

# Tarmo Annilo

## List of Publications by Year in descending order

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Version: 2024-02-01

34  
papers

2,621  
citations

279798

23  
h-index

377865

34  
g-index

36  
all docs

36  
docs citations

36  
times ranked

3998  
citing authors

#	ARTICLE	IF	CITATIONS
1	<i>ABCA3</i> Gene Mutations in Newborns with Fatal Surfactant Deficiency. <i>New England Journal of Medicine</i> , 2004, 350, 1296-1303.	27.0	621
2	EVOLUTION OF THE ATP-BINDING CASSETTE (ABC) TRANSPORTER SUPERFAMILY IN VERTEBRATES. <i>Annual Review of Genomics and Human Genetics</i> , 2005, 6, 123-142.	6.2	540
3	Meta-analysis of microRNA expression in lung cancer. <i>International Journal of Cancer</i> , 2013, 132, 2884-2893.	5.1	195
4	Evolution of the vertebrate ABC gene family: Analysis of gene birth and death. <i>Genomics</i> , 2006, 88, 1-11.	2.9	150
5	Identification of miR-374a as a prognostic marker for survival in patients with early-stage non-small cell lung cancer. <i>Genes Chromosomes and Cancer</i> , 2011, 50, 812-822.	2.8	116
6	MicroRNAs miR-124 and miR-135a are potential regulators of the mineralocorticoid receptor gene (NR3C2) expression. <i>Biochemical and Biophysical Research Communications</i> , 2010, 391, 727-732.	2.1	108
7	Identification and characterization of a novel ABCA subfamily member, ABCA12, located in the lamellar ichthyosis region on 2q34. <i>Cytogenetic and Genome Research</i> , 2002, 98, 169-176.	1.1	67
8	Natural antisense transcript of natriuretic peptide precursor A (NPPA): structural organization and modulation of NPPA expression. <i>BMC Molecular Biology</i> , 2009, 10, 81.	3.0	64
9	Methylation Markers of Early-Stage Non-Small Cell Lung Cancer. <i>PLoS ONE</i> , 2012, 7, e39813.	2.5	62
10	<i>De Novo</i> <i>SCN8A</i> Mutation Identified by Whole-Exome Sequencing in a Boy With Neonatal Epileptic Encephalopathy, Multiple Congenital Anomalies, and Movement Disorders. <i>Journal of Child Neurology</i> , 2014, 29, NP202-NP206.	1.4	59
11	Comparative genome analysis of potential regulatory elements in the ABCG5-ABCG8 gene cluster. <i>Biochemical and Biophysical Research Communications</i> , 2002, 295, 276-282.	2.1	52
12	A Missense Mutation in <i>DUSP6</i> is Associated with Class III Malocclusion. <i>Journal of Dental Research</i> , 2013, 92, 893-898.	5.2	51
13	Evolutionary analysis of a cluster of ATP-binding cassette (ABC) genes. <i>Mammalian Genome</i> , 2003, 14, 7-20.	2.2	44
14	C-reactive protein upregulates the whole blood expression of CD59 - an integrative analysis. <i>PLoS Computational Biology</i> , 2017, 13, e1005766.	3.2	44
15	Human and mouse orthologs of a new ATP-binding cassette gene, ABCG4. <i>Cytogenetic and Genome Research</i> , 2001, 94, 196-201.	1.1	42
16	The human ATP binding cassette gene ABCA13, located on chromosome 7p12.3, encodes a 5058 amino acid protein with an extracellular domain encoded in part by a 4.8-kb conserved exon. <i>Cytogenetic and Genome Research</i> , 2002, 98, 160-168.	1.1	39
17	Degeneration of an ATP-binding cassette transporter gene, ABCC13, in different mammalian lineages. <i>Genomics</i> , 2004, 84, 34-46.	2.9	36
18	Comprehensive Meta-analysis of MicroRNA Expression Using a Robust Rank Aggregation Approach. <i>Methods in Molecular Biology</i> , 2014, 1182, 361-373.	0.9	36

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19	The dMRP/CG6214 gene of <i>Drosophila</i> is evolutionarily and functionally related to the human multidrug resistance-associated protein family. <i>Insect Molecular Biology</i> , 2004, 13, 539-548.	2.0	35
20	Nuclear Import and Nucleolar Accumulation of the Human Ribosomal Protein S7 Depends on both a Minimal Nuclear Localization Sequence and an Adjacent Basic Region. <i>Biochemical and Biophysical Research Communications</i> , 1998, 249, 759-766.	2.1	34
21	miR-10a-5p is increased in atopic dermatitis and has capacity to inhibit keratinocyte proliferation. <i>Allergy: European Journal of Allergy and Clinical Immunology</i> , 2019, 74, 2146-2156.	5.7	31
22	Non-syndromic Tooth Agenesis Associated with a Nonsense Mutation in Ectodysplasin-A ( <i>EDA</i> ). <i>Journal of Dental Research</i> , 2013, 92, 507-511.	5.2	29
23	Altered Gene Expression Associated with microRNA Binding Site Polymorphisms. <i>PLoS ONE</i> , 2015, 10, e0141351.	2.5	29
24	Identifying and characterizing a five-gene cluster of ATP-binding cassette transporters mapping to human chromosome 17q24: a new subgroup within the ABCA subfamily. <i>GeneScreen</i> , 2001, 1, 157-164.	0.6	20
25	Mutation analysis and copy number alterations of <i>KIF23</i> in non-small-cell lung cancer exhibiting <i>KIF23</i> over-expression. <i>OncoTargets and Therapy</i> , 2017, Volume 10, 4969-4979.	2.0	20
26	Three ATP-binding cassette transporter genes, <i>Abca14</i> , <i>Abca15</i> , and <i>Abca16</i> , form a cluster on mouse Chromosome 7F3. <i>Mammalian Genome</i> , 2004, 15, 335-343.	2.2	17
27	The human ribosomal protein S7-encoding gene: isolation, structure and localization in 2p25. <i>Gene</i> , 1995, 165, 297-302.	2.2	16
28	Different course of lung disease in two siblings with novel <i>ABCA3</i> mutations. <i>European Journal of Pediatrics</i> , 2014, 173, 1553-1556.	2.7	16
29	Comparison of Peptide- and Lipid-Based Delivery of miR-34a-5p Mimic into PPC-1 Cells. <i>Nucleic Acid Therapeutics</i> , 2017, 27, 295-302.	3.6	13
30	A human-specific VNTR in the <i>TRIB3</i> promoter causes gene expression variation between individuals. <i>PLoS Genetics</i> , 2020, 16, e1008981.	3.5	13
31	Metagenes Associated with Survival in Non-Small Cell Lung Cancer. <i>Cancer Informatics</i> , 2011, 10, CIN.S7135.	1.9	9
32	De novo exonic mutation in <i>MYH7</i> gene leading to exon skipping in a patient with early onset muscular weakness and fiber-type disproportion. <i>Neuromuscular Disorders</i> , 2016, 26, 236-239.	0.6	8
33	Isolation and characterization of the mouse ribosomal protein S7 gene. <i>IUBMB Life</i> , 1998, 46, 287-295.	3.4	3
34	Genome-wide analysis of nuclear magnetic resonance metabolites revealed parent-of-origin effect on triglycerides in medium very low-density lipoprotein in <i>PTPRD</i> gene. <i>Biomarkers in Medicine</i> , 2018, 12, 439-446.	1.4	2