

# Dan Lipsker

## List of Publications by Year in descending order

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131  
papers

3,629  
citations

126907

33  
h-index

161849

54  
g-index

166  
all docs

166  
docs citations

166  
times ranked

3049  
citing authors

#	ARTICLE	IF	CITATIONS
1	Schnitzler's syndrome: diagnosis, treatment, and follow-up. <i>Allergy: European Journal of Allergy and Clinical Immunology</i> , 2013, 68, 562-568.	5.7	224
2	The Schnitzler Syndrome. <i>Medicine (United States)</i> , 2001, 80, 37-44.	1.0	221
3	Neutrophilic Urticarial Dermatositis. <i>Medicine (United States)</i> , 2009, 88, 23-31.	1.0	193
4	The Schnitzler syndrome. <i>Orphanet Journal of Rare Diseases</i> , 2010, 5, 38.	2.7	144
5	Trends in melanoma epidemiology suggest three different types of melanoma. <i>British Journal of Dermatology</i> , 2007, 157, 338-343.	1.5	104
6	Clinico-biological characteristics and treatment of type I monoclonal cryoglobulinaemia: a study of 64 cases. <i>British Journal of Haematology</i> , 2015, 168, 671-678.	2.5	91
7	The Cutaneous Spectrum of Lupus Erythematosus. <i>Clinical Reviews in Allergy and Immunology</i> , 2017, 53, 291-305.	6.5	83
8	Molecular genetic investigation, clinical features, and response to treatment in 21 patients with Schnitzler syndrome. <i>Blood</i> , 2018, 131, 974-981.	1.4	83
9	How not to miss autoinflammatory diseases masquerading as urticaria. <i>Allergy: European Journal of Allergy and Clinical Immunology</i> , 2012, 67, 1465-1474.	5.7	74
10	Monoclonal gammopathy of cutaneous significance: review of a relevant concept. <i>Journal of the European Academy of Dermatology and Venereology</i> , 2017, 31, 45-52.	2.4	73
11	The AESOP (Adenopathy and Extensive Skin Patch Overlying a Plasmacytoma) Syndrome. <i>Medicine (United States)</i> , 2003, 82, 51-59.	1.0	72
12	Single-cell gene expression signatures reveal melanoma cell heterogeneity. <i>Oncogene</i> , 2015, 34, 3251-3263.	5.9	72
13	Schnitzler syndrome: validation and applicability of diagnostic criteria in real-life patients. <i>Allergy: European Journal of Allergy and Clinical Immunology</i> , 2017, 72, 177-182.	5.7	66
14	Schnitzler syndrome: heterogeneous immunopathological findings involving IgM-skin interactions. <i>British Journal of Dermatology</i> , 2000, 142, 954-959.	1.5	64
15	Could Jessner's Lymphocytic Infiltrate of the Skin Be a Dermal Variant of Lupus Erythematosus? An Analysis of 210 Cases. <i>Dermatology</i> , 2006, 213, 15-22.	2.1	63
16	Schnitzler Syndrome: a Review. <i>Current Rheumatology Reports</i> , 2017, 19, 46.	4.7	59
17	Neutrophilic Cutaneous Lupus Erythematosus. <i>Dermatology</i> , 2008, 216, 283-286.	2.1	58
18	Cryopyrin-associated periodic syndrome: an autoinflammatory disease manifested as neutrophilic urticarial dermatosis with additional perieccrine involvement. <i>Journal of Cutaneous Pathology</i> , 2011, 38, 202-208.	1.3	58

