List of top level categories

CHAPTER 01 Certain infectious or parasitic diseases

1V00\_DER Gastroenteritis or colitis of infectious origin

1A6Z Syphilis

1A73 Disseminated gonococcal infection

1A80 Chlamydial lymphogranuloma

1V04\_DER Chlamydial female pelvic inflammatory disease

1V05\_DER Chlamydial epididymitis

1A90 Chancroid

1A91 Granuloma inguinale

1A94 Anogenital herpes simplex infection

1A95 Anogenital warts

1V06\_DER BCG-induced regional lymphadenopathy

1V07\_DER Tuberculosis of anal canal

1V08\_DER Tuberculosis of anus and rectum

1B12.8 Cutaneous tuberculosis

1V09\_DER Acute miliary cutaneous tuberculosis

1B20 Leprosy

Leprosy reactions (BlockL2‑1B2)

1B20.20 Type I leprosy reaction

1B20.21 Type II leprosy reaction

Complications of leprosy (BlockL2‑1V0)

1V0A\_DER Arthritis due to leprosy

1V0B\_DER Polyneuropathy due to leprosy

1V0C\_DER Involvement of eyelid in leprosy

1V0D\_DER Neuropathic skin ulceration due to leprosy

1V0E\_DER Xerosis cutis due to leprosy

1V0F\_DER Scarring alopecia due to lepromatous leprosy

1V0Z\_DER Other specified complications of leprosy

Cutaneous non-tuberculous mycobacterial infection (BlockL1‑1V0)

1V0G\_DER Mycobacterium marinum infection

1B21.20 Mycobacterium ulcerans infection

1V0H\_DER Cutaneous Mycobacterium avium-intracellulare infection

1V0J\_DER Cutaneous Mycobacterium fortuitum infection

1V0K\_DER Cutaneous Mycobacterium kansasii infection

1B21.2Y\_DER Cutaneous infection due to other specified non-tuberculous mycobacteria

1B21.2Z Cutaneous infection due to unspecified non-tuberculous mycobacteria

1B21.3 Disseminated non-tuberculous mycobacterial infection

1V0L\_DER Pneumonia in rheumatic fever without mention of heart involvement

1B50 Scarlet fever

1B52 Toxic shock syndrome

1V0M\_DER Skin infection classified elsewhere due to Panton-Valentine leucocidin producing Staphylococcus aureus

Pyogenic bacterial infections of the skin or subcutaneous tissues (BlockL1‑1B7)

1B70 Bacterial cellulitis, erysipelas or lymphangitis

1B71 Necrotising fasciitis

1B72 Impetigo

1B73 Ecthyma

1B74 Superficial bacterial folliculitis

Deep bacterial folliculitis or pyogenic abscess of the skin (BlockL2‑1B7)

1B75.0 Furuncle

1B75.1 Carbuncle

1B75.2 Furunculosis

1B75.3 Pyogenic abscess of the skin

1B75.4 Chronic deep bacterial folliculitis

Miscellaneous pyogenic bacterial infections of the skin (BlockL2‑1V0)

1V0N\_DER Cutaneous botryomycosis

Pyogenic infection of skin (BlockL3‑1V0)

1V0P\_DER Pyoderma vegetans

1V0Q\_DER Chancriform pyoderma

1V0Z\_DER Other specified pyogenic infection of skin

1V0Z\_DER Pyogenic infection of skin, unspecified

1V0Z\_DER Bacterial pyoderma

1B7Y\_DER Other specified pyogenic bacterial infection of skin and subcutaneous tissue

Rat-bite fevers (BlockL1‑1B9)

1B90.0 Spirillosis

1B90.1 Streptobacillosis

1B91 Leptospirosis

1B92 Glanders

1B93.0 Bubonic plague

1B93.1 Cellulocutaneous plague

Erysipeloid (BlockL1‑1V0)

1V0R\_DER Cutaneous erysipeloid

1V0Z\_DER Other specified erysipeloid

1V0Z\_DER Erysipeloid, unspecified

1V0S\_DER Cutaneous anthrax

1C10.2 Cervicofacial actinomycosis

1C10.3 Primary cutaneous actinomycosis

1C11.01 Verruga peruana

1C16 Gas gangrene

1V0T\_DER Primary ocular diphtheria

1V0U\_DER Diphtheritic myocarditis

1V0V\_DER Diphtheritic renal tubular necrosis

1C1A.0 Cutaneous listeriosis

1C1B.1 Cutaneous nocardiosis

1V0W\_DER Meningococcal conjunctivitis

1V0X\_DER Meningococcal retrobulbar neuritis

1C1C.0 Meningococcal meningitis

1C1C.20 Acute meningococcaemia

1V0Y\_DER Chronic meningococcaemia

1V0Z\_DER Meningococcal carditis

1V10\_DER Meningococcal endocarditis

1V11\_DER Meningococcal myocarditis

1V12\_DER Meningococcal pericarditis

1V13\_DER Postmeningococcal arthritis

Non-venereal treponematoses (BlockL1‑1C1)

1C1D Yaws

1C1E Pinta

1C1F Endemic non-venereal syphilis

1C1G.0 Early cutaneous Lyme borreliosis

1V14\_DER Acrodermatitis chronica atrophicans

Relapsing fever (BlockL1‑1C1)

1C1J.0 Tick-borne relapsing fever

1C1J.1 Louse-borne relapsing fever

1V15\_DER Pneumonia in Q fever

Systemic bacterial infection affecting skin (BlockL1‑1V1)

1V16\_DER Skin ulceration or abscess due to systemic bacterial infection

1V17\_DER Skin rash due to distant or systemic bacterial infection, not elsewhere classified

1C42 Melioidosis

1C43 Actinomycetoma

Non-pyogenic bacterial infections of the skin (BlockL1‑1V1)

1V18\_DER Erythrasma

1V19\_DER Pitted keratolysis

1V1A\_DER Axillary trichomycosis

1V1B\_DER Trichomycosis pubis

1V1Z\_DER Other specified non-pyogenic bacterial infections of the skin

Skin disorders associated with Human immunodeficiency virus infection (BlockL1‑1V1)

1V1C\_DER HIV-associated eosinophilic folliculitis

1V1D\_DER HIV-associated papular pruritic eruption

1V1E\_DER HIV-associated pruritus

1V1F\_DER HIV-modified psoriasis

1V1G\_DER HIV-associated seborrhoeic dermatitis

1V1H\_DER HIV-modified skin disease

1V1J\_DER HIV-associated lipodystrophy

1C8C Venezuelan equine encephalitis

1D2Z Dengue

Certain arthropod-borne viral fevers (BlockL1‑1D4)

1D40 Chikungunya virus disease

1D41 Colorado tick fever

1D42 O'nyong-nyong fever

1D44 Rift Valley fever

1D45 Sandfly fever

1D46 West Nile virus infection

Yellow fever (BlockL2‑1V1)

1V1K\_DER Sylvatic yellow fever

1V1L\_DER Urban yellow fever

1V1Z\_DER Other specified yellow fever

1V1Z\_DER Yellow fever, unspecified

1D49 Crimean-Congo haemorrhagic fever

1D4A Omsk haemorrhagic fever

1D4B Kyasanur Forest disease

1D4Y\_DER Other specified arthropod-borne viral fevers

1D4Z Arthropod-borne viral fever, virus unspecified

1D60.0 Ebola disease

1D60.1 Marburg disease

1D61.0 Argentinian haemorrhagic fever

1D61.1 Bolivian haemorrhagic fever

1D61.2 Lassa fever

1D62.0 Haemorrhagic fever with renal syndrome

1V1M\_DER Newcastle conjunctivitis

1V1N\_DER Encephalitis in influenza with other manifestations, other influenza virus identified

1V1P\_DER Influenzal myocarditis, other influenza virus identified

1V1Q\_DER Encephalitis in influenza with other manifestations, virus not identified

1V1R\_DER Influenzal myocarditis, virus not identified

1E71 Monkeypox

1E72 Cowpox

1E73 Vaccinia

1V1S\_DER Paravaccinia

1E74 Buffalopox

1E75 Orf

1V1T\_DER Tanapox

1E76 Molluscum contagiosum

Human papillomavirus infection of skin or mucous membrane (BlockL1‑1E8)

1E80 Common warts

1E81 Plane warts

1E82 Warts of lips or oral cavity

Wart virus proliferation in immune-deficient states (BlockL2‑1V1)

1V1U\_DER Epidermodysplasia verruciformis

1V1V\_DER Viral warts due to iatrogenic immunosuppression

1V1W\_DER Viral warts due to acquired immunodeficiency

1E8Z Viral warts, not elsewhere classified

1E90.0 Varicella without complication

1E91.0 Zoster without complications

1E91.1 Ophthalmic zoster

1E91.2 Disseminated zoster

1V1X\_DER Postherpetic neuralgia

1F00 Herpes simplex infections

Herpes simplex infection of skin or mucous membrane (BlockL2‑1F0)

1F00.00 Herpes simplex infection of skin

1F00.01 Herpes simplex labialis

1F00.02 Herpes simplex gingivostomatitis

1F00.03 Disseminated cutaneous herpes simplex infection complicating other skin diseases

1V1Y\_DER Herpes simplex pharyngotonsillitis

1V1Z\_DER Herpes simplex infection of external ear

1V20\_DER Paronychial herpes simplex infection

1F00.0Y\_DER Other specified herpes simplex infection of skin or mucous membrane

1F01 Roseola infantum

1F02.2 Rubella without complication

1F03.0 Measles without complication

1F04 Erythema infectiosum

Picornavirus infections presenting in the skin or mucous membranes (BlockL1‑1F0)

1F05.0 Enteroviral vesicular stomatitis

1F05.1 Enteroviral vesicular pharyngitis

1F05.2 Enteroviral exanthematous fever

1F05.3 Foot and mouth disease

1F05.Y\_DER Other specified picornavirus infections presenting in the skin or mucous membranes

1F20.10 Aspergillus otomycosis

1F21 Basidiobolomycosis

1V21\_DER Primary cutaneous blastomycosis

1V22\_DER Disseminated blastomycosis

Candidosis of lips or oral mucous membranes (BlockL1‑1V2)

1V23\_DER Angular cheilitis due to candidosis

1V24\_DER Acute erythematous oral candidosis

1V25\_DER Acute pseudomembranous oral candidosis

1V26\_DER Chronic erythematous oral candidosis

1V27\_DER Chronic hyperplastic oral candidosis

1V28\_DER Chronic multifocal oral candidosis

Candidosis of skin or mucous membranes (BlockL1‑1F2)

Candidosis of external genitalia (BlockL2‑1F2)

1F23.10 Vulvovaginal candidosis

1F23.11 Candida balanoposthitis

1F23.12 Flexural or intertriginous candidosis

1F23.13 Candidosis of nail or paronychium

1F23.14 Chronic mucocutaneous candidosis

1F23.15 Disseminated cutaneous candidosis

1F23.16 Candida otomycosis

1F23.1Y\_DER Candidosis of skin or mucous membrane of other specified site

1F23.1Z Candidosis of skin or mucous membranes, unspecified

1F24 Chromoblastomycosis

1V29\_DER Coccidioidal pneumonia

1F25.1 Extrathoracic coccidioidomycosis

1F26 Conidiobolomycosis

1V2A\_DER Cutaneous cryptococcosis

Dermatophytosis (BlockL1‑1V2)

Dermatophytosis due to anthropophilic dermatophytes (BlockL2‑1V2)

1V2B\_DER Dermatophytosis due to Epidermophyton floccosum

1V2C\_DER Dermatophytosis due to Microsporum audouinii

1V2D\_DER Dermatophytosis due to Microsporum ferrugineum

1V2E\_DER Dermatophytosis due to Trichophyton concentricum

1V2F\_DER Dermatophytosis due to Trichophyton gourvilii

1V2G\_DER Dermatophytosis due to Trichophyton interdigitale

1V2H\_DER Dermatophytosis due to Trichophyton megninii

1V2J\_DER Dermatophytosis due to Trichophyton rubrum

1V2K\_DER Dermatophytosis due to Trichophyton schoenleinii

1V2L\_DER Dermatophytosis due to Trichophyton soudanense

1V2M\_DER Dermatophytosis due to Trichophyton tonsurans

1V2N\_DER Dermatophytosis due to Trichophyton violaceum

1V2P\_DER Dermatophytosis due to Trichophyton yaoundei

1V2Z\_DER Dermatophytosis due to other specified anthropophilic dermatophytes

Dermatophytosis due to zoophilic dermatophytes (BlockL2‑1V2)

1V2Q\_DER Dermatophytosis due to Microsporum canis

1V2R\_DER Dermatophytosis due to Microsporum equinum

1V2S\_DER Dermatophytosis due to Microsporum gallinae

1V2T\_DER Dermatophytosis due to Microsporum nanum

1V2U\_DER Dermatophytosis due to Microsporum persicolor

1V2V\_DER Dermatophytosis due to Trichophyton equinum

1V2W\_DER Dermatophytosis due to Trichophyton mentagrophytes

1V2X\_DER Dermatophytosis due to Trichophyton simii

1V2Y\_DER Dermatophytosis due to Trichophyton verrucosum

1V3Z\_DER Dermatophytosis due to other specified zoophilic dermatophytes

Dermatophytosis due to geophilic dermatophytes (BlockL2‑1V2)

1V2Z\_DER Dermatophytosis due to Microsporum gypseum

1V30\_DER Dermatophytosis due to Microsporum praecox

1V3Z\_DER Dermatophytosis due to other specified geophilic dermatophytes

Dermatophytosis organised by site of involvement (BlockL2‑1F2)

1F28.0 Dermatophytosis of scalp

1V31\_DER Dermatophytosis of beard

1V32\_DER Dermatophytosis of face

1F28.1 Dermatophytosis of nail

1V33\_DER Dermatophytosis of hand

1F28.2 Dermatophytosis of foot

1V34\_DER Dermatophytosis of trunk or limbs

1F28.3 Genitocrural dermatophytosis

Special forms of dermatophytosis (BlockL2‑1F2)

1F28.4 Kerion

1V35\_DER Corticosteroid-modified dermatophytosis

1F28.5 Disseminated dermatophytosis

1V36\_DER Granulomatous dermatophytosis

1F28.Y\_DER Other specified dermatophytosis

1F28.Z Dermatophytosis, unspecified

1F29 Eumycetoma

1V37\_DER Acute pulmonary histoplasmosis capsulati

1V38\_DER Disseminated histoplasmosis capsulati

1F2A.1 Histoplasmosis due to Histoplasma duboisii

1F2B Lobomycosis

1V39\_DER Cutaneous mucormycosis

Non-dermatophyte superficial dermatomycoses (BlockL1‑1F2)

1F2D.0 Pityriasis versicolor

1F2D.1 Malassezia folliculitis

1F2D.2 White piedra

1F2D.3 Black piedra

1F2D.4 Tinea nigra

1V3A\_DER Neoscytalidium dermatomycosis

1F2D.5 Onychomycosis due to non-dermatophyte mould

1F2D.Y\_DER Other specified non-dermatophyte superficial dermatomycoses

1F2E.1 Disseminated paracoccidioidomycosis

1V3B\_DER Extrapulmonary paracoccidioidomycosis

1V3C\_DER Mucocutaneous paracoccidioidomycosis

1V3D\_DER Pulmonary fibrosis due to chronic paracoccidioidomycosis

1V3E\_DER Fibrosis of upper airways and oropharynx due to paracoccidioidomycosis

1V3F\_DER Hypoadrenalism and adrenal failure due to paracoccidioidomycosis

1F2F Phaeohyphomycosis

1V3G\_DER Pneumocystis colonization

1V3H\_DER Pneumocystis associated pneumothorax

1V3J\_DER Pneumocystis associated cysts or pneumatocoele

1F2J Sporotrichosis

Cutaneous or lymphocutaneous sporotrichosis (BlockL2‑1F2)

1F2J.0 Lymphocutaneous sporotrichosis

1F2J.1 Fixed cutaneous sporotrichosis

1F2K Talaromycosis

1F2L.0 Disseminated adiaspiromycosis

1V3K\_DER Glomerular disorders in plasmodium malariae malaria

1V3L\_DER Conjunctivitis due to Acanthamoeba

1V3M\_DER Megaoesophagus in Chagas disease

1V3N\_DER Post-kala-azar dermal leishmaniasis

1F54.1 Cutaneous leishmaniasis

1F54.2 Mucocutaneous leishmaniasis

1F56 Rhinosporidiosis

1F64 Dracunculiasis

1F66.0 Loiasis

1F66.1 Mansonelliasis

1V3P\_DER Dirofilariasis

1F66.3 Lymphatic filariasis

1F67 Gnathostomiasis

Hookworm diseases (BlockL1‑1F6)

1F68.0 Ancylostomiasis

1F68.1 Necatoriasis

1F68.2 Cutaneous larva migrans

Onchocerciasis of the skin (BlockL1‑1V3)

1V3Q\_DER Onchodermatitis

1V3R\_DER Onchocercoma

1V3Z\_DER Other specified onchocerciasis of the skin

1V3S\_DER Glomerular disorders in strongyloidiasis

1V3T\_DER Cutaneous strongyloidiasis

1V3U\_DER Glomerular disorders in disseminated strongyloidiasis

1F6E Trichinosis

1V3V\_DER Schistosomiasis involving the skin

1F86.4 Cercarial dermatitis

1G0Z Infestations by ectoparasites

Infestation by mites (BlockL2‑1G0)

1G07.0 Infestation by Demodex

1G07.Y\_DER Infestation of the skin by other specified parasitic mites

1G60.0 Mycetoma of unknown or unspecified type

1G60.1 Pythiosis

1G60.2 Protothecosis

1H0Z Infection, unspecified

CHAPTER 02 Neoplasms

2B3Z Neoplasms of haematopoietic or lymphoid tissues

Mature T-cell or NK-cell lymphoma, primary cutaneous specified types (BlockL2‑2B0)

2B00 Subcutaneous panniculitis-like T-cell lymphoma

2B01 Mycosis fungoides

Mycosis fungoides variants (BlockL4‑2V3)

