Diagnoses and Procedures of Inpatients with Female Genital Mutilation/Cutting in Swiss University Hospitals: A Cross-Sectional Study

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Research Article

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**Abstract**

**Background:** Female genital mutilation/Cutting (FGM/C) can result in short and long-term complications, which can impact physical, psychological and sexual health. Our objective was to obtain descriptive data about the most frequent health conditions and procedures associated with FGM/C in Swiss university hospitals inpatient women and girls with a diagnosis of FGM/C. Our research focused on the gynaecology and obstetrics departments.

**Methods:** We conducted an exploratory descriptive study to identify the health outcomes of women and girls with a coded FGM/C diagnose who had been admitted to Swiss university hospitals between 2016 and 2018. Four of the five Swiss university hospitals provided anonymized data on primary and secondary diagnoses coded with the International Classification of Diseases (ICD) and interventions coded in their medical files.

**Results:** Between 2016 and 2018, 207 inpatients had a diagnosis of FGM/C. The majority (96%) were admitted either to gynaecology or obstetrics divisions with few genito-urinary and psychosexual conditions coded.

**Conclusions:** FGM/C coding capacities in Swiss university hospitals are low, and some complications of FGM/C are probably not diagnosed. Pregnancy and delivery represent key moments to identify and offer medical care to women and girls who live with FGM/C.

**Trial registration:** This cross-sectional study (protocol number 2018-01851) was conducted in 2019, and approved by the Swiss ethics committee.

**Plain English Summary**

Female genital mutilation/Cutting (FGM/C) can result in short and long-term complications, which can impact physical, psychological and sexual health. Our objective was to obtain descriptive data about the most frequent health conditions and procedures associated with FGM/C among inpatients with a diagnosis of FGM/C in Swiss university hospitals. We asked the Swiss university hospitals for anonymized data of women and girls with a coded FGM/C diagnose who had been admitted between 2016 and 2018. Four of the five Swiss university hospitals provided the primary and secondary diagnoses coded with the International Classification of Diseases (ICD) and the interventions coded in their medical files. Only 207 inpatients had a diagnosis of FGM/C. The majority was admitted either to gynaecology or obstetrics divisions. Some complications of FGM/C are probably not diagnosed. Pregnancy and childbirth represent key moments to care for and counsel a population that might not consult or be identified otherwise.

**Keywords**

Female genital mutilation, female genital cutting, female genital mutilation/cutting, international classification of diseases, ICD, coding, Switzerland

**Introduction**

Female Genital Mutilation/Cutting (FGM/C) comprises all procedures involving partial or total removal of the external female genitalia without medical indication [1]. The World Health Organization (WHO) defines four main types of FGM/C [Table 1] [2]. 200 million women and girls have undergone the practice in 31 countries according to nationally representative household surveys, without counting female migrants with FGM/C who live high-income countries [3, 4]. According to estimates, almost 600,000 individuals living in the European Union are believed to have been exposed to genital cutting (2016) [5], and in Switzerland, approximately 21,706 women and girls are estimated to have undergone the procedure (2018) [6]. These estimates were obtained by indirect measures: multiplying the number of female migrants from an FGM/C practicing country with the FGM/C prevalence rate from the same country. This method does not account for regional and ethnic variations of the practice within countries, and does not include corrections for any changes in attitudes towards FGM/C, which have been described among migrants [7–11]. The actual prevalence of FGM/C among communities of migrants remains unknown [12, 13]. Recent studies conducted in the United Kingdom (UK) showed significantly less cases of FGM/C than expected among minors according to prevalence estimates [14, 15]. Nevertheless, the total number of women and girls who have undergone FGM/C is expected to grow in high-income countries because of increasing migration from countries where FGM/C prevalence remains high [16]. Although several interventions effectively promote
the abandonment of FGM/C, many countries are simultaneously facing population growth, with consequent increase in the absolute number of girls exposed to FGM/C [17].

Table 1

| Type    | Description                                                                 |
|---------|-----------------------------------------------------------------------------|
| Type I  | Partial or total removal of the clitoral glans (the external and visible part of the clitoris, which is a sensitive part of the female genitals, with the function of providing sexual pleasure to the woman), and/or the prepuce/clitoral hood (the fold of skin surrounding the clitoral glans). |
| Type Ia | Removal of the prepuce/clitoral hood only                                   |
| Type Ib | Removal of the clitoral glans with the prepuce/clitoral hood                |

| Type II | Partial or total removal of the clitoral glans and the labia minora, with or without removal of the labia majora. |
|---------|------------------------------------------------------------------------------------------------------------------|
| Type IIa| Removal of the labia minora only                                                                                  |
| Type IIb| Partial or total removal of the clitoris and the labia minora                                                    |
| Type IIc| Partial or total removal of the clitoris, the labia minora and the labia majora                                  |

| Type III (Inbilation) | Narrowing of the vaginal opening with the creation of a covering seal. The seal is formed by cutting and repositioning the labia minora, or labia majora. The covering of the vaginal opening is done with or without removal of the clitoral prepuce/clitoral hood and glans. |
|----------------------|------------------------------------------------------------------------------------------------------------------|
| Type IIIa            | Removal and apposition of the labia minora                                                                       |
| Type IIIb            | Removal and apposition of the labia majora                                                                        |

| Type IV | All other harmful procedures to the female genitalia for non-medical purposes, for example, pricking, piercing, incising, scraping and cauterization. |

It has been widely studied that FGM/C can result in short and long-term complications, which can impact physical, psychological and sexual health [1]. WHO has released an FGM/C cost calculator, revealing the financial burden generated by health complications of FGM/C to be 1.4 billion USD per year [18]. This estimation refers to women and girls living in countries of high-FGM/C prevalence, and not those living in diaspora countries. Systematic reviews and meta-analyses show that female individuals with FGM/C are at higher risk of dyspareunia, genito-urinary complications, prolonged labour and episiotomies [19–21]. Low quality of the studies included was frequently cited as a limitation and some subjects such as the association of FGM/C with caesarean section, infertility and HIV are still a matter of debate [19–21]. Depending on the study design, some of the available data about complications and their clinical management may be subject to self-report and recall bias or inappropriate health management due to lacking training and might therefore be biased, incomplete and/or inaccurate [22]. To our knowledge, no study has yet described FGM/C complications and associated procedures using hospital inpatient data coded with the International Classification of Diseases (ICD).

