LuÃ-sa RomÃ£o

List of Publications by Year
in descending order

40 papers	1,843 citations	331670 21 h-index	289244 40 g-index
44 all docs	44 docs citations	44 times ranked	3221 citing authors

#	Article	IF	Citations
1	Gene Variants Involved in Nonsense-Mediated mRNA Decay Suggest a Role in Autism Spectrum Disorder. Biomedicines, 2022, 10, 665.	3.2	6
2	Nonsense suppression therapies in human genetic diseases. Cellular and Molecular Life Sciences, 2021, 78, 4677-4701.	5.4	38
3	Nonsense-mediated RNA decay and its bipolar function in cancer. Molecular Cancer, 2021, 20, 72.	19.2	40
4	Translation of ABCE1 Is Tightly Regulated by Upstream Open Reading Frames in Human Colorectal Cells. Biomedicines, 2021, 9, 911.	3.2	6
5	Experimental supporting data on DIS3L2 over nonsense-mediated mRNA decay targets in human cells. Data in Brief, 2020, 28, 104943.	1.0	2
6	Perspective in Alternative Splicing Coupled to Nonsense-Mediated mRNA Decay. International Journal of Molecular Sciences, 2020, 21, 9424.	4.1	39
7	Nonsense-Mediated mRNA Decay in Development, Stress and Cancer. Advances in Experimental Medicine and Biology, 2019, 1157, 41-83.	1.6	15
8	Translational Regulation by Upstream Open Reading Frames and Human Diseases. Advances in Experimental Medicine and Biology, 2019, 1157, 99-116.	1.6	32
9	A role for DIS3L2 over natural nonsense-mediated mRNA decay targets in human cells. Biochemical and Biophysical Research Communications, 2019, 518, 664-671.	2.1	11
10	eIF3: a factor for human health and disease. RNA Biology, 2018, 15, 26-34.	3.1	70
11	More than just scanning: the importance of cap-independent mRNA translation initiation for cellular stress response and cancer. Cellular and Molecular Life Sciences, 2017, 74, 1659-1680.	5.4	98
12	Cap-independent translation ensures mTOR expression and function upon protein synthesis inhibition. Rna, 2017, 23, 1712-1728.	3.5	22
13	The role of alternative splicing coupled to nonsense-mediated mRNA decay in human disease. International Journal of Biochemistry and Cell Biology, 2017, 91, 168-175.	2.8	58
14	<scp>PROS</scp>1 novel spliceâ€site variant decreases protein S expression in patients from two families with thrombotic disease. Clinical Case Reports (discontinued), 2017, 5, 2062-2065.	0.5	4
15	Expression of Human Hemojuvelin (HJV) Is Tightly Regulated by Two Upstream Open Reading Frames in HJV mRNA That Respond to Iron Overload in Hepatic Cells. Molecular and Cellular Biology, 2015, 35, 1376-1389.	2.3	10
16	Resistance of mRNAs with AUG-proximal nonsense mutations to nonsense-mediated decay reflects variables of mRNA structure and translational activity. Nucleic Acids Research, 2015, 43, 6528-6544.	14.5	30
17	Translation of the human erythropoietin transcript is regulated by an upstream open reading frame in response to hypoxia. Rna, 2014, 20, 594-608.	3.5	28
18	Gene Expression Regulation by Upstream Open Reading Frames and Human Disease. PLoS Genetics, 2013, 9, e1003529.	3.5	455

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19	Interaction of PABPC1 with the translation initiation complex is critical to the NMD resistance of AUG-proximal nonsense mutations. Nucleic Acids Research, 2012, 40, 1160-1173.	14.5	109
20	Unspliced Precursors of NMD-Sensitive Î²-Globin Transcripts Exhibit Decreased Steady-State Levels in Erythroid Cells. PLoS ONE, 2012, 7, e38505.	2.5	5
21	A new function of ROD1 in nonsenseâ€mediated mRNA decay. FEBS Letters, 2012, 586, 1101-1110.	2.8	26
22	Alternative Polyadenylation and Nonsense-Mediated Decay Coordinately Regulate the Human HFE mRNA Levels. PLoS ONE, 2012, 7, e35461.	2.5	11
23	Control of human Â-globin mRNA stability and its impact on beta-thalassemia phenotype. Haematologica, 2011, 96, 905-913.	3.5	41
24	The mammalian nonsenseâ€mediated mRNA decay pathway: To decay or not to decay! Which players make the decision?. FEBS Letters, 2009, 583, 499-505.	2.8	121
25	Early modification of sickle cell disease clinical course by UDP-glucuronosyltransferase 1A1 gene promoter polymorphism. Journal of Human Genetics, 2008, 53, 524-528.	2.3	12
26	Proximity of the poly(A)-binding protein to a premature termination codon inhibits mammalian nonsense-mediated mRNA decay. Rna, 2008, 14, 563-576.	3.5	132
27	Hb Evora [Â2-35 (B16), Ser->Pro], a novel hemoglobin variant associated with an Â-thalassemia phenotype. Haematologica, 2007, 92, 252-253.	3.5	9
28	Epidemiology of haemoglobin disorders in Europe: an overview. Scandinavian Journal of Clinical and Laboratory Investigation, 2007, 67, 39-70.	1.2	134
29	Comment on â€˜Nonsense-mediated mRNA decay modulates clinical outcome of genetic diseaseâ€™. European Journal of Human Genetics, 2007, 15, 533-534.	2.8	4
30	Mutational spectrum of delta-globin gene in the Portuguese population. European Journal of Haematology, 2007, 79, 422-428.	2.2	27
31	Human alpha2-globin nonsense-mediated mRNA decay induced by a novel alpha-thalassaemia frameshift mutation at codon 22. British Journal of Haematology, 2006, 133, 98-102.	2.5	17
32	Hemoglobin Loves Park [Î²68 (E12) Leuâ†'Phe]: Report of five cases including one originating from a de novo mutation. American Journal of Hematology, 2006, 81, 256-261.	4.1	2
33	The canonical UPF1-dependent nonsense-mediated mRNA decay is inhibited in transcripts carrying a short open reading frame independent of sequence context. Rna, 2006, 12, 2160-2170.	3.5	40
34	HFE gene mutations are extremely rare in Western sub-Saharan Africa. Annals of Hematology, 2005, 84, 686-688.	1.8	2
35	Hb YaoundÃ© [Î²134(H12)Valâ†'Ala] in Association with Hb C [Î²6(A3)Gluâ†'Lys] in a Caucasian Portuguese Family. Hemoglobin, 2004, 28, 229-235.	0.8	3
36	Nonsense Mutations in Close Proximity to the Initiation Codon Fail to Trigger Full Nonsense-mediated mRNA Decay. Journal of Biological Chemistry, 2004, 279, 32170-32180.	3.4	116

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37	The role of HFE mutations on iron metabolism in beta-thalassemia carriers. Journal of Human Genetics, 2004, 49, 651-655.	2.3	34
38	COMPOUND HETEROZYGOSITY FOR Hb SPANISH TOWN [Î±27(B8)Gluâ†'Val], Hb S [Î²6(A3)Gluâ†'Val] AND THE â^'Î±(3.7) Tj ETQq0 0 0	0.8	2
39	Asymptomatic homozygous deletional Î²0-thalassemia in an African individual. American Journal of Hematology, 2002, 70, 232-236.	4.1	3
40	Molecular Basis of A-Thalassa in Portugal. Hemoglobin, 1995, 19, 343-352.	0.8	20