2V3X\_DER Folliculotropic mycosis fungoides

2V3Y\_DER Syringotropic mycosis fungoides

2V3Z\_DER Pagetoid reticulosis

2V40\_DER Granulomatous slack skin

2V4Z\_DER Other specified mycosis fungoides variants

2V4Z\_DER Mycosis fungoides variants, unspecified

2B02 Sézary syndrome

Primary cutaneous CD-30 positive T-cell lymphoproliferative disorders (BlockL3‑2B0)

2B03.0 Primary cutaneous CD30 positive anaplastic large cell lymphoma

2B03.1 Lymphomatoid papulosis

2V41\_DER Primary cutaneous aggressive epidermotropic CD8 positive T-cell lymphoma

2V42\_DER Primary cutaneous gamma delta T-cell lymphoma

2V43\_DER Primary cutaneous CD4+ small or intermediate pleomorphic T-cell lymphoproliferative disorder

2V44\_DER Hydroa vacciniforme-like cutaneous T-cell lymphoma

2V45\_DER Adult T-cell leukaemia or lymphoma, skin

2V46\_DER Primary cutaneous acral CD8+ T-cell lymphoma

2B0Y\_DER Other specified primary cutaneous mature T-cell or NK-cell lymphomas and lymphoproliferative disorders

2B0Z Primary cutaneous T-cell lymphoma of undetermined or unspecified type

2V47\_DER Dermatofibrosarcoma protuberans

2B56.1 Angiosarcoma of skin

2B57.1 Kaposi sarcoma of skin

2V48\_DER Cutaneous leiomyosarcoma

Malignant neoplasms of skin (BlockL1‑2C3)

2C30 Melanoma of skin

Squamous cell carcinoma of skin (BlockL2‑2C3)

2C31.0 Verrucous squamous cell carcinoma of skin

2C31.Z Cutaneous squamous cell carcinoma

Basal cell carcinoma of skin (BlockL2‑2V4)

2V49\_DER Fibroepithelial basal cell carcinoma of skin

2C32.0 Nodular basal cell carcinoma of skin

2C32.1 Sclerosing basal cell carcinoma of skin

2C32.2 Superficial basal cell carcinoma of skin

2C32.Y\_DER Other specified basal cell carcinoma of skin

2C32.Z Basal cell carcinoma of skin, unspecified

Adnexal carcinoma of skin (BlockL2‑2V4)

2V4A\_DER Aggressive digital papillary adenocarcinoma

2V4B\_DER Apocrine carcinoma

2V4C\_DER Malignant neoplasm of sebaceous glands

2V4Z\_DER Other specified adnexal carcinoma of skin

2V4Z\_DER Adnexal carcinoma of skin, unspecified

Cutaneous neuroendocrine carcinoma (BlockL2‑2V4)

2V4D\_DER Merkel cell carcinoma of skin

2V4Z\_DER Other specified cutaneous neuroendocrine carcinoma

2V4Z\_DER Cutaneous neuroendocrine carcinoma, unspecified

2C3Y\_DER Other specified malignant neoplasms of skin

2C3Z Malignant neoplasm of skin of unknown or unspecified type

2C70.2 Squamous cell carcinoma of vulva

2C81.0 Squamous cell carcinoma of penis

Metastatic malignant neoplasm involving skin (BlockL1‑2V4)

2V4E\_DER Metastatic melanoma involving skin

2V4F\_DER Metastatic carcinoma involving skin

2V4G\_DER Skin infiltration by nodal or non-cutaneous extranodal lymphoma

2V4H\_DER Leukaemic infiltration of skin

2V4Z\_DER Malignant infiltration of skin by other specified malignant neoplasm

2V4Z\_DER Malignant infiltration of skin

Melanoma in situ neoplasms (BlockL1‑2E6)

2E63.0 Melanoma in situ of skin

2E63.1 Melanoma in situ of conjunctiva

2E63.Y\_DER Other specified melanoma in situ neoplasms

2E63.Z Melanoma in situ neoplasms, unspecified

Carcinoma in situ of skin (BlockL1‑2E6)

2E64.0 Intraepidermal squamous cell carcinoma

2E64.1 Extramammary Paget disease of skin

2E64.2 Carcinoma in situ of anal margin or perianal skin

2E64.Y\_DER Other specified carcinoma in situ of skin

2E64.Z Carcinoma in situ of skin, unspecified

2E67.1 Carcinoma in situ of vulva

2E67.4 Carcinoma in situ of penis

2E80.01 Deep subfascial lipoma

2E80.02 Deep internal or visceral lipoma

2V4J\_DER Familial multiple lipomata

2V4K\_DER Haemangioma of orbit

2E81.2 Benign vascular neoplasms of infancy and childhood

Benign cutaneous melanocytic neoplasms (BlockL1‑2F2)

2F20.0 Common acquired melanocytic naevus

2V4L\_DER Blue naevus

2V4M\_DER Spindle cell melanocytic naevus

Special types of melanocytic naevus (BlockL2‑2V4)

2V4N\_DER Acral melanocytic naevus

2V4P\_DER Genital melanocytic naevi

2V4Q\_DER Combined melanocytic naevus

2V4R\_DER Halo melanocytic naevus

2V4S\_DER Melanocytic naevus with eczematous halo

2V4T\_DER Irritated melanocytic naevus

2V4U\_DER Persistent melanocytic naevus

2V4Z\_DER Other specified special types of melanocytic naevus

2V4Z\_DER Special types of melanocytic naevus, unspecified

2F20.1 Atypical melanocytic naevus

2V4V\_DER Unclassifiable benign cutaneous melanocytic proliferation

2F20.2 Congenital melanocytic naevus

2F20.Y\_DER Other specific types of melanocytic naevus

2F20.Z Melanocytic naevus, unspecified

2F21 Benign keratinocytic acanthomas

2F22 Benign neoplasms of epidermal appendages

Benign cutaneous neoplasms with follicular differentiation (BlockL2‑2V4)

2V4W\_DER Pilomatrixoma

2V4X\_DER Eruptive vellus hair cysts

2V4Y\_DER Trichoepithelioma of eyelids

2V4Z\_DER Multiple familial trichoepithelioma

2V5Z\_DER Other specified benign cutaneous neoplasms with follicular differentiation

2V5Z\_DER Benign cutaneous neoplasms with follicular differentiation, unspecified

Benign cutaneous neoplasms with sebaceous differentiation (BlockL2‑2V5)

2V50\_DER Steatocystoma simplex

2V51\_DER Steatocystoma multiplex

2V5Z\_DER Other specified benign cutaneous neoplasms with sebaceous differentiation

2V5Z\_DER Benign cutaneous neoplasms with sebaceous differentiation, unspecified

2F23 Benign dermal fibrous or fibrohistiocytic neoplasms

2V52\_DER Benign tumours of cutaneous smooth muscle

Benign proliferations or neoplasms of cutaneous blood vessels (BlockL1‑2F2)

2F25 Cherry angioma

2F26 Lobular capillary haemangioma

2V53\_DER Epithelioid haemangioma of skin

2V54\_DER Glomus tumour of the skin or nail apparatus

2V5Z\_DER Other specified benign vascular neoplasm of skin

2F24 Benign cutaneous neoplasms of neural or nerve sheath origin

2F33.0 Vulvar intraepithelial neoplasia, grade I, usual type, HPV-associated

2F72 Neoplasms of uncertain behaviour of skin

CHAPTER 03 Diseases of the blood or blood-forming organs

3V55\_DER Glomerular disorders in haemolytic uraemic syndrome

3V56\_DER Osteonecrosis due to acquired pure red cell aplasia

3V57\_DER Fanconi anaemia

3V58\_DER Dyskeratosis congenita

3B20 Disseminated intravascular coagulation

3B62 Qualitative platelet defects

3B64.10 Immune thrombocytopenic purpura

3B64.12 Drug-induced thrombocytopenic purpura

3B64.14 Thrombotic thrombocytopenic purpura

3C0Y\_DER Other specified diseases of the blood or blood-forming organs

3C0Z Diseases of the blood or blood-forming organs, unspecified

CHAPTER 04 Diseases of the immune system

4A0Z Primary immunodeficiencies

Angioedema due to disordered complement activation or kinin metabolism (BlockL2‑4A0)

4A00.14 Hereditary angioedema

4A00.15 Acquired angioedema

4A40 Lupus erythematosus

4A41.0 Dermatomyositis

4A42 Systemic sclerosis

Overlap or undifferentiated nonorgan specific systemic autoimmune disease (BlockL1‑4V5)

4V59\_DER Antisynthetase syndrome

4A43.0 IgG4 related disease

4A43.1 Mikulicz disease

4V5A\_DER Reynolds syndrome

4V5B\_DER Syndromic multisystem autoimmune disease due to ITCH deficiency

4A43.2 Sjögren syndrome

4A43.3 Mixed connective tissue disease

4A43.4 Diffuse eosinophilic fasciitis

4A43.Y\_DER Other specified overlap non-organ specific systemic autoimmune disease

4A43.Z Undifferentiated non-organ specific systemic autoimmune disease

4A44.2 Giant cell arteritis

4A44.4 Polyarteritis nodosa

4A44.5 Mucocutaneous lymph node syndrome

4A44.6 Sneddon syndrome

4A44.8 Thromboangiitis obliterans

4A44.90 Cryoglobulinaemic vasculitis

4A44.91 Hypocomplementaemic urticarial vasculitis

4A44.92 IgA vasculitis

4A44.A Antineutrophil cytoplasmic antibody-associated vasculitis

4A44.B Leukocytoclastic vasculitis

4V5C\_DER Vasculitis associated with probable aetiology

4A45 Antiphospholipid syndrome

Monogenic autoinflammatory syndromes (BlockL1‑4V5)

4V5D\_DER Autoinflammatory granulomatosis of childhood

4V5E\_DER Pyogenic arthritis - pyoderma gangrenosum - acne

4A60.0 Familial Mediterranean fever

4V5F\_DER Mevalonate kinase deficiency with recurrent fever

Cryopyrin-associated periodic syndromes (BlockL2‑4V5)

4V5G\_DER Chronic infantile neurological, cutaneous and articular syndrome

4V5H\_DER Familial cold autoinflammatory syndrome

4V5J\_DER Muckle-Wells syndrome

4V5Z\_DER Other specified cryopyrin-associated periodic syndromes

4A60.2 Tumour necrosis factor receptor 1 associated periodic syndrome

4A60.Y\_DER Other specified monogenic autoinflammatory syndromes

4A60.Z Autoimflammatory syndrome, unspecified

4A61 SAPHO syndrome

4V5K\_DER Chronic recurrent multifocal osteomyelitis

4A62 Behçet disease

4A84.1 Drug-induced anaphylaxis

4A84.30 Exercise-induced anaphylaxis

4V5L\_DER Drug-associated immune complex vasculitis

4A85.31 Cutaneous allergic or hypersensitivity reactions to Hymenoptera venom

4V5M\_DER Certain disorders involving the immune system

4V5S\_DER Castleman disease

4B4Y\_DER Other specified diseases of the immune system

4B4Z Diseases of the immune system, unspecified

CHAPTER 05 Endocrine, nutritional or metabolic diseases

5B3Z Endocrine diseases

5V5T\_DER Phrynoderma

5B5B Vitamin B2 deficiency

5B5C Vitamin B3 deficiency

5B5G Biotin deficiency

5B56.0 Scurvy

5B90.0 Hypervitaminosis A

Dermatoses resulting from defective nutrition (BlockL1‑5V5)

5V5U\_DER Skin disorder attributable to malnutrition or malabsorption

5V5V\_DER Skin disorder attributable to food fads or unhealthy diet

5V6Z\_DER Other specified dermatoses resulting from defective nutrition

5V6Z\_DER Dermatoses resulting from defective nutrition, unspecified

5C50.00 Classical phenylketonuria

5C50.10 Alkaptonuria

5C50.12 Tyrosinaemia type 2

5C50.20 Histidinaemia

5C50.A0 Argininosuccinic aciduria

5C50.A3 Citrullinaemia

5V5W\_DER Classical homocystinuria

5C50.D0 Maple-syrup-urine disease

5V5X\_DER Methylmalonic aciduria

5V5Y\_DER Propionic aciduria

5V5Z\_DER Multiple carboxylase deficiency

5V60\_DER Glutaric aciduria type 1

5C50.F0 Prolidase deficiency

5C52.03 Sjögren-Larsson syndrome

5V61\_DER Mevalonate kinase deficiency

5V62\_DER Dorfman-Chanarin disease

5V63\_DER Xanthoma in association with lipid storage diseases

5V64\_DER Hyperphosphataemic familial tumoural calcinosis

5C55.01 Lesch-Nyhan syndrome

5V65\_DER Gaucher disease type 1

5V66\_DER Gaucher disease type 2

5C56.01 Fabry disease

5V67\_DER Mucosulfatidosis

5V68\_DER Niemann-Pick disease type A

5V69\_DER Mucolipidosis type 2

5V6A\_DER Aspartylglucosaminuria

5C56.3 Mucopolysaccharidosis

5V6B\_DER Refsum disease

5V6C\_DER Arthrogryposis - renal dysfunction - cholestasis

5C58.10 Porphyria cutanea tarda

5V6D\_DER Hepatoerythropoietic porphyria

5C58.13 Variegate porphyria

5V6E\_DER Hereditary coproporphyria

Erythropoietic porphyrias (BlockL1‑5V6)

5V6F\_DER Erythropoietic protoporphyria

5V6G\_DER Congenital erythropoietic porphyria

5V6Z\_DER Other specified eythropoietic porphyria

5V6H\_DER Alpha-1 antitrypsin deficiency panniculitis

5V6J\_DER Hartnup syndrome

5V6K\_DER Renal tubulo-interstitial disorders in cystinosis

5C64.00 Wilson disease

5V6L\_DER Menkes disease

5V6M\_DER Hereditary haemochromatosis

5V6N\_DER Neonatal haemochromatosis

5C64.20 Acrodermatitis enteropathica

Zinc deficiency syndromes (BlockL1‑5V6)

5V6P\_DER Acquired zinc deficiency syndrome

5V6Q\_DER Neonatal nutritional zinc deficiency

5V6Z\_DER Other specified zinc deficiency syndromes

5V6Z\_DER Zinc deficiency syndrome of unspecified type

5V6R\_DER Fluid overload with oedema

5C80.00 Primary hypercholesterolaemia

5V6S\_DER Xanthoma in association with primary hypercholesterolaemia

5V6T\_DER Xanthoma in association with primary hypertriglyceridaemia

5V6U\_DER Xanthoma in association with primary chylomicronaemia

5V6V\_DER Xanthoma in association with primary combined hyperlipidaemia

5V6W\_DER Xanthoma in association with secondary hyperlipidaemia

5C81.0 Hypoalphalipoproteinaemia

5C81.1 Hypobetalipoproteinaemia

5V6X\_DER Primary localised cutaneous amyloidosis

5V6Y\_DER Cerebral non-neuropathic heredofamilial amyloidosis angiopathy

5V6Z\_DER Cutaneous amyloidosis

CHAPTER 06 Mental, behavioural or neurodevelopmental disorders

6B21 Body dysmorphic disorder

6B22 Olfactory reference disorder

6B25 Body-focused repetitive behaviour disorders

Self-inflicted hair-damaging disorder (BlockL2‑6B2)

6B25.0 Trichotillomania

6V70\_DER Trichoteiromania

6V71\_DER Trichotemnomania

6V7Z\_DER Other specified self-inflicted hair-damaging disorder

6V72\_DER Trichophagia

6V73\_DER Factitious skin disorder imposed on another

6E8Y\_DER Other specified mental, behavioural or neurodevelopmental disorders

6E8Z Mental, behavioural or neurodevelopmental disorders, unspecified

CHAPTER 08 Diseases of the nervous system

8B4Z Spinal cord disorders excluding trauma

8V74\_DER Melkersson syndrome

8C21 Hereditary sensory or autonomic neuropathy

8E7Y\_DER Other specified diseases of the nervous system

8E7Z Diseases of the nervous system, unspecified

CHAPTER 09 Diseases of the visual system

9V75\_DER Disorders of the visual organs

9E1Y\_DER Other specified diseases of the visual system

9E1Z Diseases of the visual system, unspecified

CHAPTER 10 Diseases of the ear or mastoid process

AA6Z Diseases of external ear

AC0Y\_DER Other specified diseases of the ear or mastoid process

AC0Z Diseases of the ear or mastoid process, unspecified

CHAPTER 11 Diseases of the circulatory system

BV76\_DER Woolly hair – palmoplantar keratoderma – dilated cardiomyopathy

BD5Z Diseases of arteries or arterioles

BD70 Superficial thrombophlebitis

BD71.4 Lower limb deep vein thrombosis

Chronic peripheral venous insufficiency of lower extremities (BlockL1‑BD7)

BD74.0 Uncomplicated lower limb venous hypertension

BD74.1 Lower limb varicose veins

BD74.2 Lipodermatosclerosis

BD74.3 Venous leg ulcer

BD74.Z Chronic peripheral venous insufficiency of lower extremities, unspecified

BD75 Venous varicosities of sites other than lower extremity

BD92.1 Cutaneous lymphangiectasia

Lymphoedema (BlockL1‑BD9)

BD93.0 Primary lymphoedema

BD93.1 Secondary lymphoedema

BD93.Y\_DER Other specified forms of lymphoedema

BD93.Z Lymphoedema, unspecified

BE1B Lymphoedema due to surgery or radiotherapy

BE2Y\_DER Other specified diseases of the circulatory system

BE2Z Diseases of the circulatory system, unspecified

CHAPTER 12 Diseases of the respiratory system

CB40 Certain diseases of the respiratory system

CB7Z Diseases of the respiratory system, unspecified

CHAPTER 13 Diseases of the digestive system

Disorders of lips (BlockL1‑DA0)

DA00.0 Cheilitis

Self-induced lip trauma (BlockL2‑DV7)

DV77\_DER Lip-biting

DV7Z\_DER Other specified self-induced lip trauma

DV7Z\_DER Self-induced lip trauma, unspecified

Pigmentary abnormalities of lips (BlockL2‑DV7)

