-85.e10. doi: https://doi.org/10.1016/j.ajog.2018.04.052
- 17. Desale M, Thinkhamrop J, Lumbiganon P, et al. Ending preventable maternal and newborn deaths

- 18. Sotunsa JO, Adeniyi AA, Imaralu JO, et al. Maternal near-miss and death among women with postpartum haemorrhage: a secondary analysis of the Nigeria Near-miss and Maternal Death Survey. *BJOG*: an international journal of obstetrics and gynaecology 2019;126 Suppl 3:19-25. doi: 10.1111/1471-0528.15624 [published Online First: 2019/03/22]
- Goad L, Rockhill K, Schwarz J, et al. Development and validation of a prediction model for postpartum hemorrhage at a single safety net tertiary care center. *American journal of obstetrics & gynecology MFM* 2021;3(5):100404. doi: 10.1016/j.ajogmf.2021.100404 [published Online First: 2021/05/29]
- 20. Albright CM, Spillane TE, Hughes BL, et al. A Regression Model for Prediction of Cesarean-Associated Blood Transfusion. Am J Perinatol 2019;36(9):879-85. doi: 10.1055/s-0039-1678604 [published Online First: 2019/02/12]
- 21. Liang J, Li X, Kang C, et al. Maternal mortality ratios in 2852 Chinese counties, 1996-2015, and achievement of Millennium Development Goal 5 in China: a subnational analysis of the Global Burden of Disease Study 2016. *Lancet* 2019;393(10168):241-52. doi: 10.1016/s0140-6736(18)31712-4 [published Online First: 2018/12/18]
- 22. Gao Y, Zhou H, Singh NS, et al. Progress and challenges in maternal health in western China: a Countdown to 2015 national case study. *Lancet Glob Health* 2017;5(5):e523-e36. doi: 10.1016/s2214-109x(17)30100-6 [published Online First: 2017/03/28]
- 23. Kleinrouweler CE, Cheong-See FM, Collins GS, et al. Prognostic models in obstetrics: available, but far from applicable. *Am J Obstet Gynecol* 2016;214(1):79-90.e36. doi: 10.1016/j.ajog.2015.06.013 [published Online First: 2015/06/14]
- 24. Tsu VD. Antenatal screening: its use in assessing obstetric risk factors in Zimbabwe. *J Epidemiol Community Health* 1994;48(3):297-305. doi: 10.1136/jech.48.3.297 [published Online First: 1994/06/01]
- 25. Biguzzi E, Franchi F, Ambrogi F, et al. Risk factors for postpartum hemorrhage in a cohort of 6011 Italian women. *Thromb Res* 2012;129(4):e1-7. doi: 10.1016/j.thromres.2011.09.010 [published Online First: 2011/10/25]
- 26. Prata N, Hamza S, Bell S, et al. Inability to predict postpartum hemorrhage: insights from Egyptian intervention data. BMC Pregnancy Childbirth 2011;11:97. doi: 10.1186/1471-2393-11-97 [published Online First: 2011/11/30]
- Rubio-Álvarez A, Molina-Alarcón M, Arias-Arias Á, et al. Development and validation of a predictive model for excessive postpartum blood loss: A retrospective, cohort study. *Int J Nurs Stud* 2018;79:114-21. doi: 10.1016/j.ijnurstu.2017.11.009 [published Online First: 2017/12/11]
- 28. Ahmadzia HK, Phillips JM, James AH, et al. Predicting peripartum blood transfusion in women undergoing cesarean delivery: A risk prediction model. *PloS one* 2018;13(12):e0208417. doi: 10.1371/journal.pone.0208417 [published Online First: 2018/12/15]
- 29. Dunkerton SE, Jeve YB, Walkinshaw N, et al. Predicting Postpartum Hemorrhage (PPH) during Cesarean Delivery Using the Leicester PPH Predict Tool: A Retrospective Cohort Study. *Am J Perinatol* 2018;35(2):163-69. doi: 10.1055/s-0037-1606332 [published Online First: 2017/08/29]
- 30. Yoon SY, You JY, Choi SJ, et al. A combined ultrasound and clinical scoring model for the prediction

- of peripartum complications in pregnancies complicated by placenta previa. *Eur J Obstet Gynecol Reprod Biol* 2014;180:111-5. doi: 10.1016/j.ejogrb.2014.06.030 [published Online First: 2014/08/01]
- 31. Chi Z, Zhang S, Wang Y, et al. Research of the assessable method of postpartum hemorrhage. *Technol Health Care* 2016;24 Suppl 2:S465-9. doi: 10.3233/thc-161169 [published Online First: 2016/05/11]
- 32. !!! INVALID CITATION !!! [73]
- 33. Moher D, Liberati A, Tetzlaff J, et al. Preferred reporting items for systematic reviews and meta-analyses: the PRISMA statement. *PLoS Med* 2009;6(7):e1000097. doi: 10.1371/journal.pmed.1000097 [published Online First: 2009/07/22]
- 34. Steyerberg EW, Vergouwe Y. Towards better clinical prediction models: seven steps for development and an ABCD for validation. *Eur Heart J* 2014;35(29):1925-31. doi: 10.1093/eurheartj/ehu207 [published Online First: 2014/06/06]
- 35. Vickers AJ, Elkin EB. Decision curve analysis: a novel method for evaluating prediction models.

 Med Decis Making 2006;26(6):565-74. doi: 10.1177/0272989x06295361 [published Online First: 2006/11/14]
- 36. Hijazi Z, Oldgren J, Lindbäck J, et al. The novel biomarker-based ABC (age, biomarkers, clinical history)-bleeding risk score for patients with atrial fibrillation: a derivation and validation study. *Lancet* 2016;387(10035):2302-11. doi: 10.1016/s0140-6736(16)00741-8 [published Online First: 2016/04/09]
- 37. Zuckerwise LC, Lipkind HS. Maternal early warning systems-Towards reducing preventable maternal mortality and severe maternal morbidity through improved clinical surveillance and responsiveness. *Semin Perinatol* 2017;41(3):161-65. doi: 10.1053/j.semperi.2017.03.005 [published Online First: 2017/04/19]
- 38. Aoyama K, D'Souza R, Pinto R, et al. Risk prediction models for maternal mortality: A systematic review and meta-analysis. *PloS* one 2018;13(12):e0208563. doi: 10.1371/journal.pone.0208563 [published Online First: 2018/12/05]

Table 1 The general view of the ma	iternal.		
	Cohort population	Validation population	
Characteristics	(N=24,833)	(N=5009)	P
Age (years) *, N (%)			0.393
<25	1,309 (5.27%)	266 (5.31%)	
25-29	11,736 (47.26%)	2340 (46.71%)	
30-34	9,445 (38.03%)	1959 (39.11%)	
≥35	2,343 (9.44%)	444 (8.86%)	
Ethnicity, N (%)			0.983
Han	22,222 (89.49%)	4,475 (89%)	
Manchu	1,872 (7.54%)	389 (7.8%)	
Other ethnic groups	739 (2.98%)	145 (2.9%)	
Educational Attainment, N (%)			0.115
High school or below	8,635 (34.77%)	1,742 (35%)	
Bachelor's degree	13,639 (54.92%)	2,703 (54%)	
Postgraduate or higher	2,559 (10.30%)	564 (11%)	
Occupation, N (%)			0.777
Unemployed	11,373 (45.80%)	2,266 (45%)	
Light physical labor	2,825 (11.38%)	569 (11%)	
Moderate physical labor	10,011 (40.31%)	2,038 (41%)	
Heavy physical labor	624 (2.51%)	136 (2.7%)	
Family Per Capita Monthly			
Income (10,000 yuan), N (%)			0.9862
< 0.5	10,325 (41.58%)	2,080 (42%)	
0.5-2.0	9,534 (38.39%)	1,922 (38%)	
2.0-5.0	3,584 (14.43%)	720 (14%)	
>5.0	1,390 (5.60%)	287 (5.7%)	
Pre-pregnancy BMI (Kg/m2)			
*, N (%)			< 0.001
<18.5 (Underweight)	7,294 (29.37%)	1475 (29.45%)	
18.5-23.9 (Normal)	15,005 (60.42%)	2963 (59.16%)	
24.0-27.9 (Overweight)	1,700 (6.85%)	307 (6.13%)	
>28.0 (Obesity)	834 (3.36%)	264 (5.29%)	
Pregnancy History, N (%)			0.565
1	14,985 (60.34%)	3,005 (60%)	
2	6,513 (26.23%)	1,303 (26%)	
≥3	3,335 (13.43%)	701 (14%)	
Parity (number of deliveries), N			
(%)			0.775
0	20,550 (82.75%)	4,127 (82%)	
1	4,152 (16.72%)	853 (17%)	
≥2	131 (0.53%)	29 (0.6%)	
Gestational Age at Delivery			
(weeks), N (%)			0.434

<38	1,507 (6.07%)	296 (5.91%)	
38-40	13,023 (52.44%)	2589 (51.59%)	
>40	10,303 (41.49%)	2129 (42.50%)	
Blood Loss (ml)	393.54±92.53	413.48±124.65	0.081
Postpartum Hemorrhage, N (%)	1,623 (6.54%)	286 (5.71%)	0.032
Due to uterine atony, N (%)	1,225 (4.93%)	266 (5.31%)	0.279
Due to placental factors, N (%)	242 (0.97%)	43 (0.86%)	0.489
Due to Cancal Trauma, N (%)	139 (0.56%)	31 (0.62%)	0.686
Due to coagulation disorders, N			
(%)	76 (0.31%)	17 (0.34%)	0.705



d by copyright, including for 36/bmjopen-2024-089734 on

Table 2: Multivariate Risk Factor Analys	is for Postpartum Her	norrhage (PPH) within S	Subgroups
Characteristics	PPH	UA-PPH	PF-P
		~	

