Marco Cicardi

List of Publications by Year
in descending order

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250 papers	15,782 citations	11651 70 h-index	18647 119 g-index
261 all docs	261 docs citations	261 times ranked	4789 citing authors

#	Article	IF	Citations
1	Therapeutic monoclonal antibodies with a focus on hereditary angioedema. Allergology International, 2023, 72, 54-62.	3.3	4
2	Effects of Continuous Plasma-Derived Subcutaneous C1-Esterase Inhibitor on Coagulation and Fibrinolytic Parameters. Thrombosis and Haemostasis, 2021, 121, 690-693.	3.4	9
3	Anti-C1-Inhibitor Autoantibody Detection by ELISA. Methods in Molecular Biology, 2021, 2227, 115-120.	0.9	0
4	Long-term health-related quality of life in patients treated with subcutaneous C1-inhibitor replacement therapy for the prevention of hereditary angioedema attacks: findings from the COMPACT open-label extension study. Orphanet Journal of Rare Diseases, 2021, 16, 86.	2.7	10
5	Hereditary angioedema due to C1 inhibitor deficiency in Belarus: epidemiology, access to diagnosis and seven novel mutations in SERPING1 gene. Clinical and Molecular Allergy, 2021, 19, 3.	1.8	14
6	Handling shock in idiopathic systemic capillary leak syndrome (Clarkson's disease) less is more: reply. Internal and Emergency Medicine, 2020, 15, 349-350.	2.0	1
7	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.	3.8	43
8	ACE inhibitor-mediated angioedema. International Immunopharmacology, 2020, 78, 106081.	3.8	55
9	Impaired control of the contact system in hereditary angioedema with normal C1â€inhibitor. Allergy: European Journal of Allergy and Clinical Immunology, 2020, 75, 1394-1403.	5.7	23
10	Long-Term Efficacy of Subcutaneous C1 Inhibitor in Pediatric Patients with Hereditary Angioedema. Pediatric, Allergy, Immunology, and Pulmonology, 2020, 33, 136-141.	0.8	10
11	Perioperative Management of Patients With Hereditary Angioedema With Special Considerations for Cardiopulmonary Bypass. Anesthesia and Analgesia, 2020, 131, 155-169.	2.2	5
12	The central role of endothelium in hereditary angioedema due to C1 inhibitor deficiency. International Immunopharmacology, 2020, 82, 106304.	3.8	15
13	Short-term prophylaxis in patients with angioedema due to C1-inhibitor deficiency undergoing dental procedures: An observational study. PLoS ONE, 2020, 15, e0230128.	2.5	7
14	Life expectancy in Italian patients with hereditary angioedema due to C1-inhibitor deficiency. Journal of Allergy and Clinical Immunology: in Practice, 2020, 8, 1772-1774.	3.8	5
15	Pediatric angioedema: Essential features and preliminary results from the Hereditary Angioedema Global Registry in Italy. Pediatric Allergy and Immunology, 2020, 31, 22-24.	2.6	6
16	Patterns of Treatment and Retreatment of Acute Attacks of Hereditary Angioedema (HAE) with Standard of Care (SOC) On-Demand Medication: Results from the APeX-2 Study. Journal of Allergy and Clinical Immunology, 2020, 145, AB107.	2.9	0
17	Long-term efficacy and safety of subcutaneous C1-inhibitor in women with hereditary angioedema: subgroup analysis from an open-label extension of a phase 3 trial. Allergy, Asthma and Clinical Immunology, 2020, 16, 8.	2.0	16
18	Hereditary Deficiency of C1 Inhibitor and Angioedema. , 2020, , 341-345.		0

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19	Gastrointestinal (GI) Adverse Events (AEs) Observed With Berotralstat (BCX7353) Treatment for Hereditary Angioedema (HAE) are Primarily Mild, Self-limited, and Diminish with Time on Treatment. Journal of Allergy and Clinical Immunology, 2020, 145, AB102.	2.9	4
20	Title is missing!. , 2020, 15, e0230128.		0
21	Title is missing!. , 2020, 15, e0230128.		0
22	Title is missing!. , 2020, 15, e0230128.		0
23	Title is missing!. , 2020, 15, e0230128.		0
24	Driving towards Precision Medicine for angioedema without wheals. Journal of Autoimmunity, 2019, 104, 102312.	6.5	9
25	Lanadelumab for the prevention of attacks in hereditary angioedema. Expert Review of Clinical Immunology, 2019, 15, 1239-1248.	3.0	1
26	Subcutaneous C1 inhibitor for prevention of attacks of hereditary angioedema: additional outcomes and subgroup analysis of a placebo-controlled randomized study. Allergy, Asthma and Clinical Immunology, 2019, 15, 49.	2.0	10
27	Clinical Features of Patients With Primary Angioedema With Normal Levels of C1-Inhibitor. Journal of Allergy and Clinical Immunology, 2019, 143, AB47.	2.9	0
28	Handling shock in idiopathic systemic capillary leak syndrome (Clarksonâ€™s disease): less is more. Internal and Emergency Medicine, 2019, 14, 723-730.	2.0	13
29	Paroxysmal Permeability Disorders: Development of a Microfluidic Device to Assess Endothelial Barrier Function. Frontiers in Medicine, 2019, 6, 89.	2.6	15
30	Oral Plasma Kallikrein Inhibitor BCX7353 is Safe and Effective as an On-Demand Treatment of Angioedema Attacks in Hereditary Angioedema (HAE) Patients: Results of the ZENITH-1 Trial. Journal of Allergy and Clinical Immunology, 2019, 143, AB36.	2.9	4
31	Two Cases Of Peculiar Hypereosinophilic Syndrome Treated With Mepolizumab. Journal of Allergy and Clinical Immunology, 2019, 143, AB292.	2.9	0
32	Long-Term Outcomes with Subcutaneous C1-Inhibitor Replacement Therapy for Prevention of Hereditary Angioedema Attacks. Journal of Allergy and Clinical Immunology: in Practice, 2019, 7, 1793-1802.e2.	3.8	58
33	Current and emerging biologics for the treatment of hereditary angioedema. Expert Opinion on Biological Therapy, 2019, 19, 517-526.	3.1	16
34	Treatment effect of switching from intravenous to subcutaneous C1-inhibitor for prevention of hereditary angioedema attacks: COMPACT subgroup findings. Journal of Allergy and Clinical Immunology: in Practice, 2019, 7, 2035-2038.	3.8	9
35	Efficacy of Lanadelumab in Hereditary Angioedema Patients Switching From C1 Inhibitor Long-Term Prophylaxis: Interim Results From the HELP Open-Label Extension Study. Journal of Allergy and Clinical Immunology, 2019, 143, AB37.	2.9	3
36	The International/Canadian Hereditary Angioedema Guideline. Allergy, Asthma and Clinical Immunology, 2019, 15, 72.	2.0	112

