

Agnieszka Fiszler

List of Publications by Year in descending order

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Version: 2024-02-01

22
papers

878
citations

623574

14
h-index

713332

21
g-index

24
all docs

24
docs citations

24
times ranked

1265
citing authors

#	ARTICLE	IF	CITATIONS
1	Triplet repeat RNA structure and its role as pathogenic agent and therapeutic target. <i>Nucleic Acids Research</i> , 2012, 40, 11-26.	6.5	182
2	Ribonuclease Dicer Cleaves Triplet Repeat Hairpins into Shorter Repeats that Silence Specific Targets. <i>Molecular Cell</i> , 2007, 25, 575-586.	4.5	176
3	The panorama of miRNA-mediated mechanisms in mammalian cells. <i>Cellular and Molecular Life Sciences</i> , 2014, 71, 2253-2270.	2.4	88
4	Inhibition of mutant huntingtin expression by RNA duplex targeting expanded CAG repeats. <i>Nucleic Acids Research</i> , 2011, 39, 5578-5585.	6.5	64
5	RNA toxicity in polyglutamine disorders: concepts, models, and progress of research. <i>Journal of Molecular Medicine</i> , 2013, 91, 683-691.	1.7	51
6	Oligonucleotide-based strategies to combat polyglutamine diseases. <i>Nucleic Acids Research</i> , 2014, 42, 6787-6810.	6.5	48
7	Mouse Ataxin-3 Functional Knock-Out Model. <i>NeuroMolecular Medicine</i> , 2011, 13, 54-65.	1.8	46
8	An evaluation of oligonucleotide-based therapeutic strategies for polyQ diseases. <i>BMC Molecular Biology</i> , 2012, 13, 6.	3.0	33
9	Self-duplexing CUG repeats selectively inhibit mutant huntingtin expression. <i>Nucleic Acids Research</i> , 2013, 41, 10426-10437.	6.5	30
10	Generation of New Isogenic Models of Huntingtonâ€™s Disease Using CRISPR-Cas9 Technology. <i>International Journal of Molecular Sciences</i> , 2020, 21, 1854.	1.8	25
11	What, When and How to Measureâ€”Peripheral Biomarkers in Therapy of Huntingtonâ€™s Disease. <i>International Journal of Molecular Sciences</i> , 2021, 22, 1561.	1.8	21
12	A potential role of extended simple sequence repeats in competing endogenous RNA crosstalk. <i>RNA Biology</i> , 2018, 15, 1399-1409.	1.5	20
13	Artificial miRNAs targeting CAG repeat expansion in ORFs cause rapid deadenylation and translation inhibition of mutant transcripts. <i>Cellular and Molecular Life Sciences</i> , 2021, 78, 1577-1596.	2.4	19
14	RAN Translation of the Expanded CAG Repeats in the SCA3 Disease Context. <i>Journal of Molecular Biology</i> , 2020, 432, 166699.	2.0	17
15	Reduction of Huntingtonâ€™s Disease RNA Foci by CAG Repeat-Targeting Reagents. <i>Frontiers in Cellular Neuroscience</i> , 2017, 11, 82.	1.8	15
16	Silencing of genes responsible for polyQ diseases using chemically modified single-stranded siRNAs. <i>Acta Biochimica Polonica</i> , 2017, 63, 759-764.	0.3	14
17	Mutant CAG Repeats Effectively Targeted by RNA Interference in SCA7 Cells. <i>Genes</i> , 2016, 7, 132.	1.0	13
18	Regulatory Potential of Competing Endogenous RNAs in Myotonic Dystrophies. <i>International Journal of Molecular Sciences</i> , 2021, 22, 6089.	1.8	6

#	ARTICLE	IF	CITATIONS
19	Generation of human iPS cell line IBCHi001-A from dentatorubralâ€“pallidoluysian atrophy patient's fibroblasts. Stem Cell Research, 2019, 39, 101512.	0.3	5
20	Implications of Poly(A) Tail Processing in Repeat Expansion Diseases. Cells, 2022, 11, 677.	1.8	4
21	Generation of human iPS cell line IBCHi002-A from spinocerebellar ataxia type 3/Machado-Joseph disease patient's fibroblasts. Stem Cell Research, 2020, 45, 101796.	0.3	1
22	103â€“...Mutant HTT mRNA as therapeutic target in allele-selective cag repeat-directed rnai approach and putative pathogenic agent in hd. , 2018, , .		0