

Terminal Complement Inhibitor Eculizumab in Atypica

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Citation Report

#	ARTICLE	IF	CITATIONS
1	Guidelines on the Use of Therapeutic Apheresis in Clinical Practice—Evidence-Based Approach from the Writing Committee of the American Society for Apheresis: The Sixth Special Issue. <i>Journal of Clinical Apheresis</i> , 2013, 28, 145-284.	0.7	520
2	Clopidogrel with Aspirin in Minor Stroke or Transient Ischemic Attack. <i>New England Journal of Medicine</i> , 2013, 369, 1375-1377.	13.9	15
3	Eculizumab: A Review of Its Use in Atypical Haemolytic Uraemic Syndrome. <i>Drugs</i> , 2013, 73, 2053-2066.	4.9	52
4	HUS and TTP in Children. <i>Pediatric Clinics of North America</i> , 2013, 60, 1513-1526.	0.9	32
5	Atypical Hemolytic Uremic Syndrome. <i>Seminars in Nephrology</i> , 2013, 33, 508-530.	0.6	287
6	The immune system and kidney disease: basic concepts and clinical implications. <i>Nature Reviews Immunology</i> , 2013, 13, 738-753.	10.6	522
7	Targeted strategies in the prevention and management of atypical HUS recurrence after kidney transplantation. <i>Transplantation Reviews</i> , 2013, 27, 117-125.	1.2	74
8	Recurrence of glomerulonephritis after renal transplantation. <i>Transplantation Reviews</i> , 2013, 27, 126-134.	1.2	38
9	Complement therapy in atypical haemolytic uraemic syndrome (aHUS). <i>Molecular Immunology</i> , 2013, 56, 199-212.	1.0	70
12	Eculizumab in an anephric patient with atypical haemolytic uraemic syndrome and advanced vascular lesions. <i>Nephrology Dialysis Transplantation</i> , 2013, 28, 2899-2907.	0.4	25
13	A systematic review of eculizumab for atypical haemolytic uraemic syndrome (aHUS). <i>BMJ Open</i> , 2013, 3, e003573.	0.8	80
14	Managing and preventing atypical hemolytic uremic syndrome recurrence after kidney transplantation. <i>Current Opinion in Nephrology and Hypertension</i> , 2013, 22, 704-712.	1.0	58
15	Eculizumab in Atypical Hemolytic Uremic Syndrome. <i>New England Journal of Medicine</i> , 2013, 369, 1377-1380.	13.9	61
16	Eculizumab for atypical haemolytic uraemic syndrome: what next?. <i>Nature Reviews Nephrology</i> , 2013, 9, 495-496.	4.1	10
18	Successful Long-Term Treatment of TMA with Eculizumab in a Transplanted Patient with Atypical Hemolytic Uremic Syndrome Due to MCP Mutation. <i>Transplantation</i> , 2013, 96, e74-e76.	0.5	10
19	Anti-C5 as Prophylactic Therapy in Atypical Hemolytic Uremic Syndrome in Living-Related Kidney Transplantation. <i>Transplantation</i> , 2013, 96, e26-e29.	0.5	14
21	Endotheliopathies: Hemolytic Uremic Syndrome, Thrombotic Thrombocytopenic Purpura, and Preeclampsia. , 2014, , 2767-2787.		1
22	The Platelet as a Target for Damage. , 2014, , 3115-3138.		1

