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# Prediction of emergency cesarean section by measurable maternal and fetal characteristics

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**ABSTRACT**

**Objective** To analyze the maternal and fetal factors affecting emergency cesarean section (EmCS) and establish a risk scoring system to quantitatively predict the risk of EmCS.

**Design** A total of 10,295 pregnant women were enrolled in this study. The influence of maternal and fetal factors on the risk of EmCS was analyzed.

**Results** 991 (9.63%) cases of failed vaginal delivery received EmCS. The two main causes of EmCS were fetal distress (67.21%) and abnormal fetal position (14.93%). There were significant differences in 17 maternal and fetal factors between the normal vaginal delivery (NVD) and EmCS groups ( $p < 0.05$  for all). Univariate and multivariate regression analyses showed that nine maternal and infant factors were independent risk factors ( $p < 0.05$  for all). The major factors were abnormal quantity of amniotic fluid (OR 6.867, 95% CI 4.442 to 10.618), nulliparous (OR 4.336, 95% CI 3.074 to 6.115), induction of labor (OR 5.300, 95% CI 4.514 to 6.224) and abnormal characters of amniotic fluid (OR 3.126, 95% CI 2.708 to 3.608). A risk scoring system (six grades) was established based on those factors which showed high discriminative power. The rate of EmCS was 1.30%, 2.57%, 5.83%, 13.94%, 21.75% and 39.71% in grades 1, 2, 3, 4, 5 and 6, respectively. The area under the curve of the risk scoring system was 0.787, indicating that the regression model of the risk factors had a good predictive ability.

**Conclusion** An effective risk scoring system has been developed to quantitatively assess the risk of EmCS based on measurable maternal and fetal factors. The system is simple, easy to operate and has good repeatability in clinical practice.

**INTRODUCTION**

Although there is major geographic variation in rates and trends over time, with rates declining in some of the largest urban areas, the overall annual rate of cesarean section (CS) has increased in China, reaching 34.9%.<sup>1</sup> Such data are comparable to those in the USA,<sup>2</sup> but more than twice as high as the 10–15% recommended by the World Health Organization.<sup>3,4</sup> This also brings a series of adverse maternal and fetal outcomes caused by CS.<sup>5–7</sup> Therefore, CS should be done only when there are definite medical indications. However, we have to admit that CS is an effective means to solve the

**Significance of this study****What is already known about this subject?**

- ▶ Emergency cesarean section (EmCS) can be a life-saving procedure if pregnant women experience abnormal conditions in the process of vaginal delivery.
- ▶ Deciding to perform EmCS is a complicated process, occurring only in specific obstetric conditions, and requires awareness and rapid assessment of the risky situation.

**What are the new findings?**

- ▶ Maternal and fetal factors differ in normal vaginal delivery and EmCS groups, with >20 maternal and infant characteristics differing between the two groups.
- ▶ About 12 maternal and fetal factors were associated with risk of EmCS.
- ▶ A risk scoring system for EmCS based on nine distinct factors that could be determined before delivery has been established with strong objectivity and good repeatability for Chinese women.

**Significance of this study****How might these results change the focus of research or clinical practice?**

- ▶ In clinical circumstances, the information of factors included in the risk scoring system can be evaluated before delivery. Thus, this risk scoring system is simple, easy to operate and has good repeatability in clinical practice. If the score is too high, we should pay close attention to the situation of mothers and babies should be monitored closely during childbirth.

medical and surgical complications of dystocia and severe complications of pregnancy, and has an irreplaceable role.<sup>8</sup> Emergency CS (EmCS) can be a life-saving procedure if pregnant women experience abnormal conditions in the process of vaginal delivery such as fetal distress, eclampsia or severe pre-eclampsia.<sup>9</sup> It has been reported that about 14.9% of all deliveries are transferred to EmCS during vaginal delivery for identified indications in China.<sup>10</sup>