We sought to describe the most frequent health conditions and procedures associated with FGM/C in inpatient women and girls identified from ICD diagnoses of FGM/C from five Swiss university hospitals.

**Materials And Methods**

This cross-sectional study (protocol number 2018-01851) was conducted in 2019, and approved by the Swiss ethics committee. We invited all five Swiss university hospitals (Geneva, Lausanne, Bern, Basel and Zurich) to provide anonymized data for all inpatient adult women and girls (<18 years) with a nationality from any of the 30 FGM/C practicing countries [3] in addition to all inpatients who had a coded diagnosis of FGM/C between January 1, 2016 and December 31, 2018. We did not include inpatients from the Maldives, where FGM/C has been recently reported [23], because no nationally representative survey was available when the study began.

In Swiss university hospitals, healthcare professionals record the diagnosis responsible for the hospitalization (primary diagnosis); eventual complications that arise during the patient’s hospital stay, as well as any additional diseases treated (secondary diagnoses) in the patients’ electronic medical charts. Professional coders in Switzerland code this information with the German Modification of the tenth edition of the International Classification of Diseases (ICD-10-GM), and interventions are coded with the Swiss Classification of Surgical Interventions (CHOP) [24].
We received the requested data from four university hospitals: Geneva (HUG), Lausanne (CHUV), Bern (Inselspital), and Zurich (USZ). The university hospital of Basel (USB) did not participate due to logistical difficulties in data provision. All data were then merged in a single database using STATA version 15.

The data for all inpatient women and girls from the 30 targeted FGM/C countries and all primary and secondary diagnoses of FGM/C coded between January 1, 2016 and December 31, 2018 was anonymized. The university hospital of Bern did not provide data on the interventions performed. Lausanne and Zürich provided CHOP codes of the interventions performed, and Geneva provided the name of the CHOP interventions. We analyzed all diagnoses and interventions in patients’ records with a coded primary or secondary diagnosis of FGM/C. We provided descriptive statistics with mean, ±standard deviation, and median for continuous variables, numbers by categorical variables. We compared all diagnoses from our sample with the FGM/C ICD “tip-sheet” for FGM/C associated health conditions (full methods available in another manuscript) [25]. We focused our analysis on the gynaecology and obstetrics divisions, where most of the inpatients with an FGM/C code were admitted.

The Swiss Federal Office of Public Health, the Swiss Network against Female Circumcision, and Caritas Switzerland funded the study. They had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript.

Results

In four of the five Swiss university hospitals, 207 inpatients received a primary (n=22, 10.6%) or a secondary (n=185) diagnosis of FGM/C during the study period [Table 2]. Of these 207 women and girls, 199 (96%, 89.4%) were admitted either to gynaecology or obstetrics divisions. The remaining women and girls were admitted to other departments (surgery, internal medicine, emergency, and paediatrics).
Table 2
Description of inpatients with a FGM/C (n=207) as primary or secondary diagnosis between 2016 and 2018 followed in one of four Swiss university hospitals (Geneva, Lausanne, Bern and Zurich).

| Variables                        | 2016 (n=42) | 2017 (n=69) | 2018 (n=96) |
|----------------------------------|-------------|-------------|-------------|
| Center, n (%)                    |             |             |             |
| Geneva                           | 20 (47.6)   | 24 (34.8)   | 67 (69.8)   |
| Lausanne                         | 13 (31.0)   | 10 (14.5)   | 19 (19.8)   |
| Bern                             | 3 (7.1)     | 23 (33.3)   | 6 (6.3)     |
| Zurich                           | 6 (14.3)    | 12 (17.4)   | 4 (4.2)     |
| Country of origin, n (%)         |             |             |             |
| Benin                            | 0 (0)       | 0 (0)       | 1 (1.0)     |
| Burkina Faso                     | 1 (2.4)     | 2 (2.9)     | 0 (0)       |
| Cameroon                         | 1 (2.4)     | 0 (0)       | 0 (0)       |
| Côte d’Ivoire                    | 1 (2.4)     | 1 (1.5)     | 1 (1.0)     |
| Egypt                            | 0 (0)       | 0 (0)       | 5 (5.2)     |
| Eritrea                          | 12 (28.6)   | 37 (53.6)   | 36 (37.5)   |
| Ethiopia                         | 2 (4.8)     | 3 (4.4)     | 2 (2.1)     |
| Guinea                           | 0 (0)       | 0 (0)       | 6 (6.2)     |
| Guinea-Bissau                    | 0 (0)       | 0 (0)       | 2 (2.1)     |
| Mali                             | 0 (0)       | 0 (0)       | 1 (1.0)     |
| Mauritania                       | 0 (0)       | 0 (0)       | 1 (1.0)     |
| Nigeria                          | 1 (2.4)     | 1 (1.5)     | 3 (3.1)     |
| Senegal                          | 0 (0)       | 0 (0)       | 3 (3.1)     |
| Somalia                          | 14 (33.3)   | 18 (26.1)   | 22 (22.9)   |
| Sudan and South Sudan            | 1 (2.4)     | 1 (1.5)     | 3 (3.1)     |
| Unknown or other                 | 9 (21.4)    | 6 (8.7)     | 10 (10.4)   |
| Service, n (%)                   |             |             |             |
| Gynecology                       | 13 (31.0)   | 12 (17.4)   | 9 (9.4)     |
| Gynecology or Obstetricsa        | 1 (2.4)     | 23 (33.3)   | 6 (6.3)     |
| Obstetrics                       | 23 (54.8)   | 33 (47.8)   | 79 (82.3)   |
| Others                           | 5 (11.9)    | 1 (1.5)     | 2 (2.1)     |
| Mean age at first visit (±SD, median) | 30.7 (±12.0, 27) | 27.7 (±6.1, 27.4) | 29.8 (±6.7, 30) |

* Data obtained from Bern did not specify whether patients were admitted in gynecology or obstetrics.
| Variables                  | 2016 (n=42) | 2017 (n=69) | 2018 (n=96) |
|---------------------------|-------------|-------------|-------------|
| FGM/C type, n (%)         |             |             |             |
| Type I                    | 3 (7.1)     | 13 (18.8)   | 10 (10.4)   |
| Type II                   | 8 (19.1)    | 16 (23.2)   | 33 (34.4)   |
| Type III                  | 21 (50.0)   | 33 (47.8)   | 39 (40.6)   |
| Type IV                   | 0 (0)       | 1 (1.5)     | 2 (2.1)     |
| Unspecified or other      | 10 (23.8)   | 6 (8.7)     | 12 (12.5)   |

* Data obtained from Bern did not specify whether patients were admitted in gynecology or obstetrics.