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19	Hidradenitis suppurativa (HS): An unrecognized paradoxical effect of biologic agents (BA) used in chronic inflammatory diseases. <i>Journal of the American Academy of Dermatology</i> , 2016, 74, 1153-1159.	1.2	56
20	The combination of complement deficiency and cigarette smoking as risk factor for cutaneous lupus erythematosus in men; a focus on combined C2/C4 deficiency. <i>British Journal of Dermatology</i> , 2005, 152, 265-270.	1.5	54
21	Efficacy of Immunotherapy in Patients with Metastatic Mucosal or Uveal Melanoma. <i>Journal of Oncology</i> , 2018, 2018, 1-9.	1.3	53
22	High Frequency of Genital Lichen Sclerosus in a Prospective Series of 76 Patients With Morphea. <i>Archives of Dermatology</i> , 2012, 148, 24.	1.4	49
23	Cutaneous manifestations of complement deficiencies. <i>Lupus</i> , 2010, 19, 1096-1106.	1.6	48
24	Chloroquine-Quinacrine Association in Resistant Cutaneous Lupus. <i>Dermatology</i> , 1995, 190, 257-258.	2.1	48
25	Association of Cigarette Smoking but Not Alcohol Consumption With Cutaneous Lupus Erythematosus. <i>Archives of Dermatology</i> , 2009, 145, 1012-6.	1.4	46
26	Proteasome subunit <i>PSMC3</i> variants cause neurosensory syndrome combining deafness and cataract due to proteotoxic stress. <i>EMBO Molecular Medicine</i> , 2020, 12, e11861.	6.9	43
27	Disease Expression of Lyme Borreliosis in Northeastern France. <i>European Journal of Clinical Microbiology and Infectious Diseases</i> , 2001, 20, 0225-0230.	2.9	42
28	Long-term prognosis of patients treated for erythema migrans in France. <i>British Journal of Dermatology</i> , 2002, 146, 872-876.	1.5	42
29	The need to revisit the nosology of cutaneous lupus erythematosus. <i>Lupus</i> , 2010, 19, 1047-1049.	1.6	39
30	Acitretin- and Tumor Necrosis Factor Inhibitor “Resistant Acrodermatitis Continua of Hallopeau Responsive to the Interleukin 1 Receptor Antagonist Anakinra. <i>Archives of Dermatology</i> , 2012, 148, 297.	1.4	39
31	A New Concept: Paraviral Eruptions. <i>Dermatology</i> , 2005, 211, 309-311.	2.1	36
32	The Role of Circumstances of Diagnosis and Access to Dermatological Care in Early Diagnosis of Cutaneous Melanoma. <i>Archives of Dermatology</i> , 2010, 146, 240-6.	1.4	36
33	Neutrophilic urticarial dermatosis: an entity bridging monogenic and polygenic autoinflammatory disorders, and beyond. <i>Journal of the European Academy of Dermatology and Venereology</i> , 2020, 34, 685-690.	2.4	36
34	Unexplained recurrent fever: when is autoinflammation the explanation?. <i>Allergy: European Journal of Allergy and Clinical Immunology</i> , 2013, 68, 285-296.	5.7	35
35	Anakinra for Difficult-to-Treat Neutrophilic Panniculitis: IL-1 Blockade as a Promising Treatment Option for Neutrophil-Mediated Inflammatory Skin Disease. <i>Dermatology</i> , 2010, 220, 264-267.	2.1	34
36	Lupus Erythematosus and Neutrophilic Urticarial Dermatitis. <i>Medicine (United States)</i> , 2014, 93, e351.	1.0	34

