Sirkku Peltonen

List of Publications by Year in descending order

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201658 223791 2,228 50 27 46 citations h-index g-index papers 51 51 51 3224 docs citations times ranked citing authors all docs

#	Article	IF	Citations
1	Distinctive Cancer Associations in Patients With Neurofibromatosis Type 1. Journal of Clinical Oncology, 2016, 34, 1978-1986.	1.6	271
2	Incidence and Mortality of Neurofibromatosis: A Total Population Study in Finland. Journal of Investigative Dermatology, 2015, 135, 904-906.	0.7	189
3	Autosomal Dominant Diabetes Arising From a Wolfram Syndrome 1 Mutation. Diabetes, 2013, 62, 3943-3950.	0.6	100
4	Barriers of the peripheral nerve. Tissue Barriers, 2013, 1, e24956.	3.2	97
5	Early Dental Epithelial Transcription Factors Distinguish Ameloblastoma from Keratocystic Odontogenic Tumor. Journal of Dental Research, 2015, 94, 101-111.	5.2	82
6	Complement Factor H: A Biomarker for Progression of Cutaneous Squamous Cell Carcinoma. Journal of Investigative Dermatology, 2014, 134, 498-506.	0.7	73
7	Breast cancer in neurofibromatosis type 1: overrepresentation of unfavourable prognostic factors. British Journal of Cancer, 2017, 116, 211-217.	6.4	69
8	Complement Factor I Promotes Progression of Cutaneous Squamous Cell Carcinoma. Journal of Investigative Dermatology, 2015, 135, 579-588.	0.7	68
9	Class III \hat{I}^2 -Tubulin Is a Component of the Mitotic Spindle in Multiple Cell Types. Journal of Histochemistry and Cytochemistry, 2008, 56, 1113-1119.	2.5	64
10	The Development of Cutaneous Neurofibromas. American Journal of Pathology, 2011, 178, 500-505.	3.8	63
11	Complement Component C3 and Complement Factor B Promote Growth of Cutaneous Squamous Cell Carcinoma. American Journal of Pathology, 2017, 187, 1186-1197.	3.8	63
12	A role for caveolin-1 in desmoglein binding and desmosome dynamics. Oncogene, 2012, 31, 1636-1648.	5.9	62
13	Long Noncoding RNA PICSAR Promotes Growth of Cutaneous Squamous Cell Carcinoma by Regulating ERK1/2 Activity. Journal of Investigative Dermatology, 2016, 136, 1701-1710.	0.7	61
14	Compound Heterozygous Desmoplakin Mutations Result in a Phenotype with a Combination of Myocardial, Skin, Hair, and Enamel Abnormalities. Journal of Investigative Dermatology, 2010, 130, 968-978.	0.7	57
15	Reevaluation of the Normal Epidermal Calcium Gradient, and Analysis of Calcium Levels and ATP Receptors in Hailey–Hailey and Darier Epidermis. Journal of Investigative Dermatology, 2009, 129, 1379-1387.	0.7	55
16	A controlled register-based study of 460 neurofibromatosis 1 patients: Increased fracture risk in children and adults over 41 years of age. Journal of Bone and Mineral Research, 2012, 27, 2333-2337.	2.8	55
17	Congenital pseudarthrosis of neurofibromatosis type 1: Impaired osteoblast differentiation and function and altered NF1 gene expression. Bone, 2009, 44, 243-250.	2.9	49
18	Osteoclasts in neurofibromatosis type 1 display enhanced resorption capacity, aberrant morphology, and resistance to serum deprivation. Bone, 2010, 47, 583-590.	2.9	49

