

LuÃ-sa RomÃ£o

List of Publications by Year in descending order

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Version: 2024-02-01

40
papers

1,843
citations

331538

21
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289141

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. <i>Biomedicines</i> , 2022, 10, 665. | 1.4 | 6 |
| 2 | Nonsense suppression therapies in human genetic diseases. <i>Cellular and Molecular Life Sciences</i> , 2021, 78, 4677-4701. | 2.4 | 38 |
| 3 | Nonsense-mediated RNA decay and its bipolar function in cancer. <i>Molecular Cancer</i> , 2021, 20, 72. | 7.9 | 40 |
| 4 | Translation of ABCE1 Is Tightly Regulated by Upstream Open Reading Frames in Human Colorectal Cells. <i>Biomedicines</i> , 2021, 9, 911. | 1.4 | 6 |
| 5 | Experimental supporting data on DIS3L2 over nonsense-mediated mRNA decay targets in human cells. <i>Data in Brief</i> , 2020, 28, 104943. | 0.5 | 2 |
| 6 | Perspective in Alternative Splicing Coupled to Nonsense-Mediated mRNA Decay. <i>International Journal of Molecular Sciences</i> , 2020, 21, 9424. | 1.8 | 39 |
| 7 | Nonsense-Mediated mRNA Decay in Development, Stress and Cancer. <i>Advances in Experimental Medicine and Biology</i> , 2019, 1157, 41-83. | 0.8 | 15 |
| 8 | Translational Regulation by Upstream Open Reading Frames and Human Diseases. <i>Advances in Experimental Medicine and Biology</i> , 2019, 1157, 99-116. | 0.8 | 32 |
| 9 | A role for DIS3L2 over natural nonsense-mediated mRNA decay targets in human cells. <i>Biochemical and Biophysical Research Communications</i> , 2019, 518, 664-671. | 1.0 | 11 |
| 10 | eIF3: a factor for human health and disease. <i>RNA Biology</i> , 2018, 15, 26-34. | 1.5 | 70 |
| 11 | More than just scanning: the importance of cap-independent mRNA translation initiation for cellular stress response and cancer. <i>Cellular and Molecular Life Sciences</i> , 2017, 74, 1659-1680. | 2.4 | 98 |
| 12 | Cap-independent translation ensures mTOR expression and function upon protein synthesis inhibition. <i>Rna</i> , 2017, 23, 1712-1728. | 1.6 | 22 |
| 13 | The role of alternative splicing coupled to nonsense-mediated mRNA decay in human disease. <i>International Journal of Biochemistry and Cell Biology</i> , 2017, 91, 168-175. | 1.2 | 58 |
| 14 | <i>PROS</i> novel splice site variant decreases protein S expression in patients from two families with thrombotic disease. <i>Clinical Case Reports (discontinued)</i> , 2017, 5, 2062-2065. | 0.2 | 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. <i>Molecular and Cellular Biology</i> , 2015, 35, 1376-1389. | 1.1 | 10 |
| 16 | Resistance of mRNAs with AUG-proximal nonsense mutations to nonsense-mediated decay reflects variables of mRNA structure and translational activity. <i>Nucleic Acids Research</i> , 2015, 43, 6528-6544. | 6.5 | 30 |
| 17 | Translation of the human erythropoietin transcript is regulated by an upstream open reading frame in response to hypoxia. <i>Rna</i> , 2014, 20, 594-608. | 1.6 | 28 |
| 18 | Gene Expression Regulation by Upstream Open Reading Frames and Human Disease. <i>PLoS Genetics</i> , 2013, 9, e1003529. | 1.5 | 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. <i>Nucleic Acids Research</i> , 2012, 40, 1160-1173. | 6.5 | 109 |