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23	Clinical and Renal Biopsy Findings Predicting Outcome in Renal Thrombotic Microangiopathy: A Large Cohort Study from a Single Institute in China. <i>Scientific World Journal</i> , The, 2014, 2014, 1-9.	0.8	25
24	Comprehensive Genetic Analysis of Complement and Coagulation Genes in Atypical Hemolytic Uremic Syndrome. <i>Japanese Journal of Pediatric Nephrology</i> , 2014, 27, 43-44.	0.0	1
26	Meningococcal disease and the complement system. <i>Virulence</i> , 2014, 5, 98-126.	1.8	189
27	Postpartum thrombotic microangiopathy revealed as atypical hemolytic uremic syndrome successfully treated with eculizumab: a case report. <i>Journal of Medical Case Reports</i> , 2014, 8, 307.	0.4	31
28	First-line therapy in atypical hemolytic uremic syndrome: consideration on infants with a poor prognosis. <i>Italian Journal of Pediatrics</i> , 2014, 40, 101.	1.0	8
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32	Atypical Hemolytic Uremic Syndrome: Differential Diagnosis from TTP/HUS and Management. <i>Turkish Journal of Haematology</i> , 2014, 31, 216-25.	0.2	8
33	Pharmacological treatment of atypical hemolytic-uremic syndrome. <i>Expert Opinion on Orphan Drugs</i> , 2014, 2, 123-135.	0.5	1
34	Complement Mutations in Diacylglycerol Kinase-Associated Atypical Hemolytic Uremic Syndrome. <i>Clinical Journal of the American Society of Nephrology: CJASN</i> , 2014, 9, 1611-1619.	2.2	61
35	Natural History of Thrombotic Thrombocytopenic Purpura and Hemolytic Uremic Syndrome. <i>Seminars in Thrombosis and Hemostasis</i> , 2014, 40, 866-873.	1.5	33
36	Differing Tales of Two Patients after Receiving a Kidney Transplant from a Donor with Disseminated Intravascular Coagulation. <i>Case Reports in Transplantation</i> , 2014, 2014, 1-5.	0.1	0
37	Atypical haemolytic uraemic syndrome associated with a CD46 mutation triggered by <i>Shigella flexneri</i> . <i>CKJ: Clinical Kidney Journal</i> , 2014, 7, 286-288.	1.4	12
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40	Anti-factor H autoantibody-associated hemolytic uremic syndrome: the earlier diagnosed and treated, the better. <i>Kidney International</i> , 2014, 85, 1019-1022.	2.6	12
42	Use of eculizumab in refractory gemcitabine-induced thrombotic microangiopathy. <i>British Journal of Haematology</i> , 2014, 164, 894-896.	1.2	26

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45	Targeted Therapy with Eculizumab for Inherited CD59 Deficiency. New England Journal of Medicine, 2014, 370, 90-92.	13.9	55
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59	Complement Factor B Mutations in Atypical Hemolytic Uremic Syndrome—Disease-Relevant or Benign?. Journal of the American Society of Nephrology: JASN, 2014, 25, 2053-2065.	3.0	107
60	Eculizumab Therapy Leads to Rapid Resolution of Thrombocytopenia in Atypical Hemolytic Uremic Syndrome. Advances in Hematology, 2014, 2014, 1-7.	0.6	19
61	Insights From the Use in Clinical Practice of Eculizumab in Adult Patients With Atypical Hemolytic Uremic Syndrome Affecting the Native Kidneys: An Analysis of 19 Cases. American Journal of Kidney Diseases, 2014, 63, 40-48.	2.1	74
62	Anti-Complement Therapy for Glomerular Diseases. Advances in Chronic Kidney Disease, 2014, 21, 152-158.	0.6	15

