Pierre Wolkenstein

List of Publications by Year in descending order

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184 papers 10,333 citations

³⁸⁷⁴² 50 h-index

97 g-index

198 all docs

198 docs citations

198 times ranked 6968 citing authors

#	Article	IF	CITATIONS
1	SCORTEN: A Severity-of-Illness Score for Toxic Epidermal Necrolysis. Journal of Investigative Dermatology, 2000, 115, 149-153.	0.7	850
2	Prevalence and factors associated with hidradenitis suppurativa: Results from two case-control studies. Journal of the American Academy of Dermatology, 2008, 59, 596-601.	1.2	559
3	Randomised comparison of thalidomide versus placebo in toxic epidermal necrolysis. Lancet, The, 1998, 352, 1586-1589.	13.7	414
4	Repair of the lower and middle parts of the face by composite tissue allotransplantation in a patient with massive plexiform neurofibroma: a 1-year follow-up study. Lancet, The, 2008, 372, 639-645.	13.7	329
5	Quality of life impairment in hidradenitis suppurativa: A study of 61 cases. Journal of the American Academy of Dermatology, 2007, 56, 621-623.	1.2	325
6	Severe cutaneous adverse reactions to drugs. Lancet, The, 2017, 390, 1996-2011.	13.7	293
7	Association between benign and malignant peripheral nerve sheath tumors in NF1. Neurology, 2005, 65, 205-211.	1.1	274
8	Elevated Risk for MPNST in NF1 Microdeletion Patients. American Journal of Human Genetics, 2003, 72, 1288-1292.	6.2	271
9	Patch testing in severe cutaneous adverse drug reactions, including Stevensâ€Johnson syndrome and toxic epidermal necrolysis. Contact Dermatitis, 1996, 35, 234-236.	1.4	260
10	Neurological complications of neurofibromatosis type 1 in adulthood. Brain, 1999, 122, 473-481.	7.6	245
11	Clinical characteristics of a series of 302 French patients with hidradenitis suppurativa, with an analysis of factors associated with disease severity. Journal of the American Academy of Dermatology, 2009, 61, 51-57.	1.2	244
12	Combination Therapy with Clindamycin and Rifampicin for Hidradenitis Suppurativa: A Series of 116 Consecutive Patients. Dermatology, 2009, 219, 148-154.	2.1	222
13	NF1 microdeletions in neurofibromatosis type 1 : from genotype to phenotype. Human Mutation, $2010, 31, E1506-E1518$.	2.5	208
14	Open trial of ciclosporin treatment for Stevens-Johnson syndrome and toxic epidermal necrolysis. British Journal of Dermatology, 2010, 163, 847-853.	1.5	204
15	Identification of Three Hidradenitis Suppurativa Phenotypes: Latent Class Analysis of a Cross-Sectional Study. Journal of Investigative Dermatology, 2013, 133, 1506-1511.	0.7	187
16	Quality-of-Life Impairment in Neurofibromatosis Type 1. Archives of Dermatology, 2001, 137, 1421-5.	1.4	177
17	Pulmonary complications in toxic epidermal necrolysis: a prospective clinical study. Intensive Care Medicine, 1997, 23, 1237-44.	8.2	163
18	<i>NF1</i> Molecular Characterization and Neurofibromatosis Type I Genotype-Phenotype Correlation: The French Experience. Human Mutation, 2013, 34, 1510-1518.	2.5	140

