Sylvie Fraitag

List of Publications by Year
in descending order

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72	3,939	257450 24	123424 61
papers	citations	h-index	g-index

84	84	84	5612
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Diagnostic performance of artificial intelligence for histologic melanoma recognition compared to 18 international expert pathologists. Journal of the American Academy of Dermatology, 2022, 86, 640-642.	1.2	35
2	Rubella vaccineâ€“induced granulomas are a novel phenotype with incomplete penetrance of genetic defects in cytotoxicity. Journal of Allergy and Clinical Immunology, 2022, 149, 388-399.e4.	2.9	11
3	Agminated Spitz naevus with an activating HRAS Q61R mutation. Pathology, 2022, 54, 374-376.	0.6	4
4	Establishing diagnostic criteria for mastocytosis in skin biopsies. Histopathology, 2022, 80, 501-514.	2.9	9
5	ALK-positiveÂhistiocytosis: a new clinicopathologic spectrum highlighting neurologic involvement and responses to ALK inhibition. Blood, 2022, 139, 256-280.	1.4	60
6	RASGRF1-rearranged Cutaneous Melanocytic Neoplasms With Spitzoid Cytomorphology. American Journal of Surgical Pathology, 2022, 46, 655-663.	3.7	8
7	When extended genetics rescues diagnosis: a patient with CANDLE-like phenotype and de novo mutation in theSAMD9Lgene. Annals of the Rheumatic Diseases, 2022, 81, 447-448.	0.9	0
8	Erythroderma in a neonate. JAAD Case Reports, 2022, 21, 97-100.	0.8	0
9	Bone kaposiform hemangioendothelioma: A rare entity dramatically improved by sirolimus. Journal of Paediatrics and Child Health, 2022, , .	0.8	1
10	Cutaneous histiocytoses in children. Histopathology, 2022, 80, 196-215.	2.9	14
11	Malignant Superficial Mesenchymal Tumors in Children. Cancers, 2022, 14, 2160.	3.7	1
12	Eosinophilic Pustular Folliculitis of Infancy: A Histologic Assessment of 43 Cases. American Journal of Dermatopathology, 2022, 44, 395-403.	0.6	2
13	Dermatofibrosarcoma protuberans, fibrosarcomatous variant: A rare tumor in children. Pediatric Dermatology, 2021, 38, 217-222.	0.9	9
14	Uncommon preputial localization of Langerhans cell histiocytosis. Pediatric Dermatology, 2021, 38, 500-501.	0.9	1
15	Necrotizing infundibular (ostial) crystalline folliculitis. Clinics in Dermatology, 2021, 39, 194-198.	1.6	3
16	Major response to imatinib and chemotherapy in a newborn patient prenatally diagnosed with generalized infantile myofibromatosis. Pediatric Blood and Cancer, 2021, 68, e28576.	1.5	8
17	Novel COL4A1â€“VEGFD gene fusion in myofibroma. Journal of Cellular and Molecular Medicine, 2021, 25, 4387-4394.	3.6	8
18	Clinical spectrum of MTOR-related hypomelanosis of Ito with neurodevelopmental abnormalities. Genetics in Medicine, 2021, 23, 1484-1491.	2.4	14