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Thus, a trial of labor via the vagina should be carried out for pregnant women without specific indications of CS. At the same time, close attention should be paid to the changes in maternal and fetal health status in order to ensure the safety of mothers and infants, and to prepare for EmCS. Deciding to perform EmCS is a complicated process, occurring only in specific obstetric conditions, and requires awareness and rapid assessment of the risky situation.<sup>11</sup> Failure to perform EmCS in time may lead to postpartum mental disorders<sup>12</sup> and even serious adverse maternal and fetal outcomes.<sup>13</sup> Recognizing an acute situation which may be potentially life-threatening is considered one of the most challenging tasks in obstetrics.<sup>14</sup> Most studies focus on the related factors and prognosis of elective CS, but few on the factors of vaginal delivery failure.<sup>13 15</sup> Several studies have found that anxiety and fear about the delivery process increased the risk of EmCS twofold.<sup>16</sup> Furthermore, advanced age, obesity and overweight are also risk factors for failure of vaginal delivery and EmCS.<sup>17 18</sup> Unfortunately, these single-factor studies do not combine other factors of mother and infant, and cannot quantify the risk of EmCS.

The purpose of this study is to systematically analyze 22 maternal and fetal factors in terms of demographic characteristics of pregnant women, pregnancy history, perinatal complications, neonatal features and labor process, in order to identify the high-risk factors associated with EmCS before delivery. A risk scoring system was then established to quantitatively predict the risk of EmCS during vaginal delivery.

## MATERIALS AND METHODS

### Study population

In this study we retrospectively analyzed the medical records in the Department of Obstetrics, Maternal and Child Health Hospital of Hubei Province, China. The inclusion and exclusion criteria were as follows. Inclusion criteria: (1) all patients delivered in our hospital from January 1, 2016 to December 31, 2016; (2) gestation age (weeks) <42; (3) vaginal examination was performed before vaginal delivery and there were no abnormalities in pelvic measurements or complications endangering the lives of mothers and fetuses; (4) there was no fetal distress and no obvious head-pelvic disproportion before the trial of vaginal delivery. A total of 19,816 cases met the criteria. Exclusion criteria: (1) multiple birth (n=402); (2) breech position (n=810) and transverse/oblique position (n=161), (3) received CS after full-term pregnancy due to severe complications of fetal malformation (n=20); (4) those who failed in vaginal delivery in the external hospital and were transferred to our hospital (n=20); (5) direct CS with certain indications (n=8108). Thus, a total of 10,295 cases were included in this study. EmCS was defined as a case in which the procedure of vaginal delivery was discontinued due to the occurrence of fetal distress or other reasons. Fetal distress was defined as depletion of oxygen and accumulation of carbon dioxide, leading to a state of 'hypoxia and acidosis' during intrauterine life. Continuous electronic fetal heart rate (FHR) monitoring was performed for each woman during the process of labor to evaluate the status of fetal distress based on a three-tiered system categorization of FHR patterns which was recommended by the American College

of Obstetricians and Gynecologists (ACOG). The research protocol for this study was approved by the Medical Ethics Committee of Maternal and Child Health Hospital of Hubei Province. This study was a retrospective study. We only extracted clinically relevant information, did not affect the treatment and health of patients, and the patients' privacy was protected, so written informed consent was not applicable for this study.

### Outcome measures

All perinatal data were extracted from medical records and classified into five items: demographic characteristics of pregnant women (maternal age, height, weight before pregnancy, body mass index (BMI) before pregnancy, weight at delivery, BMI at delivery, gestational age), pregnancy history (gravidity, parity, previous CS), perinatal complications (characters of amniotic fluid, umbilical cord status, hypertensive disorders of pregnancy, low lying placenta, placental abruption, oligohydramnios, hydramnion), neonatal features (birth weight, macrosomia, fetal gender) and labor process (induction of labor, epidural anesthesia). According to the outcome of vaginal delivery, all women were classified into two groups: normal vaginal delivery (NVD) and EmCS. The primary outcome of this study was to identify the possible independent factors associated with the failure of vaginal delivery and to quantify the risk of EmCS.

### Statistical analysis

Statistical analyses were performed using SPSS version 19.0 (SPSS Inc, Chicago, Illinois, USA). Quantitative variables were expressed as mean±SD. Qualitative variables were expressed as absolute and relative frequencies. Univariate analysis was performed to explore the relationship between EmCS rate and perinatal factors. In order to find factors independently associated with the risk for EmCS, a multiple logistic regression analysis was performed in a stepwise fashion. Statistical significance was considered to be the case when  $p < 0.05$  or if the 95% CI of odds ratios (OR) did not include 1. In order to better evaluate the risk coefficient in the development of the risk scoring system, the 75% quantile was chosen as the cut-off value and the integer of the OR value was used as the score for each risk factor.