DV78\_DER Labial melanin incontinence

DV79\_DER Labial melanotic macule

DA00.3 Lip fissure

DV7A\_DER Hypertrophy of lip

DV7B\_DER Ectropion of lip

DA00.Y\_DER Other specified disorders of lips

Disorders of oral mucosa (BlockL1‑DA0)

Disturbances of oral epithelium (BlockL2‑DA0)

DA01.01 Hairy leukoplakia

DA01.02 Wandering rash of the mouth

DA01.0Y\_DER Other specified disturbances of oral epithelium

Noninfectious erosive or ulcerative disorders of oral mucosa (BlockL2‑DV7)

Oral aphthae or aphtha-like ulceration (BlockL3‑DV7)

DV7C\_DER Recurrent aphthous stomatitis

DV7D\_DER Periodic fever - aphthous stomatitis - pharyngitis - adenopathy

DV7E\_DER Oral aphthae

DV7Z\_DER Other specified oral aphthae or aphtha-like ulceration

DA01.11 Oral mucositis

DA01.12 Chronic ulcerative stomatitis

DA01.13 Erythema multiforme with oral ulceration

DA01.15 Mouth ulcers

Oral ulceration due to immunobullous disease (BlockL3‑DV7)

DV7F\_DER Oral mucosal involvement by immunobullous disorder classified elsewhere

DV7Z\_DER Other specified oral ulceration due to immunobullous disease

DV7Z\_DER Oral ulceration due to immunobullous disease, unspecified

Oral ulceration due to physical injury (BlockL3‑DV7)

DV7G\_DER Mechanical oral ulceration

Thermal oral ulceration (BlockL4‑DV7)

DV7H\_DER Cold burn of oral mucosa

DV7J\_DER Heat burn of oral mucosa

DV7Z\_DER Other specified thermal oral ulceration

DV7Z\_DER Thermal oral ulceration, unspecified

DV7K\_DER Radiation burn of oral mucosa

DV7L\_DER Chemical burn of oral mucosa

DV7Z\_DER Other specified oral ulceration due to physical injury

DV7Z\_DER Oral ulceration due to physical injury, unspecified

DA01.1Y\_DER Other specified noninfectious erosive or ulcerative disorders of oral mucosa

Pigmentary disturbances of oral mucosa (BlockL2‑DV7)

DV7M\_DER Oral melanin incontinence

DV7N\_DER Melanotic macule of oral mucosa

DV7Z\_DER Other specified pigmentary disturbances of oral mucosa

Granuloma or granuloma-like lesions of oral mucosa (BlockL2‑DV7)

DV7P\_DER Lobular capillary haemangioma of oral mucosa

DV7Q\_DER Traumatic ulcerative granuloma with stromal eosinophilia

DV7R\_DER Verruciform xanthoma of oral mucosa

DA01.2Y\_DER Other specified granuloma or granuloma-like lesions of oral mucosa

Oral mucosal disorders attributable to mechanical damage (BlockL2‑DA0)

DA01.4 Irritative hyperplasia of oral mucosa

DA0Z\_DER Other specified oral mucosal disorders attributable to mechanical damage

DA01.Y\_DER Other specified disorders of oral mucosa

Miscellaneous specified disorders of lips or oral mucosa (BlockL1‑DA0)

DA02.1 Xerostomia

DA02.2 Oral submucous fibrosis

DA02.3 Contact gingivostomatitis

Diseases of tongue (BlockL1‑DA0)

DA03.5 Macroglossia

DA03.Y\_DER Other specified diseases of tongue

DA09.61 Periapical abscess with sinus

DV7S\_DER Drug-induced gingival hyperplasia

DA0F.0 Burning mouth syndrome

DV7T\_DER Oesophageal varices without bleeding in alcoholic liver disease

DV7U\_DER Oesophageal varices without bleeding in hepatic fibrosis

DV7V\_DER Oesophageal varices without bleeding in primary biliary cholangiopathy

DV7W\_DER Oesophageal varices without bleeding in toxic liver disease

DV7X\_DER Noninfective enteritis or ulcer due to graft-versus-host disease

DV7Y\_DER Perianal threadworm dermatosis

DB9Z Diseases of liver

DV7Z\_DER Crohn disease of lips or oral mucosa

DD70.4 Crohn disease of anal region

DE2Y\_DER Other specified diseases of the digestive system

DE2Z Diseases of the digestive system, unspecified

CHAPTER 14 Diseases of the skin

Certain skin disorders attributable to infection or infestation (BlockL1‑EA0)

Certain skin disorders attributable to viral infection (BlockL2‑EA0)

Viral exanthems (BlockL3‑EA0)

EA00 Viral exanthem due to unknown or unspecified agent

EA0Y\_DER Viral exanthem due to other specified virus

Certain dermatoses with suspected viral aetiology (BlockL3‑EA1)

EA10 Pityriasis rosea

EA11 Papular purpuric gloves and socks syndrome

EA12 Infantile papular acrodermatitis

Dermatoses from distant or systemic viral infection (BlockL3‑EA2)

EA20 Necrolytic acral erythema

EA3Z Unspecified skin disorder attributable to viral infection

Certain skin disorders attributable to bacterial infection (BlockL2‑EA4)

Predominantly tropical or subtropical bacterial infections affecting skin (BlockL3‑EA4)

EA40 Tropical phagedaenic ulcer

Certain mycobacterial infections affecting the skin (BlockL3‑EV8)

Skin complications of BCG immunisation (BlockL4‑EV8)

EV80\_DER BCG granuloma

EV81\_DER Tuberculids

EV8Z\_DER Other specified certain mycobacterial infections affecting the skin

Toxin-mediated cutaneous reactions to distant or systemic bacterial infection (BlockL3‑EA5)

Streptococcal toxin-mediated reactions involving skin (BlockL4‑EA5)

EA50.0 Erythema marginatum rheumaticum

EA50.1 Streptococcal toxin-mediated perineal erythema

Staphylococcal toxin-mediated reactions involving skin (BlockL4‑EA5)

EA50.2 Staphylococcal scalded skin syndrome

EA50.3 Staphylococcal scarlatina

EV82\_DER Staphylococcal toxin-mediated recalcitrant erythema and desquamation

EV83\_DER Staphylococcal toxin-mediated perineal erythema

EV8Z\_DER Other specified staphylococcal toxin-mediated reactions involving skin

EA50.Y\_DER Other specified toxin-mediated cutaneous reactions to distant or systemic bacterial infection

Reactive dermatoses due to distant bacterial infection (BlockL3‑EV8)

EV84\_DER Erythema multiforme provoked by other specified bacterial infection

EV8Z\_DER Other specified reactive dermatoses due to distant bacterial infection

EV8Z\_DER Reactive dermatoses due to distant bacterial infection, unspecified

EV85\_DER Skin infection classified elsewhere due to abnormally virulent or therapy resistant bacteria

EA5Y\_DER Cutaneous involvement by other specified bacterial infection

EA5Z Cutaneous involvement by unspecified bacterial infection

Certain skin disorders attributable to fungal infection (BlockL2‑EV8)

Dermatoses resulting from distant or systemic fungal infection (BlockL3‑EV8)

EV86\_DER Dermatophytide

EV8Z\_DER Dermatoses resulting from distant or systemic fungal infection, unspecified

EA60.Y\_DER Skin involvement in other specified fungal infection

EA60.Z Fungal infection of the skin, unspecified

EA6Y\_DER Cutaneous involvement by other specified infection or infestation

Inflammatory dermatoses (BlockL1‑EA8)

Dermatitis and eczema (BlockL2‑EA8)

EA80 Atopic eczema

Seborrhoeic dermatitis and related conditions (BlockL3‑EA8)

EA81.0 Seborrhoeic dermatitis of face

EA81.1 Seborrhoeic dermatitis of the scalp

EV87\_DER Seborrhoeic dermatitis of the flexures

EV88\_DER Generalised seborrhoeic dermatitis

EV89\_DER Seborrhoeic dermatitis of unspecified location

EA81.Y\_DER Seborrhoeic dermatosis of other specified type or distribution

EA81.Z Seborrhoeic dermatitis, unspecified

EA82 Nummular dermatitis

Lichen simplex or lichenification (BlockL3‑EV8)

Lichen simplex (BlockL4‑EV8)

EV8A\_DER Lichen simplex of nape of neck

EA83.00 Lichen simplex of vulva

EA83.01 Lichen simplex of male genitalia

EA83.02 Perianal lichen simplex

EA83.0Y\_DER Lichen simplex of other specified site

EA83.0Z Lichen simplex of unspecified site

EA83.1 Secondary lichenification

EA84 Asteatotic eczema

EA85 Dermatitis or eczema of hands and feet

EA86 Dermatitis and eczema of lower legs

EA87 Dermatitis or eczema of anogenital region

EV8B\_DER Dermatitis or eczema of certain specified sites

Miscellaneous specified eczematous dermatoses (BlockL3‑EA8)

EA88.0 Infectious dermatitis

EA88.1 Post traumatic eczema

EA88.2 Disseminated secondary eczema

EA88.3 Secondary eczema

EA88.4 Pityriasis alba

EA8Y\_DER Other specified eczematous dermatosis

EA8Z Dermatitis or eczema, unspecified

Papulosquamous dermatoses (BlockL2‑EA9)

EA90 Psoriasis

Psoriasis of specified site or distribution (BlockL4‑EA9)

EA90.50 Scalp psoriasis

EV8C\_DER Seborrhoeic psoriasis

EV8D\_DER Non-pustular psoriasis of the palms or soles

EA90.51 Nail psoriasis

EA90.52 Flexural and intertriginous psoriasis

EA90.53 Anogenital psoriasis

EA90.5Y\_DER Psoriasis of other specified site or distribution

Miscellaneous specified forms of psoriasis (BlockL4‑EV8)

EV8E\_DER Kobner psoriasis

EV8F\_DER Drug-exacerbated psoriasis

EV8G\_DER Photoaggravated psoriasis

EV8H\_DER Follicular psoriasis

EV8Z\_DER Other specified miscellaneous specified forms of psoriasis

EV8Z\_DER Miscellaneous specified forms of psoriasis, unspecified

EA91 Lichen planus

Lichen planus and lichenoid reactions of oral mucosa (BlockL4‑EA9)

EA91.40 Non-erosive lichen planus of oral mucosa

EA91.41 Erosive oral lichen planus

EA91.42 Oral lichen planus, unspecified

EV8J\_DER Drug-induced oral lichenoid reaction

EV8K\_DER Amalgam-associated oral lichenoid reaction

EA91.43 Lichenoid mucositis

EA91.4Y\_DER Other specified lichenoid reactions of oral mucosa

Miscellaneous specified forms of lichen planus (BlockL4‑EV8)

EV8L\_DER Annular lichen planus

EV8M\_DER Atrophic lichen planus

EV8N\_DER Bullous lichen planus

EV8P\_DER Flexural lichen planus

EV8Q\_DER Lichen planus pigmentosus

EV8R\_DER Ulcerative lichen planus

EV8S\_DER Linear lichen planus

EV8T\_DER Lichen planus of lips

EV8U\_DER Actinic lichen planus

EV8Z\_DER Other specified miscellaneous specified forms of lichen planus

EV8Z\_DER Miscellaneous specified forms of lichen planus, unspecified

Lichenoid dermatoses (BlockL3‑EV8)

EV8V\_DER Lichen nitidus

EV8W\_DER Lichen striatus

EV8X\_DER Keratosis lichenoides chronica

EV9Z\_DER Other specified lichenoid dermatoses

EA93 Pityriasis lichenoides

EA94 Pityriasis rubra pilaris

EA95 Small plaque parapsoriasis

Urticaria, angioedema and other urticarial disorders (BlockL2‑EB0)

EB00 Spontaneous urticaria

Inducible urticaria and angioedema (BlockL3‑EB0)

EB01.0 Dermographism

EB01.1 Cold urticaria

EV8Y\_DER Heat contact urticaria

EB01.2 Delayed pressure urticaria

EV8Z\_DER Vibratory angioedema

EV90\_DER Solar urticaria

EB01.3 Contact urticaria

EB01.Y\_DER Other specified forms of inducible urticaria and angioedema

Cholinergic urticaria and related conditions (BlockL3‑EB0)

EB02.0 Cholinergic urticaria

EV91\_DER Cholinergic pruritus

EV92\_DER Cholinergic erythema

EB02.Y\_DER Other conditions mediated by cholinergic activation

Syndromes with urticarial reactions or angioedema (BlockL3‑EV9)

EV93\_DER Schnitzler syndrome

EV94\_DER Episodic angioedema with eosinophilia

EV9Z\_DER Other specified syndromes with urticarial reactions or angioedema

EV9Z\_DER Syndromes with urticarial reactions or angioedema, unspecified

Miscellaneous urticarial disorders (BlockL3‑EV9)

EV95\_DER Aquagenic urticaria

EB04 Idiopathic angioedema

EB05 Urticaria of unspecified type

EB0Z\_DER Other specified urticarial disorders

EB0Y\_DER Other specified urticarial disorders

Inflammatory erythemas and other reactive inflammatory dermatoses (BlockL2‑EV9)

Diffuse inflammatory erythemas (BlockL3‑EV9)

EV96\_DER Erythroderma

EV97\_DER Toxic erythema

EV98\_DER Papuloerythroderma of Ofuji

EV9Z\_DER Other specified diffuse inflammatory erythemas

EB11 Annular erythema

EB12 Erythema multiforme

Erythema multiforme classified by aetiology (BlockL4‑EV9)

EV99\_DER Erythema multiforme provoked by viral infection

EV9A\_DER Erythema multiforme provoked by Mycoplasma pneumoniae

EV9B\_DER Erythema multiforme provoked by systemic fungal infection

Stevens-Johnson syndrome or toxic epidermal necrolysis (BlockL3‑EB1)

EB13.0 Stevens-Johnson syndrome

EB13.1 Toxic epidermal necrolysis

EB13.2 Stevens-Johnson and toxic epidermal necrolysis overlap syndrome

Neutrophilic dermatoses (BlockL3‑EB2)

EB20 Acute febrile neutrophilic dermatosis

EV9C\_DER Subcorneal pustular dermatosis

EB21 Pyoderma gangrenosum

Pyoderma gangrenosum classified by aetiological associations (BlockL5‑EV9)

EV9D\_DER Pyoderma gangrenosum associated with inflammatory bowel disease

EV9E\_DER Pyoderma gangrenosum linked to haematological disorder

EV9F\_DER Pyoderma gangrenosum associated with inflammatory polyarthropathy

EV9G\_DER Pyoderma gangrenosum linked to certain specified identified triggers

EV9H\_DER Pyoderma gangrenosum without identifiable trigger

EV9Z\_DER Other specified pyoderma gangrenosum classified by aetiological associations

EV9Z\_DER Pyoderma gangrenosum classified by aetiological associations, unspecified

EV9J\_DER Rheumatoid neutrophilic dermatosis

EV9K\_DER Neutrophilic eccrine hidradenitis

Pustular vasculitic reactions (BlockL4‑EV9)

EV9L\_DER Bowel-associated dermatosis-arthritis syndrome

EV9M\_DER Pustular vasculitis of undetermined cause

EV9Z\_DER Other specified pustular vasculitic reactions

EV9Z\_DER Pustular vasculitic reactions, unspecified

EV9N\_DER Amicrobial pustulosis of the folds

EB2Y\_DER Other specified neutrophilic dermatoses

EB30 Eosinophilic cellulitis

EB31 Erythema nodosum

Immunobullous diseases of the skin (BlockL2‑EB4)

EB40 Pemphigus

EB41 Pemphigoid

EB42 Linear IgA bullous dermatosis

EB43 Epidermolysis bullosa acquisita

EV9P\_DER Immunobullous systemic lupus erythematosus

EB44 Dermatitis herpetiformis

EV9Q\_DER Immunobullous disorder of undefined type

EB4Y\_DER Other specified immunobullous disorder

Cutaneous lupus erythematosus (BlockL2‑EB5)

EB50 Subacute cutaneous lupus erythematosus

EB51 Chronic cutaneous lupus erythematosus

EB5Z Cutaneous lupus erythematosus of unspecified type

Scarring or sclerosing inflammatory dermatoses (BlockL2‑EB6)

EB60 Lichen sclerosus

EB61 Morphoea

EB7Y\_DER Other specified inflammatory dermatoses

Metabolic and nutritional disorders affecting the skin (BlockL1‑EB9)

EB90 Dermatoses resulting from disturbed metabolic processes

Porphyria or pseudoporphyria affecting the skin (BlockL3‑EV9)

Pseudoporphyria (BlockL4‑EV9)

EV9R\_DER Drug-induced pseudoporphyria

EV9S\_DER Pseudoporphyria due to excessive exposure to UV radiation

EV9T\_DER Haemodialysis-associated pseudoporphyria

EB90.3Y\_DER Other specified porphyria or pseudoporphyria affecting the skin

Calcification of skin or subcutaneous tissue (BlockL3‑EV9)

Dystrophic calcification of the skin due to inflammatory disease (BlockL4‑EV9)

EV9U\_DER Calcinosis cutis due to systemic sclerosis

EV9Z\_DER Calcinosis cutis secondary to other specified inflammatory process

EV9V\_DER Dystrophic calcification of the skin due to localised skin injury

EV9W\_DER Calcification of skin and subcutis due to metabolic disturbance

EB90.40 Dystrophic calcification of the skin of uncertain or unspecified aetiology

EB90.4Y\_DER Other specified calcification of skin or subcutaneous tissue

EB9Y\_DER Other specified metabolic and nutritional disorders affecting the skin

Genetic and developmental disorders affecting the skin (BlockL1‑EV9)

Genetic syndromes affecting the skin (BlockL2‑EV9)

Genetic syndromes with poikiloderma (BlockL3‑EV9)

EV9X\_DER Poikiloderma with neutropaenia

EV9Y\_DER Hereditary sclerosing poikiloderma

EV9Z\_DER Congenital or hereditary poikiloderma of uncertain or unspecified type

EW0Z\_DER Other specified genetic syndromes with poikiloderma

EW00\_DER Miscellaneous specified genetic syndromes affecting the skin

EC1Y\_DER Other specified genetic syndromes affecting the skin

Genetic disorders of keratinisation (BlockL2‑EC2)

