

(RESEARCH ARTICLE)



Postpartum hemorrhage among women delivered at Tumbi Regional Referral Hospital, Coast Region, Tanzania Prevalence and Risk factors

Sylvester Wanyaseleli Mkama *

Department of Obstetrics and Gynecology St. Francis University College of Health and allied Science Ifakara, Tanzania.

International Journal of Biological and Pharmaceutical Sciences Archive, 2024, 08(02), 047–053

Publication history: Received on 15 September 2024; revised on 24 October 2024; accepted on 26 October 2024

Article DOI: <https://doi.org/10.53771/ijbpsa.2024.8.2.0068>

Abstract

Background: Postpartum hemorrhage (PPH) is generally defined as blood loss greater than or equal to 500mls within 24 hours after birth, while severe PPH is blood loss greater than or equal to 1000mls within 24 hours. The aim of this study is to access the prevalence, causes and risk factors associated with postpartum hemorrhage among women delivered at Tumbi Regional Referral Hospital, Coast Region, and Tanzania.

Keywords: Postpartum hemorrhage; Delivery; Women Tumbi Regional Referral Hospital Coast Region; Tanzania

1. Introduction

Postpartum Hemorrhage (PPH) is generally defined as blood loss greater than or equal to 500mls within 24 hours after birth, while severe PPH is blood loss greater than or equal to 1000mls within 24hours. PPH is the most common cause of maternal death worldwide. Most cases of morbidity and mortality due to PPH occur in the first 24 hours following delivery and these are regarded as primary PPH whereas any abnormal or excessive bleeding from the birth canal occurring between 24 hours and 12 weeks postnatally is regarded as secondary PPH. Worldwide about half a million women dies as results of complications of pregnancy and child birth [1]. In both developed and developing countries, postpartum hemorrhage is a leading cause of severe maternal morbidity and mortality. Approximately 14 million women suffer primary postpartum hemorrhage annually and at least 128, 000 of these women bleed to death [2]. Postpartum hemorrhage is the most preventable and treatable problem through active management of the third stage of labor AMTSL [3]. However, the use of oxytocin is not feasible in many low-income countries, where most births take place at home with untrained birth attendants [4]. The magnitude of PPH in Sub-Saharan Africa is high at 10.5% [5]. Ethiopia is one of the countries with the highest maternal mortality rate (MMR) and almost all of these deaths were due to direct obstetric complications [6]. PPH is a leading direct causes of maternal morbidity and mortality in all region of the country [6]. World Health Organization reported in 2012 that PPH affects 2% of childbirth [7]. Because two-thirds of pregnant women that develop PPH have no known risk factors, preventive measures must be offered to all pregnant women [8]. The main PPH prevention measure recommended in low-and middle-income countries (LMICs) is active management of the third stage of labor (AMTSL), which WHO recommends for all deliveries in LMIC settings [9]. The highest prevalence of PPH revealed in Africa (25.7%) followed by Latin America and Asia (18%), North America and Europe (13%) and Oceania (7.2%) [10]. Study done in Netherlands shows that the prevalence was 4.1% (2000) [11] which rose to 6.4% in 2013 [12]. In Senegal PPH rose between 10% in 2007 to 23% in 2008 [10] while in Quarite PPH was 5.4% in 2015 [13]. In Sub-Saharan Africa, where 1 in 16 women die of pregnancy and childbirth-related conditions, PPH is estimated to account for between 25%-30% of these deaths [14], thus making severe bleeding the single most important cause of maternal mortality worldwide [15]. Primary postpartum hemorrhage is traditionally considered as a disorder of one or more of four processes: uterine atony, retained clots or placental debris, genital lesions or trauma, and disorders of blood coagulation. Uterine atony which can occur after normal vaginal delivery or abdominal delivery is said to account for up to 75-90% of cases of Postpartum Hemorrhage [16]. Secondary (late) PPH occurs between 24