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19	Case â€“ Penile verrucous epithelial hyperplasia. Canadian Urological Association Journal, 2021, 15, E614-E616.	0.6	0
20	Humans with inherited TÂcell CD28 deficiency are susceptible to skin papillomaviruses but are otherwise healthy. Cell, 2021, 184, 3812-3828.e30.	28.9	53
21	Histiocytosis. Lancet, The, 2021, 398, 157-170.	13.7	58
22	Update on Superficial Spindle Cell Mesenchymal Tumors in Children. Dermatopathology (Basel,) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 62	1.5	4
23	Pseudomalignancies in Children: Histological Clues, and Pitfalls to Be Avoided. Dermatopathology (Basel, Switzerland), 2021, 8, 376-389.	1.5	0
24	What to Look Out for in a Newborn with Multiple Papulonodular Skin Lesions at Birth. Dermatopathology (Basel, Switzerland), 2021, 8, 390-417.	1.5	4
25	The association of Greig syndrome and mastocytosis reveals the involvement of the hedgehog pathway in advanced mastocytosis. Blood, 2021, 138, 2396-2407.	1.4	5
26	Skin cancer classification via convolutional neural networks: systematic review of studies involving human experts. European Journal of Cancer, 2021, 156, 202-216.	2.8	115
27	New Insights in Paediatric Dermatopathology. Dermatopathology (Basel, Switzerland), 2021, 8, 531-534.	1.5	0
28	Soft tissue angiomatosis: another PIK3CA â€related disorder. Histopathology, 2020, 76, 540-549.	2.9	12
29	Low Prevalence of GSC Gene Mutations in a Large Cohort of Predominantly Caucasian Patients with Hidradenitis Suppurativa. Journal of Investigative Dermatology, 2020, 140, 2085-2088.e14.	0.7	47
30	Non-congenital dorsal tumefaction with rapid growth in a young child identified as an intramuscular hemangioma. JAAD Case Reports, 2020, 6, 616-618.	0.8	1
31	Cutis marmorata telangiectatica congenitaâ€like lesion with fibrotic appearance. Pediatric Dermatology, 2020, 37, 204-206.	0.9	1
32	Clinical and pathological dermatological features of deficiency of adenosine deaminase 2: A multicenter, retrospective, observational study. Journal of the American Academy of Dermatology, 2020, 83, 1794-1798.	1.2	13
33	HAVCR2 mutations are associated with severe hemophagocytic syndrome in subcutaneous panniculitis-like T-cell lymphoma. Blood, 2020, 135, 1058-1061.	1.4	29
34	A toxic palmitoylation of Cdc42 enhances NF-ÎºB signaling and drives a severe autoinflammatory syndrome. Journal of Allergy and Clinical Immunology, 2020, 146, 1201-1204.e8.	2.9	33
35	Outcome and clinicophenotypical features of acute lymphoblastic leukemia/lymphoblastic lymphoma with cutaneous involvement: A multicenter case series. Journal of the American Academy of Dermatology, 2020, 83, 1166-1170.	1.2	6
36	Local Inhibition of MEK/Akt Prevents Cellular Growth in Human Congenital Melanocytic Nevi. Journal of Investigative Dermatology, 2019, 139, 2004-2015.e13.	0.7	14

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37	Concomitant PIK3CD and TNFRSF9 deficiencies cause chronic active Epstein-Barr virus infection of T cells. Journal of Experimental Medicine, 2019, 216, 2800-2818.	8.5	59
38	Association of PDGFRB Mutations With Pediatric Myofibroma and Myofibromatosis. JAMA Dermatology, 2019, 155, 946.	4.1	43
39	Malignant melanoma withâ€areas ofâ€rhabdomyosarcomatousâ€differentiation arising in a giant congenital nevus with RAF1 gene fusion. Pigment Cell and Melanoma Research, 2019, 32, 708-713.	3.3	22
40	Langerhans Cell Histiocytoma: A Benign Histiocytic Neoplasm of Diverse Lines of Terminal Differentiation. American Journal of Dermatopathology, 2019, 41, 29-36.	0.6	12
41	Novel KHDRBS1-NTRK3 rearrangement in a congenital pediatric CD34-positive skin tumor: a case report. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2019, 474, 111-115.	2.8	15
42	Highly sensitive methods are required to detect mutations in histiocytoses. Haematologica, 2019, 104, e97-e99.	3.5	27
43	Self-healing juvenile cutaneous mucinosis: Clinical and histopathologic findings of 9 patients. Journal of the American Academy of Dermatology, 2018, 78, 1164-1170.	1.2	24
44	Necrotizing Infundibular Crystalline Folliculitis: An Unusual Clinical Presentation and Demonstration of the Presence of Calcium Palmitate. American Journal of Dermatopathology, 2018, 40, e9-e11.	0.6	4
45	Severe dermatophytosis in solid organ transplant recipients: A French retrospective series and literature review. Transplant Infectious Disease, 2018, 20, e12799.	1.7	44
46	Verrucous hemangioma (also known as verrucous venous malformation): A vascular anomaly frequently misdiagnosed as a lymphatic malformation. Pediatric Dermatology, 2018, 35, e378-e381.	0.9	16
47	A recessive form of hyper-IgE syndrome by disruption of ZNF341-dependent STAT3 transcription and activity. Science Immunology, 2018, 3, .	11.9	132
48	Pharmacological modulators of autophagy activate a parallel noncanonical pathway driving unconventional LC3 lipidation. Autophagy, 2017, 13, 854-867.	9.1	122
49	Clinical and pathological significance of cutaneous manifestations in ANCA-associated vasculitides. Autoimmunity Reviews, 2017, 16, 1138-1146.	5.8	32
50	New somatic BRAF splicing mutation in Langerhans cell histiocytosis. Molecular Cancer, 2017, 16, 115.	19.2	37
51	BRAF Mutation Correlates With High-Risk Langerhans Cell Histiocytosis and Increased Resistance to First-Line Therapy. Journal of Clinical Oncology, 2016, 34, 3023-3030.	1.6	233
52	Varying proliferative and clonogenic potential in <scp>NRAS</scp>â€mutated congenital melanocytic nevi according to size. Experimental Dermatology, 2016, 25, 789-796.	2.9	12
53	Langerhans cell histiocytosis: therapeutic strategy and outcome in a 30â€year nationwide cohort of 1478 patients under 18Âyears of age. British Journal of Haematology, 2016, 174, 887-898.	2.5	83
54	Pseudoangiomatous xanthelasmoid mastocytosis: two case reports showing the hypervascularity of this rare variant of cutaneous mastocytosis. Journal of Cutaneous Pathology, 2016, 43, 388-393.	1.3	4