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37	The role of genetics in the current diagnostic workup of idiopathic nonâ€histaminergic angioedema. Allergy: European Journal of Allergy and Clinical Immunology, 2019, 74, 810-812.	5.7	12
38	Opioid Utilization and Perception of Pain Control in Hospitalized Patients: A Crossâ€Sectional Study of 11 Sites in 8 Countries. Journal of Hospital Medicine, 2019, 14, 737-745.	1.4	19
39	C1 esterase inhibitor concentrates and attenuated androgens â€“ Authors' reply. Lancet, The, 2018, 391, 1356.	13.7	1
40	Intermittent C1-Inhibitor Deficiency Associated with Recessive Inheritance: Functional and Structural Insight. Scientific Reports, 2018, 8, 977.	3.3	22
41	Exposureâ€Response Model of Subcutaneous C1â€Inhibitor Concentrate to Estimate the Risk of Attacks in Patients With Hereditary Angioedema. CPT: Pharmacometrics and Systems Pharmacology, 2018, 7, 158-165.	2.5	11
42	Mutation of the angiopoietin-1 gene (ANGPT1) associates with a new type of hereditary angioedema. Journal of Allergy and Clinical Immunology, 2018, 141, 1009-1017.	2.9	223
43	Effect of Lanadelumab Compared With Placebo on Prevention of Hereditary Angioedema Attacks. JAMA - Journal of the American Medical Association, 2018, 320, 2108.	7.4	174
44	The physician and hereditary angioedema friend or foe: 62-year diagnostic delay and iatrogenic procedures. Allergy, Asthma and Clinical Immunology, 2018, 14, 75.	2.0	4
45	Subcutaneous (SC) vs Intravenous (IV) C1-esterase-inhibitor (C1-INH) Replacement Treatment For The Prevention Of Attacks Of Hereditary Angioedema (HAE): A Population-based Exposure-response Analysis. Journal of Allergy and Clinical Immunology, 2018, 141, AB52.	2.9	0
46	Secreted Phospholipases A2 in Hereditary Angioedema With C1-Inhibitor Deficiency. Frontiers in Immunology, 2018, 9, 1721.	4.8	19
47	Costs and effects of on-demand treatment of hereditary angioedema in Italy: a prospective cohort study of 167 patients. BMJ Open, 2018, 8, e022291.	1.9	5
48	Emotional processes and stress in children affected by hereditary angioedema with C1-inhibitor deficiency: a multicenter, prospective study. Orphanet Journal of Rare Diseases, 2018, 13, 115.	2.7	24
49	Recurrent Retroperitoneal Angioedema. Journal of Allergy and Clinical Immunology: in Practice, 2018, 6, 1384-1385.	3.8	3
50	A transcriptomics study of hereditary angioedema attacks. Journal of Allergy and Clinical Immunology, 2018, 142, 883-891.	2.9	18
51	Splenic marginal zone lymphomas in acquired C1-inhibitor deficiency: clinical and molecular characterization. Medical Oncology, 2018, 35, 118.	2.5	18
52	Oral Plasma Kallikrein Inhibitor for Prophylaxis in Hereditary Angioedema. New England Journal of Medicine, 2018, 379, 352-362.	27.0	89
53	The Role of Failing Autonomic Nervous System on Life-Threatening Idiopathic Systemic Capillary Leak Syndrome. Frontiers in Medicine, 2018, 5, 111.	2.6	6
54	Angioedema Due to Bradykinin Dysregulation. Journal of Allergy and Clinical Immunology: in Practice, 2018, 6, 1132-1141.	3.8	94

