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## List of Publications by Year in descending order

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Version: 2024-02-01

50  
papers

612  
citations

758635

12  
h-index

610482

24  
g-index

55  
all docs

55  
docs citations

55  
times ranked

806  
citing authors

#	ARTICLE	IF	CITATIONS
1	Cutaneous vascular anomalies associated with a mosaic variant of AKT3: Genetic analysis continues to refine the diagnosis, nomenclature, and classification of vascular anomalies. <i>Journal of the American Academy of Dermatology</i> , 2022, 87, 162-164.	0.6	7
2	A retrospective analysis of diagnostic testing in a large North American cohort of patients with epidermolysis bullosa. <i>Journal of the American Academy of Dermatology</i> , 2021, . .	0.6	3
3	An erythematous patch on the ear and face of a young boy. <i>Pediatric Dermatology</i> , 2021, 38, e45-e47.	0.5	0
4	A pediatric case of pigmented epithelioid melanocytoma with chromosomal copy number alterations in 15q and 17q and a novel <i>NTRK3</i> gene fusion. <i>Journal of Cutaneous Pathology</i> , 2020, 47, 70-75.	0.7	9
5	Congenital midline nodules on the chin and sternum. <i>Pediatric Dermatology</i> , 2020, 37, 1145-1146.	0.5	0
6	Proliferative nodule resembling angiomatoid Spitz tumor with degenerative atypia arising within a giant congenital nevus. <i>Journal of Cutaneous Pathology</i> , 2020, 47, 1200-1204.	0.7	1
7	Melanoma arising in a patient with ataxia-telangiectasia: A call for full skin examinations in this patient population. <i>Pediatric Dermatology</i> , 2020, 37, 767-768.	0.5	2
8	Retinal Vascular Abnormalities in Phakomatosis Pigmentovascularis. <i>Ophthalmology Retina</i> , 2019, 3, 1098-1104.	1.2	6
9	ABCA 12 homozygous mutation in harlequin ichthyosis: Survival without systemic retinoids. <i>Pediatric Dermatology</i> , 2019, 36, 339-341.	0.5	3
10	Assessment of the Timing of Milestone Clinical Events in Patients With Epidermolysis Bullosa From North America. <i>JAMA Dermatology</i> , 2019, 155, 196.	2.0	27
11	Large hemorrhagic plaque with central crusting. <i>Cutis</i> , 2019, 103, 68;79;80.	0.4	0
12	Managing sleep disturbances in children with atopic dermatitis. <i>Pediatric Dermatology</i> , 2018, 35, 428-433.	0.5	9
13	Asymptomatic bands on an infant's extremity. <i>Pediatric Dermatology</i> , 2018, 35, 139-140.	0.5	3
14	Cantharidin for treatment of facial molluscum contagiosum: A retrospective review. <i>Journal of the American Academy of Dermatology</i> , 2018, 78, 198-200.	0.6	20
15	Poikiloderma with neutropenia and associated squamous cell carcinoma: A case report. <i>Pediatric Dermatology</i> , 2018, 35, e366-e367.	0.5	3
16	Cutaneous manifestations of metastatic Crohn's disease. <i>Pediatric Dermatology</i> , 2018, 35, 566-574.	0.5	40
17	Gotttron papules mimicking dermatomyositis: an unusual manifestation of systemic lupus erythematosus. <i>Cutis</i> , 2018, 102, E16-E18.	0.4	2
18	Scalp-to-toes application of permethrin for patients with scabies. <i>Dermatology Online Journal</i> , 2018, 24, .	0.2	0

