Obstetric outcome of female genital mutilation in the Gambia - an observational study

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Abstract:

Background: A 2010 survey in The Gambia among women of reproductive age put the prevalence rate of FGM/C at 76.3%. FGM/C was banned in 2015, but there is no real effort at enforcement of the ban. This study aimed to provide national data on obstetric outcomes to support advocacy and health education.

A multicentre observational study to assess the obstetric and neonatal outcomes of parturient women with and without FGM/C was carried out across 4 healthcare facilities in The Gambia. The primary outcome was postpartum haemorrhage (>500ml) and secondary outcomes were caesarean section, perineal tears (including episiotomy), neonatal resuscitation and perinatal death.

Of the 1,569 participants recruited into the study, 23% had no FGM/C while 77% had FGM/C of varying severity. The risk of postpartum haemorrhage was doubled for women with type I FGM/C, tripled in type II FGM/C and increased by 5-fold for those with type III and IV FGM/C. Caesarean section and perineal tears were also increased. FGM/C was associated with increased risk for neonatal resuscitation and perinatal death.

FGM/C is associated with poor obstetric and neonatal outcomes in the Gambia with degree of risk correlating with the severity of FGM/C.

Keywords: Female genital mutilation, obstetric outcome, Gambia.

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Background

The World Health Organization (WHO) definition of Female Genital Mutilation/Cutting (FGM/C) comprises all procedures involving partial or total removal of the external female genitalia or other injury to the female genital organs for non-medical reasons¹. An estimated 200 million women and girls worldwide have been subjected to this practice, the majority living in African countries². FGM/C spans cultural and ethnic groups and occurs among Muslim, Christian and secular communities. The WHO classifies FGM/C by type and severity of mutila-

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Patrick Idoko, School of Medical and Allied Health Sciences, University of The Gambia, P O Box 3031 Banjul, The Gambia. Email: patidoko@gmail.com tion, Type I being the least mutilating and Type III, also known as infibulation, being the most severe.

Reasons given for the practice of FGM/C are based on cultural and religious beliefs and have no basis in science. These include: prevention of promiscuity in the female, enhancement of male sexual performance and pleasure, maintenance of cleanliness of the genital area, aesthetic reasons, enhancement of fertility and improving a woman's marriage prospects³. However, the practice of FGM/C is fraught with adverse health consequences and is frequently performed by traditional practitioners who have had no formal medical training, thus increasing the associated risks ⁴. Immediate complications include bleeding leading to shock, transmission of infection and injuries to adjacent organs like the urethra and the rectum. Long term sequelae of FGM/C include dyspareunia, anorgasmia, Para clitoral cyst and chronic pain³. Obstetric risks include prolonged labour, increased cae-

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sarean section rates, increased rates of episiotomy, perineal tears and postpartum haemorrhage³⁻⁵. Women with FGM/C also suffer increased rates of perinatal complications including need for neonatal resuscitation, still birth and early neonatal death⁵.

In part because of these negative health consequences of FGM/C, many governments around the world have outlawed the practice. However, FGM/C continues to be practiced at high rates across many African countries.

The WHO classifies FGM/C into 4 types^{1,5}. Type I is a partial or complete removal of the clitoris and/or the prepuce (clitoridectomy). In type II, the clitoris and labia minora are partially or completely removed with or without excision of the labia majora. Type III FGM/C involves a narrowing of the vaginal orifice by cutting the labia minora and /or the labia majora and creating a covering seal by close apposition of the cut surfaces. Type IV refers to all other harmful procedures to the female genitalia for non-medical purposes like pricking, cauterization and piercing^{1,5}. The severity of the FGM/C and its complications worsens from type I to type III.

In 2010 a survey in the Gambia found the national prevalence rate of FGM/C among women aged 15-49 to be 76.3%⁶. The Foundation for Research on Women's Health, Productivity and the Environment (BAFROW) reports that seven of The Gambia's nine ethnic groups practice FGM/C⁴. Nearly all Mandinkas, Jolas and Hausas (together 52% of the population) practice Type II on girls between 10 years and 15 years of age. The Sarahulis (9% of the population) practice Type I on girls one week after birth. The Bambaras (1% of the population) practice Type III, which takes place when girls are between 10 years and 15 years of age. The Fulas (18% of the population) engage in a practice analogous to Type III that is described as "vaginal sealing" on girls anywhere between one week and 18 years of age.