## RESULTS

### Characteristics of study population

A total of 10,295 women met the inclusion and exclusion criteria, of which 9304 (90.37%) cases were successfully managed with vaginal delivery and 991 (9.63%) cases were transferred to EmCS during the process of vaginal delivery. There were some differences between the two groups in terms of the demographic characteristics of the pregnant women, pregnancy history, perinatal complications, neonatal features and labor process (table 1). The mean±SD age of the pregnant women in the EmCS group (28.59±3.22) was significantly higher than that in the NVD group (28.02±3.53) ( $p < 0.001$ ), and the proportion of women of advanced age was significantly higher in the EmCS group (5.45%) than in the NVD group (4.61%) ( $p < 0.001$ ). There was no significant difference in gestational age and maternal height between the two groups

**Table 1** Characteristics of study population

Variable	Study population (n=10,295) (%)	Groups		
		NVD (n=9304) (%)	EmCS (n=991) (%)	P value*
Demographic characteristics of pregnant women				
Mean age (years)	28.08±3.51	28.02±3.53	28.59±3.22	<0.001
Age categories (years)				
<25	2121 (20.60)	1986 (21.35)	135 (13.62)	<0.001
25–29	5179 (47.30)	4649 (49.96)	530 (53.48)	
30–35	2512 (24.40)	2240 (24.08)	272 (27.45)	
≥35	483 (4.70)	429 (4.61)	54 (5.45)	
Height (cm)	161.6±0.046	161.6±0.045	161.8±0.048	0.313
Gestational age (weeks)	39.41±1.41	39.41±1.43	39.43±1.28	0.555
Weight before pregnancy (kg)	51.35±5.41	51.17±5.29	53.03±6.19	<0.001
BMI before pregnancy (kg/m <sup>2</sup> )	19.67±2.10	19.61±2.03	20.31±2.58	<0.001
Weight at delivery (kg)	68.71±8.18	68.48±8.06	70.89±8.88	<0.001
BMI at delivery (kg/m <sup>2</sup> )	26.30±2.92	26.21±2.83	27.13±3.56	<0.001
Pregnancy history				
Gravidity				
Primigravidity	6165 (59.88)	5512 (59.24)	653 (65.89)	<0.001
Multigravidas	4130 (40.12)	3792 (40.76)	338 (34.11)	
Parity				
Nulliparous	8775 (85.24)	7830 (84.16)	946 (95.46)	<0.001
Multiparous	1519 (14.76)	1474 (15.84)	45 (4.54)	
Previous cesarean section				
Yes	40 (0.39)	36 (0.39)	4 (0.39)	0.936
No	10255 (99.61)	9268 (99.61)	987 (99.61)	
Perinatal complications				
Characters of amniotic fluid				
Clear	7459 (72.45)	6977 (74.99)	482 (48.64)	<0.001
Stage I	239 (2.33)	214 (2.30)	25 (2.52)	
Stage II	194 (1.88)	185 (1.99)	9 (0.91)	
Stage III/bloody	2403 (23.34)	1928 (20.72)	475 (47.93)	
Umbilical cord status				
Normal	6452 (62.67)	5928 (63.71)	524 (52.88)	<0.001
Cord entanglement	3547 (34.45)	3136 (33.71)	411 (41.47)	
Torsion of umbilical cord	296 (2.88)	240 (2.58)	56 (5.65)	
HDP*				
Yes	429 (4.17)	362 (3.89)	67 (6.76)	<0.001
No	9866 (95.83)	8942 (96.11)	924 (93.24)	
Low lying placenta				
Yes	40 (0.39)	38 (0.41)	2 (0.20)	0.320
No	10255 (99.61)	9266 (99.59)	989 (99.80)	
Placental abruption				
Yes	40 (0.39)	36 (0.39)	4 (0.40)	0.936
No	10255 (99.61)	9268 (99.61)	987 (99.60)	
Oligohydramnios				
Yes	72 (0.70)	40 (0.43)	32 (3.23)	<0.001
No	10223 (99.3)	9264 (99.57)	959 (96.77)	
Hydramnion				
Yes	50 (0.49)	36 (0.39)	14 (1.41)	<0.001
No	10245 (99.51)	9268 (99.61)	977 (98.59)	
Neonatal features				
Birth weight (g)	3293.78±415.51	3275.88±413.38	3461.84±397.55	<0.001
Neonatal weight (g)				
<2500	260 (2.53)	252 (2.71)	8 (0.81)	<0.001
2500–4000	9581 (93.06)	8698 (93.48)	883 (89.10)	