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19	Neurofibromatosis type 1: from genotype to phenotype. Journal of Medical Genetics, 2012, 49, 483-489.	3.2	133
20	Cardiovascular disease risk factors in patients with hidradenitis suppurativa: a systematic review and meta-analysis of observational studies. British Journal of Dermatology, 2015, 173, 1142-1155.	1.5	132
21	Unravelling the genetic basis of variable clinical expression in neurofibromatosis 1. Human Molecular Genetics, 2009, 18, 2768-2778.	2.9	129
22	Intrinsic Defect in Keratinocyte Function Leads to Inflammation in Hidradenitis Suppurativa. Journal of Investigative Dermatology, 2016, 136, 1768-1780.	0.7	129
23	Impact of neurofibromatosis 1 on Quality of Life: A crossâ€sectional study of 176 American cases. American Journal of Medical Genetics, Part A, 2006, 140A, 1893-1898.	1.2	121
24	Systemic involvement of acute generalized exanthematous pustulosis: a retrospective study on 58 patients. British Journal of Dermatology, 2013, 169, 1223-1232.	1.5	121
25	Acne prevalence and associations with lifestyle: a crossâ€sectional online survey of adolescents/young adults in 7 European countries. Journal of the European Academy of Dermatology and Venereology, 2018, 32, 298-306.	2.4	115
26	Therapeutic management of DRESS: A retrospective study of 38 cases. Journal of the American Academy of Dermatology, 2015, 72, 246-252.	1.2	110
27	Role of Noncoding RNA ANRIL in Genesis of Plexiform Neurofibromas in Neurofibromatosis Type 1. Journal of the National Cancer Institute, 2011, 103, 1713-1722.	6.3	106
28	Symptoms associated with malignancy of peripheral nerve sheath tumours: a retrospective study of 69 patients with neurofibromatosis 1. British Journal of Dermatology, 2005, 153, 79-82.	1.5	104
29	Histopathology of drug rash with eosinophilia and systemic symptoms syndrome: a morphological and phenotypical study. British Journal of Dermatology, 2015, 173, 50-58.	1.5	104
30	SPRED1 germline mutations caused a neurofibromatosis type 1 overlapping phenotype. Journal of Medical Genetics, 2009, 46, 425-430.	3.2	103
31	Linear IgA bullous dermatosis: comparison between the drug-induced and spontaneous forms. British Journal of Dermatology, 2013, 169, 1041-1048.	1.5	99
32	Neurofibromatosis type 1 molecular diagnosis: what can NGS do for you when you have a large gene with loss of function mutations?. European Journal of Human Genetics, 2015, 23, 596-601.	2.8	97
33	Mortality Associated with Neurofibromatosis 1: A Cohort Study of 1895 Patients in 1980-2006 in France. Orphanet Journal of Rare Diseases, 2011, 6, 18.	2.7	96
34	Toxic epidermal necrolysis, DRESS, AGEP: Do overlap cases exist?. Orphanet Journal of Rare Diseases, 2012, 7, 72.	2.7	96
35	Neurofibromatosis 1 French national guidelines based on an extensive literature review since 1966. Orphanet Journal of Rare Diseases, 2020, 15, 37.	2.7	96
36	Bacteremia in Stevens-Johnson Syndrome and Toxic Epidermal Necrolysis. Medicine (United States), 2010, 89, 28-36.	1.0	80

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37	Cutaneous neurofibromas. Neurology, 2018, 91, S5-S13.	1.1	79
38	Psoriasis in France and Associated Risk Factors: Results of a Case-Control Study Based on a Large Community Survey. Dermatology, 2009, 218, 103-109.	2.1	76
39	Subcutaneous neurofibromas are associated with mortality in neurofibromatosis 1: A cohort study of 703 patients. American Journal of Medical Genetics, Part A, 2005, 132A, 49-53.	1.2	73
40	Health-Related Quality of Life in Patients with Neurofibromatosis Type 1. Dermatology, 2009, 218, 215-220.	2.1	73
41	Acute Respiratory Failure in Patients With Toxic Epidermal Necrolysis. Critical Care Medicine, 2014, 42, 118-128.	0.9	72
42	Association Between Mediterranean Anti-inflammatory Dietary Profile and Severity of Psoriasis. JAMA Dermatology, 2018, 154, 1017.	4.1	70
43	mTORC1 inhibition delays growth of neurofibromatosis type 2 schwannoma. Neuro-Oncology, 2014, 16, 493-504.	1.2	67
44	Evolving Pattern with Age of Cutaneous Signs in Neurofibromatosis Type 1: A Cross-Sectional Study of 728 Patients. Dermatology, 2011, 222, 269-273.	2.1	64
45	The role of prior corticosteroid use on the clinical course of Stevens-Johnson syndrome and toxic epidermal necrolysis: a case-control analysis of patients selected from the multinational EuroSCAR and RegiSCAR studies. British Journal of Dermatology, 2012, 167, 555-562.	1.5	64
46	Histiocytoid Sweet Syndrome Is More Frequently Associated With Myelodysplastic Syndromes Than the Classical Neutrophilic Variant. Medicine (United States), 2016, 95, e3033.	1.0	63
47	Neurofibromatosis 1-associated neuropathies: a reappraisal. Brain, 2004, 127, 1993-2009.	7.6	61
48	Impact of neurofibromatosis 1 upon quality of life in childhood: a cross-sectional study of 79 cases. British Journal of Dermatology, 2009, 160, 844-848.	1.5	57
49	Cellular Origin, Tumor Progression, and Pathogenic Mechanisms of Cutaneous Neurofibromas Revealed by Mice with <i>Nf1 </i> /i>Knockout in Boundary Cap Cells. Cancer Discovery, 2019, 9, 130-147.	9.4	57
50	Epidermal necrolysis French national diagnosis and care protocol (PNDS; protocole national de) Tj ETQq0 0 0 rgE	3T Oyerloo	ck <u>19</u> Tf 50 22
51	First-line Treatment of Pemphigus Vulgaris With a Combination of Rituximab and High-Potency Topical Corticosteroids. JAMA Dermatology, 2015, 151, 200.	4.1	48
52	Smoking and Dietary Factors Associated with Moderate-to-Severe Acne in French Adolescents and Young Adults: Results of a Survey Using a Representative Sample. Dermatology, 2015, 230, 34-39.	2.1	48
53	Clinical Risk Factors for Mortality in Patients With Neurofibromatosis 1. Archives of Dermatology, 2003, 139, 187-91.	1.4	47
54	Dual mTORC1/2 inhibition induces anti-proliferative effect in NF1-associated plexiform neurofibroma and malignant peripheral nerve sheath tumor cells. Oncotarget, 2016, 7, 35753-35767.	1.8	46