Characteristics	PPH	UA-PPH	PF-PPH	ू Е Т-РРН	CA-PPH
		General view		anu:	
Age (years)	*	*	-	- ary Eras	-
Ethnicity	**	**	**	* 202 d to	-
Educational Attainment	· -	-	-	5. D	-
Occupation	**	**	**	t ges	*
Family Per Capita Monthly Income (10,000 yuan)	, O _C	-	-	iloade chool d data	-
Pre-pregnancy BMI (Kg/m2)	**	**	**	mini **	*
Smoking	-	\ / In	-	ining,	*
Alcohol Consumption	-	(O)	-	ttp://bmjope	-
	Ob	stetric and Gynecologic	History	ain bm	
Pregnancy History	-	-	O	ing,	-
Parity (number of deliveries)	**	**	**	and *	*
History of miscarriage	-	-	<u> </u>	ν <mark>Ξ</mark>	-
Spontaneous abortion	-	-	-() ₆	com -	-
Induced abortion or medication				on lar tec	
abortion	-	-	-	chn -	-
Induced labor	-	-	_	ne -	-
Assisted reproductive technology	-	-	-	.com/ on June 7, 20	-
Gestational Age at Delivery (weeks)	**	**	*	025 8.	-
Diabetes	**	**	**	a * D	-
Hypertension	**	**	**)ера *	-
Anemia	**	**	**	# **	*

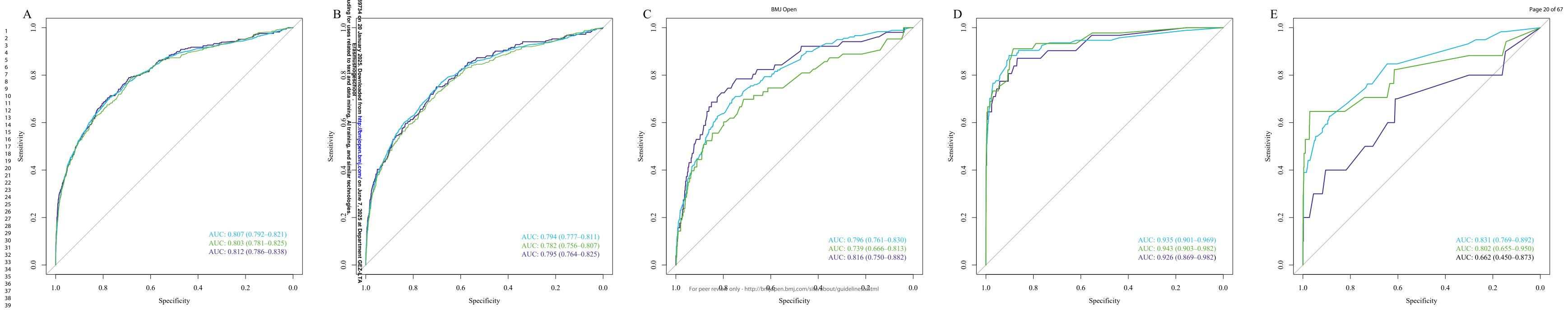
7		BMJ Open		6/bmjopen-20; by copyright,	
				n-2024-089: right, includ	
Coagulation disorder	**	-	-		*
Uterine fibroids/adenomyosis	**	**	**	for *	-
Polyhydramnios	**	**	*	20 Ja	-
Umbilical cord entanglement	-	-	-) Janu	-
Premature rupture of membra		**	**	Hary Hate	**
Placental abruption	**	**	-	2025 smus	-
Vaginal bleeding during preg	gnancy -	-	-	- 25. D sho	-
Scarred uterus	**	**	-	200 × -	*
	Deliver	y Process and Neonatal	Conditions	nlos scho	
Time of delivery		-	-	- lool . data r	-
Total duration of labor	**	*	*	5	**
First stage of labor - Latent p	ohase **	**	-		-
First stage of labor - Active p		**	-	, t -	-
Second stage of labor	**	**	*	** traii	-
Third stage of labor	**	-	**	njor *	-
Placental retention/Placenta	**	4.4		http://bmjopen.bmj.com/ y, Al training, and similar	didi
accreta/Placental implantation	** •n	**	**	nd si	**
Analgesia during labor	**	**	-0	imi.co *	**
Instrumental assistance in de	livery **	-	- 1	** ar tec	-
Lacerations of the cervix, vag	gina. or			ttp://bmjopen.bmj.com/ on June 7, 20	
perineum	**	-	-	June	-
Newborn weight (grams)	**	**	-	- 7, 2,	-
Newborn length (centimeters	-	-	-	 2025 es.	-
*: P<0.05; **: P<0.001				—— <u>a</u>	
- 1 0.00, 1 0.001				—— p	
				r t m	
				ıt Department GEZ-LTA	
				GEZ	_
				Ä	1
				> `	

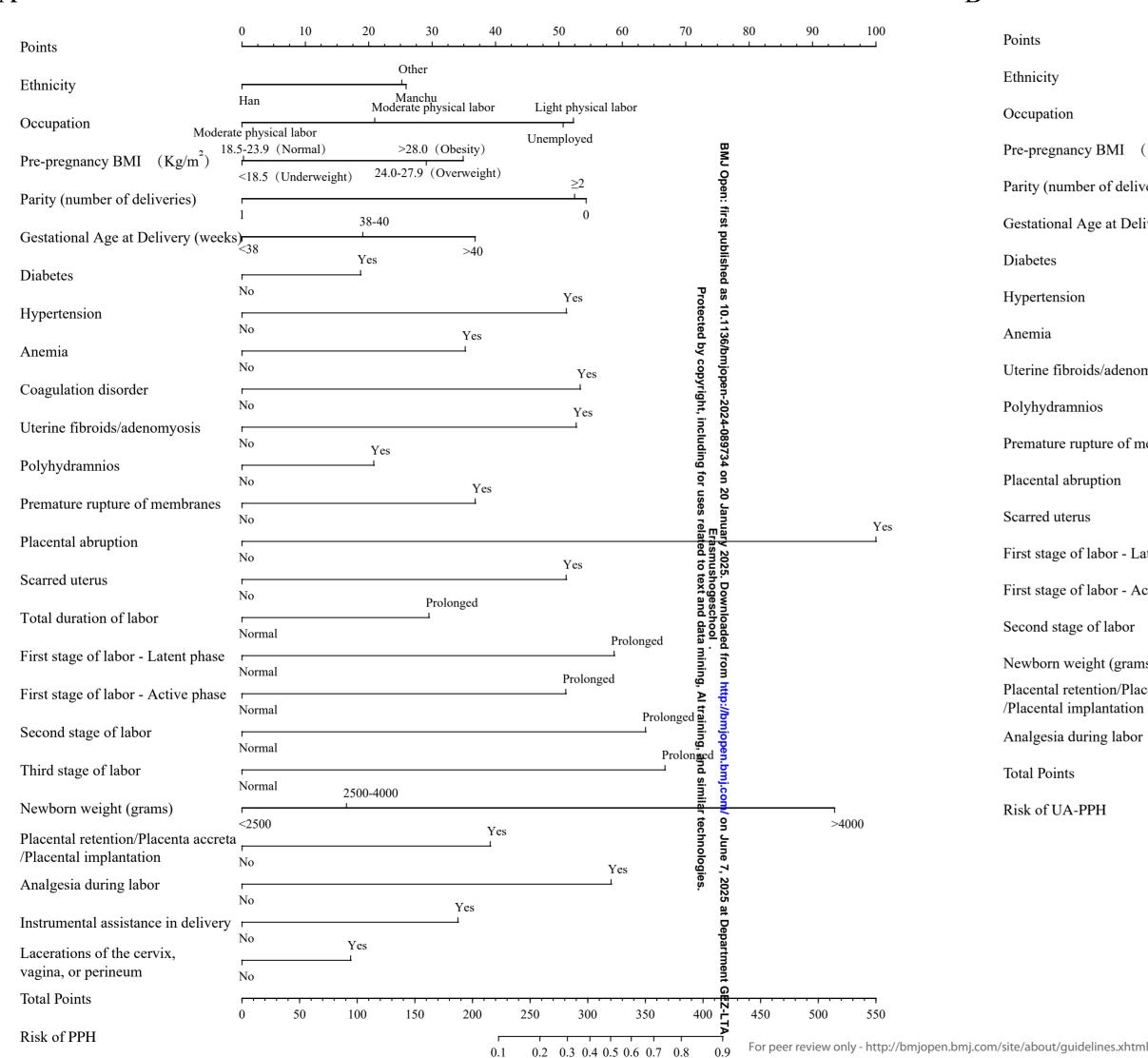
d by copyright, including for 36/bmjopen-2024-089734 on