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55	Population pharmacokinetics of subcutaneous C1â€inhibitor for prevention of attacks in patients with hereditary angioedema. Clinical and Experimental Allergy, 2018, 48, 1325-1332.	2.9	9
56	Recombinant human C1 esterase inhibitor (Conestat alfa) for prophylaxis to prevent attacks in adult and adolescent patients with hereditary angioedema. Expert Review of Clinical Immunology, 2018, 14, 707-718.	3.0	9
57	Effectiveness of icatibant for treatment of hereditary angioedema attacks is not affected by body weight: findings from the Icatibant Outcome Survey, a cohort observational study. Clinical and Translational Allergy, 2018, 8, 11.	3.2	3
58	Initiation of Prophylactic Treatment with Subcutaneous C1-Esterase Inhibitor (C1-INH [SC]) for Prevention of Hereditary Angioedema (HAE) Attacks and Onset of Effect: Findings from the Phase III COMPACT Study. Journal of Allergy and Clinical Immunology, 2018, 141, AB45.	2.9	1
59	Inhibiting Plasma Kallikrein for Hereditary Angioedema Prophylaxis. New England Journal of Medicine, 2017, 376, 717-728.	27.0	138
60	Reply. Journal of Allergy and Clinical Immunology, 2017, 139, 1720-1721.	2.9	4
61	Hereditary Angioedema with Normal C1 Inhibitor: An Italian Case Series. Journal of Allergy and Clinical Immunology, 2017, 139, AB231.	2.9	0
62	Risk for Attacks in Hereditary Angioedema (HAE) Population Correlates with C1-inhibitor Functional Activity (C1-INHact). Journal of Allergy and Clinical Immunology, 2017, 139, AB233.	2.9	1
63	The Icatibant Outcome Survey: experience of hereditary angioedema management from six European countries. Journal of the European Academy of Dermatology and Venereology, 2017, 31, 1214-1222.	2.4	21
64	Diagnosis, Course, and Management of Angioedema in Patients With Acquired C1-Inhibitor Deficiency. Journal of Allergy and Clinical Immunology: in Practice, 2017, 5, 1307-1313.	3.8	62
65	Prevention of Hereditary Angioedema Attacks with a Subcutaneous C1 Inhibitor. New England Journal of Medicine, 2017, 376, 1131-1140.	27.0	169
66	Catabolism of C1 inhibitor influences the response to replacement therapy in hereditary angioedema. Journal of Allergy and Clinical Immunology, 2017, 139, 2005-2007.e1.	2.9	2
67	Longâ€term safety of icatibant treatment of patients with angioedema in realâ€world clinical practice. Allergy: European Journal of Allergy and Clinical Immunology, 2017, 72, 994-998.	5.7	16
68	Complement and contact system activation in acute congestive heart failure patients. Clinical and Experimental Immunology, 2017, 190, 251-257.	2.6	7
69	Recombinant human C1 esterase inhibitor for prophylaxis of hereditary angio-oedema: a phase 2, multicentre, randomised, double-blind, placebo-controlled crossover trial. Lancet, The, 2017, 390, 1595-1602.	13.7	55
70	Cleaved kininogen as a biomarker for bradykinin release in hereditary angioedema. Journal of Allergy and Clinical Immunology, 2017, 140, 1700-1703.e8.	2.9	34
71	International consensus on the diagnosis and management of pediatric patients with hereditary angioedema with C1 inhibitor deficiency. Allergy: European Journal of Allergy and Clinical Immunology, 2017, 72, 300-313.	5.7	153
72	VP55 Health Technology Assessment Of Orphan Drugs: The Case Of Hereditary Angioedema In Italy. International Journal of Technology Assessment in Health Care, 2017, 33, 173-174.	0.5	0

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73	Recombinant human C1 esterase inhibitor for acute hereditary angioedema attacks with upper airway involvement. Allergy and Asthma Proceedings, 2017, 38, 462-466.	2.2	13
74	An open-label study to evaluate the long-term safety and efficacy of lanadelumab for prevention of attacks in hereditary angioedema: design of the HELP study extension. Clinical and Translational Allergy, 2017, 7, 36.	3.2	28
75	Hereditary angioedema: Assessing the hypothesis for underlying autonomic dysfunction. PLoS ONE, 2017, 12, e0187110.	2.5	10
76	Novelties in the Diagnosis and Treatment of Angioedema. Journal of Investigational Allergology and Clinical Immunology, 2016, 26, 212-221.	1.3	59
77	Elevated plasma levels of vascular permeability factors in C1 inhibitorâ€deficient hereditary angioedema. Allergy: European Journal of Allergy and Clinical Immunology, 2016, 71, 989-996.	5.7	46
78	Plasmin is a natural trigger for bradykinin production in patients with hereditary angioedema with factor XII mutations. Journal of Allergy and Clinical Immunology, 2016, 138, 1414-1423.e9.	2.9	146
79	Angioedema Phenotypes: Disease Expression and Classification. Clinical Reviews in Allergy and Immunology, 2016, 51, 162-169.	6.5	60
80	High rate of hepatitis B viral breakthrough in elderly non-Hodgkin lymphomas patients treated with Rituximab based chemotherapy. Digestive and Liver Disease, 2016, 48, 1394-1397.	0.9	19
81	Misdiagnosis trends in patients with hereditary angioedema from the real-world clinical setting. Annals of Allergy, Asthma and Immunology, 2016, 117, 394-398.	1.0	78
82	Refined Method for Collection of Plasma Samples to Evaluate the Role of Plasma Kallikrein in Various Disease States. Journal of Allergy and Clinical Immunology, 2016, 137, AB248.	2.9	1
83	Modeling and Analyses to Identify Potential Dosing Regimens of DX-2930 for the Long-Term Prophylaxis of Hereditary Angioedema. Journal of Allergy and Clinical Immunology, 2016, 137, AB252.	2.9	2
84	High prevalence of splenic marginal zone lymphoma among patients with acquired C1 inhibitor deficiency. British Journal of Haematology, 2016, 172, 902-908.	2.5	41
85	Current treatment options for hereditary angioedema due to C1 inhibitor deficiency. Expert Opinion on Pharmacotherapy, 2016, 17, 27-40.	1.8	20
86	Phase II study results of a replacement therapy for hereditary angioedema with subcutaneous C1â€inhibitor concentrate. Allergy: European Journal of Allergy and Clinical Immunology, 2015, 70, 1319-1328.	5.7	59
87	Differential diagnosis and management issues of idiopathic angiooedema and their resolution. The Journal of Critical Care Medicine, 2015, 1, 55-60.	0.7	0
88	Efficacy of onâ€demand treatment in reducing morbidity in patients with hereditary angioedema due to <scp>C</scp>1 inhibitor deficiency. Allergy: European Journal of Allergy and Clinical Immunology, 2015, 70, 1553-1558.	5.7	23
89	Use of subcutaneousâ€C1 INH for acute therapy and prophylaxis of a child with HAE. Pediatric Allergy and Immunology, 2015, 26, 296-297.	2.6	5
90	Treatment of hereditary angioedema with recombinant human C1 Inhibitor in a real-life setting: the experience of the HAE Centre in Milan. World Allergy Organization Journal, 2015, 8, A173.	3.5	1