Evidence on healthcare outcomes of FGM/C in the Gambia is currently limited. One study found an association between FGM/C and higher rates of bacterial vaginosis and herpes simplex virus 2⁷, but a second study did not uphold this finding⁸. A further study of Gambian women with FGM/C presenting with gynaecological problems attributed many of these to FGM/C⁹. This study did not have a control group and relied on retrospective data in a selected group of women. The same group performed another study of obstetric outcomes of FGM/C, which included women without FGM/C as a control group¹⁰. This study relied on retrospective

reporting of complications by the women but showed significantly worse outcomes for women with FGM/C including prolonged labour, episiotomy, perineal tears, caesarean section, need for neonatal resuscitation and stillbirth. There have been no prospective studies of obstetric outcome of FGM/C in the Gambia.

With a population of just 1.8 million people The Gambia is a small country that falls low on the human development index (165 out of 187 in 2012¹¹), with high rates of absolute poverty and poor access to healthcare ¹². Sadly, it faces multiple challenges in the field of health and typifies many of the issues surrounding the current crisis in human resources for health in Africa. As such, the additional health needs and risks posed by FGM/C are happening against a background of malnutrition, poor maternal and child health indicators and low rates of gender equality ¹¹. In 2015, Gambia's parliament passed a bill outlawing FGM/C for the first time. But with high rates among the current population, it may be many years before the health impact of this ban is felt. National data on healthcare outcomes may be a key component in the advocacy and health education needed to eliminate this practice. The aims of this study therefore are to determine the obstetric and neonatal outcomes of parturient women with FGM/C in the Gambia.

Methodology

This was a prospective stratified observational study. Consenting parturient women in the first stage of labour were included in the study. Recruitment was carried out at 4 health facilities across the country: Brikama District Hospital, Jammeh Foundation for Peace Hospital (now Bundung Maternal and child hospital), Bansang Hospital and Edward Francis Small Teaching Hospital.

Four midwives and a medical doctor were employed from each study site and trained in obtaining consent, identifying types of FGM/C, assessing blood loss, Apgar scoring, perineal tear assessment as well as using the questionnaire to gather other relevant data. The midwives were responsible for data collection with supervision from medical doctors and the investigators.

Women who presented in the first stage of labour (before full cervical dilatation) at the 4 participating health facilities during the study period were approached and informed of the study by the study medical doctor or midwife. Trained midwives or medical doctors then examined consenting women who met the eligibility criteria for the presence or absence of FGM/C. Where present, FGM/C was categorized on examination by the WHO classification of the type of FGM/C⁵. Structured questionnaire-based interviews were conducted to obtain demographic information as well as relevant medical and obstetric history. The women were followed up throughout the course of labour and outcome of labour was recorded in the questionnaire.

The eligibility criteria were pregnant women with a singleton foetus presenting in the first stage of labour giving informed consent to be included in the study. Any woman with multiple pregnancy, presenting in the second stage of labour or who was booked for elective caesarean section for any reason was excluded from the study.

The primary outcome was postpartum haemorrhage of >500ml, measured according to the protocol for measuring blood loss in the WHO multicentre randomized trial of misoprostol in the management of the third stage of labour¹³. Secondary obstetric outcomes were: unplanned caesarean section, perineal tears or need for episiotomy. Secondary neonatal outcomes included perinatal death, need for neonatal resuscitation and rates of low birth weight.

The sample size calculation was done using postpartum haemorrhage (binary variable) as the primary outcome. Kaplan et al. reported a proportion of 66.2% of FGM/C I, 26.3% of FGM/C II and 7.5% of FGM/C III⁹. The proportion of women with no FGM/C with postpartum haemorrhage was assumed to be 6% ⁵. The risk ratio for FGM/C I vs. no FGM/C and FGM/C II vs. no FGM/C was set at 2 while that of FGM/C III vs. no FGM/C was set at 2.5. The type I error was set at 5% and the power at 80%. A sample size ratio of 1 for comparing FGM/C I to no FGM/C and FGM/C II to no FGM/C but a ratio of 1/3 for comparing FGM/C III or IV to no FGM/C because of the low prevalence of FG/C III or IV was considered. A minimum of 1178 parturient women (a minimum of 356 in no FGM/C, FGM/C I and FGM/C II and of 110 in FGM/C III or IV) was the calculated sample size.

