Maternal and fetal outcome among patients requiring high-dependency unit admission: a five-year prospective study

Abstract

Objective: to test the hypothesis whether the cause of admission to high-dependency unit (HDU) could negatively affect the maternal and fetal outcome or not.

Methods: This five-year prospective study included 346 pregnant patients who were required admission to HDU. Patients were divided into two groups based on the cause of admission to HDU: group one with obstetric complications (n=204) and group two with medical complications (n=142). Obstetric outcome was the main outcome measure. Data was collected and analyzed.

Results: Patients in the medical group were more prone for preterm delivery (p<0.001), development of organ failure (p<0.05), persistent hypertension (p<0.001), persistent left ventricle dysfunction (p<0.001), pulmonary embolism (p<0.05), longer time of stay at HDU, more transfer to specialized ICU and longer time for recovery (p<0.001) with higher maternal mortality (p=0.001) compared to patients in the obstetric group. Neonates of patients in the medical group were more prone for development of fetal growth restriction (p=0.001), prematurity (p=0.001), intraterine fetal demise (p=0.05), admission to NICU (P<0.001) and neonatal death (p=0.05) compared to those in the obstetric group.

Conclusion: Although more patients with obstetric complications required admission to HDU, yet they still experienced better obstetric outcome than those with underlying medical disorders.

Keywords: high dependency unit, maternal outcome, obstetric outcome, fetal outcome

Abbreviations: HDU, high dependency unit; ICU, intensive care unit; SGA, small for gestational age; IUFD, intrauterine fetal demise

Introduction

Pregnancy and delivery have a potential for maternal life threatening complications which indicate the provision of a higher level of health care than that traditionally provided in standard wards.

High dependency unit (HDU) was defined as a level of care which lies in between the general ward and an intensive care unit (ICU) and provides health care to patients with single organ failure or those who are at a high risk of developing severe maternal morbidity.

The aim of this study was to test the hypothesis whether the cause of admission to high-dependency unit (HDU) could negatively affect the maternal and fetal outcome or not.

Materials and methods

This prospective observational study was conducted at the department of Obstetrics and Gynecology in collaboration with the Anesthesiology & Critical Care and Pediatrics departments at Menoufia University hospital, Menoufia governorate, Egypt during the period between September 2012 and September 2017. Menoufia University hospital is the biggest tertiary health care and referral hospital in Menoufia governorate with well equipped high dependency unit (HDU) containing 25 beds complying with the standard international recommendations.

Ethical approval was granted from Menoufia Faculty of Medicine ethical committee and an informed consent was obtained from all participants prior to commencement of the study.

Out of 21,364 parturients in the hospital during the five-year study period, 352 patients with single organ failure had required HDU admission during their course of treatment were included in this study. During the follow up period, six patients dropped out and 346 patients completed the study (Figure 1).

Patients with multiple organ failure and those required primary admission to intensive care unit (ICU) were excluded from the study. For better interpretation of results, patients were divided into two groups regarding the cause of admission to HDU as follows:

a. Obstetric complications group (n=204): Obstetric complications were defined as maternal morbidity originating from any cause related directly to pregnancy or its management...
during pregnancy, delivery or postpartum till the end of the puerperium.

b. Medical complications group (n=142): Medical complications were defined as maternal morbidity arising from any pre-existing medical disorders.

Patients were followed from the time of admission to HDU throughout delivery till three months after discharge from the hospital. All patients were managed individually with standard invasive or non-invasive measures as indicated.

Figure 1 Flow diagram of recruitment and retention of patients in the study.

Outcome measures

Maternal outcome: included interventions at HDU (cause of admission, investigations, length of stay, eventual outcomes as transfer to specialized intensive care unit, ICU), mode of delivery, postpartum hemorrhage, defective lactation, venous thromboembolism and maternal mortality.

Fetal and neonatal outcome: small for gestational age (SGA) defined as a birth weight < 5th percentile, preterm labour (delivery<37 weeks), intrauterine fetal demise (IUD), admission to neonatal intensive care unit (NICU) and neonatal death (defined as death during the first four weeks after delivery).

Statistical analysis

Data was analyzed by SPSS (statistical package for the social science software) version 22. Quantitative data was analyzed by applying student t-test or Mann-Whitney test as required while qualitative data was analyzed by applying Chi-square test and Fisher’s exact test as required. P-value less than 0.05 are significant and P-value less than 0.001 are highly significant.