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91	A nationwide survey of hereditary angioedema due to C1 inhibitor deficiency in Italy. Orphanet Journal of Rare Diseases, 2015, 10, 11.	2.7	102
92	High Molecular Weight Kininogen Cleavage in Idiopathic Angioedema. Journal of Allergy and Clinical Immunology, 2015, 135, AB198.	2.9	1
93	Clinical and Pathological Findings of a Fatal Systemic Capillary Leak Syndrome (Clarkson Disease). Medicine (United States), 2015, 94, e591.	1.0	14
94	The safety of treatments for angioedema with hereditary C1 inhibitor deficiency. Expert Opinion on Drug Safety, 2015, 14, 1725-1736.	2.4	8
95	Recombinant replacement therapy for hereditary angioedema due to C1 inhibitor deficiency. Immunotherapy, 2015, 7, 739-752.	2.0	24
96	Presentation, diagnosis and treatment of angioedema without wheals: a retrospective analysis of a cohort of 1058 patients. Journal of Internal Medicine, 2015, 277, 585-593.	6.0	86
97	Canadian hereditary angioedema guideline. Allergy, Asthma and Clinical Immunology, 2014, 10, 50.	2.0	68
98	Non-invasive ventilation in the treatment of sleep-related breathing disorders: A review and update. Revista Portuguesa De Pneumologia, 2014, 20, 324-335.	0.7	22
99	Multimorbidity and polypharmacy in the elderly: lessons from REPOSI. Internal and Emergency Medicine, 2014, 9, 723-734.	2.0	121
100	Pathophysiology of Hereditary Angioedema. Pediatric, Allergy, Immunology, and Pulmonology, 2014, 27, 159-163.	0.8	59
101	Highâ€molecularâ€weight kininogen cleavage correlates with disease states in the bradykininâ€mediated angioedema due to hereditary <scp>C</scp>1â€inhibitor deficiency. Clinical and Experimental Allergy, 2014, 44, 1503-1514.	2.9	78
102	Guidance for diagnosis and treatment of acute angioedema in the emergency department: consensus statement by a panel of Italian experts. Internal and Emergency Medicine, 2014, 9, 85-92.	2.0	38
103	Classification, diagnosis, and approach to treatment for angioedema: consensus report from the <scp>H</scp>ereditary <scp>A</scp>ngioedema <scp>I</scp>nternational <scp>W</scp>orking <scp>G</scp>roup. Allergy: European Journal of Allergy and Clinical Immunology, 2014, 69, 602-616.	5.7	538
104	C1 Inhibitor Autoantibodies. , 2014, , 699-705.		0
105	Gout, allopurinol intake and clinical outcomes in the hospitalized multimorbid elderly. European Journal of Internal Medicine, 2014, 25, 847-852.	2.2	1
106	Pharmacokinetics of plasmaâ€derived <scp>C</scp>1â€esterase inhibitor after subcutaneous versus intravenous administration in subjects with mild or moderate hereditary angioedema: the <scp>PASSION</scp> study. Transfusion, 2014, 54, 1552-1561.	1.6	40
107	A young man with cough, fever and epigastric pain. Internal and Emergency Medicine, 2014, 9, 569-573.	2.0	0
108	Recombinant human C1-esterase inhibitor relieves symptoms of hereditary angioedema attacks: phase 3, randomized, placebo-controlled trial. Annals of Allergy, Asthma and Immunology, 2014, 112, 163-169.e1.	1.0	70

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109	Lung ultrasonography for the assessment of rapid extravascular water variation: evidence from hemodialysis patients. Internal and Emergency Medicine, 2013, 8, 409-415.	2.0	97
110	Acquired C1-inhibitor deficiency and lymphoproliferative disorders: A tight relationship. Critical Reviews in Oncology/Hematology, 2013, 87, 323-332.	4.4	59
111	Profile of infective endocarditis observed from 2003 - 2010 in a single center in Italy. BMC Infectious Diseases, 2013, 13, 545.	2.9	24
112	Diagnosing Angioedema. Immunology and Allergy Clinics of North America, 2013, 33, 449-456.	1.9	15
113	The management of paediatric allergy. Current Opinion in Allergy and Clinical Immunology, 2013, 13, S1-S50.	2.3	2
114	Review of Recent Guidelines and Consensus Statements on Hereditary Angioedema Therapy with Focus on Self-Administration. International Archives of Allergy and Immunology, 2013, 161, 3-9.	2.1	32
115	Therapeutic management of hereditary angioedema due to C1 inhibitor deficiency. Expert Review of Clinical Immunology, 2013, 9, 477-488.	3.0	10
116	Ecallantide for treatment of acute attacks of acquired C1 esterase inhibitor deficiency. Allergy and Asthma Proceedings, 2013, 34, 72-77.	2.2	25
117	Ongoing Contact Activation in Patients with Hereditary Angioedema. PLoS ONE, 2013, 8, e74043.	2.5	25
118	A Biomarker Assay For The Detection Of Contact System Activation. Blood, 2013, 122, 2347-2347.	1.4	0
119	Long-term prophylaxis in hereditary angio-oedema: a systematic review. BMJ Open, 2012, 2, e000524.	1.9	17
120	Hereditary and Acquired Complement Component 1 Esterase Inhibitor Deficiency: A Review for the Hematologist. Acta Haematologica, 2012, 127, 208-220.	1.4	32
121	Hereditary angioedema with normal C1 inhibitor function: Consensus of an international expert panel. Allergy and Asthma Proceedings, 2012, 33, 145-156.	2.2	142
122	Hereditary angio-oedema. Lancet, The, 2012, 379, 474-481.	13.7	294
123	Content Validity of Visual Analog Scales to Assess Symptom Severity of Acute Angioedema Attacks in Adults with Hereditary Angioedema. Patient, 2012, 5, 113-126.	2.7	18
124	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, 308-320.	2.9	207
125	Clinical Impact of Peripheral Attacks in Hereditary Angioedema Patients. American Journal of Medicine, 2012, 125, 937.e17-937.e24.	1.5	22
126	Target levels of functional C1â€inhibitor in hereditary angioedema. Allergy: European Journal of Allergy and Clinical Immunology, 2012, 67, 123-130.	5.7	51

