Anette Bygum

List of Publications by Year in Descending Order

Source: https://exaly.com/author-pdf/3834071/anette-bygum-publications-by-year.pdf
Version: 2024-04-09

This document has been generated based on the publications and citations recorded by exaly.com. For the latest version of this publication list, visit the link given above.
The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

91 papers	1,405 citations	20 h-index	34 g-index
93 ext. papers	1,794 ext. citations	4.9 avg, IF	4.69 L-index

#	Paper	IF	Citations
91	The international WAO/EAACI guideline for the management of hereditary angioedema - the 2021 revision and update.. *Allergy: European Journal of Allergy and Clinical Immunology*, **2022**,	9.3	6
90	The international WAO/EAACI guideline for the management of hereditary angioedema - The 2021 revision and update.. *World Allergy Organization Journal*, **2022**, 15, 100627	5.2	6
89	Outcomes of Systemic Treatment in Children and Adults With Netherton Syndrome: A Systematic Review.. *Frontiers in Immunology*, **2022**, 13, 864449	8.4	2
88	A girl with unruly locks: molecular genetics makes a diagnosis of uncombable hair syndrome.. *Lancet, The*, **2022**, 399, 1079	40	
87	Spontaneous pneumothorax as a clinical manifestation of neurofibromatosis type 1. *BMJ Case Reports*, **2021**, 14,	0.9	1
86	Acceptance of Teledermoscopy by General Practitioners and Dermatologists in Denmark. *Dermatology Practical and Conceptual*, **2021**, 11, e2021033	1.5	2
85	Consensus on treatment goals in hereditary angioedema: A global Delphi initiative. *Journal of Allergy and Clinical Immunology*, **2021**, 148, 1526-1532	11.5	3
84	Long-term safety and effectiveness of berotralstat for hereditary angioedema: The open-label APeX-S study. *Clinical and Translational Allergy*, **2021**, 11, e12035	5.2	5
83	Cercarial Dermatitis at Public Bathing Sites (Region Zealand, Denmark): A Case Series and Literature Review. *Case Reports in Dermatology*, **2021**, 13, 360-365	1.1	0
82	Meta-Analysis of Mutations in or Identified in a Large Cohort of 224 Patients. *Genes*, **2021**, 12,	4.2	5
81	Hereditary leiomyomatosis and renal cell carcinoma: a case series and literature review. *Orphanet Journal of Rare Diseases*, **2021**, 16, 34	4.2	5
80	Introducing teledermoscopy of possible skin cancers in general practice in Southern Denmark. *Family Practice*, **2020**, 37, 513-518	1.9	4
79	Hereditary angioedema: the challenges of cross-border family investigation and treatment. *BMJ Case Reports*, **2020**, 13,	0.9	
78	Exome Sequencing Reveals Common and Rare Variants in F5 Associated With ACE Inhibitor and Angiotensin Receptor Blocker-Induced Angioedema. *Clinical Pharmacology and Therapeutics*, **2020**, 108, 1195-1202	6.1	5
77	Long-term safety and efficacy of subcutaneous C1-inhibitor in older patients with hereditary angioedema. *Annals of Allergy, Asthma and Immunology*, **2020**, 125, 334-340.e1	3.2	2
76	Modeling-based bone formation transforms trabeculae to cortical bone in the sclerotic areas in Buschke-Ollendorff syndrome. A case study of two females with LEMD3 variants. *Bone*, **2020**, 135, 115313	4.7	4
75	Genome-wide association study of angioedema induced by angiotensin-converting enzyme inhibitor and angiotensin receptor blocker treatment. *Pharmacogenomics Journal*, **2020**, 20, 770-783	3.5	9