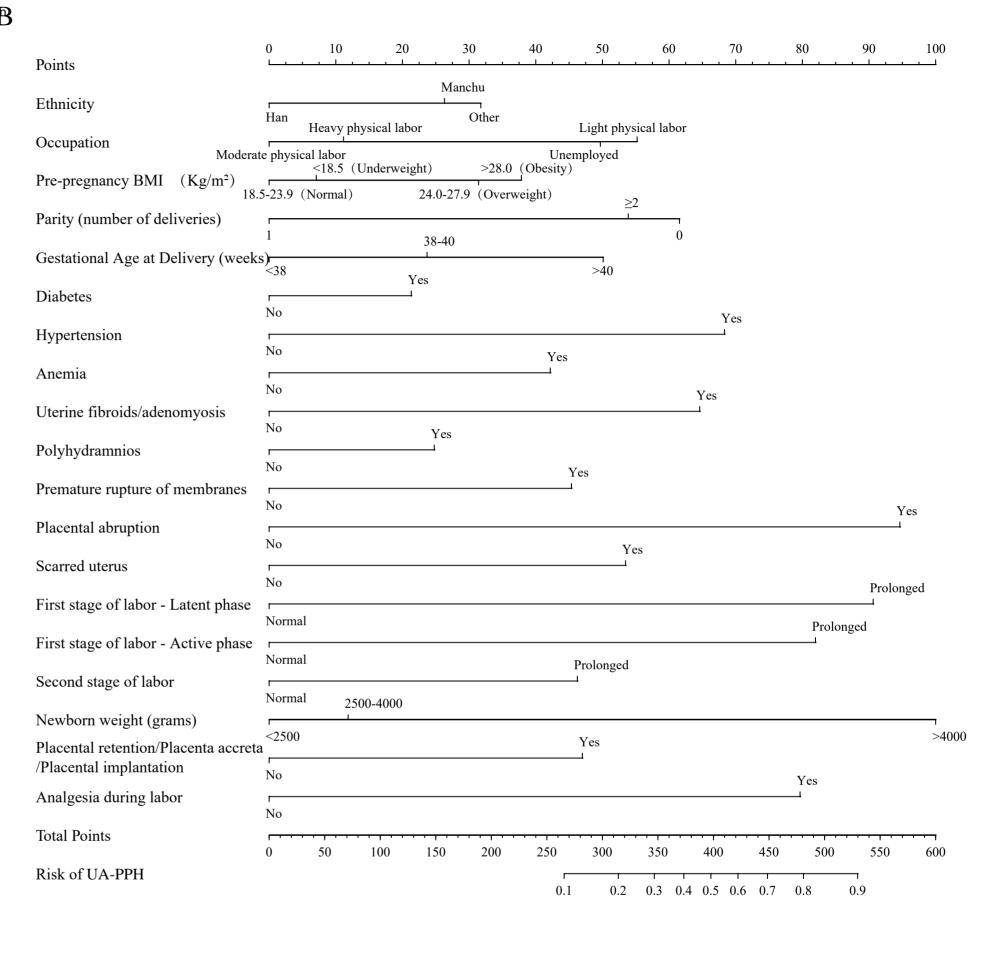
Figure legend

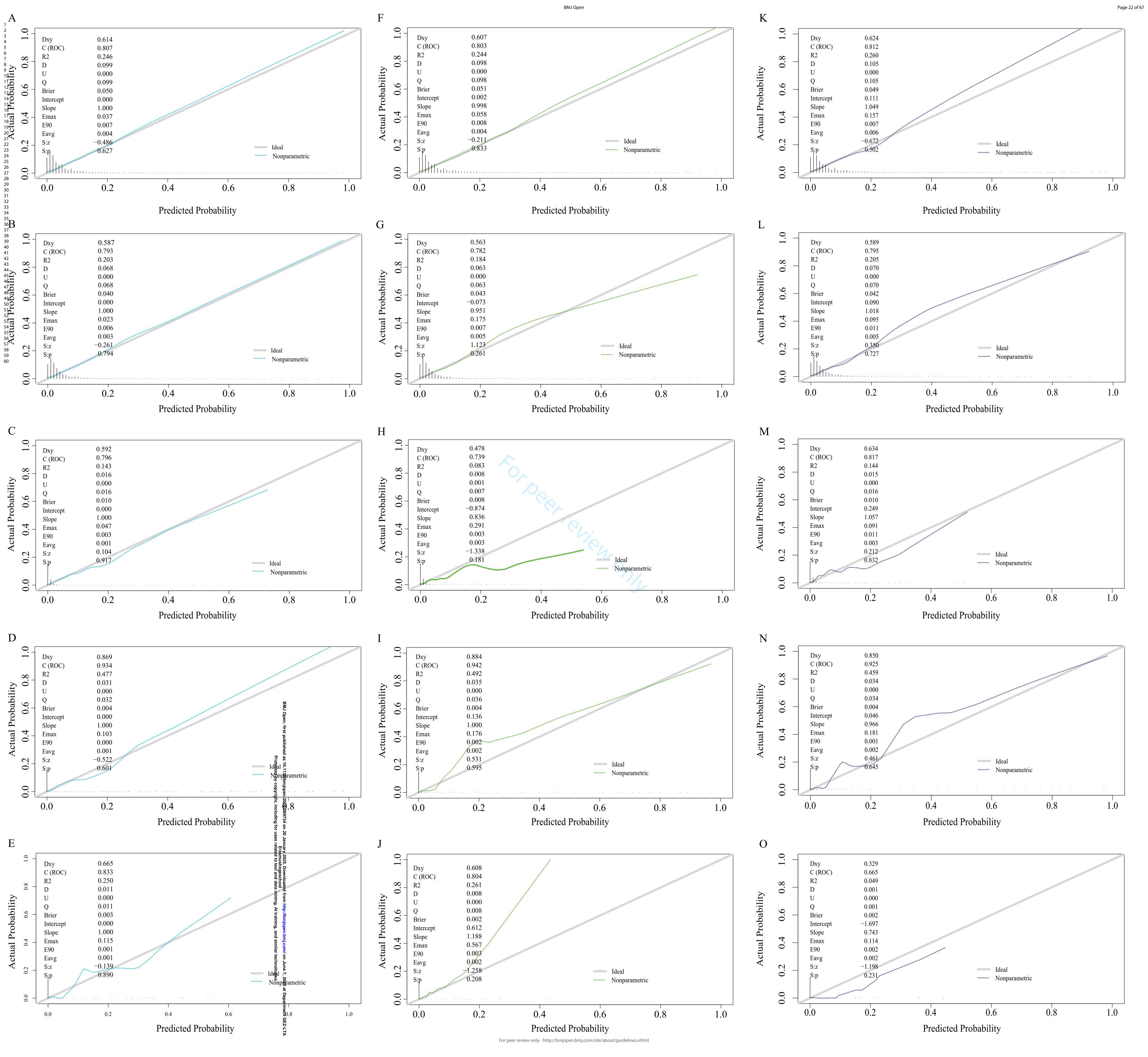
- Figure 1. Patient Selection Criteria Flowchart.
- Figure 2. AUC Curve for Postpartum Hemorrhage (PPH) Group and four subgroups. A. PPH Group; B. Uterine Atopy PPH Group; C. Placental Factors PPH Group; D. Cervical Trauma PPH Group; E. Coagulation Abnormalities PPH Group. The blue line signifies the training dataset, which is employed to evaluate the model's predictive capabilities following the training phases. The green line corresponds to the internal validation data votal for refining model parameters and for conducting initial assessments of the model's accuracy. The purple line denotes the external validation datase which is utilized to ascertain the model's generalizability and to verify its performance in an independent dataset. Figure 3. Nomograms for Postpartum Hemorrhage (PPH) and Uterine Atony PPH Group. A. PPH Group; B. Uterang Atony PPH Group.
- Supplemental Figure 1. Calibration curves for Postpartum Hemorrhage (PPH) Group and four subgroups. A PH Group; B\G\L. Uterine Atony PPH dataset; The green line signifies the internal validation dataset; The purple line signifies the external validation dataset.

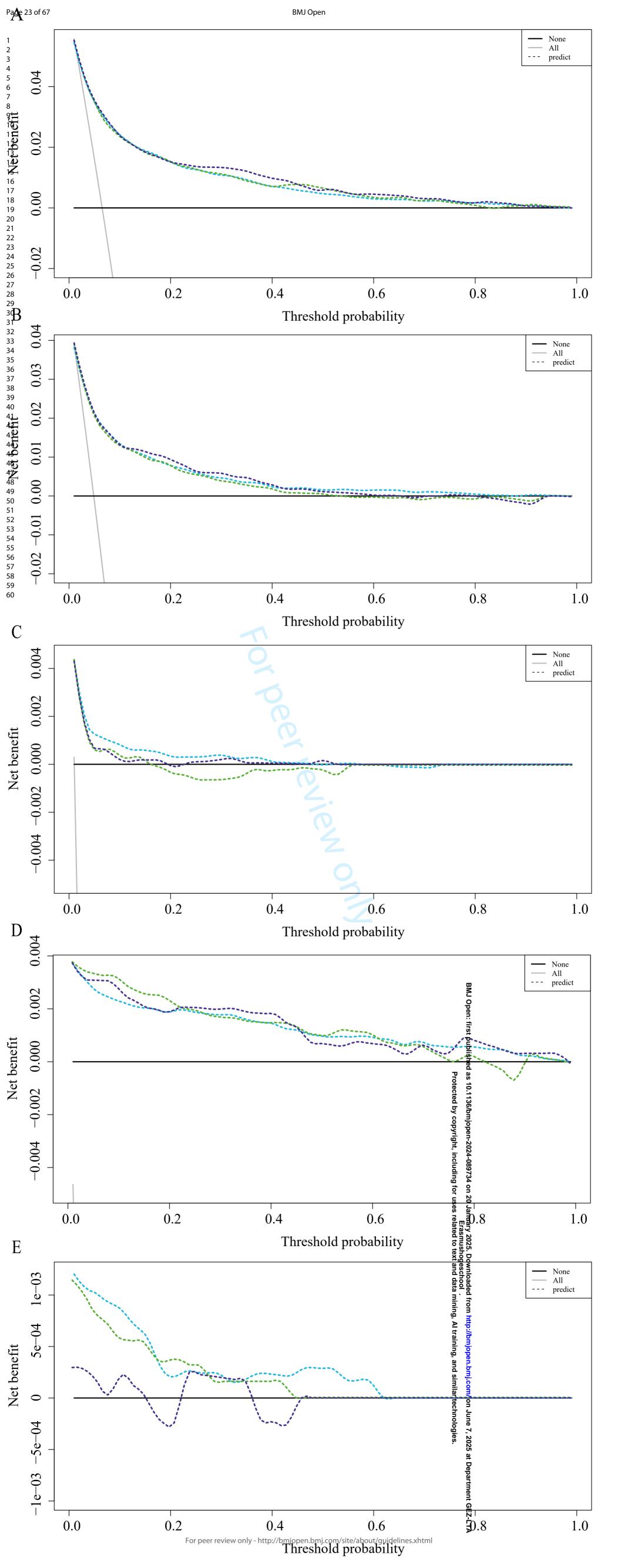
Supplemental Figure 2. Decision Curve Analysis for Postpartum Hemorrhage (PPH) Group and four subgroups. E PH Group; B. Uterine Atony PPH Group; C. Placental Factors PPH Group; D. Cervical Trauma PPH Group; E. Coagulation Abnormalities PPH Group. The blu in signifies the training dataset; The green training, and similar technologies line signifies the internal validation dataset; The purple line signifies the external validation dataset. bmjopen.bmj.com/ on June 7, 2025 at Department GEZ-LTA











