Andrea Diociaiuti

List of Publications by Year
in descending order

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105	1,794	304743	330143
papers	citations	22 h-index	37 g-index

107	107	107	2289
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Multicentre consensus recommendations for skin care in inherited epidermolysis bullosa. Orphanet Journal of Rare Diseases, 2014, 9, 76.	2.7	124
2	A clinical, histopathological and laboratory study of 19 consecutive Italian paediatric patients with chilblainâ€like lesions: lights and shadows on the relationship with COVIDâ€19 infection. Journal of the European Academy of Dermatology and Venereology, 2020, 34, 2620-2629.	2.4	121
3	RISK OF NONMELANOMA SKIN CANCER IN ITALIAN ORGAN TRANSPLANT RECIPIENTS. A REGISTRY-BASED STUDY. Transplantation, 2000, 70, 1479-1484.	1.0	104
4	Efficacy and Adverse Events During Janus Kinase Inhibitor Treatment of SAVI Syndrome. Journal of Clinical Immunology, 2019, 39, 476-485.	3.8	85
5	Management of congenital ichthyoses: European guidelines of care, part one. British Journal of Dermatology, 2019, 180, 272-281.	1.5	70
6	Management of congenital ichthyoses: European guidelines of care, part two. British Journal of Dermatology, 2019, 180, 484-495.	1.5	68
7	Consensus Conference on Clinical Management of pediatric Atopic Dermatitis. Italian Journal of Pediatrics, 2016, 42, 26.	2.6	67
8	Management of chronic urticaria in children: a clinical guideline. Italian Journal of Pediatrics, 2019, 45, 101.	2.6	63
9	Transient cutaneous manifestations after administration of Pfizerâ€BioNTech COVIDâ€19 Vaccine: an Italian singleâ€centre case series. Journal of the European Academy of Dermatology and Venereology, 2021, 35, e483-e485.	2.4	62
10	Skin manifestations of COVIDâ€19 in children: Part 1. Clinical and Experimental Dermatology, 2021, 46, 444-450.	1.3	61
11	Topical corticosteroid phobia in parents of pediatric patients with atopic dermatitis: a multicentre survey. Italian Journal of Pediatrics, 2017, 43, 22.	2.6	48
12	Mosaic abnormalities of the skin: review and guidelines from the European Reference Network for rare skin diseases. British Journal of Dermatology, 2020, 182, 552-563.	1.5	45
13	Kaposiâ€™s Sarcoma in Renal-Transplant Recipients: Experience at the Catholic University in Rome, 1988â€“1996. Dermatology, 1997, 194, 229-233.	2.1	44
14	Epidermolysis bullosa simplex withPLECmutations: new phenotypes and new mutations. British Journal of Dermatology, 2013, 168, 808-814.	1.5	44
15	Ichthyosis with confetti: clinics, molecular genetics and management. Orphanet Journal of Rare Diseases, 2015, 10, 115.	2.7	30
16	Common Community-acquired Bacterial Skin and Soft-tissue Infections in Children: an Intersociety Consensus on Impetigo, Abscess, and Cellulitis Treatment. Clinical Therapeutics, 2019, 41, 532-551.e17.	2.5	30
17	Congenital Lupus Erythematosus: Case Report and Review of the Literature. Pediatric Dermatology, 2005, 22, 240-242.	0.9	28
18	Phakomatosis pigmentovascularis type IIIb: A case associated with Sturge-Weber and Klippel-Trenaunay syndromes. Journal of the American Academy of Dermatology, 2005, 53, 535-538.	1.2	28

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19	Xâ€linked ichthyosis: Clinical and molecular findings in 35 Italian patients. Experimental Dermatology, 2019, 28, 1156-1163.	2.9	28
20	Skin manifestations of COVIDâ€19 in children: Part 3. Clinical and Experimental Dermatology, 2021, 46, 462-472.	1.3	27
21	Relapsing polychondritis: new therapeutic strategies with biological agents. Rheumatology International, 2010, 30, 691-693.	3.0	25
