ΣΠΑΝΙΕΣ ΡΕΥΜΑΤΟΛΟΓΙΚΕΣ ΠΑΘΗΣΕΙΣ ΜΕ ΚΩΔΙΚΟ ΑΠΟ ΤΟ ORPHANET

* **Rare systemic or rheumatologic disease ORPHA:98023**

**Non-histaminic angioedema ORPHA:658**

* + - Acquired angioedema ORPHA:91385
		- Acquired angioedema with C1Inh deficiency ORPHA:528663
* Renin-angiotensin-aldosterone system-blocker-induced angioedema ORPHA:100057
	+ - [Hereditary angioedema](https://www.orpha.net/consor/cgi-bin/Disease_Search.php?lng=EN&data_id=12136&disease=Hereditary-angioedema&search=Disease_Search_Simple) ORPHA:91378
		- Hereditary angioedema with C1Inh deficiency ORPHA:528623
	+ Hereditary angioedema with normal C1Inh ORPHA:528647

**Rare rheumatologic disease ORPHA:182231**

* + - [Adult-onset Still disease](https://www.orpha.net/consor/cgi-bin/Disease_Search.php?lng=EN&data_id=5525&disease=Adult-onset-Still-disease&search=Disease_Search_Simple) ORPHA:829
		- [Autoinflammatory syndrome](https://www.orpha.net/consor/cgi-bin/Disease_Search.php?lng=EN&data_id=12458&disease=Autoinflammatory-syndrome&search=Disease_Search_Simple) ORPHA:93665
			* Granulomatous autoinflammatory syndrome ORPHA:324930
			* Mixed autoinflammatory and autoimmune syndrome ORPHA:324933
			* Periodic fever syndrome ORPHA:101995
			* Pyogenic autoinflammatory syndrome ORPHA:324927
			* Type 1 interferonopathy ORPHA:477647
			* Unclassified autoinflammatory syndrome ORPHA:324936
		- [Camptodactyly-arthropathy-coxa-vara-pericarditis syndrome](https://www.orpha.net/consor/cgi-bin/Disease_Search.php?lng=EN&data_id=2583&disease=Camptodactyly-arthropathy-coxa-vara-pericarditis-syndrome&search=Disease_Search_Simple) ORPHA:2848
		- [Familial calcium pyrophosphate deposition](https://www.orpha.net/consor/cgi-bin/Disease_Search.php?lng=EN&data_id=1614&disease=Familial-calcium-pyrophosphate-deposition&search=Disease_Search_Simple) ORPHA:1416
		- [Felty syndrome](https://www.orpha.net/consor/cgi-bin/Disease_Search.php?lng=EN&data_id=10611&disease=Felty-syndrome&search=Disease_Search_Simple) ORPHA:47612
		- [Fibroblastic rheumatism](https://www.orpha.net/consor/cgi-bin/Disease_Search.php?lng=EN&data_id=25135&disease=Fibroblastic-rheumatism&search=Disease_Search_Simple) ORPHA:477650
		- [Gorham-Stout disease](https://www.orpha.net/consor/cgi-bin/Disease_Search.php?lng=EN&data_id=8684&disease=Gorham-Stout-disease&search=Disease_Search_Simple) ORPHA:73
		- [Hajdu-Cheney syndrome](https://www.orpha.net/consor/cgi-bin/Disease_Search.php?lng=EN&data_id=1276&disease=Hajdu-Cheney-syndrome&search=Disease_Search_Simple) ORPHA:955
		- [Idiopathic juvenile osteoporosis](https://www.orpha.net/consor/cgi-bin/Disease_Search.php?lng=EN&data_id=11651&disease=Idiopathic-juvenile-osteoporosis&search=Disease_Search_Simple) ORPHA:85193
		- [Immune dysregulation-inflammatory bowel disease-arthritis-recurrent infections-lymphopenia syndrome](https://www.orpha.net/consor/cgi-bin/Disease_Search.php?lng=EN&data_id=27635&disease=Immune-dysregulation-inflammatory-bowel-disease-arthritis-recurrent-infections-lymphopenia-syndrome&search=Disease_Search_Simple) ORPHA:529977