The primary diagnoses of women with a secondary diagnosis of FGM/C (n=185) spanned 11 chapters of the ICD-10 [Table 3]. 156 inpatients had a primary diagnosis related to pregnancy and childbirth. The most frequent diagnoses were perineal laceration during delivery (n=29, 18.6%), labour and delivery complicated by fetal heart rate anomaly (n=16, 10.3%), prolonged second stage of labour (n=13, 8.3%) and premature rupture of membranes (n=13, 8.3%). Nine patients were admitted for some type of anaemia: anaemia complicating pregnancy, childbirth and the puerperium (n=5), iron deficiency anaemia (n=3), and post-haemorrhagic anaemia (n=1). Primary diagnoses of genitourinary diseases included vulvar cysts (n=4), and infectious diseases such as abscess of vulva (n=2), chronic salpingitis and oophoritis (n=1) and pyonephrosis (n=1).

The mean number of secondary diagnoses coded among women with a primary or secondary diagnosis of FGM/C was 2.59 (median 2, range 0-15), spanning 16 chapters of the ICD-10 [Table 4]. There were 281 secondary diagnoses related to pregnancy and childbirth, including 114 codes describing duration of pregnancy (O09.1-O09.7, O48). Other frequent codes were perineal laceration during delivery (n=21), prolonged second stage of labour (n=8), and anaemia complicating pregnancy, childbirth and the puerperium (n=24).
Table 3
Primary diagnoses of inpatients with a secondary diagnosis of FGM/C (n=185) presented by chapter of the ICD-10

| Variables | N  |
|-----------|----|
| ICD-10 chapter and codes | ICD-10 diagnoses |
| Neoplasms | 3 |
| C77.4, C90.00 | Malignant neoplasms |
| D25.9 | Leiomyoma of uterus, unspecified |
| Diseases of the blood and blood-forming organs and certain disorders involving the immune mechanism | 4 |
| D50.8, D50.9 | Iron deficiency anemias |
| D62 | Acute posthemorrhagic anemia |
| Endocrine, nutritional and metabolic diseases | 1 |
| E55.9 | Vitamin D deficiency, unspecified |
| Diseases of the circulatory system | 1 |
| I05.0 | Rheumatic mitral stenosis |
| Diseases of the skin and subcutaneous tissue | 1 |
| L02.2 | Cutaneous abscess, furuncle and carbuncle of trunk |
| Diseases of the genitourinary system | 20 |
| N13.1 | Hydronephrosis with ureteral stricture, not elsewhere classified |
| N13.6 | Pyonephrosis |
| N39.3 | Stress incontinence |
| N70.1 | Chronic salpingitis and oophoritis |
| N76.4 | Abscess of vulva |
| N84.0 | Polyp of corpus uteri |
| N90.7 | Vulvar cyst |
| Pregnancy, childbirth and the puerperium | 156 |
| O00.1 | Tubal pregnancy |
| O02.1 | Missed abortion |
| O09.6 | "Duration of pregnancy 37 to 41 completed weeks, 253 to 287 completed days" |
| O09.7 | "Duration of pregnancy More than 41 completed weeks More than 287 completed days" |
| O12.1 | Gestational proteinuria |
| O14.0, O14.1, O14.9 | Pre-eclampsia |
| O24.0 | Pre-existing type 1 diabetes mellitus, in pregnancy, childbirth and the puerperium |
| O24.4 | Gestational diabetes mellitus |
| O30.0 | Twin pregnancy |
| O32.1 | Maternal care for breech presentation |
| O33.5 | Maternal care for disproportion due to unusually large fetus |
| Variables | N |
|-----------|---|
| O34.2 Maternal care due to uterine scar from previous surgery | 3 |
| O34.30 Maternal care for cervical incompetence, unspecified trimester | 1 |
| O34.7 Maternal care for abnormality of vulva and perineum | 1 |
| O36.5 Maternal care for known or suspected poor fetal growth | 4 |
| O36.6 Maternal care for excessive fetal growth | 2 |
| 041.0 Oligohydramnios | 2 |
| 041.1 Infection of amniotic sac and membranes | 1 |
| O42.0, 042.11, 042.12 Premature rupture of membranes, onset of labor within 24 hours of rupture | 13 |
| O43.21 Placenta accreta | 1 |
| O44.11 Complete placenta previa with hemorrhage, first trimester | 2 |
| 048 Post-term pregnancy | 4 |
| O60.1 Preterm labor with preterm delivery | 2 |
| O61.0 Failed medical induction of labor | 2 |
| O62.8 Other abnormalities of forces of labor | 2 |
| O63.0 Prolonged first stage (of labor) | 4 |
| O63.1 Prolonged second stage (of labor) | 13 |
| O64.8 Obstructed labor due to other malposition and malpresentation | 1 |
| O65.4 Obstructed labor due to fetopelvic disproportion, unspecified | 1 |
| O66.2 Obstructed labor due to unusually large fetus | 1 |
| O66.5 Attempted application of vacuum extractor and forceps | 1 |
| O68.0 Labour and delivery complicated by fetal heart rate anomaly | 16 |
| O68.2 Labour and delivery complicated by fetal heart rate anomaly with meconium in amniotic fluid | 3 |
| O70.0 First degree perineal laceration during delivery | 13 |
| O70.1 Second degree perineal laceration during delivery | 11 |
| O70.2 Third degree perineal laceration during delivery | 1 |
| O70.3 Fourth degree perineal laceration during delivery | 2 |
| O70.9 Perineal laceration during delivery, unspecified | 2 |
| O71.1 Rupture of uterus during labour | 1 |
| O72.0, 072.1 Third-stage haemorrhage | 3 |
| O75.6 Delayed delivery after spontaneous or unspecified rupture of membranes | 1 |
| O75.7 Vaginal delivery following previous caesarean section | 2 |
| 080 Single spontaneous delivery | 3 |
| O98.8 Other maternal infectious and parasitic diseases complicating pregnancy, childbirth and the puerperium | 1 |
| O99.0 Anaemia complicating pregnancy, childbirth and the puerperium | 5 |
| O99.1 Other diseases of the blood and blood-forming organs and certain disorders involving the immune mechanism complicating pregnancy, childbirth and the puerperium | 1 |
| Variables                                                                 | N  |
|---------------------------------------------------------------------------|----|
| O99.2 Endocrine, nutritional and metabolic diseases complicating pregnancy, childbirth and the puerperium | 1  |
| O99.8 Other specified diseases and conditions complicating pregnancy, childbirth and the puerperium | 4  |
| **Congenital malformations, deformations and chromosomal abnormalities**   | 1  |
| Q50.5 Embryonic cyst of broad ligament                                     | 1  |
| **Symptoms, signs and abnormal clinical and laboratory findings, not elsewhere classified** | 1  |
| R74.0 Elevation of levels of transaminase and lactic acid dehydrogenase [LDH] | 1  |
| **Injury, poisoning and certain other consequences of external causes**    | 1  |
| S72.3 Fracture of shaft of femur                                           | 1  |
| **Factors influencing health status and contact with health services**     | 18 |
| Z37.0 Single live birth                                                    | 4  |
| Z65 Problems related to other psychosocial circumstances                   | 1  |
| Z91.70 Personal history of female genital mutilation, type unspecified     | 1  |
| Z91.71 Personal history of female genital mutilation, type 1               | 1  |
| Z91.72 Personal history of female genital mutilation, type 2               | 2  |
| Z91.73 Personal history of female genital mutilation, type 3               | 9  |
Table 4
Secondary diagnoses of inpatients with a diagnosis of FGM/C presented by chapter of the ICD-10