74	Contact activation-induced complex formation between complement factor H and coagulation factor XIIa. *Journal of Thrombosis and Haemostasis*, **2020**, 18, 876-884	15.4	4
73	Definition, aims, and implementation of GA LEN/HAEi Angioedema Centers of Reference and Excellence. *Allergy: European Journal of Allergy and Clinical Immunology*, **2020**, 75, 2115-2123	9.3	10
72	Increased mortality in patients with porphyria cutanea tarda-A nationwide cohort study. *Journal of the American Academy of Dermatology*, **2020**, 83, 817-823	4.5	2
71	International Consensus on the Use of Genetics in the Management of Hereditary Angioedema. *Journal of Allergy and Clinical Immunology: in Practice*, **2020**, 8, 901-911	5.4	28
70	Intake of dietary fibre, red and processed meat and risk of late-onset Chronic Inflammatory Diseases: A prospective Danish study on the "diet, cancer and health" cohort. *International Journal of Medical Sciences*, **2020**, 17, 2487-2495	3.7	2
69	The impact of adolescentsReveryday life experiences on their primary hyperhidrosis treatment - a qualitative study. *Journal of Dermatological Treatment*, **2020**, 1-7	2.8	1
68	Clinical Features and Disease Course of Primary Angioedema Patients in a Tertiary Care Hospital. *Journal of Asthma and Allergy*, **2020**, 13, 225-236	3.1	3
67	In hyperhidrosis quality of life is even worse than in acne, eczema, or psoriasis. A comparison of Skindex-16 and Dermatology Life Quality Index (DLQI). *International Journal of Dermatology*, **2020**, 59, e392-e393	1.7	3
66	Temperature and light effects on Trichobilharzia szidati cercariae with implications for a risk analysis. *Acta Veterinaria Scandinavica*, **2020**, 62, 54	2	6
65	A spot diagnosis! Aplasia cutis congenita in monozygotic twins. *Lancet, The*, **2019**, 394, 868	40	1
64	Chinese herbal remedy found to contain steroids and antifungals. *Lancet, The*, **2019**, 393, 446	40	6
63	Reporting through smartphone application results in detailed data on acquired and hereditary angioedema attacks. *Allergy: European Journal of Allergy and Clinical Immunology*, **2019**, 74, 1800-1802	9.3	2
62	Genetic Variation of Kallikrein-Kinin System and Related Genes in Patients With Hereditary Angioedema. *Frontiers in Medicine*, **2019**, 6, 28	4.9	8
61	Gene Editing-Mediated Disruption of Epidermolytic Ichthyosis-Associated KRT10 Alleles Restores Filament Stability in Keratinocytes. *Journal of Investigative Dermatology*, **2019**, 139, 1699-1710.e6	4.3	15
60	Elderly versus younger patients with hereditary angioedema type I/II: patient characteristics and safety analysis from the Icatibant Outcome Survey. *Clinical and Translational Allergy*, **2019**, 9, 37	5.2	7
59	Hereditary Angio-Oedema for Dermatologists. *Dermatology*, **2019**, 235, 263-275	4.4	2
58	Genetic susceptibility to angiotensin-converting enzyme-inhibitor induced angioedema: A systematic review and evaluation of methodological approaches. *PLoS ONE*, **2019**, 14, e0224858	3.7	4
57	Dominant-negative SERPING1 variants cause intracellular retention of C1 inhibitor in hereditary angioedema. *Journal of Clinical Investigation*, **2019**, 129, 388-405	15.9	24

56	ACE-Inhibitor Related Angioedema Is Not Sufficiently Reported to the Danish Adverse Drug Reactions Database. *Drug, Healthcare and Patient Safety*, **2019**, 11, 105-113	1.6	1
55	Angiotensin II receptor blockers are safe in patients with prior angioedema related to angiotensin-converting enzyme inhibitors - a nationwide registry-based cohort study. *Journal of Internal Medicine*, **2019**, 285, 553-561	10.8	14
54	Pediatricians diagnosed few patients with childhood-presented hereditary angioedema: Icatibant Outcome Survey findings. *Journal of Allergy and Clinical Immunology: in Practice*, **2019**, 7, 1078-1080	5.4	3
53	Genetic susceptibility to angiotensin-converting enzyme-inhibitor induced angioedema: A systematic review and evaluation of methodological approaches **2019**, 14, e0224858		
52	Genetic susceptibility to angiotensin-converting enzyme-inhibitor induced angioedema: A systematic review and evaluation of methodological approaches **2019**, 14, e0224858		
51	Genetic susceptibility to angiotensin-converting enzyme-inhibitor induced angioedema: A systematic review and evaluation of methodological approaches **2019**, 14, e0224858		
50	Genetic susceptibility to angiotensin-converting enzyme-inhibitor induced angioedema: A systematic review and evaluation of methodological approaches **2019**, 14, e0224858		
49	Diagnosing diabetes mellitus in patients with porphyria cutanea tarda. *International Journal of Dermatology*, **2018**, 57, 763-769	1.7	1
48	Impact of red and processed meat and fibre intake on treatment outcomes among patients with chronic inflammatory diseases: protocol for a prospective cohort study of prognostic factors and personalised medicine. *BMJ Open*, **2018**, 8, e018166	3	11
47	Improvement in diagnostic delays over time in patients with hereditary angioedema: findings from the Icatibant Outcome Survey. *Clinical and Translational Allergy*, **2018**, 8, 42	5.2	15
46	Atrophoderma Vermiculatum in a 12-Year-Old Girl. *Skinmed*, **2018**, 16, 421-422	0.2	
45	Drug-induced cutaneous lupus erythematosus: 88 new cases. *European Journal of Dermatology*, **2017**, 27, 28-33	0.8	44
44	Complement factor C4 activation in patients with hereditary angioedema. *Clinical Biochemistry*, **2017**, 50, 816-821	3.5	16
43	Clinical characteristics and real-life diagnostic approaches in all Danish children with hereditary angioedema. *Orphanet Journal of Rare Diseases*, **2017**, 12, 55	4.2	19
42	Assessment of 105 Patients with Angiotensin Converting Enzyme-Inhibitor Induced Angioedema. *International Journal of Otolaryngology*, **2017**, 2017, 1476402	1.4	10
41	A Proposal for a Study on Treatment Selection and Lifestyle Recommendations in Chronic Inflammatory Diseases: A Danish Multidisciplinary Collaboration on Prognostic Factors and Personalised Medicine. *Nutrients*, **2017**, 9,	6.7	16
40	Disease Severity, Activity, Impact, and Control and How to Assess Them in Patients with Hereditary Angioedema. *Frontiers in Medicine*, **2017**, 4, 212	4.9	32
39	Safety and Usage of C1-Inhibitor in Hereditary Angioedema: Berinert Registry Data. *Journal of Allergy and Clinical Immunology: in Practice*, **2016**, 4, 963-71	5.4	40

