

Konrad Bork

List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

156 papers	10,699 citations	51 h-index	101 g-index
206 ext. papers	12,361 ext. citations	5.9 avg, IF	6.28 L-index

#	Paper	IF	Citations
156	Attenuated androgen discontinuation in patients with hereditary angioedema: a commented case series.. <i>Allergy, Asthma and Clinical Immunology</i> , 2022 , 18, 4	3.2	0
155	The international WAO/EAACI guideline for the management of hereditary angioedema - the 2021 revision and update.. <i>Allergy: European Journal of Allergy and Clinical Immunology</i> , 2022 ,	9.3	6
154	The international WAO/EAACI guideline for the management of hereditary angioedema - The 2021 revision and update.. <i>World Allergy Organization Journal</i> , 2022 , 15, 100627	5.2	6
153	Uncommon Signs Associated With Hereditary Angioedema With Normal C1 Inhibitor. <i>Journal of Investigational Allergology and Clinical Immunology</i> , 2021 , 31, 257-258	2.3	1
152	Assessment and management of disease burden and quality of life in patients with hereditary angioedema: a consensus report. <i>Allergy, Asthma and Clinical Immunology</i> , 2021 , 17, 40	3.2	5
151	Mitigating Disparity in Health-care Resources Between Countries for Management of Hereditary Angioedema. <i>Clinical Reviews in Allergy and Immunology</i> , 2021 , 61, 84-97	12.3	3
150	The Expanding Spectrum of Mutations in Hereditary Angioedema. <i>Journal of Allergy and Clinical Immunology: in Practice</i> , 2021 , 9, 2229-2234	5.4	11
149	A germline mutation in the platelet-derived growth factor receptor beta gene may be implicated in hereditary progressive mucinous histiocytosis. <i>British Journal of Dermatology</i> , 2021 , 184, 967-970	4	5
148	Current and Prospective Targets of Pharmacologic Treatment of Hereditary Angioedema Types 1 and 2. <i>Clinical Reviews in Allergy and Immunology</i> , 2021 , 61, 66-76	12.3	5
147	Novel hereditary angioedema linked with a heparan sulfate 3-O-sulfotransferase 6 gene mutation. <i>Journal of Allergy and Clinical Immunology</i> , 2021 , 148, 1041-1048	11.5	18
146	Analysis of cold activation of the contact system in hereditary angioedema with normal C1 inhibitor. <i>Molecular Immunology</i> , 2021 , 136, 150-160	4.3	0
145	Hereditary angioedema in children and adolescents - A consensus update on therapeutic strategies for German-speaking countries. <i>Pediatric Allergy and Immunology</i> , 2020 , 31, 974-989	4.2	7
144	Hereditary angioedema in a single family with specific mutations in both plasminogen and SERPING1 genes. <i>JDDG - Journal of the German Society of Dermatology</i> , 2020 , 18, 215-223	1.2	5
143	Measurement of Bradykinin Formation and Degradation in Blood Plasma: Relevance for Acquired Angioedema Associated With Angiotensin Converting Enzyme Inhibition and for Hereditary Angioedema Due to Factor XII or Plasminogen Gene Variants. <i>Frontiers in Medicine</i> , 2020 , 7, 358	4.9	7
142	sgp120 and the contact system in hereditary angioedema: A diagnostic tool in HAE with normal C1 inhibitor. <i>Molecular Immunology</i> , 2020 , 119, 27-34	4.3	6
141	Definition, aims, and implementation of GA LEN/HAEi Angioedema Centers of Reference and Excellence. <i>Allergy: European Journal of Allergy and Clinical Immunology</i> , 2020 , 75, 2115-2123	9.3	10
140	Treatment of patients with hereditary angioedema with the c.988A>G (p.Lys330Glu) variant in the plasminogen gene. <i>Orphanet Journal of Rare Diseases</i> , 2020 , 15, 52	4.2	10

