ESPE Code	***Diagnosis of disorders of the ovaries and female reproductive tract and breasts according to the European society of Pediatric Endocrinology (ESPE)***	**OMIM**	**ICD-10**
10A	**OVARY**		E28.3
10A.1	**Primary ovarian failure (hypergonadotrophic hypogonadism)** *Possible supplementary codes:*		
3E.1a.0	*primary amenorrhoea*		
3E.1b.0	*secondary amenorrhoea*		E28.3
10A.1a	Due to disorder classified elsewhere: *Steroidogenic block: CAH* (8A.1) *Aromatase deficiency* (4C.2b) *Turner syndrome* (14A.5) *Autoimmune polyglandular syndrome* (14C.4a)		
10A.1b	Gonadal agenesis	600171	Q99.1
10A.1c	Gonadal dysgenesis		Q99.1
10A.1c.1	Pure 46,XX gonadal dysgenesis (complete or incomplete)		Q99.1
10A.1c.2	Gonadal dysgenesis with other specified chromosomal/genetic abnormality (e.g. trisomy 13, trisomy 18, trisomy 21, Denys Drash syndrome in XX individual)		
10A.1c.3	Mixed gonadal dysgenesis		Q99.8
10A.1c.8	Other, specified gonadal dysgenesis, e.g. 47,XXX, etc.		Q97.0 Q97.1 Q97.2 Q97.8
10A.1c.9	Gonadal dysgenesis, unspecified		Q98.9
10A.1d	Post-ablative ovarian failure, e.g. post-irradiation, post-surgical, post-chemotherapy		E89.4 E28.3
10A.1e	Resistant ovary syndrome (Savage syndrome, mutation of FSH receptor gene)	*136435	E28.3
10A.1y	Due to other specified disorder, e.g. infection/oophoritis, autoimmune SLE		E28.3
10A.1z	Idiopathic/unspecified		E28.3
10A.2	**Ovarian androgen excess**		E28.1
10A.2a	Polycystic ovary syndrome	#184700	E28.2
10A.2b	Other causes		E28.2
10A.3	**Ovarian cysts and tumours**		
10A.3a	Ovarian follicular cyst *Note: if associated with precocious pseudopuberty:* 3A.2c.1		N83.0
10A.3b	Corpus luteum cyst		N83.1
10A.3c	Cysts, unspecified		N83.2
10A.3d	Germ cell tumours		Benign: D27 Malignant: C56
10A.3e	Non-germ cell tumours		Benign: D27 Malignant: C56
10A.3e.1	Granulosa tumour		
10A.3e.2	Other specified tumours		

10B	**DISORDERS OF THE UTERUS AND CERVIX** *Functional disorders are classified in* 3E, *menstrual disorders*		
10B.1	**Congenital malformations**		Q51
10B.1a	Agenesis and aplasia of the uterus (Müllerian agenesis/Mayer-Rokitansky-Kuster-Hauser syndrome, Müllerian-renal-cervical spine (MURCS) syndrome)	#277000	Q51.0
10B.1b	Congenital absence of the cervix (isolated)		Q51.5
10B.1c	Endometrial hypoplasia/aplasia		Q51.8
10B.1d	Incomplete Müllerian fusion [includes: double uterus (uterus didelphy), half uterus (uterus unicornis), partial duplication (uterus bicornis, Fryns syndrome), partial or complete uterine septum (uterus septus and subseptus)]	192050	Q51.2–Q51.4
10B.1y	Other specified congenital malformations of uterus and cervix		Q51.8 Q51.6 Q51.7 Q51.1
10B.1z	Other congenital malformations of uterus and cervix, unspecified		Q51.9
10B.2	**Acquired disorders of the uterus and cervix**		
10B.2a	Uterine synechiae/Asherman syndrome		N85.6
10B.2b	Cervical stenosis		N88.2
10B.2y	Other specified acquired malformations of uterus and cervix		
10B.2z	Acquired malformations of uterus and cervix, unspecified		
10B.3	**Tumours of uterus and cervix**		Benign: D25, D26 Malignant: C53, C54
10B.8	**Other specified disorder of uterus and cervix, e.g. polyps**		N84.0 N84.1
10B.9	**Disorder of uterus and cervix, unspecified**		
10C	**DISORDERS OF THE VAGINA AN D EXTERNAL FEMALE GENITALIA** *Possible secondary codes:*		
3E.1a.0	*primary amenorrhoea*		
3E.1b.0	*secondary amenorrhoea*		
10C.1	**Congenital malformations**		Q52
10C.1a	Vaginal agenesis (isolated)		Q52.0
10C.1b	Imperforate hymen (can be part of McKusick-Kaufman syndrome)	#236700	Q52.3
10C.1c	Transverse vaginal septum		Q52.8
10C.1d	Labial fusion/agglutination		Q52.5
10C.1e	Congenital malformation of the clitoris *Excluded:* Clitoromegaly due to endocrine causes/virilisation (4C.2)		Q52.6
10C.1y	Other specified congenital malformations of the female external genitalia and vagina		Q52.8
10C.1z	Congenital malformations of female external genitalia and vagina, unspecified		Q52.9
10C.2	**Acquired disorders of the vagina and external female genitalia**		
10C.2a	Adhaesions (vaginal, labial, vulval)		N89.5 N90.8
10C.2b	Acquired disorders of the clitoris *Excluded:* Clitoromegaly due to endocrine causes (3C.2)		N90.8
10C.2c	Trauma		S30.2
10C.2z	Other disorders, unspecified		S30.2

10C.3	**Tumours of the vagina and external female genitalia**		Benign: D28.0 D28.1 D28.7 D28.9 Malignant: C51, C52
10C.8	**Other, specified, disorders of the vagina and external female genitalia** E.g. polyps		N84.2– N84.9
10C.9	**Disorders of the vagina and external female genitalia, unspecified**		
10D	**DISORDERS OF THE BREAST**		
10D.1	**Galactorrhoea (not associated with childbirth)** *Excluded:* Galactorrhoea in the male (3C.1)		N64.3
10D.2	**Disorders of size**		
10D.2a	Hypoplasia/aplasia/hypomastia/micromastia		Q83.8
10D.2b	Macromastia		N62
10D.3	**Disorders in numbers**		
10D.3a	Polythelia, polymastia	163700%	Q83.1
10D.3b	Absence of breast and nipple (athelia)	113700	Q83.0
10D.4	**Tumours of breasts**		Benign: D24 Malignant: C50
10D.8	**Other specified disorders of the breast**		N60, N61 N63– N64.8 Q83.2 Q83.3 Q83.8
10D.9	**Other disorder of the breast, unspecified**		N64.9 Q83.9