Poisson regression was performed to assess the association between FGM/C and binary outcomes. Linear re-

gression model was performed to assess the association between FGM/C and continuous outcomes. Confounders adjusted for were selected based on known associations with the outcome under study or factors that were hypothesized to play a role in the outcome under study in our setting. Data was analysed for test of association using STATA.

The care provided to women and babies was in accordance with normal standards and protocols at the participating health facilities. The Gambian Government Ethics Committee gave approval for the study.

Results

The total number of deliveries during the 5-month study period (1st May 2016 to 30th September 2016) from all of the participating health facilities was 3,867 out of which 2,197 (56.8%) met the inclusion criteria. However only 1,821 (82.9%) were informed of the study out of which 1,569 (83.7%) consented and were recruited into the study. Of the 1,569 participants that were recruited into the study, 23% had nFGM/C while 77% had FGM/C. Table 1 shows the baseline characteristics of study participants stratified by the presence and type of FGM/C.

Table 2 shows the obstetric and neonatal outcomes of FGM/C and table 3 shows the association between duration of labour and the type of FGM/C.

These data show that all forms of FGM/C increase the risk of poor obstetric outcomes: postpartum haemorrhage (blood loss of over 500ml), caesarean section and episiotomy or perineal tear. Neonatal outcomes are similarly shown to be poor for women with FGM/C; there was increased risk of need for neonatal resuscitation with all forms of FGM/C and perinatal death increased in type II FGM/C. In types I and III FGM/C an increased number of perinatal deaths were observed compared with no FGM/C but these numbers were too small to demonstrate statistical significance. There was no statistically significant difference in risk of having a baby with birth weight under 2500g in women with FGM/C.

Variable	No FGM	WHO type 1	WHO type 2	WHO type 3 or 4	p-value
	(n=361)	(n=372)	(n=704)	(n=132) °	
Age in years, n (%)					
Median (1si-3 rd quartiles)	27 (22-31)	25 (21-30)	25 (22-30)	24 (20-29)	<0.001#
<20	32 (8.9)	61 (16.4)	89 (12.6)	25 (18.9)	
20-24	94 (26.0)	107 (28.8)	225 (32.0)	50 (37.9)	
25-29	104 (28.8)	84 (22.6)	185 (26.3)	27 (20.5)	
30-34 35 or greater	81 (22.4) 50 (13.9)	75 (20.2) 45 (12.1)	118 (16.8) 87 (12.4)	13 (9.9) 17 (12.9)	
Tribe, n (%)					
Mandinka	32 (8.9)	150 (40.3)	329 (46.7)	60 (45.5)	< 0.001*
Fula	45 (12.5)	144 (38.7)	202 (28.7)	32 (24.2)	
Wollof	180 (49.9)	15 (4.0)	23 (3.3)	4 (3.0)	
Jola Other ª	25 (6.9) 79 (21.9)	29 (7.8) 84 (9.1)	75 (10.7) 75 (10.7)	13 (9.9) 23 (17.4)	
Education level, n (%)	4.40 (44.0)				
None	148 (41.0)	171 (46.0)	284 (4.03)	50 (37.9)	0.08*
Primary/non-formal	93 (25.8)	102 (27.4)	210 (29.8)	36 (27.3)	
Secondary Tertiary	94 (26.0) 26 (7.2)	82 (22.0) 17 (4.6)	147 (20.9) 63 (9.0)	38 (28.8) 8 (6.1)	
Residential area, n (%)					
Urban	248 (68.7)	201 (54.0)	431 (61.2)	87 (65.9)	<0.001*
Rural	113 (31.3)	171 (46.0)	273 (38.8)	45 (34.1)	
Height in cm, median (1st-3rd quartiles) ^b	162 (158-167)	160 (155-165)	160 (156-165)	160 (156-165)	0.003#
Parity					
Median (1 st -3 rd quartiles)	2 (0-3)	2 (0-3)	1 (0-3)	0 (0-2)	<0.001#
0	102 (28.3)	121 (32.5)	212 (30.1)	67 (50.8)	
1	67 (18.6)	64 (17.2)	157 (22.3)	28 (21.2)	
2	62 (17.2)	61 (16.4)	121 (17.2)	11 (8.3)	
3	49 (13.6)	45 (12.1)	79 (11.2)	9 (6.8)	
4	40 (11.1)	35 (9.4)	49 (7.0)	5 (3.8)	

Table 1: Baseline characteristics of study participants stratified by type of FGM

