Nora Veszeli

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/6055116/publications.pdf

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33	537	623574	677027
papers	citations	14	22
		h-index	g-index

35	35	35	660
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Pathways of Neutrophil Granulocyte Activation in Hereditary Angioedema with C1 Inhibitor Deficiency. Clinical Reviews in Allergy and Immunology, 2021, 60, 383-395.	2.9	6
2	A Novel Homozygous In-Frame Deletion in Complement Factor 3 Underlies Early-Onset Autosomal Recessive Atypical Hemolytic Uremic Syndrome - Case Report. Frontiers in Immunology, 2021, 12, 608604.	2.2	1
3	FHR-5 Serum Levels and CFHR5 Genetic Variations in Patients With Immune Complex-Mediated Membranoproliferative Glomerulonephritis and C3-Glomerulopathy. Frontiers in Immunology, 2021, 12, 720183.	2.2	12
4	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.	2.0	43
5	Hereditary angioedema attack: what happens to vasoactive mediators?. International Immunopharmacology, 2020, 78, 106079.	1.7	7
6	Validation of Early Increase in Complement Activation Marker sC5b-9 as a Predictive Biomarker for the Development of Thrombotic Microangiopathy After Stem Cell Transplantation. Frontiers in Medicine, 2020, 7, 569291.	1.2	14
7	Patterns of C1-Inhibitor/Plasma Serine Protease Complexes in Healthy Humans and in Hereditary Angioedema Patients. Frontiers in Immunology, 2020, 11, 794.	2.2	29
8	Clinical Characteristics and Safety of Plasma-Derived C1-Inhibitor Therapy in Children and Adolescents with Hereditary Angioedemaâ€"A Long-Term Survey. Journal of Allergy and Clinical Immunology: in Practice, 2020, 8, 2379-2383.	2.0	6
9	Changes of coagulation parameters during erythema marginatum in patients with hereditary angioedema. International Immunopharmacology, 2020, 81, 106293.	1.7	7
10	Evaluation of the efficacy and safety of home treatment with the recombinant human C1-inhibitor in hereditary angioedema resulting from C1-inhibitor deficiency. International Immunopharmacology, 2020, 80, 106216.	1.7	3
11	Serum fetuin-A, tumor necrosis factor alpha and C-reactive protein concentrations in patients with hereditary angioedema with C1-inhibitor deficiency. Orphanet Journal of Rare Diseases, 2019, 14, 67.	1.2	3
12	Complete kinetic followâ€up of symptoms and complement parameters during a hereditary angioedema attack. Allergy: European Journal of Allergy and Clinical Immunology, 2018, 73, 516-520.	2.7	6
13	Secreted Phospholipases A2 in Hereditary Angioedema With C1-Inhibitor Deficiency. Frontiers in Immunology, 2018, 9, 1721.	2.2	19
14	Idiopathic Nonhistaminergic Acquired Angioedema Versus Hereditary Angioedema. Journal of Allergy and Clinical Immunology: in Practice, 2018, 6, 1205-1208.	2.0	10
15	Healthâ€related quality of life among children with hereditary angioedema. Pediatric Allergy and Immunology, 2017, 28, 370-376.	1.1	28
16	A novel prophylaxis with C1-inhibitor concentrate in hereditary angioedema during erythema marginatum. Immunology Letters, 2017, 189, 90-93.	1.1	10
17	The role of the complement system in hereditary angioedema. Molecular Immunology, 2017, 89, 59-68.	1.0	35
18	The relationship between anxiety and quality of life in children with hereditary angioedema. Pediatric Allergy and Immunology, 2017, 28, 692-698.	1.1	20

#	Article	IF	Citations
19	Cleaved kininogen as a biomarker for bradykinin release in hereditary angioedema. Journal of Allergy and Clinical Immunology, 2017, 140, 1700-1703.e8.	1.5	34
20	Glucocorticoid receptor gene polymorphisms in hereditary angioedema with C1-inhibitor deficiency. Orphanet Journal of Rare Diseases, 2017, 12, 5.	1.2	5
21	Risk of thromboembolism in patients with hereditary angioedema treated with plasma-derived C1-inhibitor. Allergy and Asthma Proceedings, 2016, 37, 164-170.	1.0	25
22	Thyroid hormones and complement parameters in hereditary angioedema with C1-inhibitor deficiency. Annals of Allergy, Asthma and Immunology, 2016, 117, 175-179.	0.5	2
23	First report of icatibant treatment in a pregnant patient with hereditary angioedema. Journal of Obstetrics and Gynaecology Research, 2016, 42, 1026-1028.	0.6	23
24	â€œNuts and Boltsâ€•of Laboratory Evaluation of Angioedema. Clinical Reviews in Allergy and Immunology, 2016, 51, 140-151.	2.9	43
25	Bacteriuria increases the risk of edematous attacks in hereditary angioedema with C1-inhibitor deficiency. Allergy: European Journal of Allergy and Clinical Immunology, 2016, 71, 1791-1793.	2.7	8
26	The effect of long-term danazol treatment on haematological parameters in hereditary angioedema. Orphanet Journal of Rare Diseases, 2016, 11, 18.	1.2	9
27	Comprehensive study into the activation of the plasma enzyme systems during attacks of hereditary angioedema due to C1-inhibitor deficiency. Orphanet Journal of Rare Diseases, 2015, 10, 132.	1.2	39
28	Neutrophil activation during attacks in patients with hereditary angioedema due to C1-inhibitor deficiency. Orphanet Journal of Rare Diseases, 2015, 10, 156.	1.2	24
29	Management of pregnancies in a hereditary angioedema patient after treatment with attenuated androgens since childhood. Journal of Obstetrics and Gynaecology, 2015, 35, 89-90.	0.4	6
30	Successful prophylaxis with recombinant human C1 inhibitor in a patient with hereditary angioedema. Annals of Allergy, Asthma and Immunology, 2015, 114, 64-65.	0.5	5
31	Frequency of the virilising effects of attenuated androgens reported by women with hereditary angioedema. Orphanet Journal of Rare Diseases, 2014, 9, 205.	1.2	24
32	A systematic analysis of the complement pathways in patients with neuromyelitis optica indicates alteration but no activation during remission. Molecular Immunology, 2014, 57, 200-209.	1.0	19
33	Home treatment of attacks with conestat alfa in hereditary angioedema due to C1-inhibitor deficiency. Allergy and Asthma Proceedings, 2014, 35, 255-259.	1.0	10