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55	Adverse events associated with JAK inhibitors in 126,815 reports from the WHO pharmacovigilance database. Scientific Reports, 2022, 12, 7140.	3.3	45
56	Usefulness of Screening Investigations in Neurofibromatosis Type 1. Archives of Dermatology, 1996, 132, 1333.	1.4	42
57	Idiopathic linear IgA bullous dermatosis: prognostic factors based on a case series of 72 adults. British Journal of Dermatology, 2017, 177, 212-222.	1.5	42
58	Usefulness of screening investigations in neurofibromatosis type 1. A study of 152 patients. Archives of Dermatology, 1996, 132, 1333-1336.	1.4	42
59	Cyclosporine for Epidermal Necrolysis: Absence of Beneficial Effect in aÂRetrospective Cohort of 174 Patientsâ€"Exposed/Unexposed and Propensity Score-Matched Analyses. Journal of Investigative Dermatology, 2018, 138, 1293-1300.	0.7	41
60	Clinical and histologic features of Mycoplasma pneumoniae –related erythema multiforme: A single-center series of 33 cases compared with 100 cases induced by other causes. Journal of the American Academy of Dermatology, 2018, 79, 110-117.	1.2	41
61	Consequences of Acne on Stress, Fatigue, Sleep Disorders and Sexual Activity: A Population-based Study. Acta Dermato-Venereologica, 2015, 95, 485-488.	1.3	38
62	Schwannomatosis: A Clinical Entity Distinct from Neurofibromatosis Type 2. Dermatology, 1997, 195, 228-231.	2.1	37
63	Hair follicle stem cell replication stress drives IFI16/STING-dependent inflammation in hidradenitis suppurativa. Journal of Clinical Investigation, 2020, 130, 3777-3790.	8.2	35
64	Prognostic value of histologic features of toxic epidermal necrolysis. Journal of the American Academy of Dermatology, 2013, 68, e29-e35.	1.2	34
65	Acute generalized exanthematous pustulosis: a retrospective audit of practice between 1994 and 2011 at a single centre. British Journal of Dermatology, 2015, 172, 1455-1457.	1.5	34
66	Detection and Characterization of NF1 Microdeletions by Custom High Resolution Array CGH. Journal of Molecular Diagnostics, 2009, 11, 524-529.	2.8	31
67	NF-1Score: A Prediction Score for Internal Neurofibromas in Neurofibromatosis-1. Journal of Investigative Dermatology, 2010, 130, 2173-2178.	0.7	31
68	Stevens-Johnson Syndrome and Toxic Epidermal Necrolysis. JAMA Dermatology, 2015, 151, 302.	4.1	31
69	Treatment of prurigo with methotrexate: a multicentre retrospective study of 39 cases. Journal of the European Academy of Dermatology and Venereology, 2018, 32, 437-440.	2.4	30
70	Different Patterns of Mast Cells Distinguish Diffuse from Encapsulated Neurofibromas in Patients with Neurofibromatosis 1. Journal of Histochemistry and Cytochemistry, 2011, 59, 584-590.	2.5	29
71	Sirolimus Improves Pain in NF1 Patients With Severe Plexiform Neurofibromas. Pediatrics, 2014, 133, e1792-e1797.	2.1	29
72	Patch testing in nonâ€immediate cutaneous adverse drug reactions: value of extemporaneous patch tests. Contact Dermatitis, 2017, 77, 297-302.	1.4	29

