

Paola Fortugno

List of Publications by Year in descending order

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Version: 2024-02-01

40
papers

2,595
citations

394421

19
h-index

302126

39
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41
all docs

41
docs citations

41
times ranked

3603
citing authors

#	ARTICLE	IF	CITATIONS
1	RIPK4 regulates cell-cell adhesion in epidermal development and homeostasis. <i>Human Molecular Genetics</i> , 2022, , .	2.9	1
2	A Novel Phenotype of Junctional Epidermolysis Bullosa with Transient Skin Fragility and Predominant Ocular Involvement Responsive to Human Amniotic Membrane Eyedrops. <i>Genes</i> , 2021, 12, 716.	2.4	5
3	Multiple Skin Squamous Cell Carcinomas in Junctional Epidermolysis Bullosa Due to Altered Laminin-332 Function. <i>International Journal of Molecular Sciences</i> , 2020, 21, 1426.	4.1	3
4	Measles skin rash: Infection of lymphoid and myeloid cells in the dermis precedes viral dissemination to the epidermis. <i>PLoS Pathogens</i> , 2020, 16, e1008253.	4.7	13
5	Recessive mutations in the neuronal isoforms of <i>DST</i> , encoding dystonin, lead to abnormal actin cytoskeleton organization and HSAN type VI. <i>Human Mutation</i> , 2019, 40, 106-114.	2.5	30
6	Microprocessor-dependent processing of splice site overlapping microRNA exons does not result in changes in alternative splicing. <i>Rna</i> , 2018, 24, 1158-1171.	3.5	12
7	A compound synonymous mutation c.474C>A with p.Arg578X mutation in <i>SPINK5</i> causes splicing disorder and mild phenotype in Netherton syndrome. <i>Experimental Dermatology</i> , 2016, 25, 568-570.	2.9	6
8	Ichthyosis Linearis Circumflexa as the Only Clinical Manifestation of Netherton Syndrome. <i>Acta Dermato-Venereologica</i> , 2015, 95, 720-724.	1.3	8
9	Betapapillomavirus in multiple non-melanoma skin cancers of Netherton syndrome: Case report and published work review. <i>Journal of Dermatology</i> , 2015, 42, 786-794.	1.2	15
10	Whole-exome sequencing in patients with ichthyosis reveals modifiers associated with increased IgE levels and allergic sensitizations. <i>Journal of Allergy and Clinical Immunology</i> , 2015, 135, 280-283.e15.	2.9	9
11	Reference genes for gene expression analysis in proliferating and differentiating human keratinocytes. <i>Experimental Dermatology</i> , 2015, 24, 314-316.	2.9	13
12	Kindler syndrome with severe mucosal involvement in a large Palestinian pedigree. <i>European Journal of Dermatology</i> , 2015, 25, 14-19.	0.6	11
13	TFIIH-dependent <i>MMP-1</i> overexpression in trichothiodystrophy leads to extracellular matrix alterations in patient skin. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2015, 112, 1499-1504.	7.1	282
14	Exon-Specific U1s Correct <i>SPINK5</i> Exon 11 Skipping Caused by a Synonymous Substitution that Affects a Bifunctional Splicing Regulatory Element. <i>Human Mutation</i> , 2015, 36, 504-512.	2.5	33
15	p63-dependent and independent mechanisms of nectin1 and nectin4 regulation in the epidermis. <i>Experimental Dermatology</i> , 2015, 24, 114-119.	2.9	25
16	Early Immunopathological Diagnosis of Ichthyosis with Confetti in Two Sporadic Cases with New Mutations in Keratin 10. <i>Acta Dermato-Venereologica</i> , 2014, 94, 579-582.	1.3	17
17	A truncating mutation in the laminin-332 chain highlights the role of the LG45 proteolytic domain in regulating keratinocyte adhesion and migration. <i>British Journal of Dermatology</i> , 2014, 170, 1056-1064.	1.5	11
18	Nectin-4 Mutations Causing Ectodermal Dysplasia with Syndactyly Perturb the Rac1 Pathway and the Kinetics of Adherens Junction Formation. <i>Journal of Investigative Dermatology</i> , 2014, 134, 2146-2153.	0.7	33

