

Walter Rossmanith

List of Publications by Year in descending order

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59
papers

3,919
citations

147566

31
h-index

128067

60
g-index

64
all docs

64
docs citations

64
times ranked

3746
citing authors

#	ARTICLE	IF	CITATIONS
1	RNase P without RNA: Identification and Functional Reconstitution of the Human Mitochondrial tRNA Processing Enzyme. <i>Cell</i> , 2008, 135, 462-474.	13.5	546
2	The m1A landscape on cytosolic and mitochondrial mRNA at single-base resolution. <i>Nature</i> , 2017, 551, 251-255.	13.7	440
3	Deciphering the m6A Code via Antibody-Independent Quantitative Profiling. <i>Cell</i> , 2019, 178, 731-747.e16.	13.5	341
4	A subcomplex of human mitochondrial RNase P is a bifunctional methyltransferase that extensively moonlights in mitochondrial tRNA biogenesis. <i>Nucleic Acids Research</i> , 2012, 40, 11583-11593.	6.5	208
5	A single Arabidopsis organellar protein has RNase P activity. <i>Nature Structural and Molecular Biology</i> , 2010, 17, 740-744.	3.6	203
6	Human Mitochondrial tRNA Processing. <i>Journal of Biological Chemistry</i> , 1995, 270, 12885-12891.	1.6	135
7	Maternally Inherited Essential Hypertension Is Associated With the Novel 4263A>C Mutation in the Mitochondrial tRNA ^{Leu} Gene in a Large Han Chinese Family. <i>Circulation Research</i> , 2011, 108, 862-870.	2.0	114
8	Of P and Z: Mitochondrial tRNA processing enzymes. <i>Biochimica Et Biophysica Acta - Gene Regulatory Mechanisms</i> , 2012, 1819, 1017-1026.	0.9	102
9	The coffee components kahweol and cafestol induce γ -glutamylcysteine synthetase, the rate limiting enzyme of chemoprotective glutathione synthesis, in several organs of the rat. <i>Archives of Toxicology</i> , 2002, 75, 685-694.	1.9	90
10	Biology of transforming growth factor β in hepatocarcinogenesis. <i>Microscopy Research and Technique</i> , 2001, 52, 430-436.	1.2	78
11	Localization of Human RNase Z Isoforms: Dual Nuclear/Mitochondrial Targeting of the ELAC2 Gene Product by Alternative Translation Initiation. <i>PLoS ONE</i> , 2011, 6, e19152.	1.1	75
12	Characterization of Human Mitochondrial RNase P: Novel Aspects in tRNA Processing. <i>Biochemical and Biophysical Research Communications</i> , 1998, 247, 234-241.	1.0	74
13	Localization and quantification of Cd- and Cu-specific metallothionein isoform mRNA in cells and organs of the terrestrial gastropod <i>Helix pomatia</i> . <i>Toxicology and Applied Pharmacology</i> , 2003, 190, 25-36.	1.3	72
14	Nuclear RNase P of <i>Trypanosoma brucei</i> : A Single Protein in Place of the Multicomponent RNA-Protein Complex. <i>Cell Reports</i> , 2012, 2, 19-25.	2.9	71
15	Overexpression of Activin β C or Activin β E in the Mouse Liver Inhibits Regenerative Deoxyribonucleic Acid Synthesis of Hepatic Cells. <i>Endocrinology</i> , 2003, 144, 3497-3504.	1.4	69
16	Impairment of tRNA processing by point mutations in mitochondrial tRNA ^{Leu} (UUR) associated with mitochondrial diseases. <i>FEBS Letters</i> , 1998, 433, 269-274.	1.3	68
17	Effects of coffee and its chemopreventive components kahweol and cafestol on cytochrome P450 and sulfotransferase in rat liver. <i>Food and Chemical Toxicology</i> , 2008, 46, 1230-1238.	1.8	63
18	Expression and dimerization of the rat activin subunits β C and β E: evidence for the formation of novel activin dimers. <i>Journal of Molecular Endocrinology</i> , 2002, 28, 137-148.	1.1	58

