Hydroxyurea (hydroxycarbamide) for sickle cell disease

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Citation Report

#	Article	IF	CITATIONS
1	Effects of Hydroxyurea Exposure on the Rat Cerebellar Neuroepithelium: an Immunohistochemical and Electron Microscopic Study Along the Anteroposterior and Mediolateral Axes. Neurotoxicity Research, 2017, 32, 671-682.	1.3	6
2	Pharmacotherapeutical strategies in the prevention of acute, vaso-occlusive pain in sickle cell disease: a systematic review. Blood Advances, 2017, 1, 1598-1616.	2.5	34
3	Phytomedicines (medicines derived from plants) for sickle cell disease. The Cochrane Library, 2018, 2, CD004448.	1.5	6
4	Hydroxyurea (hydroxycarbamide) genotoxicity in pediatric patients with sickle cell disease. Pediatric Blood and Cancer, 2018, 65, e27022.	0.8	9
5	Improving routine outpatient monitoring for patients with sickle-cell disease on hydroxyurea. BMJ Open Quality, 2018, 7, e000218.	0.4	2
6	Sickle cell crisis: A crisis of a different sort?. Archives of Disease in Childhood: Education and Practice Edition, 2018, 103, edpract-2017-313899.	0.3	1
7	Endothelin type A receptors mediate pain in a mouse model of sickle cell disease. Haematologica, 2018, 103, 1124-1135.	1.7	25
8	Sickle cell disease: a malady beyond a hemoglobin defect in cerebrovascular disease. Expert Review of Hematology, 2018, 11, 45-55.	1.0	15
9	Association between Participants' Characteristics, Patient-Reported Outcomes, and Clinical Outcomes in Youth with Sickle Cell Disease. BioMed Research International, 2018, 2018, 1-8.	0.9	29
10	TRIAMF: A New Method for Delivery of Cas9 Ribonucleoprotein Complex to Human Hematopoietic Stem Cells. Scientific Reports, 2018, 8, 16304.	1.6	36
11	Metal Antagonists and Metals. Side Effects of Drugs Annual, 2018, 40, 279-288.	0.6	3
12	Targeted Hydroxyurea Education after an Emergency Department Visit Increases Hydroxyurea Use in Children with Sickle Cell Anemia. Journal of Pediatrics, 2018, 201, 221-228.e16.	0.9	12
13	Treatment patterns and economic burden of sickle-cell disease patients prescribed hydroxyurea: a retrospective claims-based study. Health and Quality of Life Outcomes, 2019, 17, 155.	1.0	40
14	Effect of N(Epsilon)-(carboxymethyl)lysine on Laboratory Parameters and Its Association withÎ <sup>2</sup> SHaplotype in Children with Sickle Cell Anemia. Disease Markers, 2019, 2019, 1-8.	0.6	2
15	cGMP modulation therapeutics for sickle cell disease. Experimental Biology and Medicine, 2019, 244, 132-146.	1.1	21
17	Association between clinical outcomes and metformin use in adults with sickle cell disease and diabetes mellitus. Blood Advances, 2019, 3, 3297-3306.	2.5	9
18	Interventions for treating neuropathic pain in people with sickle cell disease. The Cochrane Library, 2019, 7, CD012943.	1.5	6
19	Phytomedicines (medicines derived from plants) for sickle cell disease. The Cochrane Library, 2020, 9, CD004448.	1.5	2

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20	Polymorphisms in Inflammatory Genes Modulate Clinical Complications in Patients With Sickle Cell Disease. Frontiers in Immunology, 2020, 11, 2041.	2.2	10
21	Current and novel therapies for the prevention of vaso-occlusive crisis in sickle cell disease. Therapeutic Advances in Hematology, 2020, 11, 204062072095500.	1.1	22
22	Hydroxyurea treatment is associated with reduced degree of oxidative perturbation in children and adolescents with Asickle cell anemia. Scientific Reports, 2020, 10, 18982.	1.6	5
23	Crizanlizumab and comparators for adults with sickle cell disease: a systematic review and network meta-analysis. BMJ Open, 2020, 10, e034147.	0.8	7
24	Association of HIV infection with clinical and laboratory characteristics of sickle cell disease. BMC Infectious Diseases, 2020, 20, 638.	1.3	4
25	Foetal haemoglobin inducers for reducing blood transfusion in non-transfusion dependent beta thalassaemias. The Cochrane Library, 0, , .	1.5	0
26	Patientâ€reported outcomes in sickle cell disease and association with clinical and psychosocial factors: Report from the sickle cell disease implementation consortium. American Journal of Hematology, 2020, 95, 1066-1074.	2.0	24
27	Recent Advances in the Treatment of Sickle Cell Disease. Frontiers in Physiology, 2020, 11, 435.	1.3	114
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35	PROFILE OF WOMEN WITH LEG ULCERS DUE TO SICKLE CELL DISEASE. ESTIMA Brazilian Journal of Enterostomal Therapy, $0, , .$	0.1	1
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40	Innovative Treatments for Rare Anemias. HemaSphere, 2021, 5, e576.	1.2	13
41	Time to rethink haemoglobin threshold guidelines in sickle cell disease. British Journal of Haematology, 2021, 195, 518-522.	1.2	7
42	Hydroxyurea—The Good, the Bad and the Ugly. Genes, 2021, 12, 1096.	1.0	49
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49	Hydroxyurea: Pattern of Use, Patient Adherence, and Safety Profile in Patients with Sickle Cell Disease in Oman. Oman Medical Journal, 2019, 34, 327-335.	0.3	12
50	Homozygous sickle cell disease related mortality in Senegal (2011–2020). EJHaem, 2021, 2, 711-715.	0.4	2
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54	Sickle Cell Retinopathy., 2020,, 1-21.		0
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58	Higher hydroxyurea adherence among young adults with sickle cell disease compared to children and adolescents. Annals of Medicine, 2022, 54, 683-693.	1.5	4
59	Testicular Tissue Banking for Fertility Preservation in Young Boys: Which Patients Should Be Included?. Frontiers in Endocrinology, 2022, 13, 854186.	1.5	15
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61	Life-Threatening Acute Chest Syndrome in a Patient With Sickle Cell Disease After Switching From Hydroxyurea Therapy to Partial Exchange Transfusions: A Case Report. Cureus, 2021, 13, e20236.	0.2	0
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73	Health-Related Quality of Life and Adherence to Hydroxyurea and Other Disease-Modifying Therapies among Individuals with Sickle Cell Disease: A Systematic Review. BioMed Research International, 2022, 2022, 1-8.	0.9	6
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85	Transcranial Doppler Ultrasonography-Related Research in the Caribbean Region. Cureus, 2023, , .	0.2	0
86	Allogeneic hematopoietic stem cell transplantation to cure sickle cell disease: A review. Frontiers in Medicine, 0, $10$ , .	1.2	7
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