

Ingrid van der Pluijm

List of Publications by Year in descending order

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Version: 2024-02-01

26
papers

1,063
citations

623734

14
h-index

552781

26
g-index

26
all docs

26
docs citations

26
times ranked

1842
citing authors

| # | ARTICLE | IF | CITATIONS |
|----|---|-----|-----------|
| 1 | Impaired Genome Maintenance Suppresses the Growth Hormone–Insulin-Like Growth Factor 1 Axis in Mice with Cockayne Syndrome. <i>PLoS Biology</i> , 2006, 5, e2. | 5.6 | 200 |
| 2 | Nucleotide Excision DNA Repair Is Associated With Age-Related Vascular Dysfunction. <i>Circulation</i> , 2012, 126, 468-478. | 1.6 | 153 |
| 3 | Priming of microglia in a DNA-repair deficient model of accelerated aging. <i>Neurobiology of Aging</i> , 2014, 35, 2147-2160. | 3.1 | 111 |
| 4 | The renin–angiotensin system and its involvement in vascular disease. <i>European Journal of Pharmacology</i> , 2015, 763, 3-14. | 3.5 | 94 |
| 5 | Age-related motor neuron degeneration in DNA repair-deficient <i>Ercc1</i> mice. <i>Acta Neuropathologica</i> , 2010, 120, 461-475. | 7.7 | 86 |
| 6 | Broad segmental progeroid changes in short-lived <i>Ercc1</i> ^{Δ7} mice. <i>Pathobiology of Aging & Age Related Diseases</i> , 2011, 1, 7219. | 1.1 | 79 |
| 7 | Brain Renin–Angiotensin System. <i>Hypertension</i> , 2017, 69, 1136-1144. | 2.7 | 69 |
| 8 | Cell-Autonomous Progeroid Changes in Conditional Mouse Models for Repair Endonuclease XPG Deficiency. <i>PLoS Genetics</i> , 2014, 10, e1004686. | 3.5 | 54 |
| 9 | Local endothelial DNA repair deficiency causes aging-resembling endothelial-specific dysfunction. <i>Clinical Science</i> , 2020, 134, 727-746. | 4.3 | 25 |
| 10 | Tissue-Specific Suppression of Thyroid Hormone Signaling in Various Mouse Models of Aging. <i>PLoS ONE</i> , 2016, 11, e0149941. | 2.5 | 23 |
| 11 | Transforming Growth Factor- β^2 and the Renin-Angiotensin System in Syndromic Thoracic Aortic Aneurysms: Implications for Treatment. <i>Cardiovascular Drugs and Therapy</i> , 2021, 35, 1233-1252. | 2.6 | 22 |
| 12 | Cockayne Syndrome Group B (Csb) and Group A (Csa) Deficiencies Predispose to Hearing Loss and Cochlear Hair Cell Degeneration in Mice. <i>Journal of Neuroscience</i> , 2015, 35, 4280-4286. | 3.6 | 19 |
| 13 | Extracellular Matrix Defects in Aneurysmal Fibulin-4 Mice Predispose to Lung Emphysema. <i>PLoS ONE</i> , 2014, 9, e106054. | 2.5 | 17 |
| 14 | Vascular Ageing Features Caused by Selective DNA Damage in Smooth Muscle Cell. <i>Oxidative Medicine and Cellular Longevity</i> , 2021, 2021, 1-13. | 4.0 | 15 |
| 15 | Dietary restriction but not angiotensin II type 1 receptor blockade improves DNA damage-related vasodilator dysfunction in rapidly aging <i>Ercc1</i> ^{Δ7} mice. <i>Clinical Science</i> , 2017, 131, 1941-1953. | 4.3 | 14 |
| 16 | Fibulin-4 deficiency differentially affects cytoskeleton structure and dynamics as well as TGF β^2 signaling. <i>Cellular Signalling</i> , 2019, 58, 65-78. | 3.6 | 14 |
| 17 | Accelerated loss of hearing and vision in the DNA-repair deficient <i>Ercc1</i> ^{Δ7} mouse. <i>Mechanisms of Ageing and Development</i> , 2012, 133, 59-67. | 4.6 | 13 |
| 18 | Chronic Sildenafil Treatment Improves Vasomotor Function in a Mouse Model of Accelerated Aging. <i>International Journal of Molecular Sciences</i> , 2020, 21, 4667. | 4.1 | 10 |

| # | ARTICLE | IF | CITATIONS |
|----|--|-----|-----------|
| 19 | Multi-Omics Profiling in Marfan Syndrome: Further Insights into the Molecular Mechanisms Involved in Aortic Disease. <i>International Journal of Molecular Sciences</i> , 2022, 23, 438. | 4.1 | 10 |
| 20 | Selective Phosphodiesterase 1 Inhibition Ameliorates Vascular Function, Reduces Inflammatory Response, and Lowers Blood Pressure in Aging Animals. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2021, 378, 173-183. | 2.5 | 9 |
| 21 | Molecular phenotyping and functional assessment of smooth muscle-like cells with pathogenic variants in aneurysm genes <i>ACTA2</i> , <i>MYH11</i> , <i>SMAD3</i> and <i>FBN1</i> . <i>Human Molecular Genetics</i> , 2021, 30, 2286-2299. | 2.9 | 7 |
| 22 | Slc2a10 knock-out mice deficient in ascorbic acid synthesis recapitulate aspects of arterial tortuosity syndrome and display mitochondrial respiration defects. <i>Human Molecular Genetics</i> , 2020, 29, 1476-1488. | 2.9 | 5 |
| 23 | Abnormal Aortic Wall Properties in Women with Turner Syndrome. <i>Aorta</i> , 2020, 08, 121-131. | 0.5 | 5 |
| 24 | Molecular Imaging of Aortic Aneurysm and Its Translational Power for Clinical Risk Assessment. <i>Frontiers in Medicine</i> , 2022, 9, 814123. | 2.6 | 4 |
| 25 | Pre-Operative Fasting Provides Long Term Protection Against Chronic Renal Damage Induced by Ischaemia Reperfusion Injury in Wild Type and Aneurysm Prone Fibulin-4 Mice. <i>European Journal of Vascular and Endovascular Surgery</i> , 2020, 60, 905-915. | 1.5 | 3 |
| 26 | In Vivo Renin Activity Imaging in the Kidney of Progeroid <i>Ercc1</i> Mutant Mice. <i>International Journal of Molecular Sciences</i> , 2021, 22, 12433. | 4.1 | 2 |