*Corresponding author: Sylvester Wanyaseleli Mkama

hours after delivery of the infant and 6 weeks postpartum. Most late PPH is due to retained products of conception infection, or both. In the rural areas of Tanzania, the use of a 'Kanga' has been adopted as a valid measurement too [17]. Convenient because it is produced and sold locally, the pre-cut kanga is a standard sized rectangle (100 cm X 155 cm) of local cotton fabric. When three to four soaked kangas are observed at a delivery, the trained, the trained traditional birth attendant (TBA) is entrusted to transfer patients to a health center. Even when a good measurement methodology is in place, there is still difficulty in defining PPH simply as blood loss greater than 500ml because it fails to take into account predisposing health factors that are reflected in such a definition. Since the quantity of blood loss is less often important than the actual effect that it has on the laboring women, it has been suggested that the definition take into account any blood loss that causes a major physiological change, such as low blood pressure, which threatens the women's life. Tanzania is a substantial contributor to maternal mortality, with a maternal mortality ratio of 556 per 100,000 live births during the 10-year period before the 2015 / 2016 Demographic and Health Survey. Maternal mortality accounts for 21% of deaths of women reproductive age [18]. In Uganda PPH causes 25% of all maternal deaths [5]. In Kenya maternal mortality has declined by nearly 50% over the past thirty years, it remains high in low-and middle-income countries [19]. Therefore the aim of this study was to assess the prevalence, causes and risk factors associated with postpartum hemorrhage among women delivered at Tumbi Regional Referral Hospital, Coast Region, Tanzania. The findings of this study are important for national policy makers and health institutions because it sought to open up new perspectives in early identification of women at risk for PPH, early diagnosis and might provide new strategies to prevent and control PPH.

2. Material and methods

2.1. Study area and period

This study was conducted at Tumbi Regional Referral Hospital, Kibaha District, Coast Region, Tanzania from 01 January to December 31, 2023.



Figure 1 Coast Region and its 6 districts

The study was conducted to assess the prevalence, causes and risk factors associated with postpartum hemorrhage among women delivered at Tumbi Kibaha district. Kibaha is one of the six administrative districts of Coast Region in Tanzania (Figure 1). The district covers an area of 1,502 km² (580 sq mi). Kibaha District is bordered to the northeast by Kibaha Urban District and the north by Chalinze District. The district is bordered to the southeast by the Kisarawe District, On the western side the district is bordered by Morogoro District of Morogoro Region. According to the 2012 census, the district has a total population of 70,209.

The study was conducted to access the prevalence, cause and risk factors postpartum hemorrhage among women attending antenatal clinic at Tumbi, Kibaha district.

2.2. Study design

The study design was across –sectional study which prevalence, causes and risk factors of postpartum hemorrhage among pregnant women attending antenatal clinic at Tumbi Regional Referral Hospital, Kibaha District, and Coast Region was studied at a time. Structured pretested questionnaire with key information was used to collect the desired data.

2.3. Study population

Pregnant women attending clinic at Tumbi Regional Referral Hospital were registered in the study.

2.4. Sampling size

The sample size in this study was 212 participants. The sample size calculation by Kirkwood formula

$N = \text{Sample size}$

$Z = \text{Confidence interval level } 95\% \text{ in this study which is } 1.96$

$P = \text{Proportional of study prevalence (estimated prevalence) } 1.4\% \text{ } 2021$

$D = \text{Absolute error or precision } 0.05 \text{ has to be decided by researcher.}$

$$N = Z^2 P(1-P) / D^2$$

$$N = 1.96^2 \times 1.4\% (1 - 1.4\%) / 0.05^2$$

$$N = 0.053029 / 0.0025$$

$$N = 212$$

2.5. Sampling Technique

Simple randomly technique was employed when participants attending at Tumbi Regional Referral Hospital antenatal clinic were allocated numbers (even and odd numbers). Participants who had even numbers were involved in the study.

2.6. Data collection

The data collected by structure guided questionnaires. The questionnaire prepared in English and translated into Swahili to maintain the consistency and content of the questionnaire, confidentiality of information, participant's rights and voluntarily informed consent were secured. The participants were asked the questions and their answers filled in the questionnaire by the researcher.