139	International Consensus on the Use of Genetics in the Management of Hereditary Angioedema. <i>Journal of Allergy and Clinical Immunology: in Practice</i> , 2020 , 8, 901-911	5.4	28
138	Hereditäres Angioödem. <i>Padiatrie Und Padologie</i> , 2020 , 55, 12-20	0	
137	Hereditäres Angioödem. <i>Hautnah</i> , 2020 , 19, 30-37	0.2	0
136	Clinical features of genetically characterized types of hereditary angioedema with normal C1 inhibitor: a systematic review of qualitative evidence. <i>Orphanet Journal of Rare Diseases</i> , 2020 , 15, 289	4.2	26
135	Subcutaneous C1 inhibitor for prevention of attacks of hereditary angioedema: additional outcomes and subgroup analysis of a placebo-controlled randomized study. <i>Allergy, Asthma and Clinical Immunology</i> , 2019 , 15, 49	3.2	7
134	Guideline: Hereditary angioedema due to C1 inhibitor deficiency. <i>Allergo Journal International</i> , 2019 , 28, 16-29	1.5	17
133	Hereditary angioedema cosegregating with a novel kininogen 1 gene mutation changing the N-terminal cleavage site of bradykinin. <i>Allergy: European Journal of Allergy and Clinical Immunology</i> , 2019 , 74, 2479-2481	9.3	72
132	Long-Term Outcomes with Subcutaneous C1-Inhibitor Replacement Therapy for Prevention of Hereditary Angioedema Attacks. <i>Journal of Allergy and Clinical Immunology: in Practice</i> , 2019 , 7, 1793-1802	5.4	42
131	Angioedema due to acquired C1-inhibitor deficiency: spectrum and treatment with C1-inhibitor concentrate. <i>Orphanet Journal of Rare Diseases</i> , 2019 , 14, 65	4.2	24
130	Leitlinie: Hereditäres Angioödem durch C1-Inhibitor-Mangel. <i>Allergo Journal</i> , 2019 , 28, 31-47	0	1
129	Hereditary angioedema: an update on causes, manifestations and treatment. <i>British Journal of Hospital Medicine (London, England: 2005)</i> , 2019 , 80, 391-398	0.8	34
128	Elderly versus younger patients with hereditary angioedema type I/II: patient characteristics and safety analysis from the Icatibant Outcome Survey. <i>Clinical and Translational Allergy</i> , 2019 , 9, 37	5.2	7
127	Neue Mittel in der Pipeline. <i>Der Deutsche Dermatologe</i> , 2019 , 67, 872-881	0	
126	The International/Canadian Hereditary Angioedema Guideline. <i>Allergy, Asthma and Clinical Immunology</i> , 2019 , 15, 72	3.2	68
125	Management of patients with hereditary angioedema in Germany: comparison with other countries in the Icatibant Outcome Survey. <i>Journal of the European Academy of Dermatology and Venereology</i> , 2019 , 33, 163-169	4.6	5
124	Genotype-phenotype correlations in Brazilian patients with hereditary angioedema due to C1 inhibitor deficiency. <i>Allergy: European Journal of Allergy and Clinical Immunology</i> , 2019 , 74, 1013-1016	9.3	8
123	The international WAO/EAACI guideline for the management of hereditary angioedema-The 2017 revision and update. <i>Allergy: European Journal of Allergy and Clinical Immunology</i> , 2018 , 73, 1575-1596	9.3	260
122	The international WAO/EAACI guideline for the management of hereditary angioedema The 2017 revision and update. <i>World Allergy Organization Journal</i> , 2018 , 11, 5	5.2	34

121	Antibody-mediated inhibition of FXIIa blocks downstream bradykinin generation. <i>Journal of Allergy and Clinical Immunology</i> , 2018 , 142, 1355-1358	11.5	19