| Variables | N |
|-----------|---|
| **ICD-10 chapter and codes** | **ICD-10 diagnoses** |
| Certain infectious and parasitic diseases | 27 |
| A39.0 | Meningococcal meningitis |
| A60.9 | Anogenital herpesviral infection, unspecified |
| B18.1 | Chronic viral hepatitis B without Delta virus |
| B65.0 | Schistosomiasis due to Schistosoma haematobium [urinary schistosomiasis] |
| B68.1 | Taenia saginata taeniasis |
| B95.1 | Streptococcus, group B, as the cause of diseases classified elsewhere |
| B95.91 | Other specified gram-positive anaerobic, non-spore forming pathogens causing diseases, classified elsewhere |
| B96.2 | Escherichia coli [E. coli ] as the cause of diseases classified elsewhere |
| B98.0 | Helicobacter pylori [H. pylori] as the cause of diseases classified in other chapters |
| Diseases of the blood and blood-forming organs and certain disorders involving the immune mechanism | 33 |
| D50.0 | Iron deficiency anemia secondary to blood loss (chronic) |
| D50.8, D50.9 | Iron deficiency anemias |
| D52.9 | Folate deficiency anemia, unspecified |
| D57.1 | Sickle-cell disease without crisis |
| D62 | Acute posthemorrhagic anemia |
| D64.8, D64.9 | Anemia, other or unspecified |
| D68.4, D68.9 | Coagulation defect |
| D90 | Immune compromise after radiation, chemotherapy and other immunosuppressive measures |
| Endocrine, nutritional and metabolic diseases | 16 |
| E03.8, E03.9 | Hypothyroidism |
| E11.60 | Type 2 diabetes mellitus with other specified complications |
| E44.0 | Moderate protein-calorie malnutrition |
| E53.8 | Deficiency of other specified B group vitamins |
| E55.9 | Vitamin D deficiency, unspecified |
| E66.00, E66.91 | Obesity |
| E83.38 | Other disorders of phosphorus metabolism and phosphatase |
| E87.6 | Hypokalemia |
| Mental, Behavioral and Neurodevelopmental disorders | 3 |
| F32.8 | Other depressive episodes |
| F43.1 | Post-traumatic stress disorder (PTSD) |
| F53.8 | Other postpartum mental and behavioral disorders, not elsewhere classified |
| Variables                                      | N  |
|-----------------------------------------------|----|
| **Diseases of the nervous system**            |    |
| G01 Meningitis in bacterial diseases classified elsewhere | 2  |
| G57.2 Lesion of femoral nerve                  | 1  |
| **Diseases of the circulatory system**        | 5  |
| I05.0, I07.1 Heart valve diseases              | 2  |
| I10.90 Essential hypertension, unspecified: No indication of a hypertensive crisis | 1  |
| I48.9 Unspecified atrial fibrillation and atrial flutter | 1  |
| I95.8 Other hypotension                        | 1  |
| **Diseases of the respiratory system**        | 1  |
| J90 Pleural effusion, not elsewhere classified | 1  |
| **Diseases of the digestive system**          | 5  |
| K21.9 Gastro-esophageal reflux disease without esophagitis | 1  |
| K59.0 Constipation                             | 1  |
| K64.3 Fourth degree hemorrhoids                | 1  |
| K66.1 Hemoperitoneum                           | 1  |
| K66.8 Other specified disorders of peritoneum  | 1  |
| **Diseases of the skin and subcutaneous tissue** | 2  |
| L20.8 Dermatitis                               | 2  |
| **Diseases of the musculoskeletal system and connective tissue** | 1  |
| M54.2 Cervicalgia                              | 1  |
| **Diseases of the genitourinary system**      | 49 |
| N06.8 Isolated proteinuria with other morphologic lesion | 1  |
| N13.6 Pyonephrosis                             | 1  |
| N18.9 Chronic kidney disease, unspecified      | 1  |
| N39.0 Urinary tract infection, site not specified | 1   |
| N73.6 Female pelvic peritoneal adhesions       | 1  |
| N80.3 Endometriosis of pelvic peritoneum       | 1  |
| N83.8 Other noninflammatory disorders of ovary, fallopian tube and broad ligament | 1  |
| N87.0 Mild cervical dysplasia                  | 1  |
| N90.7 Vulvar cyst                              | 1  |
| N90.8 Other specified noninflammatory disorders of vulva and perineum | 1  |
| N90.80 Female genital mutilation, type unspecified | 3  |
| N90.81 Female Genital Mutilation, Type 1       | 3  |
| N90.82 Female Genital Mutilation, Type 2       | 5  |
| N90.83 Female Genital Mutilation, Type 3       | 16 |
### Variables

| Code   | Description                                                                 | N  |
|--------|-----------------------------------------------------------------------------|----|
| N90.88 | Other specified non-inflammatory diseases of the vulva and perineum (FGM, Unspecified or other) | 6  |
| N92.0  | Excessive and frequent menstruation with regular cycle                      | 2  |
| N94.1  | Dyspareunia                                                                 | 2  |
| N94.4  | Primary dysmenorrhea                                                        | 1  |
| N97.1  | Female infertility of tubal origin                                          | 1  |