Sunnlementer	v tahla 1	Comper	ison of	Rasje Cho	ractorist	ics and Dis	BMJ Open	eening between	Non-PPH o	36/bmjopen-2024-0897	trouns		
Supplementary		-PPH		PH	1 acter ist	ics and Kis	sk ractor Sci	cening between	TOII-I I II al	on for	roups		
Characteristi	(N=2	3,210)	(N=	1,623)			Univa	riate Logistic R	egression	~ 2	Multiv	ariate Logistic I	Regression
			Sam		χ^2	P				ശയ			
cs	Sampl e size	n%	ple size	n%			OR	95% CI	P	nuary 202 Erasmu related to	OR	95% CI	P
General view										5. D sho tex			
Age (years)					77.07	<0.001*				ownloaded geschool . and data !			
<25	1,251	5.39%	58	3.57%			Ref.			ded ol . ata r	Ref.		
25-29	11,047	47.60%	689	42.45%			1.35	1.03, 1.79	0.034*	mining,		0.85, 1.49	0.430
30-34	8,719	37.57%	726	44.73%			1.80	1.38, 2.39	<0.001**	بَةِ 1. 4 ِ5		1.11, 1.94	0.009*
≥35	2,193	9.45%	150	9.24%			1.48	1.09, 2.03	0.014*	≥ 1. 2 5		0.92, 1.72	0.169
						<0.001*				aini			
Ethnicity					86.05	*				ope ng,			
Han	20,848	89.82%	1,374	84.66%			Ref.			n.bi	Ref.		
Manchu	1,693	7.29%	179	11.03%			1.60	1.36, 1.88	<0.001**	<u>s</u> 1.59		1.34, 1.87	<0.001**
Other ethnic		• 0							U A	om/ iilar			
groups	669	2.88%	70	4.31%		0.00-*	1.59	1.22, 2.03	<0.001**	<u>e</u> 1.62		1.24, 2.07	<0.001**
Educational					20.46	<0.001*	D.C			tth./bmjopen.bran.com/on June	D. C		
Attainment					39.46		Ref.			9,7	Ref.		
High school or below	8,142	35.08%	493	30.38%			1.20	1.08 1.35	0.001*	202∯ >>s.		0.84, 1.07	0.392
Bachelor's	0,142	33.0070	473	30.3676			1.20	1.08, 1.35	0.001	0.90		0.04, 1.0/	0.374
degree	12,713	54.77%	926	57.05%			1.43	1.21, 1.69	<0.001**	2025 at Degar les. 1		0.88, 1.27	0.520
Postgraduate	2,355	10.15%	204	12.57%			1.15	1.21, 1.07	-0.001	artment GEZ-L		0.00, 1.27	0.520

>28.0										jopen-2024-089734 c opyright, including 1		
(Obesity)	725	3.12%	109	6.72%			2.43	1.93, 3.03	<0.001**	34 o § 20 Jai	1.93, 3.04	<0.001**
Smoking					8.56	0.036*				0 Ja		
No	23,128	99.65%	1,612	99.32%			Ref.					
Yes	82	0.35%	11	0.68%			1.92	0.97, 3.46	0.042*	nuas 2 Frasi related	0.93, 3.35	0.058
Alcohol										2025 smus ed to t		
Consumption					0.46	0.927				. Do hog lext		
No	23,153	99.75%	1,620	99.82%			-	-	-	wnlo Jesch	-	-
Yes	57	0.25%	3	0.18%			-	-	-		-	-
Obstetric and	Gynecolo	ogic Histor	'y							aded fool .		
Pregnancy						<0.001*				from http://Enjeps		
History					46.64	*	<i>/</i> -			P Ref.		
1	13,914	59.95%	1,071	65.99%			Ref.			P		
2	6,153	26.51%	360	22.18%			0.76	0.67, 0.86	<0.001**		0.88, 1.15	0.922
≥3	3,143	13.54%	192	11.83%			0.79	0.68, 0.93	0.004*	licepen.bmj.com/ on Ref.	0.96, 1.37	0.118
Parity						0 0 0 4 4				n.br and		
(number of						<0.001*				sim sim		
deliveries)	10.060	0.0		0.1.0.10./	203.39	_	5 .0			om/		
0	19,060	82.12%	1,490	91.81%			Ref.		0.00	Ref.		0 0 0 4 4 4
1	4,027	17.35%	125	7.70%			0.40	0.33, 0.48	<0.001**	0.40	0.32, 0.49	<0.001**
≥2	123	0.53%	8	0.49%			0.83	0.37, 1.60	0.615	Ref. Ref. O. O. 7, 20 and similar technologies	0.29, 1.39	0.331
history of					0.4.5					2025 ies.		
miscarriage	15.050	70. 7 00.	1 100	50.0 501	0.16	0.984				25 a		
No	17,078	73.58%	1,189	73.26%			-	-	-	at Dep	-	-
Yes	6,132	26.42%	434	26.74%	0.46	0.04=	-	-	-	par	-	-
spontaneous					0.36	0.947				tment GEZ-LTA		

BMJ Open

 Page 26 of 67

BMJ Open

Page 27 of 67

2 3 4

5 6

7

8

9 10

11

12

13

14

15

16

17

18 19

20

21

22 23

24

25

26 27

28

29

30

31 32

33

34

35

36

37

										897 Iudi		
						*				ıσω		
No	20,207	87.06%	1,300	80.10%			Ref.			Ref. Ref. 1. Leras 1 Eras		
Yes	3,003	12.94%	323	19.90%			1.67	1.47, 1.90	<0.001**	us 1.62 1.62	1.42, 1.84	<0.001**
						<0.001*				· —		
Hypertension					284.54	*				nuary 2 Erasr		
No	21,649	93.27%	1,385	85.34%			Ref.			ry 2025 rasmust		
Yes	1,561	6.73%	238	14.66%			2.38	2.05, 2.75	<0.001**	tex SA	2.30, 3.12	<0.001**
						<0.001*				t and		
Anemia					223.87	*				vnloade school		
No	19,381	83.50%	1,189	73.26%			Ref.			a −g Kei.		
Yes	3,829	16.50%	434	26.74%			1.85	1.65, 2.07	<0.001**	1. 8 5	1.73, 2.20	<0.001**
Coagulation						<0.001*				ing. The second		
disorder					122.64	*				AI t		
No	23,084	99.46%	1,588	97.84%			Ref.			f. Ref. Ref. Ref. Ref. Ref. Ref. Ref. Re		
Yes	126	0.54%	35	2.16%			4.04	2.73, 5.82	<0.001**	ā 3 🕏	2.60, 5.77	<0.001**
Uterine										en. an		
fibroids/aden						<0.001*				d si		
omyosis					249.58	*				mil:		
No	22,595	97.35%	1,501	92.48%			Ref.			Ref.		
Yes	615	2.65%	122	7.52%			2.99	2.43, 3.64	<0.001**	3.08	2.49, 3.79	<0.001**
Polyhydram						<0.001*				ne i		
nios					58.05	*				7, 2025 Ref.		
No	20,967	90.34%	1,399	86.20%			Ref.			Ref.		
Yes	2,243	9.66%	224	13.80%			1.50	1.29, 1.73	<0.001**	a⊗Departm	1.44, 1.96	<0.001**
Umbilical)epa		
cord					0.23	0.973				irt T		

BMJ Open

Page 29 of 67

2 3 4

5 6

8

9 10

11

12

13

14

15

16

17

18 19

20

21

22 23

24

25

26 27

28

29

30

31 32

33

34

35 36

37

5 6

7

8

9 10

11

12

13

14

15

16

17

18 19

20

21

22 23

24

25

26 27

28

29

30

31 32

33

34

35

36

37

										.0897		
Yes	2,071	8.92%	225	13.86%			1.64	1.41, 1.90	<0.001**	f. Re Re On 20 January 2025∕Downloaded from http:// Erasmushogeschool . 0 ing for uses related to text and data mining, AI t	1.26, 1.74	<0.001**
Newborn										on 2		
weight					1510.0	<0.001*				use:		
grams)					2	*				anu: B		
<2500	221	0.95%	7	0.43%			Ref.			ary Ref.		
2500-40										202! d to		
00	21,944	94.55%	1,281	78.93%			1.84	0.94, 4.34	0.112	es S €	0.84, 3.94	0.187
>4000	1,045	4.50%	335	20.64%			10.1	5.09, 24.0	<0.001**	oge Ges tan	4.68, 22.40	<0.001**
Newborn										iloai cho d da		
ength						<0.001*				ded ol.		
centimeters)					52.57	* (*),				fror		
≤55	22,566	97.23%	1,542	95.01%			Ref.			Ref.		
>55	644	2.77%	81	4.99%			1.84	1.44, 2.32	<0.001**	00.//bm	0.63, 1.08	0.172
										翰///bmjopen.bmj.com/ on June 7, 2025 at Department GEZ-LTA O Al training, and similar technologies.		
				For peer	review or	nly - http://b	mjopen.bmj	.com/site/about/	guidelines.xht			