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73	Postâ€traumatic stress disorder in Stevens–Johnson syndrome and toxic epidermal necrolysis: prevalence and risk factors. A prospective study of 31 patients. British Journal of Dermatology, 2019, 180, 1206-1213.	1.5	29
74	Incidence of and mortality from epidermal necrolysis (Stevens–Johnson syndrome/toxic epidermal) Tj ETQq0 0 C Dermatology, 2020, 182, 618-624.) rgBT /Ove 1.5	erlock 10 Tf 29
75	Clinical Characteristics of Pruritus in Neurofibromatosis 1. Acta Dermato-Venereologica, 2016, 96, 398-399.	1.3	28
76	The biology of cutaneous neurofibromas. Neurology, 2018, 91, S14-S20.	1.1	27
77	Dermatological emergencies: a comparative study of activity in 2000 and 2010. Journal of the European Academy of Dermatology and Venereology, 2013, 27, 916-918.	2.4	26
78	Stevens-Johnson syndrome and toxic epidermal necrolysis: follow-up of pulmonary function after remission. British Journal of Dermatology, 2015, 172, 400-405.	1.5	26
79	Breast cancer risk in neurofibromatosis type 1 is a function of the type of $\langle i \rangle NF1 \langle j \rangle$ gene mutation: a new genotype-phenotype correlation. Journal of Medical Genetics, 2019, 56, 209-219.	3.2	26
80	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
81	Cutaneous neurofibromas: patients' medical burden, current management and therapeutic expectations: results from an online European patient community survey. Orphanet Journal of Rare Diseases, 2019, 14, 286.	2.7	25
82	Visibility of Neurofibromatosis 1 and Psychiatric Morbidity. Archives of Dermatology, 2003, 139, 103.	1.4	24
83	Healthâ€related quality of life and longâ€term sequelae in survivors of epidermal necrolysis: an observational study of 57 patients. British Journal of Dermatology, 2020, 182, 916-926.	1.5	24
84	Current status and recommendations for biomarkers and biobanking in neurofibromatosis. Neurology, 2016, 87, S40-8.	1.1	23
85	Vulvovaginal sequelae in toxic epidermal necrolysis. Journal of reproductive medicine, The, 1997, 42, 153-6.	0.2	23
86	Association Between Severe Acute Contact Dermatitis Due to <i>Nigella sativa</i> Oil and Epidermal Apoptosis. JAMA Dermatology, 2018, 154, 1062.	4.1	22
87	Optimal oncologic management and mTOR inhibitor introduction are safe and improve survival in kidney and liver allograft recipients with <i>de novo</i> carcinoma. International Journal of Cancer, 2019, 144, 886-896.	5.1	22
88	Supportive care in the acute phase of Stevens–Johnson syndrome and toxic epidermal necrolysis: an international, multidisciplinary Delphiâ€based consensus. British Journal of Dermatology, 2021, 185, 616-626.	1.5	22
89	Clinical characteristics predicting internal neurofibromas in 357 children with neurofibromatosis-1: results from a cross-selectional study. Orphanet Journal of Rare Diseases, 2012, 7, 62.	2.7	20
90	Absence of Efficacy of Everolimus in Neurofibromatosis 1-Related Plexiform Neurofibromas: Results from a Phase 2a Trial. Journal of Investigative Dermatology, 2019, 139, 718-720.	0.7	19