### Pregnancy, childbirth and the puerperium

| Code   | Description                                                                                   | N  |
|--------|-----------------------------------------------------------------------------------------------|----|
| O08.1  | Delayed or excessive haemorrhage following abortion and ectopic and molar pregnancy          | 1  |
| O09.1  | "Duration of pregnancy 5 to 13 completed weeks, 35 to 91 completed days"                      | 2  |
| O09.2  | "Duration of pregnancy 14 to 19 completed weeks, 92 to 133 completed days"                    | 4  |
| O09.3  | "Duration of pregnancy 20 to 25 completed weeks, 134 to 175 completed days"                   | 2  |
| O09.4  | "Duration of pregnancy 26 to 33 completed weeks, 176 to 231 completed days"                   | 6  |
| O09.5  | "Duration of pregnancy 34 to 36 completed weeks, 232 to 252 completed days"                   | 3  |
| O09.6  | "Duration of pregnancy 37 to 41 completed weeks, 253 to 287 completed days"                   | 72 |
| O09.7  | "Duration of pregnancy more than 41 completed weeks, more than 287 completed days"           | 15 |
| O13    | Gestational [pregnancy-induced] hypertension without significant proteinuria                 | 1  |
| O14.9  | Unspecified pre-eclampsia                                                                      | 1  |
| O16    | Unspecified maternal hypertension                                                              | 2  |
| O24.1, O24.3 | Pre-existing diabetes mellitus, in pregnancy, childbirth and the puerperium               | 4  |
| O24.4  | Gestational diabetes mellitus                                                                  | 6  |
| O32.2  | Maternal care for transverse and oblique lie                                                  | 2  |
| O33.4  | Maternal care for disproportion of mixed maternal and fetal origin                            | 3  |
| O34.2  | Maternal care due to uterine scar from previous surgery                                       | 4  |
| O34.6  | Maternal care for abnormality of vagina                                                       | 3  |
| O34.7  | Maternal care for abnormality of vulva and perineum                                            | 3  |
| O36.0  | Maternal care for rhesus isoimmunization                                                      | 2  |
| O36.5  | Maternal care for known or suspected poor fetal growth                                         | 1  |
| O36.6  | Maternal care for excessive fetal growth                                                      | 1  |
| O41.0  | Oligohydramnios                                                                               | 2  |
| O42.0, O42.11 | Premature rupture of membranes, onset of labor within 24 hours of rupture          | 3  |
| O43.20 | Placenta accreta                                                                              | 2  |
| O44.11 | Complete placenta previa with hemorrhage, first trimester                                    | 1  |
| O45.9  | Premature detachment of the placenta, unspecified                                             | 1  |
| O48    | Post-term pregnancy                                                                           | 10 |
| O60.1  | Preterm labor with preterm delivery                                                           | 1  |
| O60.3  | Preterm delivery without spontaneous labour                                                    | 5  |
| Variables | N |
|-----------|---|
| 061.0     | Failed medical induction of labor | 2 |
| 062.1     | Secondary uterine inertia | 3 |
| 063.0     | Prolonged first stage (of labor) | 2 |
| 063.1     | Prolonged second stage (of labor) | 8 |
| 064.1     | Obstructed labor due to breech presentation | 1 |
| 064.8     | Obstructed labor due to other malposition and malpresentation | 1 |
| 066.8     | Other specified obstructed labor | 2 |
| 068.0     | Labour and delivery complicated by fetal heart rate anomaly | 2 |
| 069.8     | Labour and delivery complicated by other cord complications | 1 |
| 070.0     | First degree perineal laceration during delivery | 14 |
| 070.1     | Second degree perineal laceration during delivery | 7 |
| 071.3     | Obstetric laceration of cervix | 1 |
| 071.8     | Other specified obstetric trauma | 2 |
| 072.0, 072.1 | Third-stage haemorrhage | 9 |
| 072.3     | Postpartum coagulation defects | 1 |
| 073.0     | Retained placenta without haemorrhage | 1 |
| 073.1     | Retained portions of placenta and membranes, without haemorrhage | 1 |
| 075.7     | Vaginal delivery following previous caesarean section | 3 |
| 085       | Puerperal sepsis | 1 |
| 086.2     | Urinary tract infection following delivery | 1 |
| 087.2     | Haemorrhoids in the puerperium | 1 |
| 090.2     | Haematoma of obstetric wound | 1 |
| 098.3     | Other infections with a predominantly sexual mode of transmission complicating pregnancy, childbirth and the puerperium | 1 |
| 098.4     | Viral hepatitis complicating pregnancy, childbirth and the puerperium | 1 |
| 098.8, 098.9 | Maternal infectious and parasitic diseases complicating pregnancy, childbirth and the puerperium | 2 |
| 099.0     | Anaemia complicating pregnancy, childbirth and the puerperium | 24 |
| 099.2     | Endocrine, nutritional and metabolic diseases complicating pregnancy, childbirth and the puerperium | 7 |
| 099.3     | Mental disorders and diseases of the nervous system complicating pregnancy, childbirth and the puerperium | 2 |
| 099.6     | Diseases of the digestive system complicating pregnancy, childbirth and the puerperium | 2 |
| 099.7     | Diseases of the skin and subcutaneous tissue complicating pregnancy, childbirth and the puerperium | 2 |
| 099.8     | Other specified diseases and conditions complicating pregnancy, childbirth and the puerperium | 12 |
| Symptoms, signs and abnormal clinical and laboratory findings, not elsewhere classified | 3 |
| R30.0     | Dysuria | 1 |
| Variables | N |
|-----------|---|
| R74.8     | Abnormal levels of other serum enzymes | 1 |
| R82.3     | Abnormal findings on cytological and histological examination of urine | 1 |
|            | **Definition of HIV infection stages** | 2 |
| U60.9, U61.9 | HIV classification | 2 |
|            | **External causes of morbidity** | 2 |
| Y57.9     | Drug or medicament, unspecified | 1 |
| Y84.9     | Medical procedure, unspecified | 1 |
|            | **Factors influencing health status and contact with health services** | 290 |
| Z21       | Asymptomatic human immunodeficiency virus (HIV) infection status | 1 |
| Z22.3, Z22.8 | Carrier of other infectious diseases | 17 |
| Z25.8, Z27.3, Z27.4 | Need for immunization against specified viral diseases | 8 |
| Z30.4     | Surveillance of contraceptive drugs | 1 |
| Z34       | Supervision of normal pregnancy | 3 |
| Z35.2, Z35.4, Z35.8 | Supervision of high-risk pregnancy | 4 |
| Z37.0     | Single live birth | 91 |
| Z37.2     | Twins, both liveborn | 1 |
| Z59       | Problems related to housing and economic circumstances | 1 |
| Z64.8, Z65 | Problems related to certain psychosocial circumstances | 4 |
| Z86.1     | Personal history of infectious and parasitic diseases | 1 |
| Z86.7     | Personal history of diseases of the circulatory system | 2 |
| Z87.8     | Personal history of other specified conditions | 1 |
| Z91.70    | Personal history of female genital mutilation, type unspecified | 17 |
| Z91.71    | Personal history of female genital mutilation, type 1 | 22 |
| Z91.72    | Personal history of female genital mutilation, type 2 | 47 |
| Z91.73    | Personal history of female genital mutilation, type 3 | 63 |
| Z91.74    | Personal history of female genital mutilation, type 4 | 3 |
| Z92.1     | Personal history of long-term (current) use of anticoagulants | 2 |
| Z94.0     | Kidney transplant status | 1 |

