

# Genetic Variants in C5 and Poor Response to Eculizuma

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Citation Report

#	ARTICLE	IF	CITATIONS
1	Complement in paroxysmal nocturnal hemoglobinuria: exploiting our current knowledge to improve the treatment landscape. <i>Expert Review of Hematology</i> , 2014, 7, 583-598.	1.0	43
2	Control of innate immunological mechanisms as a route to drug minimization. <i>Current Opinion in Organ Transplantation</i> , 2014, 19, 342-347.	0.8	5
3	Report of the Inefficacy of Eculizumab in Two Cases of Severe Antibody-Mediated Rejection of Renal Grafts. <i>Transplantation</i> , 2014, 98, 1056-1059.	0.5	61
4	Syndromes of Thrombotic Microangiopathy. <i>New England Journal of Medicine</i> , 2014, 371, 654-666.	13.9	972
5	Efficacy and safety of eculizumab in children and adolescents with paroxysmal nocturnal hemoglobinuria. <i>Pediatric Blood and Cancer</i> , 2014, 61, 1544-1550.	0.8	35
8	Paroxysmal nocturnal hemoglobinuria. <i>Blood</i> , 2014, 124, 2804-2811.	0.6	424
9	Mutations in C5 explain eculizumab resistance. <i>Nature Reviews Nephrology</i> , 2014, 10, 182-182.	4.1	7
10	Complement in hemolytic anemia. <i>Hematology American Society of Hematology Education Program</i> , 2015, 2015, 385-391.	0.9	9
11	Atypical haemolytic uraemic syndrome treated with the complement inhibitor eculizumab: the experience of the <scp>A</scp>ustralian compassionate access cohort. <i>Internal Medicine Journal</i> , 2015, 45, 1054-1065.	0.5	25
12	Complement inhibition for paroxysmal nocturnal hemoglobinuria: where we stand and where we are going. <i>Expert Opinion on Orphan Drugs</i> , 2015, 3, 691-704.	0.5	2
13	Complement in hemolytic anemia. <i>Blood</i> , 2015, 126, 2459-2465.	0.6	84
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16	Complement activation, regulation, and molecular basis for complement-related diseases. <i>EMBO Journal</i> , 2015, 34, 2735-2757.	3.5	302
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18	Paroxysmal nocturnal hemoglobinuria: new concepts in pathophysiology and treatment. <i>Orphan Drugs: Research and Reviews</i> , 2015, , 75.	0.6	1
19	Diagnostic and therapeutic guidelines of thrombotic microangiopathies of the Spanish Apheresis Group. <i>Medicina Clínica (English Edition)</i> , 2015, 144, 331.e1-331.e13.	0.1	2
21	Eculizumab effect on the hemostatic state in patients with paroxysmal nocturnal hemoglobinuria. <i>Blood Cells, Molecules, and Diseases</i> , 2015, 54, 144-150.	0.6	18
22	Kidney Disease Caused by Dysregulation of the Complement Alternative Pathway. <i>Journal of the American Society of Nephrology: JASN</i> , 2015, 26, 2917-2929.	3.0	84

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23	Paroxysmal Nocturnal Hemoglobinuria. <i>Hematology/Oncology Clinics of North America</i> , 2015, 29, 479-494.	0.9	52
24	Paroxysmal nocturnal hemoglobinuria revisited: news on pathophysiology, clinical course and treatment. <i>Laboratoriums Medizin</i> , 2015, 39, 87-96.	0.1	0
25	Oxidative stress in paroxysmal nocturnal hemoglobinuria and other conditions of complement-mediated hemolysis. <i>Free Radical Biology and Medicine</i> , 2015, 88, 63-69.	1.3	15
26	Thrombotic Microangiopathy. <i>Hematology/Oncology Clinics of North America</i> , 2015, 29, 541-559.	0.9	25
27	Current and Future Pharmacologic Complement Inhibitors. <i>Hematology/Oncology Clinics of North America</i> , 2015, 29, 561-582.	0.9	30
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42	Update on the diagnosis and management of paroxysmal nocturnal hemoglobinuria. <i>Hematology American Society of Hematology Education Program</i> , 2016, 2016, 208-216.	0.9	85
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56	Emerging Therapies in Antiphospholipid Syndrome. <i>Current Rheumatology Reports</i> , 2016, 18, 22.	2.1	37
57	Complement therapeutics in inflammatory diseases: promising drug candidates for C3-targeted intervention. <i>Molecular Oral Microbiology</i> , 2016, 31, 3-17.	1.3	36
58	Structural basis for therapeutic inhibition of complement C5. <i>Nature Structural and Molecular Biology</i> , 2016, 23, 378-386.	3.6	94

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