		- [Immune dysregulation-inflammatory bowel disease-arthritis-recurrent infections syndrome](https://www.orpha.net/consor/cgi-bin/Disease_Search.php?lng=EN&data_id=19277&disease=Immune-dysregulation-inflammatory-bowel-disease-arthritis-recurrent-infections-syndrome&search=Disease_Search_Simple) ORPHA:238569
		- [Intermittent hydrarthrosis](https://www.orpha.net/consor/cgi-bin/Disease_Search.php?lng=EN&data_id=21950&disease=Intermittent-hydrarthrosis&search=Disease_Search_Simple) ORPHA:329967
		- [Interstitial granulomatous dermatitis with arthritis](https://www.orpha.net/consor/cgi-bin/Disease_Search.php?lng=EN&data_id=11141&disease=Interstitial-granulomatous-dermatitis-with-arthritis&search=Disease_Search_Simple) ORPHA:79099
		- [Isolated sternocostoclavicular hyperostosis](https://www.orpha.net/consor/cgi-bin/Disease_Search.php?lng=EN&data_id=18011&disease=Isolated-sternocostoclavicular-hyperostosis&search=Disease_Search_Simple) ORPHA:178311
		- [Juvenile idiopathic arthritis](https://www.orpha.net/consor/cgi-bin/Disease_Search.php?lng=EN&data_id=720&disease=Juvenile-idiopathic-arthritis&search=Disease_Search_Simple) ORPHA:92
* Enthesitis-related juvenile idiopathic arthritis ORPHA:85438
* Oligoarticular juvenile idiopathic arthritis ORPHA:85410
* Polyarticular juvenile idiopathic arthritis ORPHA:404580
* Psoriasis-related juvenile idiopathic arthritis ORPHA:85436
* Systemic-onset juvenile idiopathic arthritis ORPHA:85414
* Unspecified juvenile idiopathic arthritis ORPHA:91140
	+ - [Osteonecrosis](https://www.orpha.net/consor/cgi-bin/Disease_Search.php?lng=EN&data_id=22750&disease=Osteonecrosis&search=Disease_Search_Simple) ORPHA:399158
* Avascular necrosis ORPHA:399164
* Epiphysiolysis of the hip ORPHA:399329
* Familial osteochondritis dissecans ORPHA:251262
* Idiopathic phalangeal acro-osteolysis ORPHA:444316
* [Mueller-Weiss syndrome](https://www.orpha.net/consor/cgi-bin/Disease_Search.php?lng=EN&data_id=28560&disease=Mueller-Weiss-syndrome&search=Disease_Search_Simple) ORPHA:566943
* Osteochondritis dissecans ORPHA:2764
* Osteochondrosis ORPHA:399319
* Osteoradionecrosis of the mandible ORPHA:521127
	+ - [Polymyalgia rheumatica](https://www.orpha.net/consor/cgi-bin/Disease_Search.php?lng=EN&data_id=12411&disease=Polymyalgia-rheumatica&search=Disease_Search_Simple) ORPHA:93569
		- [Reactive arthritis](https://www.orpha.net/consor/cgi-bin/Disease_Search.php?lng=EN&data_id=8778&disease=Reactive-arthritis&search=Disease_Search_Simple) ORPHA:29207
		- [Sweet syndrome](https://www.orpha.net/consor/cgi-bin/Disease_Search.php?lng=EN&data_id=3057&disease=Sweet-syndrome&search=Disease_Search_Simple) ORPHA:3243
		- [Tenosynovial giant cell tumor](https://www.orpha.net/consor/cgi-bin/Disease_Search.php?lng=EN&data_id=10885&disease=Tenosynovial-giant-cell-tumor&search=Disease_Search_Simple) ORPHA:66627