Among diseases of the genitourinary system, coded diagnoses featured vulvar cyst (n=1), urinary tract infection (n=1) and mild cervical dysplasia (n=1). Other secondary diagnoses related to infections were Streptococcus group B (n=17), possibly describing a carrier-state in pregnant women, and carrier of other specified bacterial or infectious diseases (n=17), and asymptomatic HIV status (n=1). Eight women required immunization against viral diseases such as measles, diphtheria, and other viral diseases.

Mental disorders and sexual health conditions were rarely coded as either primary or secondary conditions. “Problems related to psychosocial and/or economic circumstances” appeared five times as secondary diagnosis, and once as a primary diagnosis for a minor inpatient that was admitted in paediatrics. Out of the other four minors with a code of FGM/C (n=5), another was admitted in paediatrics to undergo surgery for mitral valve stenosis, and the remaining two were admitted in gynaecology for surgical treatment of a vulvar cyst. The only minor inpatient with a primary diagnosis of FGM/C underwent defibulation and had secondary codes related to pregnancy.
In total, there were 62 primary and secondary diagnoses of anaemia in 36 patients admitted in gynaecology or obstetrics. Among them, six had third-stage haemorrhage, six a first- or second-degree perineal tear, and nine underwent caesarean section. 27 of 135 patients admitted in obstetrics (19%), had a primary or secondary diagnosis of anemia complicating pregnancy and childbirth.

Several coded diagnoses in our sample might be possible long-term complications of FGM/C found in the FGM/C “tip-sheet” (25) [Table 5]. The most frequently coded diagnoses (primary and secondary combined) were: perineal laceration during delivery (n=50), prolonged second stage of labour (n=21), postpartum haemorrhage (n=12), and vulvar cysts (n=5).
Table 5
Specific codes for long-term complications to FGM/C when FGM/C was coded as primary or secondary diagnosis.

| Variables                                                                 | Primary diagnosis of FGM/C (n=22) | Secondary diagnosis of FGM/C (n=185) |
|--------------------------------------------------------------------------|------------------------------------|--------------------------------------|
| ICD-10 chapters and diagnoses                                           | ICD-10 code                        |                                      |
| **Certain infectious and parasitic diseases**                           |                                    |                                      |
| Human immunodeficiency virus (HIV) disease                              | B20.24 0                          | 0                                    |
| **Mental, Behavioral and Neurodevelopmental disorders**                 |                                    |                                      |
| Recurrent depressive disorder                                           | F32-33 0                           | 1                                    |
| Generalized anxiety disorder                                            | F41.1 0                            | 0                                    |
| Post-traumatic stress disorder                                          | F43.1 0                            | 1                                    |
| Sexual dysfunction, not due to an organic condition                     | F52 0                              | 0                                    |
| **Diseases of the genitourinary system**                                |                                    |                                      |
| Cystitis                                                                | N30 0                              | 0                                    |
| Urinary tract infection, site not specified                             | N39.0 0                            | 1                                    |
| Other inflammation of vagina and vulva                                  | N76 0                              | 0                                    |
| Dysplasia of cervix uteri                                              | N87 0                              | 1                                    |
| Other specified non-inflammatory disorders of vagina                    | N89.8 0                            | 0                                    |
| Vulvar cyst                                                             | N90.7 4                            | 1                                    |
| Non-inflammatory disorder of vulva and perineum, unspecified            | N90.9 0                            | 0                                    |
| Dyspareunia                                                             | N94.1 0                            | 2                                    |
| Dysmenorrhoea, unspecified                                              | N94.6 0                            | 0                                    |
| Other specified conditions associated with female genital organs and menstrual cycle | N94.8 0                            | 0                                    |
| **Pregnancy, childbirth and the puerperium**                            |                                    |                                      |
| Prolonged second stage of labour                                       | O63.1 13                           | 8                                    |
| First degree perineal laceration during delivery                        | O70.0 13                           | 14                                   |
| Second degree perineal laceration during delivery                       | O70.1 11                           | 7                                    |
| Third degree perineal laceration during delivery                        | O70.2 1                            | 0                                    |
| Fourth degree perineal laceration during delivery                       | O70.3 2                            | 0                                    |
| Perineal laceration during delivery, unspecified                        | O70.9 2                            | 0                                    |
| Obstetric high vaginal laceration                                       | O71.4 0                            | 0                                    |
| Other specified obstetric trauma                                       | O71.8 0                            | 2                                    |
| Obstetric trauma, unspecified                                           | O71.9 0                            | 0                                    |
| Postpartum haemorrhage                                                  | O72.0, O72.1 3                     | 9                                    |
| Low forceps delivery                                                    | O81.0 0                            | 0                                    |
| Other and unspecified forceps delivery                                  | O81.3 0                            | 0                                    |
Medical or surgical interventions were carried out in 110 (56.5%) patients with FGM/C: 47 interventions in Geneva, 42 in Lausanne and 22 in Zürich [Table 6]. The most frequent obstetrical intervention was caesarean sections (n=29). 14 patients had an episiotomy and 15 required unspecified manual assistance during delivery. The most frequent intervention aimed at treating complications of FGM/C was surgery of the clitoris (n=11). In Geneva, four inpatients underwent defibulation.
# Table 6