36/bmjopen-2024-0897 d by copyright, includi

	Non-U	A-PPH	UA	-PPH						on 2		
Characteristi	(N=2	3,608)	(N=	1,225)			Univa	riate Logistic R	egression	us es Multi	variate Logistic l	Regression
cs characteristi			Sam		χ^2	P				anu: Fel		
CS	Sampl		ple							OR nuary 202 Erasmu related to		
	e size	n%	size	n%			OR	95% CI	P	5 2 8 OR	95% CI	P
General view										M R O 1 20 January 2025. Do Erasmushog or uses related to text		
						<0.001*						
Age (years)					67.01	*				wnloaded frems eschool . 1. and data mining,		
<25	1,269	5.38%	40	3.27%			Ref.			Ref.		
25-29	11,212	47.49%	524	42.78%			1.48	1.08, 2.09	0.018*	mi 1. 2 31	0.88, 1.71	0.266
30-34	8,891	37.66%	554	45.22%			1.98	1.45, 2.78	<0.001**	 1	1.13, 2.19	0.009*
≥35	2,236	9.47%	107	8.73%			1.52	1.06, 2.22	0.027*	<u>≥</u> 1. 2 4	0.86, 1.82	0.259
						<0.001*				/bm rain		
Ethnicity					39.45	*				ing		
Han	21,172	89.68%	1,050	85.71%			Ref.			Ref.		
Manchu	1,749	7.41%	123	10.04%			1.42	1.16, 1.71	<0.001**	<u>യ്.</u> 1. 2 40	1.15, 1.70	<0.001**
Other ethnic										mil _s		
groups	687	2.91%	52	4.24%			1.53	1.13, 2.02	0.004*	1.65	1.15, 2.05	0.003*
Educational						<0.001*				ef. R R নি নি নি Inining, Al training, and similar technologies		
Attainment					54.87	*				ne olo		
High school										7, 2()gies		
or below	8,278	35.06%	357	29.14%			Ref.			. % Ref.		
Bachelor's										at D		
degree	12,939	54.81%	700	57.14%			1.25	1.10, 1.43	<0.001**	0. 2 9	0.86, 1.13	0.831
Postgraduate	2,391	10.13%	168	13.71%			1.63	1.35, 1.96	<0.001**	1. #	0.98, 1.47	0.075

5 6

7

8

9 10

11

12

13

14

15

16

17

18 19

20

21

22 23

24

25

26

27

28

29

30

31 32

33

34

35 36

37

38

44 45 46 18.5-23.9

14,339

60.74%

54.37%

666

Page 34 of 67

0.82, 1.07

0.326

ent GEZ-LTA

0.82, 1.07

0.313

0.94

BMJ Open

Page 35 of 67

2 3 4

5 6

7

8

9 10

11

12

13

14 15

16

17

18 19

20

21

22 23

24

25

26

27

28

29

30

31 32

33

34

35

36

37

							BMJ Open			6/bmjopen-2024-089734 on 20 Ja by copyright, including for uses		
miscarriage										-08973 <i>4</i>		
No	17,369	73.57%	898	73.31%			-	_	-	for -	_	_
Yes	6,239	26.43%	327	26.69%			-	-	-	20 . use	-	-
spontaneous										_		
abortion					1.55	0.671				າuary 2 Erasn related		
No	21,228	89.92%	1,111	90.69%			-	-	-	y 2025 asmus	-	-
Yes	2,380	10.08%	114	9.31%			-	-	-	≅ 	-	-
induced										ownl gescl		
abortion or										nload choo id dat		
medication										ded ol .		
abortion					2.07	0.559				fror ninii		
No	19,011	80.53%	972	79.35%			/ -	-	-	ng,	-	-
Yes	4,597	19.47%	253	20.65%			10,	-	-	A tr .	-	-
induced					10.41	0.015*				ai bi ni		
labor							Ref.			Ref.		
No	23,115	97.91%	1,211	98.86%			0.74		0.00.74	n.br		
Yes	493	2.09%	14	1.14%			0.54	0.30, 0.89	0.025*	<u>s</u> i 1. 0 9	0.59, 1.88	0.774
assisted										om/ iilar		
reproductive					6.77	0.080				on tech		
technology No	22,981	97.34%	1,203	98.20%	0.77	0.080				r Downloaded from http://bmjopen.b∰.com/ on June 7, 20 nogeschool . ∍xt and data mining, Al training, and similar technologies		
Yes	627	2.66%	22	1.80%			_	_	_	ogie	_	_
Gestational	027	2.0070	22	1.0070						r, 2025 9gies.		
Age at												
Delivery						<0.001*				at Department GEZ-LTA		
(weeks)					189.66	*				artn		

5

6

8

9 10

11

12

13

14 15

16

17

18 19

20

21

22 23

24

25

26 27

28

29

30

31 32

33

34

35 36

37

BMJ Open

2 3 4

5

6

8

9 10

11

12

13 14

15

16

17

18 19

20

21

22 23

24

25

26 27

28

29

30

31

32

33

34

35 36

37

38

44 45 46 Page 38 of 67

BMJ Open

2 3 4

5 6

8

9 10

11

12

13

14

15

16

17

18 19

20

21

22 23

24

25

26

27

28

29

30

31 32

33

34

35 36

37

44 45 46 Page 40 of 67

Supplementary							BMJ Open		;	6/bmjopen-2024-0897 by copyright, includi		
Supplementary	y table 3	Compar F-PPH	ison of	Basic Cha -PPH	racterist	ics and Risk	Factor Scree	ening between l	Non-PF-PPH (and KF-PPH g 한 음	roups	
		4,591)		(=242)			Univari	ate Logistic Re		~ N	ariate Logistic R	egression
Characteristi		1,0010	Sam	2127	χ^2	P	01111111	are Logistic Ite	81 0351011	Janu es re	ilimie Logistie It	ogi ession
cs	Sampl		ple		λ.					anuary 2025 Erasmus		
	e size	n%	size	n%			OR	95% CI	P	or September 1981	95% CI	P
General view										te sh D		
Age (years)				11.46	0.120					ownloaded		
<25	1,297	5.27%	12	4.96%			-	-	-	- Noa Cho d da	-	-
25-29	11,639	47.33%	97	40.08%			-	-	-	oaded	-	-
30-34	9,340	37.98%	105	43.39%			-	-	-	nini	-	-
≥35	2,315	9.41%	28	11.57%			/ -	-	- (d from ht	-	-
						<0.001*				A t		
Ethnicity					29.18	*				/bm		
Han	22,023	89.56%	199	82.23%			Ref.			Ref.		
Manchu	1,839	7.48%	33	13.64%			1.99	1.35, 2.84	<0.001**	1.94	1.32, 2.78	<0.001**
Other ethnic	729	2.96%	10	4.13%			1.52	0.75, 2.72	0.201	ttp://bmjopen.bmj.com/ on June 7, 20	0.76.2.79	0.181
groups	129	4.90%	10	4.13%			1.32	0.75, 2.73	0.201	nila B 1.33	0.76, 2.78	0.181
Educational					3.44	0.632				r teg		
Attainment					3.44	0.032				L Ju		
High school	8,559	34.81%	76	31.40%						ne 7 olog		
or below	0,339	34.0170	70	31.4070			-	-	- '	, 20 , 20 jies	-	-
Bachelor's	13,496	54.88%	143	59.09%						r on http://bmjopen.bmj.com/ on June 7, 2025 at Departmining, Al training, and similar technologies.		
degree	13,490	34.0070	143	37.0770			-	-	-	# - □	-	-
Postgraduate	2,536	10.31%	23	9.50%						ера		
or higher	2,330	10.5170	23	9.3070			-	-	_	rtment	_	-