120	On the pathogenicity of the plasminogen K330E mutation for hereditary angioedema. <i>Allergy: European Journal of Allergy and Clinical Immunology</i> , 2018 , 73, 1751-1753	9.3	21
119	Acquired and hereditary forms of recurrent angioedema: Update of treatment. <i>Allergologie Select</i> , 2018 , 2, 121-131	4.1	4
118	Hereditary angioedema with a mutation in the plasminogen gene. <i>Allergy: European Journal of Allergy and Clinical Immunology</i> , 2018 , 73, 442-450	9.3	148
117	Tamoxifen may cause life-threatening angioedema attacks in patients with hereditary angioedema. <i>Journal of the European Academy of Dermatology and Venereology</i> , 2017 , 31, e237-e239	4.6	4
116	Treatment for hereditary angioedema with normal C1-INH and specific mutations in the F12 gene (HAE-FXII). <i>Allergy: European Journal of Allergy and Clinical Immunology</i> , 2017 , 72, 320-324	9.3	53
115	Reply. <i>Journal of Allergy and Clinical Immunology</i> , 2017 , 139, 1720-1721	11.5	3
114	Blindness, tetraspasticity, and other signs of irreversible brain damage in hereditary angioedema. <i>Annals of Allergy, Asthma and Immunology</i> , 2017 , 118, 520-521	3.2	10
113	Prevention of Hereditary Angioedema Attacks with a Subcutaneous C1 Inhibitor. <i>New England Journal of Medicine</i> , 2017 , 376, 1131-1140	59.2	118
112	International consensus on the diagnosis and management of pediatric patients with hereditary angioedema with C1 inhibitor deficiency. <i>Allergy: European Journal of Allergy and Clinical Immunology</i> , 2017 , 72, 300-313	9.3	109
111	Shortened Activated Partial Thromboplastin Time May Help in Diagnosing Hereditary and Acquired Angioedema. <i>International Archives of Allergy and Immunology</i> , 2016 , 170, 101-7	3.7	11
110	Misdiagnosis trends in patients with hereditary angioedema from the real-world clinical setting. <i>Annals of Allergy, Asthma and Immunology</i> , 2016 , 117, 394-398	3.2	51
109	Deficiency of plasminogen activator inhibitor 2 in plasma of patients with hereditary angioedema with normal C1 inhibitor levels. <i>Journal of Allergy and Clinical Immunology</i> , 2016 , 137, 1822-1829.e1	11.5	33
108	Acute Edema Blisters on a Skin Swelling: An Unusual Manifestation of Hereditary Angioedema. <i>Acta Dermato-Venereologica</i> , 2016 , 96, 556-7	2.2	5
107	A Decade of Change: Recent Developments in Pharmacotherapy of Hereditary Angioedema (HAE). <i>Clinical Reviews in Allergy and Immunology</i> , 2016 , 51, 183-92	12.3	29
106	Efficacy of Different Medical Therapies for the Treatment of Acute Laryngeal Attacks of Hereditary Angioedema due to C1-esterase Inhibitor Deficiency. <i>Journal of Emergency Medicine</i> , 2016 , 50, 567-80.e1	1.5	14
105	Coagulation Factor XII Gene Mutation in Brazilian Families with Hereditary Angioedema with Normal C1 Inhibitor. <i>International Archives of Allergy and Immunology</i> , 2015 , 166, 114-20	3.7	14
104	Currently available treatments and future treatment options for hereditary angioedema. <i>Allergo Journal International</i> , 2015 , 24, 122-128	1.5	

103	Pathogenesis of Hereditary Angioedema with Normal C1 Inhibitor: Evidence for Abnormalities in Plasminogen Activator Inhibitors. <i>Journal of Allergy and Clinical Immunology</i> , 2015 , 135, AB277	11.5	2
102	Currently available treatments and future treatment options for hereditary angioedema. <i>Allergo Journal</i> , 2015 , 24, 32-38	0	