Main intervention reported among patients with FGM/C according to hospital.

| Variables                                      | Geneva (n=111) | Lausanne (n=42) | Zürich (n=22) |
|------------------------------------------------|----------------|-----------------|---------------|
| **Obstetrical interventions**                  |                |                 |               |
| Cerclage of the cervix                         | 1              | 0               | 0             |
| Pharmaceutical induction of labour             | 0              | 1               | 0             |
| Manual assistance during delivery:            |                |                 |               |
| - with episiotomy and instrumentation          | 0              | 1               | 2             |
| - with episiotomy only                         | 0              | 5               | 6             |
| - unspecified                                  | 0              | 13              | 2             |
| Caesarean section                              | 20             | 8               | 1             |
| Perineal tear repair                           | 2              | 6               | 0             |
| Curettage for retained placenta                | 1              | 1               | 0             |
| **Gynaecological interventions**               |                |                 |               |
| Ovarian cyst excision                          | 1              | 0               | 0             |
| Myomectomy                                     | 1              | 0               | 0             |
| Salpingectomy                                  | 1              | 0               | 1             |
| **Interventions related to FGM/C**             |                |                 |               |
| Clitoral surgery                               | 8              | 3               | 0             |
| Vulvar cyst excision                           | 2              | 0               | 0             |
| Vulvar abscess incision and drainage           | 1              | 0               | 0             |
| Defibulation                                   | 4              | 0               | 0             |
| **Interventions possibly related to FGM/C**    |                |                 |               |
| Hymenectomy<sup>a</sup>                       | 0              | 0               | 1             |
|                                              | 0              | 0               | 5             |
|                                              | 0              | 0               | 4             |
|                                              | 0              | 1               | 0             |
|                                              | 0              | 1               | 0             |
| Repair of vulva and perineum<sup>a</sup>       | 0              | 1               | 0             |
| Incision of vulva and perineum<sup>b</sup>    | 1              | 0               | 0             |
| **Other interventions**                        |                |                 |               |
| Femoral fracture repair                        | 1              | 0               | 0             |
| Hematopoietic stem cell transplant             | 0              | 1               | 0             |
| Lymph node biopsy                              | 1              | 0               | 0             |

<sup>a</sup> Patients with FGM/C type III

<sup>b</sup> Three patients with FGM/C type III, one patient with FGM/C type II
Variables | Geneva (n=111) | Lausanne (n=42) | Zürich (n=22)
--- | --- | --- | ---
Mitral valvuloplasty |  |  |  
Retrograde ureteropyelography |  |  |  
Ureteral pigtail placement |  |  |  
Transvaginal suspension for urinary incontinence |  |  |  
Trunk abscess incision and drainage |  |  |  

Discussion

Main Findings

In four Swiss university hospitals, 207 inpatients had a primary (n=22, 10.6%) or secondary (n=185, 89.4%) diagnosis of FGM/C coded at admission between 2016 and 2018 [26]. As discussed in another paper, this was much less than expected when compared with the number of inpatients who could have undergone FGM/C based on their nationality and indirect estimates (n=4947) [26]. Either fewer women than expected have undergone FGM/C, or healthcare professionals did not identify and/or record the practice, resulting in suboptimal coding. Nearly all patients with a coded diagnosis of FGM/C were admitted to an obstetrics and/or gynaecology division, and most of their primary and secondary diagnoses were related to pregnancy and delivery.

Limitations and Strengths

Limitations included the absence of participation from Basel, of interventions’ data from Bern, and the exclusion of non-university hospitals, where most deliveries of women in the cantons of Bern and Zürich occur [Tables 7 and 8] [27–34]. Future studies could assess the prevalence of FGM/C and associated health outcomes in all cantons and hospitals, and study regional variations, such as in areas near asylum centres. We did not study outpatients’ data, which would provide information on the health conditions treated and interventions performed (e.g. defibulation) in ambulatory care. The application of our method is mostly limited by undercoding of FGM/C, which most likely results from insufficient training about FGM/C [26]. Future prospective and case-control studies could assess the coding of FGM/C associated health outcomes according to training resources and possibility of referral to a specialised clinic.
Table 7
Number of deliveries between 2016 and 2018 according to center\textsuperscript{53–59}.

|                | 2016 | 2017 | 2018 |
|----------------|------|------|------|
| Geneva (HUG)   | 4101 | 4182 | 4213 |
| Vaud (CHUV)    | 3230 | 3227 | 3375 |
| Bern (Inselspital) | 1810 | 1827 | 2004 |
| Zürich (USZ)   | 2960 | 2971 | 2969 |

Table 8
Living births according to canton and nationality category of the mother\textsuperscript{60}.

|                | 2016 | 2017 | 2018 |
|----------------|------|------|------|
|                | Total\textsuperscript{a} | Foreigners\textsuperscript{b} | Total | Foreigners | Total | Foreigners |
| Geneva (Geneva) | 5361 | 3108 | 5441 | 3091 | 5353 | 3022 |
| Vaud (Lausanne) | 8730 | 4329 | 8686 | 4336 | 10145 | 4266 |
| Bern (Bem)      | 10113 | 3075 | 10141 | 3026 | 10145 | 2956 |
| Zürich (Zürich) | 17051 | 7449 | 17070 | 7580 | 16919 | 7517 |

\textsuperscript{a} Infants born to women with a Swiss or another nationality

\textsuperscript{b} Infants born to women with another nationality than Switzerland

This study’s main strength was the use of ICD-10 codes to identify health complications of FGM/C, an affordable and objective method, easily reproducible over time and across centres, at national and international level, with good comparability of data. Implementation of training, specific care, and services, as well as financial costs resulting from health complications of FGM/C might also be assessed over time using ICD codes.

