## Jeffrey M Lipton

List of Publications by Year in descending order

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567281 713466 1,411 25 15 21 citations h-index g-index papers 26 26 26 2652 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	A new system for naming ribosomal proteins. Current Opinion in Structural Biology, 2014, 24, 165-169.	5.7	481
2	The Genetic Landscape of Diamond-Blackfan Anemia. American Journal of Human Genetics, 2018, 103, 930-947.	6.2	184
3	Diamond-Blackfan Anemia: Diagnosis, Treatment, and Molecular Pathogenesis. Hematology/Oncology Clinics of North America, 2009, 23, 261-282.	2.2	174
4	Erythropoiesis: insights into pathophysiology and treatments in 2017. Molecular Medicine, 2018, 24, 11.	4.4	76
5	Pomalidomide reverses $\hat{I}^3$ -globin silencing through the transcriptional reprogramming of adult hematopoietic progenitors. Blood, 2016, 127, 1481-1492.	1.4	75
6	Increased risk of colon cancer and osteogenic sarcoma in Diamond-Blackfan anemia. Blood, 2018, 132, 2205-2208.	1.4	64
7	Diamond Blackfan anemia 2008–2009: broadening the scope of ribosome biogenesis disorders. Current Opinion in Pediatrics, 2010, 22, 12-19.	2.0	50
8	HMGB1 Mediates Anemia of Inflammation in Murine Sepsis Survivors. Molecular Medicine, 2015, 21, 951-958.	4.4	45
9	Late Effects Screening Guidelines after Hematopoietic Cell Transplantation for Inherited Bone Marrow Failure Syndromes: Consensus Statement From the Second Pediatric Blood and Marrow Transplant Consortium International Conference on Late Effects After Pediatric HCT. Biology of Blood and Marrow Transplantation. 2017. 23. 1422-1428.	2.0	43
10	p53-Independent Cell Cycle and Erythroid Differentiation Defects in Murine Embryonic Stem Cells Haploinsufficient for Diamond Blackfan Anemia-Proteins: RPS19 versus RPL5. PLoS ONE, 2014, 9, e89098.	2.5	33
11	Increased Prevalence of Congenital Heart Disease in Children With Diamond Blackfan Anemia Suggests Unrecognized Diamond Blackfan Anemia as a Cause of Congenital Heart Disease in the General Population. Circulation Genomic and Precision Medicine, 2018, 11, e002044.	3.6	32
12	Current Knowledge and Priorities for Future Research in Late Effects after Hematopoietic Cell Transplantation for Inherited Bone Marrow Failure Syndromes: Consensus Statement from the Second Pediatric Blood and Marrow Transplant Consortium International Conference on Late Effects after Pediatric Hematopoietic Cell Transplantation. Biology of Blood and Marrow Transplantation,	2.0	31
13	2017, 23, 726-735.  Tropomodulin 1 controls erythroblast enucleation via regulation of F-actin in the enucleosome. Blood, 2017, 130, 1144-1155.	1.4	31
14	Molecular convergence in ex vivo models of Diamond-Blackfan anemia. Blood, 2017, 129, 3111-3120.	1.4	30
15	Varying presentations and favourable outcomes of COVIDâ€19 infection in children and young adults with sickle cell disease: an additional case series with comparisons to published cases. British Journal of Haematology, 2020, 190, e221-e224.	2.5	26
16	Lâ€leucine improves anemia and growth in patients with transfusionâ€dependent Diamondâ€Blackfan anemia: Results from a multicenter pilot phase I/II study from the Diamondâ€Blackfan Anemia Registry. Pediatric Blood and Cancer, 2020, 67, e28748.	1.5	12
17	A functional assay for the clinical annotation of genetic variants of uncertain significance in Diamond-Blackfan anemia. Human Mutation, 2018, 39, 1102-1111.	2.5	9
18	Ribosomopathy Association With Colorectal Cancer. Gastroenterology, 2015, 148, 258.	1.3	6

#	Article	IF	CITATIONS
19	SNP Array Genotyping Reveals Constitutional and Mosaic Losses of Ribosomal Protein Gene Regions In Patients with Diamond Blackfan Anemia without Ribosomal Protein Gene Mutations Blood, 2010, 116, 1168-1168.	1.4	3
20	Autosomal Recessive Diamond-Blackfan Anemia: Identification Of Mutations In MCM2 and Flnb. Blood, 2013, 122, 589-589.	1.4	3
21	Inherited thrombocytopenia and Occam's razor. Blood, 2017, 130, 839-840.	1.4	2
22	HMGB1 Causes Anemia of Inflammation By Modulating Erythropoietin Signal Transduction. Blood, 2018, 132, 628-628.	1.4	1
23	Shwachman Diamond syndromeâ€"phenotypes and genotypes: When clinical research informs biology. Pediatric Blood and Cancer, 2008, 51, 449-450.	1.5	O
24	Identification of New Rare Sequence Changes in RP Genes in Diamond-Blackfan Anemia and Association of the RPL5 and RPL11 Mutations with Craniofacial and Thumb Malformations. Blood, 2008, 112, 39-39.	1.4	0
25	Inhibition of Human Erythropoiesis during Inflammation Is Mediated By High Mobility Group Box Protein 1 (HMGB1) through Decreased Commitment of Hematopoietic Stem Cells to the Erythroid Lineage and By Increased Apoptosis of Terminally Differentiating Erythroblasts. Blood, 2016, 128, 702-702.	1.4	0