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37	Distinctive cutaneous and systemic features associated with specific antimyositis antibodies in adults with dermatomyositis: a prospective multicentric study of 117 patients. <i>Journal of the European Academy of Dermatology and Venereology</i> , 2018, 32, 1164-1172.	2.4	33
38	Growth Rate, Early Detection, and Prevention of Melanoma. <i>Archives of Dermatology</i> , 2006, 142, 1638-40.	1.4	32
39	International and multidisciplinary expert recommendations for the use of biologics in systemic lupus erythematosus. <i>Autoimmunity Reviews</i> , 2017, 16, 650-657.	5.8	32
40	Livedoid Vasculopathy: A French Observational Study Including Therapeutic Options. <i>Acta Dermato-Venereologica</i> , 2018, 98, 842-847.	1.3	31
41	Examination of cutaneous macroglobulinosis by immunoelectron microscopy. <i>British Journal of Dermatology</i> , 1996, 135, 287-291.	1.5	30
42	Epidemiology of Merkel cell carcinoma. A population-based study from 1985 to 2013, in northeastern of France. <i>International Journal of Cancer</i> , 2019, 144, 741-745.	5.1	29
43	Between Light and Dark, the Chimera Comes Out. <i>Archives of Dermatology</i> , 2008, 144, 327-30.	1.4	28
44	Neutrophilic urticarial dermatosis: A review. <i>Annales De Dermatologie Et De Venereologie</i> , 2018, 145, 735-740.	1.0	28
45	How Accurate Is a Clinical Diagnosis of Erythema Chronicum Migrans? Prospective Study Comparing the Diagnostic Accuracy of General Practitioners and Dermatologists in an Area Where Lyme Borreliosis Is Endemic. <i>Archives of Dermatology</i> , 2004, 140, 620-1.	1.4	27
46	Expanding the clinicopathological spectrum of late cutaneous Lyme borreliosis (acrodermatitis) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 38 (PCR)-documented cases. <i>Journal of the American Academy of Dermatology</i> , 2016, 74, 685-692.	1.2	27
47	Cutaneous Vasculitis: Review on Diagnosis and Clinicopathologic Correlations. <i>Clinical Reviews in Allergy and Immunology</i> , 2021, 61, 181-193.	6.5	27
48	Blisters and Loss of Epidermis in Patients With Lupus Erythematosus. <i>Medicine (United States)</i> , 2015, 94, e2102.	1.0	26
49	Amicrobial pustulosis of the folds: Where have we gone 25 years after its original description?. <i>Annales De Dermatologie Et De Venereologie</i> , 2017, 144, 169-175.	1.0	25
50	Expanding phenotype of hereditary fibrosing poikiloderma with tendon contractures, myopathy, and pulmonary fibrosis caused by FAM111B mutations: Report of an additional family raising the question of cancer predisposition and a short review of early-onset poikiloderma. <i>JAAD Case Reports</i> , 2017, 3, 143-150.	0.8	24
51	Clinical and Sociodemographic Characteristics Associated With Thick Melanomas. <i>Archives of Dermatology</i> , 2012, 148, 1370.	1.4	23
52	Rosacea and demodicidosis associated with gain-of-function mutation in <scp>STAT</scp>1. <i>Journal of the European Academy of Dermatology and Venereology</i> , 2017, 31, e542-e544.	2.4	22
53	Paraviral eruptions in the era of COVID-19: Do some skin manifestations point to a natural resistance to SARS-CoV-2?. <i>Clinics in Dermatology</i> , 2020, 38, 757-761.	1.6	22
54	Classification of Specific Cutaneous Manifestations in Patients with Lupus Erythematosus: A Time for Change?. <i>Dermatology</i> , 2006, 212, 324-326.	2.1	21

