## Jordan A Shavit

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Biobank-driven genomic discovery yields new insight into atrial fibrillation biology. Nature Genetics, 2018, 50, 1234-1239.	21.4	547
2	The world according to Maf. Nucleic Acids Research, 1997, 25, 2953-2959.	14.5	248
3	Positive or Negative MARE-Dependent Transcriptional Regulation Is Determined by the Abundance of Small Maf Proteins. Cell, 2000, 103, 865-876.	28.9	136
4	<i>pak2a</i> mutations cause cerebral hemorrhage in <i>redhead</i> zebrafish. Proceedings of the National Academy of Sciences of the United States of America, 2007, 104, 13996-14001.	7.1	89
5	Zebrafish as a model system for the study of hemostasis and thrombosis. Current Opinion in Hematology, 2014, 21, 418-422.	2.5	69
6	Structure/Function Analysis of Recurrent Mutations in SETD2 Protein Reveals a Critical and Conserved Role for a SET Domain Residue in Maintaining Protein Stability and Histone H3 Lys-36 Trimethylation. Journal of Biological Chemistry, 2016, 291, 21283-21295.	3.4	64
7	Functions of the COPII gene paralogs SEC23A and SEC23B are interchangeable in vivo. Proceedings of the National Academy of Sciences of the United States of America, 2018, 115, E7748-E7757.	7.1	58
8	Effects of MYBPC3 loss-of-function mutations preceding hypertrophic cardiomyopathy. JCI Insight, 2020, 5, .	5.0	58
9	Targeted mutagenesis of zebrafish antithrombin III triggers disseminated intravascular coagulation and thrombosis, revealing insight into function. Blood, 2014, 124, 142-150.	1.4	52
10	Loss of Fibrinogen in Zebrafish Results in Symptoms Consistent with Human Hypofibrinogenemia. PLoS ONE, 2013, 8, e74682.	2.5	48
11	Porphyrin-Induced Protein Oxidation and Aggregation as a Mechanism of Porphyria-Associated Cell Injury. Cellular and Molecular Gastroenterology and Hepatology, 2019, 8, 535-548.	4.5	44
12	Characterization of Zebrafish von Willebrand Factor Reveals Conservation of Domain Structure, Multimerization, and Intracellular Storage. Advances in Hematology, 2012, 2012, 1-9.	1.0	30
13	Modeling Disorders of Blood Coagulation in the Zebrafish. Current Pathobiology Reports, 2015, 3, 155-161.	3.4	29
14	Thrombocyte Inhibition Restores Protective Immunity to Mycobacterial Infection in Zebrafish. Journal of Infectious Diseases, 2019, 220, 524-534.	4.0	28
15	Efficacy of emicizumab in a pediatric patient with type 3 von Willebrand disease and alloantibodies. Blood Advances, 2019, 3, 2748-2750.	5.2	26
16	Genome editing of factor X in zebrafish reveals unexpected tolerance of severe defects in the common pathway. Blood, 2017, 130, 666-676.	1.4	22
17	A precursorâ€inducible zebrafish model of acute protoporphyria with hepatic protein aggregation and multiorganelle stress. FASEB Journal, 2016, 30, 1798-1810.	0.5	21
18	Enhanced VWF biosynthesis and elevated plasma VWF due to a natural variant in the murine Vwf gene. Blood, 2006, 108, 3061-3067.	1.4	20