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127	Evidenceâ€based recommendations for the therapeutic management of angioedema owing to hereditary C1 inhibitor deficiency: consensus report of an International Working Group. Allergy: European Journal of Allergy and Clinical Immunology, 2012, 67, 147-157.	5.7	294
128	Alterations of coagulation and fibrinolysis in patients with angioedema due to C1-inhibitor deficiency. Clinical and Experimental Immunology, 2012, 167, 472-478.	2.6	60
129	Efficacy and safety of recombinant human C1â€inhibitor for the treatment of attacks of hereditary angioedema: European openâ€label extension study. Clinical and Experimental Allergy, 2012, 42, 929-935.	2.9	50
130	Icatibant treatment for acquired C1â€inhibitor deficiency: a realâ€world observational study. Allergy: European Journal of Allergy and Clinical Immunology, 2012, 67, 1074-1077.	5.7	43
131	Long-term follow-up of 111 patients with angiotensin-converting enzyme inhibitor-related angioedema. Journal of Hypertension, 2011, 29, 2273-2277.	0.5	98
132	Standard care impact on angioedema because of hereditary C1 inhibitor deficiency: a 21-month prospective study in a cohort of 103 patients. Allergy: European Journal of Allergy and Clinical Immunology, 2011, 66, 192-196.	5.7	73
133	New topics in bradykinin research. Allergy: European Journal of Allergy and Clinical Immunology, 2011, 66, 1397-1406.	5.7	146
134	Treatment of acquired angioedema with icatibant: a case report. Internal and Emergency Medicine, 2011, 6, 279-280.	2.0	23
135	Determinants of Lung Function, COPD, and Asthma. New England Journal of Medicine, 2011, 364, 86-87.	27.0	3
136	18 Kallikrein-kinin system in angioedema. , 2011, , 289-306.		0
137	Lymphoproliferative Disorder and Acquired C1-INH Deficiency. A Case Series of 48 Patients. Blood, 2011, 118, 1596-1596.	1.4	0
138	The Acquired Deficiency of C1-Inhibitor: Lymphoproliferation and Angioedema. Current Molecular Medicine, 2010, 10, 354-360.	1.3	38
139	A simple endocrinological cause of a complex cardiovascular picture. Internal and Emergency Medicine, 2010, 5, 269-271.	2.0	0
140	Activation of blood coagulation in chronic urticaria: pathophysiological and clinical implications by Dr. Massimo Cugno et al.. Internal and Emergency Medicine, 2010, 5, 95-96.	2.0	1
141	Angioedema due to C1 inhibitor deficiency in 2010. Internal and Emergency Medicine, 2010, 5, 481-486.	2.0	31
142	Acquired angioedema. Allergy, Asthma and Clinical Immunology, 2010, 6, 14.	2.0	151
143	HAE international home therapy consensus document. Allergy, Asthma and Clinical Immunology, 2010, 6, 22.	2.0	149
144	2010 International consensus algorithm for the diagnosis, therapy and management of hereditary angioedema. Allergy, Asthma and Clinical Immunology, 2010, 6, 24.	2.0	443