2.7. Data analysis

Questionnaire filled with irrelevant information were removed. The data from questionnaire with relevant information were analyzed with Statistical Package for Social Sciences (SPSS version 20)

2.8. Inclusion criteria

Women who had vaginal delivery and whose blood losses was 500mls or more willing to participate were included in the study.

Those who had primary postpartum hemorrhage within the study willing to participate were included in the study.

2.9. Exclusion criteria

Women who had caesarean delivery and unwilling to participate in the study.

Those who delivered outside the hospital without any referral letter.

Those who were brought in dead following bleeding after delivery.

2.10. Ethical clearance

A letter from St. Francis University College of Health and Allied Sciences ethical committee was obtained. The letter submitted to Regional Medical Officer, Coast Regional who forwarded the letter to the Medical Officer In charge who give permission to use participant at Tumbi Referral Hospital.

3. Results

3.1. Socio-demographic characteristic of participants pregnant women attending antenatal clinic at Tumbi Referral hospital

A total of 212 pregnant women participated in this study. Out of these 20 (9.4%) aged < 19 years , 100 (47.2%) aged 20-29 years, 60 (28.3%) aged 30-39 years, and 32 (15.1%) aged > 40 years. Education status of pregnant women who participated in the study, 15 (7.1%) were not educated, 30 (14.2%) were primary education, 100 (47.1%) were secondary education, 67 (31.6) were higher education. Primipara 35(16.5%), para 2- 4 105 (49.5), para 5 and above 72 (34.0%) , as shown in Table 1

Table 1 Social demographic characteristics of participants

Variable	Frequency	Percentage
Age(in years)		
< 19 years	20	9.4
20-29 years	100	47.2
30-39 years	60	28.3
>40years	32	15.1
Education Status		
Not educated	15	7.1
Primary school	30	14.2
Secondary school	100	47.1
Higher education	67	31.6
Parity		
Primepara	35	16.5
Para 2-4	105	49.5
Para 5 and above	72	34.0

3.2. Causes of primary postpartum hemorrhage of participant women attending antenatal clinic at Tumbi Referral Hospital

Among 212 pregnant women; uterine atony 40 (18.9%), laceration 15 (7.1%), retained placenta 140 (66.0%), coagulopathy 17(8.0%) as shown in table 2

Table 2 Causes of primary post partum hemorrhage among participants

Variable	Frequency	Percentage
Uterine atony	40	18.9
Laceration	15	7.1
Retained placenta	140	66.0
Coagulopathy	17	8.0

3.3. Causes of secondary postpartum hemorrhage of participant women attending antenatal clinic at Tumbi Referral Hospital

A total of 200 (94.3%) participants had retained placenta, 7 (3.3%) had puerperal sepsis, and 5 (2.4%) had episiotomy as shown Table 3.

Table 3 Causes of secondary post partum hemorrhage among participants

Variables	Frequency	Percentage
Retained placenta	200	94.3
Puerperal sepsis	7	3.3
Broken- down/episiotomy	5	2.4

4. Discussion

This study was aimed to identify the prevalence, causes and risk factors of pregnant women among women attending antenatal clinic in Tumbi Regional Referral Hospital, Coast Region, and Tanzania. It was observed that 47.2% were between 20-29 years. In comparison to women in the age range between 20 and 34 years old who had a vaginal delivery, women aged 35 and older were more likely to have experienced PPH, according to research by Kebede et al. [20] and Shahbazi Sighaldehy et al [21]. However, Bazirete et al. and Onong et al. were unable to discover a relationship between age and bleeding [22] and their findings are consistent with those of the current investigation. In this study 49.5% were para 2-4. A study done by Ononge et al. suggested that grand multi-parity may be tangentially related to PPH [23], and some research showed that the majority of PPH cases were in mothers who were nullparous [24]. Similar to the Ononge study, the current investigation found no evidence of a substantial impact of parity on the occurrence of PPH. In this study retained placenta 66.0% and 94.3% primary postpartum hemorrhage and secondary postpartum haemorrhage respectively, other studies was reported that abnormal placentation was responsible for majority 55.83% of severe PPH, which was much higher than the previously reported prevalence of 10% [25].