Hereditary ichthyosis (BlockL3‑EC2)

Non-syndromic ichthyosis (BlockL4‑EC2)

EC20.00 Ichthyosis vulgaris

EC20.01 X-linked ichthyosis

EC20.02 Autosomal recessive congenital ichthyosis

Keratinopathic ichthyoses (BlockL5‑EW0)

EW01\_DER Epidermolytic ichthyosis

EW02\_DER Superficial epidermolytic ichthyosis

EW03\_DER Annular epidermolytic ichthyosis

EW04\_DER Autosomal recessive epidermolytic ichthyosis

EW05\_DER Naevoid epidermolytic ichthyosis

EW06\_DER Ichthyosis Curth-Macklin

EW0Z\_DER Other specified keratinopathic ichthyoses

EW07\_DER Erythrokeratoderma variabilis et progressiva

EW08\_DER Keratosis linearis – ichthyosis congenita – keratoderma

EW09\_DER Congenital reticular ichthyosiform erythroderma

EC20.0Y\_DER Other specified or unclassifiable non-syndromic ichthyosis

EC20.1 Hereditary skin peeling

Hereditary acantholytic dermatoses (BlockL3‑EW0)

EW0A\_DER Darier disease

EW0B\_DER Hailey-Hailey disease

Hereditary palmoplantar keratodermas (BlockL3‑EW0)

Diffuse palmoplantar keratodermas (BlockL4‑EW0)

Non-syndromic diffuse palmoplantar keratodermas (BlockL5‑EW0)

EW0C\_DER Diffuse epidermolytic palmoplantar keratoderma

EW0D\_DER Diffuse non-epidermolytic palmoplantar keratoderma

EW0E\_DER Mal de Meleda

EW0F\_DER Nagashima-type palmoplantar keratoderma

EW0G\_DER Keratolytic winter erythema

EW0Z\_DER Other specified non-syndromic diffuse palmoplantar keratodermas

Syndromic diffuse palmoplantar keratodermas (BlockL5‑EW0)

EW0H\_DER Papillon-Lefèvre syndrome

EW0J\_DER Congenital palmoplantar or perioral keratoderma

EW0K\_DER Curly hair – acral keratoderma – caries syndrome

EW0L\_DER Palmoplantar keratoderma with scleroatrophy of the extremities

EW0M\_DER Connexin palmoplantar keratoderma with sensorineural deafness

EW0N\_DER Loricrin palmoplantar keratoderma with ichthyosis

EW0P\_DER Palmoplantar keratoderma – digital clubbing – acro-osteolysis

EW0Z\_DER Other specified syndromic diffuse palmoplantar keratodermas

??? Diffuse palmoplantar keratoderma

Focal palmoplantar keratodermas (BlockL4‑EW0)

EW0Q\_DER Non-syndromic nummular or linear palmoplantar keratodermas

EW0R\_DER Syndromic nummular or linear palmoplantar keratodermas

EW0Z\_DER Focal palmoplantar keratoderma

Papular palmoplantar keratodermas (BlockL4‑EW0)

EW0S\_DER Punctate palmoplantar keratoderma

EW0T\_DER Marginal papular palmoplantar keratoderma

EC20.3Z Hereditary palmoplantar keratoderma of unspecified type

EW0U\_DER Certain specified inherited disorders of keratinization

EC20.Y\_DER Other specified genetic disorders of keratinisation

Genetic defects of hair or hair growth (BlockL2‑EW0)

Genetic defects of the hair shaft (BlockL3‑EW0)

Genetic defects of the hair shaft with increased fragility (BlockL4‑EW0)

EW0V\_DER Monilethrix

EW0W\_DER Pili torti

EW0X\_DER Loose anagen syndrome

EW1Z\_DER Other specified genetic defects of the hair shaft with increased fragility

EW1Z\_DER Genetic defects of the hair shaft with increased fragility, unspecified

Genetic defects of the hair shaft without increased fragility (BlockL4‑EW0)

EW0Y\_DER Uncombable hair syndrome

EW0Z\_DER Woolly hair

EW10\_DER Pseudomonilethrix

EW11\_DER Pili annulati

EW1Z\_DER Other specified genetic defects of the hair shaft without increased fragility

EW1Z\_DER Genetic defects of the hair shaft without increased fragility, unspecified

Genetic abnormalities of hair colour (BlockL4‑EW1)

EW12\_DER Familial premature canities

EW13\_DER Familial poliosis

EW1Z\_DER Other specified genetic abnormalities of hair colour

EW1Z\_DER Genetic abnormalities of hair colour, unspecified

??? Other specified genetic defects of the hair shaft

??? Genetic defects of the hair shaft, unspecified

EW14\_DER Björnstad syndrome

EW15\_DER Trichorrhexis nodosa syndrome

EW16\_DER Skin fragility-woolly hair syndrome

Hereditary alopecia or hypotrichosis (BlockL3‑EW1)

EW17\_DER Atrichia congenita

EW18\_DER Atrichia with keratin cysts

EW19\_DER Hereditary generalised hypotrichosis

EW1A\_DER Hereditary hypotrichosis of scalp

EW1B\_DER Marie Unna congenital hypotrichosis

EW1C\_DER Ciliary or superciliary hypotrichosis

EC21.4 Genetically-determined hypertrichosis

EC21.Y\_DER Other specified genetic defects of hair or hair growth

Genetic defects of nails or nail growth (BlockL2‑EC2)

EC22.0 Inherited deformities of nails

Genetic disorders of skin pigmentation (BlockL2‑EW1)

Non-syndromic genetically-determined hypermelanosis or lentiginosis (BlockL3‑EW1)

EW1D\_DER Familial progressive hyperpigmentation

EW1E\_DER Familial generalised lentiginosis

EW1F\_DER Inherited patterned lentiginosis

EW1G\_DER Centrofacial lentiginosis

EW1H\_DER Familial multiple café-au-lait macules

EW1J\_DER Dowling-Degos disease

EW1K\_DER Hypermelanotic pigmentary mosaicism

EW1Z\_DER Other specified non-syndromic genetically-determined hypermelanosis or lentiginosis

EW1Z\_DER Non-syndromic genetically-determined hypermelanosis or lentiginosis, unspecified

Syndromic genetically-determined hypermelanosis or lentiginosis (BlockL3‑EW1)

EW1L\_DER Gastrocutaneous syndrome

EW1M\_DER Watson syndrome

EW1Z\_DER Other specified syndromic genetically-determined hypermelanosis or lentiginosis

EW1Z\_DER Syndromic genetically-determined hypermelanosis or lentiginosis, unspecified

EW1N\_DER Genetically-determined mixed hyper- and hypomelanotic disorders of skin pigmentation

Albinism or other specified genetically-determined hypomelanotic disorders (BlockL3‑EC2)

EC23.20 Oculocutaneous albinism

Hypomelanotic pigmentary mosaicism (BlockL4‑EW1)

EW1P\_DER Hypomelanosis of Ito

EW1Q\_DER Phylloid hypomelanosis

EW1Z\_DER Other specified hypomelanotic pigmentary mosaicism

EW1Z\_DER Hypomelanotic pigmentary mosaicism, unspecified

EW1R\_DER Partial albinism

EW1S\_DER Piebaldism

EW1T\_DER Waardenburg syndrome

EW1U\_DER Tietz hypomelanosis – deafness syndrome

EC23.2Y\_DER Other specified genetically-determined hypomelanotic disorders

EC23.Y\_DER Other specified genetic disorders of skin pigmentation

Genetically-determined epidermolysis bullosa (BlockL2‑EW1)

Epidermolysis bullosa simplex (BlockL3‑EW1)

Suprabasal epidermolysis bullosa simplex (BlockL4‑EW1)

EW1V\_DER Acantholytic epidermolysis bullosa

EW1W\_DER Epidermolysis bullosa simplex superficialis

EW1X\_DER Ectodermal dysplasia – skin fragility syndrome

Basal epidermolysis bullosa simplex (BlockL4‑EW1)

Autosomal dominant basal epidermolysis bullosa simplex (BlockL5‑EW1)

EW1Y\_DER Epidermolysis bullosa simplex, localised

EW1Z\_DER Epidermolysis bullosa simplex with mottled pigmentation

EW20\_DER Epidermolysis bullosa simplex, Ogna

EW21\_DER Epidermolysis bullosa simplex, generalised severe

EW22\_DER Epidermolysis bullosa simplex, generalised intermediate

Autosomal recessive basal epidermolysis bullosa simplex (BlockL5‑EW2)

EW23\_DER Epidermolysis bullosa simplex, migratory circinate

EW24\_DER Epidermolysis bullosa simplex with muscular dystrophy

EW25\_DER Epidermolysis bullosa simplex with pyloric atresia

EW26\_DER Autosomal recessive epidermolysis bullosa simplex due to BP230 deficiency

EW27\_DER Epidermolysis bullosa simplex, autosomal recessive due to exophilin 5 deficiency

EW2Z\_DER Other specified autosomal recessive basal epidermolysis bullosa simplex

Junctional epidermolysis bullosa (BlockL3‑EW2)

EW28\_DER Junctional epidermolysis bullosa, generalised severe

EW29\_DER Junctional epidermolysis bullosa, generalised intermediate type

EW2A\_DER Junctional epidermolysis bullosa with pyloric atresia

EW2B\_DER Junctional epidermolysis bullosa, late onset

EW2C\_DER Junctional epidermolysis bullosa, localised

EW2D\_DER Junctional epidermolysis bullosa inversa

EW2E\_DER Junctional epidermolysis bullosa, LOC

EW2Z\_DER Other specified junctional epidermolysis bullosa

EW2Z\_DER Junctional epidermolysis bullosa, unspecified

EC32 Dystrophic epidermolysis bullosa

EC3Z Epidermolysis bullosa

Genetic disorders affecting dermal collagen, elastin or other matrix proteins (BlockL2‑EW2)

EW2F\_DER Buschke-Ollendorff syndrome

EW2G\_DER Dermochondrocorneal dystrophy

EC40 Pseudoxanthoma elasticum

EC4Y\_DER Other specified genetic disorders affecting dermal matrix proteins

Certain genetically-determined metabolic disorders with skin involvement (BlockL2‑EW2)

EW2H\_DER Sphingolipidoses with skin manifestations

Specified developmental anomalies affecting the skin (BlockL2‑EC5)

EC50 Developmental anomalies of the umbilicus

EC5Y\_DER Other specified developmental anomalies affecting the skin

EC7Y\_DER Other specified genetic and developmental disorders affecting the skin

Sensory and psychological disorders affecting the skin (BlockL1‑EC9)

Disturbances of cutaneous sensation (BlockL2‑EC9)

Pruritus (BlockL3‑EC9)

EC90.0 Pruritus due to skin disorder

Pruritus due to systemic disorder (BlockL4‑EC9)

EC90.10 Uraemic pruritus

EC90.11 Cholestatic pruritus

EW2J\_DER Pruritus due to endocrine or metabolic disorder

EW2K\_DER Pruritus due to myeloproliferative or lymphoproliferative disorder

EC90.12 Haemodialysis-associated pruritus

EC90.1Y\_DER Pruritus due to other specified systemic disorder

EC90.2 Drug-induced pruritus

Pruritus due to neurological disorder (BlockL4‑EW2)

EW2L\_DER Brachioradial pruritus

EW2M\_DER Notalgia paraesthetica

EW2N\_DER Postherpetic pruritus

EW2P\_DER Pruritus due to central nervous system injury or degeneration

EW2Q\_DER Pruritus due to nerve damage or compression

EC90.4 Psychogenic pruritus

EW2R\_DER Pruritus of multifactorial aetiology

EW2S\_DER Aquagenic pruritus

EW2T\_DER Scalp pruritus

EC90.5 Anogenital pruritus

EC90.6 Pruritus of unknown cause

EC90.Y\_DER Pruritus of other specified type or aetiology

EC90.Z Pruritus, unspecified

Prurigo (BlockL3‑EW2)

EW2U\_DER Papular prurigo

EC91.0 Nodular prurigo

EC91.1 Atopic prurigo

EW2V\_DER Prurigo of undetermined type

EC91.Z Prurigo, unspecified

Mucocutaneous or cutaneous pain syndromes (BlockL3‑EC9)

EC92.0 Penoscrotodynia

EC92.1 Scalp dysaesthesia

EC9Y\_DER Other specified disturbances of cutaneous sensation

Psychological or psychiatric conditions affecting the skin (BlockL2‑ED0)

Self-inflicted skin disorders (BlockL3‑ED0)

ED00 Artefactual skin disorder

ED01 Simulated skin disease

ED02 Painful bruising syndrome

ED0Y\_DER Other specified self-inflicted skin disorders

ED2Y\_DER Other specified psychological or psychiatric conditions affecting the skin

Neurological conditions affecting the skin (BlockL2‑EW2)

Neuropathic skin damage (BlockL3‑EW2)

EW2W\_DER Skin damage resulting from congenital or hereditary neuropathy

EW2X\_DER Skin damage resulting from acquired neuropathy

ED30.0 Neuropathic skin ulceration

ED30.Y\_DER Other specified neuropathic skin damage

EW2Y\_DER Spinal cord defect affecting the skin

Miscellaneous neurological conditions affecting the skin (BlockL3‑ED3)

ED31 Burning feet syndrome

ED3Z\_DER Other specified miscellaneous neurological conditions affecting the skin

ED3Z\_DER Miscellaneous neurological conditions affecting the skin, unspecified

ED3Y\_DER Cutaneous involvement in other specified neurological condition

Skin disorders involving specific cutaneous structures (BlockL1‑ED5)

Disorders of the epidermis and epidermal appendages (BlockL2‑ED5)

Disorders of epidermal keratinisation (BlockL3‑ED5)

Ichthyoses (BlockL4‑ED5)

ED50.0 Acquired ichthyosis

ED50.Z Ichthyosis of unspecified type

Diffuse epidermal hyperkeratosis and acanthosis (BlockL4‑ED5)

ED51.0 Acanthosis nigricans

EW2Z\_DER Confluent and reticulated papillomatosis

EW30\_DER Pityriasis rotunda

ED51.Y\_DER Other specified hyperkeratotic and acanthotic dermatoses

Follicular digitate keratoses (BlockL4‑ED5)

ED56 Keratosis pilaris

EW31\_DER Lichen spinulosus

EW32\_DER Keratosis circumscripta

EW33\_DER Trichodysplasia spinulosa

EW3Z\_DER Other specified follicular digitate keratoses

Non-follicular punctate or digitate keratoses (BlockL4‑EW3)

EW34\_DER Flegel disease

EW35\_DER Multiple minute digitate keratoses

EW3Z\_DER Other specified non-follicular punctate or digitate keratoses

EW3Z\_DER Non-follicular punctate or digitate keratoses, unspecified

ED52 Porokeratoses

Acantholytic dermatoses (BlockL4‑EW3)

EW36\_DER Transient acantholytic dermatosis

Skin peeling (BlockL4‑EW3)

EW37\_DER Keratolysis exfoliativa

Xerosis cutis or asteatosis (BlockL4‑EW3)

EW38\_DER Atopic xeroderma

EW39\_DER Asymptomatic xerosis cutis

EW3A\_DER Pruritic xerosis cutis

EW3B\_DER Asteatosis cutis

EW3C\_DER Xerosis senilis

Palmoplantar keratodermas (BlockL4‑ED5)

ED55.0 Acquired palmoplantar keratodermas

ED55.Z Palmoplantar keratoderma, unspecified

ED5Y\_DER Other specified disorders of epidermal keratinisation

Disorders of skin colour (BlockL3‑ED6)

Acquired hypermelanosis (BlockL4‑ED6)

ED60.0 Physiological hypermelanosis

ED60.1 Melasma

EW3D\_DER Hypermelanosis due to endocrine disorder

ED60.2 Postinflammatory hypermelanosis

EW3E\_DER Paraneoplastic hypermelanosis

EW3F\_DER Ashy dermatosis

EW3G\_DER Post traumatic hypermelanosis

ED60.Y\_DER Hypermelanosis of other specified aetiology

ED60.Z Hypermelanosis of unspecified aetiology

Acquired melanotic macules or lentigines (BlockL4‑ED6)

ED61.0 Freckles

ED61.1 Mucosal melanosis

EW3H\_DER Generalised eruptive lentiginosis

ED61.Y\_DER Other specified acquired melanotic macules or lentigines

ED62 Endogenous non-melanin pigmentation

EW3J\_DER Non-melanin pigmentation due to ingested or injected substance

EW3K\_DER Non-melanin pigmentation due to exogenous substances

Acquired hypomelanotic disorders (BlockL4‑ED6)

ED63.0 Vitiligo

Syndromic acquired hypomelanosis (BlockL5‑EW3)

EW3L\_DER Alezzandrini syndrome

EW3Z\_DER Other specified syndromic acquired hypomelanosis

EW3Z\_DER Syndromic acquired hypomelanosis, unspecified

ED63.1 Hypomelanosis due to exposure to chemicals

ED63.2 Postinflammatory hypomelanosis

EW3M\_DER Postinfective hypomelanosis

EW3N\_DER Idiopathic guttate hypomelanosis

EW3P\_DER Leukoderma acquisitum centrifugum

EW3Q\_DER Other specified form of acquired hypomelanosis

EW3R\_DER Punctate leukoderma

ED63.Y\_DER Acquired hypomelanosis due to other specified disorder

ED63.Z Acquired hypomelanosis of unknown or unspecified aetiology

Acquired mixed hyper- and hypomelanotic disorders of skin pigmentation (BlockL4‑EW3)

EW3S\_DER Dyschromatosis due to chronic arsenic toxicity

EW3Z\_DER Other specified acquired mixed hyper- and hypomelanotic disorders of skin pigmentation

EW3Z\_DER Acquired mixed hyper- and hypomelanotic disorders of skin pigmentation, unspecified

ED64 Abnormal skin pigmentation

ED6Y\_DER Other specified disorders of skin pigmentation

Disorders of hair (BlockL3‑ED7)

Alopecia or hair loss (BlockL4‑ED7)