101	Analysis of characteristics associated with reinjection of icatibant: Results from the icatibant outcome survey. <i>Allergy and Asthma Proceedings</i> , 2015 , 36, 399-406	2.6	15
100	Efficacy of C1 esterase inhibitor concentrate in treatment of cutaneous attacks of hereditary angioedema. <i>Allergy and Asthma Proceedings</i> , 2015 , 36, 218-24	2.6	4
99	Hereditary angioedema with normal C1-INH with versus without specific F12 gene mutations. <i>Allergy: European Journal of Allergy and Clinical Immunology</i> , 2015 , 70, 1004-12	9.3	79
98	Hereditäres Angioödem durch C1-Inhibitor-Mangel 2015 , G3.1-G3.14		
97	Classification, diagnosis, and approach to treatment for angioedema: consensus report from the Hereditary Angioedema International Working Group. <i>Allergy: European Journal of Allergy and Clinical Immunology</i> , 2014 , 69, 602-16	9.3	410
96	Angioedema. <i>Immunology and Allergy Clinics of North America</i> , 2014 , 34, 23-31	3.3	27
95	The reply. <i>American Journal of Medicine</i> , 2014 , 127, e9	2.4	
94	Canadian hereditary angioedema guideline. <i>Allergy, Asthma and Clinical Immunology</i> , 2014 , 10, 50	3.2	61
93	Pasteurized and nanofiltered, plasma-derived C1 esterase inhibitor concentrate for the treatment of hereditary angioedema. <i>Immunotherapy</i> , 2014 , 6, 533-51	3.8	23
92	Characterization of a partial exon 9/intron 9 deletion in the coagulation factor XII gene (F12) detected in two Turkish families with hereditary angioedema and normal C1 inhibitor. <i>Haemophilia</i> , 2014 , 20, e372-5	3.3	23
91	An evidence based therapeutic approach to hereditary and acquired angioedema. <i>Current Opinion in Allergy and Clinical Immunology</i> , 2014 , 14, 354-62	3.3	19
90	Antihistamine-resistant angioedema in women with negative family history: estrogens and F12 gene mutations. <i>American Journal of Medicine</i> , 2013 , 126, 1142.e9-14	2.4	16
89	Hereditary angioedema with normal C1 inhibitor. <i>Immunology and Allergy Clinics of North America</i> , 2013 , 33, 457-70	3.3	39
88	Treatment with C1-esterase inhibitor concentrate in type I or II hereditary angioedema: a systematic literature review. <i>Allergy and Asthma Proceedings</i> , 2013 , 34, 312-27	2.6	26
87	Current status of implementation of self-administration training in various regions of Europe, Canada and the USA in the management of hereditary angioedema. <i>International Archives of Allergy and Immunology</i> , 2013 , 161 Suppl 1, 10-6	3.7	28
86	Overview of hereditary angioedema caused by C1-inhibitor deficiency: assessment and clinical management. <i>European Annals of Allergy and Clinical Immunology</i> , 2013 , 45, 7-16	1.3	26

85	WAO Guideline for the Management of Hereditary Angioedema. <i>World Allergy Organization Journal</i> , 2012 , 5, 182-99	5.2	222
84	Hereditäres Angioödem durch C1-Inhibitor-Mangel. <i>Allergo Journal</i> , 2012 , 21, 109-120	0	16
83	International consensus and practical guidelines on the gynecologic and obstetric management of female patients with hereditary angioedema caused by C1 inhibitor deficiency. <i>Journal of Allergy and Clinical Immunology</i> , 2012 , 129, 308-20	11.5	159
82	Review of the Long-Term Safety of a Human Pasteurized C1 Inhibitor Concentrate. <i>Journal of Allergy and Clinical Immunology</i> , 2012 , 129, AB222	11.5	3
81	Fatal laryngeal attacks and mortality in hereditary angioedema due to C1-INH deficiency. <i>Journal of Allergy and Clinical Immunology</i> , 2012 , 130, 692-7	11.5	272