Results

There was no significant difference between the two groups regarding maternal age, parity and body mass index (p>0.05) with more patients in the medical group booked for antenatal care (p<0.05) as depicted in Table 1.

Table 1 Maternal characteristics and indications of admission to HDU

|                      | Obstetric group (n=204) | Medical group (n=142) | Student t-test | P-value |
|----------------------|-------------------------|-----------------------|----------------|---------|
| Age (years)          | 31.4±4.3                | 30.9±4.6              | 1.03           | >0.05   |
| Parity               | 2.4±1.3                 | 2.2±1.2               | 1.45           | >0.05   |
| Body mass index (Kg/m²) | 26.2±3.3              | 25.8±3.6              | 1.07           | >0.05   |
| Booking for antenatal care | 168                  | 102†                 | 4.81           | <0.05   |

Indications of admission to HDU: n(%) Patients in the obstetric group were more commonly presented with pallor, pyrexia and tachycardia (p<0.001) while patients in the medical group were more commonly presented with cyanosis, dyspnea, hypertension and splenomegaly (p<0.001). Patients in the obstetric group required more blood gases and blood cultures performed (p<0.001) while patients in the medical group required more ECG, Echocardiography, CT brain, Chest X ray and Angiography (p<0.001). During their course of treatment, patients in the obstetric group received more antibiotics, magnesium sulfate and blood products transfusions (p<0.001) while those in the medical group received inotropic drugs, central venous line insertion, hemodialysis and mechanical ventilation (p<0.001).

Table 2 reveals clinical features, investigations and interventions at HDU. Patients in the obstetric group were more commonly presented with pallor, pyrexia and tachycardia (p<0.001) while patients in the medical group were more commonly presented with cyanosis, dyspnea, hypertension and splenomegaly (p<0.001). Patients in the obstetric group required more blood gases and blood cultures performed (p<0.001) while patients in the medical group required more ECG, Echocardiography, CT brain, Chest X ray and Angiography (p<0.001). During their course of treatment, patients in the obstetric group received more antibiotics, magnesium sulfate and blood products transfusions (p<0.001) while those in the medical group received inotropic drugs, central venous line insertion, hemodialysis and mechanical ventilation (p<0.001).

In regard to maternal outcome, patients in the medical group were more prone for preterm delivery (p<0.001), development of organ

Citation: Masood A, Gaballah K, Omar Z. Maternal and fetal outcome among patients requiring high-dependency unit admission: a five-year prospective study. Obstet Gynecol Int J. 2018;9(1): 00310. DOI: 10.15406/ogij.2018.09.00310
failure (p<0.05), persistent hypertension (p<0.001), persistent left ventricle dysfunction (p<0.001) and pulmonary embolism (p<0.05). Also, patients in the medical group have longer time of stay at HDU, more transfer to specialized ICU and longer time for recovery (p<0.001) with higher maternal mortality (p<0.001) compared to patients in the obstetric group. The most common causes of maternal mortality in the obstetric group were obstetric hemorrhage and sepsis while rheumatic heart disease and chronic liver disease were in the medical group as shown in (Table 3).

Table 2 Clinical features, investigations and interventions at HDU

| Clinical features:         | Obstetric group (n=204) | Medical group (n=142) | Chi square test | P-value |
|---------------------------|-------------------------|-----------------------|-----------------|---------|
| Pallor.                   | 104                     | 26                    | 36.7            | <0.001  |
| Jaundice.                 | 14                      | 18                    | 2.71            | >0.05   |
| Cyanosis.                 | 6                       | 24                    | 18.8            | <0.001  |
| Dyspnea.                  | 8                       | 28                    | 20.7            | <0.001  |
| Pyrexia (>38°C).          | 56                      | 12                    | 17.9            | <0.001  |
| Bradycardia.              | 110                     | 48                    | 12.8            | <0.001  |
| Hypertension (>160/100mmHg). | 6               | 8                     | 0.95            | >0.05   |
| Hypotension (<90/60mmHg). | 44                      | 48                    | 5.8             | <0.05   |
| Abnormal bleeding.        | 28                      | 32                    | 3.9             | <0.05   |

