Martina RinÄiÄ

List of Publications by Year
in descending order

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33 papers	294 citations	1039406 9 h-index	940134 16 g-index
33 all docs	33 docs citations	33 times ranked	442 citing authors

#	Article	IF	Citations
1	Somatic Mosaicism in Cases with Small Supernumerary Marker Chromosomes. Current Genomics, 2010, 11, 432-439.	0.7	44
2	Aporrectodea caliginosa, a suitable earthworm species for field based genotoxicity assessment?. Environmental Pollution, 2011, 159, 841-849.	3.7	31
3	Heteromorphic variants of chromosome 9. Molecular Cytogenetics, 2013, 6, 14.	0.4	31
4	Comprehensive chronic lymphocytic leukemia diagnostics by combined multiplex ligation dependent probe amplification (MLPA) and interphase fluorescence in situ hybridization (iFISH). Molecular Cytogenetics, 2014, 7, 79.	0.4	27
5	High rates of submicroscopic aberrations in karyotypically normal acute lymphoblastic leukemia. Molecular Cytogenetics, 2015, 8, 45.	0.4	17
6	European registration process for Clinical Laboratory Geneticists in genetic healthcare. European Journal of Human Genetics, 2017, 25, 515-519.	1.4	13
7	Regarding the rights and duties of Clinical Laboratory Geneticists in genetic healthcare systems; results of a survey in over 50 countries. European Journal of Human Genetics, 2019, 27, 1168-1174.	1.4	12
8	Complex intrachromosomal rearrangement in 1q leading to 1q32.2 microdeletion: a potential role of SRGAP2 in the gyrification of cerebral cortex. Molecular Cytogenetics, 2016, 9, 19.	0.4	11
9	Association of new deletion/duplication region at chromosome 1p21 with intellectual disability, severe speech deficit and autism spectrum disorder-like behavior: an all-in approach to solving the DPYD enigma. Translational Neuroscience, 2015, 6, 59-86.	0.7	9
10	Thoughts about SLC16A2, TSIX and XIST gene like sites in the human genome and a potential role in cellular chromosome counting. Molecular Cytogenetics, 2016, 9, 56.	0.4	9
11	Deletion 2q31.2â€q31.3 in a 4â€yearâ€old girl with microcephaly and severe mental retardation. American Journal of Medical Genetics, Part A, 2011, 155, 1476-1482.	0.7	8
12	Molecular Cytogenomic Characterization of the Murine Breast Cancer Cell Lines C-127I, EMT6/P and TA3 Hauschka. International Journal of Molecular Sciences, 2020, 21, 4716.	1.8	8
13	Presence of harmless small supernumerary marker chromosomes hampers molecular genetic diagnosis: a case report. Molecular Medicine Reports, 2010, 3, 571-4.	1.1	7
14	A Novel Cryptic Three-Way Translocation t(2;9;18)(p23.2;p21.3;q21.33) with Deletion of Tumor Suppressor Genes in 9p21.3 and 13q14 in a T-Cell Acute Lymphoblastic Leukemia. Leukemia Research and Treatment, 2014, 2014, 1-7.	2.0	7
15	Novel Cryptic Rearrangements in Adult B-Cell Precursor Acute Lymphoblastic Leukemia Involving the MLL Gene. Journal of Histochemistry and Cytochemistry, 2015, 63, 384-390.	1.3	7
16	Genotype-phenotype Correlation of Î²-Thalassemia in Croatian Patients: A Specific HBB Gene Mutations. Journal of Pediatric Hematology/Oncology, 2018, 40, e77-e82.	0.3	7
17	First Molecular Cytogenetic Characterization of Murine Malignant Mesothelioma Cell Line AE17 and In Silico Translation to the Human Genome. Current Bioinformatics, 2017, 12, 11-18.	0.7	7
18	Molecular Cytogenetic Characterization Identified the Murine B-Cell Lymphoma Cell Line A-20 as a Model for Sporadic Burkittâ€™s Lymphoma. Journal of Histochemistry and Cytochemistry, 2017, 65, 669-677.	1.3	5

#	Article	IF	Citations
19	Molecular Cytogenetic Characterization of Two Murine Colorectal Cancer Cell Lines. OBM Genetics, 2018, 2, 1-1.	0.2	5
20	Is There a Yet Unreported Unbalanced Chromosomal Abnormality without Phenotypic Consequences in Proximal 4p?. Cytogenetic and Genome Research, 2011, 132, 121-123.	0.6	4
21	MLLT10 and IL3 rearrangement together with a complex four-way translocation and trisomy 4 in a patient with early T-cell precursor acute lymphoblastic leukemia: A case report. Oncology Reports, 2015, 33, 625-630.	1.2	4
22	A novel IGH@ gene rearrangement associated with CDKN2A/B deletion in young adult B-cell acute lymphoblastic leukemia. Oncology Letters, 2016, 11, 2117-2122.	0.8	4
23	Cytogenomic characterization of three murine malignant mesothelioma tumor cell lines. Molecular Cytogenetics, 2020, 13, 43.	0.4	4
24	Identification of metastasis-related genes by genomic and transcriptomic studies in murine melanoma. Life Sciences, 2021, 267, 118922.	2.0	4
25	Molecular cytogenetic characterization of two murine cancer cell lines derived from salivary gland. Biological Communications, 2018, 63, 243-255.	0.4	4
26	7p21.3 Together With a 12p13.32 Deletion in a Patient With Microcephalyâ€”Does 12p13.32 Locus Possibly Comprises a Candidate Gene Region for Microcephaly?. Frontiers in Molecular Neuroscience, 2021, 14, 613091.	1.4	2
27	Isochromosome 17q in Chronic Lymphocytic Leukemia. Leukemia Research and Treatment, 2015, 2015, 1-6.	2.0	1
28	Cytogenomic characteristics of murine breast cancer cell line JC. Molecular Cytogenetics, 2021, 14, 7.	0.4	1
29	Trisomy 21 with a Small Supernumerary Marker Chromosome Derived from Chromosomes 13/21 and 18. Balkan Journal of Medical Genetics, 2010, 13, 55-58.	0.5	1
30	First molecular cytogenetic characterisation of tracheal squamous cell carcinoma cell line KLN 205. , 0, , .		0
31	Molecular Cytogenetic Characterization of the Murine Melanoma Cell Lines S91 Clone M3 and B16-F1 with Variant B16-4A5. Cytogenetic and Genome Research, 2021, 161, 82-92.	0.6	0
32	Cytogenomics of murine melanoma cell lines C57/B1 and B16-F0. Molecular and Experimental Biology in Medicine, 2020, 3, 39-44.	0.1	0
33	Molecular cytogenetic characterization of the urethane-induced murine lung cell line LA-4 as a model for human squamous cell lung cancer. Molecular and Clinical Oncology, 2021, 16, 9.	0.4	0