

# 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	Processing single-cell RNA-seq datasets using SingCellaR. STAR Protocols, 2022, 3, 101266.	1.2	5
2	Epigenome-wide association study of acute lymphoblastic leukemia in children with Down syndrome. Blood Advances, 2022, 6, 4132-4136.	5.2	1
3	In utero origin of myelofibrosis presenting in adult monozygotic twins. Nature Medicine, 2022, 28, 1207-1211.	30.7	26
4	H3K79me2/3 controls enhancer-promoter interactions and activation of the pan-cancer stem cell marker PROM1/CD133 in MLL-AF4 leukemia cells. Leukemia, 2021, 35, 90-106.	7.2	35
5	The genome-wide impact of trisomy 21 on DNA methylation and its implications for hematopoiesis. Nature Communications, 2021, 12, 821.	12.8	32
6	The Origin of B-cells: Human Fetal B Cell Development and Implications for the Pathogenesis of Childhood Acute Lymphoblastic Leukemia. Frontiers in Immunology, 2021, 12, 637975.	4.8	22
7	A KMT2A-AFF1 gene regulatory network highlights the role of core transcription factors and reveals the regulatory logic of key downstream target genes. Genome Research, 2021, 31, 1159-1173.	5.5	16
8	Single-cell profiling of human bone marrow progenitors reveals mechanisms of failing erythropoiesis in Diamond-Blackfan anemia. Science Translational Medicine, 2021, 13, eabf0113.	12.4	32
9	Transitions in lineage specification and gene regulatory networks in hematopoietic stem/progenitor cells over human development. Cell Reports, 2021, 36, 109698.	6.4	38
10	Blood and immune development in human fetal bone marrow and Down syndrome. Nature, 2021, 598, 327-331.	27.8	73
11	Heterogeneous disease-propagating stem cells in juvenile myelomonocytic leukemia. Journal of Experimental Medicine, 2021, 218, .	8.5	25
12	A human fetal liver-derived infant MLL-AF4 acute lymphoblastic leukemia model reveals a distinct fetal gene expression program. Nature Communications, 2021, 12, 6905.	12.8	28
13	Base Editing Repairs the HbE Mutation Restoring the Production of Normal Globin Chains in Severe HbE/ $\beta^2$ -Thalassemia Patient Hematopoietic Stem Cells and Erythroid Cells. Blood, 2021, 138, 2935-2935.	1.4	0
14	A "œgut feeling" about precursor B-ALL. Blood, 2020, 136, 1995-1996.	1.4	2
15	MLL-rearranged infant leukaemia: A "thorn in the side" of a remarkable success story. Biochimica Et Biophysica Acta - Gene Regulatory Mechanisms, 2020, 1863, 194564.	1.9	13
16	The BET inhibitor CPI203 promotes ex vivo expansion of cord blood long-term repopulating HSCs and megakaryocytes. Blood, 2020, 136, 2410-2415.	1.4	18
17	Germline variants in predisposition genes in children with Down syndrome and acute lymphoblastic leukemia. Blood Advances, 2020, 4, 672-675.	5.2	5
18	Discovery of a CD10-negative B-progenitor in human fetal life identifies unique ontogeny-related developmental programs. Blood, 2019, 134, 1059-1071.	1.4	62

#	ARTICLE	IF	CITATIONS
19	Decoding human fetal liver haematopoiesis. <i>Nature</i> , 2019, 574, 365-371.	27.8	392
20	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
21	Impaired human hematopoiesis due to a cryptic intronic GATA1 splicing mutation. <i>Journal of Experimental Medicine</i> , 2019, 216, 1050-1060.	8.5	27
22	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
23	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
24	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
25	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
26	High Resolution Igh Repertoire Analysis Reveals the Human Fetal Liver As the Origin of Life-Long, Innate B Lymphopoiesis. <i>Blood</i> , 2016, 128, 127-127.	1.4	0
27	Stem and progenitor cell dysfunction in human trisomies. <i>EMBO Reports</i> , 2015, 16, 44-62.	4.5	38
28	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
29	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
30	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
31	The impact of trisomy 21 on foetal haematopoiesis. <i>Blood Cells, Molecules, and Diseases</i> , 2013, 51, 277-281.	1.4	21
32	The impact of trisomy 21 on early human hematopoiesis. <i>Cell Cycle</i> , 2013, 12, 533-534.	2.6	8
33	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
34	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
35	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
36	Trilineage Perturbation of Hematopoiesis In Neonates with Down Syndrome. <i>Blood</i> , 2010, 116, 876-876.	1.4	1

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37	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
38	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