Intersex Genital Mutilations in ICD-10

Zwischengeschlecht.org / StopIGM.org 2014 (v2.1)

ICD-10 Codes and Descriptions: http://apps.who.int/classifications/icd10/browse/2010/en

1. Reference: 17 Most Common IGM Procedures (a–q)

Source: 2014 CRC NGO Report, **Supplement 2 "Most Common Forms of IGMs"**, p. 63–76:
http://intersex.shadowreport.org/public/2014-CRC-Swiss-NGO-Zwischengeschlecht-Intersex-IGM_v2.pdf

2. IGMs: List of Relevant ICD-10 Codes

Chapter IV. Endocrine, nutritional and metabolic diseases
Disorders of other endocrine glands (E20—E35)

E23.0 Hypopituitarism
- Fertile eunuch syndrome
- **Hypogonadotropic hypogonadism**
- Idiopathic growth hormone deficiency
- Isolated deficiency of:
  - gonadotropin
  - growth hormone
  - pituitary hormone
- **Kallmann syndrome**
- Lorain-Levi short stature
- Necrosis of pituitary gland (postpartum)
- Panhypopituitarism
- Pituitary:
  - cachexia
  - insufficiency NOS
  - short stature
- Sheehan syndrome
- Simmonds disease

E25 Adrenogenital disorders **a, b, c, d, e, f, h, i, j, k, l, m, n, o, p, q**
E25.0 Congenital adrenogenital disorders associated with enzyme deficiency
Congenital adrenal hyperplasia
21-Hydroxylase deficiency
Salt-losing congenital adrenal hyperplasia
E25.8 Other adrenogenital disorders
Idiopathic adrenogenital disorder
E25.9 Adrenogenital disorder, unspecified
Adrenogenital syndrome NOS

E28. Androgen excess
Hypersecretion of ovarian androgens

E29. Testicular dysfunction
E29.0 Testicular hyperfunction
Hypersecretion of testicular hormones
E29.1 Testicular hypofunction **a, b, c, d, e, g, h, i, j, k, l, m, n, o, p, q**
5-Alpha-reductase deficiency (with male pseudohermaphroditism)
Defective biosynthesis of testicular androgen NOS
Testicular hypogonadism NOS
Use additional external cause code (Chapter XX), if desired, to identify drug, if drug-induced.
E29.8 Other testicular dysfunction
E29.9 Testicular dysfunction, unspecified

E30 Disorders of puberty, not elsewhere classified
E30.0 Delayed puberty
Constitutional delayed puberty
Delayed sexual development

E34 Other endocrine disorders
E34.5 Androgen resistance syndrome **a, b, c, d, e, f, g, h, i, j, k, l, m, n, o, p, q**
Male pseudohermaphroditism with androgen resistance
Peripheral hormonal receptor disorder
Reifenstein syndrome
Testicular feminization (syndrome)

--> N46, N48, N50.0

Chapter XVII Congenital malformations, deformations and chromosomal abnormalities (Q00-Q99)
Congenital malformations of genital organs (Q50-Q56)
Q50 Congenital malformations of ovaries, fallopian tubes and broad ligaments
Q50.0 Congenital absence of ovary
Q50.3 Other congenital malformations of ovary
Accessory ovary
Congenital malformation of ovary NOS
Congenital malformation of fallopian tube or broad ligament NOS

Q51 Congenital malformations of uterus and cervix

Q51.0Agenesis and aplasia of uterus
Congenital absence of uterus
Q51.5 Agenesis and aplasia of cervix
Congenital absence of cervix
Q51.6 Embryonic cyst of cervix
Q51.7 Congenital fistulae between uterus and digestive and urinary tracts
Q51.8 Other congenital malformations of uterus and cervix
Hypoplasia of uterus and cervix
Q51.9 Congenital malformation of uterus and cervix, unspecified

Q52 Other congenital malformations of female genitalia
Q52.0 Congenital absence of vagina **d, e, i, j, k, l, m, n, o**
Q52.1 Doubling of vagina
Q52.4 Other congenital malformations of vagina
Congenital malformation of vagina NOS
Q52.5 Fusion of labia
Q52.6 Congenital malformation of clitoris
Q52.7 Other congenital malformations of vulva

Q52.8 Other specified congenital malformations of female genitalia
Q52.9 Congenital malformation of female genitalia, unspecified

Q53Undescended testicle **g, h, i, j, k, l, m, n, o**
Q53.0 Ectopic testis
Unilateral or bilateral ectopic testes
Q53.1 Undescended testicle, unilateral
Q53.2 Undescended testicle, bilateral
Q53.9 Undescended testicle, unspecified
Cryptorchism NOS