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91	At-Risk Phenotype of Neurofibromatose-1 Patients: A Multicentre Case-Control Study. Orphanet Journal of Rare Diseases, 2011, 6, 51.	2.7	18
92	Antitumour necrosis factor- \hat{l}_{\pm} therapy for hidradenitis suppurativa: results from a national cohort study between 2000 and 2013. British Journal of Dermatology, 2016, 174, 667-670.	1.5	18
93	Severe cutaneous adverse reactions due to inappropriate medication use. British Journal of Dermatology, 2018, 179, 329-336.	1.5	17
94	Trends in mortality rates for Stevens–Johnson syndrome and toxic epidermal necrolysis: experience of a single centre in France between 1997 and 2017. British Journal of Dermatology, 2020, 182, 247-248.	1.5	16
95	Contact dermatitis caused by ascorbyl tetraisopalmitate in a cream used for the management of atopic dermatitis. Contact Dermatitis, 2014, 71, 60-61.	1.4	15
96	NF1 single and multi-exons copy number variations in neurofibromatosis type 1. Journal of Human Genetics, 2015, 60, 221-224.	2.3	15
97	Rituximab, a new treatment for difficultâ€ŧoâ€ŧreat chronic erythema multiforme major? Five cases. Journal of the European Academy of Dermatology and Venereology, 2016, 30, 1140-1143.	2.4	15
98	Dermatological emergencies: evolution from 2008 to 2014 and perspectives. Journal of the European Academy of Dermatology and Venereology, 2017, 31, 274-279.	2.4	15
99	Acne across Europe: an online survey on perceptions and management of acne. Journal of the European Academy of Dermatology and Venereology, 2018, 32, 463-466.	2.4	15
100	Severe Phenotype in Patients with Large Deletions of NF1. Cancers, 2021, 13, 2963.	3.7	15
101	Idiopathic Stevens-Johnson syndrome and toxic epidermal necrolysis: Prevalence and patients' characteristics. Journal of the American Academy of Dermatology, 2019, 80, 1453-1455.	1.2	14
102	Individual―and hospital―evel factors associated with epidermal necrolysis mortality: a nationwide multilevel study, France, 2012–2016. British Journal of Dermatology, 2020, 182, 900-906.	1.5	13
103	Agreement Between Self-reported Inflammatory Skin Disorders and Dermatologists' Diagnosis: A Cross-sectional Diagnostic Study. Acta Dermato-Venereologica, 2017, 97, 1243-1244.	1.3	12
104	Crossâ€reactivity in betaâ€lactams after a nonâ€lmmediate cutaneous adverse reaction: experience of a reference centre for toxic bullous diseases and severe cutaneous adverse reactions. Journal of the European Academy of Dermatology and Venereology, 2020, 34, 787-794.	2.4	12
105	Phenotype and Outcomes of Pulmonary Hypertension Associated with Neurofibromatosis Type 1. American Journal of Respiratory and Critical Care Medicine, 2020, 202, 843-852.	5.6	12
106	Creating a comprehensive research strategy for cutaneous neurofibromas. Neurology, 2018, 91, S1-S4.	1.1	11
107	Incidence of bloodstream infections and predictive value of qualitative and quantitative skin cultures of patients with overlap syndrome or toxic epidermal necrolysis: A retrospective observational cohort study of 98 cases. Journal of the American Academy of Dermatology, 2019, 81, 342-347.	1.2	11
108	Severe contact allergy to mupirocin in a polysensitized patient. Contact Dermatitis, 2019, 80, 397-398.	1.4	11