80	Current management options for hereditary angioedema. <i>Current Allergy and Asthma Reports</i> , 2012 , 12, 273-80	5.6	19
79	Evidence-based recommendations for the therapeutic management of angioedema owing to hereditary C1 inhibitor deficiency: consensus report of an International Working Group. <i>Allergy: European Journal of Allergy and Clinical Immunology</i> , 2012 , 67, 147-57	9.3	254
78	New drug targets and future direction: icatibant 2012 , 58-69		
77	Hereditary angioedema with normal C1 inhibitor function: consensus of an international expert panel. <i>Allergy and Asthma Proceedings</i> , 2012 , 33 Suppl 1, S145-56	2.6	121
76	Risk of laryngeal edema and facial swellings after tooth extraction in patients with hereditary angioedema with and without prophylaxis with C1 inhibitor concentrate: a retrospective study. <i>Oral Surgery Oral Medicine Oral Pathology Oral Radiology and Endodontics</i> , 2011 , 112, 58-64		96
75	A single nucleotide deletion at the C1 inhibitor gene as the cause of hereditary angioedema: insights from a Brazilian family. <i>Allergy: European Journal of Allergy and Clinical Immunology</i> , 2011 , 66, 1384-90	9.3	12
74	Long-term efficacy of danazol treatment in hereditary angioedema. <i>European Journal of Clinical Investigation</i> , 2011 , 41, 256-62	4.6	45
73	A novel mutation in the coagulation factor 12 gene in subjects with hereditary angioedema and normal C1-inhibitor. <i>Clinical Immunology</i> , 2011 , 141, 31-5	9	98
72	Hereditary angioedema: long-term treatment with one or more injections of C1 inhibitor concentrate per week. <i>International Archives of Allergy and Immunology</i> , 2011 , 154, 81-8	3.7	48
71	Human pasteurized C1-inhibitor concentrate for the treatment of hereditary angioedema due to C1-inhibitor deficiency. <i>Expert Review of Clinical Immunology</i> , 2011 , 7, 723-33	5.1	5
70	ALDEN, an algorithm for assessment of drug causality in Stevens-Johnson Syndrome and toxic epidermal necrolysis: comparison with case-control analysis. <i>Clinical Pharmacology and Therapeutics</i> , 2010 , 88, 60-8	6.1	359
69	Icatibant, a new bradykinin-receptor antagonist, in hereditary angioedema. <i>New England Journal of Medicine</i> , 2010 , 363, 532-41	59.2	387
68	Economic costs associated with acute attacks and long-term management of hereditary angioedema. <i>Annals of Allergy, Asthma and Immunology</i> , 2010 , 104, 314-20	3.2	119

67	Diagnosis and treatment of hereditary angioedema with normal C1 inhibitor. <i>Allergy, Asthma and Clinical Immunology</i> , 2010 , 6, 15	3.2	75
66	HAE international home therapy consensus document. <i>Allergy, Asthma and Clinical Immunology</i> , 2010 , 6, 22	3.2	129
65	2010 International consensus algorithm for the diagnosis, therapy and management of hereditary angioedema. <i>Allergy, Asthma and Clinical Immunology</i> , 2010 , 6, 24	3.2	333
64	Recurrent angioedema and the threat of asphyxiation. <i>Deutsches A&#x0308;rztblatt International</i> , 2010 , 107, 408-14	2.5	24
63	Hereditary angioedema with normal c1 inhibition. <i>Current Allergy and Asthma Reports</i> , 2009 , 9, 280-5	5.6	17
62	Hereditary angioedema: increased number of attacks after frequent treatments with C1 inhibitor concentrate. <i>American Journal of Medicine</i> , 2009 , 122, 780-3	2.4	39