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145	Ecallantide for the Treatment of Acute Attacks in Hereditary Angioedema. New England Journal of Medicine, 2010, 363, 523-531.	27.0	266
146	Icatibant, a New Bradykinin-Receptor Antagonist, in Hereditary Angioedema. New England Journal of Medicine, 2010, 363, 532-541.	27.0	477
147	Recombinant human C1-inhibitor for the treatment of acute angioedema attacks in patients with hereditary angioedema. Journal of Allergy and Clinical Immunology, 2010, 126, 821-827.e14.	2.9	203
148	Replacement therapy with C1 esterase inhibitors for hereditary angioedema. Drugs of Today, 2010, 46, 867.	1.1	13
149	Plasma biomarkers of acute attacks in patients with angioedema due to C1â€inhibitor deficiency. Allergy: European Journal of Allergy and Clinical Immunology, 2009, 64, 254-257.	5.7	83
150	C1-inhibitor deficiency and angioedema: molecular mechanisms and clinical progress. Trends in Molecular Medicine, 2009, 15, 69-78.	6.7	207
151	Novel pathogenic mechanism and therapeutic approaches to angioedema associated with C1 inhibitor deficiency. Journal of Allergy and Clinical Immunology, 2009, 124, 1303-1310.e4.	2.9	94
152	Disease expression in women with hereditary angioedema. American Journal of Obstetrics and Gynecology, 2008, 199, 484.e1-484.e4.	1.3	134
153	Angioedema due to acquired C1-inhibitor deficiency: A bridging condition between autoimmunity and lymphoproliferation. Autoimmunity Reviews, 2008, 8, 156-159.	5.8	96
154	Functional C1-Inhibitor diagnostics in hereditary angioedema: Assay evaluation and recommendations. Journal of Immunological Methods, 2008, 338, 14-20.	1.4	84
155	Mutation screening of C1 inhibitor gene in 108 unrelated families with hereditary angioedema: Functional and structural correlates. Molecular Immunology, 2008, 45, 3536-3544.	2.2	116
156	A randomised, placebo-controlled, double blind phase III study of the efficacy and safety of recombinant human C1 inhibitor for the treatment of acute attacks in patients with hereditary angioedema. Molecular Immunology, 2008, 45, 4118-4119.	2.2	1
157	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. Annals of Allergy, Asthma and Immunology, 2008, 100, S30-S40.	1.0	181
158	Successful resolution of bowel obstruction in a patient with hereditary angioedema. European Journal of Gastroenterology and Hepatology, 2008, 20, 583-587.	1.6	12
159	Lymphoproliferative disease and acquired C1 inhibitor deficiency. Haematologica, 2007, 92, 716-718.	3.5	73
160	C1 INHIBITOR AUTOANTIBODIES. , 2007, , 695-701.		0
161	The deficiency of C1 inhibitor and its treatment. Immunobiology, 2007, 212, 325-331.	1.9	18
162	Established and new treatments for hereditary angioedema: An update. Molecular Immunology, 2007, 44, 3858-3861.	2.2	17

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163	The use of plasma-derived C1 inhibitor in the treatment of hereditary angioedema. Expert Opinion on Pharmacotherapy, 2007, 8, 3173-3181.	1.8	38
164	Natural History and Clinical Impact of Cryoglobulins in Chronic Hepatitis C: 10-Year Prospective Study of 343 Patients. Gastroenterology, 2007, 133, 835-842.	1.3	34
165	Helicobacter pylori Infection as a Triggering Factor of Attacks in Patients with Hereditary Angioedema. Helicobacter, 2007, 12, 251-257.	3.5	53
166	Anaphylaxis in Response to C1 Esterase Inhibitor in a Patient with Hereditary Angioedema. Allergy and Clinical Immunology International, 2007, 19, 159-163.	0.3	3
167	Acquired Deficiency of the Inhibitor of the First Complement Component: Presentation, Diagnosis, Course, and Conventional Management. Immunology and Allergy Clinics of North America, 2006, 26, 669-690.	1.9	101
168	Angioedema without urticaria: a large clinical survey. Cmaj, 2006, 175, 1065-1070.	2.0	170
169	Angiotensin-converting enzyme inhibitor-related angioedema: how to deal with it. Expert Opinion on Drug Safety, 2006, 5, 643-649.	2.4	56
170	Nickel and Sulfites Food Allergy in Patients With Angioedema Associated With ACE Inhibitor Use. Archives of Internal Medicine, 2005, 165, 814.	3.8	3
171	C1 inhibitor: molecular and clinical aspects. Seminars in Immunopathology, 2005, 27, 286-298.	4.0	92
172	Laboratory diagnostics for hereditary angioedema: An economic, evidence-based standpoint. Journal of Allergy and Clinical Immunology, 2005, 115, 878-879.	2.9	3
173	Angioedema Associated With Angiotensin-Converting Enzyme Inhibitor Use. Archives of Internal Medicine, 2004, 164, 910.	3.8	184
174	C1 inhibitor gene expression in patients with hereditary angioedema: Quantitative evaluation by means of real-time RT-PCR. Journal of Allergy and Clinical Immunology, 2004, 114, 638-644.	2.9	26
175	Canadian 2003 International Consensus Algorithm for the Diagnosis, Therapy, and Management of Hereditary Angioedema. Journal of Allergy and Clinical Immunology, 2004, 114, 629-637.	2.9	177
176	Hereditary and acquired angioedema: Problems and progress: Proceedings of the third C1 esterase inhibitor deficiency workshop and beyond. Journal of Allergy and Clinical Immunology, 2004, 114, S51-S131.	2.9	582
177	Swelling isn't swell. Clinical Immunology, 2004, 113, 231-233.	3.2	0
178	Increased expression of C1-inhibitor mRNA in patients with hereditary angioedema treated with Danazol. Immunology Letters, 2003, 86, 271-276.	2.5	66
179	Recent advances in the use of C1 inhibitor as a therapeutic agent. Molecular Immunology, 2003, 40, 155-158.	2.2	15
180	Research on complement: old issues revisited and a novel sphere of influence. Trends in Immunology, 2003, 24, 292-295.	6.8	12