5 or greater	41 (11.4)	46 (12.4)	86 (12.2)	12 (9.1)	
Chronic medical conditions, n (%)					
Yes No	18 (5.0) 343 (95.0)	29 (7.8) 343 (92.2)	34 (4.8) 670 (95.2)	6 (4.6) 126 (95.4)	0.19*
Previous caesarean section, n (%)	5 (4 1)				
Yes No	5 (1.4) 356 (98.6)	21 (5.7) 351 (94.3)	26 (3.7) 678 (96.3)	4 (3.0) 128 (97.0)	0.02
Number of antenatal care visits, n (%)					0.40#
Median (1 st -3 nd quartiles)	3 (3-4)	3 (3-4)	3 (3-4)	3 (3-4)	0.42#
0	0 (0.0)	0 (0.0)	2 (0.3)	1 (0.8)	
1	14 (3.9)	18 (4.8)	32 (4.6)	10 (7.6)	
2	50 (13.9)	56 (15.1)	104 (14.8)	15 (11.4)	
3	100 (27.7)	108 (29.0)	187 (26.6)	46 (34.9)	
4 or greater	153 (42.4)	156 (41.9)	308 (43.8)	49 (37.1)	

^a include Jahanka, Konyagi, Manjago, Serere, Serahule, Woyinko and other minority tribes.

^bData were missing for 130 women who were excluded from the descriptive analysis.

^cOnly six FGM type IV were found and recruited into the study.

Kruskal-Willi's test.

* Pearson chi-square test.

Obstetric outcomes Post-partum blood loss >500ml -72 No FGM 34/361 9.4 1.0 <0.001 1.0 <0.0011 FGM I 67/372 18.0 1.9 (1.3. 2.1 (1.3- 2.1 (1.3- FGM II 67/372 18.0 1.9 (1.3. 2.1 (1.3- 2.3 (1.7- FGM III 58/126 46.0 3.1 (2.3- 2.9 (2.1- 4.2 Caesarean section 1.0 0.004 1.0 0.03601 No FGM 4.4 1.5 2.6 (1.5- 2.4 (1.1- FGM II 16/361 1.0 0.004 1.0 0.03601 No FGM 4.4 1.5 2.6 (1.5- 2.4 (1.1- 4.4 4.8) 1.10 0.00014 1.0 0.03601 FGM II 16/126 12.7 2.9 (1.5- 2.7 (1.2- 1.10 5.6) 6.1) 1.0 0.0014 Perinatal tear 76/361 21.1 1.0 <0.001 1.0 <0.0014 No FGM 140/372 37.6 1.8	Outcome and FGM status	Cases/population	Prevalence (%)	Crude RR (95% CI) *	р	Adjusted RR (95% CI) *	р
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$\begin{array}{cccccccccccccccccccccccccccccccccccc$	FGM III	58/126	46.0			(
$\begin{array}{cccccccccccccccccccccccccccccccccccc$	Caesarean section						
$\begin{array}{c ccccccccccccccccccccccccccccccccccc$	No FGM	16/361	4.4	1.0	0.004	1.0	0.036#
$\begin{array}{cccccccccccccccccccccccccccccccccccc$	FGM I	36/372	9.7			•	
$\begin{array}{c ccccccccccccccccccccccccccccccccccc$	FGM II	81/704	11.5			2.4 (1.1-	
or perineal tear $76/361$ 21.1 1.0 <0.001 1.0 <0.0014 No FGM $140/372$ 37.6 1.8 (1.4 1.7 (1.3-2.3) 2.1) FGM I 283/704 40.2 1.9 (1.5- 1.8 (1.4-2.4) 2.2) 101/126 80.2 3.8 (3.1- 2.4 (2.1-3.4) 2.4) FGM III 4.7) 3.4) Fetal outcomes Perinatal death $7/361$ 1.9 1.0 0.13 1.0 0.11* No FGM $13/372$ 3.5 1.8 (0.7- 1.9 (0.7-4.5) 4.6) 33/704 4.7 2.4 (1.1- 2.5 (1.1-5.4) 4.6) 33/704 4.7 2.4 (1.1- 2.5 (1.1-5.4) 5.7) 3/126 2.4 1.2 (0.3-1.3 (0.3-5.7) 5.1) Need for resuscitation $31/361$ 8.6 1.0 <0.001 1.0 <0.001* No FGM $13/372$ 1.3.4 1.6 (1.0-1.9 (1.2-2.4) 3.2) FGM I $21/704$ 17.2 2.0 (1.4- 2.5 (1.6-1.4)	FGM III	16/126	12.7			•	
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No FGM 50/372 13.4 1.6 (1.0- 1.9 (1.2- 2.4) 3.2) 121/704 17.2 2.0 (1.4- 2.5 (1.6-	Need for resuscitation	31/361	8.6	1.0	< 0.001	1.0	<0.001**
FGM I 2.4) 3.2) 121/704 17.2 2.0 (1.4- 2.5 (1.6-	No FGM	·					
	FGM I			2.4)		3.2)	
	FGM II	121/ /04	1/.2			•	