61	Hereditary angioedema caused by missense mutations in the factor XII gene: clinical features, trigger factors, and therapy. <i>Journal of Allergy and Clinical Immunology</i> , 2009 , 124, 129-34	11.5	144
60	Kallikrein-kinin system and fibrinolysis in hereditary angioedema due to factor XII gene mutation Thr309Lys. <i>Blood Coagulation and Fibrinolysis</i> , 2009 , 20, 325-32	1	38
59	Treatment of skin swellings with C1-inhibitor concentrate in patients with hereditary angio-oedema. <i>Allergy: European Journal of Allergy and Clinical Immunology</i> , 2008 , 63, 751-7	9.3	66
58	Functional C1-inhibitor diagnostics in hereditary angioedema: assay evaluation and recommendations. <i>Journal of Immunological Methods</i> , 2008 , 338, 14-20	2.5	63
57	Mutational spectrum of the C1INH (SERPING1) gene in patients with hereditary angioedema. <i>Cytogenetic and Genome Research</i> , 2008 , 121, 181-8	1.9	57
56	Quel avenir pour les nouvelles th&eap;apeutiques ?. <i>Revue Francaise D&allergologie Et D&immunologie Clinique</i> , 2008 , 48, 159-161		
55	Benefits and risks of danazol in hereditary angioedema: a long-term survey of 118 patients. <i>Annals of Allergy, Asthma and Immunology</i> , 2008 , 100, 153-61	3.2	171
54	Hereditary angiodema: a current state-of-the-art review, VII: Canadian Hungarian 2007 International Consensus Algorithm for the Diagnosis, Therapy, and Management of Hereditary Angioedema. <i>Annals of Allergy, Asthma and Immunology</i> , 2008 , 100, S30-40	3.2	162
53	Pasteurized C1 inhibitor concentrate in hereditary angioedema: pharmacology, safety, efficacy and future directions. <i>Expert Review of Clinical Immunology</i> , 2008 , 4, 13-20	5.1	35
52	Disease expression in women with hereditary angioedema. <i>American Journal of Obstetrics and Gynecology</i> , 2008 , 199, 484.e1-4	6.4	97
51	The European Register of Hereditary Angioedema: Experience and Preliminary Results. <i>Journal of Allergy and Clinical Immunology</i> , 2007 , 119, S276	11.5	4
50	Treatment of acute edema attacks in hereditary angioedema with a bradykinin receptor-2 antagonist (Icatibant). <i>Journal of Allergy and Clinical Immunology</i> , 2007 , 119, 1497-503	11.5	172

49	Helicobacter pylori infection as a triggering factor of attacks in patients with hereditary angioedema. <i>Helicobacter</i> , 2007 , 12, 251-7	4.9	46
48	Hereditary angioedema with normal C1 inhibitor: clinical symptoms and course. <i>American Journal of Medicine</i> , 2007 , 120, 987-92	2.4	101
47	Symptoms, course, and complications of abdominal attacks in hereditary angioedema due to C1 inhibitor deficiency. <i>American Journal of Gastroenterology</i> , 2006 , 101, 619-27	0.7	192
46	Hereditary angioedema with normal C1 inhibitor activity including hereditary angioedema with coagulation factor XII gene mutations. <i>Immunology and Allergy Clinics of North America</i> , 2006 , 26, 709-24 ^{3.3}		50
45	Increased activity of coagulation factor XII (Hageman factor) causes hereditary angioedema type III. <i>American Journal of Human Genetics</i> , 2006 , 79, 1098-104	11	264
44	Hereditary angioedema: new findings concerning symptoms, affected organs, and course. <i>American Journal of Medicine</i> , 2006 , 119, 267-74	2.4	494
43	Missense mutations in the coagulation factor XII (Hageman factor) gene in hereditary angioedema with normal C1 inhibitor. <i>Biochemical and Biophysical Research Communications</i> , 2006 , 343, 1286-9	3.4	288