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55	Species of <i>Borrelia burgdorferi</i> complex that cause borrelial lymphocytoma in France. British Journal of Dermatology, 2009, 161, 174-176.	1.5	19
56	The role of general practitioners in diagnosis of cutaneous melanoma: a population-based study in France. British Journal of Dermatology, 2012, 167, 1351-1359.	1.5	19
57	A systematic review of terms used to describe hidradenitis suppurativa. British Journal of Dermatology, 2015, 173, 1298-1300.	1.5	19
58	Increased severity and epidermal alterations in persistent versus evanescent skin lesions in adult-onset Still disease. Journal of the American Academy of Dermatology, 2018, 79, 969-971.	1.2	18
59	Mechanics hands in patients with antisynthetase syndrome: 25 cases. Annales De Dermatologie Et De Venereologie, 2019, 146, 19-25.	1.0	16
60	A Unified Concept of Acne in the PAPA Spectrum Disorders. Dermatology, 2021, 237, 827-834.	2.1	16
61	A chilblain epidemic during the COVID-19 pandemic. A sign of natural resistance to SARS-CoV-2?. Medical Hypotheses, 2020, 144, 109959.	1.5	15
62	Plasma cell-directed therapies in monoclonal gammopathy-associated scleromyxedema. Blood, 2020, 135, 1101-1110.	1.4	15
63	Extracorporeal photopheresis in recalcitrant lupus erythematosus. Clinical and Experimental Dermatology, 2009, 34, e295-e296.	1.3	14
64	A Review and Proposed Approach to the Neutrophilic Dermatoses of Childhood. Pediatric Dermatology, 2015, 32, 437-446.	0.9	12
65	Cytokine Signature in Schnitzler Syndrome: Proinflammatory Cytokine Production Associated to Th Suppression. Frontiers in Immunology, 2020, 11, 588322.	4.8	12
66	The history of lupus throughout the ages. Journal of the American Academy of Dermatology, 2022, 87, 1361-1369.	1.2	12
67	Lupus erythematosus: Significance of dermatologic findings. Annales De Dermatologie Et De Venereologie, 2021, 148, 6-15.	1.0	11
68	Cutis laxa associated with monoclonal gammopathy: 14 new cases and review of the literature. Journal of the American Academy of Dermatology, 2018, 79, 945-947.	1.2	10
69	MRI-Guided Cryoablation of In-Transit Metastases from Cutaneous Melanoma: A Brief Report on a Preliminary Experience. CardioVascular and Interventional Radiology, 2017, 40, 1285-1289.	2.0	9
70	Livedoid vasculopathy: how to diagnose and how to treat?. Journal of the European Academy of Dermatology and Venereology, 2019, 33, 1627-1628.	2.4	9
71	Clinical Images: Toe dactylitis revealing late Lyme borreliosis. Arthritis and Rheumatism, 2012, 64, 1293-1293.	6.7	8
72	Management of Schnitzler's syndrome. Expert Opinion on Orphan Drugs, 2014, 2, 947-955.	0.8	8

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73	Ehlers-Danlos syndrome related to FKBP14 mutations: detailed cutaneous phenotype. Clinical and Experimental Dermatology, 2017, 42, 64-67.	1.3	8
74	The molluscum pendulum necklace sign in tuberous sclerosis complex: a case series A pathognomonic finding?. Journal of the European Academy of Dermatology and Venereology, 2017, 31, e507-e508.	2.4	8
75	Cytokine levels in persistent skin lesions of adult-onset Still disease. Journal of the American Academy of Dermatology, 2018, 79, 947-949.	1.2	8
76	Human babesiosis in Alsace. Médecine Et Maladies Infectieuses, 2020, 50, 486-491.	5.0	8
77	AESOP syndrome: a potential life-saving and early clue to the diagnosis of POEMS syndrome. Clinics in Dermatology, 2021, 39, 215-219.	1.6	8
78	Schnitzler Syndrome: the paradigm of an acquired adult-onset auto-inflammatory disease. Giornale Italiano Di Dermatologia E Venereologia, 2020, 155, 567-573.	0.8	8
79	Retiform Purpura. New England Journal of Medicine, 2008, 358, e1.	27.0	7
80	Eruptive nevi under tocilizumab: first case report and data analysis. Journal of the European Academy of Dermatology and Venereology, 2018, 32, e253-e254.	2.4	7
81	The Importance of Thrombosis in Patients with Lupus Erythematosus. Dermatology, 2006, 212, 214-215.	2.1	6
82	Cutaneous haemorrhage induced by minimal trauma as a sign of light chain-associated amyloidosis. British Journal of Haematology, 2012, 159, 383-383.	2.5	5
83	Efficacy of interferon in recurrent valaciclovir-refractory erythema multiforme in a patient not infected with hepatitis C virus. Clinical and Experimental Dermatology, 2016, 41, 648-650.	1.3	5
84	Granular pemphigus-like IgM deposition around keratinocytes in a patient with Waldenström's macroglobulinaemia: a so far unreported finding. Journal of the European Academy of Dermatology and Venereology, 2017, 31, e47-e49.	2.4	4
85	Small infantile haemangioma and breast hypoplasia. Journal of the European Academy of Dermatology and Venereology, 2017, 31, e355-e356.	2.4	4
86	An Unusual Cause of Recurrent Gastric Bleeding. Gastroenterology, 2018, 155, e9-e11.	1.3	4
87	Prospective evaluation of the frequency of genital lichen sclerosis in 79 patients with systemic sclerosis. British Journal of Dermatology, 2018, 179, 999-1000.	1.5	4
88	Multilocus sequence typing of clinical Borrelia afzelii strains: population structure and differential ability to disseminate in humans. Parasites and Vectors, 2018, 11, 374.	2.5	4
89	Adenopathy and extensive skin patch overlying a plasmacytoma (AESOP): Two morphologic variants can be outlined. Journal of the American Academy of Dermatology, 2021, 85, 1286-1287.	1.2	4
90	Impulse control disorder-linked hypersexuality complicated by disseminated gonococcal infection in a patient with Parkinson's disease. Revue Neurologique, 2020, 176, 292-293.	1.5	4