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19	Levels of transforming growth factor β^2 and transforming growth factor β^2 receptors in rat liver during growth, regression by apoptosis and neoplasia. <i>Hepatology</i> , 1998, 28, 717-726.	3.6	56
20	Distribution of Ribonucleoprotein and Protein-Only RNase P in Eukarya. <i>Molecular Biology and Evolution</i> , 2015, 32, msv187.	3.5	56
21	Molecular insights into HSD10 disease: impact of SDR5C1 mutations on the human mitochondrial RNase P complex. <i>Nucleic Acids Research</i> , 2015, 43, 5112-5119.	6.5	55
22	Expression of activins C and E induces apoptosis in human and rat hepatoma cells. <i>Carcinogenesis</i> , 2003, 24, 1801-1809.	1.3	53
23	Minimal and RNA-free RNase P in <i>Aquifex aeolicus</i> . <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2017, 114, 11121-11126.	3.3	46
24	A novel <i>HSD17B10</i> mutation impairing the activities of the mitochondrial RNase P complex causes X-linked intractable epilepsy and neurodevelopmental regression. <i>RNA Biology</i> , 2016, 13, 477-485.	1.5	42
25	Cellular Gene Dose and Kinetics of Gene Expression in Mouse Livers Transfected by High-Volume Tail-Vein Injection of Naked DNA. <i>DNA and Cell Biology</i> , 2002, 21, 847-853.	0.9	40
26	Follistatin overexpression in rodent liver tumors: A possible mechanism to overcome activin growth control. <i>Molecular Carcinogenesis</i> , 2002, 35, 1-5.	1.3	38
27	Functional characterization of the human tRNA methyltransferases TRMT10A and TRMT10B. <i>Nucleic Acids Research</i> , 2020, 48, 6157-6169.	6.5	38
28	RNase MRP and RNase P share a common substrate. <i>Nucleic Acids Research</i> , 1993, 21, 3239-3243.	6.5	35
29	Expression of cytochrome P450 2A5 in preneoplastic and neoplastic mouse liver lesions. <i>Molecular Carcinogenesis</i> , 1998, 22, 229-234.	1.3	35
30	Substrate recognition and cleavage-site selection by a single-subunit protein-only RNase P. <i>Nucleic Acids Research</i> , 2016, 44, 2323-2336.	6.5	35
31	Accumulation of very long-chain fatty acids does not affect mitochondrial function in adrenoleukodystrophy protein deficiency. <i>Human Molecular Genetics</i> , 2005, 14, 1127-1137.	1.4	32
32	tRNA Processing by Protein-Only versus RNA-Based RNase P: Kinetic Analysis Reveals Mechanistic Differences. <i>ChemBioChem</i> , 2012, 13, 2270-2276.	1.3	31
33	Processing of human mitochondrial tRNAGCUGSer(AGY): a novel pathway in tRNA biosynthesis. <i>Journal of Molecular Biology</i> , 1997, 265, 365-371.	2.0	30
34	A novel mechanism for mitogenic signaling via pro-transforming growth factor β^2 within hepatocyte nuclei. <i>Hepatology</i> , 2002, 35, 1372-1380.	3.6	29
35	Isolated cytochrome c oxidase deficiency as a cause of MELAS. <i>Journal of Medical Genetics</i> , 2007, 45, 117-121.	1.5	26
36	Bi-allelic variants in the mitochondrial RNase P subunit PRORP cause mitochondrial tRNA processing defects and pleiotropic multisystem presentations. <i>American Journal of Human Genetics</i> , 2021, 108, 2195-2204.	2.6	26