#	Article	IF	Citations
181	How do we treat patients with hereditary angioedema. Transfusion and Apheresis Science, 2003, 29, 221-227.	1.0	51
182	Bradykinin and the pathophysiology of angioedema. International Immunopharmacology, 2003, 3, 311-317.	3.8	197
183	Title is missing!. Medicine (United States), 2003, 82, 274-281.	1.0	9
184	Autoantibodies and Lymphoproliferative Diseases in Acquired C1-Inhibitor Deficiencies. Medicine (United States), 2003, 82, 274-281.	1.0	121
185	Bradykinin-Mediated Angioedema. New England Journal of Medicine, 2002, 347, 621-622.	27.0	213
186	Mechanisms of C1-Inhibitor Deficiency. Immunobiology, 2002, 205, 542-551.	1.9	46
187	Acquired C1 esterase inhibitor deficiency in two patients presenting with a lupus-like syndrome and anticardiolipin antibodies. Arthritis and Rheumatism, 2002, 47, 223-226.	6.7	10
188	Drug-Induced Angioedema without Urticaria. Drug Safety, 2001, 24, 599-605.	3.2	138
189	C1-inhibitor deficiency and angioedema. Molecular Immunology, 2001, 38, 161-173.	2.2	95
190	Rapid detection by fluorescent multiplex PCR of exon deletions and duplications in theC1 inhibitorgene of hereditary angioedema patients. Human Mutation, 2001, 17, 61-70.	2.5	45
191	Dental experience and self-perceived dental care needs of patients with angioedema. Special Care in Dentistry, 2001, 21, 27-31.	0.8	6
192	Prevalence and risk factors for the presence of serum cryoglobulins in patients with chronic hepatitis C. Journal of Viral Hepatitis, 2000, 7, 138-143.	2.0	71
193	Does acquired angioedema increase the risk of surgery with cardiopulmonary bypass?. Journal of Thoracic and Cardiovascular Surgery, 2000, 120, 609-610.	0.8	1
194	Frequent de novo mutations and exon deletions in the C1inhibitor gene of patients with angioedema. Journal of Allergy and Clinical Immunology, 2000, 106, 1147-1154.	2.9	208
195	Acquired C1 Esterase Inhibitor Deficiency. Annals of Internal Medicine, 2000, 133, 837.	3.9	2
196	C1 Inhibitor Function and Anti-C1 Inhibitor Autoantibodies in Patients with HIV Type 1 Infection. AIDS Research and Human Retroviruses, 1999, 15, 95-96.	1.1	3
197	Consumption of C4b-binding protein (C4BP) duringin vivoactivation of the classical complement pathway. Clinical and Experimental Immunology, 1999, 116, 220-224.	2.6	7
198	Angioedema due to angiotensin-converting enzyme inhibitors. Immunopharmacology, 1999, 44, 21-25.	2.0	137

#	Article	IF	Citations
199	Local bradykinin generation in hereditary angioedemaâ˜†â˜†â˜†â˜…. Journal of Allergy and Clinical Immunology, 1999, 104, 1321-1322.	2.9	168
200	Idiopathic nonhistaminergic angioedema. American Journal of Medicine, 1999, 106, 650-654.	1.5	98
201	Pathogenetic and Clinical Aspects of C1 Inhibitor Deficiency. Immunobiology, 1998, 199, 366-376.	1.9	85
202	Plasma bradykinin in angio-oedema. Lancet, The, 1998, 351, 1693-1697.	13.7	681
203	Idiopathic capillary leak syndrome: Evidence of CD8-positive lymphocytes surrounding damaged endothelial cellsâ˜†, â˜†â˜†, â˜…. Journal of Allergy and Clinical Immunology, 1997, 99, 417-419.	2.9	39
204	Side effects of long-term prophylaxis with attenuated androgens in hereditary angioedema: Comparison of treated and untreated patientsâ˜†â˜†â˜†â˜…. Journal of Allergy and Clinical Immunology, 1997, 99, 194-196.	2.9	148
205	Activation of the Coagulation Cascade in C1-Inhibitor Deficiencies. Blood, 1997, 89, 3213-3218.	1.4	149
206	Effects of Radiographic Contrast Media. Investigative Radiology, 1997, 32, 648.	6.2	1
207	Activation of factor XII and cleavage of high molecular weight kininogen during acute attacks in hereditary and acquired Cl-inhibitor deficiencies. Immunopharmacology, 1996, 33, 361-364.	2.0	62
208	Relevance of lymphoproliferative disorders and of anti-C1 inhibitor autoantibodies in acquired angio-oedema. Clinical and Experimental Immunology, 1996, 106, 475-480.	2.6	115
209	Hereditary Angioedema. New England Journal of Medicine, 1996, 334, 1666-1667.	27.0	204
210	Non Neutralizing Antibodies to Tissue Type Plasminogen Activator in the Serum of Acute Myocardial Infarction Patients Treated with the Recombinant Protein. Thrombosis and Haemostasis, 1996, 76, 234-238.	3.4	17
211	C1 Inhibitor Autoantibodies. , 1996, , 126-131.		2
212	Reduction in transmission of hepatitis C after the introduction of a heat-treatment step in the production of C1-inhibitor concentrate. Transfusion, 1995, 35, 209-212.	1.6	45
213	Infusion of C1-inhibitor plasma concentrate prevents hyperamylasemia induced by endoscopic sphincterotomy. Gastrointestinal Endoscopy, 1995, 42, 301-305.	1.0	31
214	Activation of complement and kinin systems after thrombolytic therapy in patients with acute myocardial infarction. A comparison between streptokinase and recombinant tissue-type plasminogen activator.. Circulation, 1994, 90, 2666-2670.	1.6	69
215	Activation of the contact system and fibrinolysis in autoimmune acquired angioedema: A rationale for prophylactic use of tranexamic acid. Journal of Allergy and Clinical Immunology, 1994, 93, 870-876.	2.9	90
216	A cluster of mutations within a short triplet repeat in the C1 inhibitor gene.. Proceedings of the National Academy of Sciences of the United States of America, 1994, 91, 9622-9625.	7.1	26