ED70.0 Male pattern hair loss

ED70.1 Female pattern hair loss

ED70.2 Alopecia areata

ED70.3 Telogen effluvium

ED70.4 Anagen effluvium

ED70.5 Scarring alopecia

Scarring alopecia due to inflammatory disorders (BlockL6‑EW3)

EW3T\_DER Pseudopelade

ED70.50 Folliculitis decalvans

ED70.51 Dissecting cellulitis

ED7Z\_DER Scarring alopecia due to other specified inflammatory disorder

ED70.Y\_DER Other specified alopecia or hair loss

ED70.Z Alopecia, unspecified

Hirsutism and syndromes with hirsutism (BlockL4‑ED7)

ED72.0 Constitutional hirsutism

ED72.1 Hirsutism associated with hyperandrogenaemia

ED72.Z Hirsutism, unspecified

Acquired disorders of the hair shaft (BlockL4‑ED7)

ED73.0 Weathered hair

EW3U\_DER Trichorrhexis nodosa

EW3V\_DER Pili bifurcati

EW3W\_DER Pili multigemini

ED73.1 Acquired changes in hair colour

ED73.Y\_DER Other specified acquired disorders of the hair shaft

ED7Y\_DER Other specified disorders of hair

Disorders of the hair follicle (BlockL3‑ED8)

Acne and related disorders (BlockL4‑ED8)

ED80 Acne

ED81 Acneform inflammatory disorders

Rosacea and related disorders (BlockL4‑ED9)

ED90.0 Rosacea

ED90.1 Periorificial dermatitis

EW3X\_DER Acne agminata

EW3Y\_DER Idiopathic facial aseptic granuloma

EW3Z\_DER Corticosteroid-induced rosacea

ED90.Y\_DER Other specified rosacea-like disorders

Disorders of the sebaceous gland (BlockL4‑ED9)

ED91.0 Heterotopic sebaceous glands

ED91.1 Sebaceous gland hyperplasia

ED91.2 Seborrhoea

Disorders involving the apocrine follicular unit (BlockL4‑ED9)

ED92.0 Hidradenitis suppurativa

Miscellaneous disorders involving the hair follicle (BlockL4‑EW4)

EW40\_DER Pseudofolliculitis

EW41\_DER Pseudofolliculitis barbae

EW42\_DER Classical adult eosinophilic pustular folliculitis

EW43\_DER Immunosuppression-associated eosinophilic pustular folliculitis

EW44\_DER Eosinophilic pustular folliculitis

EW45\_DER Disseminate or recurrent infundibulofolliculitis

ED9Y\_DER Other specified disorders involving the hair follicle

Disorders of eccrine sweat glands or sweating (BlockL3‑EE0)

Hyperhidrosis (BlockL4‑EE0)

EE00.0 Localised hyperhidrosis

EE00.1 Primary generalised hyperhidrosis

EE00.Z Hyperhidrosis, unspecified

Hypohidrosis (BlockL4‑EW4)

Hypohidrosis attributable to defective sudomotor innervation or function (BlockL5‑EW4)

Primary autonomic disorders with acquired anhidrosis (BlockL6‑EW4)

Acquired idiopathic generalised anhidrosis (BlockL7‑EW4)

EW46\_DER Idiopathic pure sudomotor failure

EW4Z\_DER Other specified acquired idiopathic generalised anhidrosis

EW4Z\_DER Acquired idiopathic generalised anhidrosis, unspecified

EW47\_DER Progressive isolated segmental anhidrosis

EW48\_DER Ross syndrome

EW4Z\_DER Other specified primary autonomic disorders with acquired anhidrosis

EW49\_DER Hypohidrosis due to acquired neurological injury or disorder

EW4A\_DER Neuropathic hypohidrosis

EE01.1 Hypohidrosis due to genetic abnormalities of eccrine gland structure or function

EE01.2 Hypohidrosis of undetermined aetiology

EE01.Y\_DER Other specified forms of hypohidrosis

EE01.Z Hypohidrosis, unspecified

EE02 Miliaria

EE0Y\_DER Other specified disorders of eccrine sweat glands or sweating

Disorders of the nail or perionychium (BlockL3‑EE1)

Acquired deformities of the nail plate (BlockL4‑EE1)

EE10.0 Abnormality of nail shape

EE10.1 Abnormality of nail surface

EE10.2 Onycholysis

EW4B\_DER Nail pterygium

EE10.3 Nail hypertrophy

EE10.4 Nail atrophy

EE10.5 Nail dystrophy, not otherwise specified

EE10.Y\_DER Other specified acquired deformities of the nail plate

Acquired abnormalities of nail colour (BlockL4‑EW4)

EW4C\_DER Acquired leukonychia

EE11.0 Melanonychia

EW4D\_DER Longitudinal erythronychia

EE11.1 Yellow nail syndrome

EE11.Y\_DER Other abnormalities of nail colour

Infections of the nail or perionychium (BlockL4‑EE1)

EE12.0 Acute bacterial paronychia

EW4E\_DER Pseudomonas infection of nail

EE12.1 Onychomycosis

Onychomycosis classified according to clinical pattern (BlockL6‑EW4)

EW4F\_DER Superficial white onychomycosis

EW4G\_DER Distal and lateral subungual onychomycosis

EW4H\_DER Proximal subungual onychomycosis

EW4J\_DER Total dystrophic onychomycosis

EW4Z\_DER Other specified onychomycosis classified according to clinical pattern

EW4Z\_DER Onychomycosis classified according to clinical pattern, unspecified

EE12.Y\_DER Other specified infections of the nail or perionychium

Certain disorders affecting the nails or perionychium (BlockL4‑EE1)

EE13.0 Nail fragility

EE13.1 Ingrowing nail

EE13.4 Nail disorder associated with specified systemic disease

EW4K\_DER Painful dorsolateral fissure of the fingertip

EW4L\_DER Subungual exostosis

EE13.5 Eczematous nail dystrophy

EE13.Y\_DER Other specified nail disorder

EE1Y\_DER Other specified disorders of the nail or perionychium

EE1Z Disorders of the nail or perionychium, unspecified

EW4M\_DER Disorders of epidermal integrity

Disorders of the dermis and subcutis (BlockL2‑EW4)

Disorders of cutaneous connective tissue (BlockL3‑EW4)

Atrophy or degeneration of dermal or subcutaneous connective tissue (BlockL4‑EW4)

EW4N\_DER Intrinsic ageing of the skin

EW4P\_DER Premature ageing of the skin

EE40.0 Corticosteroid-induced skin atrophy

EE40.1 Stretch marks

EE40.2 Atrophic scarring of the skin

EE40.3 Skin fragility

EE40.Y\_DER Other specified atrophy or degeneration of dermal or subcutaneous connective tissue

EE40.Z Atrophy or degeneration of dermal or subcutaneous connective tissue, unspecified type

Abnormalities of dermal elastin (BlockL4‑EW4)

EW4Q\_DER Acquired pseudoxanthoma elasticum

EE41.0 Cutis laxa

EE41.1 Anetoderma

EW4R\_DER Dermal elastolysis

EW4S\_DER Papular elastorrhexis

EW4T\_DER Elastoderma

EE41.Y\_DER Other specified dermatoses characterised by abnormal dermal elastin

EW4U\_DER Atrophoderma

EW4V\_DER Poikiloderma

Fibromatoses and keloids (BlockL4‑EE6)

Keloid or hypertrophic scars (BlockL5‑EE6)

EE60.0 Keloid

EE60.1 Hypertrophic scar

EW4W\_DER Keloidal acne

EW4X\_DER Folliculitis keloidalis

EE60.Y\_DER Other specified keloidal disorders

Superficial fibromatoses (BlockL5‑EW4)

EW4Y\_DER Pachydermodactyly

Camptodactyly or streblodactyly (BlockL6‑EW4)

EW4Z\_DER Familial camptodactyly

EW50\_DER Sporadic camptodactyly

EW51\_DER Streblodactyly

EW5Z\_DER Other specified camptodactyly or streblodactyly

EW5Z\_DER Camptodactyly or streblodactyly, unspecified

EW5Z\_DER Other specified superficial fibromatoses

EW52\_DER Fibrous or myofibroblastic proliferations in infants or children

EW53\_DER Infantile stiff skin syndromes

EE6Y\_DER Other specified fibromatous disorders of skin and soft tissue

Perforating dermatoses (BlockL4‑EW5)

EW54\_DER Familial reactive perforating collagenosis

EW55\_DER Elastosis perforans serpiginosa

EE70.0 Acquired perforating dermatosis

EW56\_DER Perforating periumbilical calcific elastosis

EW57\_DER Perforating folliculitis

EE70.Y\_DER Other specified perforating dermatoses

Scleroedema and miscellaneous scleroderma-like conditions (BlockL4‑EW5)

EW58\_DER Scleroedema

EW59\_DER Sclerodactyly

Miscellaneous conditions affecting dermal connective tissue (BlockL4‑EW5)

EW5A\_DER Ainhum

EW5B\_DER Cutis verticis gyrata

EE7Y\_DER Other specified disorders of cutaneous connective tissue

Histiocytic-granulomatous disorders of the skin (BlockL3‑EE8)

EE80.0 Granuloma annulare

EE80.1 Necrobiosis lipoidica

EW5C\_DER Cutaneous or mucocutaneous Crohn disease

EW5D\_DER Miscellaneous specified granulomatous disorders of skin

Dermal dendrocyte, Class IIa histiocytoses (BlockL4‑EW5)

EW5E\_DER Benign cephalic histiocytosis

EW5F\_DER Generalised eruptive histiocytosis

EW5G\_DER Papular xanthoma

EW5H\_DER Progressive nodular histiocytosis

EW5J\_DER Xanthoma disseminatum

EW5K\_DER Diffuse plane xanthomatosis

EW5L\_DER Miscellaneous specified, Class IIb non-Langerhans cell histiocytoses involving the skin

EE8Y\_DER Other specified histiocytic and granulomatous disorders of the skin

Benign dermal lymphocytic or lymphoplasmacytic infiltrations or proliferations (BlockL3‑EE9)

EE90 Benign lymphocytic infiltration of the skin

EE91 Lymphocytoma cutis

Disorders of subcutaneous fat (BlockL3‑EF0)

Panniculitis (BlockL4‑EF0)

EF00.0 Pancreatic enzyme panniculitis

EW5M\_DER Lipoatrophic panniculitis

EW5N\_DER Cold panniculitis

EW5P\_DER Artefactual panniculitis

EF00.Y\_DER Panniculitis of other specified aetiology

EF00.Z Panniculitis of undetermined or unspecified etiology

Lipoatrophy or lipodystrophy (BlockL4‑EF0)

EF01.1 Localised lipoatrophy and lipodystrophy

EF01.Y\_DER Other specified forms of lipodystrophy and lipoatrophy

EF01.Z Lipodystrophy of unspecified type

EF02 Certain noninflammatory disorders of subcutaneous fat

Miscellaneous disorders of subcutaneous fat (BlockL5‑EF0)

EF02.3 Cellulite

EF0Y\_DER Other specified disorders of subcutaneous fat

Disorders of cutaneous blood and lymphatic vessels (BlockL2‑EF2)

Malformations involving cutaneous blood vessels (BlockL3‑EF2)

Acquired malformations of cutaneous blood vessels (BlockL4‑EF2)

Discrete acquired superficial venous or capillary ectasias (BlockL5‑EF2)

EF20.0 Venous lake

EF20.1 Angiokeratoma

EF20.2 Lower limb venous telangiectases

EW5Q\_DER Diffuse acquired telangiectasia

EF20.Y\_DER Other specified acquired malformations of cutaneous blood vessels

EF20.Z Acquired malformations of cutaneous blood vessels, unspecified

EF2Z Cutaneous vascular malformation, unspecified

Purpura or bruising (BlockL3‑EF3)

EF30 Purpura or bruising due to disorders of coagulation

EW5R\_DER Orthostatic purpura

EW5S\_DER Purpura due to paroxysmal coughing or vomiting

EW5T\_DER Purpura or bruising due to friction or trauma

EF3Y\_DER Other specified purpura

EF3Z Purpura of unspecified aetiology

Vasculitis or capillaritis involving the skin (BlockL3‑EF4)

EF40.0 Capillaritis

Vasculitis affecting small cutaneous blood vessels (BlockL4‑EF4)

EF40.10 Urticarial vasculitis

EF40.1Y\_DER Other specified vasculitis affecting small cutaneous blood vessels

Localised cutaneous vasculitis (BlockL4‑EW5)

EW5U\_DER Erythema induratum

EF40.20 Granuloma faciale

EW5V\_DER Erythema elevatum diutinum

EF40.2Y\_DER Other specified localised cutaneous vasculitis

Vasculitis affecting medium or large cutaneous blood vessels (BlockL4‑EW5)

EW5W\_DER Polyarteritis nodosa involving the skin

EW6Z\_DER Other specified vasculitis affecting medium or large cutaneous blood vessels

EW6Z\_DER Vasculitis affecting medium or large cutaneous blood vessels, unspecified

EW5X\_DER Lymphocytic cutaneous vasculitis

EF40.Z Cutaneous vasculitis unspecified

Dermatoses attributable to hyperviscosity or microvascular occlusion (BlockL3‑EW5)

EW5Y\_DER Cutaneous microvascular occlusion by platelet plugs

EW5Z\_DER Cutaneous microvascular occlusion by emboli

EW60\_DER Cutaneous microvascular disturbances due to vascular coagulopathies

EW61\_DER Cutaneous microvascular occlusion due to systemic coagulopathies

EW62\_DER Cutaneous microvascular disturbances due to dysproteinaemia

EF5Y\_DER Other specified dermatoses attributable to hyperviscosity or microvascular occlusion

Dermatoses resulting from vascular insufficiency (BlockL3‑EF7)

EF70 Lower limb venous eczema

EF9Y\_DER Other specified dermatoses resulting from vascular insufficiency

Functional vascular disorders of the skin (BlockL3‑EW6)

Vasodilatation of extremities (BlockL4‑EW6)

EW63\_DER Drug-induced erythromelalgia

EW6Z\_DER Other specified forms of vasodilatation of extremities

Vasoconstriction of extremities (BlockL4‑EW6)

EW64\_DER Ergotism

EW6Z\_DER Other specified forms of vasoconstriction of extremities

EW65\_DER Exaggerated physiological flushing

EW66\_DER Emotional flushing

EW67\_DER Drug-induced flushing

EW68\_DER Gustatory flushing

Skin disorders involving certain specific body regions (BlockL1‑EG3)

Skin disorders involving the head and neck (BlockL2‑EG3)

Skin disorders localised to the scalp (BlockL3‑EG3)

EG30.0 Scalp folliculitis

EG30.1 Erosive pustular dermatosis of scalp

EG30.2 Pityriasis amiantacea

EG30.Y\_DER Other specified scalp disorders not elsewhere classifiable

Disorders of the external ear involving the skin (BlockL3‑EG4)

Inflammatory disorders of the external ear (BlockL4‑EG4)

EG40 Contact dermatitis of external ear

EG4Y\_DER Other specified inflammatory disorder of external ear

EG4Z Otitis externa

Skin disorders involving the genital and perianal regions (BlockL2‑EG6)

Dermatoses of the anus, perianal area or perineum (BlockL3‑EG6)

EG60 Anal pruritus

EG61 Infections of the anus or perianal skin

EG63 Sacrococcygeal pilonidal disease

EG7Y\_DER Other specified skin disorders involving the genital and perianal regions

EG9Y\_DER Skin disorders involving other specific body regions

EG9Z Skin disorders involving certain specific body regions, unspecified

Skin disorders associated with pregnancy, the neonatal period and infancy (BlockL1‑EW6)

Skin disorders specific to the perinatal or neonatal period (BlockL2‑EW6)

Neonatal skin infection (BlockL3‑EW6)

Neonatal pyogenic skin infections (BlockL4‑EW6)

EW69\_DER Neonatal staphylococcal pyoderma

EW6A\_DER Neonatal streptococcal pyoderma

EW6B\_DER Neonatal pseudomonas skin infection

EW6Z\_DER Other specified neonatal pyogenic skin infections

Neonatal fungal infections involving the skin (BlockL4‑EW6)

EW6C\_DER Neonatal mucocutaneous candidosis

EW6Z\_DER Neonatal infection due to other fungus classified elsewhere

EH1Z Neonatal skin infection, unspecified

EH3Y\_DER Other specified skin disorders specific to the perinatal or neonatal period

Dermatoses of infancy (BlockL2‑EH4)

EH40.0 Infantile seborrhoeic dermatitis

EW6D\_DER Infantile pustular psoriasis

Infantile napkin dermatoses (BlockL3‑EH4)

EH40.10 Primary irritant napkin dermatitis

EH40.1Y\_DER Other specified infantile napkin dermatoses

Erythrodermas of infancy (BlockL3‑EW6)

EW6E\_DER Congenital erythroderma of uncertain nature

EW6Z\_DER Other specified erythrodermas of infancy

EW6Z\_DER Erythrodermas of infancy, unspecified

Miscellaneous skin conditions arising predominantly in infancy (BlockL3‑EH4)

EH40.3 Acute haemorrhagic oedema of infancy

EW6F\_DER Infantile acropustulosis

EW6G\_DER Infantile eosinophilic pustular folliculitis

EW6H\_DER Infantile pedal papules

EW6Z\_DER Other specified miscellaneous skin conditions arising predominantly in infancy

EW6Z\_DER Miscellaneous skin conditions arising predominantly in infancy, unspecified

EH40.Y\_DER Other specified dermatoses of infancy

Adverse cutaneous reactions to medication (BlockL1‑EH6)

Drug eruptions (BlockL2‑EH6)

EH60 Exanthematic drug eruption

Drug-induced urticaria, angioedema and anaphylaxis (BlockL3‑EH6)