5. Conclusion

In this study if postpartum haemorrhage is reduced, maternal death will be greatly reduced since PPH is the main cause of maternal death. It is important to remember that we have to prepare for all mothers giving birth, as some get severe PPH without any known risk factors.

Recommendation

Health professional attending labor and delivery should give more attention for high risk mothers older age, grand multiparity and history of PPH mothers during delivery. Continue to encourage ANC visit and prevent prolonged labor should be advice to reduce the occurrence of postpartum haemorrhage.

Compliance with ethical standards

Acknowledgments

I wish to thank the management of St. Francis University College of Health and Allied Sciences for the support to this study. Also, I thank the Regional Medical Officer, Coast Region, Medical Officer In charge Tumbi Regional Referral Hospital, antenatal clinic staff for their support in this study and all pregnant women attending antenatal clinic who participate in this study.

Disclosure of conflict of interest

No conflict of interest in this study.

Statement of ethical approval

In this study no animal was used but human was used. The only study tool used to collect data was questionnaire. However, ethical clearance was obtained from the respective authorities to conduct the study. The research committee of St. Francis University College of Health and Allied Sciences, Regional Medical Officer, Coast Region, Medical Officer in charge Tumbi Regional Referral Hospital gave permission the study to be conducted.

Statement of informed consent

Written informed consent was obtained from all antenatal pregnant women who consented to the study, records were coded and participant / Researcher names were not used. All the information collected remained confidential and was used for purposes of the study only. Participation was voluntary and no incentives were given.

References

- [1] World Health Organization, author. The WorldReport 2005. Attending to 136 million births, every year. 2005. Make every mother and child count. Geneva: the World Health Organization; 2005. pp. 62-63.
- [2] WHO. Attending to 136 Million births everyyear, make every mother and child count. World Health Organization. Geneva: 2005:62-63.
- [3] Evensen A, Anderson JM, Fontaine P. Postpartum hemorrhage: prevention and treatment .Am Fam Physician . 2017; 95 (7):442-9.
- [4] MobeenN, Durocher J, Zuberi N. Jahan N, Blum J, Wasim S, Walraven G, Hatcher J. Administration of misoprostol by trained traditional birth attendants to prevent postpartum hemorrhage in home births in Pakistan a randomizedplacebo – controlled trial. BJOG Int J ObstetGynaecol. 2011;118 (3)353-61.
- [5] Carroli G, Cuesta C, Ablalos E. Gulmezoglu AM. Epidemiology of postpartum hemorrhage : a systematic review Best Pract Res ClinObsteGynaecol 2008; 22. 999-1012.
- [6] CSA. 17. Ethiopia demographic and health survey. Addis Ababa: Central Statistical Agency; 2016.
- [7] WHO. WHO recommendations for the prevention and treatment of postpartum hemorrhage. In: WHO, editor . Maternal, newborn, child and adolescent health, Geneva: WHO; 2012.
- [8] Mpemba F, Kampo S, Zhang X. Towards 2015: post-partum hemorrhage in Sub- Saharan Africa still on the rise. J ClinNurs. 2014; 23 (5 / 6): 774-83. 10p
- [9] Tuncalpõ, Souza JP, Gülmezoglu M. New WHO recommendations on prevention and treatment of postpartum hemorrhage. Int J Gynecol Obstet. 2013; 123 (3):254-6.
- [10] Calvert C, Thomas SL, Ronsmans C, Wagner KS, Adler AJ, Filippi V. Identifying regional variation in the prevalence of postpartum hemorrhage: A systematic review and meta-analysis. PLoS One. 2012; 7 (7).
- [11] Prual A, Bouvier-Colle MH, De Bernis L, Brèart G. Severe maternal morbidity from direct obstetric causes in West Africa: Incidence and case fatality rates. Bull WorldHealth Organ. 2000; 78(5):593-602.
- [12] Ngowa JDK, Ngassam AN, Dohbit JS, Nzedjom C, Kasia JM. Pregnancy outcome at advanced maternal age in a group of African women in two teaching hospitals in Yaounde, Cameroon. Pan Afr Med J. 2013; 14.