38	The Humanistic, Societal, and Pharmaco-economic Burden of Angioedema. *Clinical Reviews in Allergy and Immunology*, **2016**, 51, 230-9	12.3	58
37	Mutations in Three Genes Encoding Proteins Involved in Hair Shaft Formation Cause Uncombable Hair Syndrome. *American Journal of Human Genetics*, **2016**, 99, 1292-1304	11	72
36	The first Danish family reported with an AQP5 mutation presenting diffuse non-epidermolytic palmoplantar keratoderma of Bothnian type, hyperhidrosis and frequent Corynebacterium infections: a case report. *BMC Dermatology*, **2016**, 16, 7	2.1	10
35	Use of a C1 Inhibitor Concentrate in Adults ≥65 Years of Age with Hereditary Angioedema: Findings from the International Berinert (C1-INH) Registry. *Drugs and Aging*, **2016**, 33, 819-827	4.7	7
34	Unmasking sarcoidosis following surgery for Cushing disease. *Dermato-Endocrinology*, **2016**, 8, e983688		4
33	Psychometric Field Study of Hereditary Angioedema Quality of Life Questionnaire for Adults: HAE-QoL. *Journal of Allergy and Clinical Immunology: in Practice*, **2016**, 4, 464-473.e4	5.4	34
32	Estimation of EuroQol 5-Dimensions health status utility values in hereditary angioedema. *Patient Preference and Adherence*, **2016**, 10, 1699-707	2.4	31
31	The Use of Plasma-Derived Complement C1-Esterase Inhibitor Concentrate (Berinert®) in the Treatment of Angiotensin Converting Enzyme-Inhibitor Related Angioedema. *Case Reports in Emergency Medicine*, **2016**, 2016, 3930923	0.6	5
30	Eczema Coxsackium Caused by Coxsackievirus A6. *Pediatric Dermatology*, **2016**, 33, e230-1	1.9	6
29	Real-life experience with long-term prophylactic C1 inhibitor concentrate treatment of patients with hereditary angioedema: Effectiveness and cost. *Annals of Allergy, Asthma and Immunology*, **2016**, 116, 476-7	3.2	6
28	Second episode of near-fatal angioedema in a patient treated with everolimus. *Annals of Allergy, Asthma and Immunology*, **2015**, 115, 152-3	3.2	5
27	Case report: a novel KERA mutation associated with cornea plana and its predicted effect on protein function. *BMC Medical Genetics*, **2015**, 16, 40	2.1	7
26	Burden of Illness in Hereditary Angioedema: A Conceptual Model. *Acta Dermato-Venereologica*, **2015**, 95, 706-10	2.2	55
25	Idiopathic Thrombocytopenic Purpura Misdiagnosed as Hereditary Angioedema. *Case Reports in Dermatological Medicine*, **2015**, 2015, 934247	0.8	
24	Safety of C1-esterase inhibitor in acute and prophylactic therapy of hereditary angioedema: findings from the ongoing international Berinert patient registry. *Journal of Allergy and Clinical Immunology: in Practice*, **2015**, 3, 213-9	5.4	24
23	Pathogenicity of POFUT1 in Dowling-Degos disease: additional mutations and clinical overlap with reticulate acropigmentation of kitamura. *Journal of Investigative Dermatology*, **2015**, 135, 615-618	4.3	22
22	Angioedema in pediatric heart transplant recipients-reporting C1-esterase inhibitor deficiency without analysing protein. *Pediatric Radiology*, **2014**, 44, 899	2.8	
21	Mutations in POGLUT1, encoding protein O-glucosyltransferase 1, cause autosomal-dominant Dowling-Degos disease. *American Journal of Human Genetics*, **2014**, 94, 135-43	11	110