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91	Isomorphic and symmetric adult-onset generalized morphea are associated with distinctive clinical features: A retrospective multicenter study. <i>Journal of the American Academy of Dermatology</i> , 2021, 84, 1701-1703.	1.2	4
92	Absence of <i>NLRP3</i> somatic mutations and VEXAS-related <i>UBA1</i> mutations in a large cohort of patients with Schnitzler syndrome. <i>Allergy: European Journal of Allergy and Clinical Immunology</i> , 2022, 77, 3435-3436.	5.7	4
93	Aortitis: A new feature of Schnitzler syndrome. <i>JAAD Case Reports</i> , 2017, 3, 454-456.	0.8	3
94	Bedside reasoning on causes and mechanisms of diseases in the era of precision medicine: a timeless story?. <i>Journal of the European Academy of Dermatology and Venereology</i> , 2018, 32, 1436-1440.	2.4	3
95	9q22.3 Microdeletion Syndrome with Multiple Basal Cell Carcinomas Treated with Vismodegib: Three Key Messages in One Patient. <i>Acta Dermato-Venereologica</i> , 2018, 98, 287-288.	1.3	3
96	Diagnostic value of skin biopsy in autoinflammatory diseases for patients with recurrent fever and urticarial eruption. <i>Clinical and Experimental Dermatology</i> , 2021, 46, 728-730.	1.3	3
97	A White Hand and a Red Hand â€” Erythromelalgia. <i>New England Journal of Medicine</i> , 2010, 363, 1463-1463.	27.0	2
98	Haemorrhage and amyloidosis â€” response to Veneriet al. <i>British Journal of Haematology</i> , 2013, 160, 854-855.	2.5	2
99	Eruptive pigmentation around naevi and seborrheic keratoses in a patient with stage III melanoma: hyperpigmented halo or â€”Nottusâ€™ phenomenon. <i>British Journal of Dermatology</i> , 2013, 168, 1140-1141.	1.5	2
100	Occurrence of Rheumatoid Arthritis in a Patient Treated with Anakinra for Schnitzler Syndrome: A Case Report. <i>Journal of Rheumatology</i> , 2016, 43, 1447.1-1447.	2.0	2
101	The Spectrum of Cutaneous Manifestations in Systemic Lupus Erythematosus and Novel Classification. <i>Rare Diseases of the Immune System</i> , 2016, , 77-94.	0.1	2
102	Haemophagocytic lymphohistiocytosis and silvery hair in Griscelli syndrome. <i>British Journal of Haematology</i> , 2016, 175, 11-11.	2.5	2
103	Bendamustine conditioning for refractory type I cryoglobulinemia. <i>Joint Bone Spine</i> , 2016, 83, 591-592.	1.6	2
104	Delayed effort-induced swelling with myofasciitis and systemic manifestations. <i>Medicine (United Tj ETQq0 0 0 rgBTJ /Overlock 10 Tf 50</i>	1.0	2
105	Efficacy and safety of low-dose oral lenalidomide in refractory cutaneous lupus erythematosus: an open series of 19 cases. <i>Journal of the European Academy of Dermatology and Venereology</i> , 2021, 35, e113-e115.	2.4	2
106	Prospective evaluation of frequency of signs of systemic sclerosis in 76 patients with morphea. <i>Clinical and Experimental Rheumatology</i> , 2015, 33, S23-5.	0.8	2
107	Cutaneous collagenous vasculopathy leading to the diagnosis of an advanced pancreatic cancer. <i>Journal of the European Academy of Dermatology and Venereology</i> , 2022, 36, .	2.4	2
108	More about molluscum pendulum/acrochordons in tuberous sclerosis complex. <i>Journal of the European Academy of Dermatology and Venereology</i> , 2018, 32, e144.	2.4	1