Continued

Table 1 Continued

Variable	Study population (n=10,295) (%)	Groups		
		NVD (n=9304) (%)	EmCS (n=991) (%)	P value*
≥4000 (macrosomia)	454 (4.41)	354 (3.81)	100 (10.09)	
Gender				
Male	5372 (52.18)	4806 (51.66)	566 (57.11)	<b>0.001</b>
Female	4923 (47.82)	4498 (48.34)	425 (42.89)	
Labor process				
Induction of labor				
Yes	1180 (11.46)	822 (8.83)	358 (36.13)	<b>&lt;0.001</b>
No	9115 (88.54)	8482 (91.17)	633 (63.87)	
Epidural anesthesia				
Yes	6485 (62.99)	5855 (62.92)	630 (63.57)	0.052
No	3810 (37.01)	3449 (37.08)	361 (36.43)	

\*Two-sided  $p < 0.05$  was considered statistically significant, and the bold typeface shows the significant data.

BMI, body mass index; EmCS, emergency cesarean section; HDP, hypertensive disorders of pregnancy; NVD, normal vaginal delivery.

( $p > 0.05$  for all). Compared with the NVD group, maternal weight before pregnancy, maternal BMI before pregnancy, maternal weight at delivery and maternal BMI at delivery in the EmCS group were significantly higher ( $p < 0.05$  for all).

The proportion of the pregnant women who conceived and gave birth for the first time in the EmCS group was higher than that in the NVD group ( $p < 0.01$ ). There was no difference in the proportion of pregnant women with previous CS between the two groups ( $p = 0.936$ ). Compared with the NVD group, pregnant women with abnormal amniotic fluid characteristics, abnormal quantity of amniotic fluid (oligohydramnios, hydramnion), abnormal umbilical cord status (normal, cord entanglement, torsion) and hypertensive disorders of pregnancy (HDP) was significantly higher in the EmCS group ( $p < 0.05$  for all). The differences in the proportion of pregnant women with low lying placenta and placental abruption between the two groups were not statistically significant ( $p > 0.05$  for all).

In the EmCS group, the proportion of male infants (57.11%) was significantly higher than in the NVD group (51.66%), the weight of neonates in the EmCS group ( $3461.84 \pm 397.55$ ) was significantly higher than in the NVD group ( $3275.88 \pm 413.38$ ), and the proportion of macrosomia in the EmCS group (10.09%) was significantly higher than in the NVD group (3.81%) ( $p < 0.001$ ). In addition, the proportion of pregnant women receiving induction of labor in the EmCS group (36.13%) was higher than that in the NVD group (8.83%), and the difference was statistically significant ( $p < 0.001$ ). There was no significant difference in the proportion of pregnant women receiving epidural anesthesia between the two groups ( $p = 0.052$ ). Altogether, differences were observed in 17 variables between the two groups.

### Analysis of causes of EmCS

Among 991 pregnant women who received EmCS, the major direct causes were fetal distress ( $n = 666$ , 67.21%), persistent occipital transverse position or occipital posterior position ( $n = 148$ , 14.93%), and relative cephalopelvic disproportion ( $n = 81$ , 8.17%). Sixty-four (6.46%) women underwent EmCS because of their unwillingness to cooperate with vaginal delivery due to personal factors. In particular, 11

(1.11%) pregnant women suffered from prenatal hemorrhage during delivery and EmCS was performed to ensure the safety of mothers and infants (table 2).