22	Unusual Father-to-Daughter Transmission of Incontinentia Pigmenti Due to Mosaicism in IP Males. Pediatrics, 2017, 140, .	2.1	24
23	Pretransplantation human herpes virus 8 seropositivity as a risk factor for Kaposiâ€™s sarcoma in kidney transplant recipients. Transplantation Proceedings, 2000, 32, 526-527.	0.6	23
24	HHV8 in renal transplant recipients. Transplant International, 2000, 13, S410-S412.	1.6	21
25	Role of molecular testing in the multidisciplinary diagnostic approach of ichthyosis. Orphanet Journal of Rare Diseases, 2016, 11, 4.	2.7	21
26	Multidisciplinary consensus recommendations from a European network for the diagnosis and practical management of patients with incontinentia pigmenti. Journal of the European Academy of Dermatology and Venereology, 2020, 34, 1415-1424.	2.4	20
27	Early Immunopathological Diagnosis of Ichthyosis with Confetti in Two Sporadic Cases with New Mutations in Keratin 10. Acta Dermato-Venereologica, 2014, 94, 579-582.	1.3	17
28	Lethal Netherton Syndrome Due to Homozygous p.<scp>A</scp>rg371<scp>X</scp> Mutation in <scp>SPINK</scp>5. Pediatric Dermatology, 2013, 30, e65-7.	0.9	15
29	Long-term Follow-up of a Spontaneously Improving Patient with Junctional Epidermolysis Bullosa Associated with ITGB4 c.3977-19T>A Splicing Mutation. Acta Dermato-Venereologica, 2013, 93, 116-118.	1.3	14
30	A Case of Neonatal Linear IgA Bullous Dermatosis with Severe Eye Involvement. Acta Dermato-Venereologica, 2015, 95, 1015-1017.	1.3	14
31	The Role of KRAS Mutations in Cortical Malformation and Epilepsy Surgery: A Novel Report of Nevus Sebaceous Syndrome and Review of the Literature. Brain Sciences, 2021, 11, 793.	2.3	14
32	Safety and effectiveness of oral propranolol for infantile hemangiomas started before 5Âweeks and after 5Âmonths of age: an Italian multicenter experience. Italian Journal of Pediatrics, 2017, 43, 40.	2.6	13
33	Are SARSâ€CoVâ€2 IgA antibodies in paediatric patients with chilblainâ€like lesions indicative of COVIDâ€19 asymptomatic or paucisymptomatic infection?. Journal of the European Academy of Dermatology and Venereology, 2021, 35, e10-e13.	2.4	13
34	T-lymphocytes are Directly Involved in the Clinical Expression of Migratory Circinate Erythema in Epidermolysis Bullosa Simplex Patients. Acta Dermato-Venereologica, 2014, 94, 307-311.	1.3	12
35	A truncating mutation in the laminin-332Î± chain highlights the role of the LG45 proteolytic domain in regulating keratinocyte adhesion and migration. British Journal of Dermatology, 2014, 170, 1056-1064.	1.5	11
36	Kindler syndrome with severe mucosal involvement in a large Palestinian pedigree. European Journal of Dermatology, 2015, 25, 14-19.	0.6	11

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37	Cutaneous Crohn disease mimicking anal condylomata in a child. Journal of the American Academy of Dermatology, 2010, 63, 165-166.	1.2	10
38	Naevoid Basal Cell Carcinoma Syndrome in a 22-month-old Child Presenting with Multiple Basal Cell Carcinomas and a Fetal Rhabdomyoma. Acta Dermato-Venereologica, 2015, 95, 243-244.	1.3	10
39	Cardiopulmonary anomalies in incontinentia pigmenti patients. International Journal of Dermatology, 2018, 57, 40-45.	1.0	10
40	First Case of KRT2 Epidermolytic Nevus and Novel Clinical and Genetic Findings in 26 Italian Patients with Keratinopathic Ichthyoses. International Journal of Molecular Sciences, 2020, 21, 7707.	4.1	10
41	Immunofluorescence mapping, electron microscopy and genetics in the diagnosis and subâ€classification of inherited epidermolysis bullosa: a singleâ€centre retrospective comparative study of 87 cases with longâ€term followâ€up. Journal of the European Academy of Dermatology and Venereology, 2021, 35, 1007-1016.	2.4	10