#	Article	IF	Citations
217	Autoimmune C1 inhibitor deficiency: Report of eight patients. American Journal of Medicine, 1993, 95, 169-175.	1.5	94
218	The Complement System. International Journal of Immunopathology and Pharmacology, 1992, 5, 123-130.	2.1	8
219	Hereditary and Acquired C1-Inhibitor Deficiency. Medicine (United States), 1992, 71, 206-215.	1.0	534
220	Replacement therapy in hereditary and acquired angioedema. Pharmacological Research, 1992, 26, 148-149.	7.1	5
221	A dysfunctional Cl inhibitor protein with a new reactive center mutation (Arg-444â†'Leu). FEBS Letters, 1992, 301, 34-36.	2.8	29
222	Complement Deficiency and Antibody Profile in Survivors of Meningococcal Meningitis due to common Serogroups in Italy. Scandinavian Journal of Immunology, 1992, 35, 589-596.	2.7	31
223	Peripheral edema due to increased vascular permeability: a clinical appraisal. International Journal of Clinical and Laboratory Research, 1992, 21, 241-246.	1.0	12
224	Contraindications to the use of ace inhibitors in patients with c1 esterase inhibitor deficiency. American Journal of Medicine, 1991, 90, 278.	1.5	34
225	Contraindications to the use of ace inhibitors in patients with c1 esterase inhibitor deficiency. American Journal of Medicine, 1991, 90, 278.	1.5	17
226	Long-term treatment of hereditary angioedema with attenuated androgens: A survey of a 13-year experience. Journal of Allergy and Clinical Immunology, 1991, 87, 768-773.	2.9	119
227	Nonsense mutations affect C1 inhibitor messenger RNA levels in patients with type I hereditary angioneurotic edema.. Journal of Clinical Investigation, 1991, 88, 755-759.	8.2	33
228	Type II hereditary angioneurotic edema that may result from a single nucleotide change in the codon for alanine-436 in the C1 inhibitor gene.. Proceedings of the National Academy of Sciences of the United States of America, 1990, 87, 265-268.	7.1	62
229	Increased levels of soluble interleukin-2 receptors in serum of patients with lung cancer. British Journal of Cancer, 1990, 61, 434-435.	6.4	34
230	The Systemic Capillary Leak Syndrome: Appearance of Interleukin-2-Receptor-Positive Cells during Attacks. Annals of Internal Medicine, 1990, 113, 475.	3.9	78
231	Identification of a new P1 residue mutation (444Argâ†'Ser) in a dysfunctional C1 inhibitor protein contained in a type II hereditary angioedema plasma. FEBS Letters, 1990, 266, 13-16.	2.8	25
232	Plasma levels of C1- inhibitor complexes and cleaved C1- inhibitor in patients with hereditary angioneurotic edema.. Journal of Clinical Investigation, 1990, 85, 1215-1220.	8.2	60
233	Type I C1 inhibitor deficiency with a small messenger RNA resulting from deletion of one exon.. Journal of Clinical Investigation, 1989, 83, 1888-1893.	8.2	43
234	Molecular basis for the deficiency of complement 1 inhibitor in type I hereditary angioneurotic edema.. Journal of Clinical Investigation, 1987, 79, 698-702.	8.2	61

#	Article	IF	Citations
235	Restriction fragment length polymorphism of the C1 inhibitor gene in hereditary angioneurotic edema.. Journal of Clinical Investigation, 1987, 80, 1640-1643.	8.2	35
236	In vivo study of the complement system during infusion of radiographic contrast media. Journal of Allergy and Clinical Immunology, 1986, 77, 690-692.	2.9	12
237	The metabolism of C1 inhibitor and C1q in patients with acquired C1-inhibitor deficiency. Journal of Allergy and Clinical Immunology, 1986, 77, 322-326.	2.9	59
238	Human inhibitor of the first component of complement, C1: characterization of cDNA clones and localization of the gene to chromosome 11.. Proceedings of the National Academy of Sciences of the United States of America, 1986, 83, 3161-3165.	7.1	104
239	Acquired C1 Inhibitor Deficiency with Angioedema Symptoms in a Patient Infected with Echinococcus granulosus. Complement (Basel, Switzerland), 1985, 2, 133-139.	0.9	32
240	Acquired C1-Inhibitor Deficiency Associated with Antiidiotypic Antibody to Monoclonal Immunoglobulins. New England Journal of Medicine, 1985, 312, 534-540.	27.0	166
241	C1 INH Concentrate in the Therapy of Hereditary Angioedema. Allergy: European Journal of Allergy and Clinical Immunology, 1983, 38, 81-84.	5.7	48
242	Morphologic evaluation of the liver in hereditary angioedema patients on long-term treatment with androgen derivatives. Journal of Allergy and Clinical Immunology, 1983, 72, 294-298.	2.9	46
243	Behavior in vivo of normal and dysfunctional C1 inhibitor in normal subjects and patients with hereditary angioneurotic edema.. Journal of Clinical Investigation, 1983, 71, 1041-1046.	8.2	119
244	Effect of treatment with 17 alpha-alkylated androgens on C4 conversion products in hereditary angioedema studied by crossed immunoelectrophoresis.. Journal of Clinical Pathology, 1982, 35, 728-731.	2.0	0
245	Hereditary Angioedema: An Appraisal of 104 Cases. American Journal of the Medical Sciences, 1982, 284, 2-9.	1.1	156
246	Danazol and stanozolol in long-term prophylactic treatment of hereditary angioedema. Journal of Allergy and Clinical Immunology, 1980, 65, 75-79.	2.9	89
247	Hepatic Function and Fibrinolysis in Patients with Hereditary Angioedema Undergoing Long-Term Treatment with Tranexamic Acid. Allergy: European Journal of Allergy and Clinical Immunology, 1978, 33, 216-221.	5.7	25
248	Treatment of hereditary angioedema. Klinische Wochenschrift, 1978, 56, 819-823.	0.6	37
249	INTERMITTENT THERAPY WITH DANAZOL IN HEREDITARY ANGIÅ'DEMA. Lancet, The, 1978, 311, 453.	13.7	11
250	Hereditary angioneurotic oedema. Klinische Wochenschrift, 1975, 53, 679-684.	0.6	5