Investigations:

| Investigations:           | Obstetric group (n=204) | Medical group (n=142) | Chi square test | P-value |
|--------------------------|-------------------------|-----------------------|-----------------|---------|
| Blood gases.             | 42                      | 12                    | 8.5             | <0.05   |
| Blood culture.           | 44                      | 8                     | 15.4            | <0.001  |
| ECG.                     | 22                      | 78                    | 77.2            | <0.001  |
| Echocardiography.        | 10                      | 46                    | 44.6            | <0.001  |
| CT brain.                | 6                       | 18                    | 10.8            | <0.001  |
| CT abdomen.              | 12                      | 8                     | 0.02            | >0.05   |
| Chest X ray.             | 24                      | 56                    | 34.5            | <0.001  |
| Angiography.             | 14                      | 38                    | 24.4            | <0.001  |

Interventions:

| Interventions:           | Obstetric group (n=204) | Medical group (n=142) | Chi square test | P-value |
|--------------------------|-------------------------|-----------------------|-----------------|---------|
| Antibiotics.             | 112                     | 36                    | 28.6            | <0.001  |
| Inotropics.              | 22                      | 48                    | 26.1            | <0.001  |
| Magnesium sulfate.       | 50                      | 14                    | 10.9            | <0.001  |
| Central venous line.     | 54                      | 98                    | 59.8            | <0.001  |
| Blood products.          | 92                      | 28                    | 22.7            | <0.001  |
| transfusion. Hemodialysis. | 8                  | 26                    | 17.9            | <0.001  |
| Mechanical ventilation.  | 12                      | 32                    | 19.4            | <0.001  |

Table 3 Maternal outcome

| Organ failure:            | Obstetric group (n=204) | Medical group (n=142) | Chi square test | P-value | Odd’s ratio at 95% Confidence interval |
|--------------------------|-------------------------|-----------------------|-----------------|---------|---------------------------------------|
| Gestational age at delivery | 36.5±2.3               | 34.3±3.4              | 7.18†           | <0.001  | -                                     |
| Heart failure.           | 5                       | 26                    | 23.9            | <0.001  | 0.11(0.04-0.3)                        |
| Respiratory failure.     | 3                       | 12                    | 8.2*            | <0.05   | 0.16(0.04-0.58)                       |
| Renal failure.           | 8                       | 24                    | 15.3            | <0.001  | 0.2(0.09-0.46)                        |
| Hepatic encephalopathy.  | 4                       | 10                    | 4.34*           | <0.05   | 0.26(0.08-0.86)                       |
| Disseminated intravascular coagulopathy. | 18   | 10                    | 0.16            | >0.05   | 1.28(0.57-2.86)                       |
| Persistent hypertension  | 14                      | 58                    | 56.6            | <0.001  | 0.11(0.06-0.2)                        |
| Persistent left ventricle dysfunction. | 3   | 19                    | 17.9*           | <0.001  | 0.1(0.03-0.3)    |
Table Continued.....

|                                | Obstetric group (n=204) | Medical group (n=142) | Chi square test | P-value | Odd's ratio at 95% Confidence interval |
|--------------------------------|-------------------------|-----------------------|-----------------|---------|----------------------------------------|
| Pulmonary embolism             | 6                       | 16                    | 8.4             | <0.05   | 0.24(0.09-0.63)                        |
| **Outcome at HDU:**            |                         |                       |                 |         |                                        |
| Duration of stay (days)        | 5.6±4.3                 | 10.4±4.2              | 10.31†          | <0.001  |                                        |
| Transfer to specialized ICU    | 8                       | 28                    | 20.7            | <0.001  |                                        |
| **Maternal mortality**         |                         |                       |                 |         |                                        |
| Obstetric hemorrhage           | 4                       | 5                     | 14.4†           | <0.001  | 0.14(0.05-0.42)                        |
| Sepsis                         | 3                       | 5                     | 9.05*           | <0.05   | 0.18(0.06-0.57)                        |
| Pulmonary edema                | 5                       | 18                    |                 |         |                                        |
| Pulmonary embolism             | 5                       | 16                    |                 |         |                                        |
| AF                             | 3                       | 1                     |                 |         |                                        |
| Stroke                         | 1                       | 1                     |                 |         |                                        |
| Hepatic encephalopathy         | 2                       | 2                     |                 |         |                                        |
| Bleeding varices               | 2                       | 2                     |                 |         |                                        |

†Student t-test, *Fisher’s exact test, ICU, intensive care unit; AF, atrial fibrillation

Table 4 revealed fetal and neonatal outcome. Neonates of patients in the medical group were more prone for development of fetal growth restriction (p<0.001), prematurity (p<0.001), intrauterine fetal demise (p<0.05), admission to NICU (P<0.001) and neonatal death (p<0.05) compared to those in the obstetric group.