In the course of vaginal delivery, the cervical dilatation of five women was 0 cm and about 10 women had EmCS even though the cervical dilatation was 10 cm. Of the 991 cases, the mean cervical dilatation was  $3.11 \pm 1.92$  cm (median 2.5 cm). There were 518 (52.27%) parturients with cervical dilatation less than 3 cm in whom EmCS was performed. The cervical dilatation was significantly higher in women who underwent EmCS for abnormal fetal position ( $p < 0.001$ ).

### Univariate and multivariate analysis of the risk of EmCS

Variables with differences between the NVD and EmCS groups were included in the univariate analysis. The results showed that the risk of EmCS increased if the pregnant woman had one of the following conditions: maternal age  $\geq 30$  years, BMI  $\geq 20.5$  before pregnancy, BMI  $\geq 28$  at delivery, nulliparous, abnormal characteristics of amniotic fluid, abnormal umbilical cord, HDP, abnormal quantity of amniotic fluid, neonatal weight  $> 3550$  g, male newborn and need to induce labor ( $p < 0.05$  for all). However,

Table 2 Reasons for emergency cesarean section

Reasons	Value	P value	Cervical dilatation (cm)*
Fetal distress	666 (67.21)	$<0.001$	$2.85 \pm 1.63$
Abnormal fetal position	148 (14.93)		$4.93 \pm 2.38$
Relative cephalopelvic disproportion	81 (8.17)		$2.71 \pm 1.73$
Social factors	64 (6.46)		$2.16 \pm 1.30$
Others†	21 (2.12)		$2.83 \pm 1.93$
Antepartum hemorrhage	11 (1.11)		$2.82 \pm 1.47$

\*Compared with other groups, the cervical dilatation was significantly higher in women who received EmCS for abnormal fetal position ( $p < 0.001$  for all).

†Others included pathologic retraction rings in 6 cases (0.61%), pre-eclampsia in 9 (0.91%), threatened uterine rupture in 5 (0.50%) and presentation of umbilical cord in 1 case (0.10%).

**Table 3** Univariate and multivariate analyses of factors associated with EmCS

Variables	N (%)	Univariate		Multivariate	
		OR (95% CI)	P value	OR (95% CI)	P value
Demographic characteristics					
Age (years)					
<30	7300 (70.91)	Ref		Ref	
≥30	2995 (29.09)	1.129 (1.060 to 1.402)	0.006	1.619 (1.381 to 1.898)	<0.001
BMI before pregnancy (kg/m <sup>2</sup> )					
<20.5	7793 (75.70)	Ref		Ref	
≥20.5	2502 (24.30)	1.936 (1.687 to 2.23)	<0.001	1.868 (1.576 to 2.214)	<0.001
BMI at delivery (kg/m <sup>2</sup> )					
<28	7713 (74.92)	Ref		Ref	
≥28	2582 (25.08)	1.765 (1.537 to 2.027)	<0.001	1.244 (1.048 to 1.478)	0.016
Pregnancy history					
Gravidity					
Primigravidity	6165 (59.88)	Ref		Ref	
Multigravidity	4130 (40.12)	0.752 (0.656 to 0.863)	<0.001	0.918 (0.783 to 1.077)	0.235
Parity					
Multiparous	1519 (14.76)	Ref		Ref	
Nulliparous	8775 (85.24)	3.957 (2.292 to 5.364)	<0.001	4.336 (3.074 to 6.115)	<0.001
Perinatal complications					
Characters of amniotic fluid					
Clear	7459 (72.45)	Ref		Ref	
Abnormal*	2836 (27.55)	3.166 (2.772 to 3.617)	<0.001	3.126 (2.708 to 3.608)	<0.001
Umbilical cord					
Normal	6452 (62.67)	Ref		Ref	
Abnormal†	3843 (37.3)	1.565 (1.372 to 1.785)	<0.001	1.414 (1.225 to 1.632)	<0.001
HDP					
No	9866 (95.83)	Ref		Ref	
Yes	429 (4.17)	1.791 (1.368 to 2.345)	<0.001	1.372 (1.012 to 1.860)	0.050
Abnormal quantity of amniotic fluid					
No	10173 (98.81)	Ref		Ref	
Yes‡	122 (1.19)	5.910 (4.074 to 8.576)	<0.001	6.867 (4.442 to 10.618)	<0.001
Neonatal features					
Neonatal weight (g)					
<3550	7454 (72.40)	Ref		Ref	
≥3550	2841 (27.60)	2.181 (1.908 to 2.494)	<0.001	2.070 (1.786 to 2.400)	<0.001
Gender					
Female	4923 (47.82)	Ref		Ref	
Male	5372 (52.18)	1.246 (1.092 to 1.423)	0.001	1.216 (1.052–1.404)	0.008
Labor process					
Induction of labor					
No	9115 (88.54)	Ref		Ref	
Yes	1180 (11.46)	5.836 (5.033 to 6.767)	<0.001	5.300 (4.514 to 6.224)	<0.001