42	The Burden of Autosomal Recessive Congenital Ichthyoses on Patients and their Families: An Italian Multicentre Study. Acta Dermato-Venereologica, 2021, 101, adv00477.	1.3	10
43	Human Herpesvirus-8 Serology in Pediatric Organ Transplantation. Transplantation Proceedings, 2008, 40, 3683-3684.	0.6	9
44	Multi-drugs resistant acne rosacea in a child affected by Ataxia-Telangiectasia: successful treatment with Isotretinoin. Italian Journal of Pediatrics, 2015, 41, 23.	2.6	9
45	An additional patient with a homozygous mutation in DCPS contributes to the delination of Alâ€Raqad syndrome. American Journal of Medical Genetics, Part A, 2018, 176, 2781-2786.	1.2	9
46	Vascular Anomalies Guidelines by the Italian Society for the study of Vascular Anomalies (SISAV). International Angiology, 2015, 34, 1-45.	0.9	9
47	Increased S100B protein serum levels in psoriasis. Journal of Dermatological Science, 2007, 48, 148-150.	1.9	8
48	Melanoma in a 5-year-old Child with a Giant Congenital Melanocytic Naevus. Acta Dermato-Venereologica, 2012, 92, 607-608.	1.3	8
49	Necrosis of the tongue as first symptom of Polyarteritis Nodosa (PAN): unusual presentation of a rare disease in children. Rheumatology International, 2013, 33, 1071-1073.	3.0	8
50	Electrochemotherapy, a potential new treatment for the management of squamous cell carcinoma in patients with recessive dystrophic epidermolysis bullosa: report of three cases. Journal of the European Academy of Dermatology and Venereology, 2016, 30, 1195-1196.	2.4	8
51	Severe osteoarticular involvement in isotretinoinâ€triggered acne fulminans: two cases successfully treated with anakinra. Journal of the European Academy of Dermatology and Venereology, 2017, 31, e277-e279.	2.4	8
52	Novel clinical features associated with Clouston syndrome. International Journal of Dermatology, 2019, 58, e143-e146.	1.0	8
53	Vascular Birthmarks as a Clue for Complex and Syndromic Vascular Anomalies. Frontiers in Pediatrics, 2021, 9, 730393.	1.9	8
54	Herpetic Whitlow as a Harbinger of Pediatric HIV-1 Infection. Pediatric Dermatology, 2005, 22, 119-121.	0.9	7

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55	Ichthyosis Prematurity Syndrome due to a Novel SLC27A4 Homozygous Mutation in an Italian Patient. Acta Dermato-Venereologica, 2018, 98, 803-804.	1.3	7
56	Phenotypic Features of Epidermolysis Bullosa Simplex due to KLHL24 Mutations in 3 Italian Cases. Acta Dermato-Venereologica, 2019, 99, 238-239.	1.3	7
57	Two Italian Patients with ELOVL4-Related Neuro-Ichthyosis: ÂExpanding the Genotypic and Phenotypic Spectrum and Ultrastructural Characterization. Genes, 2021, 12, 343.	2.4	7
58	Proposal for a 6â€step approach for differential diagnosis of neonatal erythroderma. Journal of the European Academy of Dermatology and Venereology, 2022, 36, 973-986.	2.4	7
59	Multidisciplinary care for patients with epidermolysis bullosa from birth to adolescence: experience of one Italian reference center. Italian Journal of Pediatrics, 2022, 48, 58.	2.6	7
60	Narrative review on the management of moderate-severe atopic dermatitis in pediatric age of the Italian Society of Pediatric Allergology and Immunology (SIAIP), of the Italian Society of Pediatric Dermatology (SIDerP) and of the Italian Society of Pediatrics (SIP). Italian Journal of Pediatrics, 2022, 48, .	2.6	7
61	HHV8 in renal transplant recipients. Transplant International, 2000, 13, S410-S412.	1.6	6
62	Increased S100B serum levels in diffuse dermatitis. Melanoma Research, 2002, 12, 633.	1.2	6
63	Frequent Occurrence of Aplasia Cutis Congenita in Bullous Dermolysis of the Newborn. Acta Dermato-Venereologica, 2014, 96, 784-7.	1.3	6
