

## Maternal near miss: The crown of death in obstetrics

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### ABSTRACT

**Aim:** Maternal near-miss is defined as a woman who survives a life-threatening condition during pregnancy, delivery, or postpartum period. Assessing near-miss cases offers a critical opportunity to apply standardized care before a fatal outcome occurs. However, there is currently no centralized database for near-miss cases in Turkey. The present study aimed to estimate the incidence and causes of near-miss cases at a tertiary hospital setting in Turkey.

**Methods:** This retrospective observational study was carried out at a university-affiliated research and training hospital between January 2019 and December 2022. A total of 150 maternal near-miss cases and 4 maternal deaths were analyzed. Demographic, obstetrics and laboratory characteristics were presented.

**Results:** The near miss ratio was 4.25 per 1,000 live births, while the maternal mortality ratio was 11.34 per 100,000 live births. It was found that 38% of patients in the maternal near-miss group had received regular prenatal care and 20% of the near-miss cases were referred to our center after delivering in an external hospital. Hypertensive disorders of pregnancy accounted for 40% of near-miss cases, while hemorrhagic conditions comprised 38%. Placental abruption and obstetric-related disseminated intravascular coagulation made up 12.6% of all cases, and other systemic diseases accounted for 9.3%.

**Conclusion:** Similar to maternal mortality, hypertensive disorders and hemorrhagic conditions were the leading causes of near-miss events. However, given the higher incidence of near-miss cases compared to maternal deaths, we suggest that identifying near-miss events could play a crucial role in preventing maternal mortality.

**Keywords:** Maternal morbidity, maternal mortality, near-miss.

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Received: 2025-07-12

Accepted: 2025-08-14 / Published: 2025-08-20

### Introduction

Maternal mortality rate is one of the key indicators reflecting the overall developmental status of a country. However, focusing solely

on maternal deaths overlooks the fact that many women experience severe health complications without a fatal outcome [1]. Therefore, in 2009, the World Health Organization introduced a standardized definition for near-miss cases to accurately identify such instances and establish a standardized approach. According to this definition, a near-miss case involves a woman who survives life-threatening conditions during pregnancy, childbirth, or within 42 days

postpartum [2]. Assessing near-miss cases is particularly valuable, as they occur more frequently than maternal deaths and, due to their clinical similarity, offer a critical opportunity to apply standardized care before a fatal outcome occurs [3]. Nowadays, the rate of near-miss cases is used as a measure to evaluate the quality of maternal healthcare [4].

The defining criteria include three main approaches for maternal near miss cases: clinical criteria based on specific diseases, intervention-based criteria, and criteria based on organ system dysfunction [2]. The primary pathologies contributing to maternal near-miss cases include hemorrhage, hypertensive disorders of pregnancy, sepsis, abruption placenta, placental invasion anomalies and uterine rupture [5].

It is estimated that each year, approximately 536,000 women die due to pregnancy- or childbirth-related complications, and more than 250,000 of these deaths are considered largely preventable [6,7]. Furthermore, around 99% of these cases are believed to occur in developing countries [6]. According to WHO criteria, the global prevalence of life-threatening maternal conditions in the general population is estimated to exceed 18.67 per 1,000 live births [8]. Prevalence rates vary depending on the criteria used. A systematic review utilizing disease-specific criteria reported that the frequency of maternal near-miss events ranged from 0.80% to 8.23%, whereas studies based on organ-system dysfunction criteria found rates between 0.38% and 1.09% [2]. Similarly, studies using management-based criteria reported prevalence rates ranging from 0.01% to 2.99% [2,9]. Depending on the region and the specific criteria applied, the prevalence of near miss cases has been shown to vary from 0.5% to over 40% of all live birth-related hospital admissions [10]. However, there is

currently no centralized database for near miss cases in Turkey.

The present study aimed to estimate the incidence of maternal near-miss cases and identify the main contributing factors within a tertiary hospital setting in Turkey, with the goal of improving the quality of maternal care.

### Materials and Methods

This retrospective observational study was carried out at a university-affiliated research and training hospital between January 2019 and December 2022. Ethical approval was obtained from the local ethics committee (Approval No: KAEK-25 2022/12-21).

All pregnant women and those within 42 days postpartum who were admitted to the intensive care unit of our hospital were identified using the hospital's electronic medical records. A total of 150 maternal near-miss cases and 4 maternal deaths were recorded. Near-miss cases were defined based on the organ dysfunction criteria established by the WHO in 2009, incorporating clinical, laboratory, and management-based indicators (Table 1) [2].