Table 4 Fetal and neonatal outcome:

|                                | Obstetric group (n=204) | Medical group (n=142) | Chi square test | P-value | Odds ratio at 95% confidence interval |
|--------------------------------|-------------------------|-----------------------|-----------------|---------|--------------------------------------|
| Small for gestational age      | 12                      | 38                    | 27.8            | <0.001  | 0.17(0.09-0.34)                      |
| Prematurity                    | 10                      | 26                    | 14.7            | <0.001  | 0.23(0.11-0.49)                      |
| Intrauterine fetal demise      | 4                       | 14                    | 9.05*           | <0.05   | 0.18(0.06-0.57)                      |
| Admission to NICU              | 12                      | 34                    | 22.2            | <0.001  | 0.21(0.1-0.4)                        |
| Neonatal sepsis                | 8                       | 8                     | 0.24            | >0.05   | 0.68(0.25-1.87)                      |
| Neonatal death                 | 6                       | 16                    | 8.4             | <0.05   | 0.24(0.09-0.63)                      |

Discussion

During the five-year period of the study, 21,364 patients delivered at our hospital with 352 of them required admission to HDU. Thus obstetric admission in HDU at our institution was 16.5 per 1000 deliveries.

The incidence varies widely among developing and developed countries with range of 10.2 in four years in Dublin in Ireland (3), 11.2 per 1000 deliveries over four years in India (5), and 26.7 in 23 years in UK.4

In this study, about half of the admitted patients in the obstetric group suffered major obstetric hemorrhage (52.8%) followed by sepsis which constitutes 26.3%. On the other hand, cardiovascular disease (chronic hypertension and rheumatic heart disease) was the most common cause of maternal admission to HDU in the medical group (45.2%).

Previous studies ranked obstetric hemorrhage and hypertensive disorders as the most common causes of admission to HDU.5−8

At our institution, the rate of PPH as recently reported was 2.4% 9 while chronic hypertension affected 7%10 and rheumatic heart disease complicates pregnancy in 2.5% of patients attended our antenatal care clinic.11

In the current study, patients in the medical group were more prone for poorer obstetric outcome in terms of preterm delivery, development of organ failure, persistent hypertension, persistent left ventricle dysfunction, pulmonary embolism, longer time of stay at HDU, more transfer to specialized ICU and longer time for recovery with higher maternal mortality with their neonates developed fetal growth restriction, intrauterine fetal demise, admission to NICU and neonatal death compared to patients in the obstetric group.

Maternal mortality among patients admitted to HDU in this study was 6.4% and perinatal mortality was 11.6%.

A previous Indian studies reported maternal and perinatal mortality of 12.3% and 12.3% among patients required admission to HDU9 and 52% and 21.6% among patients admitted to ICU respectively,12 while in developed countries maternal mortality of HDU admissions was nil and perinatal mortality was 1.05% to 8%.3,7

The poorer obstetric outcome among patients admitted to HDU was mostly attributed to the underlying medical disorders10,11,13,14
rather than obstetric complications which were readily responded to proper treatment strategies for obstetric hemorrhage and sepsis.\textsuperscript{3}

Maternal mortality in this study was secondary to rheumatic heart disease and chronic liver disease. In the presence of maternal heart disease, the circulatory changes of pregnancy may result in decompensation or death of the mother and/or the fetus secondary to development of pulmonary edema or atrial fibrillation particularly in patients with severe mitral stenosis.\textsuperscript{15–17}

The physiological increase in plasma volume during pregnancy can worsen portal hypertension in pregnant women with chronic hepatitis, resulting in increased risk of variceal bleeding and maternal mortality in up to 18\% with higher fetal loss rate.\textsuperscript{18}

Emphasis on the importance of multidisciplinary preconception and antenatal care of patients with medical disorders can minimize the incidence of maternal and fetal compromise as well as the need for HDU admission.