Q54Hypospadias **b, i, j, k, l, m, n, o**
Q54.0 Hypospadias, balanic
Hypospadias:
- coronal
- glandular

Q54.1 Hypospadias, penile
Q54.2 Hypospadias, penoscrotal
Q54.3 Hypospadias, perineal
Q54.4 Congenital chordee
Q54.8 Other hypospadias
Q54.9 Hypospadias, unspecified

Q55 Other congenital malformations of male genital organs
Q55.0 Absence and aplasia of testis
Monorchism
Q55.1 Hypoplasia of testis and scrotum
Fusion of testes
Q55.2 Other congenital malformations of testis and scrotum
Congenital malformation of testis or scrotum NOS
Polyorchism

Retractile testis **g, h, i, j, k, l, m, n, o**
Testis migrans **g, h, i, j, k, l, m, n, o**
Q55.3 Atresia of vas deferens
Q55.4 Other congenital malformations of vas deferens, epididymis, seminal vesicles and prostate
Q55.5 Congenital absence and aplasia of penis
Q55.6 Other congenital malformations of penis
Congenital malformation of penis NOS
Curvature of penis (lateral)
Hypoplasia of penis
Q55.8 Other specified congenital malformations of male genital organs
Q55.9 Congenital malformation of male genital organ, unspecified

Q56 Indeterminate sex and pseudohermaphroditism
a, b, c, d, e, f, g, h, i, j, k, l, m, n, o
Q56.0 Hermaphroditism, not elsewhere classified
Ovotestis
Q56.1 Male pseudohermaphroditism, not elsewhere classified
Male pseudohermaphroditism NOS
Q56.2 Female pseudohermaphroditism, not elsewhere classified
Female pseudohermaphroditism NOS
Q56.3 Pseudohermaphroditism, unspecified
Q56.4 Indeterminate sex, unspecified
Ambiguous genitalia

Other congenital malformations (Q80-89)
Congenital malformations of breast
Q83.0 Congenital absence of breast with absent nipple
Q83.1 Accessory breast
Supernumerary breast
Q83.2 Absent nipple
Q83.3 Accessory nipple
Supernumerary nipple
Q83.8 Other congenital malformations of breast
Hypoplasia of breast
Q83.9 Congenital malformation of breast, unspecified

Other congenital malformations, not elsewhere classified (Q90-99)
Q96Turner syndrome **p, q**
Q96.0 Karyotype 45,X
Q96.1 Karyotype 46,X iso (Xq)
Q96.2 Karyotype 46,X with abnormal sex chromosome, except iso (Xq)
Q96.3 Mosaicism, 45,X/46,XX or XY
Q96.4 Mosaicism, 45,X/other cell line(s) with abnormal sex chromosome
Q96.8 Other variants of Turner syndrome
Q96.9 Turner syndrome, unspecified

Q97Other sex chromosome abnormalities, female phenotype, not elsewhere classified
Q97.0 Karyotype 47,XXX
Q97.1 Female with more than three X chromosomes

Q97.2 Mosaicism, lines with various numbers of X chromosomes
Q97.3 Female with 46,XY karyotype
Q97.8 Other specified sex chromosome abnormalities, female phenotype
Q97.9 Sex chromosome abnormality, female phenotype, unspecified

Q98 Other sex chromosome abnormalities, male phenotype, not elsewhere classified
f, h, p, q
Q98.0 Klinefelter syndrome karyotype 47,XXY
Q98.1 Klinefelter syndrome, male with more than two X chromosomes
Q98.2 Klinefelter syndrome, male with 46,XX karyotype
Q98.3 Other male with 46,XX karyotype
Q98.4 Klinefelter syndrome, unspecified
Q98.5 Karyotype 47,XYY
Q98.6 Male with structurally abnormal sex chromosome
Q98.7 Male with sex chromosome mosaicism
Q98.8 Other specified sex chromosome abnormalities, male phenotype
Q98.9 Sex chromosome abnormality, male phenotype, unspecified

Q99 Other chromosome abnormalities, not elsewhere classified
a, b, c, d, e, f, g, h, i, j, k, l, m, n, o, p, q
Q99.0 Chimera 46,XX/46,XY
Chimera 46,XX/46,XY true hermaphrodite
Q99.1 46,XX true hermaphrodite
46,XX with streak gonads
46,XY with streak gonads
Pure gonadal dysgenesis
Q99.2 Fragile X chromosome
Fragile X syndrome