Outcome of the use of bakri balloon in obstetric hemorrhage

Abstract

Objective: To present the results obtained by using the Bakri Balloon to obstetric control hemorrhage.

Material and method: Prospective, observational study within two inpatient medical care units since January 1 to December 31 2016. Twenty patients with post-partum and trans cesarean hemorrhage. All of them were applied the Bakri Balloon because of failure to respond to uterotonic drug therapy. The following were analyzed:

a. Clinical data
b. Amount of bleeding before and after the balloon
c. Amount of postpartum or trans cesarean bleeding
d. Time between diagnosis and insertion, insufflation time and volume supplied
e. Hemoglobin levels and coagulation tests results initially, post- hemorrhage and post insertion
f. Use of and amount of haemo components ministered
g. The time use balloons, success, and complications

Results: The amount of bleeding after insertion, both in postpartum and trans cesarean was reduced and there was an adequate response in hemoglobin levels. The average time between hemorrhage diagnosis and balloon insertion was thirty minutes; average insufflation time, five minutes, and average volume supplied 400mL. Balloon’s average use time, five minutes, and average volume supplied 400mL. Successful post-surgery and after its application.

Conclusions: The Bakri Balloon proved to be a useful, quick and complication-free therapy for controlling obstetric hemorrhage.

Keywords: obstetric hemorrhage, Bakri Balloon

Background

Obstetric hemorrhage has increased its frequency1-3 for different reasons, among which are the increase in cesarean section and inadequate use of uterotonic, among others.4-6. It occupies the second cause of maternal deaths in our country and in some States, the first.5,9 Several strategies have been carried out for its reduction, without obtaining the expected results.3,4,9 Several decades ago the most practiced surgical treatment, once failed ls medical measures, was obstetric hysterecromy, which besides being a limiting fertility procedure in many patients, and was not without many complications.10,11 Nowadays, this behavior is increasingly questioned as the first therapeutic procedure and even in many countries it is the subject of medical litigation.12,13 Others, though effective, surgical techniques such as hypogastric artery ligation or devascularization of the uterine vessels require significant medical training and good experience, both often absent in our hospital means.14-16 The advent of procedures considered more conservative, such as the application of the Bakri intrauterine balloon, which requires a low and rapid learning curve, has demonstrated its usefulness for the effective control of obstetric hemorrhage and with minimal complications.17,18 Up to this date there are multiple reports on their use.19,20

The objective of this report is to present the results obtained in two hospitals with the use of Bakri’s intrauterine balloon to control obstetric hemorrhage both in the postpartum and in the trans-cesarean, its relation to the amount of hemorrhage and hemoglobin levels before and after its application.

Material and method

Prospective, longitudinal study performed in two hospital units: Hospital Silao General of the Ministry of Health of the State of Guanajuato and General Hospital Los Reyes, Secretary of Health of the State of Michoacan, in a span of 1. From January to December 31, 2016, of consecutive cases that went to the medical units in the period indicated. We included 20 patients who presented both postpartum and trans cesarean hemorrhage. Hemorrhage was considered from 500mL,2 and / or when the patient presented data on hypovolemia, characterized by hypotension (equal to or less than 90/60 for systolic / diastolic), tachycardia (equal to or greater than 100), 1/min), tachypnea (equal to or greater than 20/min), neurological changes (confusion, restlessness, decreased alertness), decreased urine output (less than 20mL/hour) and decreased capillary refill.4,22 The amount of bleeding was calculated in two ways: 1) by gravimetry, weighing with gauze and compresses during the procedures and 2) with collection bag type Brass -V, Calibrated obstetric drape (Maternova®). All the patients were initially treated with uterotonics. First choice oxytocin was used at a rate of 10U IV, diluted in 10 ml of Hartman solution.
passing slowly (1 min), if no response was obtained; Carbetocin was continued\(^{20}\) (Lab. Ferring) 100mcgs, 1ml IV in 1 minute. In all patients the hemoglobin admission unit determined, a post-hemorrhagic event twenty - four hours time after use with the ball or before removal thereof.\(^{23,24}\)

All patients without response to medical treatment were given an intrauterine Bakri balloon (Cook Medical, Bloomington, IN. USA, Demesa, Mexico City) (Figure 1).

The result is divided into:

Positive response: Control of bleeding no greater than 150-200mL March 2 in 24hours and recovered ban signs of hypovolemia. In these cases the balloon was removed between 24 to 48hours

Negative response: When control of bleeding was not obtained and hypovolemia data persisted. In these cases, another therapeutic measure was taken with the immediate removal of the balloon.