	BMJ Open									36/bmjopen-2024-089734 d by copyright, including		
					77.00	<0.001*				ing 73		
Occupation Unemployed	11,218	45.62%	155	64.05%			Ref.			n 20 Daf		
Light physical labor	2,796	11.37%	29	11.98%			0.75	0.49, 1.10	0.159	Ref. Ref. 14 on 20 January 20 Erasn g for uses related	0.49, 1.08	0.137
Moderate physical labor	9,955	40.48%	56	23.14%			0.41	0.30, 0.55	<0.001**	025 to te	0.30, 0.55	<0.001**
Heavy physical labor	622	2.53%	2	0.83%			0.23	0.04, 0.73	0.041*	5. Downloaded shogeschool .	0.04, 0.71	0.038*
Family Per										paded nool .		
Capita										nini		
Monthly					2.01	0.959				ng,		
Income										Al tı		
(10,000 yuan)										aini		
< 0.5	10,222	41.57%	103	42.56%			-	10 ;	-	ing,	-	-
0.5-2.0	9,440	38.39%	94	38.84%			-	1/1/	-	n.b	-	-
2.0-5.0	3,549	14.43%	35	14.46%			-	-	-	sin 3.	-	-
>5.0	1,380	5.61%	10	4.13%			-	-		nila	-	-
Pre-pregnan cy BMI					64.09	<0.001*				I from http://bmjopen.bmj.com/ on June 7, 20 mining, Al training, and similar technologies.		
(Kg/m2)										ne 7		
<18.5										7, 2025 Ref.		
(Underweig ht)	7,235	29.42%	59	24.38%			Ref.			% Ref.		
18.5-23.9 (Normal)	14,873	60.48%	132	54.55%			1.09	0.80, 1.49	0.590	at Department GEZ	0.81, 1.50	0.578

										jop Op		
										njopen-2024-0897 copyright, includi		
										024-1 t, inc		
										0897.		
24.0-27.9										ing for 2.45		
(Overweight	1,667	6.78%	33	13.64%			2.43	1.56, 3.70	<0.001**	on 2.45	1.58, 3.75	<0.001
)										Ja		
>28.0	816	3.32%	18	7.44%			2.71	1.54, 4.51	<0.001**	January 2 Erası Erası res related	1.55, 4.55	<0.001
(Obesity)					2 (0	0.444				ry 2 rasr rasr		
Smoking	24.500	00.6207	2.40	00.150/	2.68	0.444				2025 smus		
No	24,500	99.63%	240	99.17%			-	-	-	hog	-	-
Yes	91	0.37%	2	0.83%			-	-	-	Downloa	-	-
Alcohol					0.60	0.897	-	-	-	oade hool data	-	_
Consumption	24.522	00.760/	241	00.500/								
No V	24,532	99.76%	241	99.59%						in rom		
Yes	59 C	0.24%	1	0.41%						g, A		
Obstetric and	Gynecolo	gic Histor	y			<0.001*				p://b		
Pregnancy History					14.33	~0.001 *				in in jo		
History 1	14,822	60.27%	163	67.36%	14.33		Ref.			رق کے Ref.		
2	6,454	26.25%	59	24.38%			0.83	0.61, 1.11	0.226	nd 1 81	0.80, 1.51	0.519
≥ ≥3	3,315	13.48%	20	8.26%			0.83	0.33, 0.85	0.012*	Si . 1. S i.	0.47, 1.28	0.319
Parity	3,313	13.40/0	20	0.2070			0.55	0.55, 0.65	0.012	llar 1	0.47, 1.20	0.507
(number of						<0.001*				on J		
deliveries)					39.16	*				lune		
0	20,324	82.65%	226	93.39%	37.10		Ref.			f. Ref. Ref. Ref. Ref. Ref. Ref. Ref. Re		
1	4,137	16.82%	15	6.20%			0.33	0.18, 0.53	<0.001**	s. 0 %	0.20, 0.63	<0.001**
≥2	130	0.53%	1	0.41%			0.69	0.04, 3.11	0.714	0. 3 0	0.05, 4.55	0.933
history of	100	0.0070	•	0.11/0			···	,	V., - 1	•	,	0.555
					0.17	0.982				əpartm		

 Page 44 of 67

							ымь орен		3	njopen		
										mjopen-2024-089734 on 20 Ja		
No	18,087	73.55%	180	74.38%			_	-	- (9734 9734	_	
Yes	6,504	26.45%	62	25.62%			-	-	_ :	on 20 January 2025 Erasmus	-	-
spontaneous										20 J		
abortion					0.04	0.998				lant S re		
No	22,122	89.96%	217	89.67%			-	-	-	uary 2 Erasr	-	_
Yes	2,469	10.04%	25	10.33%			-	-		y 2025 asmus	-	-
induced												
abortion or										. Downloaded from http://bmjopen.bmj.com/ on June hogeschool .		
medication												
abortion					0.27	0.965				oadec 1001		
No	19,786	80.46%	197	81.40%			-	-	-	mir de fro	-	-
Yes	4,805	19.54%	45	18.60%			<u> </u>	-	- 0		-	-
induced										<u>≥</u> #		
labor					6.48	0.090						
No	24,085	97.94%	241	99.59%			-	0:	-	I from http://bmjopen.bmj.com/ on June 7, 20	-	-
Yes	506	2.06%	1	0.41%			-	C1/	-	_ <u> </u>	-	-
assisted												
reproductive												
technology					2.34	0.504			1/1/	ar 2 t 0		
No	23,951	97.40%	233	96.28%			-	-		ה ה ה	-	-
Yes	640	2.60%	9	3.72%			-	-	-		-	-
Gestational									9	7, 2 gies		
Age at									:	025		
Delivery						<0.001*				7, 2025 at D		
(weeks)					26.69	*				Department GEZ-LTA		
<38	1,499	6.10%	8	3.31%			Ref.			Ref.		

Page 45 of 67

										njopen-2024-0897 		
										4-0897 nclud		
38-40	12,916	52.52%	107	44.21%			1.55	0.81, 3.47	0.231		0.98, 4.24	0.085
>40	10,176	41.38%	127	52.48%			2.34	1.22, 5.21	0.020*	on 20 January 2825 3 Erasmus for uses related to t	1.60, 6.97	0.002*
						<0.001*				Ses of		
Diabetes					30.50	*				s rel		
No	21,318	86.69%	189	78.10%			Ref.			ated Ref.		
Yes	3,273	13.31%	53	21.90%			1.83	1.33, 2.46	<0.001**	2025.	1.28, 2.38	<0.001**
						<0.001*				~ ~ .		
hypertension					26.00	*				Downl Downl hogesc		
No	22,824	92.81%	210	86.78%			Ref.			d Choa Ref.		
Yes	1,767	7.19%	32	13.22%			1.97	1.33, 2.82	<0.001**	oaded data r	1.43, 3.06	<0.001**
						<0.001*				f. ef. Re		
anemia					29.60	*				n <mark>n</mark>		
No	20,392	82.92%	178	73.55%			Ref.			Ref.		
Yes	4,199	17.08%	64	26.45%			1.75	1.30, 2.31	<0.001**	a i. 1. 3	1.33, 2.39	<0.001**
coagulation										ing,		
disorder					0.24	0.971				and and		
No	24,432	99.35%	240	99.17%			-	-	-	d sii -	-	-
Yes	159	0.65%	2	0.83%			-	-	O _A	nila	-	-
uterine										r te		
fibroids/aden						<0.001*				chn Ju		
omyosis					27.94	*				ne 7		
No	23,871	97.07%	225	92.98%			Ref.			9 Ref. 2.21		
Yes	720	2.93%	17	7.02%			2.50	1.47, 4.00	<0.001**	· 2 10/21	1.42, 3.92	<0.001**
polyhydramn										at Department GEZ-LTA		
ios					9.25	0.026*				ера		
No	22,158	90.11%	208	85.95%			Ref.			₹ Ref.		

 Page 46 of 67

Page 47 of 67

2 3 4

5

6

7

8

9 10

11

12

13

14 15

16

17

18 19

20

21

22 23

24

25

26 27

28

29

30

31

32

33

34

35 36

37

2 3 4

5 6

7

8

9 10

11

12

13

14 15

16

17

18 19

20

21

22 23

24

25

26 27

28

29

30

31 32

33

34

35 36

37

38

44 45 46 Page 48 of 67

5 6

7

8

9 10

11

12

13

14 15

16

17

18 19

20

21

22 23

24

25

26 27

28

29

30

31 32

33

34

35 36

37

milar technologies.

com/ on June 7, 2025 at Department GEZ-LTA

0.860

0.059

vagina, or										9734 on	
perineum										for on ;	
No	22,326	90.79%	211	87.19%			-	-	-	20 Ja	-
Yes	2,265	9.21%	31	12.81%			-	-	-	- Du	-
Newborn										uary : Eras related	
weight						<0.001*				2025 smus	
(grams)					212.47	*					
<2500	226	0.92%	2	0.83%			Ref.			Ref. Download&di hogeschoot. ext and data m	
2500-40										nloa ichc id d	
00	23,035	93.67%	190	78.51%			0.93	0.30, 5.65	0.922		0.27, 5.39
>4000	1,330	5.41%	50	20.66%			4.25	1.31, 26.1	<i>0.046</i> *	min 4.68 nin m	1.21, 24.90
Newborn										ing, m	
length										≱ <mark>#</mark>	
(centimeters)					1.10	0.777				//bm	
≤55	23,875	97.09%	233	96.28%			_	10:	-	http://bmjope 3, Al training,	-
>55	716	2.91%	9	3.72%			-		-	deom http://bmjopen.bmj . 4 mining, Al training, and si	-
										d si	

							BMJ Open			PH P- T- 36/bmjopen-2024-089734 o d d by copyright, including fu		
Supplementary	y table 4	Compar	ison of	Basic Cha	racterist	ics and Ris	sk Factor Scr	eening between	Non-CT-PP	E S Hagnd LCT-PPH ⇒ O	I groups	
	Non-C	T-PPH	Ci	I'-PPH						윽 ㅋ		
Characteristi		4,694)	Sam	(=139)	χ^2	P	Univa	iate Logistic R	egression	is a	variate Logistic l	Kegression
cs	Sampl		ple		χ	Γ				Inuary 20 Erasn		
	e size	n%	size	n%			OR	95% CI	P	2025. Do ed to text	95% CI	P
General view										15. D		
Age (years)					11.23	0.129				ownloaded geschool .		
<25	1,303	5.28%	6	4.32%			-	-	-	ا اloa cho d da	-	-
25-29	11,682	47.31%	54	38.85%			-	-	-	oaded	-	-
30-34	9,379	37.98%	66	47.48%			-	-	-	from h	-	-
≥35	2,330	9.44%	13	9.35%			<u> </u>	-	-	from http://bmjope	-	-
						<0.001*				A t		
Ethnicity					24.67	*				/bm rain		
Han	22,109	89.53%	113	81.29%			Ref.			Ref.		
Manchu	1,856	7.52%	16	11.51%			1.69	0.96, 2.77	0.051	an 1. <mark>6</mark> 7	0.95, 2.75	0.057
Other ethnic										y sir		
groups	729	2.95%	10	7.19%			2.68	1.31, 4.89	0.003*	ef. R R R 1.2.2.1 Al training, and similar technologies	1.32, 4.95	0.003*
Educational										/ on		
Attainment					4.25	0.515				hn Jur		
High school										ne 7 plog		
or below	8,585	34.77%	50	35.97%			-	-	-	7, 2025 ogies.	-	-
Bachelor's										25 a		
degree	13,569	54.95%	70	50.36%			-	-	-	at Departm	-	-
Postgraduate										≯par		
or higher	2,540	10.29%	19	13.67%			-	-	-	-tment	-	-