[**Rare systemic disease**](https://www.orpha.net/consor/cgi-bin/Disease_Search.php?lng=EN&data_id=18224&disease=Rare-systemic-disease&search=Disease_Search_Simple)**ORPHA:182222**

* [Amyloidosis](https://www.orpha.net/consor/cgi-bin/Disease_Search.php?lng=EN&data_id=632&disease=Amyloidosis&search=Disease_Search_Simple) ORPHA:69
* [Calciphylaxis](https://www.orpha.net/consor/cgi-bin/Disease_Search.php?lng=EN&data_id=20422&disease=Calciphylaxis&search=Disease_Search_Simple) ORPHA:280062
* [CAR T cell therapy-associated cytokine release syndrome](https://www.orpha.net/consor/cgi-bin/Disease_Search.php?lng=EN&data_id=28040&disease=CAR-T-cell-therapy-associated-cytokine-release-syndrome&search=Disease_Search_Simple) ORPHA:542323
* [Connective tissue disorder due to lysyl hydroxylase-3 deficiency](https://www.orpha.net/consor/cgi-bin/Disease_Search.php?lng=EN&data_id=21104&disease=Connective-tissue-disorder-due-to-lysyl-hydroxylase-3-deficiency&search=Disease_Search_Simple) ORPHA:300284
* [Drug-induced lupus erythematosus](https://www.orpha.net/consor/cgi-bin/Disease_Search.php?lng=EN&data_id=19154&disease=Drug-induced-lupus-erythematosus&search=Disease_Search_Simple) ORPHA:231111
* [Ehlers-Danlos/osteogenesis imperfecta syndrome](https://www.orpha.net/consor/cgi-bin/Disease_Search.php?lng=EN&data_id=19145&disease=Ehlers-Danlos-osteogenesis-imperfecta-syndrome&search=Disease_Search_Simple) ORPHA:230857
* [Ehlers-Danlos syndrome](https://www.orpha.net/consor/cgi-bin/Disease_Search.php?lng=EN&data_id=13266&disease=Ehlers-Danlos-syndrome&search=Disease_Search_Simple) ORPHA:98249
* [Familial articular hypermobility syndrome](https://www.orpha.net/consor/cgi-bin/Disease_Search.php?lng=EN&data_id=2162&disease=Familial-articular-hypermobility-syndrome&search=Disease_Search_Simple) ORPHA:2295
* [Hereditary hemorrhagic telangiectasia](https://www.orpha.net/consor/cgi-bin/Disease_Search.php?lng=EN&data_id=236&disease=Hereditary-hemorrhagic-telangiectasia&search=Disease_Search_Simple) ORPHA:774
* [Hypoplasminogenemia](https://www.orpha.net/consor/cgi-bin/Disease_Search.php?lng=EN&data_id=8694&disease=Hypoplasminogenemia&search=Disease_Search_Simple) ORPHA:722
* [Kikuchi-Fujimoto disease](https://www.orpha.net/consor/cgi-bin/Disease_Search.php?lng=EN&data_id=10660&disease=Kikuchi-Fujimoto-disease&search=Disease_Search_Simple) ORPHA:50918
* [Kimura disease](https://www.orpha.net/consor/cgi-bin/Disease_Search.php?lng=EN&data_id=8612&disease=Kimura-disease&search=Disease_Search_Simple) ORPHA:482
* [LAMA5-related multisystemic syndrome](https://www.orpha.net/consor/cgi-bin/Disease_Search.php?lng=EN&data_id=27333&disease=LAMA5-related-multisystemic-syndrome&search=Disease_Search_Simple) ORPHA:521450
* [Langerhans cell histiocytosis](https://www.orpha.net/consor/cgi-bin/Disease_Search.php?lng=EN&data_id=3768&disease=Langerhans-cell-histiocytosis&search=Disease_Search_Simple) ORPHA:389
* [Macrophage activation syndrome](https://www.orpha.net/consor/cgi-bin/Disease_Search.php?lng=EN&data_id=17186&disease=Macrophage-activation-syndrome&search=Disease_Search_Simple) ORPHA:158061
* [Malignant atrophic papulosis](https://www.orpha.net/consor/cgi-bin/Disease_Search.php?lng=EN&data_id=8660&disease=Malignant-atrophic-papulosis&search=Disease_Search_Simple) ORPHA:679
* [Marfan syndrome and Marfan-related disorders](https://www.orpha.net/consor/cgi-bin/Disease_Search.php?lng=EN&data_id=20632&disease=Marfan-syndrome-and-Marfan-related-disorders&search=Disease_Search_Simple) ORPHA:284993
* [Multisystem inflammatory syndrome in children and adults](https://www.orpha.net/consor/cgi-bin/Disease_Search.php?lng=EN&data_id=29878&disease=Multisystem-inflammatory-syndrome-in-children-and-adults&search=Disease_Search_Simple) ORPHA:598363
* [Myalgia-eosinophilia syndrome associated with tryptophan](https://www.orpha.net/consor/cgi-bin/Disease_Search.php?lng=EN&data_id=3134&disease=Myalgia-eosinophilia-syndrome-associated-with-tryptophan&search=Disease_Search_Simple) ORPHA:2582
* [Non-Langerhans cell histiocytosis](https://www.orpha.net/consor/cgi-bin/Disease_Search.php?lng=EN&data_id=17169&disease=Non-Langerhans-cell-histiocytosis&search=Disease_Search_Simple) ORPHA:157987
* [Postorgasmic illness syndrome](https://www.orpha.net/consor/cgi-bin/Disease_Search.php?lng=EN&data_id=20421&disease=Postorgasmic-illness-syndrome&search=Disease_Search_Simple) ORPHA:279947
* [Progeroid and marfanoid aspect-lipodystrophy syndrome](https://www.orpha.net/consor/cgi-bin/Disease_Search.php?lng=EN&data_id=21117&disease=Progeroid-and-marfanoid-aspect-lipodystrophy-syndrome&search=Disease_Search_Simple) ORPHA:300382
* [Pyomyositis](https://www.orpha.net/consor/cgi-bin/Disease_Search.php?lng=EN&data_id=8609&disease=Pyomyositis&search=Disease_Search_Simple) ORPHA:764
* [Retinal ischemic syndrome-digestive tra](https://www.orpha.net/consor/cgi-bin/Disease_Search.php?lng=EN&data_id=2714&disease=Retinal-ischemic-syndrome-digestive-tract-small-vessel-hyalinosis-diffuse-cerebral-calcifications-syndrome&search=Disease_Search_Simple)

