Anette Bygum

List of Publications by Citations

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

#	Paper	IF	Citations
91	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
90	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
89	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
88	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
87	The Humanistic, Societal, and Pharmaco-economic Burden of Angioedema. *Clinical Reviews in Allergy and Immunology*, **2016**, 51, 230-9	12.3	58
86	Burden of Illness in Hereditary Angioedema: A Conceptual Model. *Acta Dermato-Venereologica*, **2015**, 95, 706-10	2.2	55
85	Drug-induced cutaneous lupus erythematosus: 88 new cases. *European Journal of Dermatology*, **2017**, 27, 28-33	0.8	44
84	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
83	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
82	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
81	Ichthyosis prematurity syndrome: a well-defined congenital ichthyosis subtype. *Journal of the American Academy of Dermatology*, **2008**, 59, S71-4	4.5	32
80	Angiotensin-converting enzyme inhibitor-induced angioedema--a dangerous new epidemic. *Acta Dermato-Venereologica*, **2014**, 94, 260-4	2.2	31
79	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
78	Estimation of EuroQol 5-Dimensions health status utility values in hereditary angioedema. *Patient Preference and Adherence*, **2016**, 10, 1699-707	2.4	31
77	Hereditary angioedema - consequences of a new treatment paradigm in Denmark. *Acta Dermato-Venereologica*, **2014**, 94, 436-41	2.2	29
76	The hereditary angioedema burden of illness study in Europe (HAE-BOIS-Europe): background and methodology. *BMC Dermatology*, **2012**, 12, 4	2.1	28
75	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

74	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
73	Dominant-negative SERPING1 variants cause intracellular retention of C1 inhibitor in hereditary angioedema. *Journal of Clinical Investigation*, **2019**, 129, 388-405	15.9	24
72	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
71	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
70	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
69	Complement factor C4 activation in patients with hereditary angioedema. *Clinical Biochemistry*, **2017**, 50, 816-821	3.5	16
68	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
67	Systemic epidermal nevus with involvement of the oral mucosa due to FGFR3 mutation. *BMC Medical Genetics*, **2011**, 12, 79	2.1	16
66	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
65	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
64	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
63	Angioedema--assessment and treatment. *Tidsskrift for Den Norske Laegeforening*, **2012**, 132, 2391-5	3.5	14
62	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
61	Presence of C1-inhibitor polymers in a subset of patients suffering from hereditary angioedema. *PLoS ONE*, **2014**, 9, e112051	3.7	13
60	KID Syndrome: report of a Scandinavian patient with connexin-26 gene mutation. *Acta Dermato-Venereologica*, **2005**, 85, 152-5	2.2	12
59	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	5.5	12
58	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
57	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

56	Assessment of 105 Patients with Angiotensin Converting Enzyme-Inhibitor Induced Angioedema. *International Journal of Otolaryngology*, **2017**, 2017, 1476402	1.4	10
55	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
54	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
53	Generalized and naevoid epidermolytic ichthyosis in Denmark: clinical and mutational findings. *Acta Dermato-Venereologica*, **2013**, 93, 309-13	2.2	9
52	Genetic Variation of Kallikrein-Kinin System and Related Genes in Patients With Hereditary Angioedema. *Frontiers in Medicine*, **2019**, 6, 28	4.9	8
51	Isolated oedema of the uvula induced by intense snoring and ACE inhibitor. *BMJ Case Reports*, **2014** , 2014,	0.9	8
50	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
49	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
48	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
47	Xeroderma Pigmentosum-Trichothiodystrophy overlap patient with novel XPD/ERCC2 mutation. *Rare Diseases (Austin, Tex)*, **2013**, 1, e24932		7
46	Chinese herbal remedy found to contain steroids and antifungals. *Lancet, The*, **2019**, 393, 446	40	6
45	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
44	Temperature and light effects on Trichobilharzia szidati cercariae with implications for a risk analysis. *Acta Veterinaria Scandinavica*, **2020**, 62, 54	2	6
43	Eczema Coxsackium Caused by Coxsackievirus A6. *Pediatric Dermatology*, **2016**, 33, e230-1	1.9	6
42	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
41	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
40	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
39	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