Table 2: Crude and adjusted relative risk (RR) of adverseoutcomes for any type of FGM vs No FGM.

FGM III	37/126	29.4	3.4 (2.2- 5.3)	3.9 (2.4- 6.5)
Birth weight <2500g				
	30/361	8.3	1.0 0.99	1.0 0.98***
No FGM	29/372	7.8	0.9 (0.6-	0.9 (0.6-
FGM I	,		1.5)	1.5)
	57/704	8.1	1.0 (0.6-	0.9 (0.6-
FGM II			1.5)	1.5)
	11/126	8.7	1.1 (0.5-	0.9 (0.5-
FGM III			2.0)	1.7)

*95% confidence interval of the relative risk.

* Adjusted for parity and birth weight

#Adjusted for parity, education, birth weight and previous caesarean section

Adjusted for parity, maternal age, birth weight and instrumental delivery

*Adjusted for maternal age, residence and history of chronic illness.

**Adjusted for maternal age, residence, parity and history of chronic illness

*** Adjusted for parity, residence and history of chronic illness

Table 3: Duration of labour (in hours) and FGM/C type

FGM status	Geometric	Crude geometric	Adjusted
	mean	mean ratio (95%	geometric
		CI) (95% CI) *	mean ratio
			(95% CI) **
No FGM	5.9	1.0	1.0
FGM I	8.3	1.4 (1.3-1.5)	1.4 (1.3-1.6)
FGM II	8.9	1.5 (1.4-1.6)	1.6 (1.4-1.7)
FGM III	10.6	1.8 (1.6-2.0)	1.8 (1.6-2.1)

* p-value = <0.001

** p-value = <0.001; Adjusted for parity,

birth weight and maternal body mass index

Discussion

Across all outcomes, as the type of FGM/C moves from type I to type III (corresponding with an increase in the severity of mutilation) then the risks increase further, adding weight to the argument that the association is causal. These findings are in keeping with international data on outcomes of FGM/C in African countries⁵.

Although no statistical significance in perinatal death among women with FGM/C type I and III was observed, this has been shown in a large study with similar settings⁵. Despite being a relatively small study, it was possible to show increased perinatal death in FGM/C type II, with a two-fold increase in risk. It may be that the study was underpowered to demonstrate increased perinatal death in the other types of FGM/C. The increased rate of need for resuscitation is similarly striking and is likely to correlate to increased rates of hypoxic ischaemic encephalopathy (HIE) in babies who do survive with subsequent long-term disability.

This study was carried out at four health-care facilities across the country of the Gambia. As such there was a good representation of the tribal spread and the aim was to proportionally represent the varied population of the country.

There are multiple challenges around conducting research into FGM/C in this setting. Among communities where this is practiced there is an understandable reluctance to discuss the issue. There may be stigma in acknowledging that FGM/C has been performed, or that health consequences occur. As such it is difficult to exclude the risk of recruitment bias for those women willing to join the study. However, the use of local midwives consenting women and performing data collection at each site aimed to ameliorate this risk.

Many of the population lack the means to travel or seek emergency help when required. As such richer women and those with access to healthcare may be over-represented in this study. Similarly, women with pregnancy complications or risk of complex delivery may preferentially seek out healthcare settings and be overrepresented in this study. This is important in interpreting our findings because while 86.2% of pregnant women in the Gambia receive antenatal care from a skilled health professional, only 57.2% of all deliveries are conducted by a skilled health professional¹⁴. A weakness of this study was that not all eligible women presenting to the healthcare facilities during the study window were approached potentially introducing selection bias. Limited resources and availability of staff made this challenging.

