# Yuji Nakamachi

List of Publications by Year
in descending order



| 24 | 598 | 1040056 9 | 642732 23 |
|---|---|---|---|
| papers | citations | h-index | g-index |

| 25 | 25 | 25 | 936 |
|---|---|---|---|
| all docs | docs citations | times ranked | citing authors |

| # | Article | IF | Citations |
|---|---|---|---|
| 1 | MicroRNAâ€124a is a key regulator of proliferation and monocyte chemoattractant protein 1 secretion in fibroblastâ€like synoviocytes from patients with rheumatoid arthritis. Arthritis and Rheumatism, 2009, 60, 1294-1304. | 6.7 | 290 |
| 2 | MicroRNA-124 inhibits the progression of adjuvant-induced arthritis in rats. Annals of the Rheumatic Diseases, 2016, 75, 601-608. | 0.9 | 80 |
| 3 | Neurological outcomes in symptomatic congenital cytomegalovirus-infected infants after introduction of newborn urine screening and antiviral treatment. Brain and Development, 2016, 38, 209-216. | 1.1 | 45 |
| 4 | Efficacy of Valganciclovir Treatment Depends on the Severity of Hearing Dysfunction in Symptomatic Infants with Congenital Cytomegalovirus Infection. International Journal of Molecular Sciences, 2019, 20, 1388. | 4.1 | 23 |
| 5 | MicroRNA-124 inhibits TNF-Î±- and IL-6-induced osteoclastogenesis. Rheumatology International, 2019, 39, 689-695. | 3.0 | 23 |
| 6 | A new transcriptional variant and small azurophilic granules in an acute promyelocytic leukemia case with NPM1/RARA fusion gene. International Journal of Hematology, 2015, 102, 713-718. | 1.6 | 19 |
| 7 | Low total IgM values and high cytomegalovirus loads in the blood of newborns with symptomatic congenital cytomegalovirus infection. Journal of Perinatal Medicine, 2015, 43, 239-243. | 1.4 | 16 |
| 8 | Identification of a heterozygous p.Gly568Val missense mutation in the <scp>TRPV</scp>3 gene in a Japanese patient with Olmsted syndrome: In silico analysis of <scp>TRPV</scp>3. Journal of Dermatology, 2017, 44, 1059-1062. | 1.2 | 11 |
| 9 | Prediction of poor neurological development in patients with symptomatic congenital cytomegalovirus diseases after oral valganciclovir treatment. Brain and Development, 2019, 41, 743-750. | 1.1 | 11 |
| 10 | Determination of the antimicrobial susceptibility and molecular profile of clarithromycin resistance in the Mycobacterium abscessus complex in Japan by variable number tandem repeat analysis. Diagnostic Microbiology and Infectious Disease, 2018, 91, 256-259. | 1.8 | 9 |
| 11 | Diagnostic Value of Cytomegalovirus IgM Antibodies at Birth in PCR-Confirmed Congenital Cytomegalovirus Infection. International Journal of Molecular Sciences, 2019, 20, 3239. | 4.1 | 9 |
| 12 | Fluorescent Signaling of Molecularly Imprinted Nanogels Prepared via Postimprinting Modifications for Specific Protein Detection. Advanced NanoBiomed Research, 2021, 1, 2000079. | 3.6 | 9 |
| 13 | Quantitative evaluation of ventricular dilatation using computed tomography in infants with congenital cytomegalovirus infection. Brain and Development, 2014, 36, 10-15. | 1.1 | 8 |
| 14 | Sequence analyses of variable cytomegalovirus genes for distinction between breast milkâ€“ and transfusionâ€transmitted infections in veryâ€lowâ€birthâ€weight infants. Transfusion, 2018, 58, 2894-2902. | 1.6 | 8 |
| 15 | Expression of a novelZMYND11/MBTD1fusion transcript in CD7<sup>+</sup>CD56<sup>+</sup>acute myeloid leukemia with t(10;17)(p15;q21). Leukemia and Lymphoma, 2018, 59, 2706-2710. | 1.3 | 8 |
| 16 | Prenatal genetic testing for familial severe congenital protein C deficiency. Human Genome Variation, 2015, 2, 15017. | 0.7 | 7 |
| 17 | Telomeric Region of the Spinal Muscular Atrophy Locus Is Susceptible to Structural Variations. Pediatric Neurology, 2016, 58, 83-89. | 2.1 | 7 |
| 18 | ETV6â€“ABL1 fusion combined with monosomy 7 in childhood B-precursor acute lymphoblastic leukemia. International Journal of Hematology, 2018, 107, 604-609. | 1.6 | 6 |

| # | Article | IF | Citations |
|---|---|---|---|
| 19 | Associations between left ventricular diastolic function and right ventricular function in patients with and without preserved left ventricular ejection fraction. Journal of Echocardiography, 2018, 16, 81-86. | 0.8 | 2 |
| 20 | Coexpression ofETV6/MDS1/EVI1andETV6/EVI1fusion transcripts in acute myeloid leukemia with t(3;12)(q26.2;p13) and thrombocytosis. Leukemia and Lymphoma, 2019, 60, 1294-1298. | 1.3 | 2 |
| 21 | Detection of a novel CBFB-MYH11 fusion transcript in acute myeloid leukemia M1 with inv(16)(p13q22). Cancer Genetics, 2020, 241, 72-76. | 0.4 | 2 |
| 22 | Lymphoplasmacytic lymphoma in a patient with Birtâ€“Hoggâ€“DubÃ© syndrome. International Journal of Hematology, 2020, 112, 864-870. | 1.6 | 2 |
| 23 | Genetic Analysis of UGT1A1 Polymorphisms Using Preserved Dried Umbilical Cord for Assessing the Potential of Neonatal Jaundice as a Risk Factor for Autism Spectrum Disorder in Children. Journal of Autism and Developmental Disorders, 2022, 52, 483-489. | 2.7 | 1 |
| 24 | HLA-Matched and HLA-Haploidentical Hematopoietic Stem Cell Transplantation for Acute Myelogenous Leukemia with t (16;21) (p11.2;q22). Journal of Hematopoietic Cell Transplantation, 2020, 9, 65-69. | 0.1 | 0 |