**Interpretation**

Women with FGM/C might consult, be admitted or referred more frequently when pregnant, resulting in better FGM/C coding in obstetrics divisions. Furthermore, Swiss basic health insurance covers most costs related to pregnancy, facilitating access to healthcare [35]. Obstetricians and gynaecologists routinely perform genital examinations and are more likely trained to diagnose FGM/C [26]. FGM/C is also more likely to be recorded in obstetrics charts, because it can influence childbirth [1]. For instance, UK’s report on FGM/C prevalence in the National Health System (NHS) showed that 1630 women and girls had a consultation where FGM/C was recorded between October and December 2020, with 74.9% of attendances in midwifery or obstetrical units [36]. Antenatal consultations provide major opportunities to identify and care for individuals with FGM/C who might not seek or receive medical attention otherwise [1, 37].

Meta-analyses including studies from FGM/C practicing countries, and diaspora countries showed that FGM/C was significantly associated with prolonged labour [20, 21], perineal tears, episiotomy, but not with caesarean section [21]. Obstetric outcomes coded in our study were mainly prolonged second stage of labour (n=21) and perineal lacerations (n=50) especially of first or second degree (90%). 29 inpatients required a caesarean section, 14 an episiotomy, and 15 assistance during delivery.

Among 85,990 deliveries in 2017 in Swiss medical institutions, 54.7% of women had a perineal tear mainly of first or second degree (94.7%); 32.3% a caesarean section; 11.1% an assisted delivery and, 17% an episiotomy [38]. In comparison, our data do not suggest high rates of obstetric complications in the studied population of women with a coded FGM/C. However we cannot draw any conclusion, because if common obstetric complications were likely correctly coded, FGM/C was undercoded [26].
A prospective study conducted in six African countries found a significant association between obstetric complications and FGM/C, especially type III [39]. However, retrospective studies from high-income countries such as Sweden, the UK, and Switzerland have shown that women with and without FGM/C have similar obstetric outcomes, likely because of overall quality and availability of obstetric care and specific management of FGM/C [40–42]. The relationship between FGM/C and obstetric complications is influenced by other factors than FGM/C alone. For example, studies on the association between FGM/C and caesarean section have shown different results: a Swiss and Australian studies found higher rates of caesarean sections among women with FGM/C compared with women from the general population without FGM/C [43, 44]. One possible explanation was that health professionals unfamiliar with FGM/C could perform caesarean sections for inappropriate reasons, especially in case of infibulation [45]. In contrast, a retrospective study that included only Somali women who delivered between 1990 and 2014 in Norway showed that women with FGM/C did not undergo more caesarean sections than non-migrants [46]. Migrant women in high-income countries often have higher rates of caesarean sections than non-migrants [47]. Communication barriers, economic difficulties, and exposure to violence have been found to result in poor maternal health and/or insufficient care quality for some migrants regardless of FGM/C [48–52].

Only five minor inpatients had an FGM/C code. Outpatient clinics may attend more children with FGM/C than university hospitals, but paediatricians may also miss and/or not record it, because they lack knowledge on FGM/C, or rarely perform a genital examination [53–55]. Alternatively, abandonment of the practice could explain why few minors had an FGM/C code. A UK study confirmed 55 out of 148 suspected cases of FGM/C after referral to specialized clinics over four years, representing considerably fewer cases than previously suggested by UK prevalence estimates [14]. 21% of 55 children suffered from mental health symptoms such as anxiety, sleep and behaviour disorders, 13% from physical symptoms such as problems with micturition, menstruation and genital pain [14]. Except one post-traumatic stress disorder, mental health symptoms were not coded in our minor population, and rarely among adults. Swiss university hospitals’ health professionals may lack time or training on how to detect and treat such symptoms and other FGM/C complications. Or, they may identify and manage psychophysical complications, without however identifying or documenting the FGM/C as an associated condition [54–60].

Coding of surgical interventions was incomplete. Perineal tears were more coded (n=50) than perineal tears repairs (n=8). Other repairs were either not coded, or coded as secondary interventions, which were not provided to us. Because no CHOP codes exist for defibulation and clitoral reconstruction, we had to hypothesize that codes such as repair (n=5), or incision (n=4) of vulva and perineum had been used to indicate these surgeries. Geneva provided the interventions’ names instead of codes, and reported 8 clitoral surgeries and 4 defibulations among inpatients, and additionally reported 12 clitoral surgeries, 25 defibulations and 8 other surgeries for scar complications of FGM/C in outpatient care. Unfortunately some Swiss insurance companies have tried to refuse to reimburse these surgeries. Specific CHOP codes would facilitate medical coding and reimbursement.

Sensitisation and training of healthcare professionals and professional coders on FGM/C could improve identification, documentation and coding of FGM/C and its complications in Swiss university hospitals; inform and improve the quality of future policies, services and interventions.

**Conclusion**

Most of the 207 women and girls admitted to Swiss university hospitals between 2016 and 2018 with a primary or secondary diagnosis of FGM/C were admitted to obstetrics divisions. Pregnancy and delivery seem to be key moments to care for and counsel a population that might not consult or be identified otherwise. FGM/C coding capacities in Swiss university hospitals are low, and some complications of FGM/C are probably not diagnosed, or diagnosed alone, without FGM/C.

**Abbreviations**

CHOP
Swiss Classification of Surgical Interventions

CHUV
University Hospital of Lausanne

FGM/C
Female Genital Mutilation/Cutting

HUG
Declarations

Ethics approval

This cross-sectional study was approved in December, 2018 by the Swiss Ethics Committees (SwissEthics) with the protocol number 2018–01851, and conducted according to the protocol, the Swiss legal requirements, and the World Medical Association Declaration of Helsinki. An exemption of informed consent was granted by the state of Geneva Swiss Ethics committee for the use of anonymized data extracted from the university hospitals databases.

Consent for publication

Not applicable.

Availability of data and materials

The datasets used and/or analysed during the current study available from the corresponding author on reasonable request.

Competing interests

No competing interests to declare.

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Authors’ contributions

JA designed and directed the project. SCC collected the data, together with JA and MH. SCC did the data analysis, in collaboration with MH. MH, SCC and JA wrote the manuscript. All the authors reviewed and approved the manuscript.

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