EH61.0 Drug-induced urticaria

EH61.1 Drug-induced angioedema

EH62 Lichenoid drug eruption

EH63 Stevens-Johnson syndrome and toxic epidermal necrolysis due to drug

EW6J\_DER Eczematous drug eruption

EH64 Drug-induced erythroderma

EH65 DRESS syndrome

EH66 Fixed drug eruption

EW6K\_DER Pseudolymphomatous drug hypersensitivity syndrome

EH67 Acne or acneform reactions attributable to drugs

EW6L\_DER Miscellaneous specified cutaneous eruptions due to drug

EH6Y\_DER Drug eruption of other specified type

EH6Z Drug eruption of unspecified type

EH70 Pigmentary abnormalities of skin due to drug

EH71 Dermatoses precipitated by drug therapy

EH72 Drug-induced hair abnormalities

EH73 Drug-induced nail abnormalities

EH74 Drug-induced oral conditions

EH75 Photosensitivity due to drug

Dermatoses associated with specific classes of medication (BlockL2‑EW6)

Dermatoses resulting from cytotoxic or cancer chemotherapy (BlockL3‑EW6)

EW6M\_DER Cytotoxic chemotherapy-induced recall reaction

EW6N\_DER Drug-related adverse cutaneous reactions to epidermal growth factor receptor inhibitor therapy

EW6P\_DER Eccrine syringosquamous metaplasia

EW6Q\_DER Palmoplantar erythrodysaesthesia due to cytotoxic chemotherapy

EW6Z\_DER Other specified dermatoses resulting from cytotoxic or cancer chemotherapy

EW6Z\_DER Dermatoses resulting from cytotoxic or cancer chemotherapy, unspecified

Dermatoses resulting from immunosuppressive therapy (BlockL3‑EW6)

EW6R\_DER Immunosuppressive therapy as contributor to skin infection classified elsewhere

EW6S\_DER Immunosuppressive therapy as contributor to malignant skin neoplasm classified elsewhere

EW6T\_DER Immunosuppressive therapy as contributor to dermatosis classified elsewhere

EW6Z\_DER Other specified dermatoses resulting from immunosuppressive therapy

EW6Z\_DER Dermatoses resulting from immunosuppressive therapy, unspecified

Dermatoses attributable to corticosteroid therapy (BlockL3‑EW6)

EW6U\_DER Corticosteroid-induced acne

EW6Z\_DER Other specified dermatoses attributable to corticosteroid therapy

EW6Z\_DER Dermatoses attributable to corticosteroid therapy, unspecified

EW6V\_DER Dermatoses associated with other specific medications

EH76.Y\_DER Other dermatoses associated with specific classes of medication

Localised adverse cutaneous reactions to administration of drug (BlockL2‑EW6)

EW6W\_DER BCG ulcer

EH78 Adverse cutaneous reactions to herbal, homoeopathic or other alternative therapies

EH7Y\_DER Other specified adverse cutaneous reactions to medication

EH7Z Unspecified adverse cutaneous reactons to medication

Skin disorders provoked by external factors (BlockL1‑EW6)

EW6X\_DER Dermatoses provoked by physical or environmental factors

Miscellaneous specified dermatoses provoked by pressure (BlockL3‑EW6)

EW6Y\_DER Piezogenic pedal papules

EW7Z\_DER Other specified pressure-induced skin disorder

Dermatoses due to foreign bodies (BlockL3‑EW7)

Tattoos or tattoo reactions (BlockL4‑EW7)

EW73\_DER Decorative tattoo

EW74\_DER Iatrogenic tattoo

EW75\_DER Traumatic tattoo

EW76\_DER Occupational tattoo

EW77\_DER Tattoo reaction

EW78\_DER Tattoo infection

EW7Z\_DER Other specified complications of tattoos

Foreign body reaction to inorganic matter in the skin (BlockL4‑EW7)

EW79\_DER Silica granuloma

EW7A\_DER Talc granuloma

EW7B\_DER Cutaneous suture granuloma

EW7C\_DER Oleogranuloma

EW7Z\_DER foreign body reaction to other specified inorganic matter in the skin

EH93.2 Foreign body reaction to organic matter in the skin

EH93.3 Foreign body granuloma of skin

EH93.Y\_DER Other specified reaction to foreign body in the skin

Dermatoses provoked or exacerbated by exposure to cold (BlockL3‑EW7)

EW7D\_DER Skin or soft tissue injury due to exposure to cold

Abnormal vascular reactivity to cold (BlockL4‑EW7)

EW7E\_DER Acrocyanosis

EW7F\_DER Erythrocyanosis

EW7G\_DER Physiological cutis marmorata

EW7Z\_DER Other specified abnormal vascular reactivity to cold

EW7Z\_DER Abnormal vascular reactivity to cold, unspecified

EJ0Y\_DER Other specified dermatoses provoked or exacerbated by exposure to cold

Dermatoses provoked by heat or electricity (BlockL3‑EJ1)

EJ10 Erythema ab igne

EJ1Y\_DER Other specified dermatoses provoked by heat or electricity

Dermatoses provoked by light or UV radiation (BlockL3‑EJ2)

Chronic effects of ultraviolet radiation on the skin (BlockL4‑EJ2)

EJ20 Photoaging of the skin

EW7H\_DER Photodistributed telangiectasia

EJ2Y\_DER Other specified chronic effects of ultraviolet radiation on the skin

Autoimmune or other photodermatoses (BlockL4‑EJ3)

EJ30.0 Polymorphic light eruption

EW7J\_DER Actinic prurigo

EW7K\_DER Hydroa vacciniforme

EJ30.1 Chronic actinic dermatitis

EW7L\_DER Actinic folliculitis

EJ30.Y\_DER Other specified photodermatoses

Photoaggravated skin disease (BlockL4‑EW7)

EW7M\_DER Atopic eczema, photoaggravated

Acute effects of ultraviolet radiation on normal skin (BlockL4‑EJ4)

EJ40 Sunburn

EJ41 Burn from exposure to artificial source of ultraviolet radiation

EJ4Z Acute effects of ultraviolet radiation on normal skin, unspecified

EJ6Y\_DER Other specified dermatoses provoked by light or UV radiation

Dermatoses due to ionizing radiation (BlockL3‑EW7)

Acute effects of ionizing radiation on the skin (BlockL4‑EW7)

EW7N\_DER Acute radiodermatitis, not elsewhere classified

EW7Z\_DER Other specified acute effects of ionizing radiation on the skin

EW7Z\_DER Acute effects of ionizing radiation on the skin, unspecified

Chronic effects of ionizing radiation on the skin (BlockL4‑EW7)

EW7P\_DER Chronic radiodermatitis, not elsewhere classified

EW7Z\_DER Other specified chronic effects of ionizing radiation on the skin

EW7Z\_DER Chronic effects of ionizing radiation on the skin, unspecified

EJ7Z Dermatoses due to ionizing radiation, unspecified

Dermatoses resulting from exposure to water (BlockL3‑EW7)

EW7Q\_DER Aquagenic wrinkling of the fingers or palms

Dermatitis due to exogenous factors (BlockL2‑EK0)

Allergic contact dermatitis (BlockL3‑EK0)

Allergic contact dermatitis organised by allergen class (BlockL4‑EK0)

EK00.0 Allergic contact dermatitis due to clothing or footwear

EK00.1 Allergic contact dermatitis due to cosmetics or fragrances

EK00.2 Allergic contact dermatitis due to dental materials

EK00.3 Allergic contact dermatitis due to food flavours or additives

EK00.4 Allergic contact dermatitis due to hairdressing products

EK00.5 Allergic contact dermatitis due to industrial biocides, cutting oils or disinfectants

EK00.6 Allergic contact dermatitis due to metals or metal salts

EK00.7 Allergic contact dermatitis due to allergenic haptens derived from plants or organic matter

EK00.8 Allergic contact dermatitis due to plastics, glues or resin systems

EK00.9 Allergic contact dermatitis due to preservatives or biocides

EK00.A Allergic contact dermatitis due to rubber chemicals

EK00.B Allergic contact dermatitis due to systemic medicaments

EK00.C Allergic contact dermatitis due to topical medicaments

EW7R\_DER Allergic contact eczema due to low molecular weight allergens not elsewhere classifiable

EW7Z\_DER Allergic contact dermatitis due to other specified allergens not elsewhere classifiable

EW7Z\_DER Allergic contact dermatitis due to unknown or unspecified allergen

Allergic contact dermatitis organised by site (BlockL4‑EW7)

EW7S\_DER Allergic contact dermatitis of scalp

EW7T\_DER Allergic contact cheilitis

EW7U\_DER Allergic contact dermatitis of face or neck

EW7V\_DER Allergic contact dermatitis of hands

EW7W\_DER Allergic contact dermatitis of male genitalia

EW7X\_DER Allergic contact dermatitis of female genitalia

EW8Z\_DER Other specified allergic contact dermatitis organised by site

EK00.Y\_DER Other specified allergic contact dermatitis

EK00.Z Allergic contact dermatitis, unspecified

Photo-allergic contact dermatitis (BlockL3‑EW7)

Photo-allergic contact dermatitis organised by photo-allergen class (BlockL4‑EW7)

EW7Y\_DER Photo-allergic contact dermatitis due to fragrances

EW7Z\_DER Photo-allergic contact dermatitis due to sunscreens

EW80\_DER Photo-allergic contact dermatitis due to plants or vegetable matter

EW81\_DER Photo-allergic contact dermatitis due to preservatives or biocides

EW82\_DER Photo-allergic contact dermatitis due to nonsteroidal anti-inflammatory drugs

EW83\_DER Photo-allergic contact dermatitis due to medicaments other than NSAIDs

EW8Z\_DER Photo-allergic contact dermatitis due to other specified low molecular weight photoallergens

EW84\_DER Occupational photo-allergic contact dermatitis

Irritant contact dermatitis (BlockL3‑EK0)

Irritant contact dermatitis from specified external agents (BlockL4‑EK0)

EK02.00 Irritant contact dermatitis due to wet work

EK02.01 Irritant contact dermatitis due to solvents

EK02.02 Irritant contact dermatitis due to exposure to acids, alkalis or other specified chemical irritants

EK02.03 Irritant contact dermatitis due to cosmetics or emollients

EK02.04 Irritant contact dermatitis due to topical medicaments or antimicrobials

EK02.05 Irritant contact dermatitis due to plants or other vegetable matter

EK02.06 Irritant contact dermatitis due to foods

Irritant contact dermatitis of specified site (BlockL4‑EW8)

EW85\_DER Irritant contact dermatitis of scalp

EK02.10 Irritant contact dermatitis of external ear

EK02.11 Irritant contact blepharoconjunctivitis

EW86\_DER Irritant contact cheilitis

EW87\_DER Irritant contact dermatitis of face or neck

EK02.12 Irritant contact dermatitis of hands

EW88\_DER Irritant contact dermatitis of lower leg

EW89\_DER Irritant contact dermatitis of feet

EW8A\_DER Irritant contact dermatitis of male genitalia

EK02.13 Irritant contact dermatitis of vulva

EW8B\_DER Irritant contact dermatitis of perianal skin

EK02.1Y\_DER Irritant contact dermatitis of other specified site

EW8C\_DER Contact dermatitis due to skin damage from friction or micro-trauma

Irritant contact dermatitis due to friction, sweating or contact with body fluids (BlockL4‑EK0)

EK02.20 Intertriginous dermatitis due to friction, sweating or contact with body fluids

EK02.21 Irritant contact dermatitis due to saliva

EK02.22 Irritant contact dermatitis due to incontinence

EK02.23 Irritant contact dermatitis related to stoma or fistula

EK02.24 Irritant contact dermatitis related to skin contact with prostheses or surgical appliances

EW8D\_DER Occupational irritant contact dermatitis

EK02.Y\_DER Irritant contact dermatitis due to other specified cause

EK02.Z Irritant contact dermatitis, unspecified

Contact dermatitis of specified site (BlockL3‑EW8)

EW8E\_DER Contact dermatitis of scalp

EW8F\_DER Contact dermatitis of face or neck

EW8G\_DER Contact dermatitis of lower leg

EW8H\_DER Contact dermatitis of feet

EW8Z\_DER Contact dermatitis of other specified type or location

EW8Z\_DER Contact dermatitis of unspecified type or location

EW8J\_DER Occupational contact dermatitis

EK0Y\_DER Dermatitis due to other specified exogenous factors

EK0Z Contact dermatitis, unspecified

Allergy to substances in contact with the skin (BlockL2‑EK1)

EK10 Allergic contact urticaria

EK11 Protein contact dermatitis

EW8K\_DER Exacerbation of constitutional dermatitis due to exposure to contact allergens

Cutaneous reactions to systemic exposure to contact allergens (BlockL3‑EW8)

EW8L\_DER Systemic contact dermatitis due to ingested allergen

EW8M\_DER Systemic contact dermatitis due to implanted allergen

EK12 Allergic contact sensitisation

EK1Y\_DER Other specified forms of cutaneous allergy

Dermatoses provoked by contact with irritant or noxious substance (BlockL2‑EW8)

EW8N\_DER Skin injury due to exposure to corrosive substances

EW8P\_DER Exacerbation of constitutional dermatitis due to exposure to skin irritants

Phototoxic reactions to skin contact with photoactive agents (BlockL3‑EW8)

EW8Q\_DER Phototoxic reaction to skin contact with plant matter

EK20 Phototoxic reaction to fragrance or cosmetics

EW8R\_DER Phototoxic reaction to skin contact with tar or tar derivatives

EK2Y\_DER Phototoxic reaction to skin contact with other specified photoactive agent

EK2Z Phototoxic dermatitis, unspecified

EK4Y\_DER Other specified dermatoses provoked by contact with irritant or noxious substance

Cutaneous reactions to venomous or noxious animals (BlockL2‑EK5)

Cutaneous insect bite reactions (BlockL3‑EK5)

EK50.00 Papular urticaria

EK50.01 Bullous insect bite reaction

EK50.02 Persistent insect bite reaction

EK50.0Y\_DER Other specified cutaneous insect bite reactions

EK50.0Z Cutaneous insect bite reactions, unspecified

Dermatoses arising through work or occupation (BlockL2‑EW8)

Occupational contact dermatitis, contact urticaria or allergy (BlockL3‑EW8)

EW8S\_DER Occupational contact urticaria

EW8Z\_DER Other specified occupational contact dermatitis, contact urticaria or allergy

EW8Z\_DER Occupational contact dermatitis, contact urticaria or allergy, unspecified

EW8T\_DER Exacerbation of constitutional dermatitis due to occupation

EW8U\_DER Skin injury due to occupational exposure to corrosive substances

EW8V\_DER Occupational phototoxic reactions to skin contact with exogenous photoactive agents

EW8W\_DER Occupational acne or folliculitis

Occupationally-acquired dermatoses due to exposure to cold or heat (BlockL3‑EW8)

EW8X\_DER Occupationally-acquired dermatosis due to exposure to cold

EW8Y\_DER Occupationally-acquired dermatosis due to heat

EW9Z\_DER Other specified occupationally-acquired dermatoses due to exposure to cold or heat

EW9Z\_DER Occupationally-acquired dermatoses due to exposure to cold or heat, unspecified

Occupationally-acquired skin infections or infestations (BlockL3‑EW8)

EW8Z\_DER Occupationally-acquired skin infection, classified elsewhere

EW90\_DER Occupationally-acquired parasitic skin infestation, classified elsewhere

Skin manifestations of work-related poisoning (BlockL3‑EW9)

EW91\_DER Skin disorder resulting from occupational exposure to poison

EW9Z\_DER Other specified skin manifestations of work-related poisoning

EW9Z\_DER Skin manifestations of work-related poisoning, unspecified

Occupationally-acquired disorders of skin pigmentation (BlockL3‑EW9)

EW92\_DER Occupational leukoderma

EW93\_DER Occupational melanosis

EW94\_DER Miscellaneous occupationally-acquired skin disorders

EW9Z\_DER Other specified skin disorder caused or exacerbated by occupation

EK5Y\_DER Other specified skin disorders provoked by external factors

Benign proliferations, neoplasms and cysts of the skin (BlockL1‑EW9)

EW95\_DER Benign adipocytic neoplasms of skin or soft tissue

EK70 Cutaneous cysts

EK71 Skin tags or polyps

Disorders of the skin of uncertain or unpredictable malignant potential (BlockL1‑EK9)

Actinic keratosis and other discrete epidermal dysplasias (BlockL2‑EK9)

EK90.0 Actinic keratosis

EW96\_DER Actinic cheilitis

EK90.1 Diffuse actinic keratinocyte dysplasia

EW97\_DER Photochemotherapy-induced keratosis

EW98\_DER Thermal keratosis

EW99\_DER Chronic radiation keratosis

EW9A\_DER Arsenical keratosis

EK90.Y\_DER Other discrete epidermal dysplasias

Dermatoses which may presage cutaneous lymphoma (BlockL2‑EK9)

EK91.0 Large plaque parapsoriasis

EK91.1 Poikiloderma vasculare atrophicans

EK91.2 Primary cutaneous plasmacytosis

EK92 Histiocytoses of uncertain malignant potential

Cutaneous markers of internal disorders (BlockL1‑EW9)

EW9B\_DER Cutaneous markers of internal malignancy

Miscellaneous specific cutaneous signs of internal malignancy (BlockL3‑EW9)

EW9C\_DER Paraneoplastic pruritus

EW9Z\_DER Other specified miscellaneous specific cutaneous signs of internal malignancy

EW9Z\_DER Miscellaneous specific cutaneous signs of internal malignancy, unspecified

EW9D\_DER Cutaneous markers of endocrine and metabolic disorders

EW9E\_DER Cutaneous markers of disorders of the digestive system

EW9F\_DER Cutaneous markers of disorders of the respiratory system

EW9G\_DER Cutaneous markers of disorders of the kidney or urinary tract

EW9H\_DER Cutaneous disorders attributable to acquired or iatrogenic immunodeficiency

EL3Y\_DER Other specified cutaneous markers of internal disorders

Postprocedural disorders of the skin (BlockL1‑EW9)

Cutaneous complications of surgical, laser or other interventional procedures (BlockL2‑EW9)

Cutaneous complications of surgical procedures (BlockL3‑EW9)

EW9J\_DER Surgical wound of skin

EL50 Unsatisfactory surgical scar of skin

Complications of cutaneous flaps or grafts (BlockL4‑EL5)

EL51 Cutaneous flap necrosis

EL52 Myocutaneous flap necrosis

EL53 Skin graft failure

EL54 Composite graft failure

EL5Z\_DER Other specified complications of cutaneous flaps or grafts

Miscellaneous complications of cutaneous surgical procedures (BlockL4‑EW9)