A limitation of all studies of medical outcomes of FG-M/C is that they are observational (i.e., it is not possible to randomize). As such it is not possible to draw definitive conclusions about causality and to exclude confounding factors. However, it was only after recruitment into the study that the women were examined for the presence of FGM/C thus ensuring that the groups were as similar as possible. Another limitation of our study is the absence of data on de-infibulation before delivery in type III FGM/C. WHO guidelines suggests that antepartum or intrapartum de-infibulation be done for women with type III FGM/C¹⁵. However, de-infibulation is not mentioned in antenatal care guidelines in the Gambia and there is no data on its routine practice.

The outcomes of need for episiotomy and caesarean section may represent a source of bias. The medical decision to progress to such measures had a subjective element and may show baseline variation in practice between settings. Healthcare worker's experience of FGM/C obstructing labour may lead to increased willingness to progress to these measures when FGM/C is noted to be present.

Studies on obstetric outcomes of FGM/C show that outcomes are worse in resource poor settings whereas there was no significant difference in obstetric outcomes in resource rich countries¹⁶⁻¹⁹. This is probably due to a lack of knowledge and access to antenatal care services in resource poor countries¹⁶. Even though The Gambia

is a resource constrained country, more than 85% of pregnant women receive antenatal care from a skilled car provider¹⁴ and the country has more than the required WHO minimum number of recommended comprehensive emergency obstetric care centres^{20,21}. However, the quality of care available in these health centres needs to be evaluated.

In conclusion, FGM/C in The Gambia was found to be associated with adverse obstetric outcomes, risk of postpartum haemorrhage, unplanned caesarean section, risk of perineal tears and episiotomy. In terms of neonatal outcomes FGM/C is associated with increased risk of need for neonatal resuscitation and FGM/C type II is associated with increased perinatal death. Across all outcomes, as the type of FGM/C moves from type I to type III (corresponding with an increase in the severity of mutilation) then the risks increase further, adding weight to the argument that the association is causal. FGM/C in The Gambia was not shown to be associated with lowbirth-weight babies in this study. We recommend that routine vaginal examination be included as part of the first antenatal care visit in The Gambia and women who are found to have type III FGM/C be referred to health facilities were de-infibulation can be done before delivery. FGM/C has been banned in The Gambia but there is fear that it may be driven underground and continue at high rates. It is hoped that the results of this study will be useful in the advocacy and sensitization of Gambians needed to end this practice. More qualitative research will be needed to understand the factors that are driving the practice in order to finally eliminate it from the Gambia.

Declarations

Ethics approval and consent to participate

The study was approved by the Gambian Government /Medical research Council Joint Ethics Committee with reference number SCC 1474V2.

Sensitization about the study started at the antenatal clinics in all the participating hospitals about 6 weeks before the start of the study and continued until the end of the study. Parturient women in labour were told about the study and a written informed consent was obtained. The consent form containing information about the study was translated into the local languages for those who could not read or understand English. Participants who agreed to be a part of the study were required to sign an informed consent form. Those who could not read or write were allowed to thumbprint this form in the presence of a witness.

Consent to publish

Not applicable.

Availability of data and material

The datasets generated and/or analysed during this study are available from Action Aid International The Gambia on reasonable request.

Competing interests

All the authors declare that they have no competing interests.

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Authors' contribution

PI & MB conceived the study. All the authors contributed to the study design, data collection and manuscript write-up. AA and PI contributed to the data analysis. All the authors read and approved the final version of the manuscript.

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List Of Abbreviations

EFSTH: Edward Francis Small teaching Hospital. FGM: Female genital mutilation. FGM/C: Female genital mutilation or cutting. HIV: Human immunodeficiency virus. LBW: Low birth weight. WHO: World Health Organization.

References

1. World Health Organization. An update on WHO's

work on female genital mutilation (FGM): progress report. Progress Report. Geneva: World Health Organization, Reserch DoRHa; 2011.

2. United Nations Children's Fund. Female Genital Mutilation and Cutting 2016 [cited 2016. Available from: https://data.unicef.org/topic/child-protection/female-genital-mutilation-and-cutting/.

3. El-Shawarby SA, Rymer J. Female Genital Cutting. *Obstetric Gynaecology and Reproductive Medicine*. 2008;18(9):253-5.

4. United States Department of State. The Gambia- Report on Female Genital Mutilation (FGM) or Female Genital Cutting (FGC): US States Department; 2001 [updated 2009; cited 1999. Available from: https://2001-2009. state.gov/g/wi/rls/rep/crfgm/10099.htm. [Accessed 21 May 2019]

5. WHO study group on female genital mutilation and obstetric outcome. Female genital mutilation and obstetric outcome: WHO collaborative prospective study in six African countries. *Lancet.* 2006;367:1835-41 PubMed .