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19	•Marks the Spot: An Injection Technique for Alopecia Areata. <i>Pediatric Dermatology</i> , 2017, 34, 214-215.	0.5	0
20	Punch and Scoop Technique for Removing Pilomatricoma. <i>Pediatric Dermatology</i> , 2017, 34, 622-623.	0.5	2
21	Teenage boy with thickened dorsal hands and feet. <i>Pediatric Dermatology</i> , 2017, 34, 719-720.	0.5	3
22	Transient Porphyrinemia in a Neonate: A Case Report. <i>Pediatric Dermatology</i> , 2016, 33, e375-e376.	0.5	3
23	Lymphoplasmacytic Plaque in Children: A Demonstrative Case of an Emerging Clinicopathologic Entity. <i>Pediatric Dermatology</i> , 2016, 33, e349-e350.	0.5	1
24	PHACE Syndrome: Consensus-Derived Diagnosis and Care Recommendations. <i>Journal of Pediatrics</i> , 2016, 178, 24-33.e2.	0.9	186
25	Congenital Ichthyosiform Erythroderma Superimposed with Chronic Dermatophytosis: A Report of Three Siblings. <i>Pediatric Dermatology</i> , 2016, 33, e6-9.	0.5	9
26	Photodermatoses: Kids are not just little people. <i>Clinics in Dermatology</i> , 2016, 34, 724-735.	0.8	8
27	Melanoma Gene Expression Markers for Surveillance of Epidermolysis Bullosa Nevi Malignant Transformation. <i>JAMA Dermatology</i> , 2016, 152, 584.	2.0	1
28	Development of Sinus Tracts within a Connective Tissue Nevus. <i>Pediatric Dermatology</i> , 2015, 32, e298-9.	0.5	0
29	Suspected rapid•onset neutropenia following etanercept use: disproved following rechallenge and lessons to be learned. <i>International Journal of Dermatology</i> , 2013, 52, 1287-1288.	0.5	1
30	Multiple pigmented eccrine hidrocystomas within scars in a patient with recessive dystrophic epidermolysis bullosa. <i>Journal of the American Academy of Dermatology</i> , 2013, 69, e245-e246.	0.6	2
31	Dermatological Complications of Circumcision: Lesson Learned from Cases In a Pediatric Dermatology Practice. <i>Pediatric Dermatology</i> , 2013, 30, 519-528.	0.5	9
32	Bullous Dermolysis of the Newborn: Four New Cases and Clinical Review. <i>Pediatric Dermatology</i> , 2013, 30, 736-740.	0.5	8
33	Allogeneic Hematopoietic Cell Transplantation Is Effective In Patients With Advanced Systemic Mastocytosis: A Multicenter Retrospective Analysis. <i>Blood</i> , 2013, 122, 2145-2145.	0.6	0
34	Poikiloderma with Neutropenia: Report of Three Cases Including One with Calcinosis Cutis. <i>Pediatric Dermatology</i> , 2012, 29, 463-472.	0.5	22
35	An Adolescent Boy with Persistent Penile and Scrotal Erythema and Swelling. <i>Pediatric Dermatology</i> , 2012, 29, 765-766.	0.5	5
36	A young boy with bullae. <i>Journal of the American Academy of Dermatology</i> , 2011, 65, e71.	0.6	0

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37	Phacomatosis Pigmentokeratolica: A Further Case without Extracutaneous Anomalies and Review of the Condition. <i>Pediatric Dermatology</i> , 2011, 28, 715-719.	0.5	17
38	Identification of a novel <i>C16orf57</i> mutation in Athabaskan patients with Poikiloderma with Neutropenia. <i>American Journal of Medical Genetics, Part A</i> , 2011, 155, 337-342.	0.7	34
39	Neonatal skin barrier: structure, function, and disorders. <i>Dermatologic Therapy</i> , 2005, 18, 87-103.	0.8	79
40	Disorders of Keratinization. <i>American Journal of Clinical Dermatology</i> , 2004, 5, 17-29.	3.3	39
41	Herlitz Junctional Epidermolysis Bullosa Presenting at Birth with Anonychia: A Case Report and Review of H&EB. <i>Pediatric Dermatology</i> , 2001, 18, 217-222.	0.5	13
42	Congenital Ichthyosiform Erythroderma: Particulate Staining Pattern of TCK. <i>Journal of Dermatology</i> , 1999, 26, 791-796.	0.6	0
43	Congenital CD34-positive granular cell dendrocytosis. <i>Journal of Cutaneous Pathology</i> , 1999, 26, 253-258.	0.7	15
44	Generalized Atrophic Benign Epidermolysis Bullosa: A Case of Severe Hemidesmosomal Deficiency. <i>Journal of Dermatology</i> , 1999, 26, 512-517.	0.6	1
45	Ichthyosis in a Nutshell. <i>Pediatrics in Review</i> , 1999, 20, 5-8.	0.2	1
46	What Syndrome Is This?. <i>Pediatric Dermatology</i> , 1997, 14, 239-240.	0.5	4
47	Netherton Syndrome Presenting as Congenital Psoriasis. <i>Pediatric Dermatology</i> , 1997, 14, 473-476.	0.5	12
48	Nodule on the Forearm in an Adolescent. <i>Pediatric Dermatology</i> , 1989, 6, 53-54.	0.5	0
49	Symptem on Alopecia Areata. <i>Pediatric Dermatology</i> , 1988, 5, 69-69.	0.5	0
50	Microcystic lymphatic malformation presenting as firm, skin-colored papules of the lips. <i>Pediatric Dermatology</i> , 0, , .	0.5	1