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19	Lethal Netherton Syndrome Due to Homozygous p.<sc>A</sc>rg371<sc>X</sc> Mutation in <sc>SPINK</sc>5. <i>Pediatric Dermatology</i> , 2013, 30, e65-7.	0.9	15
20	Long-term Follow-up of a Spontaneously Improving Patient with Junctional Epidermolysis Bullosa Associated with ITGB4 c.3977-19T<sc>A Splicing Mutation. <i>Acta Dermato-Venereologica</i> , 2013, 93, 116-118.	1.3	14
21	A synonymous mutation in SPINK5 exon 11 causes Netherton syndrome by altering exonic splicing regulatory elements. <i>Journal of Human Genetics</i> , 2012, 57, 311-315.	2.3	12
22	The 420K LEKTI variant alters LEKTI proteolytic activation and results in protease deregulation: implications for atopic dermatitis. <i>Human Molecular Genetics</i> , 2012, 21, 4187-4200.	2.9	84
23	Full Sequencing of the FLG Gene in Italian Patients with Atopic Eczema: Evidence of New Mutations, but Lack of an Association. <i>Journal of Investigative Dermatology</i> , 2011, 131, 982-984.	0.7	49
24	Proteolytic Activation Cascade of the Netherton Syndromeâ€œDefective Protein, LEKTI, in the Epidermis: Implications for Skin Homeostasis. <i>Journal of Investigative Dermatology</i> , 2011, 131, 2223-2232.	0.7	56
25	Mutations in PVRL4, Encoding Cell Adhesion Molecule Nectin-4, Cause Ectodermal Dysplasia-Syndactyly Syndrome. <i>American Journal of Human Genetics</i> , 2010, 87, 265-273.	6.2	98
26	Intracellular targets of RGDS peptide in melanoma cells. <i>Molecular Cancer</i> , 2010, 9, 84.	19.2	27
27	Chemerin expression marks early psoriatic skin lesions and correlates with plasmacytoid dendritic cell recruitment. <i>Journal of Experimental Medicine</i> , 2009, 206, 249-258.	8.5	268
28	Downregulation of β 6 in keratinocytes by p14ARF-mediated SUMO-conjugation and degradation. <i>Cell Cycle</i> , 2009, 8, 3545-3551.	2.6	28
29	Rational design of shepherdin, a novel anticancer agent. <i>Cancer Cell</i> , 2005, 7, 457-468.	16.8	311
30	Identification of tumor-associated antigens by screening phage-displayed human cDNA libraries with sera from tumor patients. <i>International Journal of Cancer</i> , 2003, 106, 534-544.	5.1	80
31	Regulation of survivin function by Hsp90. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2003, 100, 13791-13796.	7.1	311
32	Antigenicity and immunogenicity of phage library-selected peptide mimics of the major surface proteophosphoglycan antigens of <i>Entamoeba histolytica</i> . <i>Parasite Immunology</i> , 2002, 24, 321-328.	1.5	17
33	Survivin exists in immunochemically distinct subcellular pools and is involved in spindle microtubule function. <i>Journal of Cell Science</i> , 2002, 115, 575-585.	2.0	255
34	Survivin exists in immunochemically distinct subcellular pools and is involved in spindle microtubule function. <i>Journal of Cell Science</i> , 2002, 115, 575-85.	2.0	198
35	ADAM-HCV, a new-concept diagnostic assay for antibodies to hepatitis C virus in serum. <i>FEBS Journal</i> , 2001, 268, 4758-4768.	0.2	10
36	Colony Assay for Phage-Displayed Libraries. <i>Analytical Biochemistry</i> , 2000, 284, 412-415.	2.4	1

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37	Isolation of Phage Mimotopes Mimicking a Protective Epitope of GPI-Linked Proteophosphoglycan Antigens of <i>Entamoeba histolytica</i> . <i>Archives of Medical Research</i> , 2000, 31, S309-S310.	3.3	0
38	“Affinity maturation” of ligands for HCV-specific serum antibodies. <i>Journal of Immunological Methods</i> , 2000, 236, 167-176.	1.4	13
39	Induction of anti-carbohydrate antibodies by phage library-selected peptide mimics. <i>European Journal of Immunology</i> , 1997, 27, 2620-2625.	2.9	108
40	Selection of biologically active peptides by phage display of random peptide libraries. <i>Current Opinion in Biotechnology</i> , 1996, 7, 616-621.	6.6	113