

Junko Oshima

List of Publications by Year in descending order

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Version: 2024-02-01

9
papers

612
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1163117
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docs citations

9
times ranked

768
citing authors

#	ARTICLE	IF	CITATIONS
1	DNA methylation signatures in Blood DNA of Hutchinson-Gilford Progeria syndrome. <i>Aging Cell</i> , 2022, 21, e13555.	6.7	18
2	SMAD4 mutations and cross-talk between TGF- β ² /IFN γ ³ signaling accelerate rates of DNA damage and cellular senescence, resulting in a segmental progeroid syndrome—the Myhre syndrome. <i>GeroScience</i> , 2021, 43, 1481-1496.	4.6	9
3	<i>ERCC4</i> variants identified in a cohort of patients with segmental progeroid syndromes. <i>Human Mutation</i> , 2018, 39, 255-265.	2.5	23
4	<i>CTC1</i> mutations in a Brazilian family with progeroid features and recurrent bone fractures. <i>Molecular Genetics & Genomic Medicine</i> , 2018, 6, 1148-1156.	1.2	19
5	<i>WRN</i> Mutation Update: Mutation Spectrum, Patient Registries, and Translational Prospects. <i>Human Mutation</i> , 2017, 38, 7-15.	2.5	79
6	WRN mutations in Werner syndrome patients: genomic rearrangements, unusual intronic mutations and ethnic-specific alterations. <i>Human Genetics</i> , 2010, 128, 103-111.	3.8	87
7	Accelerated telomere shortening and replicative senescence in human fibroblasts overexpressing mutant and wild-type lamin A. <i>Experimental Cell Research</i> , 2008, 314, 82-91.	2.6	110
8	The spectrum of <i>WRN</i> mutations in Werner syndrome patients. <i>Human Mutation</i> , 2006, 27, 558-567.	2.5	198
9	Correction of cellular phenotypes of Hutchinson-Gilford Progeria cells by RNA interference. <i>Human Genetics</i> , 2005, 118, 444-450.	3.8	69