S and performed the following analysis:

i. Semenate gestation, gynecology-obstetric history, diagnosis of the cause of bleeding and medical treatment prior to application of the ball.

ii. The amount of bleeding before and after the application of the balloon.

iii. The amount of bleeding between the cases of delivery and cesarean, before and after the placement of the balloon in the same groups.

iv. The time between diagnosis and the affixing of the balloon, filling time and volume delivered.

v. Hemoglobin levels and coagulation tests 1 entrance to unit, the posterior bleeding hour and 24 hours after of treatment with blood components, at the moment of removal of the balloon.

vi. The cases that required the use of heme components and the amount of them used during the event.

vii. The percentage of success or failure of the procedure and time of use of the ball. Likewise complications analyzed by the use of the ball and if present arons cases of maternal death.

Descriptive statistics of numerical (mean and standard deviation) and categorical (proportions) data were used. Due to the small size of the sample, since the distribution of the data was not normal, non-parametric statistics were used. For the comparison of numerical variables before and after the use of the Bakri balloon in patients with uterine bleeding, the Wilcoxon test of the signed ranges was used. To compare numerical data of the patients with cesarean section and delivery, the amount of bleeding comparing them and the hemoglobin values at the different times indicated, the Mann Whitney U test was used. A value of p <0.05 was considered statistically significant. For the analysis of the data, the IBM SPSS version 22 statistical package was used.

Results

Twenty patients were treated during the study period. According to the seven points poured into the material and method, the following results were obtained:

a. The weeks of gestation , gynecological-obstetric history , cause of hemorrhage and the treatment instituted before the application of the Bakri balloon , appear and n Table 1.

b. The mean and standard deviation for age and weeks of gestation were respectively 24.6 (7.0) and 38.2 (2.7).

c. In 15 patients (75%) labor occurred and in 5 (25%) the resolution of the pregnancy was by cesarean section.

d. The amount of bleeding before and after Bakri balloon application
was 1570 ± 687mL and 130.7 ± 112mL, respectively (Z = -3.924, p = 0.000087). The total amount of bleeding during the event, the time between diagnosis and placement of the balloon, time of filling, amount of solution for filling and time of permanence of the balloon appear in Table 2.

e. The amount of bleeding between the cases of delivery and cesarean section before the application of the Bakri balloon was 1407 + 672 mL and 2060 + 513mL (Z = -2.08, p = 0.037). The amount of bleeding between the same groups after balloon placement was 128 + 123mL and 140 + 82.1mL (Z = -3.189, p = 0.001).

f. The average time between diagnosis and Bakri balloon placement was 30 minutes (5-120minutes). This was the time between which the bleeding was presented, the medical treatment was performed, the result was expected and the balloon application was decided. Once he proceeded to the application of the balloon into the uterine cavity e l time it was made and l filling averaged 5 minutes (3-10minutes). The average amount of volume administered was 400mL (250-500mL) Table 2.

g. Average levels of hemoglobin at admission to the unit, one post-adult hour and twenty - four hours after the treatment with blood components and before the balloon removal, were 11.8g/dL , 8.1g/dL and 9.6g/dL , respectively, with significant differences in the first two cases (on admission and post- graduation ) with Z = -3.92; p = 0.00008. There was also a difference between hemoglobin at admission and post - treatment with hemo components before the removal of the balloon, with Z = -3.03; p = 0.002 (Table 3).

h. In 19 patients (95%) the coagulation tests in the same periods of time were normal. Only in one case (5%, patient 4) the coagulation times were prolonged in the immediate post- acute (Table 4).

i. In 85% of the cases the use of a globular package with an average of 2.7units was required, with a range of 1 to 6 units. In case number 4, 8units of fresh plasma and 6 platelet concentrates were required.

j. In 19 patients (95%) a favorable response was obtained with the application of the Bakri balloon. In none of these cases was required the application of a second balloon during the time of use, nor after the withdrawal of the same. The mean and standard deviation of the time of use of the balloon was 29.5hours with an SD of 13.5hours.

