

Martina RinÄiÄ

List of Publications by Year in descending order

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33
papers

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times ranked

442
citing authors

| # | ARTICLE | IF | CITATIONS |
|----|---|-----|-----------|
| 1 | Cytogenomic characteristics of murine breast cancer cell line JC. <i>Molecular Cytogenetics</i> , 2021, 14, 7. | 0.4 | 1 |
| 2 | Identification of metastasis-related genes by genomic and transcriptomic studies in murine melanoma. <i>Life Sciences</i> , 2021, 267, 118922. | 2.0 | 4 |
| 3 | 7p21.3 Together With a 12p13.32 Deletion in a Patient With Microcephaly "Does 12p13.32 Locus Possibly Comprises a Candidate Gene Region for Microcephaly?. <i>Frontiers in Molecular Neuroscience</i> , 2021, 14, 613091. | 1.4 | 2 |
| 4 | Molecular Cytogenetic Characterization of the Murine Melanoma Cell Lines S91 Clone M3 and B16-F1 with Variant B16-4A5. <i>Cytogenetic and Genome Research</i> , 2021, 161, 82-92. | 0.6 | 0 |
| 5 | Molecular cytogenetic characterization of the urethane-induced murine lung cell line LA-4 as a model for human squamous cell lung cancer. <i>Molecular and Clinical Oncology</i> , 2021, 16, 9. | 0.4 | 0 |
| 6 | Molecular Cytogenomic Characterization of the Murine Breast Cancer Cell Lines C-1271, EMT6/P and TA3 Hauschka. <i>International Journal of Molecular Sciences</i> , 2020, 21, 4716. | 1.8 | 8 |
| 7 | Cytogenomic characterization of three murine malignant mesothelioma tumor cell lines. <i>Molecular Cytogenetics</i> , 2020, 13, 43. | 0.4 | 4 |
| 8 | Cytogenomics of murine melanoma cell lines C57/B1 and B16-F0. <i>Molecular and Experimental Biology in Medicine</i> , 2020, 3, 39-44. | 0.1 | 0 |
| 9 | Regarding the rights and duties of Clinical Laboratory Geneticists in genetic healthcare systems; results of a survey in over 50 countries. <i>European Journal of Human Genetics</i> , 2019, 27, 1168-1174. | 1.4 | 12 |
| 10 | Genotype-phenotype Correlation of β^2 -Thalassemia in Croatian Patients: A Specific HBB Gene Mutations. <i>Journal of Pediatric Hematology/Oncology</i> , 2018, 40, e77-e82. | 0.3 | 7 |
| 11 | Molecular cytogenetic characterization of two murine cancer cell lines derived from salivary gland. <i>Biological Communications</i> , 2018, 63, 243-255. | 0.4 | 4 |
| 12 | Molecular Cytogenetic Characterization of Two Murine Colorectal Cancer Cell Lines. <i>OBM Genetics</i> , 2018, 2, 1-1. | 0.2 | 5 |
| 13 | European registration process for Clinical Laboratory Geneticists in genetic healthcare. <i>European Journal of Human Genetics</i> , 2017, 25, 515-519. | 1.4 | 13 |
| 14 | Molecular Cytogenetic Characterization Identified the Murine B-Cell Lymphoma Cell Line A-20 as a Model for Sporadic Burkitt's Lymphoma. <i>Journal of Histochemistry and Cytochemistry</i> , 2017, 65, 669-677. | 1.3 | 5 |
| 15 | First Molecular Cytogenetic Characterization of Murine Malignant Mesothelioma Cell Line AE17 and In Silico Translation to the Human Genome. <i>Current Bioinformatics</i> , 2017, 12, 11-18. | 0.7 | 7 |
| 16 | A novel IGH@ gene rearrangement associated with CDKN2A/B deletion in young adult B-cell acute lymphoblastic leukemia. <i>Oncology Letters</i> , 2016, 11, 2117-2122. | 0.8 | 4 |
| 17 | Thoughts about SLC16A2, TSIX and XIST gene like sites in the human genome and a potential role in cellular chromosome counting. <i>Molecular Cytogenetics</i> , 2016, 9, 56. | 0.4 | 9 |
| 18 | Complex intrachromosomal rearrangement in 1q leading to 1q32.2 microdeletion: a potential role of SRGAP2 in the gyrification of cerebral cortex. <i>Molecular Cytogenetics</i> , 2016, 9, 19. | 0.4 | 11 |

| # | ARTICLE | IF | CITATIONS |
|----|--|-----|-----------|
| 19 | 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. <i>Translational Neuroscience</i> , 2015, 6, 59-86. | 0.7 | 9 |
| 20 | High rates of submicroscopic aberrations in karyotypically normal acute lymphoblastic leukemia. <i>Molecular Cytogenetics</i> , 2015, 8, 45. | 0.4 | 17 |
| 21 | Isochromosome 17q in Chronic Lymphocytic Leukemia. <i>Leukemia Research and Treatment</i> , 2015, 2015, 1-6. | 2.0 | 1 |
| 22 | 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. <i>Oncology Reports</i> , 2015, 33, 625-630. | 1.2 | 4 |
| 23 | Novel Cryptic Rearrangements in Adult B-Cell Precursor Acute Lymphoblastic Leukemia Involving the MLL Gene. <i>Journal of Histochemistry and Cytochemistry</i> , 2015, 63, 384-390. | 1.3 | 7 |
| 24 | Comprehensive chronic lymphocytic leukemia diagnostics by combined multiplex ligation dependent probe amplification (MLPA) and interphase fluorescence in situ hybridization (iFISH). <i>Molecular Cytogenetics</i> , 2014, 7, 79. | 0.4 | 27 |
| 25 | 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. <i>Leukemia Research and Treatment</i> , 2014, 2014, 1-7. | 2.0 | 7 |
| 26 | Heteromorphic variants of chromosome 9. <i>Molecular Cytogenetics</i> , 2013, 6, 14. | 0.4 | 31 |
| 27 | <i>Aporrectodea caliginosa</i> , a suitable earthworm species for field based genotoxicity assessment?. <i>Environmental Pollution</i> , 2011, 159, 841-849. | 3.7 | 31 |
| 28 | Deletion 2q31.2â€“q31.3 in a 4â€“year-old girl with microcephaly and severe mental retardation. <i>American Journal of Medical Genetics, Part A</i> , 2011, 155, 1476-1482. | 0.7 | 8 |
| 29 | Is There a Yet Unreported Unbalanced Chromosomal Abnormality without Phenotypic Consequences in Proximal 4p?. <i>Cytogenetic and Genome Research</i> , 2011, 132, 121-123. | 0.6 | 4 |
| 30 | Somatic Mosaicism in Cases with Small Supernumerary Marker Chromosomes. <i>Current Genomics</i> , 2010, 11, 432-439. | 0.7 | 44 |
| 31 | Presence of harmless small supernumerary marker chromosomes hampers molecular genetic diagnosis: a case report. <i>Molecular Medicine Reports</i> , 2010, 3, 571-4. | 1.1 | 7 |
| 32 | Trisomy 21 with a Small Supernumerary Marker Chromosome Derived from Chromosomes 13/21 and 18. <i>Balkan Journal of Medical Genetics</i> , 2010, 13, 55-58. | 0.5 | 1 |
| 33 | First molecular cytogenetic characterisation of tracheal squamous cell carcinoma cell line KLN 205. , 0, , . | | 0 |