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109	Long-term stability of 0.1% rapamycin hydrophilic gel in the treatment of facial angiofibromas. European Journal of Hospital Pharmacy, 2020, 27, e48-e52.	1.1	11
110	Treatment of cutaneous neurofibromas with carbon dioxide laser: Technique and patient experience. European Journal of Medical Genetics, 2022, 65, 104386.	1.3	11
111	ERN GENTURIS clinical practice guidelines for the diagnosis, treatment, management and surveillance of people with schwannomatosis. European Journal of Human Genetics, 2022, 30, 812-817.	2.8	11
112	Neurofibromatosis type 1: neurofibromas and sex. British Journal of Dermatology, 2016, 174, 402-404.	1.5	10
113	Renal replacement therapy during Stevens-Johnson syndrome and toxic epidermal necrolysis: a retrospective observational study of 238 patients. British Journal of Dermatology, 2017, 176, 1370-1372.	1.5	10
114	Gastrointestinal involvement in Stevens–Johnson syndrome and toxic epidermal necrolysis: a retrospective case series. British Journal of Dermatology, 2019, 180, 1234-1235.	1.5	10
115	A process modelling approach to assess the impact of teledermatology deployment onto the skin tumor care pathway. International Journal of Medical Informatics, 2021, 146, 104361.	3.3	10
116	Chronic pain: a longâ€ŧerm sequela of epidermal necrolysis (Stevens–Johnson syndrome/toxic epidermal) Tj ETQ of Dermatology and Venereology, 2021, 35, 188-194.)q0 0 0 rg 2.4	BT /Overlocl 10
117	High-concentration topical capsaicin in the management of refractory neuropathic pain in patients with neurofibromatosis type 1: a case series. Current Medical Research and Opinion, 2018, 34, 887-891.	1.9	9
118	Severe sequelae of erythema multiforme: three cases. Journal of the European Academy of Dermatology and Venereology, 2018, 32, e34-e36.	2.4	9
119	Disabling ocular sequelae of epidermal necrolysis: risk factors during the acute phase and associated sequelae. British Journal of Dermatology, 2019, 181, 421-422.	1.5	9
120	Bladder Dysfunction in Children with Neurofibromatosis Type I: Report of Four Cases and Review of the Literature. Urologia Internationalis, 2018, 100, 339-345.	1.3	8
121	Lenalidomide as an Alternative to Thalidomide for Treatment of Recurrent Erythema Multiforme. JAMA Dermatology, 2018, 154, 487.	4.1	8
122	Eruption of lymphocyte recovery with atypical lymphocytes mimicking a primary cutaneous T-cell lymphoma: a series of 12 patients. Human Pathology, 2018, 71, 100-108.	2.0	8
123	Usefulness of screening investigations in neurofibromatosis type 1. A study of 152 patients. Archives of Dermatology, 1996, 132, 1333-6.	1.4	8
124	Epidermolysis bullosa acquisita-like eruption with anticollagen VII autoantibodies induced by <scp>d < /scp>-penicillamine in Wilson disease. British Journal of Dermatology, 2014, 171, 1574-1576.</scp>	1.5	7
125	Facial Scars following Toxic Epidermal Necrolysis: Role of Adnexal Involvement?. Dermatology, 2016, 232, 220-223.	2.1	7
126	Epidermal necrolysis and autoimmune diseases: two more observations supporting the concept that †toxic†toxic†epidermal necrolysis can be †non†toxicâ€. Journal of the European Academy of Dermatology ar Venereology, 2018, 32, e360-e361.	n d .4	7

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127	A large epidemiological study of erythema multiforme in France, with emphasis on treatment choices. British Journal of Dermatology, 2018, 179, 1009-1011.	1.5	7
128	Polysensitivity in delayed cutaneous adverse drug reactions to macrolides, clindamycin and pristinamycin: clinical history and patch testing. British Journal of Dermatology, 2018, 179, 978-979.	1.5	7
129	Neurofibromatosis I and multiple sclerosis. Orphanet Journal of Rare Diseases, 2020, 15, 186.	2.7	7
130	Identifying challenges in neurofibromatosis: a modified Delphi procedure. European Journal of Human Genetics, 2021, 29, 1625-1633.	2.8	7
131	Clinical and histological features of fixed drug eruption: a single-centre series of 73 cases with comparison between bullous and non-bullous forms. European Journal of Dermatology, 2021, 31, 372-380.	0.6	7
132	Quality of life in neurofibromatosis 1: development and validation of a tool dedicated to cutaneous neurofibromas in adults. Journal of the European Academy of Dermatology and Venereology, 2022, 36, 1359-1366.	2.4	7
133	Febrile ulceronecrotic Mucha Habermann disease mimicking aggressive epidermotropic CD8+ cytotoxic T-cell lymphoma: a diagnostic challenge. European Journal of Dermatology, 2018, 28, 834-835.	0.6	7
134	Bullous pemphigoid: Three main clusters defining 3 outcome profiles. Journal of the American Academy of Dermatology, 2022, 87, 359-365.	1.2	7
135	Facial transplantation: facing the limits, planning the future. Lancet, The, 2017, 389, 1293-1294.	13.7	6
136	Face transplantation: A longitudinal histological study focusing on chronic active and mucosal rejection in a series with long-term follow-up. American Journal of Transplantation, 2021, 21, 3088-3100.	4.7	6
137	S100B and neurofibromin immunostaining and Xâ€inactivation patterns of laserâ€microdissected cells indicate a multicellular origin of some NF1â€associated neurofibromas. Journal of Neuroscience Research, 2011, 89, 1451-1460.	2.9	5
138	Severe Cutaneous Adverse Reactions to Drugs: From Patients to the National Office for Compensation of Medical Accidents. Dermatology, 2014, 228, 338-343.	2.1	5
139	Toxic epidermal necrolysis: the past, the guidelines and challenges for the future. British Journal of Dermatology, 2016, 174, 1171-1173.	1.5	5
140	Skin biopsy polymerase chain reaction for rapid microbiological diagnosis in patients with purpura fulminans. British Journal of Dermatology, 2017, 177, e154-e155.	1.5	5
141	Dermatitis herpetiformis and bone mineral density: analysis of a French cohort of 53 patients. European Journal of Dermatology, 2017, 27, 353-358.	0.6	5
142	Very late sensitization to parabens induced by repeated applications of an anaesthetic therapeutic plaster to nonâ€damaged skin. Contact Dermatitis, 2018, 79, 194-195.	1.4	5
143	Cutaneous tests and interest of iobitridol in nonâ€immediate hypersensitivity to contrast media: a case series of 43 patients. Journal of the European Academy of Dermatology and Venereology, 2020, 34, e178-e180.	2.4	5
144	Combined Methotrexate and Alitretinoin for the treatment of difficultâ€toâ€treat generalized prurigo nodularis: a case series. Journal of the European Academy of Dermatology and Venereology, 2021, 35, e516-e519.	2.4	5