Occupation					10.34	0.170				jopen-2024-089734 on 20 Jaı opyright, including for uses		
Unemployed	11,307	45.79%	66	47.48%			_	_	-	on .	_	_
Light physical	,									20 c		
labor	2,802	11.35%	23	16.55%			_	_	-	Janu	-	_
Moderate										January 2 Erasr ses related		
physical labor	9,963	40.35%	48	34.53%			_	-	-	2025. smusl	-	_
Heavy										o - -		
physical labor	622	2.52%	2	1.44%			_	_	-	Downlo	-	_
Family Per												
Capita										aded ool . data .		
Monthly										from http://bmjopen.bmj.com/ on June 7, 20		
Income										ing,		
(10,000 yuan)					4.07	0.772				http://bmjopen.bmj.com/ 3, Al training, and similar		
< 0.5	10,261	41.55%	64	46.04%				<u> </u>	-	//bm	-	-
0.5-2.0	9,481	38.39%	53	38.13%			-	10:	-	ing -	-	-
2.0-5.0	3,569	14.45%	15	10.79%			-	1	-	en.t	-	-
>5.0	1,383	5.60%	7	5.04%			-	- /-	-	d si	-	-
Pre-pregnan										con mila		
cy BMI						<0.001*				n tec		
(Kg/m2)					270.37	*				1 June		
<18.5										ne l		
(Underweig										7, 20 gies		
ht)	7,274	29.46%	20	14.39%			Ref.			Ref.		
18.5-23.9										at D		
(Normal)	14,939	60.50%	66	47.48%			1.61	0.99, 2.72	0.064	1.6.	0.99, 2.72	0.064
24.0-27.9	1,671	6.77%	29	20.86%			6.31	3.58, 11.30	<0.001**	6. 3 3	3.59, 11.40	<0.001**

 Page 52 of 67

Page 53 of 67

2 3 4

5 6 7

8

9 10

11

12

13

14 15

16

17

18 19

20

21

22 23

24

25

26 27

28

29

30

31 32

33

34

35

36

37

							BMJ Open			/bmjop			
										pen-20 yright			
										5/bmjopen-2024-089734 on 20 Ja by copyright, including for uses			
miscarriage										9734 ding			
No	18,167	73.57%	100	71.94%			-	-	-	for	-	-	-
Yes	6,527	26.43%	39	28.06%			-	-	-	20 J use	-	-	-
spontaneous										_			
abortion					0.67	0.881				nuary Eras relate			
No	22,216	89.97%	123	88.49%			-	-	-	202 d to	-	-	-
Yes	2,478	10.03%	16	11.51%			-	-	-	5. D sho tex	-	-	-
induced										Download ogeschoo xt and dat			
abortion or										าloa cho d da			
medication										ded ol .			
abortion					0.07	0.996				mini			
No	19,872	80.47%	111	79.86%			<u> </u>	-	-	ing,	-	-	-
Yes	4,822	19.53%	28	20.14%			10.	-	-	from http://bmjopen.bmj.com/ on June 7, 20 nining, Al training, and similar technologies	-	-	-
induced										/bm			
labor					0.51	0.917				ing			
No	24,189	97.95%	137	98.56%			-		-	, an	-	-	-
Yes	505	2.05%	2	1.44%			-	- /	-	d si	-	-	-
assisted										con mila			
reproductive										ur te			
technology					1.52	0.679				chn			
No	24,047	97.38%	137	98.56%			-	-	- =		-	-	-
Yes	647	2.62%	2	1.44%			-	-	-	7, 2025 gies.	-	-	-
Gestational										. 25			
Age at										at D			
Delivery						<0.001*				t Department GEZ-L			
(weeks)					22.32	*				ir tr			

 Page 54 of 67

5 6

7

8

9 10

11

12

13

14 15

16

17

18 19

20

21

22 23

24

25

26 27

28

29

30

31 32

33

34

35 36

37

38

umbilical										jopen-2024-089734 on 20 Ja opyright, including for uses		
cord										on 20 for us		
entanglement					4.01	0.261				0 Ja		
No	17,292	70.03%	105	75.54%			-	-	-		-	-
Yes	7,402	29.97%	34	24.46%			-	-	-		-	-
premature										2025. smus		
rupture of						<0.001*				® ⊅ .		
membranes	10.207	77.700/	0.1	65.450/	24.20		D (ext and diagrams		
No	19,207	77.78%	91	65.47%			Ref.	1.20. 2.61	-0.001**		1 10 2 41	0.002*
Yes	5,487	22.22%	48	34.53%			1.85	1.29, 2.61	<0.001**	. O.	1.19, 2.41	0.003*
placental					4.01	0.260				y from h mining,		
abruption	24 (47	00.010/	120	00.200/	4.01	0.260				g, A		
No Yes	24,647 47	99.81% 0.19%	138	99.28% 0.72%			(9)	-	-	p://b	-	-
	47	0.1970	1	0.7270					-	inir -	-	-
vaginal bleeding										og, a		
during										nd:		
pregnancy					0.28	0.964				http://bmjopen.bmj.com/ on June 7, 20 , Al training, and similar technologies.		
No	23,445	94.94%	131	94.24%	0.20	0.704	_	_		lar t	_	_
Yes	1,249	5.06%	8	5.76%			_	_	<u> </u>	on J	_	_
scarred	1,219	3.0070	O	3.7070						June		
uterus					0.41	0.937				7, 2 ogie		
No	24,436	98.96%	137	98.56%			_	-	_	2025 les.	_	_
Yes	258	1.04%	2	1.44%			-	_	-	at Dep	_	_
Delivery Proce										Dep		
Time of					0.30	0.960				partm		

 Page 56 of 67

5 6

7

8

9 10

11

12

13

14 15

16

17

18 19

20

21

22 23

24

25

26

27

28

29

30

31 32

33

34

35 36

37

44 45 46 Page 58 of 67

by copyright, 36/bmjopen-202

)24-08 , inclu		
perineum										9734 Jaing		
No	22,506	91.14%	31	22.30%			Ref.			of S Ref.		
Yes	2,188	8.86%	108	77.70%			35.82	24.3, 54.4	<0.001**	us 37 2 42	23.80, 61.01	<0.001**
Newborn										Janu Ps re		
weight						<0.001*				lary Eras		
(grams)					669.48	*				202 Smu d to		
<2500	227	0.92%	1	0.72%			Ref.			sho Ref.		
2500-40										owr ges		
00	23,144	93.72%	81	58.27%			0.79	0.18, 14.0	0.820	d မေ ရွာ စီး	0.12, 12.3	0.689
>4000	1,323	5.36%	57	41.01%			9.78	2.14, 173	0.024*	: <u> </u>	1.15, 120	0.089
Newborn										fror		
length						<0.001*				n ht		
(centimeters)					18.02	*				A t		
≤55	23,979	97.10%	129	92.81%			Ref.			Ref.		
>55	715	2.90%	10	7.19%			2.60	1.27, 4.72	0.004*	ng, ar	0.47, 2.50	0.762
										ef. R 2 124-089734 on 20UJanuary 2025. Downloaded from http://bmjöpen.bmj.com/ on June 7, 2025 at Department GEZ-LTA S Erasmushogeschoob. , including for uses related to text and data mining, Al training, and similar technologies.		
				For peer	review or	nly - http://l	bmjopen.bmj.	com/site/about/	′guidelines.xh			

							BMJ Open			6/bmjopen-2 l by copyrigh		
Supplementary Characteristi	Non-C	Compar A-PPH 4,757)	CA	Basic Cha A-PPH N=76)	racteristi χ²	cs and Risk P		ening between No iate Logistic Reg		or c	groups uriate Logistic Re	gression
cs	Sampl		ple		λ.	1				uary Erag		
	e size	n%	size	n%			OR	95% CI	P	ö m 20 OR	95% CI	P
General view										5. D sho tex		
Age (years)					6.09	0.530				own ges t an		
<25	1,308	5.28%	1	1.32%			-	-	-	lload choo	-	-
25-29	11,702	47.27%	34	44.74%			-	-	-	ded ol . nta n	-	-
30-34	9,412	38.02%	33	43.42%			-	-	-	fror	-	-
≥35	2,335	9.43%	8	10.53%			<u> </u>	-	-	ng,	-	-
Ethnicity					6.92	0.227				:tp:// Al tı		
Han	22,158	89.50%	64	84.21%				-	-	/bm/	-	-
Manchu	1,862	7.52%	10	13.16%			-	10.	-	ing, -	-	-
Other ethnic										n.b anc		
groups	737	2.98%	2	2.63%			-	- /	-	isin ∃.	-	-
Educational										iom, nila		
Attainment					1.65	0.895				on tec		
High school										Jun		
or below	8,611	34.78%	24	31.58%			-	-	-	1e 7,	-	-
Bachelor's										, 2025 ies.		
degree	13,597	54.92%	42	55.26%			-	-	-	25 - 25 -	-	-
Postgraduate										at Departm		
or higher	2,549	10.30%	10	13.16%			-	-	-	par	-	-
Occupation					21.85	0.003*				tment		