42	Hereditary angioedema: causes, manifestations and treatment. <i>British Journal of Hospital Medicine (London, England: 2005)</i> , 2006 , 67, 654-7	0.8	22
41	Hereditary angio-oedema with normal C1 inhibitor in a family with affected women and men. <i>British Journal of Dermatology</i> , 2006 , 154, 542-5	4	57
40	Missense Mutations in the Proline-Rich Region of Coagulation Factor XII in Hereditary and Idiopathic Angioedema.. <i>Blood</i> , 2006 , 108, 1619-1619	2.2	
39	Treatment with C1 inhibitor concentrate in abdominal pain attacks of patients with hereditary angioedema. <i>Transfusion</i> , 2005 , 45, 1774-84	2.9	112
38	Pruritus precipitated by hydroxyethyl starch: a review. <i>British Journal of Dermatology</i> , 2005 , 152, 3-12	4	129
37	Canadian 2003 International Consensus Algorithm For the Diagnosis, Therapy, and Management of Hereditary Angioedema. <i>Journal of Allergy and Clinical Immunology</i> , 2004 , 114, 629-37	11.5	155
36	Hereditary and acquired angioedema: problems and progress: proceedings of the third C1 esterase inhibitor deficiency workshop and beyond. <i>Journal of Allergy and Clinical Immunology</i> , 2004 , 114, S51-131 ^{11.5}		490
35	Hereditary angioedema type III, angioedema associated with angiotensin II receptor antagonists, and female sex. <i>American Journal of Medicine</i> , 2004 , 116, 644-5	2.4	26
34	Laryngeal edema and death from asphyxiation after tooth extraction in four patients with hereditary angioedema. <i>Journal of the American Dental Association</i> , 2003 , 134, 1088-94	1.9	120
33	Clinical studies of sudden upper airway obstruction in patients with hereditary angioedema due to C1 esterase inhibitor deficiency. <i>Archives of Internal Medicine</i> , 2003 , 163, 1229-35		183
32	Sudden upper airway obstruction in patients with hereditary angioedema. <i>Transfusion and Apheresis Science</i> , 2003 , 29, 235-8	2.4	51

31	Recurrent episodes of skin angioedema and severe attacks of abdominal pain induced by oral contraceptives or hormone replacement therapy. <i>American Journal of Medicine</i> , 2003 , 114, 294-8	2.4	125
30	Danazol-induced hepatocellular adenoma in patients with hereditary angio-oedema. <i>Journal of Hepatology</i> , 2002 , 36, 707-9	13.4	63
29	Treatment of 193 episodes of laryngeal edema with C1 inhibitor concentrate in patients with hereditary angioedema. <i>Archives of Internal Medicine</i> , 2001 , 161, 714-8		170
28	Episodes of severe dyspnea caused by snoring-induced recurrent edema of the soft palate in hereditary angioedema. <i>Journal of the American Academy of Dermatology</i> , 2001 , 45, 968-9	4.5	15
27	Transfer of aciclovir from plasma to human breast milk. <i>Arzneimittelforschung</i> , 2000 , 50, 656-8		4
26	Hereditary angioedema and normal C1-inhibitor activity in women. <i>Lancet, The</i> , 2000 , 356, 1440-1441	4.0	4
25	Hereditary angioedema with normal C1-inhibitor activity in women. <i>Lancet, The</i> , 2000 , 356, 213-7	4.0	414
24	Successful Extubation With Use of C1 Esterase Inhibitor Concentrate in a Patient With Hereditary Angioedema: In Response. <i>Mayo Clinic Proceedings</i> , 2000 , 75, 870	6.4	2
23	Asphyxiation by laryngeal edema in patients with hereditary angioedema. <i>Mayo Clinic Proceedings</i> , 2000 , 75, 349-54	6.4	204