#	Article	IF	CITATIONS
19	EphB2 Promotes Progression of Cutaneous Squamous Cell Carcinoma. Journal of Investigative Dermatology, 2015, 135, 1882-1892.	0.7	48
20	Tight Junction Proteins in Human Schwann Cell Autotypic Junctions. Journal of Histochemistry and Cytochemistry, 2009, 57, 523-529.	2.5	46
21	c-Src/Cav1-dependent activation of the EGFR by Dsg2. Oncotarget, 2016, 7, 37536-37555.	1.8	46
22	Radiographic Findings in the Jaws of Patients With Neurofibromatosis 1. Journal of Oral and Maxillofacial Surgery, 2012, 70, 1351-1357.	1.2	42
23	Legius syndrome in fourteen families. Human Mutation, 2011, 32, E1985-E1998.	2.5	40
24	Neurofibromatosis type 1 (NF1) gene: Beyond caf \tilde{A} © au lait spots and dermal neurofibromas. Experimental Dermatology, 2017, 26, 645-648.	2.9	39
25	Oral soft tissue alterations in patients with neurofibromatosis. Clinical Oral Investigations, 2012, 16, 551-558.	3.0	37
26	Pediatric malignancies in neurofibromatosis type 1: A populationâ€based cohort study. International Journal of Cancer, 2019, 145, 2926-2932.	5.1	36
27	Hailey–Hailey disease and tight junctions: Claudins 1 and 4 are regulated by <scp>ATP</scp> 2C1 gene encoding Ca ²⁺ /Mn ²⁺ <scp>ATP</scp> ase <scp>SPCA</scp> 1 in cultured keratinocytes. Experimental Dermatology, 2012, 21, 586-591.	2.9	33
28	Speech characteristics in neurofibromatosis type 1. American Journal of Medical Genetics, Part A, 2010, 152A, 42-51.	1.2	26
29	Breast cancer in neurofibromatosis 1: survival and risk of contralateral breast cancer in a five country cohort study. Genetics in Medicine, 2020, 22, 398-406.	2.4	26
30	Keratinocyte Growth Factor Induces Gene Expression Signature Associated with Suppression of Malignant Phenotype of Cutaneous Squamous Carcinoma Cells. PLoS ONE, 2012, 7, e33041.	2.5	24
31	Congenital anomalies in neurofibromatosis 1: a retrospective register-based total population study. Orphanet Journal of Rare Diseases, 2018, 13, 5.	2.7	23
32	Short mandible, maxilla and cranial base are common in patients with neurofibromatosis 1. European Journal of Oral Sciences, 2011, 119, 121-127.	1.5	22
33	Neurofibromatosis 1-Related Osteopenia Often Progresses to Osteoporosis in 12ÂYears. Calcified Tissue International, 2013, 92, 23-27.	3.1	20
34	Follow-Up of Six Patients with Neurofibromatosis 1-Related Osteoporosis Treated with Alendronate for 23 Months. Calcified Tissue International, 2014, 94, 608-612.	3.1	19
35	Osteoclasts derived from patients with neurofibromatosis 1 (NF1) display insensitivity to bisphosphonates in vitro. Bone, 2012, 50, 798-803.	2.9	18
36	p38δ mitogen-activated protein kinase regulates the expression of tight junction protein ZO-1 in differentiating human epidermal keratinocytes. Archives of Dermatological Research, 2014, 306, 131-141.	1.9	18

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37	The pregnancy in neurofibromatosis 1: A retrospective registerâ€based total population study. American Journal of Medical Genetics, Part A, 2017, 173, 2641-2648.	1.2	17
38	Neurofibromatosis 1 and dental caries. Clinical Oral Investigations, 2011, 15, 119-121.	3.0	15
39	Diversity of actin architecture in human osteoclasts: network of curved and branched actin supporting cell shape and intercellular micrometer-level tubes. Molecular and Cellular Biochemistry, 2017, 432, 131-139.	3.1	13
40	A rare disease and education: Neurofibromatosis type 1 decreases educational attainment. Clinical Genetics, 2021, 99, 529-539.	2.0	13
41	Dental age in patients with neurofibromatosis 1. European Journal of Oral Sciences, 2012, 120, 549-552.	1.5	12
42	Expression of claudinâ€11 by tumor cells in cutaneous squamous cell carcinoma is dependent on the activity of p38l´. Experimental Dermatology, 2017, 26, 771-777.	2.9	12
43	Treatment of cutaneous neurofibromas with carbon dioxide laser: Technique and patient experience. European Journal of Medical Genetics, 2022, 65, 104386.	1.3	11
44	Intestinal tumors in neurofibromatosis 1 with special reference to fatal gastrointestinal stromal tumors (GIST). Molecular Genetics & Enomic Medicine, 2019, 7, e927.	1.2	10
45	Phenotypic characterization of transgenic mice harboring Nf1 ^{+/â^'} or Nf1 ^{â^'/â^'} osteoclasts in otherwise Nf1 ^{+/+} background. Journal of Cellular Biochemistry, 2012, 113, 2136-2146.	2.6	9
46	Neurofibromatosis Type 1 Gene Mutation Analysis Using Sequence Capture and High-throughput Sequencing. Acta Dermato-Venereologica, 2014, 94, 663-666.	1.3	8
47	Neurofibromatosis type 1 of the child increases birth weight. American Journal of Medical Genetics, Part A, 2019, 179, 1173-1183.	1.2	6
48	Brooke-Spiegler Syndrome Associated with Ulcerative Rectosigmoiditis. Acta Dermato-Venereologica, 2013, 93, 112-113.	1.3	4
49	Circulating free <scp>DNA</scp> in the plasma of individuals with neurofibromatosis type 1. American Journal of Medical Genetics, Part A, 2021, 185, 1098-1104.	1.2	4
50	Celebrating the 50th Anniversary of ESDR. Journal of Investigative Dermatology, 2020, 140, S145-S146.	0.7	O