Women admitted to the intensive care unit during pregnancy or within 42 days postpartum who did not meet the WHO near-miss criteria were excluded from the study.

Participants were categorized into four groups: Group 1 – Hypertensive disorders; Group 2 – Hemorrhagic conditions; Group 3 – Placental abruption and disseminated intravascular coagulation (DIC); and Group 4 – Other systemic diseases. Group 1 comprised patients with HELLP syndrome (hemolysis, elevated liver enzymes, and low platelets), eclampsia, or severe preeclampsia. Group 2 included cases of postpartum hemorrhage, placental invasion anomalies, and uterine rupture. Group 3 consisted of patients who

**Table 1.** The WHO maternal near miss criteria.

| Clinical Criteria  | Laboratory-based Criteria                                    | Management-based Criteria  |
|--|--|--|
| Shock  | pH< 7.1 (severe acidosis)                                    | Cardio-pulmonary resuscitation                                       |
| Gasping  | PaO <sub>2</sub> /FiO <sub>2</sub> <200 mmHg                 | Continuous use of vasoactive drugs                                   |
| Acute cyanosis   | Lactate >5   | Dialysis for acute renal failure                                     |
| Clotting failure   | Oxygen saturation <90% for ≥60 minutes                       | Transfusion of ≥5 units of red blood cells                           |
| Respiratory rate >40/min (severe tachypnea) or <6/min (severe bradypnea) | Loss of consciousness and the presence of ketoacids in urine | Intubation and ventilation for ≥60 minutes not related to anesthesia |
| Oliguria non-responsive to fluids or diuretics                           | Creatinine ≥300 µmol/L or ≥3.5 mg/dl                         | Hysterectomy due to infection or hemorrhage                          |
| Loss of consciousness lasting ≥12 hours                                  | Bilirubin >100 µmol/L or >6 mg/dl                            |  |
| Loss of consciousness and absence of heart beta                          | Acute thrombocytopenia (<50x10 <sup>3</sup> /µL)             |  |
| Stroke   |  |  |
| Uncontrollable fit/status epilepticus                                    |  |  |
| Jaundice in the presence of preeclampsia                                 |  |  |

developed placental abruption and obstetric-related DIC. Group 4 encompassed individuals diagnosed with pulmonary edema, sepsis, shock, cardiac arrest, or systemic lupus erythematosus.

Demographic characteristics such as age, parity, gravida, body mass index, gestational week, birth week, delivery mode, chronic illnesses, laboratory results, hospitalization time, the amount of blood and blood product transfusion, surgical interventions were collected from the hospital's electronic medical records.

### Statistical analysis

Shapiro Wilk test was used to assess the normality of the distribution of variables. Descriptive statistics were expressed as mean ±

standard deviation with minimum and maximum values for continuous variables and frequency (percentage) for categorical variables. The SPSS 22.0 (SPSS Inc; Chicago, IL, USA) software was used for all statistical analyses.

### Results

During the study period, a total of 35,246 live births and 473 stillbirths occurred at our hospital. The study included 150 maternal near-miss cases and 4 maternal deaths. The near miss ratio was calculated as 4.25 per 1,000 live births, while the maternal mortality ratio was 11.34 per 100,000 live births. The demographic and peripartum characteristics of near-miss and maternal mortality cases were presented in Table 2.

**Table 2.** The demographic and peripartum characteristics of near-miss and maternal mortality cases.

| Parameters                 | Near-miss Cases |                 | Maternal Mortality |                 |
|----------------------------|-----------------|-----------------|--------------------|-----------------|
|                            | Min-max         | Mean $\pm$ SD   | Min-max            | Mean $\pm$ SD   |
| Age (years)                | 20-44           | 30.7 $\pm$ 5.9  | 23-38              | 31.8 $\pm$ 6.5  |
| Gravida (n)                | 1-15            | 3 $\pm$ 1.6     | 1-4                | 2 $\pm$ 1.4     |
| Parity (n)                 | 0-7             | 1.6 $\pm$ 1.39  | 0-2                | 0.8 $\pm$ 0.9   |
| BMI (kg/m <sup>2</sup> )   | 25-44           | 31.6 $\pm$ 2.11 | 30-32              | 30.8 $\pm$ 0.9  |
| Gestational week (week)    | 13-41           | 33.8 $\pm$ 4.3  | 25-40              | 32.3 $\pm$ 7.8  |
| Birth weight (gram)        | 555-4500        | 2340 $\pm$ 915  | 640-3645           | 2152 $\pm$ 1490 |
| Transfused unit (n)        |                 |                 |                    |                 |
| -Erythrocyte transfusion   | 1-14            | 4.6 $\pm$ 2.5   | 0-13               | 7 $\pm$ 6.6     |
| -Fresh frozen plasma       | 1-27            | 4 $\pm$ 3.4     | 2-13               | 8.7 $\pm$ 5.9   |
| -Thrombocyte suspension    | 1-6             | 2.4 $\pm$ 1.4   | 0-8                | 4 $\pm$ 4       |
| Fibrinogen use (gram)      | 1-8             | 2.7 $\pm$ 1.8   | 0-8                | 5.3 $\pm$ 4.6   |
| Hospitalization time (day) | 2-25            | 7 $\pm$ 3.8     | 2-8                | 5.5 $\pm$ 3     |