[**Vasculitis**](https://www.orpha.net/consor/cgi-bin/Disease_Search.php?lng=EN&data_id=10704&disease=Vasculitis&search=Disease_Search_Simple)**ORPHA:52759**

* + [Predominantly large-vessel vasculitis](https://www.orpha.net/consor/cgi-bin/Disease_Search.php?lng=EN&data_id=17095&disease=Predominantly-large-vessel-vasculitis&search=Disease_Search_Simple) ORPHA:156140
	+ [Predominantly medium-vessel vasculitis](https://www.orpha.net/consor/cgi-bin/Disease_Search.php?lng=EN&data_id=17096&disease=Predominantly-medium-vessel-vasculitis&search=Disease_Search_Simple) ORPHA:156143
	+ [Predominantly small-vessel vasculitis](https://www.orpha.net/consor/cgi-bin/Disease_Search.php?lng=EN&data_id=17097&disease=Predominantly-small-vessel-vasculitis&search=Disease_Search_Simple) ORPHA:156146
	+ [Secondary vasculitis](https://www.orpha.net/consor/cgi-bin/Disease_Search.php?lng=EN&data_id=23521&disease=Secondary-vasculitis&search=Disease_Search_Simple) ORPHA:445197
	+ [Unclassified vasculitis](https://www.orpha.net/consor/cgi-bin/Disease_Search.php?lng=EN&data_id=19629&disease=Unclassified-vasculitis&search=Disease_Search_Simple) ORPHA:251328