

# Julie Curtin

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/5508561/publications.pdf>

Version: 2024-02-01

10  
papers

118  
citations

1478505

6  
h-index

1872680

6  
g-index

10  
all docs

10  
docs citations

10  
times ranked

141  
citing authors

#	ARTICLE	IF	CITATIONS
1	Long-acting recombinant fusion protein linking coagulation factor IX with albumin (rIX-FP) in children. <i>Thrombosis and Haemostasis</i> , 2016, 116, 659-668.	3.4	50
2	ALPK1 missense pathogenic variant in five families leads to ROSAH syndrome, an ocular multisystem autosomal dominant disorder. <i>Genetics in Medicine</i> , 2019, 21, 2103-2115.	2.4	28
3	A Practical, One-Clinic Visit Protocol for Pharmacokinetic Profile Generation with the ADVATE myPKFiT Dosing Tool in Severe Hemophilia A Subjects. <i>Thrombosis and Haemostasis</i> , 2021, 121, 1326-1336.	3.4	12
4	Final results of the PUPs B-LONG study: evaluating safety and efficacy of rFIXFc in previously untreated patients with hemophilia B. <i>Blood Advances</i> , 2021, 5, 2732-2739.	5.2	11
5	Immune tolerance induction using a factor VIII/von Willebrand factor concentrate (BIOSTATEÂ®), with or without immunosuppression, in Australian paediatric severe haemophilia A patients with high titre inhibitors: A multicentre, retrospective study. <i>Thrombosis Research</i> , 2014, 134, 1046-1051.	1.7	9
6	Simplifying surgery in haemophilia B: Low factor IX consumption and infrequent infusions in surgical procedures with rIX-FP. <i>Thrombosis Research</i> , 2020, 188, 85-89.	1.7	8
7	Renal tubular dysfunction and lactic acidosis: Answers. <i>Pediatric Nephrology</i> , 2012, 27, 2215-2216.	1.7	0
8	Congenital Macrothrombocytopenia and Defective Localization of the Nonmuscle Myosin Heavy Chain IIA in Leukocytes and Megakaryocytes with a Normal MYH9 Gene.. <i>Blood</i> , 2004, 104, 3032-3032.	1.4	0
9	Use of Rituximab in Patients with Congenital Bleeding Disorders and High Titre Inhibitors.. <i>Blood</i> , 2009, 114, 3501-3501.	1.4	0
10	Extended Molecular and Clinical Phenotype of Human G6PC3 Deficiency.. <i>Blood</i> , 2010, 116, 1495-1495.	1.4	0