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145	Lymphoproliferative malignancies in patients with neurofibromatosis 1. Orphanet Journal of Rare Diseases, 2021, 16, 230.	2.7	5
146	Patch tests in nonâ€immediate cutaneous adverse drug reactions: the importance of late readings on day 4. Contact Dermatitis, 2021, , .	1.4	5
147	Toxic epidermal necrolysis: The past, the guidelines and challenges for the future. Journal of Plastic, Reconstructive and Aesthetic Surgery, 2016, 69, 733-735.	1.0	4
148	Patient-hospital communication: A platform to improve outpatient chemotherapy., 2016,,.		4
149	Beard dermatitis induced by coloration. Contact Dermatitis, 2019, 81, 471-473.	1.4	4
150	Acute generalized exanthematous pustulosis and epidermal necrolysis differ in innate cytokine patterns. Clinical and Experimental Allergy, 2019, 49, 1258-1261.	2.9	4
151	Strong reactions to diltiazem patch tests: Plea for a low concentration. Contact Dermatitis, 2020, 83, 224-225.	1.4	4
152	Acute generalized exanthematous pustulosis induced by enoxaparin: 2 cases. Contact Dermatitis, 2021, 84, 280-282.	1.4	4
153	Relapsing generalized bullous fixed drug eruption: A severe and avoidable cutaneous drug reaction. Three case reports. Therapie, 2021, , .	1.0	4
154	Towards a better understanding of adult idiopathic epidermal necrolysis: a retrospective study of 19 cases. Journal of the European Academy of Dermatology and Venereology, 2021, 35, 1569-1576.	2.4	4
155	Dermatological emergency unit, dayâ€care hospital and consultations in time of COVIDâ€19: the impact of teledermatology. Journal of the European Academy of Dermatology and Venereology, 2022, 36, .	2.4	4
156	Necrotizing fasciitis of the thigh revealing a Crohn's disease. Journal of the European Academy of Dermatology and Venereology, 2015, 29, 1648-1649.	2.4	3
157	Pemphigoid gestationis revealing a denial of pregnancy. Journal of the European Academy of Dermatology and Venereology, 2016, 30, 1411-1413.	2.4	3
158	Are Idiopathic Stevens-Johnson Syndrome/Toxic Epidermal Necrolysis Related to Drugs in Food? The Example of Phenylbutazone. Journal of Investigative Dermatology, 2017, 137, 1179-1181.	0.7	3
159	Hemostasis and Type 1 Neurofibromatosis. Plastic and Reconstructive Surgery - Global Open, 2017, 5, e1414.	0.6	3
160	Epidemiology of fragile skin: Internet-based surveys in Mexico and Russia. Clinical, Cosmetic and Investigational Dermatology, 2017, Volume 10, 221-228.	1.8	3
161	Stevens-Johnson Syndrome During Pregnancy. JAMA Dermatology, 2018, 154, 224.	4.1	3
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