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109	Extensive cutaneous necrosis in monoclonal cryoglobulinemia: an example of monoclonal gammopathy of cutaneous significance. Journal of the European Academy of Dermatology and Venereology, 2019, 33, e167.	2.4	1
110	Seborrheic area involvement in patients with systemic lupus erythematosus. Journal of the European Academy of Dermatology and Venereology, 2019, 33, e284-e286.	2.4	1
111	Abnormal lipid storage related to adipocyte shrinkage in acquired partial lipodystrophy (Barraquerâ€“Simons syndrome). Journal of the European Academy of Dermatology and Venereology, 2019, 33, 2188-2191.	2.4	1
112	Epidemiology of juvenile dermatomyositis in Alsace. British Journal of Dermatology, 2020, 182, 1307-1308.	1.5	1
113	<i>PTEN</i> hamartoma tumor syndrome in children: diagnosis based on cutaneous manifestations with a focus on translucent palmoplantar papules. Journal of the European Academy of Dermatology and Venereology, 2020, 34, e632-e633.	2.4	1
114	Pigmented spots on the lacrimal caruncle: a key to the diagnosis of Carney complex in a child with multiple lentigines. Journal of the European Academy of Dermatology and Venereology, 2020, 34, e604-e606.	2.4	1
115	Palmoplantar lichen planusâ€“like lupus erythematosus keratoderma: an underrecognized and distinctive cutaneous manifestation of systemic or subacute lupus erythematosus. Journal of the European Academy of Dermatology and Venereology, 2021, 35, e124-e126.	2.4	1
116	Gammopathies. , 2020, , 1-9.		1
117	Autoinflammatory diseases: why they should matter to the dermatologist. Giornale Italiano Di Dermatologia E Venereologia, 2020, 155, 533-536.	0.8	1
118	Mandibular sterile osteitis as a manifestation of synovitis, acne, pustulosis, hyperostosis, osteitis syndrome: a literature review. Journal of the European Academy of Dermatology and Venereology, 2021, 35, e335-e338.	2.4	1
119	Quels sont les signes cutanÃ©s que doivent connaÃ®tre les rhumatologues pour Ã©voquer une maladie systÃ©mique inflammatoire?. Revue Du Rhumatisme Monographies, 2011, 78, 201-209.	0.0	0
120	Manifestaciones cutÃ¡neas de la borreliosis de Lyme. EMC - DermatologÃ­a, 2017, 51, 1-12.	0.1	0
121	Hiperpigmentaciones. EMC - DermatologÃ­a, 2017, 51, 1-14.	0.1	0
122	Immune deficiency and rosacea. Journal of the European Academy of Dermatology and Venereology, 2018, 32, e88.	2.4	0
123	HautverÃ¤nderungen durch Gammopathien. , 2018, , 1669-1678.		0
124	Multiple brown papules at the site of mastectomy for breast cancer. Lancet Oncology, The, 2020, 21, e460.	10.7	0
125	Semeiologia cutanea. EMC - AKOS - Trattato Di Medicina, 2020, 22, 1-8.	0.0	0
126	An Abscess Is Not a Descriptive Term but an Entity With a Universally Accepted Definitionâ€”A Clarification on Semantics. JAMA Dermatology, 2021, 157, 1244-1245.	4.1	0



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127	Autoinflammatory Syndromes: Relevance to Inflammatory Skin Diseases and Personalized Medicine. , 2015, , 101-110.		0
128	Hautveränderungen durch Gammopathien. , 2017, , 1-10.		0
129	Paraproteinämies. , 2007, , 160-168.		0
130	Gammopathies. , 2022, , 1661-1669.		0
131	Congenital spilo-melanocytic nevus: a speckled lentiginous naevus and a spilus-like congenital melanocytic nevus in close proximity. Journal of the European Academy of Dermatology and Venereology, 2022, 36, .	2.4	0