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65	Haemolytic uremic syndrome following fire ant bites. <i>BMC Nephrology</i> , 2014, 15, 5.	0.8	11
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72	Eculizumab in atypical haemolytic uraemic syndrome with severe cardiac and neurological involvement. <i>Pediatric Nephrology</i> , 2014, 29, 1103-1106.	0.9	23
73	Eculizumab as First-Line Therapy for Atypical Hemolytic Uremic Syndrome. <i>Pediatrics</i> , 2014, 133, e1759-e1763.	1.0	22
74	Cryptic Activity of Atypical Hemolytic Uremic Syndrome and Eculizumab Treatment. <i>Pediatrics</i> , 2014, 133, e1769-e1771.	1.0	10
75	Genetics of Atypical Hemolytic Uremic Syndrome (aHUS). <i>Seminars in Thrombosis and Hemostasis</i> , 2014, 40, 422-430.	1.5	122
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80	Macrovascular involvement in a child with atypical hemolytic uremic syndrome. <i>Pediatric Nephrology</i> , 2014, 29, 1273-1277.	0.9	17
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83	Time to change the definition of hemolytic uremic syndrome. <i>European Journal of Internal Medicine</i> , 2014, 25, e29.	1.0	22
84	Cardiovascular complications in atypical haemolytic uraemic syndrome. <i>Nature Reviews Nephrology</i> , 2014, 10, 174-180.	4.1	63
85	Diagnosis and management of complement mediated thrombotic microangiopathies. <i>Blood Reviews</i> , 2014, 28, 67-74.	2.8	43
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87	Autoantibodies to complement components in C3 glomerulopathy and atypical hemolytic uremic syndrome. <i>Immunology Letters</i> , 2014, 160, 163-171.	1.1	50
88	Liverâ€“kidney transplantation to cure atypical HUS: still an option post-eculizumab?. <i>Pediatric Nephrology</i> , 2014, 29, 329-332.	0.9	50
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94	Factors influencing long-term outcome after kidney transplantation. <i>Transplant International</i> , 2014, 27, 19-27.	0.8	176
95	Innovative therapeutic approach: Sequential treatment with plasma exchange and eculizumab in a pregnant woman affected by atypical hemolytic-uremic syndrome. <i>Transfusion and Apheresis Science</i> , 2014, 51, 134-136.	0.5	21
96	Syndromes of Thrombotic Microangiopathy. <i>New England Journal of Medicine</i> , 2014, 371, 654-666.	13.9	972
97	Ten-year advances in immunopathology of glomerulonephritis: translated into patientsâ€™ care or lost in translation?. <i>Seminars in Immunopathology</i> , 2014, 36, 377-379.	2.8	5
98	An audit analysis of a guideline for the investigation and initial therapy of diarrhea negative (atypical) hemolytic uremic syndrome. <i>Pediatric Nephrology</i> , 2014, 29, 1967-1978.	0.9	95
99	Ecuzumab: a guide to its use in atypical haemolytic uraemic syndrome. <i>Drugs and Therapy Perspectives</i> , 2014, 30, 166-172.	0.3	1
100	How I treat: the clinical differentiation and initial treatment of adult patients with atypical hemolytic uremic syndrome. <i>Blood</i> , 2014, 123, 2478-2484.	0.6	115
101	Ecuzumab in neonatal hemolytic uremic syndrome with homozygous factor H deficiency. <i>Pediatric Nephrology</i> , 2014, 29, 2415-2419.	0.9	18

