

# Karolina Skvarova Kramarzova

## List of Publications by Year in descending order

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Version: 2024-02-01

21  
papers

274  
citations

932766

10  
h-index

1125271

13  
g-index

22  
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22  
docs citations

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times ranked

681  
citing authors

#	ARTICLE	IF	CITATIONS
1	Germ-line GATA2 p.THR354MET mutation in familial myelodysplastic syndrome with acquired monosomy 7 and ASXL1 mutation demonstrating rapid onset and poor survival. <i>Haematologica</i> , 2012, 97, 890-894.	1.7	85
2	Real-time PCR quantification of major Wilmsâ€™ tumor gene 1 (WT1) isoforms in acute myeloid leukemia, their characteristic expression patterns and possible functional consequences. <i>Leukemia</i> , 2012, 26, 2086-2095.	3.3	31
3	CRISPR/Cas9-Mediated Correction of the FANCD1 Gene in Primary Patient Cells. <i>International Journal of Molecular Sciences</i> , 2017, 18, 1269.	1.8	23
4	Prognosis of ProB ALL in Children. <i>Blood</i> , 2008, 112, 2512-2512.	0.6	19
5	<scp>TLR8</scp>/<scp>TLR7</scp> dysregulation due to a novel <i>TLR8</i> mutation causes severe autoimmune hemolytic anemia and autoinflammation in identical twins. <i>American Journal of Hematology</i> , 2022, 97, 338-351.	2.0	17
6	Wilms tumor gene 1 (WT1), TP53, RAS/BRAF and KIT aberrations in testicular germ cell tumors. <i>Cancer Letters</i> , 2016, 376, 367-376.	3.2	16
7	Low HOX gene expression in PML-RAR $\pm$ -positive leukemia results from suppressed histone demethylation. <i>Epigenetics</i> , 2018, 13, 73-84.	1.3	16
8	Homeobox gene expression in acute myeloid leukemia is linked to typical underlying molecular aberrations. <i>Journal of Hematology and Oncology</i> , 2014, 7, 94.	6.9	14
9	A homozygous deletion in the SLC19A1 gene as a cause of folate-dependent recurrent megaloblastic anemia. <i>Blood</i> , 2020, 135, 2427-2431.	0.6	13
10	Molecular Basis of Cisplatin Resistance in Testicular Germ Cell Tumors. <i>Cancers</i> , 2019, 11, 1316.	1.7	12
11	Novel SAMD9 Mutation in a Patient With Immunodeficiency, Neutropenia, Impaired Anti-CMV Response, and Severe Gastrointestinal Involvement. <i>Frontiers in Immunology</i> , 2019, 10, 2194.	2.2	12
12	Expression Pattern of WT1 Isoforms in Patients with Acute Myeloid Leukemia (AML), Myelodysplastic Syndrome (MDS) and Severe Aplastic Anemia (SAA). <i>Blood</i> , 2011, 118, 2502-2502.	0.6	10
13	Evaluation of WT1 expression in bone marrow vs peripheral blood samples of children with acute myeloid leukemiaâ€™ impact on minimal residual disease detection. <i>Leukemia</i> , 2013, 27, 1194-1196.	3.3	6
14	WT1 Expression at the Diagnosis of Childhood AML Has No Prognostic Value but Corresponds with the Biological Characteristics of Leukemic Cells - Results From European Multicenter Study.. <i>Blood</i> , 2010, 116, 1684-1684.	0.6	0
15	Changes Identified by Flow Cytometry and WT1 Expression in Consecutive Bone Marrow Samples in Refractory Cytopenia of Childhood and Aplastic Anemia Before Start of the Therapy. <i>Blood</i> , 2011, 118, 1342-1342.	0.6	0
16	Transcription Regulation of HOX Genes in Normal Hematopoiesis and Leukemogenesis in Children. <i>Blood</i> , 2012, 120, 4614-4614.	0.6	0
17	Leukemic Pattern Of HOX Gene Expression Is Driven By Genetic Aberrations Through Epigenetic Modifiers. <i>Blood</i> , 2013, 122, 2504-2504.	0.6	0
18	The Role of Histone Demethylases in the Transcription Regulation of HOX Genes in PML-RARa+ AML Patients. <i>Blood</i> , 2014, 124, 876-876.	0.6	0

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19	Wilms's tumor gene 1 (WT1) aberrations in testicular germ cell tumors (TGCTs).. Journal of Clinical Oncology, 2015, 33, 4534-4534.	0.8	0
20	The Role of Histone Demethylases and DNA Methyltransferases in the Transcription Regulation of HOX Genes in PML-RARa+ AML Patients. Blood, 2016, 128, 3921-3921.	0.6	0
21	Folate-Dependent Normocytic Anemia Caused By a Hypomorphic Mutation in SLC19A1 gene. Blood, 2018, 132, 502-502.	0.6	0