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55	Exocrine pancreatic insufficiency in a child with Netherton syndrome. European Journal of Dermatology, 2016, 26, 311-312.	0.6	5
56	Revised classification of histiocytoses and neoplasms of the macrophage-dendritic cell lineages. Blood, 2016, 127, 2672-2681.	1.4	1,040
57	Cutaneous malignant melanoma in children and adolescents treated in pediatric oncology units. Pediatric Blood and Cancer, 2016, 63, 1922-1927.	1.5	21
58	5-Hydroxymethylcytosine Expression in Proliferative Nodules Arising within Congenital Nevi Allows Differentiation from Malignant Melanoma. Journal of Investigative Dermatology, 2016, 136, 2453-2461.	0.7	22
59	Proliferative Nodules vs Melanoma Arising in Giant Congenital Melanocytic Nevi During Childhood. JAMA Dermatology, 2016, 152, 1147.	4.1	21
60	Dual T cellâ€“ and B cellâ€“intrinsic deficiency in humans with biallelic RLTPR mutations. Journal of Experimental Medicine, 2016, 213, 2413-2435.	8.5	117
61	Telangiectasia macularis eruptiva perstans (TMEP): A form of cutaneous mastocytosis with potential systemic involvement. Journal of the American Academy of Dermatology, 2016, 74, 885-891.e1.	1.2	30
62	Anal manifestation of sarcoidosis. Presse Medicale, 2016, 45, 146-147.	1.9	3
63	Efficacy of colchicine in a child with relapsing bullous Henoch-SchÃ¶nlein purpura. European Journal of Pediatrics, 2016, 175, 147-149.	2.7	15
64	Melanoma in xeroderma pigmentosum type C children: Overrepresentation of desmoplastic type?. Journal of the American Academy of Dermatology, 2015, 72, e173-e176.	1.2	9
65	Assessment and effective targeting of Interleukin-1 in multicentric reticulohistyocytosis. Joint Bone Spine, 2015, 82, 280-283.	1.6	19
66	Langerhans cell histiocytosis in children: Correlation of BRAF status with clinical characteristic.. Journal of Clinical Oncology, 2015, 33, 10003-10003.	1.6	0
67	A NewTRPV3Missense Mutation in a Patient With Olmsted Syndrome and Erythromelalgia. JAMA Dermatology, 2014, 150, 303.	4.1	51
68	Infantile myofibromatosis: A series of 28 cases. Journal of the American Academy of Dermatology, 2014, 71, 264-270.	1.2	89
69	Deep Dermatophytosis and Inherited CARD9 Deficiency. New England Journal of Medicine, 2013, 369, 1704-1714.	27.0	362
70	High prevalence of BRAF V600E mutations in Erdheim-Chester disease but not in other non-Langerhans cell histiocytoses. Blood, 2012, 120, 2700-2703.	1.4	589
71	Long Term Efficacy and Safety of Cladribine In Adult Systemic mastocytosis: a French Multicenter Study of 44 Patients. Blood, 2010, 116, 1982-1982.	1.4	12
72	Skin biopsy is helpful for the diagnosis of incontinentia pigmenti at late stage (IV): a series of 26 cutaneous biopsies. Journal of Cutaneous Pathology, 2009, 36, 966-971.	1.3	29