\*Abnormal amniotic fluid character included stage I–III, bloody.

†Abnormal umbilical cord included cord entanglement and torsion of the umbilical cord.

‡Abnormal quantity of amniotic fluid includes oligohydramnios and hydramnion.

BMI, body mass index; HDP, hypertensive disorders of pregnancy; Ref, reference category.

multigravidity (OR 0.752, 95%CI 0.656 to 0.863) decreased the risk of EmCS (table 3).

By incorporating the above 12 meaningful indicators into further multivariate analysis, it was found that gravidity was not an independent risk factor for EmCS ( $p=0.235$ ) and the significance of HDP was marginal ( $p=0.50$ ). The risk of EmCS in pregnant women with abnormal amniotic fluid quantity was more than six times higher than that in those with a normal quantity of amniotic fluid (OR 6.867,

95%CI 4.442 to 10.618,  $p<0.001$ ). The risk of EmCS in nulliparous women was about four times higher than in multiparous women (OR 4.336, 95%CI 3.074 to 6.115,  $p<0.001$ ), the risk of EmCS in pregnant women who needed induction of labor was more than five times higher than in pregnant women who had spontaneous onset (OR 5.300, 95%CI 4.514 to 6.224,  $p<0.001$ ), while the risk of EmCS in pregnant women with abnormal characters of amniotic fluid was three times higher than in pregnant

**Table 4** Development of EmCS risk scoring system

EmCS risk scoring system (scores )	N (%)	NVD (n, %)	EmCS (n, %)	P value*	Or (95% CI)	P value†
Grade 1 (0–3 points)	919 (8.93)	907 (9.75)	12 (1.21)	<0.001	Ref	
Grade 2 (4–5 points)	3389 (32.92)	3302 (35.49)	87 (8.78)		1.991 (1.084 to 3.658)	0.026
Grade 3 (6–7 points)	2605 (25.30)	2453 (26.37)	152 (15.34)		4.684 (2.589 to 8.471)	<0.001
Grade 4 (8–9 points)	1686 (16.38)	1451 (15.59)	235 (23.71)		12.241 (6.813 to 21.994)	<0.001
Grade 5 (10–11 points)	938 (9.11)	734 (7.89)	204 (20.58)		21.007 (11.641 to 37.906)	<0.001
Grade 6 (≥12 points)	758 (7.36)	457 (4.91)	301 (30.38)		49.782 (27.656 to 89.611)	<0.001

\*Chi square tests were used.

†Logistic regression analysis was used.

EmCS, emergency cesarean section; HDP, hypertensive disorders of pregnancy; NVD, normal vaginal delivery; Ref, reference category.

women with clear amniotic fluid (OR 3.126, 95% CI 2.708 to 3.608,  $p < 0.001$ ).