**Table 3.** The prepartum and postpartum laboratory characteristics of near-miss and maternal mortality cases.

| Parameters                                 | Near-miss Cases |                   | Maternal Mortality |                   |
|--|-----------------|-------------------|--------------------|-------------------|
|  | Min-max         | Mean $\pm$ SD     | Min-max            | Mean $\pm$ SD     |
| Prepartum hemoglobin (g/dl)                | 4-14            | 9.8 $\pm$ 2.4     | 11-14              | 12.3 $\pm$ 1.3    |
| Postpartum hemoglobin (g/dl)               | 4-13            | 7.8 $\pm$ 1.61    | 4-8                | 6 $\pm$ 2         |
| Prepartum hematocrit (%)                   | 12-43           | 29.7 $\pm$ 7.1    | 32-43              | 36.5 $\pm$ 4.6    |
| Postpartum hematocrit (%)                  | 14-40           | 24.2 $\pm$ 4.59   | 16-25              | 19.7 $\pm$ 4.7    |
| Prepartum platelet (x10 <sup>3</sup> /μL)  | 14-510          | 181.1 $\pm$ 91.6  | 32-191             | 121 $\pm$ 77.7    |
| Postpartum platelet (x10 <sup>3</sup> /μL) | 21-380          | 134 $\pm$ 75.3    | 14-146             | 73.3 $\pm$ 67     |
| Prepartum AST (IU/L)                       | 8-940           | 69.5 $\pm$ 148.9  | 18-357             | 116.3 $\pm$ 162   |
| Postpartum AST (IU/L)                      | 8-1785          | 90.4 $\pm$ 228.1  | 122-6592           | 2841.3 $\pm$ 3356 |
| Prepartum ALT (IU/L)                       | 2-597           | 44 $\pm$ 93.9     | 8-305              | 86.8 $\pm$ 145.6  |
| Postpartum ALT (IU/L)                      | 5-990           | 50 $\pm$ 113.8    | 36-1876            | 1252 $\pm$ 1053   |
| Prepartum creatinine (mg/dl)               | 0.1-4           | 0.61 $\pm$ 0.36   | 0.5-1.1            | 0.7 $\pm$ 0.3     |
| Postpartum creatinine (mg/dl)              | 0.1-4           | 0.63 $\pm$ 0.4    | 0.1-3              | 1.9 $\pm$ 0.9     |
| Prepartum INR                              | 0.7-8           | 0.97 $\pm$ 0.59   | 0.8-1.6            | 1.1 $\pm$ 0.3     |
| Postpartum INR                             | 0.7-1.9         | 0.9 $\pm$ 0.14    | 0.9-3              | 1.96 $\pm$ 1      |
| Prepartum aPTT (sec)                       | 2-90            | 26.9 $\pm$ 10.9   | 20-65              | 36 $\pm$ 19.9     |
| Postpartum aPTT (sec.)                     | 15-75           | 26.7 $\pm$ 8.1    | 26-56              | 44.7 $\pm$ 16.3   |
| Prepartum protrombin time (second)         | 9-22            | 11,9 $\pm$ 2,19   | 10-19              | 13 $\pm$ 4.1      |
| Postpartum protrombin time (sec)           | 9-20            | 12.1 $\pm$ 1.9    | 10-43              | 25.7 $\pm$ 16.6   |
| Prepartum fibrinogen (mg/dl)               | 30-790          | 378.3 $\pm$ 165.1 | 96-640             | 447 $\pm$ 250.8   |
| Postpartum fibrinogen (mg/dl)              | 80-890          | 322.9 $\pm$ 149.5 | 41-410             | 175 $\pm$ 204     |