Jordan A Shavit

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19	Modifiers of von Willebrand factor identified by natural variation in inbred strains of mice. Blood, 2009, 114, 5368-5374.	1.4	20
20	Analysis of factor V in zebrafish demonstrates minimal levels needed for early hemostasis. Blood Advances, 2019, 3, 1670-1680.	5.2	18
21	Simple and Rapid Quantification of Thrombocytes in Zebrafish Larvae. Zebrafish, 2015, 12, 238-242.	1.1	17
22	Nfe2 is dispensable for early but required for adult thrombocyte formation and function in zebrafish. Blood Advances, 2018, 2, 3418-3427.	5.2	16
23	The Role of Platelets and ε-Aminocaproic Acid in Arthrogryposis, Renal Dysfunction, and Cholestasis (ARC) Syndrome Associated Hemorrhage. Pediatric Blood and Cancer, 2016, 63, 561-563.	1.5	14
24	The transcription factor, Nuclear factor, erythroid 2 (Nfe2), is a regulator of the oxidative stress response during Danio rerio development. Aquatic Toxicology, 2016, 180, 141-154.	4.0	13
25	Hemophilias and Other Disorders of Hemostasis. , 2013, , 1-33.		12
26	Membrane-myofibril cross-talk in myofibrillogenesis and in muscular dystrophy pathogenesis: lessons from the zebrafish. Frontiers in Physiology, 2014, 5, 14.	2.8	12
27	Loss of fibrinogen in zebrafish results in an asymptomatic embryonic hemostatic defect and synthetic lethality with thrombocytopenia. Journal of Thrombosis and Haemostasis, 2019, 17, 607-617.	3.8	12
28	Disruption of the kringle 1 domain of prothrombin leads to late onset mortality in zebrafish. Scientific Reports, 2020, 10, 4049.	3.3	10
29	Emicizumab prophylaxis to facilitate anticoagulant therapy for management of intraâ€atrial thrombosis in severe haemophilia with an inhibitor. Haemophilia, 2019, 25, e203-e205.	2.1	9
30	Zebrafish otolith biomineralization requires polyketide synthase. Mechanisms of Development, 2019, 157, 1-9.	1.7	9
31	A genetic modifier of venous thrombosis in zebrafish reveals a functional role for fibrinogen AαE in early hemostasis. Blood Advances, 2020, 4, 5480-5491.	5.2	9
32	Agent specific effects of anticoagulant induced alopecia. Research and Practice in Thrombosis and Haemostasis, 2017, 1, 90-92.	2.3	7
33	Genomeâ€wide linkage analysis and wholeâ€exome sequencing identifies an <i><scp>ITGA</scp>2B</i> mutation in a family with thrombocytopenia. British Journal of Haematology, 2019, 186, 574-579.	2.5	7
34	Novel treatments for hemophilia through rebalancing of the coagulation cascade. Pediatric Blood and Cancer, 2021, 68, e28934.	1.5	7
35	Otx2b mutant zebrafish have pituitary, eye and mandible defects that model mammalian disease. Human Molecular Genetics, 2020, 29, 1648-1657.	2.9	6
36	Nuclear Progestin Receptor–mediated Linkage of Blood Coagulation and Ovulation. Endocrinology, 2022, 163, .	2.8	5

JORDAN A SHAVIT

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37	Phage display broadly identifies inhibitorâ€reactive regions in von Willebrand factor. Journal of Thrombosis and Haemostasis, 2021, 19, 2702-2709.	3.8	4
38	The bleeding edge of symptom assessment. Pediatric Blood and Cancer, 2012, 58, 657-658.	1.5	3
39	Acitretin mitigates uroporphyrin-induced bone defects in congenital erythropoietic porphyria models. Scientific Reports, 2021, 11, 9601.	3.3	2
40	Conservation of Hemostatic System Component Function Between Zebrafish and Mammals Blood, 2009, 114, 3165-3165.	1.4	1
41	Factor X Mutant Zebrafish Tolerate a Severe Hemostatic Defect in Early Development Yet Develop Lethal Hemorrhage in Adulthood. Blood, 2015, 126, 426-426.	1.4	1
42	A Zebrafish Model Of Antithrombin III Deficiency Displays Bleeding and Thrombosis Secondary To Disseminated Intravascular Coagulation. Blood, 2013, 122, 200-200.	1.4	1
43	Nfe2 Is Dispensable for Early, but Required for Adult Thrombocyte Formation and Function in Zebrafish. Blood, 2016, 128, 2534-2534.	1.4	1
44	Genome Editing of Factor V in Zebrafish Embryos Results in a Severe Hemostatic Defect without Spontaneous Hemorrhage. Blood, 2016, 128, 2565-2565.	1.4	0