## Patricia Pérez-Vera

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	The insecticides permethrin and chlorpyrifos show limited genotoxicity and no leukemogenic potential in human and murine hematopoietic stem progenitor cells. Haematologica, 2022, 107, 544-549.	3.5	3
2	High occurrence of CRLF2 abnormalities in Mexican children with B-cell acute lymphoblastic leukemia. Cytokine, 2022, 155, 155896.	3.2	5
3	UPD(14)mat and UPD(14)mat in concomitance with mosaic small supernumerary marker chromosome 14 in two new patients with Temple syndrome. European Journal of Medical Genetics, 2021, 64, 104199.	1.3	3
4	Genetic and clinical characterization of 73 Pigmentary Mosaicism patients: revealing the genetic basis of clinical manifestations. Orphanet Journal of Rare Diseases, 2019, 14, 259.	2.7	14
5	Variants in ARID5B gene are associated with the development of acute lymphoblastic leukemia in Mexican children. Annals of Hematology, 2019, 98, 2379-2388.	1.8	11
6	Pyrethroid pesticide exposure and hematological cancer: epidemiological, biological and molecular evidence. Reviews on Environmental Health, 2019, 34, 197-210.	2.4	31
7	Low concentrations of permethrin and malathion induce numerical and structural abnormalities in <i>KMT2A</i> and <i>IGH</i> genes in vitro. Journal of Applied Toxicology, 2018, 38, 1262-1270.	2.8	12
8	"Exposure to the insecticides permethrin and malathion induces leukemia and lymphoma-associated gene aberrations in vitro ― Toxicology in Vitro, 2017, 44, 17-26.	2.4	20
9	Epigenetic alterations in acute lymphoblastic leukemia. BoletÃn Médico Del Hospital Infantil De México (English Edition), 2017, 74, 243-264.	0.0	14
10	Expression of Ik6 and Ik8 Isoforms and Their Association with Relapse and Death in Mexican Children with Acute Lymphoblastic Leukemia. PLoS ONE, 2015, 10, e0130756.	2.5	7
11	Cytogenomic and phenotypic analysis in lowâ€level monosomy 7 mosaicism with nonâ€supernumerary ring chromosome 7. American Journal of Medical Genetics, Part A, 2014, 164, 1765-1769.	1.2	8
12	Partial and complete trisomy 14 mosaicism: clinical follow-up, cytogenetic and molecular analysis. Molecular Cytogenetics, 2014, 7, 65.	0.9	15
13	Significance of <i>CASP8AP2</i> and <i>H2AFZ</i> expression in survival and risk of relapse in children with acute lymphoblastic leukemia. Leukemia and Lymphoma, 2014, 55, 2305-2311.	1.3	16
14	Expression of RUNX1 isoforms and its target gene BLK in childhood acute lymphoblastic leukemia. Leukemia Research, 2012, 36, 1105-1111.	0.8	16
15	Multiple copies of RUNX1: description of 14 new patients, follow-up, and a review of the literature. Cancer Genetics and Cytogenetics, 2008, 180, 129-134.	1.0	12
16	Analysis of gene rearrangements using a fluorescence in situ hybridization method in Mexican patients with acute lymphoblastic leukemia: experience at a single institution. Cancer Genetics and Cytogenetics, 2008, 184, 94-98.	1.0	13
17	Detection of ETV6 and RUNX1 gene rearrangements using fluorescence in situ hybridization in Mexican patients with acute lymphoblastic leukemia: experience at a single institution. Cancer Genetics and Cytogenetics, 2005, 162, 140-145.	1.0	14
18	Cytogenetics in Acute Lymphoblastic Leukemia in Mexican Children. Archives of Medical Research, 2001, 32, 202-207.	3.3	18