### Development of EmCS risk scoring system

In order to effectively identify pregnant women with a high risk of EmCS, we further scored the abovementioned independent risk factors according to OR value. Briefly, women who had a risk factor with OR value between 1 and 2 were scored as 1 point, women with a risk factor with OR value between 2 and 3 were scored as 2 points, and so on. The score was 0 if there was no risk factor. Considering that it is illegal to differentiate the sex of newborns before delivery, and the significance of HDP in this study was limited, only nine factors were included and scored as follows: maternal age  $\geq 30$  years (1 point), BMI  $\geq 20.5$  before pregnancy (1 point), BMI  $\geq 28$  before delivery (1 point), nulliparous (4 points), abnormal characters of amniotic fluid (3 points), abnormal umbilical cord (1 point), abnormal quantity of amniotic fluid (6 points), neonatal weight  $\geq 3550$  g (2 points) and induction of labor (5 points). Each woman scored the sum of the nine factors (0–24 points). The results showed that all women in this study had scores ranging from 0 to 22. According to the proportion of pregnant women with different scores in the EmCS group, the EmCS risk scoring system was further established: scores of 0–3 were categorized as grade 1, scores of 4–5 were grade 2, scores of 6–7 were grade 3, scores of 8–9 were grade 4, scores of 10–11 were grade 5 and scores of 12 points and above were grade 6 (table 4).

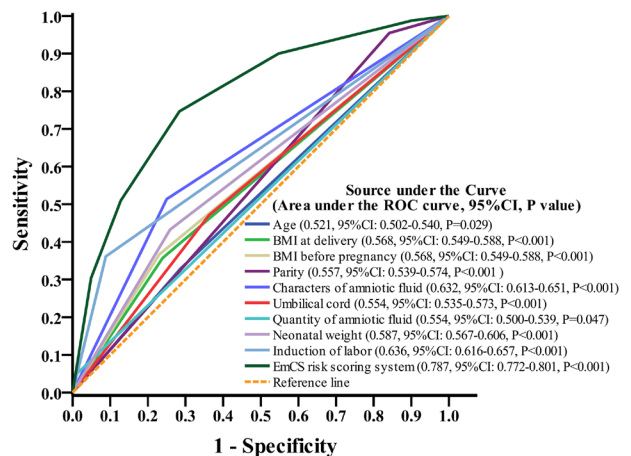
The rate of EmCS was 1.30%, 2.57%, 5.83%, 13.94%, 21.75% and 39.71% in patients with grade 1, 2, 3, 4, 5 and 6, respectively. The difference was statistically significant ( $p < 0.001$ ). Nearly 70% of pregnant women in the NVD group were grades 1–3, while over 74% of pregnant women in the EmCS group were grades 4–6; the difference was statistically significant ( $p < 0.001$ ). When the risk of the pregnant women was lower than or equal to grade 3, the rate of EmCS was lower than that of the study population (9.63%). Therefore, pregnant women with a risk score above grade 4 need to be particularly vigilant and may need EmCS during the vaginal delivery process. Compared with pregnant women in grade 1, the risk of EmCS in pregnant women increased significantly with the increase in grade, with OR (95% CI) values of 1.991 (1.084–3.658), 4.684 (2.589–8.471), 12.241 (6.813–21.994), 21.007 (11.641–37.906) and 49.782 (27.656–89.611) for grades 1, 2, 3, 4, 5 and 6, respectively. The area under the curve of the risk scoring system was 0.787 (95% CI 0.772 to 0.801,

$p < 0.001$ ), which was more valuable than any other single risk factor, indicating that the regression model of the risk factors had a good predictive ability (figure 1).

### DISCUSSION

This study aimed to answer the following questions: (1) Do the maternal and fetal factors differ in NVD and EmCS groups? (2) To what extent do maternal and fetal perinatal factors affect EmCS? (3) Is there a simple and feasible model based on measurable and repeatable maternal and fetal factors that can predict the risk of EmCS?

Globally, the CS rate is rising and medically unnecessary CSs are prevalent.<sup>19 20</sup> Increased maternal obesity, widespread assisted reproductive technology, cultural and social elements are also contributing factors.<sup>21 22</sup> However, there are some special cases such as those who fail in the course of attempting vaginal delivery and then must receive EmCS to terminate the delivery. How to identify these mothers and perform emergency surgery in time is very important to ensure the safety of mothers and infants.<sup>23</sup> In this study, our findings showed that 9.63% of pregnant women had EmCS. This result is lower than that of other researchs.<sup>24</sup> To answer the first question, we analyzed the differences in more than 20 maternal and infant characteristics between the two groups. The results showed that there was no difference in the proportion of pregnant women with low lying placenta and placental abruption between the NVD



**Figure 1** Maternal and fetal factors-based risk scoring system performed well in predicting the risk of emergency cesarean section (EmCS) of pregnant women compared with other factors.

