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International Statistical Classification of Diseases and Related Health Problems 10th Revision expanded

Chapter XII
Diseases of the skin and subcutaneous tissue
(L00-L99)

Disorders of skin appendages
(L60-L75)

Excl.: congenital malformations of integument ([Q84.-](#))

L60 **Nail disorders**
Excl.: clubbing of nails ([R68.3](#))
onychia and paronychia ([L03.0](#))

- L60.0** **Ingrowing nail**
- L60.1** **Onycholysis**
- L60.2** **Onychogryphosis**
- L60.3** **Nail dystrophy**
- L60.4** **Beau lines**
- L60.5** **Yellow nail syndrome**
- L60.8** **Other nail disorders**
- L60.9** **Nail disorder, unspecified**

- L62*** **Nail disorders in diseases classified elsewhere**
- L62.0*** **Clubbed nail pachydermoperiostosis ([M89.4†](#))**
- L62.8*** **Nail disorders in other diseases classified elsewhere**

- L63** **Alopecia areata**
- L63.0** **Alopecia (capitis) totalis**
- L63.1** **Alopecia universalis**
- L63.2** **Ophiasis**
- L63.8** **Other alopecia areata**
- L63.9** **Alopecia areata, unspecified**

- L64** **Androgenic alopecia**
Incl.: male-pattern baldness
- L64.0** **Drug-induced androgenic alopecia**
Use additional external cause code (Chapter XX), if desired, to identify drug
- L64.8** **Other androgenic alopecia**
- L64.9** **Androgenic alopecia, unspecified**

- L65** **Other nonscarring hair loss**
Use additional external cause code (Chapter XX), if desired, to identify cause
- Excl.:** trichotillomania ([F63.3](#))

- L65.0** **Telogen effluvium**
- L65.1** **Anagen effluvium**
- L65.2** **Alopecia mucinosa**
- L65.8** **Other specified nonscarring hair loss**
- L65.9** **Nonscarring hair loss, unspecified**
Alopecia NOS

- L66** **Cicatricial alopecia [scarring hair loss]**
- L66.0** **Pseudopelade**
- L66.1** **Lichen planopilaris**
Follicular lichen planus

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- L66.2 Folliculitis decalvans
- L66.3 Perifolliculitis capitis abscedens
- L66.4 Folliculitis ulerythematosae reticulata
- L66.8 Other cicatricial alopecia
- L66.9 Cicatricial alopecia, unspecified

L67 Hair colour and hair shaft abnormalities

Excl.: monilethrix ([Q84.1](#))
pili annulati ([Q84.1](#))
telogen effluvium ([L65.0](#))

L67.0 Trichorrhexis nodosa

L67.1 Variations in hair colour

Canities
Greyness, hair (premature)
Heterochromia of hair
Poliosis:

- NOS
- circumscripta, acquired

L67.8 Other hair colour and hair shaft abnormalities

Fragilitas crinium

L67.9 Hair colour and hair shaft abnormality, unspecified

L68 Hypertrichosis

Incl.: excess hair

Excl.: congenital hypertrichosis ([Q84.2](#))
persistent lanugo ([Q84.2](#))

L68.0 Hirsutism

Use additional external cause code (Chapter XX), if desired, to identify

L68.1 Acquired hypertrichosis lanuginosa

Use additional external cause code (Chapter XX), if desired, to identify

L68.2 Localized hypertrichosis

L68.3 Polytrichia

L68.8 Other hypertrichosis

L68.9 Hypertrichosis, unspecified

L70 Acne

Excl.: acne keloid ([L73.0](#))

L70.0 Acne vulgaris

L70.1 Acne conglobata

L70.2 Acne varioliformis
Acne necrotica miliaris

L70.3 Acne tropica

L70.4 Infantile acne

L70.5 Acné excoriée
Acné excoriée des jeunes filles

L70.8 Other acne

L70.9 Acne, unspecified

L71 Rosacea

L71.0 Perioral dermatitis

Use additional external cause code (Chapter XX), if desired, to identify

L71.1 Rhinophyma

L71.8 Other rosacea

L71.9 Rosacea, unspecified

L72 Follicular cysts of skin and subcutaneous tissue

L72.0 Epidermal cyst



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- L68.2 Localized hypertrichosis
- L68.3 Polytrichia
- L68.8 Other hypertrichosis
- L68.9 Hypertrichosis, unspecified

