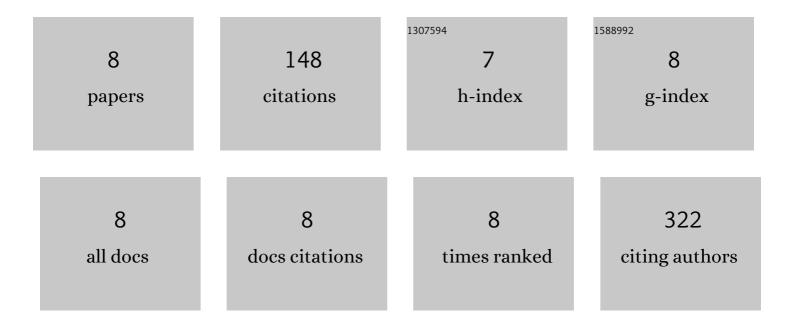
## **Clemens Stockklausner**

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/9597728/publications.pdf Version: 2024-02-01



#	Article	IF	CITATIONS
1	Bone marrow failure unresponsive to bone marrow transplant is caused by mutations in thrombopoietin. Blood, 2017, 130, 875-880.	1.4	42
2	The uORF-containing thrombopoietin mRNA escapes nonsense-mediated decay (NMD). Nucleic Acids Research, 2006, 34, 2355-2363.	14.5	41
3	The thrombopoietin receptor P106L mutation functionally separates receptor signaling activity from thrombopoietin homeostasis. Blood, 2015, 125, 1159-1169.	1.4	18
4	DNA methylation in <i>PRDM8</i> is indicative for dyskeratosis congenita. Oncotarget, 2016, 7, 10765-10772.	1.8	15
5	Hereditary thrombocythemia caused by a thrombopoietin (THPO) gain-of-function mutation associated with multiple myeloma and congenital limb defects. Annals of Hematology, 2012, 91, 1129-1133.	1.8	11
6	Long-term remission of children with relapsed and secondary anaplastic large cell non-Hodgkin lymphoma (ALCL) following treatment with pulsed dexamethasone and low dose etoposide. Pediatric Blood and Cancer, 2008, 50, 126-129.	1.5	10
7	Thrombocytosis in children and adolescents—classification, diagnostic approach, and clinical management. Annals of Hematology, 2021, 100, 1647-1665.	1.8	10
8	Mpl Gain-of-Function Mutations Can be Classified By Differential Subcellular Processing, Molecular Mechanisms, Mode of Inheritance and Clinical Impact. Blood, 2015, 126, 1634-1634.	1.4	1