20	Linear eruptions on the calf in a six-week-old boy: a quiz. *Acta Dermato-Venereologica*, **2014**, 94, 125-6	2.2	2
19	Antiviral treatment of a boy with EBV-associated hydroa vacciniforme. *BMJ Case Reports*, **2014**, 2014,	0.9	3
18	Isolated oedema of the uvula induced by intense snoring and ACE inhibitor. *BMJ Case Reports*, **2014** , 2014,	0.9	8
17	Angiotensin-converting enzyme inhibitor-induced angioedema--a dangerous new epidemic. *Acta Dermato-Venereologica*, **2014**, 94, 260-4	2.2	31
16	Hereditary angioedema - consequences of a new treatment paradigm in Denmark. *Acta Dermato-Venereologica*, **2014**, 94, 436-41	2.2	29
15	Presence of C1-inhibitor polymers in a subset of patients suffering from hereditary angioedema. *PLoS ONE*, **2014**, 9, e112051	3.7	13
14	Generalized and naevoid epidermolytic ichthyosis in Denmark: clinical and mutational findings. *Acta Dermato-Venereologica*, **2013**, 93, 309-13	2.2	9
13	Xeroderma Pigmentosum-Trichothiodystrophy overlap patient with novel XPD/ERCC2 mutation. *Rare Diseases (Austin, Tex)*, **2013**, 1, e24932		7
12	Acquired angioedema--occurrence, clinical features and associated disorders in a Danish nationwide patient cohort. *International Archives of Allergy and Immunology*, **2013**, 162, 149-55	3.7	31
11	The hereditary angioedema burden of illness study in Europe (HAE-BOIS-Europe): background and methodology. *BMC Dermatology*, **2012**, 12, 4	2.1	28
10	International consensus and practical guidelines on the gynecologic and obstetric management of female patients with hereditary angioedema caused by C1 inhibitor deficiency. *Journal of Allergy and Clinical Immunology*, **2012**, 129, 308-20	11.5	159
9	Angioedema--assessment and treatment. *Tidsskrift for Den Norske Laegeforening*, **2012**, 132, 2391-5	3.5	14
8	Systemic epidermal nevus with involvement of the oral mucosa due to FGFR3 mutation. *BMC Medical Genetics*, **2011**, 12, 79	2.1	16
7	Self-administration of intravenous C1-inhibitor therapy for hereditary angioedema and associated quality of life benefits. *European Journal of Dermatology*, **2009**, 19, 147-51	0.8	100
6	Ichthyosis prematurity syndrome: a well-defined congenital ichthyosis subtype. *Journal of the American Academy of Dermatology*, **2008**, 59, S71-4	4.5	32
5	KID Syndrome: report of a Scandinavian patient with connexin-26 gene mutation. *Acta Dermato-Venereologica*, **2005**, 85, 152-5	2.2	12
4	Atopy patch tests in young adult patients with atopic dermatitis and controls: dose-response relationship, objective reading, reproducibility and clinical interpretation. *Acta Dermato-Venereologica*, **2003**, 83, 18-23	2.2	16
3	Familial and sporadic porphyria cutanea tarda: clinical, biochemical and genetic features with emphasis on iron status. *Acta Dermato-Venereologica*, **2003**, 83, 115-20	2.2	18

List of Publications

2	Denaturing Gradient Gel Electrophoresis Analysis of the Hemochromatosis (HFE) Gene: Impact of HFE Gene Mutations on the Manifestation of Porphyria Cutanea Tarda. *Clinical Chemistry*, **1999**, 45, 2025-2026	55	12
1	Mosaicism in Tuberous Sclerosis Complex: A Case Report, Literature Review, and Original Data from Danish Hospitals98-105		0