- L70 Acne
 - Excl.: acne keloid ([L73.0](#))
- L70.0 Acne vulgaris
- L70.1 Acne conglobata
- L70.2 Acne varioliformis
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- L70.8 Other acne
- L70.9 Acne, unspecified

- L71 Rosacea
 - L71.0 Perioral dermatitis
 - Use additional external cause code (Chapter XX), if desired, to identify cause
 - L71.1 Rhinophyma
 - L71.8 Other rosacea
 - L71.9 Rosacea, unspecified

- L72 Follicular cysts of skin and subcutaneous tissue
 - L72.0 Epidermal cyst
 - L72.1 Trichilemmal cyst
 - Pilar cyst
 - Sebaceous cyst
 - L72.2 Steatocystoma multiplex
 - L72.8 Other follicular cysts of skin and subcutaneous tissue
 - L72.9 Follicular cyst of skin and subcutaneous tissue, unspecified

- L73 Other follicular disorders
 - L73.0 Acne keloid
 - L73.1 Pseudofolliculitis barbae
 - L73.2 Hidradenitis suppurativa
 - L73.8 Other specified follicular disorders
 - Sycosis barbae
 - L73.9 Follicular disorder, unspecified

- L74 Eccrine sweat disorders
 - Excl.: hyperhidrosis ([R61.-](#))
 - L74.0 Miliaria rubra
 - L74.1 Miliaria crystallina
 - L74.2 Miliaria profunda
 - Miliaria tropicalis
 - L74.3 Miliaria, unspecified
 - L74.4 Anhidrosis
 - Hypohidrosis
 - L74.8 Other eccrine sweat disorders
 - L74.9 Eccrine sweat disorder, unspecified
 - Sweat gland disorder NOS



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Chapter XVII
Congenital malformations, deformations and chromosomal anomalies (Q00-Q99)

Congenital malformations of eye, ear, face and neck (Q10-Q18)

- Excl.:** cleft lip and cleft palate ([Q35-Q37](#))
congenital malformation of:
- cervical spine ([Q05.0](#), [Q05.5](#), [Q67.5](#), [Q76.0-Q76.4](#))
 - larynx ([Q31.-](#))
 - lip NEC ([Q38.0](#))
 - nose ([Q30.-](#))
 - parathyroid gland ([Q89.2](#))
 - thyroid gland ([Q89.2](#))

Q10 Congenital malformations of eyelid, lacrimal apparatus and orbit

- Excl.:** cryptophthalmos:
- NOS ([Q11.2](#))
 - syndrome ([Q87.0](#))

Q10.0 Congenital ptosis

Q10.1 Congenital ectropion

Q10.2 Congenital entropion

Q10.3 Other congenital malformations of eyelid

- Ablepharon
Absence or agenesis of:
- cilia
 - eyelid
- Accessory:
- eye muscle
- Blepharophimosis, congenital
Coloboma of eyelid
Congenital malformation of eyelid NOS

Q10.4 Absence and agenesis of lacrimal apparatus

- Absence of punctum lacrimale

Q10.5 Congenital stenosis and stricture of lacrimal duct

Q10.6 Other congenital malformations of lacrimal apparatus

- Congenital malformation of lacrimal apparatus NOS

Q10.7 Congenital malformation of orbit

Q11 Anophthalmos, microphthalmos and macropthalmos

Q11.0 Cystic eyeball

Q11.1 Other anophthalmos

- Agenesis
Aplasia
- of eye

Q11.2 Microphthalmos

- Cryptophthalmos NOS
Dysplasia of eye
Hypoplasia of eye
Rudimentary eye
- Excl.:** cryptophthalmos syndrome ([Q87.0](#))

Q11.3 Macropthalmos

- Excl.:** macropthalmos in congenital glaucoma ([Q15.0](#))

