

Anindita Roy

List of Publications by Year in descending order

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Version: 2024-02-01

38
papers

1,853
citations

394421

19
h-index

395702

33
g-index

45
all docs

45
docs citations

45
times ranked

3135
citing authors

#	ARTICLE	IF	CITATIONS
1	Decoding human fetal liver haematopoiesis. <i>Nature</i> , 2019, 574, 365-371.	27.8	392
2	GATA1-mutant clones are frequent and often unsuspected in babies with Down syndrome: identification of a population at risk of leukemia. <i>Blood</i> , 2013, 122, 3908-3917.	1.4	162
3	Abnormalities in the myeloid progenitor compartment in Down syndrome fetal liver precede acquisition of GATA1 mutations. <i>Blood</i> , 2008, 112, 4507-4511.	1.4	143
4	Perturbation of fetal liver hematopoietic stem and progenitor cell development by trisomy 21. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2012, 109, 17579-17584.	7.1	138
5	Single-cell profiling of human megakaryocyte-erythroid progenitors identifies distinct megakaryocyte and erythroid differentiation pathways. <i>Genome Biology</i> , 2016, 17, 83.	8.8	124
6	Acute megakaryoblastic leukaemia (AMKL) and transient myeloproliferative disorder (TMD) in Down syndrome: a multi-step model of myeloid leukaemogenesis. <i>British Journal of Haematology</i> , 2009, 147, 3-12.	2.5	115
7	Unraveling the cellular origin and clinical prognostic markers of infant B-cell acute lymphoblastic leukemia using genome-wide analysis. <i>Haematologica</i> , 2019, 104, 1176-1188.	3.5	76
8	Blood and immune development in human fetal bone marrow and Down syndrome. <i>Nature</i> , 2021, 598, 327-331.	27.8	73
9	MLL-AF4 Spreading Identifies Binding Sites that Are Distinct from Super-Enhancers and that Govern Sensitivity to DOT1L Inhibition in Leukemia. <i>Cell Reports</i> , 2017, 18, 482-495.	6.4	69
10	Biology and management of transient abnormal myelopoiesis (TAM) in children with Down syndrome. <i>Seminars in Fetal and Neonatal Medicine</i> , 2012, 17, 196-201.	2.3	62
11	Discovery of a CD10-negative B-progenitor in human fetal life identifies unique ontogeny-related developmental programs. <i>Blood</i> , 2019, 134, 1059-1071.	1.4	62
12	Stem and progenitor cell dysfunction in human trisomies. <i>EMBO Reports</i> , 2015, 16, 44-62.	4.5	38
13	Transitions in lineage specification and gene regulatory networks in hematopoietic stem/progenitor cells over human development. <i>Cell Reports</i> , 2021, 36, 109698.	6.4	38
14	H3K79me2/3 controls enhancer-promoter interactions and activation of the pan-cancer stem cell marker PROM1/CD133 in MLL-AF4 leukemia cells. <i>Leukemia</i> , 2021, 35, 90-106.	7.2	35
15	The genome-wide impact of trisomy 21 on DNA methylation and its implications for hematopoiesis. <i>Nature Communications</i> , 2021, 12, 821.	12.8	32
16	Single-cell profiling of human bone marrow progenitors reveals mechanisms of failing erythropoiesis in Diamond-Blackfan anemia. <i>Science Translational Medicine</i> , 2021, 13, eabf0113.	12.4	32
17	A human fetal liver-derived infant MLL-AF4 acute lymphoblastic leukemia model reveals a distinct fetal gene expression program. <i>Nature Communications</i> , 2021, 12, 6905.	12.8	28
18	Impaired human hematopoiesis due to a cryptic intronic GATA1 splicing mutation. <i>Journal of Experimental Medicine</i> , 2019, 216, 1050-1060.	8.5	27

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19	In utero origin of myelofibrosis presenting in adult monozygotic twins. <i>Nature Medicine</i> , 2022, 28, 1207-1211.	30.7	26
20	Heterogeneous disease-propagating stem cells in juvenile myelomonocytic leukemia. <i>Journal of Experimental Medicine</i> , 2021, 218, .	8.5	25
21	The Origin of B-cells: Human Fetal B Cell Development and Implications for the Pathogenesis of Childhood Acute Lymphoblastic Leukemia. <i>Frontiers in Immunology</i> , 2021, 12, 637975.	4.8	22
22	The impact of trisomy 21 on foetal haematopoiesis. <i>Blood Cells, Molecules, and Diseases</i> , 2013, 51, 277-281.	1.4	21
23	Single-cell analysis of bone marrow-derived CD34+ cells from children with sickle cell disease and thalassemia. <i>Blood</i> , 2019, 134, 2111-2115.	1.4	21
24	The BET inhibitor CPI203 promotes ex vivo expansion of cord blood long-term repopulating HSCs and megakaryocytes. <i>Blood</i> , 2020, 136, 2410-2415.	1.4	18
25	A KMT2A-AFF1 gene regulatory network highlights the role of core transcription factors and reveals the regulatory logic of key downstream target genes. <i>Genome Research</i> , 2021, 31, 1159-1173.	5.5	16
26	High resolution IgH repertoire analysis reveals fetal liver as the likely origin of life-long, innate B lymphopoiesis in humans. <i>Clinical Immunology</i> , 2017, 183, 8-16.	3.2	15
27	MLL-rearranged infant leukaemia: A "thorn in the side"™ of a remarkable success story. <i>Biochimica Et Biophysica Acta - Gene Regulatory Mechanisms</i> , 2020, 1863, 194564.	1.9	13
28	The impact of trisomy 21 on early human hematopoiesis. <i>Cell Cycle</i> , 2013, 12, 533-534.	2.6	8
29	Germline variants in predisposition genes in children with Down syndrome and acute lymphoblastic leukemia. <i>Blood Advances</i> , 2020, 4, 672-675.	5.2	5
30	Processing single-cell RNA-seq datasets using SingCellaR. <i>STAR Protocols</i> , 2022, 3, 101266.	1.2	5
31	A "œgut feeling" about precursor B-ALL. <i>Blood</i> , 2020, 136, 1995-1996.	1.4	2
32	Trilineage Perturbation of Hematopoiesis In Neonates with Down Syndrome. <i>Blood</i> , 2010, 116, 876-876.	1.4	1
33	Developmental Stage Specific B-Progenitor Expansion in Normal Fetal Bone Marrow Is Absent in Down Syndrome: Implications for Infant ALL. <i>Blood</i> , 2014, 124, 4331-4331.	1.4	1
34	Epigenome-wide association study of acute lymphoblastic leukemia in children with Down syndrome. <i>Blood Advances</i> , 2022, 6, 4132-4136.	5.2	1
35	Clinical and Hematologic Impact of Fetal and Perinatal Variables on Mutant GATA1 Clone Size in Neonates with Down Syndrome. <i>Blood</i> , 2014, 124, 2349-2349.	1.4	0
36	Trisomy 21-Associated Abnormalities in IGF Signalling and the Fetal Microenvironment Both Contribute to Disruption of Fetal Hematopoiesis in Down Syndrome. <i>Blood</i> , 2014, 124, 1885-1885.	1.4	0

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37	High Resolution Igh Repertoire Analysis Reveals the Human Fetal Liver As the Origin of Life-Long, Innate B Lymphopoiesis. Blood, 2016, 128, 127-127.	1.4	0
38	Base Editing Repairs the HbE Mutation Restoring the Production of Normal Globin Chains in Severe HbE/ β^0 -Thalassemia Patient Hematopoietic Stem Cells and Erythroid Cells. Blood, 2021, 138, 2935-2935.	1.4	0