The large sample size and the prospective nature of this study constituted the meaningful importance of its results.

Inability to recruit and report the obstetric outcome among patients with multiple organ failure who were primarily admitted to ICU was unintended limitation of this study.

Future research should explore new strategies to improve the obstetric outcome among mothers with medical disorders.

Conclusion

Although more patients with obstetric complications required admission to HDU, yet they still experienced better obstetric outcome than those with underlying medical disorders.

Acknowledgements

The authors would like to acknowledge the contribution of the residents and nursing staff of the delivery ward, NICU and HDU of Menoufia University Hospital.

Conflict of interest

Authors declare there is no conflict of interest in publishing the article.

References

1. Zeeman GG. Obstetric critical care: a blueprint for improved outcomes. Crit Care Med. 2006;34(9 Suppl):S208–14.
2. Guidelines on admission to and discharge from Intensive care and High Dependency units. Department of Health/NHS Executive, London: DOH; 1996.
3. Ryan M, Hamilton V, Bowen M, et al. The role of high-dependency unit in a regional obstetric hospital. Anaesthesia. 2000;55(12):1155–8.
4. Hazelgrove JF, Price C, Pappachan VJ, et al. Multicenter study of obstetric admissions to 14 intensive care units in southern England. Crit Care Med. 2001;29(4):770–5.
5. Dattaray C, Mandal D, Shankar U, et al. Obstetric patients requiring high-dependency unit admission in a tertiary referral centre. Int J Crit Illn Inf Sci. 2013;3(1):31–5.
6. Saravankumar K, Davies L, Lewis M, et al. High dependency care in obstetric setting in the UK. Anaesthesia. 2008;63(10):1081–6.
7. Leung NY, Lau AC, Chan KK, et al. Clinical characteristics and outcomes of obstetric patients admitted to Intensive care unit: A 10–year retrospective review. Hong Kong Med J. 2010;16(1):18–25.
8. Zwart JJ, Dupis JR, Ory F, et al. Obstetric intensive care unit admission: A 2 year national wide population–based cohort study. Intensive Care Med. 2010;36(2):256–63.
9. Kandeel M, Sanad Z, Ellakwa H, et al. Management of postpartum hemorrhage with intrauterine balloon tamponade using a condom catheter in an Egyptian setting. Int J Gynaecol Obstet. 2016;135(3):272–275.
10. Rezk M, Ellakwa H, Gamal A, et al. Maternal and fetal morbidity following discontinuation of antihypertensive drugs in mild to moderate chronic hypertension: A 4-year observational study. Pregnancy Hypertens. 2016;6(4):291–294.
11. Rezk M, Gamal A. Maternal and fetal outcome in women with rheumatic heart disease: a 3-year observational study. Arch Gynecol Obstet. 2016;294(2):273–8.
12. Karnad DR, Lapsia V, Krishnan A, et al. Prognostic factors in obstetric patients admitted to an Indian intensive care unit. Crit Care Med. 2004;32(6):1294–9.
13. Rezk M, Dawood R, Badr H. Maternal and fetal outcome in women with antiphospholipid syndrome: a three-year observational study. J Matern Fetal Neonatal Med. 2016;29(24):4015–9.
14. Rezk M, Ellakwa H, Al-Halaby A, et al. Predictors of poor obstetric outcome in women with systemic lupus erythematosus: a 10-year experience of a university hospital. J Matern Fetal Neonatal Med. 2017;30(17):2031–2035.
15. Scisicione AC, Ivester T, Largoza M, et al. Acute Pulmonary Oedema in pregnancy. Obstet Gynecol. 2003;101(3):511–15.
16. Essop MR, Nkomo VT. Rheumatic and Nonrheumatic Valvular Heart Disease: Epidemiology, Management, and Prevention in Africa. Circulation. 2005;112:3584–3591.
17. Konar H, Chaudhuri S. Pregnancy complicated by maternal heart disease: a review of 281 women. J Obstet Gynaecol India. 2012;62(3):301–6.
18. Sandhu BS, Sanyal AJ. Pregnancy and liver disease. Gastroenterol Clin North Am. 2003;32(1):407–436.

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