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37	Difference between Mitochondrial RNase P and Nuclear RNase P. <i>Molecular and Cellular Biology</i> , 2001, 21, 8236-8237.	1.1	25
38	CPEO associated with a single nucleotide deletion in the mitochondrial tRNA ^{Tyr} gene. <i>Neurology</i> , 2001, 57, 2298-2301.	1.5	25
39	Playing RNase P Evolution: Swapping the RNA Catalyst for a Protein Reveals Functional Uniformity of Highly Divergent Enzyme Forms. <i>PLoS Genetics</i> , 2014, 10, e1004506.	1.5	24
40	YBEY is an essential biogenesis factor for mitochondrial ribosomes. <i>Nucleic Acids Research</i> , 2020, 48, 9762-9786.	6.5	24
41	The expanding mutational spectrum of MERRF substitution G8361A in the mitochondrial tRNA ^{Lys} gene. <i>Annals of Neurology</i> , 2003, 54, 820-823.	2.8	23
42	tRNA recognition, processing, and disease: Hypotheses around an unorthodox type of RNase P in human mitochondria. <i>Mitochondrion</i> , 2009, 9, 284-288.	1.6	21
43	Protein-only RNase P function in <i>Escherichia coli</i> : viability, processing defects and differences between PRORP isoenzymes. <i>Nucleic Acids Research</i> , 2017, 45, 7441-7454.	6.5	21
44	Definition of the Th/ To ribonucleoprotein by RNase P and RNase MRP. <i>Molecular Biology Reports</i> , 1993, 18, 29-35.	1.0	20
45	Processing mitochondrial (t)RNAs: New enzyme, old job. <i>Cell Cycle</i> , 2009, 8, 1650-1653.	1.3	20
46	Apoptosis and Hepatocarcinogenesis. <i>Digestion</i> , 1998, 59, 64-65.	1.2	18
47	Positioning Europe for the EPITRANSCRIPTOMICS challenge. <i>RNA Biology</i> , 2018, 15, 1-3.	1.5	18
48	Mitochondrial poly(A) polymerase is involved in tRNA repair. <i>Nucleic Acids Research</i> , 2015, 43, gkv891.	6.5	17
49	Estrogen Receptor-Alpha and Estrogen Receptor-Beta Are Present in the Human Growth Plate in Childhood and Adolescence, in Identical Distribution. <i>Hormone Research in Paediatrics</i> , 2002, 58, 99-103.	0.8	16
50	Mitochondrial genotype and risk for Alzheimer's disease: cross-sectional data from the Vienna-Transdanube-Aging ?VITA? study. <i>Journal of Neural Transmission</i> , 2004, 111, 1155-65.	1.4	13
51	Diversity and Evolution of RNase P. , 2020, , 255-299.		13
52	RNase Mitochondrial RNA Processing Cleaves RNA from the Rat Mitochondrial Displacement Loop at the Origin of Heavy-Strand DNA Replication. <i>FEBS Journal</i> , 1995, 227, 657-662.	0.2	11
53	Expression of mouse RNase MRP RNA in human embryonic kidney 293 cells. <i>Molecular Biology Reports</i> , 1997, 24, 221-230.	1.0	10
54	Repairing tRNA termini: News from the 3' end. <i>RNA Biology</i> , 2016, 13, 1182-1188.	1.5	10

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55	Analysis of the Cleavage Mechanism by Protein-Only RNase P Using Precursor tRNA Substrates with Modifications at the Cleavage Site. <i>Journal of Molecular Biology</i> , 2016, 428, 4917-4928.	2.0	9
56	The Amyloid- β -SDR5C1 (ABAD) Interaction Does Not Mediate a Specific Inhibition of Mitochondrial RNase P. <i>PLoS ONE</i> , 2013, 8, e65609.	1.1	9
57	RNase Mitochondrial RNA Processing Cleaves RNA from the Rat Mitochondrial Displacement Loop at the Origin of Heavy-Strand DNA Replication. <i>FEBS Journal</i> , 1995, 227, 657-662.	0.2	7
58	Hepatocarcinogenesis in the Context of Strain Differences in Energy Metabolism Between Inbred Strains of Mice (C57BL/6J and C3H/He). <i>Advances in Experimental Medicine and Biology</i> , 2001, 500, 607-611.	0.8	4
59	Isolated cytochrome c oxidase deficiency as a cause of MELAS. <i>BMJ Case Reports</i> , 2009, 2009, bcr0820080666-bcr0820080666.	0.2	4