Page 61 of 67

2 3 4

5 6

7

8

9 10

11

12

13

14 15

16

17

18 19

20

21

22 23

24

25

26 27

28

29

30

31 32

33

34

35 36

37

38

										fi. Re 190pen-2024-089734 on 2€January 2024 Ci Erasmu: 20 Erasmu:		
)										9734 ding		
>28.0										on 2		
(Obesity)	830	3.35%	4	5.26%			2.92	0.82, 8.42	0.064	e R 2€January 2025€Downloaded from http://bmjopen.bmj.com/ on June ○ Erasmuskogeschool . uses related to text and data mining, Al training, and similar technolo	0.82, 8.43	0.064
						<0.001*				inua E		
Smoking					20.82	*				ıry 2 rası ated		
No	24,666	99.63%	74	97.37%			Ref.			mus Ref.		
Yes	91	0.37%	2	2.63%			7.33	1.19, 23.81	0.006^{*}	sac sac text	1.08, 21.70	0.009*
Alcohol					0.27	0.047				wnl esc and		
Consumption				100.00	0.37	0.947				oade hool data		
No	24.607	99.76%	76	100.00						ed fi a m		
No Yes	24,697 60	0.24%	76 0	0.00%			-	-	-	inin -	-	_
Obstetric and				0.0076				-	-	g, A	-	-
Pregnancy	Gynecolo	gic mstor	y							o://b		
History					7.08	0.215						
1	14,932	60.31%	53	69.74%	7.00	0.215	_	101	_	g, a	_	_
2	6,500	26.26%	13	17.11%			_		· <u>-</u>	d from http://bmjopen.bmj.com/ on June 7, 20 mining, Al training, and similar technologies.	-	_
≥3	3,325	13.43%	10	13.16%			_	_		j.co imii -	-	_
Parity	,									m√ o ar te		
(number of										on Ju		
deliveries)					15.86	0.007*				ine		
0	20,479	82.72%	71	93.42%			Ref.			gi 7, Ref.		
1	4,148	16.75%	4	5.26%			0.28	0.08, 0.67	0.013*	0.27	0.08, 0.68	0.014*
≥2	130	0.53%	1	1.32%			2.22	0.13, 10.1	0.431	2. 3 6	0.14, 13.3	0.375
history of										ера		
miscarriage					0.16	0.983				partment GEZ-LTA		

 Page 62 of 67

							вив орен			copyright, includ			
No	18,210	73.55%	57	75.00%			_	_		cluding		_	
Yes	6,547	26.45%	19	25.00%			_	_	_	y for	<u>-</u>	_	_
spontaneous	0,5 17	20.1570	17	25.0070						uses	3		
abortion					0.04	0.998				zo January Era uses relate	-		
No	22,271	89.96%	68	89.47%			-	_	_	Era elat	_	_	_
Yes	2,486	10.04%	8	10.53%			_	_	_	ed to)	_	_
induced	,									ush o te			
abortion or										oges ext ar	,		
medication													
abortion					0.57	0.903				nioaded school . id data i	-		
No	19,920	80.46%	63	82.89%			-	-	-	min 7		-	_
Yes	4,837	19.54%	13	17.11%			-	-	-	ing	-	-	_
induced										<u>`</u> ≥ =			
labor					3.18	0.365				mining, Al training, and similar technologies.			
				100.00						ning	•		
No	24,250	97.95%	76	%			-	C1/	-	j, ar	-	-	-
Yes	507	2.05%	0	0.00%			-	-	_	ld si	-	-	-
assisted										<u> </u>			
reproductive										ar te			
technology					0.02	0.999				ichr Ju	-		
No	24,110	97.39%	74	97.37%			-	-		lolo		-	-
Yes	647	2.61%	2	2.63%			-	-	-	r, zi	· -	-	-
Gestational										ş. 025	8		
Age at										gies.			
Delivery													
(weeks)					14.46	0.013*				epartment GEZ-LIA			

Page 63 of 67

<38	1,503	6.07%	4	5.26%			Ref.			jopen-2024-089734		
38-40	12,994	52.49%	29	38.16%			0.84	0.33, 2.83	0.742	fo on _	_	_
>40	10,260	41.44%	43	56.58%			1.57	0.64, 5.24	0.386	20 Ja	_	_
Diabetes	,				0.75	0.860		,		1 20 January Eras r uses relate		
No	21,443	86.61%	64	84.21%			-	_	_	uary Era	_	-
Yes	3,314	13.39%	12	15.79%			-	_	_	, 2025 smus	_	-
hypertension	,				2.44	0.486				요구.		
No	22,966	92.77%	68	89.47%			-	_	_	Dow -	_	_
Yes	1,791	7.23%	8	10.53%			-	_	_	nlo: scho	_	-
	,					<0.001*				ader ool		
anemia					18.39	*				- Ref. Downloaded from h ogeschool . xt and data mining,		
No	20,517	82.87%	53	69.74%			Ref.			Ref.		
Yes	4,240	17.13%	23	30.26%			2.10	1.26, 3.38	0.003*	<u>≥</u> 1. 5 3	1.13, 3.20	0.012*
coagulation					3100.8	<0.001*				://br		
disorder					0	*				njor ning		
No	24,624	99.46%	48	63.16%			Ref.			ref. Ref. Ref. 21 		
								65.10,		bmj si	67.90,	
Yes	133	0.54%	28	36.84%			108.01	176.01	<0.001**	114.21	190.00	<0.001**
uterine										n/ o		
fibroids/aden										on June technoic		
omyosis					0.51	0.917				ine		
No	24,023	97.04%	73	96.05%			-	-	-	- 7, 2: gies	-	-
Yes	734	2.96%	3	3.95%			-	-	-	7, 2025 at De gies.	-	-
polyhydramn										at D		
ios					0.62	0.892				ера		
No	22,299	90.07%	67	88.16%			-	-	-	- partm	-	_

 Page 64 of 67

Page 65 of 67

2 3 4

5 6

8

9 10

11

12

13

14 15

16

17

18 19

20

21

22 23

24

25

26 27

28

29

30

31 32

33

34

35

36

37

							J.1.13 G P G.1.			ijo Pop		
										njopen-2024-089. copyright, includ		
										024- it, in		
Time of									•	' '34 on 20 Ja		
delivery	10.274	44.000/		42.4207	0.14	0.986				n 20 or u		
Day shift	10,374	41.90%	33	43.42%			-	-	-	۔ الادر Ses	-	-
Night	1.4.202	50.100/	40	56.500/						1 20 January Era r uses relate		
shift	14,383	58.10%	43	56.58%			-	-	-		-	-
Total						-0.001*				2025. smus		
duration of					(2.15	<0.001*				h Do ext		
labor	24 205	00.540/	(0	00.700/	62.15		D.C			wnl escl		
normal	24,395	98.54%	69	90.79%			Ref.			xt and data r		
prolonge d	362	1.46%	7	9.21%			6.84	2.84, 14.0	<0.001**	a - ed mir - fc 5.57	2.18, 12.10	<0.001**
u First stage of	302	1.4070	/	9.2170			0.64	2.84, 14.0	~0.001	mining,	2.16, 12.10	~0.001
labor -										http://bmjopen.bmj.com/ g, Al training, and similar		
Latent phase					4.59	0.205				ttp://bmjopen.bmj.com/ on June 7, 20		
normal	24,337	98.30%	73	96.05%	1.09	0.200	_	/	_		_	_
prolonge	2 1,557	70.5070	75	70.0270						pen g, a		
d	420	1.70%	3	3.95%			_	-	_	nd s	_	_
First stage of												
labor -										ar t		
Active phase					8.27	0.041				on June		
normal	24,264	98.01%	72	94.74%			-	-	_	ine -	-	-
prolonge									•	7, 2 gie:		
d	493	1.99%	4	5.26%			-	-	-	. 25 -	-	-
Second stage						<0.001*				at I		
of labo					19.57	*				Department GEZ-LTA		
of labo	24565	99.22%	73	96.05%			Ref.			ع Def		

 Page 66 of 67

Page 67 of 67

2 3 4

5 6

7

8

9 10

11

12

13

14 15

16

17

18 19

20

21

22 23

24

25

26 27

28

29

30

31 32

33

34

35 36

37

										<u>a 9</u>		
vagina, or										89734 on 20 Ja uding for uses		
perineum										n 20 or us		
No	22,468	90.75%	69	90.79%			-	-	-	o Ja	-	-
Yes	2,289	9.25%	7	9.21%			-	-	-	rel	-	-
Newborn										January 2025. Erasmush ses related to te		
weight										2025 mui		
(grams)					11.83	0.037 *				5. D sho		
<2500	227	0.92%	1	1.32%			Ref.			and Ref.		
2500-40										Ref. S. Downloaded the shogeschool .		
00	23,159	93.55%	66	86.84%			0.65	0.14, 11.4	0.666	ded -	-	
>4000	1,371	5.54%	9	11.84%			1.49	0.28, 27.5	0.706	min -	-	-
Newborn										ing,		
length										≥ f		
(centimeters)					2.95	0.399				//bn		
≤55	24,036	97.09%	72	94.74%			_	/O-	-	njop ning	-	-
>55	721	2.91%	4	5.26%			-		-	י י י י י י י י י י י י י י י י י י י	-	-
										omj d si		
										mil:		
										n/o		
										n J.		
										nolo		
										7, 2 gie		
										025 s.		
										at		
										Dep		
										oart		
										7, 2025 at Department GEZ-LTA vgies.		
										o t		
										EZ-		
										Ţ		