6. Gambia Bureau of Statistics. The Gambia Multiple Indicator Cluster Survey 2010, Final Report Banjul: Gambia Bureau of Statistics; 2011 [cited 2017 19 June 2017]. Available from: www.gbos.gov.gm/uploads/survey/ UNICEFGambiaMICSIV.pdf

7. Morison L, Scherf C, Ekpo G, Paine K, West B, Coleman R, et al. The long-term reproductive health consequences of female genital cutting in rural Gambia: a community-based survey. *Trop Med Int Health.* 2001;6(8):643 PubMed -53.

8. Demba E, Morison L, van der Loeff MS, Awasana AA, Gooding E, Bailey R, et al. Bacterial vaginosis, vaginal flora patterns and vaginal hygiene practices in patients presenting with vaginal discharge syndrome in The Gambia, West Africa. *BMC Infectious Diseases.* 2005; 5:12-.

9. Kaplan A, Hechavarría S, Martín M, Bonhoure I. Health consequences of female genital mutilation/cutting in the Gambia, evidence into action. *Reproductive Health*. 2011;8(1):26 PubMed.

10. Kaplan A, Forbes M, Bonhoure I, Utzet M, Martin M, Manneh M, et al. Female genital mutilation/cutting in The Gambia: long-term health consequences and complications during delivery and for the newborn. *Int J Womens Health.* 2013;5:323-31 PubMed .

11. United Nations Development Program. The Gambia national human development report 2014. Available at: https://www.gm.undp.org/content/gambia/en/home/ library/poverty/national-human-development-report. html [Accessed 19 December 2021]

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12. Unicef. At a glace: Gambia (Statistics). Hämtad 2011-04-27 PubMed från: http://www. unicef. org/infobycountry/gambia_statistics. html; 2010.

13. Gulmezoglu AM, Villar J, Ngoc NTN, Piaggio G, Carroli G, Adetoro L, et al. WHO multicentre randomised trial of misoprostol in the management of the third stage of labour. *The Lancet.* 2001;358:690 - 5 PubMed .

14. Gambia Bureau of Statistics. The Gambia Demographic and Health Survey 2013 Preliminary Report Banjul: Gambia Bureau of Statistics; 2013. Available from: http://www.gbos.gov.gm/uploads/survey/The Gambia Demographic and Health Survey 2013 Preliminary Report.pdf.4. [Accessed 19 December 2021]

15. World Health Organization. WHO guidelines on the management of health complications from female genital mutilation, 2016, available at: https://www.refworld.org/docid/573ad6d54.html [accessed 19 December 2021]

16. Reisel D, Creighton SM. Long term health consequences of Female Genital Mutilation (FGM). *Maturitas*. 2015;80(1):48 PubMed -51.

17. Varol N, Dawson A, Turkmani S, Hall JJ, Nanayakkara S, Jenkins G, et al. Obstetric outcomes for women with female genital mutilation at an Australian hospital, 2006-2012: a descriptive study. *BMC Pregnancy Childbirth*. 2016;16(1):328.

18. Essén B, Sjöberg N-O, Gudmundsson S, Östergren PO, Lindqvist PG. No association between female circumcision and prolonged labour: a case control study of immigrant women giving birth in Sweden. *European Journal of Obstetrics and Gynecology and Reproductive Biology.* 2005;121(2):182-5.

19. Wuest S, Raio L, Wyssmueller D, Mueller MD, Stadlmayr W, Surbek DV, et al. Effects of female genital mutilation on birth outcomes in Switzerland. *BJOG*. 2009;116(9):1204-9.

20. Ministry of Health and Social Welfare TG. National assessment for emergency obstetric and newborn care Banjul: Ministry of Health and Social Welfare; 2012 Available at: http://countryoffice.unfpa.org/thegambia/ drive/NationalAssessmentforEmergency Obstetricand-NewbornCare-FinalReport2012.pdf. [Accessed 19 December 2021]

21. World Health Organization. Monitoring Emergency Obstetric Care: A Handbook. Geneva: World Health Organization; 2009. Available at: https://apps.who.int/iris/ bitstream/handle/10665/44121/9789241547734_eng. pdf?sequence=1 [Accessed on 19 December 2021].