

Long-term safety and efficacy of sustained eculizumab in patients with paroxysmal nocturnal haemoglobinuria

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Citation Report

#	ARTICLE	IF	CITATIONS
1	Paroxysmal nocturnal hemoglobinuria and the age of therapeutic complement inhibition. <i>Expert Review of Clinical Immunology</i> , 2013, 9, 1113-1124.	1.3	8
2	Learnings from over 25 years of PNH experience: The era of targeted complement inhibition. <i>Blood Reviews</i> , 2013, 27, S1-S6.	2.8	5
3	A systematic review of eculizumab for atypical haemolytic uraemic syndrome (aHUS). <i>BMJ Open</i> , 2013, 3, e003573.	0.8	80
4	Blood consult: paroxysmal nocturnal hemoglobinuria and its complications. <i>Blood</i> , 2013, 122, 2795-2798.	0.6	7
5	Meningococcal disease and the complement system. <i>Virulence</i> , 2014, 5, 98-126.	1.8	189
6	Eculizumab Therapy in Children with Severe Hematopoietic Stem Cell Transplantation-Associated Thrombotic Microangiopathy. <i>Biology of Blood and Marrow Transplantation</i> , 2014, 20, 518-525.	2.0	218
7	Treatment of Enterohemorrhagic <i>Escherichia coli</i> -Induced Hemolytic Uremic Syndrome (eHUS). <i>Seminars in Thrombosis and Hemostasis</i> , 2014, 40, 508-516.	1.5	40
8	Successful discontinuation of eculizumab therapy in a patient with aHUS. <i>Annals of Hematology</i> , 2014, 93, 1423-1425.	0.8	19
9	Diagnosis and management of complement mediated thrombotic microangiopathies. <i>Blood Reviews</i> , 2014, 28, 67-74.	2.8	43
10	Hemoglobinuria paroxística nocturna en una mujer de 86 años. <i>Revista Clinica Espanola</i> , 2014, 214, e111-e113.	0.2	0
11	Complement and cytokine response in acute thrombotic thrombocytopenic purpura. <i>British Journal of Haematology</i> , 2014, 164, 858-866.	1.2	49
12	Eculizumab for treating patients with paroxysmal nocturnal hemoglobinuria. <i>The Cochrane Library</i> , 2014, , CD010340.	1.5	13
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15	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
16	Paroxysmal nocturnal hemoglobinuria: a single Spanish center's experience over the last 40 years. <i>European Journal of Haematology</i> , 2014, 93, 309-319.	1.1	29
17	Eculizumab: a guide to its use in atypical haemolytic uraemic syndrome. <i>Drugs and Therapy Perspectives</i> , 2014, 30, 166-172.	0.3	1
18	Concerns about unapproved meningococcal vaccination for eculizumab therapy in Japan. <i>Orphanet Journal of Rare Diseases</i> , 2014, 9, 48.	1.2	3
19	New anti-complement drugs: not so far away. <i>Blood</i> , 2014, 123, 1975-1976.	0.6	13

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20	Peptide inhibitors of C3 activation as a novel strategy of complement inhibition for the treatment of paroxysmal nocturnal hemoglobinuria. <i>Blood</i> , 2014, 123, 2094-2101.	0.6	172
21	Paroxysmal nocturnal hemoglobinuria. <i>Blood</i> , 2014, 124, 2804-2811.	0.6	424
22	Complement in hemolytic anemia. <i>Hematology American Society of Hematology Education Program</i> , 2015, 2015, 385-391.	0.9	9
23	Increased eculizumab requirements during pregnancy in a patient with paroxysmal nocturnal hemoglobinuria: case report and review of the literature. <i>Clinical Case Reports (discontinued)</i> , 2015, 3, 88-91.	0.2	16
24	Atypical haemolytic uraemic syndrome treated with the complement inhibitor eculizumab: the experience of the Australian compassionate access cohort. <i>Internal Medicine Journal</i> , 2015, 45, 1054-1065.	0.5	25
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26	Complement in hemolytic anemia. <i>Blood</i> , 2015, 126, 2459-2465.	0.6	84
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30	Eculizumab in Typical Hemolytic Uremic Syndrome (HUS) With Neurological Involvement. <i>Medicine (United States)</i> , 2015, 94, e1000.	0.4	83
31	Use of Eculizumab in Patients With Allogeneic Stem Cell Transplant-Associated Thrombotic Microangiopathy. <i>Transplantation</i> , 2015, 99, 1953-1959.	0.5	110
32	Improvement of Renal Function by Long-Term Sustained Eculizumab Treatment in a Patient with Paroxysmal Nocturnal Hemoglobinuria. <i>Case Reports in Hematology</i> , 2015, 2015, 1-4.	0.3	1
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127	Paroxysmal Nocturnal Hemoglobinuria. , 2018, , 415-424.		1
128	Meningococcal B Vaccine Immunogenicity in Children With Defects in Complement and Splenic Function. <i>Pediatrics</i> , 2018, 142, .	1.0	17
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161	Intrinsic defects leading to increased erythrocyte destruction. , 2020, , 336-362.		0
162	Complement in neurological disorders and emerging complement-targeted therapeutics. Nature Reviews Neurology, 2020, 16, 601-617.	4.9	163
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