and EmCS groups. Studies have shown that only patients with low lying placenta who have a history of hospitalization due to vaginal bleeding and whose cervix continues to shorten have an increased risk of EmCS.<sup>25</sup> In addition, our results suggested that epidural anesthesia during delivery does not reduce productivity and lead to EmCS. In fact, some studies have shown that anesthetic leads to a better analgesic satisfaction without adverse effects on uterine electromyographic activities or obstetric outcomes during labor.<sup>26 27</sup> We can therefore use labor analgesia in more patients to relieve pain. As we know, childbirth may cause the most severe pain some women experience in their lifetime, and some women finally choose EmCS because they cannot tolerate the pain.<sup>26</sup>

The influence of the abovementioned maternal and fetal factors on the risk of EmCS was further analyzed to answer the second question. Univariate analysis found that 12 factors were associated with the risk of EmCS, including age, pre-pregnancy BMI, BMI at delivery, gravidity, parity, fetal weight and gender. However, multivariate analysis found that gravidity was not an independent risk factor and the impact of HDP was only marginally significant. These results are similar to other studies.<sup>17 28–30</sup> It was noteworthy that an abnormal quantity of amniotic fluid was the greatest risk factor affecting EmCS, followed by spontaneous seizure or induction of labor. Although induction of labor is the most common intervention in obstetrics, it actually reflects the underlying problems of pregnant women with an uncomplicated singleton pregnancy in the onset of labor and subsequent spontaneous vaginal delivery.<sup>31</sup> It has been documented that, compared with spontaneous onset of delivery, induction of labor is associated with an increased risk for EmCS among both nulliparous and multiparous women.<sup>32</sup> However, a recent clinical trial showed that, for certain women with large-for-date fetuses, labor would not increase the rate of EmCS and would instead have certain benefits in reducing shoulder dystocia and increase the success rate of vaginal delivery.<sup>33</sup> Interestingly, we found that when the fetus was male, women were more likely to have EmCS. Similarly, other studies also demonstrated that male babies were more likely to be delivered by instrumental delivery, as well as by EmCS for failure to progress and fetal distress, even after adjusting for birth weight.<sup>34 35</sup> It is suggested that there is a different fetoplacental response to the induction of the labor process between the fetal genders, which could explain the higher failure rates in male fetuses.<sup>36</sup>

Although some studies suggest that mental health status can predict delivery outcomes,<sup>14</sup> there is still a lack of an independent scoring system with strong objectivity and good repeatability for Chinese women. Thus, we discussed the third issue by incorporating these maternal and fetal factors, aiming to quantify the risk for each factor and identify high-risk groups of pregnant women. Because it is illegal to identify the sex of the fetus before delivery, this study included nine factors that could be determined before delivery, and a risk scoring system for EmCS has been established. The results showed that this system can effectively identify women with a high risk of EmCS, and was superior to any single factor in prediction. In clinical circumstances, the information of maternal age, BMI before pregnancy, BMI at delivery and childbirth history can be obtained

after admission, and other factors including the weight of the newborns, the quantity of amniotic fluid and umbilical cord status can be evaluated by ultrasound before delivery. Thus, this risk scoring system is simple, easy to operate and has good repeatability in clinical practice. If the score is too high, we should pay close attention to the situation of mothers and babies during childbirth. The limitation of this study was that it was a retrospectively designed study, and the conclusions were only based on the results of the present study. The clinical significance should be tested and validated in a further prospective study.

## CONCLUSION

This study analyzed the differences in 22 maternal and fetal factors between NVD and EmCS groups, then identified nine factors (maternal age  $\geq 30$ , BMI  $\geq 20.5$  before pregnancy, BMI  $\geq 28$  at delivery, nulliparous, abnormal characteristics of the amniotic fluid, abnormal umbilical cord, abnormal quantity of amniotic fluid, neonatal weight  $\geq 3550$  g, and induction of labor) that were associated with a high risk of EmCS. An effective risk scoring system based on the nine factors has been developed to quantitatively assess the risk of EmCS and classify women into different risk groups from the very beginning of labor. The practical value will be determined in further prospective observation studies.

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