Table 1 Clinical characteristics. Diagnosis and initial treatment

| No patients | Age | AUG | Weeks Gestation | Diagnosis                      | Pretreatment ball Bakri |
|-------------|-----|-----|-----------------|-------------------------------|------------------------|
|             |     |     |                 |                               |                        |
| 1           | 21  | G1, C1 | 36               | Previous placenta            | Ox, Mi, Ca             |
| 2           | 21  | G2, C2 | 40.2             | TP Extended                  | Ox, Mi, Ca             |
| 3           | 18  | G1, P1 | 38               | Uterine hypotonia            | Ox, Ca                 |
| 4           | 19  | G2, P2 | 37               | Uterine hypotonia            | Ox, Mi, Ca             |
| 5           | 21  | G2, P1, C1 | 40          | Uterine hypotonia            | Ox, Mi, Ca             |
| 6           | 20  | G2, P2 | 38.5             | Uterine hypotonia            | Ox, Mii, Ca            |
| 7           | 24  | G3, P3 | 41               | Uterine hypotonia            | Ox, Mi, Ca             |
| 8           | 17  | G2, P2 | 40               | Uterine hypotonia            | Ox, Mi, Ca             |
| 9           | 32  | G3, C2, A1 | 39.1         | Previous placenta            | Ox, Mi, Ca             |
| 10          | 2.3 | G3, P1, C1, A1 | 32.6       | RPM, Prolonged TP 16 hours   | Ox, Ca                 |
| 11          | 19  | G1, P1 | 38.6             | Prolonged TP 15 hrs. macrosomia | Ox, Mi, Ca             |
| 12          | 3.4 | G5, P4, A1 | 39.1         | Uterine hypotonia            | Ox, Mi, Ca             |
| 13          | 42  | G6, P5, A1 | 30           | Previous placenta            | Ox, Mi, C              |
| 14          | 35  | G4, P4 | 39.6             | Fortuitous birth             | Ox, Mi, Ca             |
| 15          | 2.3 | G2, P1, C1 | 39.6         | Uterine hypotonia            | Ox, Mi, Ca             |
| 16          | 32  | G1, P1 | 39.4             | severe pre-eclampsia         | Ox, Mi, Ca             |
| 17          | 18  | G1, P1 | 41.3             | Prolonged TP 19 hrs          | Ox, Mi, Ca             |
| 18          | 20  | G1, P1 | 39.4             | Uterine hypotonia            | Ox, Mi, Ca             |
| 19          | 22  | G2, P1, A1 | 39           | Uterine hypotonia            | Ox, Mi                 |
| 20          | 31  | G2, P2 | 36               | RPM, prolonged PT 17 hours   | My, Ca                 |

G, Gesta; P, Child birth; C, Cesarean; A, Abortion; TP, Labor; RPM, premature rupture of membranes; Ox, Oxytocin; Mi, Misoprostol; Ca, Carbetocin

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Table 2 Amount of bleeding, time between diagnosis and placement of the balloon, filling time, amount of solution used and dwell time

| No. | Patient | Total amount of bleeding (mL) | Time between diagnosis and placement (min) | Time of filling (min) | Amount of solution used (mL) | Time of permanence (hours) |
|-----|---------|------------------------------|-------------------------------------------|----------------------|-----------------------------|---------------------------|
| 1   |         | 2300                         | 20                                        | 5                    | 300                         | 48                        |
| 2   |         | 1800                         | 20                                        | 5                    | 400                         | 24                        |
| 3   |         | 1200                         | 15                                        | 6                    | 500                         | 48                        |
| 4   |         | 3500                         | 5                                         | 7                    | 500                         | 30                        |
| 5   |         | 1200                         | 5                                         | 5                    | 450                         | 26                        |
| 6   |         | 1200                         | 10                                        | 5                    | 500                         | 24                        |
| 7   |         | 900                          | 5                                         | 5                    | 350                         | 48                        |
| 8   |         | 1800                         | 20                                        | 5                    | 340                         | 24                        |
| 9   |         | 2300                         | 5                                         | 5                    | 500                         | 48                        |
| 10  |         | 1300                         | 5                                         | 4                    | 400                         | 24                        |
| 11  |         | 1000                         | 5                                         | 5                    | 500                         | 24                        |
| 12  |         | 800                          | 20                                        | 5                    | 300                         | 24                        |
| 13  |         | 2600                         | 5                                         | 5                    | 300                         | 48                        |
| 14  |         | 1000                         | 40                                        | 10                   | 350                         | 24                        |
| 15  |         | 2000                         | 5                                         | 5                    | 500                         | 24                        |
| 16  |         | 1500                         | 5                                         | 5                    | 400                         | 24                        |
| 17  |         | 1500                         | 5                                         | 3                    | 300                         | 24                        |
| 18  |         | 1500                         | 5                                         | 5                    | 500                         | 48                        |
| 19  |         | 1000                         | 5                                         | 5                    | 250                         | 24                        |
| 20  |         | 1000                         | 20                                        | 5                    | 350                         | 24                        |

* 1) By gravimetry and 2) with collection bag Brass-V. Calibrated obstetric drape (Maternova TM) type.