| 20 | Unspliced Precursors of NMD-Sensitive Î²-Globin Transcripts Exhibit Decreased Steady-State Levels in Erythroid Cells. <i>PLoS ONE</i> , 2012, 7, e38505. | 1.1 | 5 |
| 21 | A new function of ROD1 in nonsense-mediated mRNA decay. <i>FEBS Letters</i> , 2012, 586, 1101-1110. | 1.3 | 26 |
| 22 | Alternative Polyadenylation and Nonsense-Mediated Decay Coordinately Regulate the Human HFE mRNA Levels. <i>PLoS ONE</i> , 2012, 7, e35461. | 1.1 | 11 |
| 23 | Control of human Î²-globin mRNA stability and its impact on beta-thalassemia phenotype. <i>Haematologica</i> , 2011, 96, 905-913. | 1.7 | 41 |
| 24 | The mammalian nonsense-mediated mRNA decay pathway: To decay or not to decay! Which players make the decision?. <i>FEBS Letters</i> , 2009, 583, 499-505. | 1.3 | 121 |
| 25 | Early modification of sickle cell disease clinical course by UDP-glucuronosyltransferase 1A1 gene promoter polymorphism. <i>Journal of Human Genetics</i> , 2008, 53, 524-528. | 1.1 | 12 |
| 26 | Proximity of the poly(A)-binding protein to a premature termination codon inhibits mammalian nonsense-mediated mRNA decay. <i>Rna</i> , 2008, 14, 563-576. | 1.6 | 132 |
| 27 | Hb Evora [Î²35 (B16), Ser->Pro], a novel hemoglobin variant associated with an Î²-thalassemia phenotype. <i>Haematologica</i> , 2007, 92, 252-253. | 1.7 | 9 |
| 28 | Epidemiology of haemoglobin disorders in Europe: an overview. <i>Scandinavian Journal of Clinical and Laboratory Investigation</i> , 2007, 67, 39-70. | 0.6 | 134 |
| 29 | Comment on "Nonsense-mediated mRNA decay modulates clinical outcome of genetic disease". <i>European Journal of Human Genetics</i> , 2007, 15, 533-534. | 1.4 | 4 |
| 30 | Mutational spectrum of delta-globin gene in the Portuguese population. <i>European Journal of Haematology</i> , 2007, 79, 422-428. | 1.1 | 27 |
| 31 | Human alpha2-globin nonsense-mediated mRNA decay induced by a novel alpha-thalassaemia frameshift mutation at codon 22. <i>British Journal of Haematology</i> , 2006, 133, 98-102. | 1.2 | 17 |
| 32 | Hemoglobin Loves Park [Î²68 (E12) Leu->Phe]: Report of five cases including one originating from a de novo mutation. <i>American Journal of Hematology</i> , 2006, 81, 256-261. | 2.0 | 2 |
| 33 | The canonical UPF1-dependent nonsense-mediated mRNA decay is inhibited in transcripts carrying a short open reading frame independent of sequence context. <i>Rna</i> , 2006, 12, 2160-2170. | 1.6 | 40 |
| 34 | HFE gene mutations are extremely rare in Western sub-Saharan Africa. <i>Annals of Hematology</i> , 2005, 84, 686-688. | 0.8 | 2 |
| 35 | Hb YaoundÃ© [Î²134(H12)Val->Ala] in Association with Hb C [Î²6(A3)Glu->Lys] in a Caucasian Portuguese Family. <i>Hemoglobin</i> , 2004, 28, 229-235. | 0.4 | 3 |
| 36 | Nonsense Mutations in Close Proximity to the Initiation Codon Fail to Trigger Full Nonsense-mediated mRNA Decay. <i>Journal of Biological Chemistry</i> , 2004, 279, 32170-32180. | 1.6 | 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. | 1.1 | 34 |
| 38 | COMPOUND HETEROZYGOSITY FOR Hb SPANISH TOWN [$\beta^{27}(B8)Glu\rightarrow Val$], Hb S [$\beta^{26}(A3)Glu\rightarrow Val$] AND THE $\alpha^{+}(3.7)Tj$ $\beta^{+}TQq0$ 0 0 | 0.4 | 2 |
| 39 | Asymptomatic homozygous deletional β^{20} -thalassemia in an African individual. American Journal of Hematology, 2002, 70, 232-236. | 2.0 | 3 |
| 40 | Molecular Basis of A-Thalassa in Portugal. Hemoglobin, 1995, 19, 343-352. | 0.4 | 20 |