EW9K\_DER Anatomical distortion of skin or soft tissues following cutaneous surgical procedure

EW9L\_DER Cicatricial contracture of skin or soft tissues following cutaneous surgical procedure

EW9M\_DER Temporary cutaneous sensory disturbance resulting from cutaneous surgical procedure

EW9N\_DER Permanent cutaneous sensory impairment resulting from cutaneous surgical procedure

EW9P\_DER Impairment of motor nerve function resulting from cutaneous surgical procedure

EW9Q\_DER Cutaneous infection specified elsewhere resulting from surgical procedure

EW9Z\_DER Other specified cutaneous complications of surgical procedures

Cutaneous complications of laser surgery (BlockL3‑EW9)

EW9R\_DER Thermal burns resulting from cutaneous laser surgery

EW9S\_DER Infection specified elsewhere following cutaneous laser surgery

EW9T\_DER Dyspigmentation following cutaneous laser surgery

EW9Z\_DER Other specified cutaneous complication of laser surgery

Cutaneous complications of stomas or fistulas (BlockL3‑EW9)

EW9U\_DER Skin problems resulting from tracheostomy

EW9V\_DER Skin problem resulting from external stoma of digestive organs

EW9W\_DER Skin problem resulting from urinary diversion procedure

EW9X\_DER Parastomal ulcer

EX0Z\_DER Other specified cutaneous complications of stomas or fistulas

??? Cutaneous complications of other interventional procedures

Adverse effects of phototherapy (BlockL2‑EW9)

EW9Y\_DER Adverse effects of UVB phototherapy

EW9Z\_DER Adverse effects of psoralen photochemotherapy

EX0Z\_DER Adverse effects of other specified phototherapy

Adverse cutaneous effects of therapeutic ionizing irradiation (BlockL2‑EL6)

EL60 Acute radiodermatitis following radiotherapy

EL61 Chronic radiodermatitis following radiotherapy

EX00\_DER Radionecrosis of skin due to therapeutic ionizing irradiation

EX01\_DER Radiotherapy-induced xerostomia

EX02\_DER Scarring alopecia following radiotherapy

EL62 Radiotherapy-induced skin malignancy

EL6Y\_DER Other specified adverse cutaneous effects of therapeutic ionizing irradiation

EL6Z Adverse cutaneous reaction to radiotherapy

Complications of cutaneous cosmetic procedures (BlockL2‑EL7)

EL70 Adverse reaction to dermal or deep fillers

EL71 Adverse reaction to chemical peel

EL72 Adverse reaction to injection of neurotoxin

EL73 Unsatisfactory outcome from cutaneous cosmetic surgical procedure

EX03\_DER Unsatisfactory outcome from cosmetic laser surgery

Specified cutaneous complications of cosmetic procedures (BlockL3‑EX0)

EX04\_DER Cutaneous infection specified elsewhere following cosmetic procedure

EX05\_DER Hypomelanosis resulting from cosmetic procedure

EX06\_DER Dyspigmentation resulting from cosmetic procedure

EX07\_DER Fibrosis or scarring following cosmetic procedure

EX0Z\_DER Other specified complications of cosmetic procedures

EL7Y\_DER Other specified complications of cutaneous cosmetic procedures

EL80 Adverse cutaneous effects of diagnostic procedures

EX08\_DER Adverse cutaneous reactions to parenteral administration of proteins

EL8Y\_DER Other specified postprocedural disorders of the skin

Absence of skin condition (BlockL1‑EX0)

EX09\_DER No significant skin abnormality found

EX0A\_DER Skin condition resolved

EX0B\_DER Normal skin

EX0C\_DER Normal hair

EX0D\_DER Normal nails

EM0Y\_DER Other specified diseases of the skin

EM0Z Skin disease of unspecified nature

CHAPTER 15 Diseases of the musculoskeletal system or connective tissue

FX0E\_DER Rheumatoid pericarditis

FX0F\_DER Distal interphalangeal psoriatic arthritis

FA21.0 Psoriatic spondyloarthritis

FA25.20 Tophaceous gout

FX0G\_DER Gouty panniculitis

FB33 Secondary disorders of muscle

FB43 Secondary disorders of synovium or tendon

FB51.0 Palmar fascial fibromatosis

FB51.1 Knuckle pads

FX0H\_DER Plantar fascial fibromatosis

FX0J\_DER Nephrogenic systemic fibrosis

FX0K\_DER Fibro-osseous pseudotumour of the digit

FB8Z Osteopathies or chondropathies

FC0Y\_DER Other specified diseases of the musculoskeletal system or connective tissue

FC0Z Diseases of the musculoskeletal system or connective tissue, unspecified

CHAPTER 16 Diseases of the genitourinary system

GA00.3 Genital ulcer of vulva

GA02 Vaginitis

GA30.4 Menopausal hot flush

GA34.02 Vulvodynia

Dermatoses of female genitalia (BlockL1‑GX0)

GX0L\_DER Premalignant or malignant disorders of the vulva

Sensory disturbance of the vulva (BlockL2‑GA4)

GA42.0 Vulval pruritus

Miscellaneous dermatoses of female genitalia (BlockL2‑GX0)

GX0M\_DER Angiokeratoma of the vulva

GX0Z\_DER Other specified miscellaneous dermatoses of female genitalia

GX0Z\_DER Miscellaneous dermatoses of female genitalia, unspecified

GA4Y\_DER Other specified dermatoses of female genitalia

GB0Z Diseases of the male genital system

GC8Y\_DER Other specified diseases of the genitourinary system

GC8Z Diseases of the genitourinary system, unspecified

CHAPTER 18 Pregnancy, childbirth or the puerperium

JA22.1 Gestational oedema without hypertension

JX0Q\_DER Intrahepatic cholestasis of pregnancy

Pregnancy dermatoses (BlockL1‑JA6)

JA65.10 Gestational pemphigoid

JA65.11 Pruritus of pregnancy

JA65.12 Polymorphic eruption of pregnancy

JX0R\_DER Atopic eruption of pregnancy

JA65.1Y\_DER Other specified pregnancy dermatoses

JB6Z Certain obstetric conditions, not elsewhere classified

CHAPTER 19 Certain conditions originating in the perinatal period

Neonatal dermatoses due to maternal antibodies (BlockL1‑KA0)

KA07.0 Neonatal lupus erythematosus

KA07.1 Neonatal pemphigus

KX0S\_DER Neonatal gestational pemphigoid

KX0T\_DER Transient neonatal Behçet disease

KX0U\_DER Transient neonatal epidermolysis bullosa acquisita

KX0V\_DER Neonatal alloimmune thrombocytopenic purpura

KA07.Y\_DER Other specified neonatal dermatoses due to maternal antibodies

KA42 Birth injury to scalp

KA43 Birth injury to skin or soft tissues

KA6Z Infections of the Fetus or newborn

KA88 Disseminated intravascular coagulation of Fetus or newborn

KA8F Neonatal vitamin K deficiency

Inflammatory dermatoses of the newborn (BlockL1‑KX0)

KX0W\_DER Down syndrome leukemoid reaction-associated transient neonatal pustulosis

KC21.0 Neonatal acne

KX0X\_DER Neonatal cephalic pustulosis

KC21.1 Neonatal toxic erythema

KX0Y\_DER Transient neonatal pustular melanosis

KC21.2 Perianal dermatitis of the newborn

KC21.Y\_DER Other specified inflammatory dermatoses of the newborn

Neonatal disorders of subcutaneous fat (BlockL1‑KC2)

KC22.0 Subcutaneous fat necrosis of the newborn

KC22.1 Cold panniculitis of the newborn

KC22.Y\_DER Other specified neonatal disorders of subcutaneous fat

Neonatal nutritional disorders affecting the skin (BlockL1‑KX0)

Neonatal macronutrient deficiency affecting the skin (BlockL2‑KX0)

KX0Z\_DER Neonatal marasmus due to fetal malnutrition

KX1Z\_DER Other specified neonatal macronutrient deficiency affecting the skin

KX1Z\_DER Neonatal macronutrient deficiency affecting the skin, unspecified

Neonatal micronutrient excess affecting the skin (BlockL2‑KX1)

KX10\_DER Neonatal iron overload

KX11\_DER Skin manifestation of other specified neonatal micronutrient excess or toxicity classified elsewhere

KX1Z\_DER Other specified neonatal micronutrient excess affecting the skin

KX1Z\_DER Neonatal micronutrient excess affecting the skin, unspecified

Neonatal micronutrient deficiency affecting the skin (BlockL2‑KX1)

Neonatal vitamin deficiency affecting the skin (BlockL3‑KX1)

KX12\_DER Skin manifestation of other specified neonatal vitamin deficiency

KX1Z\_DER Other specified neonatal vitamin deficiency affecting the skin

KX1Z\_DER Neonatal vitamin deficiency affecting the skin, unspecified

Neonatal mineral deficiency affecting the skin (BlockL3‑KX1)

KX13\_DER Skin manifestation of other specified neonatal mineral deficiency

KX1Z\_DER Other specified neonatal mineral deficiency affecting the skin

KX1Z\_DER Neonatal mineral deficiency affecting the skin, unspecified

??? Other specified neonatal micronutrient deficiency affecting the skin

??? Neonatal micronutrient deficiency affecting the skin, unspecified

??? Other specified neonatal nutritional disorders affecting the skin

Skin disorders associated with prematurity (BlockL1‑KC3)

KC30 Skin fragility of prematurity

KC31 Congenital erosive or vesicular dermatosis healing with reticulated supple scarring

KC3Y\_DER Other specified skin disorders associated with prematurity

Miscellaneous skin disorders in the neonate (BlockL1‑KC4)

KC40.1 Neonatal milia

KX14\_DER Congenital extramedullary dermal haematopoiesis

KC40.Y\_DER Other specified skin disorders in the neonate

Iatrogenic injuries involving the skin of the neonate (BlockL1‑KX1)

Postnatal iatrogenic skin injury (BlockL2‑KX1)

KX15\_DER Neonatal extravasation injury

KX16\_DER Iatrogenic neonatal dystrophic calcification

KX17\_DER Neonatal pressure ulcer

KX18\_DER Neonatal scald injury

KC50 Neonatal phototherapy burn

KX19\_DER Bronze baby syndrome

KX1A\_DER Adverse reaction to substance applied to neonatal skin

KC5Y\_DER Other specified postnatal iatrogenic skin injury

KC5Z Postnatal iatrogenic skin injury, unspecified

KC7Y\_DER Other specified iatrogenic injuries involving the skin of the neonate

KD12.0 Neonatal cold injury syndrome

KD5Z Conditions originating in the perinatal or neonatal period, unspecified

CHAPTER 20 Developmental anomalies

LA14.0 Structural developmental anomalies of eyelids

LA21 Minor anomalies of pinnae

LA31 Structural developmental anomalies of mouth or tongue

Facial clefts (BlockL1‑LX1)

LX1B\_DER Median facial cleft

LX1C\_DER Oblique facial cleft

Lateral facial cleft (BlockL2‑LX1)

LX1D\_DER Commissural facial cleft

LX1Z\_DER Other specified lateral facial cleft

LX1Z\_DER Lateral facial cleft, unspecified

LX1E\_DER Paramedian facial cleft

LA90.00 Hereditary haemorrhagic telangiectasia

LB03.0 Allantoic duct remnants or cysts

LX1F\_DER Umbilical sinus

LX1G\_DER Umbilical vitelline cyst or remnants

LX1H\_DER Subcutaneous vitelline cyst or remnants

LX1J\_DER Genitoperineal median raphe cyst

LB99.8 Split hand

LB9A.6 Split foot

LX1K\_DER Split hand - split foot

LX1L\_DER Camptodactyly of fingers

LX1M\_DER Camptodactyly of toes

LX1N\_DER Developmental hamartomata of the epidermis and epidermal appendages

Developmental anomalies of skin pigmentation (BlockL1‑LC1)

LC10 Dermal melanocytosis

LX1P\_DER Achromic naevus

LC1Y\_DER Other specified developmental anomalies of skin pigmentation

Hamartomata derived from dermal connective tissue (BlockL1‑LX1)

LX1Q\_DER Collagenoma

LX1R\_DER Elastoma

LX1S\_DER Fibrous hamartoma of infancy

LC20 Connective tissue hamartoma

LC2Y\_DER Other specified hamartomata derived from dermal connective tissue

LX1T\_DER Developmental defects of hair or nails

Developmental defects of hair or hair growth (BlockL2‑LX1)

LX1U\_DER Temporal triangular alopecia

LX1V\_DER Congenital sporadic alopecia

LX1W\_DER Naevoid hypertrichosis

LX1X\_DER Developmental disorders of hair colour

LX2Z\_DER Other specified developmental defects of hair or hair growth

Developmental defects of the nail apparatus (BlockL2‑LX1)

LX1Y\_DER Congenital nail hypertrophy

LX1Z\_DER Congenital malalignment of the great toenails

LX20\_DER Congenital hypertrophy of the lateral fold of the hallux

LX2Z\_DER Other specified developmental defects of the nail apparatus

Developmental anomalies of cutaneous vasculature (BlockL1‑LC5)

Developmental capillary vascular malformations of the skin (BlockL2‑LC5)

LC50.0 Salmon patch

LC50.1 Port-wine stain

LX21\_DER Angioma serpiginosum

LX22\_DER Naevus anaemicus

LC50.Y\_DER Other specified cutaneous capillary vascular malformation

LX23\_DER Blue rubber bleb naevus syndrome

LX24\_DER Mucocutaneous venous malformations

LX25\_DER Glomuvenous malformation

LX26\_DER Cutis marmorata telangiectatica congenita

LC5Y\_DER Other specified developmental anomalies of cutaneous vasculature

LC5Z Developmental anomalies of cutaneous vasculature, unspecified

Congenital anomalies of skin development (BlockL1‑LC6)

LC60 Aplasia cutis congenita

LD2Z Multiple developmental anomalies or syndromes

Ectodermal dysplasia syndromes (BlockL2‑LX2)

LX27\_DER Ackerman syndrome

LX28\_DER ADULT syndrome

LX29\_DER Amelo-cerebro-hypohidrotic syndrome

LX2A\_DER Ankyloblepharon - ectodermal defects - cleft lip or palate

LX2B\_DER AREDYLD syndrome

LX2C\_DER Beare-Stevenson cutis gyrata syndrome

LX2D\_DER Blepharocheilodontic syndrome

LX2E\_DER Book syndrome

LX2F\_DER Cardio-facio-cutaneous syndrome

LX2G\_DER Cerebellar ataxia - ectodermal dysplasia

LX2H\_DER Congenital generalised hypertrichosis

LX2J\_DER X-linked dominant congenital generalised hypertrichosis

LX2K\_DER Deafness – enamel hypoplasia – nail defects

LX2L\_DER Anonychia with bizarre flexural pigmentation

LX2M\_DER Anonychia or onychodystrophy – hypoplasia or absence of distal phalanges

LX2N\_DER Autosomal dominant hypodontia with nail dysplasia

LX2P\_DER Amelo-onycho-hypohidrotic syndrome

LX2Q\_DER Deafness – onychodystrophy

LX2R\_DER Odonto-onycho-hypohidrotic dysplasia - midline scalp defects

LX2S\_DER Tricho-odonto-onycho-dermal syndrome

LX2T\_DER Curly hair – ankyloblepharon – nail dysplasia syndrome

LX2U\_DER Tricho-odonto-onychodysplasia - dominant syndactyly

LX2V\_DER Pili torti - onychodysplasia

LD27.00 Incontinentia pigmenti

LX2W\_DER Focal dermal hypoplasia

LX2X\_DER Choroidal atrophy - alopecia

LX2Y\_DER Coffin-Siris syndrome

LX2Z\_DER Conductive deafness - ptosis - skeletal anomalies

LX30\_DER Contractures - ectodermal dysplasia - cleft lip or palate

LX31\_DER Cote-Katsantoni syndrome

LX32\_DER Cranioectodermal dysplasia

LX33\_DER Dahlberg-Borer-Newcomer syndrome

LX34\_DER Dermatoosteolysis, Kirghizian type

LX35\_DER Dubowitz syndrome

LX36\_DER Ectodermal dysplasia - absent dermatoglyphs

LX37\_DER Ectodermal dysplasia - acanthosis nigricans

LX38\_DER Ectodermal dysplasia - arthrogryposis - diabetes mellitus

LX39\_DER Ectodermal dysplasia - blindness

LX3A\_DER Ectodermal dysplasia - cutaneous syndactyly syndrome

LX3B\_DER Ectodermal dysplasia - ectrodactyly - macular dystrophy

LX3C\_DER Ectodermal dysplasia - intellectual deficit - central nervous system malformation

LX3D\_DER Ectodermal dysplasia - sensorineural deafness

LX3E\_DER Ectodermal dysplasia - syndactyly syndrome

LX3F\_DER Ectodermal dysplasia with natal teeth, Turnpenny type

LX3G\_DER Ectodermal dysplasia, pure hair-nail type

LX3H\_DER Ectodermal dysplasia, Berlin type

LX3J\_DER Ectrodactyly - ectodermal dysplasia - cleft lip or palate

LX3K\_DER Ectrodactyly - ectodermal dysplasia without clefting

LX3L\_DER Facial ectodermal dysplasia

LX3M\_DER Focal facial dermal dysplasia

LX3N\_DER GAPO syndrome

LX3P\_DER Gingival fibromatosis - hypertrichosis

LX3Q\_DER Gorlin-Chaudhry-Moss syndrome

LX3R\_DER Hidrotic ectodermal dysplasia, Christianson-Fourie type

LX3S\_DER Hidrotic ectodermal dysplasia, Halal type

LX3T\_DER Hypertrichosis cubiti - short stature

LX3U\_DER Hypoparathyroidism - deafness - renal disease

LX3V\_DER Ichthyosis - alopecia - eclabion - ectropion - intellectual deficit

LX3W\_DER Johanson-Blizzard syndrome

LX3X\_DER Johnson neuroectodermal syndrome

LX3Y\_DER Juvenile macular degeneration - hypotrichosis

LX3Z\_DER Lacrimo-auriculo-dento-digital syndrome

LX40\_DER Limb-mammary syndrome

LX41\_DER Marshall syndrome

LX42\_DER Naegeli-Franceschetti-Jadassohn syndrome

LX43\_DER Oculo-dento-digital dysplasia

LX44\_DER Oculo-osteo-cutaneous syndrome

LX45\_DER Odonto-tricho-ungual-digito-palmar syndrome

LX46\_DER Odontotrichomelic syndrome

LX47\_DER OL-EDA-ID syndrome

LX48\_DER Oligodontia - cancer predisposition syndrome

LX49\_DER Pilodental dysplasia - refractive errors

LX4A\_DER Rapp-Hodgkin syndrome

LX4B\_DER Scalp-ear-nipple syndrome

LX4C\_DER Schinzel-Giedion syndrome

LX4D\_DER Sparse hair - short stature - skin anomalies

LX4E\_DER Stern-Lubinsky-Durrie syndrome

LX4F\_DER Taurodontia - absent teeth - sparse hair

LX4G\_DER Toriello-Lacassie-Droste syndrome

LX4H\_DER Tricho-dento-osseous syndrome

LX4J\_DER Tricho-oculo-dermo-vertebral syndrome

LX4K\_DER Tricho-odonto-onychial dysplasia with bone deficiency in fronto-parietal region