- [13] Journal JS-DESMS, 2010 undefined. Postpartum Hemorrhage Among Women Delivered at Mbeya Referral Hospital in 2008. *Ajol.info*. Published online 2010. Accessed September 21, 2021. <https://www.ajol.info/index.php/dmsj/article/view/61337>.
- [14] Claudio GS, Althabe F, Belizan JM, Buckens P. Risk factors for postpartum hemorrhages in vaginal deliveries in a Latin American population. *ObstetGynaecol* 2009; 113 (6): 1313-1319.
- [15] Lain SJ, Roberts CL, Hadfield RM, Bell JC, Morris JM. How accurate is the reporting of obstetric hemorrhage in hospital discharge data? A validation study. *Aust NZ J ObstetGynaecol*. 2008; 48(5): 481-4.
- [16] Lalonde AB, Davis B-A, Hershderfer K. Postpartum Hemorrhage Today: living in the shadow of the Taj Mahal. In: B-lynch C, Keith IG, Lalonde AB, Karoshi M (eds). *Textbook of Postpartum Hemorrhage*. Dumfriesshire, UK. Sapiens Publishing. 2006: 2-9.
- [17] Prata N, Mbaruku G, Campbell M. Using the kanga to measure postpartum blood loss. *Int J Gynaecol Obstet* 2005; 89 : 49-50.
- [18] Ministry of Health, community Development ,Gender, Elderly and Children (MoHCDGEC) [Tanzania Mainland], Ministry of Health Zanzibar, National Bureau of Statistics (NBS), Office of Chief Government Statistician , ICF. *Tanzania Demographic and Health Survey and malaria Indicator Survey (TDHS-MIS) 2015-16*. Dar es Salaam, Tanzania, and Rockville, MD, USA: MoHCDGEC, MoH, NBS, OCGS, and ICF; 2016. p. 172-3.
- [19] WHO. *Trends in Maternal Mortality: 1990-2015*. Estimates by WHO, UNICEF, UNFPA, World Bank Group and United Nations. World Health Organization; 2015. Available at : <https://www.who.int/reproductive-health/publications/monitoring/maternal-mortality-2015/en/>.
- [20] Kebede BA, Abdo RA, Anshebo AA, Gebremariam BM. Prevalence and predictors of primary postpartum hemorrhage: An implication for designing effective intervention at selected hospitals, Southern Ethiopia. *PloS One*. 2019; 14 (10):e0224579. [PMID] [PMCID] [DOI :10.1371/journal.pone.0224579]
- [21] ShahbaziSighaldehy S, Nazari A, Maasoumi R, Kazemnejad A, Mazari Z. Prevalence, related factors and maternal outcomes of primary postpartum haemorrhage in governmental hospitals haemorrhage in governmental hospital in Kabul- Afghanistan. *BMC Pregnancy Childbirth*. 2020; 20(1): 428. [PMID] [PMCID] [DOI: 10.1186/s12884-020-03123-3]
- [22] Bazirete O, Nzayirambaho M, Umubyeyi A, Karangwa I, Evans M. Risk factors for postpartum haemorrhage in the Northern Province of Rwanda: A case control study. *PloS One*. 2022; 17 (2): e0263731. [DOI : 10.1371/journal.pone.0263731] [PMID] [PMCID].
- [23] Ononge S, Mirembe F, Wandabwa J, Campbell OMR. Incidence and risk factors for postpartum haemorrhage in Uganda. *Repro Health*. 2016; 13 (1):38. [DOI :10.1186/s12978-016-0154-8] [PMID] [PMCID]
- [24] Wang C, Zhang C. Meta-analysis to assess the role of maternal characteristics and risk factors on postpartum haemorrhage. *Adv Clin Exp Med*. 2023:1-9.
- [25] Kramer MS, Berg C, Abenhaim H, Dahhou M, Rouleau J, Mehrabadi A, et al. Incidence, risk factors, and temporal trends in severe postpartum haemorrhage. *Am J Obstet Gynecol* 2013; 209 (5) : 441-9.