Tadashi Yamamoto

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Impaired proliferation of peripheral B cells and indication of autoimmune disease in lyn-deficient mice. Immunity, 1995, 3, 549-560.	14.3	454
2	miRNA-mediated deadenylation is orchestrated by GW182 through two conserved motifs that interact with CCR4–NOT. Nature Structural and Molecular Biology, 2011, 18, 1211-1217.	8.2	286
3	Oligo-astheno-teratozoospermia in mice lacking Cnot7, a regulator of retinoid X receptor beta. Nature Genetics, 2004, 36, 528-533.	21.4	127
4	Depletion of Mammalian CCR4b Deadenylase Triggers Elevation of the <i>p27</i> ^{<i>Kip1</i>} mRNA Level and Impairs Cell Growth. Molecular and Cellular Biology, 2007, 27, 4980-4990.	2.3	98
5	Multifunctional roles of the mammalian CCR4ââ,¬â€œNOT complex in physiological phenomena. Frontiers in Genetics, 2014, 5, 286.	2.3	95
6	Lyn Kinase Suppresses the Transcriptional Activity of IRF5 in the TLR-MyD88 Pathway to Restrain the Development of Autoimmunity. Immunity, 2016, 45, 319-332.	14.3	81
7	Obesity resistance and increased hepatic expression of catabolism-related mRNAs in <i>Cnot3</i> ^{+/â~`} mice. EMBO Journal, 2011, 30, 4678-4691.	7.8	71
8	CNOT2 depletion disrupts and inhibits the CCR4-NOT deadenylase complex and induces apoptotic cell death. Genes To Cells, 2011, 16, 368-379.	1.2	69
9	The role of the CNOT1 subunit of the CCR4-NOT complex in mRNA deadenylation and cell viability. Protein and Cell, 2011, 2, 755-763.	11.0	63
10	The CCR4-NOT deadenylase complex controls Atg7-dependent cell death and heart function. Science Signaling, 2018, 11, .	3.6	51
11	Post-transcriptional Stabilization of Ucp1 mRNA Protects Mice from Diet-Induced Obesity. Cell Reports, 2015, 13, 2756-2767.	6.4	46
12	CNOT3 contributes to early B cell development by controlling <i>lgh</i> rearrangement and <i>p53</i> mRNA stability. Journal of Experimental Medicine, 2015, 212, 1465-1479.	8.5	43
13	Distinct expression patterns of the subunits of the CCR4–NOT deadenylase complex during neural development. Biochemical and Biophysical Research Communications, 2011, 411, 360-364.	2.1	37
14	CNOT3 suppression promotes necroptosis by stabilizing mRNAs for cell death-inducing proteins. Scientific Reports, 2015, 5, 14779.	3.3	37
15	Cnot7-Null Mice Exhibit High Bone Mass Phenotype and Modulation of BMP Actions. Journal of Bone and Mineral Research, 2007, 22, 1217-1223.	2.8	31
16	Stability of mRNA influences osteoporotic bone mass via CNOT3. Proceedings of the National Academy of Sciences of the United States of America, 2014, 111, 2692-2697.	7.1	29
17	Fluid flow-induced left-right asymmetric decay of Dand5 mRNA in the mouse embryo requires a Bicc1-Ccr4 RNA degradation complex. Nature Communications, 2021, 12, 4071.	12.8	28
18	Tob2 Inhibits Peroxisome Proliferator-Activated Receptor γ2 Expression by Sequestering Smads and C/EBP <i>α</i> during Adipocyte Differentiation. Molecular and Cellular Biology, 2012, 32, 5067-5077.	2.3	27

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19	Essential functions of the CNOT7/8 catalytic subunits of the CCR4-NOT complex in mRNA regulation and cell viability. RNA Biology, 2020, 17, 403-416.	3.1	27
20	Genetic and chemical inhibition of IRF5 suppresses pre-existing mouse lupus-like disease. Nature Communications, 2021, 12, 4379.	12.8	24
21	Adipocyteâ€specific disruption of mouse <i>Cnot3</i> causes lipodystrophy. FEBS Letters, 2017, 591, 358-368.	2.8	20
22	Postnatal liver functional maturation requires Cnot complex-mediated decay of mRNAs encoding cell cycle and immature liver genes. Development (Cambridge), 2019, 146, .	2.5	18
23	The CCR4–NOT complex maintains liver homeostasis through mRNA deadenylation. Life Science Alliance, 2020, 3, e201900494.	2.8	17
24	Involvement of CNOT3 in mitotic progression through inhibition of MAD1 expression. Biochemical and Biophysical Research Communications, 2012, 419, 268-273.	2.1	15
25	Loss of β-cell identity and diabetic phenotype in mice caused by disruption of CNOT3-dependent mRNA deadenylation. Communications Biology, 2020, 3, 476.	4.4	13
26	The CCR4–NOT Deadenylase Complex Maintains Adipocyte Identity. International Journal of Molecular Sciences, 2019, 20, 5274.	4.1	11
27	The CCR4–NOT deadenylase complex safeguards thymic positive selection by down-regulating aberrant pro-apoptotic gene expression. Nature Communications, 2020, 11, 6169.	12.8	11
28	The mitochondrial protein PGAM5 suppresses energy consumption in brown adipocytes by repressing expression of uncoupling protein 1. Journal of Biological Chemistry, 2020, 295, 5588-5601.	3.4	9
29	ARE-binding protein ZFP36L1 interacts with CNOT1 to directly repress translation via a deadenylation-independent mechanism. Biochimie, 2020, 174, 49-56.	2.6	9
30	RNA decay machinery safeguards immune cell development and immunological responses. Trends in Immunology, 2021, 42, 447-460.	6.8	6
31	Regulation of Early Lymphocyte Development via mRNA Decay Catalyzed by the CCR4-NOT Complex. Frontiers in Immunology, 2021, 12, 715675.	4.8	5
32	Neuronal XRN1 is required for maintenance of whole-body metabolic homeostasis. IScience, 2021, 24, 103151.	4.1	5
33	CNOT7 Outcompetes Its Paralog CNOT8 for Integration into The CCR4-NOT Complex. Journal of Molecular Biology, 2022, 434, 167523.	4.2	5
34	States of decay: The systems biology of mRNA stability. Current Opinion in Systems Biology, 2019, 15, 48-57.	2.6	4
35	CNOT3 suppression promotes necroptosis by stabilizing mRNAs for cell death-inducing proteins. , 0, .		4
36	Regulation of Fetal Genes by Transitions among RNA-Binding Proteins during Liver Development. International Journal of Molecular Sciences, 2020, 21, 9319.	4.1	3

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37	CNOT1 regulates circadian behaviour through <i>Per2</i> mRNA decay in a deadenylation-dependent manner. RNA Biology, 2022, 19, 703-718.	3.1	1
38	Insufficient liver maturation affects murine early postnatal hair cycle. Biochemical and Biophysical Research Communications, 2020, 521, 172-177.	2.1	0
39	Regulation of CCR4-NOT complex deadenylase activity and cellular responses by MK2-dependent phosphorylation of CNOT2. RNA Biology, 2022, 19, 234-246.	3.1	0