LX4L\_DER Tricho-odonto-onychial ectodermal dysplasia

LX4M\_DER Tricho-retino-dento-digital syndrome

LX4N\_DER Trichodental syndrome

LX4P\_DER Trichodermodysplasia - dental alterations

LX4Q\_DER Trichodysplasia - amelogenesis imperfecta

LX4R\_DER Trichomegaly - cataract - hereditary spherocytosis

LX4S\_DER Trichomegaly - retina pigmentary degeneration - dwarfism

LX4T\_DER Xeroderma - talipes - enamel defects

LX4U\_DER Zlotogora-Ogur syndrome

LX4V\_DER Zunich-Kaye syndrome

LX4W\_DER Dermo-odontodysplasia

LX4X\_DER Oculotrichodysplasia

LX4Y\_DER Cartilage-hair hypoplasia

LX4Z\_DER Trichorhinophalangeal syndrome type 1 and 3

LX50\_DER Chondroectodermal dysplasia

LD27.01 Cronkhite-Canada syndrome

LD27.02 Hypohidrotic ectodermal dysplasia

LD27.03 Hidrotic ectodermal dysplasia, Clouston type

LX51\_DER Odonto-onycho-dermal dysplasia

LX52\_DER Woolly hair – hypotrichosis – everted lower lip – outstanding ears

LX53\_DER Autosomal dominant palmoplantar keratoderma and congenital alopecia

LX54\_DER BIDS syndrome

LX55\_DER IBIDS syndrome

LX56\_DER Sabinas brittle hair syndrome

LX57\_DER Onycho-tricho-dysplasia – neutropaenia syndrome

LX58\_DER Autosomal recessive palmoplantar keratoderma and congenital alopecia

LX59\_DER Alopecia - contractures - dwarfism - intellectual deficit

LX5A\_DER Cataract - alopecia - sclerodactyly

LX5B\_DER Odonto-onycho dysplasia - alopecia

LX5C\_DER Schöpf-Schulz-Passarge syndrome

LX5D\_DER Hypertrichosis lanuginosa congenita

LD27.0Y\_DER Other specified ectodermal dysplasia syndromes

Syndromic ichthyosis (BlockL2‑LX5)

X-linked ichthyosis syndromes (BlockL3‑LX5)

LX5E\_DER Ichthyosis follicularis – atrichia – photophobia

LX5Z\_DER Other specified x-linked ichthyosis syndromes

LX5F\_DER Autosomal ichthyosis syndromes with hair abnormalities

LX5G\_DER Autosomal ichthyosis syndromes with neurological manifestations

LX5H\_DER Autosomal ichthyoses with fatal disease course

LX5J\_DER Other autosomal ichthyosis syndromes

LX5K\_DER Ichthyosis as component of other specified genetic syndrome

LX5Z\_DER Other specified syndromic ichthyosis

Genetic syndromes affecting nails (BlockL2‑LX5)

LX5L\_DER Oto-onycho-peroneal syndrome

LX5M\_DER Congenital onychodysplasia of the index fingers

LX5N\_DER Nail dystrophy resulting from epidermolysis bullosa classified elsewhere

LX5P\_DER Nail involvement in other specified genetic disease

LX5Z\_DER Other specified genetic syndromes affecting nails

Genetic hamartoneoplastic syndromes affecting the skin (BlockL2‑LX5)

LX5Q\_DER Legius syndrome

LX5R\_DER Bazex-Dupré-Christol syndrome

LX5S\_DER Birt-Hogg-Dubé syndrome

LX5Z\_DER Other specified genetic hamartoneoplastic syndromes affecting the skin

Genetic lipodystrophy (BlockL2‑LX5)

LX5T\_DER Generalised congenital lipodystrophy with myopathy

LX5U\_DER Lipodystrophy due to peptidic growth factors deficiency

LX5V\_DER Mandibuloacral dysplasia

LX5W\_DER SHORT syndrome

LX5X\_DER Nakajo-Nishimura syndrome

LD27.6Z Genetic lipodystrophy, unspecified

Syndromes with premature ageing appearance as a major feature (BlockL2‑LX6)

LX63\_DER Cockayne syndrome

LX64\_DER Rothmund-Thomson syndrome

LX65\_DER Progeria

LX66\_DER Werner syndrome

LX67\_DER Acrogeria

LX68\_DER Wiedemann-Rautenstrauch progeroid syndrome

LX69\_DER Mulvihill-Smith progeroid syndrome

LX6A\_DER Geroderma osteodysplasticum

LX6B\_DER Wrinkly skin syndrome

LX6C\_DER Metageria

LX6D\_DER Hallermann-Streiff-François syndrome

LX6E\_DER Kindler syndrome

LX6Z\_DER Other specified syndromes with premature ageing appearance as a major feature

LX6Z\_DER Syndromes with premature ageing appearance as a major feature, unspecified

LD40.0 Complete trisomy 21

LD40.1 Complete trisomy 13

LD40.2 Complete trisomy 18

LX6F\_DER Cat-eye syndrome

LX6G\_DER Recombinant chromosome 8 syndrome

LX6H\_DER 1q21.1 deletion

LX6J\_DER 1q41q42 deletion

LX6K\_DER 2q32q33 deletion

LX6L\_DER 2q37 deletion

LX6M\_DER 2p15p16.1 deletion

LX6N\_DER 3q29 deletion

LD44.41 Deletions of the short arm of chromosome 4

LX6P\_DER Cri-du-chat syndrome

LX6Q\_DER Williams-Beuren syndrome

LX6R\_DER 9q34 deletion

LX6S\_DER Alfi syndrome

LX6T\_DER 15q26.3 deletion

LX6U\_DER 16p13.3 deletion

LX6V\_DER 17q21.31 deletion

LX6W\_DER De Grouchy syndrome

LX6X\_DER 19q13.11 deletion

LX6Y\_DER 22q13 deletion

LD44.N0 CATCH 22 phenotype

LD50.0 Turner syndrome

LD50.3 Klinefelter syndrome

LX6Z\_DER De Sanctis-Cacchione syndrome

LX70\_DER Alopecia – psychomotor epilepsy – periodontal pyorrhoea – intellectual disability syndrome

LD9Y\_DER Other specified developmental anomalies

LD9Z Developmental anomalies, unspecified

CHAPTER 21 Symptoms, signs or clinical findings, not elsewhere classified

Somatic delusion directed at the skin (BlockL1‑MX7)

MX71\_DER Delusion of infestation

MX7Z\_DER Other specified somatic delusion directed at the skin

MB27.26 Tactile hallucinations

MX72\_DER Symptoms, signs or clinical findings of the nervous system

Symptom or signs involving the skin (BlockL1‑ME6)

ME60 Skin lesion of uncertain or unspecified nature

ME61 Subcutaneous swelling, mass or lump of uncertain or unspecified nature

ME62 Acute skin eruption of uncertain or unspecified nature

ME63 Chronic skin disorder of uncertain or unspecified nature

Non-specific cutaneous vascular signs (BlockL2‑ME6)

ME64.0 Erythema

ME64.1 Cyanosis

ME64.2 Pallor

ME64.3 Petechiae

ME64.4 Flushing

Disturbances of skin sensation of unspecified aetiology (BlockL2‑ME6)

ME65.0 Burning of skin

ME65.1 Itching of skin

ME65.2 Pain or tenderness of skin

ME65.3 Stinging of skin

ME65.4 Tingling of skin

ME65.Y\_DER Other specified disturbance of skin sensation

Miscellaneous non-specific skin-related symptoms and signs (BlockL2‑ME6)

ME66.0 Abnormal sensitivity to light or UV radiation of uncertain or unspecified nature

ME66.1 Changes in skin texture

ME66.2 Excess and redundant skin

MX74\_DER Fear of skin disease

ME66.3 Symptom or complaint relating to hair or scalp

ME66.4 Symptom or complaint relating to nails

MX75\_DER Retention hyperkeratosis

ME66.6 Rash

ME66.Y\_DER Other specified skin changes

ME67 Skin disorder of uncertain or unspecified nature

ME6Y\_DER Other specified symptom or signs involving the skin

MX76\_DER Fear of cancer of skin

MX77\_DER Oedema due to dependency or immobility

MG29.10 Oedema due to increased capillary pressure

MX78\_DER Oedema due to hypoproteinaemia

MX79\_DER Skin infection classified elsewhere due to methicillin resistant Staphylococcus aureus

MX7A\_DER Skin infection classified elsewhere due to community-acquired methicillin resistant Staphylococcus aureus

MX7B\_DER Skin infection classified elsewhere due to hospital-acquired methicillin resistant Staphylococcus aureus

MH2Y\_DER Other specified symptoms, signs or clinical findings, not elsewhere classified

CHAPTER 22 Injury, poisoning or certain other consequences of external causes

NX7C\_DER Traumatic injury to nail bed or matrix of nail of hand

NX7D\_DER Subungual haematoma

NX7E\_DER Traumatic injury to nail bed or matrix of nail of foot

ND5Z Injuries to unspecified part of trunk, limb or body region

ND56 Injury of unspecified body region

ND73.20 Trichobezoar

ND90 Burn of head or neck except face

ND91 Burn of face except eye or ocular adnexa

ND98 Chemical burn due to skin contact with corrosive substance

ND99 Acute skin injury due to skin contact with corrosive substance

Frostbite (BlockL1‑NE4)

NE40 Superficial frostbite

NE41 Frostbite with tissue necrosis

NE4Z Frostbite, unspecified

NX7K\_DER Harmful effects of or exposure to noxious substances chiefly nonmedicinal as to source, Ethanol

NX7L\_DER Harmful effects of or exposure to noxious substances chiefly nonmedicinal as to source, methanol

NX7M\_DER Harmful effects of or exposure to noxious substances chiefly nonmedicinal as to source, 2-Propanol

NX7N\_DER Harmful effects of or exposure to noxious substances chiefly nonmedicinal as to source, fusel oil

NX7P\_DER Harmful effects of or exposure to noxious substances, Substances chiefly nonmedicinal as to source, Halogen derivatives of aliphatic or aromatic hydrocarbons, Tetrachloroethylene

NX7Q\_DER Harmful effects of or exposure to noxious substances, Substances chiefly nonmedicinal as to source, Other gases, fumes or vapours, Nitrogen oxides

NX7R\_DER Harmful effects of or exposure to noxious substances, Substances chiefly nonmedicinal as to source, Other gases, fumes or vapours, Chlorine gas

NX7S\_DER Harmful effects of or exposure to noxious substances, Substances chiefly nonmedicinal as to source, Metals, Lead or its compounds

NX7T\_DER Harmful effects of or exposure to noxious substances, Substances chiefly nonmedicinal as to source, Metals, Mercury or its compounds

NX7U\_DER Harmful effects of or exposure to noxious substances, Substances chiefly nonmedicinal as to source, Metals, Chromium or its compounds

NX7V\_DER Harmful effects of or exposure to noxious substances, Substances chiefly nonmedicinal as to source, Metals, Copper or its compounds

NX7W\_DER Harmful effects of or exposure to noxious substances, Substances chiefly nonmedicinal as to source, Metals, Zinc or its compounds

NX7X\_DER Harmful effects of or exposure to noxious substances, Substances chiefly nonmedicinal as to source, Metals, Tin or its compounds

NX7Y\_DER Harmful effects of or exposure to noxious substances, Substances chiefly nonmedicinal as to source, Metals, Beryllium or its compounds

NX7Z\_DER Harmful effects of or exposure to noxious substances, Substances chiefly nonmedicinal as to source, Metals, Thallium

NX80\_DER Harmful effects of or exposure to noxious substances, Substances chiefly nonmedicinal as to source, Other inorganic substances, Hydrogen cyanide

NX81\_DER Harmful effects of or exposure to noxious substances, Substances chiefly nonmedicinal as to source, Noxious substances eaten as seafood, Scombroid fish poisoning

Cutaneous reactions to arthropods (BlockL1‑NX8)

Insect bites or stings (BlockL2‑NX8)

NX82\_DER Mosquito bites

NX83\_DER Sand-fly bites

NX84\_DER Black-fly bites

NX85\_DER Midge bites

NX86\_DER Horse-fly bites

NX87\_DER Flea bites

NX88\_DER Wasp stings

NX89\_DER Bee stings

NX8A\_DER Ant bites or stings

NX8B\_DER Hornet stings

NX8Z\_DER Bites and stings due to other specified insects

NX8Z\_DER Insect bite/sting

Arachnid bites or stings (BlockL2‑NX8)

NX8C\_DER Spider bites

NX8D\_DER Tick bites

NX8Z\_DER Other specified arachnid bites or stings

NX8Z\_DER Arachnid bites or stings, unspecified

NX8E\_DER Other cutaneous reactions to arthropods

Cutaneous reactions to caterpillars (BlockL3‑NX8)

NX8F\_DER Butterfly caterpillar reaction

NX8G\_DER Erucism due to Premolis larvae

NX8H\_DER Moth caterpillar reaction

NX8Z\_DER Other specified cutaneous reactions to caterpillars

NX8Z\_DER Cutaneous reactions to caterpillars, unspecified

Cutaneous reactions to centipedes or millipedes (BlockL3‑NX8)

NX8J\_DER Cutaneous reactions to centipede bite

NX8K\_DER Cutaneous reactions to millipede sting

NX8Z\_DER Other specified cutaneous reactions to centipedes or millipedes

NX8Z\_DER Cutaneous reactions to centipedes or millipedes, unspecified

Cutaneous reactions to zoonotic mites (BlockL3‑NX8)

NX8L\_DER Cheyletiellosis

NX8M\_DER Trombiculosis

NX8N\_DER Pyemotes mite dermatitis

NX8P\_DER Bird mite dermatitis

NX8Z\_DER Cutaneous reactions to other specified mites

NX8Z\_DER Other specified cutaneous reactions to arthropods

NX8Z\_DER Cutaneous reactions to arthropods, unspecified

Cutaneous reactions to venomous or noxious aquatic invertebrates (BlockL1‑NX8)

Cutaneous reactions to Cnidaria (BlockL2‑NX8)

NX8Q\_DER Jellyfish, Scyphozoa sting

NX8R\_DER Jellyfish, Cubozoa sting

NX8S\_DER Hydrozoa envenomation

NX8T\_DER Sea anemone dermatitis

NX8U\_DER Anthozoa coral injury

NX8V\_DER Sea-bather's eruption

NX9Z\_DER Other specified cutaneous reactions to Cnidaria

NX9Z\_DER Cutaneous reactions to Cnidaria, unspecified

NX8W\_DER Cutaneous reactions to Bryozoa

NX8X\_DER Cutaneous reactions to Echinoidea

NX8Y\_DER Cutaneous reactions to Porifera

Cutaneous reactions to marine Mollusca (BlockL2‑NX8)

NX8Z\_DER Cone shell envenomation

NX90\_DER Octopus bite

NX9Z\_DER Other specified cutaneous reactions to marine Mollusca

NX9Z\_DER Cutaneous reactions to marine Mollusca, unspecified

NX9Z\_DER Other specified cutaneous reactions to venomous or noxious aquatic invertebrates

NX9Z\_DER Cutaneous reactions to venomous or noxious aquatic invertebrates, unspecified

Cutaneous reactions to venomous or noxious vertebrates (BlockL1‑NX9)

NX91\_DER Fish sting

NX9Z\_DER Other specified cutaneous reactions to venomous or noxious vertebrates

NX9Z\_DER Cutaneous reactions to venomous or noxious vertebrates, unspecified

NX92\_DER Harmful effects of or exposure to noxious substances, Substances chiefly nonmedicinal as to source, Nitroderivatives or aminoderivatives of benzene or its homologues

NX93\_DER Harmful effects of or exposure to noxious substances, Substances chiefly nonmedicinal as to source, Carbon disulfide

NX94\_DER Harmful effects of or exposure to noxious substances, Substances chiefly nonmedicinal as to source, Nitroglycerin or other nitric acids or esters

NE8Z Injury or harm arising from surgical or medical care, not elsewhere classified

NF03 Other effects of reduced temperature

NF08 Effects of certain specified external causes

NF0A.3 Post traumatic wound infection, not elsewhere classified

NF2Y\_DER Other specified injury, poisoning or certain other consequences of external causes

NF2Z Unspecified injury, poisoning or certain other consequences of external causes

CHAPTER 23 External causes of morbidity or mortality

PX95\_DER Causes of healthcare related harm or injury

PL2Y\_DER Other specified external causes of morbidity or mortality

PL2Z External causes of morbidity or mortality, unspecified

CHAPTER 24 Factors influencing health status or contact with health services

QX96\_DER Reasons for contact with the health services

QF4Y\_DER Other specified factors influencing health status or contact with health services

QF4Z Factors influencing health status or contact with health services, unspecified

CHAPTER \*\*\* Special Views

?Y66\_DER Special tabulation list of infectious agents

?Y6Z\_DER Other specified special Views

?Y6Z\_DER Special Views, unspecified