Q12 Congenital lens malformations

Q12.0 Congenital cataract

Q12.1 Congenital displaced lens

Q12.2 Coloboma of lens



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Q12.3 Congenital aphakia

Q12.4 Spherophakia

Q12.8 Other congenital lens malformations

Q12.9 Congenital lens malformation, unspecified

Q13 Congenital malformations of anterior segment of eye

Q13.0 Coloboma of iris

Coloboma NOS

Q13.1 Absence of iris

Aniridia

Q13.2 Other congenital malformations of iris

Anisocoria, congenital

Atresia of pupil

Congenital malformation of iris NOS

Corectopia

Q13.3 Congenital corneal opacity

Q13.4 Other congenital corneal malformations

Congenital malformation of cornea NOS

Microcornea

Peter anomaly

Q13.5 Blue sclera

Q13.8 Other congenital malformations of anterior segment of eye

Axenveld-Rieger syndrome

Rieger anomaly

Q13.9 Congenital malformation of anterior segment of eye, unspecified

Q14 Congenital malformations of posterior segment of eye

Q14.0 Congenital malformation of vitreous humour

Congenital vitreous opacity

Q14.1 Congenital malformation of retina

Congenital retinal aneurysm

Q14.2 Congenital malformation of optic disc

Coloboma of optic disc

Q14.3 Congenital malformation of choroid

Q14.8 Other congenital malformations of posterior segment of eye

Coloboma of the fundus

Q14.9 Congenital malformation of posterior segment of eye, unspecified

Q15 Other congenital malformations of eye

Excl.: congenital nystagmus ([H55](#))

ocular albinism ([E70.3](#))

retinitis pigmentosa ([H35.5](#))

Q15.0 Congenital glaucoma

Buphthalmos

Glaucoma of newborn

Hydrophthalmos

Keratoglobus, congenital, with glaucoma

Macrocornea with glaucoma

Macrophthalmos in congenital glaucoma

Megalocornea with glaucoma

Q15.8 Other specified congenital malformations of eye

Q15.9 Congenital malformation of eye, unspecified

Congenital:

- anomaly NOS of eye
- deformity

Q16 Congenital malformations of ear causing impairment of hearing

Excl.: congenital deafness ([H90.-](#))

Q16.0 Congenital absence of (ear) auricle



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Q16.1	Congenital absence, atresia and stricture of auditory canal (excl.) Atresia or stricture of osseous meatus
Q16.2	Absence of eustachian tube
Q16.3	Congenital malformation of ear ossicles Fusion of ear ossicles
Q16.4	Other congenital malformations of middle ear Congenital malformation of middle ear NOS
Q16.5	Congenital malformation of inner ear Anomaly: • membranous labyrinth • organ of Corti
Q16.9	Congenital malformation of ear causing impairment of hearing Congenital absence of ear NOS
Q17	Other congenital malformations of ear <i>Excl.:</i> preauricular sinus (Q18.1)
Q17.0	Accessory auricle Accessory tragus Polyotia Preauricular appendage or tag Supernumerary: • ear • lobule
Q17.1	Macrotia
Q17.2	Microtia
Q17.3	Other misshapen ear Pointed ear
Q17.4	Misplaced ear Low-set ears <i>Excl.:</i> cervical auricle (Q18.2)
Q17.5	Prominent ear Bat ear
Q17.8	Other specified congenital malformations of ear Congenital absence of lobe of ear
Q17.9	Congenital malformation of ear, unspecified Congenital anomaly of ear NOS
Q18	Other congenital malformations of face and neck <i>Excl.:</i> cleft lip and cleft palate (Q35-Q37) conditions classified to Q67.0-Q67.4 congenital malformations of skull and face bones (Q75.-) cyclopia (Q87.0) dentofacial anomalies [including malocclusion] (K07.-) malformation syndromes affecting facial appearance (Q87.0) persistent thyroglossal duct (Q89.2)
Q18.0	Sinus, fistula and cyst of branchial cleft Branchial vestige
Q18.1	Preauricular sinus and cyst Fistula (of): • auricle, congenital • cervicoaural Pretragal sinus and cyst
Q18.2	Other branchial cleft malformations Branchial cleft malformation NOS Cervical auricle Otocephaly
Q18.3	Webbing of neck Pterygium colli
Q18.4	Macrostomia
Q18.5	Microstomia