

Hemoglobin Koya Dora: high frequency of a chain term

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Citation Report

#	ARTICLE	IF	CITATIONS
1	Nucleotide sequences of the 3'-terminal untranslated region of messenger RNA for human beta globin chain.. Proceedings of the National Academy of Sciences of the United States of America, 1975, 72, 3614-3618.	7.1	31
2	Hemoglobin Cranston, an unstable variant having an elongated beta chain due to nonhomologous crossover between two normal beta chain genes.. Proceedings of the National Academy of Sciences of the United States of America, 1975, 72, 3609-3613.	7.1	69
3	Haemoglobin Tak: a $\beta^2$ -Chain Elongation. British Journal of Haematology, 1975, 31, 119-131.	2.5	41
4	Molecular Basis of Thalassaemia. British Journal of Haematology, 1975, 31, 133-141.	2.5	5
5	Hemoglobin Wayne: a frameshift mutation detected in human hemoglobin alpha chains.. Proceedings of the National Academy of Sciences of the United States of America, 1976, 73, 882-886.	7.1	67
6	Trimodality in the proportion of hemoglobin G Philadelphia in heterozygotes: evidence for heterogeneity in the number of human alpha chain loci.. Proceedings of the National Academy of Sciences of the United States of America, 1976, 73, 3633-3636.	7.1	44
7	Is haemoglobin G? Philadelphia linked to $\beta$ -thalassaemia?. Human Genetics, 1976, 31, 67-74.	3.8	14
8	Variant Lists: Variants of the Delta Chain. Hemoglobin, 1977, 1, 887-897.	0.8	2
9	Hemoglobin TAK in a Newborn Malay. Hemoglobin, 1977, 1, 747-757.	0.8	7
10	Synthesis of haemoglobin Wayne in erythroid cells. Nature, 1977, 269, 717-719.	27.8	7
11	Alpha-thalassemia. American Journal of Hematology, 1977, 2, 317-325.	4.1	11
13	Genetic and biosynthetic studies of families carrying hemoglobin J ? Mexico: Association of $\beta$ -thalassemia with Hb J. Human Genetics, 1978, 42, 189-199.	3.8	7
14	Mutation rates from rare variants of proteins in Indian tribes. Human Genetics, 1978, 43, 179-183.	3.8	9
15	Mutant Hemoglobins Having Elongated Chains. Hemoglobin, 1978, 2, 1-28.	0.8	8
16	High genetic polymorphism of hemoglobin disorders in Laos. Human Genetics, 1979, 50, 327-336.	3.8	26
17	beta 0 thalassemia, a nonsense mutation in man.. Proceedings of the National Academy of Sciences of the United States of America, 1979, 76, 2886-2889.	7.1	266
18	$\beta$ THALASSEMIA AND THE EXPRESSION OF HEMOGLOBIN G-PHILADELPHIA. Annals of the New York Academy of Sciences, 1980, 344, 62-72.	3.8	3
19	International Hemoglobin Information Center Policies - Ihic. Hemoglobin, 1981, 5, 259-323.	0.8	1

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20	Hemoglobinopathies in India. Hemoglobin, 1981, 5, 751-766.	0.8	11
21	International Hemoglobin Information Center Policies - IHIC. Hemoglobin, 1982, 6, 257-346.	0.8	8
22	Identification of hemoglobin G-Philadelphia ( $\beta^{68} \text{Asn} \rightarrow \text{Lys}$ ) and homoglobin matsue-oki ( $\beta^{75} \text{Asp} \rightarrow \text{Asn}$ ) in a black infant. BBA - Proteins and Proteomics, 1982, 707, 206-212.	2.1	8
23	Haemoglobin Constant Spring has an unstable $\beta^+$ chain messenger RNA. British Journal of Haematology, 1982, 51, 405-413.	2.5	74
24	Mapping the $\beta$ -globin genes in HB J Mexico carriers. Human Genetics, 1982, 62, 164-6.	3.8	4
25	International Hemoglobin Information Center: Policies of IHIC. Hemoglobin, 1983, 7, 331-406.	0.8	3
26	Molecular Pathology of $\beta$ -Thalassemia. Annals of the New York Academy of Sciences, 1985, 445, 28-36.	3.8	33
27	International Hemoglobin Information Center IHIC Variants List. Hemoglobin, 1986, 10, 261-327.	0.8	11
28	IHIC Variants List. Hemoglobin, 1987, 11, 243-308.	0.8	2
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34	International Hemoglobin Information Center. Hemoglobin, 1990, 14, 249-325.	0.8	2
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36	The Interactions of $\beta^+$ -Thalassemia with Hemoglobinopathies. Hematology/Oncology Clinics of North America, 1991, 5, 453-473.	2.2	22
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44	International Hemoglobin Information Center. <i>Hemoglobin</i> , 1994, 18, 77-161.	0.8	24
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49	International Hemoglobin Information Center Variant List. <i>Hemoglobin</i> , 1997, 21, 505-506.	0.8	7
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58	$\alpha$ -thalassaemia. Orphanet Journal of Rare Diseases, 2010, 5, 13.	2.7	417
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60	NEGATIVE EPISTASIS BETWEEN $\alpha$ + THALASSAEMIA AND SICKLE CELL TRAIT CAN EXPLAIN INTERPOPULATION VARIATION IN SOUTH ASIA. Evolution; International Journal of Organic Evolution, 2011, 65, 3625-3632.	2.3	20
61	First case of Hb Fontainebleau with sickle haemoglobin and other non-deletional $\alpha$ gene variants identified in neonates during newborn screening for sickle cell disorders. Journal of Clinical Pathology, 2012, 65, 654-659.	2.0	18
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69	Erythroid Cell-Specific Determinants of $\alpha$ -Globin mRNA Stability. Molecular and Cellular Biology, 1994, 14, 8123-8132.	2.3	73
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77	Thalassemic hemoglobinopathies. American Journal of Pathology, 1983, 113, 396-409.	3.8	15