38	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
37	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
36	Meta-Analysis of Mutations in or Identified in a Large Cohort of 224 Patients. *Genes*, **2021**, 12,	4.2	5
35	Hereditary leiomyomatosis and renal cell carcinoma: a case series and literature review. *Orphanet Journal of Rare Diseases*, **2021**, 16, 34	4.2	5
34	Introducing teledermoscopy of possible skin cancers in general practice in Southern Denmark. *Family Practice*, **2020**, 37, 513-518	1.9	4
33	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
32	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
31	Unmasking sarcoidosis following surgery for Cushing disease. *Dermato-Endocrinology*, **2016**, 8, e983688		4
30	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
29	Antiviral treatment of a boy with EBV-associated hydroa vacciniforme. *BMJ Case Reports*, **2014**, 2014,	0.9	3
28	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
27	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
26	Consensus on treatment goals in hereditary angioedema: A global Delphi initiative. *Journal of Allergy and Clinical Immunology*, **2021**, 148, 1526-1532	11.5	3
25	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
24	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
23	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
22	Hereditary Angio-Oedema for Dermatologists. *Dermatology*, **2019**, 235, 263-275	4.4	2
21	Linear eruptions on the calf in a six-week-old boy: a quiz. *Acta Dermato-Venereologica*, **2014**, 94, 125-6	2.2	2

20	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
19	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
18	Acceptance of Teledermoscopy by General Practitioners and Dermatologists in Denmark. *Dermatology Practical and Conceptual*, **2021**, 11, e2021033	1.5	2
17	Outcomes of Systemic Treatment in Children and Adults With Netherton Syndrome: A Systematic Review.. *Frontiers in Immunology*, **2022**, 13, 864449	8.4	2
16	A spot diagnosis! Aplasia cutis congenita in monozygotic twins. *Lancet, The*, **2019**, 394, 868	40	1
15	Diagnosing diabetes mellitus in patients with porphyria cutanea tarda. *International Journal of Dermatology*, **2018**, 57, 763-769	1.7	1
14	The impact of adolescentsReveryday life experiences on their primary hyperhidrosis treatment - a qualitative study. *Journal of Dermatological Treatment*, **2020**, 1-7	2.8	1
13	Spontaneous pneumothorax as a clinical manifestation of neurofibromatosis type 1. *BMJ Case Reports*, **2021**, 14,	0.9	1
12	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
11	Mosaicism in Tuberous Sclerosis Complex: A Case Report, Literature Review, and Original Data from Danish Hospitals98-105		0
10	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
9	Hereditary angioedema: the challenges of cross-border family investigation and treatment. *BMJ Case Reports*, **2020**, 13,	0.9	
8	Angioedema in pediatric heart transplant recipients-reporting C1-esterase inhibitor deficiency without analysing protein. *Pediatric Radiology*, **2014**, 44, 899	2.8	
7	Idiopathic Thrombocytopenic Purpura Misdiagnosed as Hereditary Angioedema. *Case Reports in Dermatological Medicine*, **2015**, 2015, 934247	0.8	
6	Atrophoderma Vermiculatum in a 12-Year-Old Girl. *Skinmed*, **2018**, 16, 421-422	0.2	
5	A girl with unruly locks: molecular genetics makes a diagnosis of uncombable hair syndrome.. *Lancet, The*, **2022**, 399, 1079	40	
4	Genetic susceptibility to angiotensin-converting enzyme-inhibitor induced angioedema: A systematic review and evaluation of methodological approaches **2019**, 14, e0224858		
3	Genetic susceptibility to angiotensin-converting enzyme-inhibitor induced angioedema: A systematic review and evaluation of methodological approaches **2019**, 14, e0224858		

List of Publications

2 Genetic susceptibility to angiotensin-converting enzyme-inhibitor induced angioedema: A systematic review and evaluation of methodological approaches **2019**, 14, e0224858

1 Genetic susceptibility to angiotensin-converting enzyme-inhibitor induced angioedema: A systematic review and evaluation of methodological approaches **2019**, 14, e0224858