**Table 4.** The distribution of near-miss and maternal mortality cases by diagnostic group.

| Parameters                                     |                           | Near-miss<br>(n) | %    | Maternal<br>mortality<br>(n) | %   |
|--|---------------------------|------------------|------|------------------------------|-----|
| <b>Hypertensive Disorders<br/>(Group 1)</b>    | Severe preeclampsia       | 27               | 45   | 0                            | 0   |
|  | Eclampsia                 | 6                | 10   | 1                            | 50  |
|  | HELLP                     | 27               | 45   | 1                            | 50  |
|  | Total                     | 60               | 100  | 2                            | 100 |
| <b>Hemorrhagic Conditions<br/>(Group 2)</b>    | Placenta accreta spectrum | 27               | 47.4 | 0                            | 0   |
|  | Postpartum haemorrhage    | 26               | 45.6 | 1                            | 25  |
|  | Uterine rupture           | 4                | 7    | 0                            | 0   |
|  | Total                     | 57               | 100  | 1                            | 100 |
| <b>Abruptio placenta and DIC<br/>(Group 3)</b> | Abruptio placenta         | 16               | 84.2 | 0                            | 0   |
|  | DIC                       | 3                | 15.8 | 1                            | 100 |
|  | Total                     | 19               | 100  | 1                            | 100 |
| <b>Other systemic conditions<br/>(Group 4)</b> | Pulmonary edema           | 3                | 21.4 | 0                            | 0   |
|  | Sepsis                    | 2                | 14.3 | 0                            | 0   |
|  | Shock                     | 7                | 50   | 0                            | 0   |
|  | Cardiac                   | 1                | 7.1  | 0                            | 0   |
|  | COVID-19                  | 1                | 7.1  | 0                            | 0   |
|  | Total                     | 14               | 100  | 0                            | 0   |
| <b>Total</b>                                   |                           | 150              | 100  | 4                            | 100 |

It was found that 38% of patients in the maternal near-miss group and 50% in the maternal mortality group had received regular prenatal care. Additionally, 20% of the near-miss cases and 25% of the maternal deaths were referred to our center after delivering in an external hospital. Blood transfusion was required in 77.2% of the near-miss group and 75% of the maternal mortality group. Fibrinogen was administered to 35.3% of patients in the near-miss group and to 50% of those in the maternal mortality group.

The preoperative and postoperative laboratory characteristics of near-miss maternal mortality cases were presented in Table 3.

The distribution of near-miss and maternal mortality cases by diagnostic group was presented in Table 4.

Hypertensive disorders of pregnancy accounted for 40% of near-miss cases, while hemorrhagic conditions comprised 38%. Placental abruption and obstetric-related DIC made up 12.6% of all cases, and other systemic diseases accounted for 9.3%. Among hypertensive disorders, severe preeclampsia and HELLP syndrome were the most common causes of maternal near-miss. In the hemorrhagic group, placenta accreta spectrum was the leading cause, followed by postpartum hemorrhage. In the maternal mortality group,

two cases were classified under hypertensive disorders. Of the remaining two maternal deaths, one was due to postpartum hemorrhage in the hemorrhagic group, and the other was attributed to DIC in the placental abruption and DIC group.

When surgical interventions in the near-miss and maternal mortality groups were analyzed, the cesarean section rate was found to be 86.7% (n=130) in the near-miss group and 75% (n=3) in the maternal mortality group. Postpartum hysterectomy was performed in 32.7% (n=49) of near-miss cases and in 50% (n=2) of maternal deaths. Major artery ligation was carried out in 18.7% (n=28) of the near-miss group and 50% (n=2) of the maternal mortality group. Balloon tamponade was applied in 19.3% (n=29) of near-miss cases, and compression sutures were used in 12.8% (n=19); in the maternal mortality group, these procedures were performed in 25% and 50% of cases, respectively. Segmental resection was conducted only in the near-miss group, in 5.3% (n=8) of patients.

### Discussion

The primary findings of this study revealed a maternal near-miss ratio of 4.25 per 1,000 live births and a maternal mortality ratio of 11.34 per 100,000 live births. Moreover, hypertensive disorders and hemorrhagic conditions emerged as the most common underlying causes of both near-miss events and maternal deaths.