64	A waterborn zoonotic helminthiase in an Italian diver: a case report of a cutaneousSparganuminfection and a review of European cases. Pathogens and Global Health, 2015, 109, 383-386.	2.3	6
65	Acral skin atrophy in an infant: an early clue to Kindler syndrome diagnosis. Journal of the European Academy of Dermatology and Venereology, 2016, 30, 1046-1049.	2.4	6
66	Novel <scp>PNPLA</scp>1 mutations in two Italian siblings with autosomal recessive congenital ichthyosis. Journal of the European Academy of Dermatology and Venereology, 2018, 32, e110-e112.	2.4	6
67	Autosomal recessive epidermolysis bullosa simplex due to EXPH5 mutation: neonatal diagnosis of the first Italian case and literature review. Journal of the European Academy of Dermatology and Venereology, 2020, 34, e694-e697.	2.4	6
68	A multicenter study on quality of life of the â€œgreater patientâ€•in congenital ichthyoses. Orphanet Journal of Rare Diseases, 2021, 16, 440.	2.7	6
69	Structural Defects of Laminin Î²3 N-terminus Underlie Junctional Epidermolysis Bullosa with Altered Granulation Tissue Response. Acta Dermato-Venereologica, 2016, 96, 954-958.	1.3	5
70	IgA tracheobronchial deposits underlie respiratory compromise in neonatal linear IgA bullous dermatosis. Journal of the European Academy of Dermatology and Venereology, 2017, 31, e333-e335.	2.4	5
71	Angioma serpiginosum: a case report and review of the literature. Italian Journal of Pediatrics, 2019, 45, 53.	2.6	5
72	Italian translation, cultural adaptation, and pilot testing of a questionnaire to assess family burden in inherited ichthyoses. Italian Journal of Pediatrics, 2019, 45, 26.	2.6	5

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73	Cutaneous Infantile Haemangiomas with Intracranial and Intraspinal Involvement: A European Multicentre Experience and Review. Acta Dermato-Venereologica, 2020, 100, adv00255.	1.3	5
74	Clinical and Molecular Spectrum of Sporadic Vascular Malformations: A Single-Center Study. Biomedicines, 2022, 10, 1460.	3.2	5
75	Congenital Myxoid and Pigmented Dermatofibrosarcoma Protuberans: A Case Report. Pediatric Dermatology, 2013, 30, e74-7.	0.9	4
76	Unclassifiable pattern of hypopigmentation in a patient with mosaic partial 12p tetrasomy without Pallisterâ€“Killian syndrome. American Journal of Medical Genetics, Part A, 2017, 173, 1943-1946.	1.2	4
77	Local anesthesia in pediatric dermatologic surgery: Evaluation of a patientâ€centered approach. Pediatric Dermatology, 2018, 35, 112-116.	0.9	4
78	A novel dermoscopic pattern observed in furuncular myiasis. Australasian Journal of Dermatology, 2019, 60, e46-e47.	0.7	4
79	Glanular Venous Malformation in Pediatric Age: An Uncommon Vascular Disorder. Urology, 2019, 131, 223-227.	1.0	4
80	Juvenile idiopathic arthritis in infants with Harlequin Ichthyosis: two cases report and literature review. Italian Journal of Pediatrics, 2020, 46, 44.	2.6	4
81	Proteasome-mediated degradation of keratins 7, 8, 17 and 18 by mutant KLHL24 in a foetal keratinocyte model: Novel insight in congenital skin defects and fragility of epidermolysis bullosa simplex with cardiomyopathy. Human Molecular Genetics, 2022, 31, 1308-1324.	2.9	4
82	Management of Upper Airway Infantile Hemangiomas: Experience of One Italian Multidisciplinary Center. Frontiers in Pediatrics, 2021, 9, 717232.	1.9	4
83	Autosomal recessive epidermolysis bullosa simplex due to <scp>KRT</scp>14 mutation: two large Palestinian families and literature review. Journal of the European Academy of Dermatology and Venereology, 2018, 32, e149-e151.	2.4	3