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103	Disorders of complement regulation. <i>Drug Discovery Today: Disease Models</i> , 2014, 11, 29-35.	1.2	0
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107	Diagnostic criteria for atypical hemolytic uremic syndrome proposed by the International Committee of the Japanese Society of Nephrology and the Japanese Pediatric Society . <i>Pediatrics International</i> , 2014, 56, 1-5.	0.2	29
108	Acquired Bleeding Disorders. <i>Emergency Medicine Clinics of North America</i> , 2014, 32, 691-713.	0.5	9
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111	Hemolytic Uremic Syndrome: Toxins, Vessels, and Inflammation. <i>Frontiers in Medicine</i> , 2014, 1, 42.	1.2	10
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116	Diagnostic and risk criteria for HSCT-associated thrombotic microangiopathy: a study in children and young adults. <i>Blood</i> , 2014, 124, 645-653.	0.6	318
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121	Syndromes of thrombotic microangiopathy associated with pregnancy. <i>Hematology American Society of Hematology Education Program</i> , 2015, 2015, 644-648.	0.9	79
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127	7. Thrombotic Microangiopathy: HUS and Atypical HUS. <i>The Journal of the Japanese Society of Internal Medicine</i> , 2015, 104, 1959-1963.	0.0	0
128	Refractory atypical hemolytic uremic syndrome with monoclonal gammopathy responsive to bortezomib-based therapy. <i>Clinical Nephrology</i> , 2015, 83 (2015), 363-369.	0.4	24
129	Progression to end-stage renal disease is reduced with eculizumab in patients with atypical haemolytic uraemic syndrome. <i>Critical Care</i> , 2015, 19, .	2.5	0
130	Early initiation of eculizumab treatment in patients with atypical haemolytic uraemic syndrome improves long-term outcomes: a pooled analysis of clinical trials. <i>Critical Care</i> , 2015, 19, .	2.5	0
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133	The global aHUS registry: methodology and initial patient characteristics. <i>BMC Nephrology</i> , 2015, 16, 207.	0.8	52
134	Solving the puzzle of posthospital recovery: What is the role of the individual physician?. <i>Journal of Hospital Medicine</i> , 2015, 10, 697-700.	0.7	4
135	Overactivation of Complement Alternative Pathway in Postpartum Atypical Hemolytic Uremic Syndrome Patients with Renal Involvement. <i>American Journal of Reproductive Immunology</i> , 2015, 74, 345-356.	1.2	17
136	Role of the Skin Biopsy in the Diagnosis of Atypical Hemolytic Uremic Syndrome. <i>American Journal of Dermatopathology</i> , 2015, 37, 349-359.	0.3	35
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144	Role of Complement and Complement Regulatory Proteins in the Complications of Diabetes. <i>Endocrine Reviews</i> , 2015, 36, 272-288.	8.9	127
145	Clinicopathologic Characteristics and Outcomes of Renal Thrombotic Microangiopathy in Anti-Neutrophil Cytoplasmic Autoantibody-Associated Glomerulonephritis. <i>Clinical Journal of the American Society of Nephrology: CJASN</i> , 2015, 10, 750-758.	2.2	30
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147	Soluble C5b-9 as a Biomarker for Complement Activation in Atypical Hemolytic Uremic Syndrome. <i>American Journal of Kidney Diseases</i> , 2015, 65, 968-969.	2.1	55
148	Plasma resistant atypical hemolytic uremic syndrome associated with a CFH mutation treated with eculizumab: a case report. <i>Journal of Medical Case Reports</i> , 2015, 9, 92.	0.4	11
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150	Characterization of a New DGKE Intronic Mutation in Genetically Unsolved Cases of Familial Atypical Hemolytic Uremic Syndrome. <i>Clinical Journal of the American Society of Nephrology: CJASN</i> , 2015, 10, 1011-1019.	2.2	47
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155	Thrombotic microangiopathy: expanding genetic, clinical and therapeutic spectra and the need for worldwide implementation of recent advances. <i>CKJ: Clinical Kidney Journal</i> , 2015, 8, 686-689.	1.4	3
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161	Efficacy and safety of eculizumab in atypical hemolytic uremic syndrome from 2-year extensions of phase 2 studies. <i>Kidney International</i> , 2015, 87, 1061-1073.	2.6	342
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163	Complement Factor C4d Is a Common Denominator in Thrombotic Microangiopathy. <i>Journal of the American Society of Nephrology: JASN</i> , 2015, 26, 2239-2247.	3.0	97
164	Molecules Great and Small. <i>Clinical Journal of the American Society of Nephrology: CJASN</i> , 2015, 10, 1636-1650.	2.2	217
165	Safe and effective use of eculizumab in the treatment of severe Shiga toxin-producing <i>Escherichia coli</i> -associated hemolytic uremic syndrome. <i>American Journal of Health-System Pharmacy</i> , 2015, 72, 117-120.	0.5	14
166	Podocyte dysfunction in atypical haemolytic uraemic syndrome. <i>Nature Reviews Nephrology</i> , 2015, 11, 245-252.	4.1	49
167	Quality of life one year post-Shiga toxin-producing <i>Escherichia coli</i> O104 infection—A prospective cohort study. <i>Neurogastroenterology and Motility</i> , 2015, 27, 370-378.	1.6	5
168	Complement regulators in human disease: lessons from modern genetics. <i>Journal of Internal Medicine</i> , 2015, 277, 294-305.	2.7	60
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170	Eculizumab hepatotoxicity in pediatric aHUS. <i>Pediatric Nephrology</i> , 2015, 30, 775-781.	0.9	39
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172	Complement regulator CD46: genetic variants and disease associations. <i>Human Genomics</i> , 2015, 9, 7.	1.4	87
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