The prevalence of maternal near-miss varies significantly based on the specific criteria applied and the region studied, with notably higher rates observed in low-income countries. A recent study from India reported a near-miss ratio of 2.95 per 1,000 births, while another Indian study found a near-miss rate of 22.7 per 1,000 births alongside a maternal mortality ratio of 899 per 100,000 live births [11, 12]. Reports from other low-income countries using

the WHO near-miss criteria revealed near-miss ratios of 19.8% in Nigeria, 1.21% in Egypt, and 3.49% in Peru [13]. Furthermore, a systematic review indicated that the median maternal near-miss ratio in low- to middle-income countries was 15.9 per 1,000 live births, with an interquartile range of 8.9 to 34.7 [14].

The maternal near-miss and mortality rates observed in Turkey are generally consistent with findings from international studies. Uygur et al. reported an overall maternal near-miss ratio of 2.47 per 1,000 live births and a maternal mortality ratio of 13.46 per 100,000 live births [15]. Similarly, Oğlak et al. found the near-miss incidence to be 5.03 per 1,000 live births, a figure comparable to those reported in Australia (7.0/1,000) and the Netherlands (7.1/1,000) [1,16,17]. In the same study, the maternal mortality ratio was 8.1 per 100,000 live births [1]. Tonyalı et al. reported a near-miss ratio of 2.04 per 1,000 live births, while another recent study documented a rate of 2.31 per 100 births [4,18]. In 2025, Erdem et al. reported a maternal near-miss prevalence of 218 per 100,000 live births [19]. A study by Akın Evsen revealed a near-miss incidence of 2.31 per 100 live births and a maternal mortality incidence of 202.4 per 100,000 births [18]. Considering these ratios, these findings demonstrate that the maternal near-miss and mortality ratios in the current study are consistent with national data.

Hemorrhagic and hypertensive disorders are considered the primary causes of maternal near-miss, largely due to delays in diagnosis, timely management of complications, and referral to appropriate healthcare facilities. Several studies conducted in different regions of India have identified preeclampsia and hemorrhage as the leading etiological factors for maternal near-miss cases, which aligns with the findings of the present study [20-22]. Similarly, Nakimuli et al. reported that the most common cause of



maternal near-miss was preeclampsia (7%), followed by postpartum hemorrhage (6.7%) in Uganda [23]. In the study by Verma et al., published in 2023, the majority of maternal near-miss cases (38.46%) were attributed to hypertensive disorders [24]. A review of studies published from Turkey also suggests that the causes of maternal near-miss are similar to those reported globally. In the study of Akın Evsen, the most common etiologic factors of maternal near miss were haemorrhagic diseases (52.1%) followed by hypertensive diseases (33.2%) [18]. Likewise, Cengiz et al revealed that the most frequently observed diagnosis was preeclampsia (41.5%), followed by obstetric hemorrhage (29.2%) [6]. In the study by Uygur et al., the analysis showed that the leading cause of maternal death was hypertensive disorders of pregnancy, followed by sepsis. In contrast, obstetric hemorrhage was identified as the most common cause of maternal near-miss, followed by hypertensive disorders [15].

One of the most important goals for healthcare professionals is to reduce maternal mortality and morbidity. In this circumstance, international organizations and local societies play a crucial role in developing strategies. We suggest that definition of maternal near-miss criteria is one of this strategies that could improve perinatal outcomes. Awareness of these cases is essential for ensuring timely and effective intervention.

The present study has several limitations. First, its retrospective design makes it susceptible to recognition bias. Second, the data were derived from a single-center, which may limit external validity. Third, the relatively small sample size restricts the generalizability of the findings. Lastly, as a tertiary care hospital, our facility may receive more severe cases, potentially leading to an overestimation of maternal near-miss and mortality ratios.

## Conclusion

In conclusion, maternal near-miss events are critical markers for assessing the effectiveness and safety of maternity services in healthcare settings. Considering that hypertensive disorders and hemorrhagic conditions are the leading causes of maternal near-miss, the role of preventive strategies, early diagnosis, and effective management in reducing both near-miss events and maternal mortality should be recognized. However, given the higher incidence of near-miss cases compared to maternal deaths, we suggest that identifying near-miss events could play a crucial role in preventing maternal mortality.

**Funding:** *The author(s) received no financial support for the research, authorship, and/or publication of this article.*

**Conflict of Interest:** *The authors declare that they have no conflict of interest.*

**Ethical statement:** *Ethical approval was obtained from the local ethics committee (Approval No: KAEK-25 2022/12-21).*

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