Junko Oshima

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/450462/publications.pdf

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9 papers	612 citations	1163117 8 h-index	1474206 9 g-index
9 all docs	9 docs citations	9 times ranked	768 citing authors

#	Article	IF	Citations
1	The spectrum ofWRNmutations in Werner syndrome patients. Human Mutation, 2006, 27, 558-567.	2.5	198
2	Accelerated telomere shortening and replicative senescence in human fibroblasts overexpressing mutant and wild-type lamin A. Experimental Cell Research, 2008, 314, 82-91.	2.6	110
3	WRN mutations in Werner syndrome patients: genomic rearrangements, unusual intronic mutations and ethnic-specific alterations. Human Genetics, 2010, 128, 103-111.	3.8	87
4	WRNMutation Update: Mutation Spectrum, Patient Registries, and Translational Prospects. Human Mutation, 2017, 38, 7-15.	2.5	79
5	Correction of cellular phenotypes of Hutchinson-Gilford Progeria cells by RNA interference. Human Genetics, 2005, 118, 444-450.	3.8	69
6	ERCC4 variants identified in a cohort of patients with segmental progeroid syndromes. Human Mutation, 2018, 39, 255-265.	2.5	23
7	<scp>CTC</scp>1 mutations in a Brazilian family with progeroid features and recurrent bone fractures. Molecular Genetics & Genomic Medicine, 2018, 6, 1148-1156.	1.2	19
8	DNA methylation signatures in Blood DNA of Hutchinsonâ€“Gilford Progeria syndrome. Aging Cell, 2022, 21, e13555.	6.7	18
9	SMAD4 mutations and cross-talk between TGF-Î²/IFNÎ³ signaling accelerate rates of DNA damage and cellular senescence, resulting in a segmental progeroid syndromeâ€”the Myhre syndrome. GeroScience, 2021, 43, 1481-1496.	4.6	9