22	Leberzelladenome nach Langzeitprophylaxe mit Danazol bei drei Patienten mit hereditärem Angioödem 2000 , 204-206		
21	Hepatocellular adenomas in patients taking danazol for hereditary angio-oedema. <i>Lancet, The</i> , 1999 , 353, 1066-7	4.0	116
20	Hereditary Progressive Mucinous Histiocytosis. <i>Archives of Dermatology</i> , 1994 , 130, 1300		15
19	Hereditary progressive mucinous histiocytosis. Immunohistochemical and ultrastructural studies in an additional family. <i>Archives of Dermatology</i> , 1994 , 130, 1300-4		18
18	Multifocal aplasia cutis congenita, distal limb hemimelia, and cutis marmorata telangiectatica in a patient with Adams-Oliver syndrome. <i>British Journal of Dermatology</i> , 1992 , 127, 160-3	4	21
17	Pathogenesis of paraneoplastic follicular hyperkeratotic spicules in multiple myeloma. Follicular and epidermal accumulation of IgG dysprotein and cryoglobulin. <i>Archives of Dermatology</i> , 1990 , 126, 509-13		6
16	Long-term prophylaxis with C1-inhibitor (C1 INH) concentrate in patients with recurrent angioedema caused by hereditary and acquired C1-inhibitor deficiency. <i>Journal of Allergy and Clinical Immunology</i> , 1989 , 83, 677-82	11.5	147
15	Hereditary angioedema associated with subacute cutaneous lupus erythematosus. <i>Dermatology</i> , 1989 , 179, 211-3	4.4	20
14	Acquired C1 inhibitor (C1-INH) deficiency type II. Replacement therapy with C1-INH and analysis of patients' C1-INH and anti-C1-INH autoantibodies. <i>Journal of Clinical Investigation</i> , 1989 , 83, 1794-9	15.9	41

13	Hereditary Progressive Mucinous Histiocytosis in Women. <i>Archives of Dermatology</i> , 1988 , 124, 1225		24
12	Hereditary progressive mucinous histiocytosis in women. Report of three members in a family. <i>Archives of Dermatology</i> , 1988 , 124, 1225-9		
11	Acquired and Hereditary Angioedema: Pathogenesis and Therapy 1988 , 96-99		
10	Hereditary progressive mucinous histiocytosis in women. Report of three members in a family. <i>Archives of Dermatology</i> , 1988 , 124, 1225-1229		25
9	Autoantibody-mediated acquired deficiency of C1 inhibitor. <i>New England Journal of Medicine</i> , 1987 , 316, 1360-6	59.2	184
8	Impaired function of numerically augmented Fc-receptors on granulocytes in a HLA B8+ patient with palmoplantar pustulosis. <i>Archives of Dermatological Research</i> , 1987 , 279, 444-8	3.3	3
7	Stimulation of T cells by autologous mononuclear leukocytes and epidermal cells in psoriasis. <i>Archives of Dermatological Research</i> , 1986 , 279, 89-94	3.3	30
6	Hereditäres angioneurotisches Ödem bei Kindern [Klinik, Diagnostik und Therapie 1986 , 388-395		
5	Effects of ethanol and acetaldehyde on phagocytic functions. <i>Archives of Dermatological Research</i> , 1985 , 277, 131-7	3.3	30
4	Interaction between C1-INA, coagulation, fibrinolysis and kinin system in hereditary angioneurotic edema (HANE) and urticaria. <i>Archives of Dermatological Research</i> , 1984 , 276, 375-80	3.3	15
3	Stimulation of T lymphocytes by isolated autologous or allogeneic human epidermal cells 1984 , 189-194		1
2	Hereditary angioneurotic oedema and blood-coagulation: interaction between C1-esterase-inhibitor and the activation factors of the proteolytic enzyme systems. <i>Klinische Wochenschrift</i> , 1983 , 61, 1131-5		11
1	Das Verhalten einiger Serumenzyme nach ganzkörper-muskelmassage. <i>Archives of Dermatological Research</i> , 1971 , 240, 342-348	3.3	5