84	Agminated Spitzoid Naevi after Remission of Langerhans Cell Histiocytosis: First Italian Case and Literature Review. Acta Dermato-Venereologica, 2019, 99, 824-825.	1.3	3
85	Next Generation Sequencing Uncovers a Rare Case of X-linked Ichthyosis in an Adopted Girl Homozygous for a Novel Nonsense Mutation in the STS Gene. Acta Dermato-Venereologica, 2019, 99, 828-830.	1.3	3
86	Ectodermal Dysplasia-Syndactyly Syndrome with Toe-Only Minimal Syndactyly Due to a Novel Mutation in NECTIN4: A Case Report and Literature Review. Genes, 2021, 12, 748.	2.4	3
87	Self-improving dominant dystrophic epidermolysis bullosa: phenotypic variability associated with COL7A1 mutation p.Gly2037Glu. European Journal of Dermatology, 2020, 30, 753-754.	0.6	3
88	PIK3CA-related overgrowth with an uncommon phenotype: case report. Italian Journal of Pediatrics, 2022, 48, 71.	2.6	3
89	Cost-utility analysis of propranolol versus corticosteroids in the treatment of proliferating infantile hemangioma in Italy. Pharmacoeconomics Italian Research Articles, 2015, 17, 1.	0.2	2
90	Borderline cognitive level in a family with Bazexâ€ÐuprÃ©â€Christol syndrome. American Journal of Medical Genetics, Part A, 2015, 167, 1637-1643.	1.2	2

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91	Lipoid Proteinosis: A Previously Unrecognized Mutation and Therapeutic Response to Acitretin. Acta Dermato-Venereologica, 2017, 97, 1249-1251.	1.3	2
92	Teledermatology diagnosis of the first Italian patient affected with restrictive dermopathy due to <scp>ZMPSTE</scp>24 homozygous mutation. Journal of the European Academy of Dermatology and Venereology, 2019, 33, e139-e140.	2.4	2
93	Facial comedonal acne in orofaciodigital syndrome type 1 caused by a novel frameshift variant in <scp>OFD</scp> 1 . Clinical and Experimental Dermatology, 2019, 44, 706-708.	1.3	2
94	Epidermolysis Bullosa in children: the central role of the pediatrician. Orphanet Journal of Rare Diseases, 2022, 17, 147.	2.7	2
95	Cost-Utility Analysis Comparing Propranolol With Corticosteroids in the Treatment of Proliferating Infantile Hemangioma in Italy. Value in Health, 2014, 17, A511.	0.3	1
96	Buschke-Ollendorff syndrome. British Journal of Dermatology, 2016, 174, 709-710.	1.5	1
97	Two novel mutations in the <scp>TSC</scp>2 gene causing severe phenotype in nervous system and skin in a patient with tuberous sclerosis complex. Journal of the European Academy of Dermatology and Venereology, 2018, 32, e243-e245.	2.4	1
98	Hoarse cry in a newborn with epidermolysis bullosa simplex, generalized severe. Pediatric Dermatology, 2020, 37, 393-395.	0.9	1
99	Acute generalized exanthematous pustulosis associated with Epstein-Barr virus infection reactivation. Giornale Italiano Di Dermatologia E Venereologia, 2019, 154, 91-92.	0.8	1
100	Prominent Follicular Keratosis in Multiple Intestinal Atresia with Combined Immune Deficiency Caused by a TTC7A Homozygous Mutation. Genes, 2022, 13, 821.	2.4	1
101	The VASCERN-VASCA working group diagnostic and management pathways for severe and/or rare infantile hemangiomas. European Journal of Medical Genetics, 2022, 65, 104517.	1.3	1
102	Auricular leishmaniasis in a child successfully treated with intralesional amphotericin B. Pediatric Dermatology, 0, , .	0.9	1
103	Fibrolipomatous hamartomas not only on the soles. Pediatric Dermatology, 2019, 36, 728-729.	0.9	0
104	A rare case of segmental neurofibromatosis with multiple blue-red pseudoatrophic plaques. Cutis, 2014, 94, 149-52.	0.3	0
105	ITGB4-mutated Junctional Epidermolysis Bullosa without Pyloric Atresia Presenting with Severe Urinary Involvement and Late-onset Minimal Skin Fragility: Diagnostic and Therapeutic Challenges. Acta Dermato-Venereologica, 2022, , .	1.3	0