Table 3 Amount of bleeding between cases of delivery and cesarean section. Before and after the application of the ball.

|                          | Bleeding before application of the ball * | Bleeding after the application of the ball ** |
|--------------------------|------------------------------------------|---------------------------------------------|
| Postpartum mL            | 1407±672                                  | 2060±513                                    |
| Cesarean section mL      | 128±123                                   | 140±82.1                                    |

* p = 0.03 ** p = 0.001

Table 4 Mean levels of hemoglobin on admission, one hour post- graduated and later to treatment with blood components and removal of the balloon

|                          | Hemoglobin to the Entrance g/dL* | Hemoglobin I hour Postsangrado g/dL* | Post- treatment hemoglobin g/dL ** |
|--------------------------|----------------------------------|-------------------------------------|-----------------------------------|
|                          | 11.8                             | 8.1                                 | 9.6                               |

* p = 0.008 ** p = 0.002

In only one case (5%), a positive response with the ball was not obtained. This patient presented a quantified bleeding in 3000mL, secondary to a uterine hypotonia. After the placement of the balloon at 30minutes, the loss was 500mL, so the balloon was removed. The patient was transferred to intensive care, where the presence of coagulopathy due to consumption was corroborated. After volume restitution, fresh plasma and platelet concentrate with improvement in coagulation times, total abdominal hysterectomy was performed. There was no maternal death in the 20 cases studied.

There were no complications in the 19 patients who successfully used the Bakri balloon.

Discussion

The successes of Bakri’s balloon in obstetric hemorrhage range from 100% 18, 24 to 80%.25 The variation in their results may depend on the cause of the bleeding, which is multiple.18,25–27 In this report predominated uterine hypotonia first cause and prolonged labor as second causes that seem to have a better response with the use of intrauterine balls.28,29 In both cases apparently the pathophysiology
may be due to difficulty of the muscle fiber to occlude the uterine vessels in the immediate postpartum period. Bakri’s balloon, by producing a hydrostatic effect on the vessels and occluding them, favors its haemostasis, as well as activating hemostatic mechanisms and favoring the formation of a stable clot. Its use in other pathologies, such as placental accreta, reduces its effect.

One issue discussed with its use is the amount of solution needed to fill the balloon. In our setting, Ortega-Vadillo correlated the volume to be infused with gestational age and the weight of the newborn, with good results, 94.2% of success. In our report the criterion for the amount of the filling was the cessation of bleeding, in quantities that were 250 to 500cc, with an average of 400cc and we observed similar results. Therefore, we consider that any of these methods for filling the balloon provides similar results.

There are currently no reports in the literature that correlate the use of the Bakri balloon in cases of delivery and caesarean section, before and after its application in relation to hemoglobin levels, also before and after its use. In our report we found a statistically significant difference with the use of the balloon, in both cases. Therefore, we cannot compare our results with other studies. More studies are required comparing these variables.

In the present report, there was no complication with the use of the balloon. Some authors have reported accidents such as inadvertent uterine rupture, perforation 33 or spontaneous expulsion. The use of tamponade in the vaginal fundus to elevate the cervix towards the retro pubis, the practice of ultrasound to corroborate its adequate position and to keep the patient in absolute rest during the time of use, seem to be favorable measures to avoid this type of complications.

The success of the Bakri balloon in our report, 95%, is similar to multiple studies already reported. Its ease of application, being a conservative, simple procedure with a fast and easily replicable learning curve make it suitable for the control of obstetric hemorrhage. In our report, the average time for its application was 5 minutes, a time that is not observed with other procedures that are required comparing these variables.

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Conclusion

Bakri’s balloon showed a favorable response for its use in obstetric hemorrhage. An adequate control of it is obtained, observing a significant decrease in the amount of bleeding, both in postpartum and trans cesarean cases, recovering the hemoglobin values before and after its use. Its level of security is adequate, easy to apply and with simple training. It has a successful result rate, in this report of 95%, very similar to other reports in the literature.

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None.

Conflicts of interest

The author declares that they do not have any conflicts of interest.

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