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List of Publications by Year in descending order

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1478505 1474206 11 161 9 6 citations h-index g-index papers 12 12 12 250 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	Strong association between acute phase reactants (high-sensitivity C-reactive protein and erythrocyte) Tj ETQq1 genotype., 2022, 33, 201045.	1 0.78431	4 rgBT /Over 0
2	An acquired stable variant of a dicentric dic(9;20) and complex karyotype in a Syrian childhood B-acute lymphoblastic leukemia case. Molecular Cytogenetics, 2020, 13, 29.	0.9	1
3	A new childhood ALL case with an extremely complex karyotype and acute spontaneous tumor lysis syndrome. Molecular Cytogenetics, 2020, 13, 44.	0.9	0
4	Geographical distribution of \hat{l}^2 -globin gene mutations in Syria. Hematology, 2018, 23, 697-704.	1.5	22
5	Mutations of familial Mediterranean fever in Syrian patients and controls: Evidence for high carrier rate. Gene Reports, 2017, 6, 87-92.	0.8	4
6	Familial Mediterranean fever in Syrian children: phenotype–genotype correlation. Rheumatology International, 2015, 35, 629-634.	3.0	17
7	Molecular Update of \hat{l}^2 -Thalassemia Mutations in the Syrian Population: Identification of Rare \hat{l}^2 -Thalassemia Mutations. Hemoglobin, 2014, 38, 272-276.	0.8	26
8	Arthritis patterns in familial Mediterranean fever patients and association with M694V mutation. Molecular Biology Reports, 2011, 38, 2033-2036.	2.3	33
9	A new case of de novo translocation (12;17;18)(q21.2;q22;q21.1) and cranioâ€eerebelloâ€eardiac (3C) syndrome. American Journal of Medical Genetics, Part A, 2011, 155, 648-651.	1.2	3
10	Familial Mediterranean fever in Syrian patients: MEFV gene mutations and genotype–phenotype correlation. Molecular Biology Reports, 2010, 37, 1-5.	2.3	42
11	Combination of conventional multiplex PCR and quantitative real-time PCR detects large rearrangements in the dystrophin gene in 59% of Syrian DMD/BMD patients. Clinical Biochemistry, 2010, 43, 836-842.	1.9	13