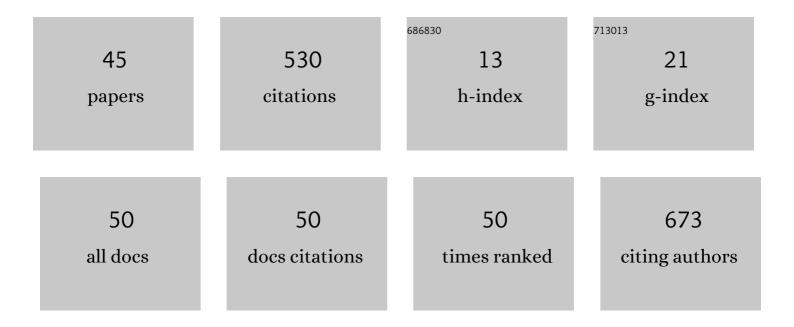
## Radoslaw Kaczmarek

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/1477658/publications.pdf Version: 2024-02-01



#	Article	IF	CITATIONS
1	Ludwik Hirszfeld: A pioneer of transfusion and immunology during the world wars and beyond. Vox Sanguinis, 2022, 117, 467-475.	0.7	4
2	Treatment-induced hemophilic thrombosis?. Molecular Therapy, 2022, 30, 505-506.	3.7	7
3	Gene therapy – are we ready now?. Haemophilia, 2022, 28, 35-43.	1.0	5
4	One of the two N-glycans on the human Gb3/CD77 synthase is essential for its activity and allosterically regulates its function. Biochemical and Biophysical Research Communications, 2022, 617, 36-41.	1.0	3
5	Management of COVIDâ€19â€associated coagulopathy in persons with haemophilia. Haemophilia, 2021, 27, 41-48.	1.0	14
6	Human Gb3/CD77 synthase produces P1 glycotope-capped N-glycans, which mediate Shiga toxin 1 but not Shiga toxin 2 cell entry. Journal of Biological Chemistry, 2021, 296, 100299.	1.6	9
7	Vaccination against COVIDâ€19: Rationale, modalities and precautions for patients with haemophilia and other inherited bleeding disorders. Haemophilia, 2021, 27, 515-518.	1.0	9
8	B cell–activating factor modulates the factor VIII immune response in hemophilia A. Journal of Clinical Investigation, 2021, 131, .	3.9	10
9	Missing the sweet spot: one of the two N-glycans on human Gb3/CD77 synthase is expendable. Glycobiology, 2021, 31, 1145-1162.	1.3	1
10	Two Paralogous Gb3/CD77 Synthases in Birds Show Different Preferences for Their Glycoprotein and Glycosphingolipid Substrates. International Journal of Molecular Sciences, 2021, 22, 9761.	1.8	0
11	Helper T Cell Response to Factor VIII <i>In Vivo</i> Requires Several Anatomically Distinct Types of Antigen Presenting Cells. Blood, 2021, 138, 440-440.	0.6	0
12	Relationship between Endogenous, Transgene FVIII Expression and Bleeding Events Following Valoctocogene Roxaparvovec Gene Transfer for Severe Hemophilia A: A Post-Hoc Analysis of the GENEr8-1 Phase 3 Trial. Blood, 2021, 138, 3972-3972.	0.6	0
13	Factor IX Delivery to the Skin Primes Inhibitor Formation and Sensitizes Hemophilia B Mice to Systemic Factor IX Administration. Blood, 2021, 138, 3194-3194.	0.6	0
14	Eliminating Panglossian thinking in development of AAV therapeutics. Molecular Therapy, 2021, 29, 3325-3327.	3.7	12
15	A Molecular Revolution in the Treatment of Hemophilia. Molecular Therapy, 2020, 28, 997-1015.	3.7	66
16	Professor Elwira Lisowska Celebrates Her 90th Birthday. Archivum Immunologiae Et Therapiae Experimentalis, 2020, 68, 1.	1.0	2
17	Curing Hemophilia: Repeated Treatments versus a One-Off Fix. Molecular Therapy, 2020, 28, 1229-1230.	3.7	1
18	Gene therapy to cure haemophilia: Is robust scientific inquiry the missing factor?. Haemophilia, 2020, 26, 931-933.	1.0	24

2

RADOSLAW KACZMAREK

#	Article	IF	CITATIONS
19	How glycosylation affects glycosylation: the role of N-glycans in glycosyltransferase activity. Glycobiology, 2020, 30, 941-969.	1.3	37
20	Towards a global multidisciplinary consensus framework on haemophilia gene therapy: Report of the 2nd World Federation of Haemophilia Gene Therapy Round Table. Haemophilia, 2020, 26, 443-449.	1.0	15
21	Alternative Approaches to Oral Tolerance Induction to Factor FVIII. Blood, 2020, 136, 8-9.	0.6	Ο
22	Revisiting the "Danger Theory": Toll-like Receptor 9 Stimulation Triggers Activation of Conventional CD8α+ and Plasmacytoid Dendritic Cells <i>En Route</i> to Enhancing FVIII Inhibitor Formation. Blood, 2020, 136, 1-1.	0.6	1
23	Safety and efficacy of emicizumab and other novel agents inÂnewborns andÂinfants. Haemophilia, 2019, 25, e334-e335.	1.0	16
24	Erythrocyte glycophorins as receptors for Plasmodium merozoites. Parasites and Vectors, 2019, 12, 317.	1.0	43
25	The patient's view on rare disease trial design – a qualitative study. Orphanet Journal of Rare Diseases, 2019, 14, 31.	1.2	34
26	Toll-like Receptor 9 Activation Accelerates Inhibitor Formation in Response to Factor VIII. Blood, 2019, 134, 1113-1113.	0.6	0
27	The Gerbich blood group system: old knowledge, new importance. Transfusion Medicine Reviews, 2018, 32, 111-116.	0.9	16
28	The POWER-tool: Recommendations for involving patient representatives in choosing relevant outcome measures during rare disease clinical trial design. Health Policy, 2018, 122, 1287-1294.	1.4	11
29	Single nucleotide polymorphisms in A4GALT spur extra products of the human Gb3/CD77 synthase and underlie the P1PK blood group system. PLoS ONE, 2018, 13, e0196627.	1.1	11
30	RT-qPCR analysis of human melanoma progression-related genes – A novel workflow for selection and validation of candidate reference genes. International Journal of Biochemistry and Cell Biology, 2018, 101, 12-18.	1.2	6
31	Do adventitious viruses carried by insect cell lines producing <scp>AAV</scp> vectors pose a safety risk in gene therapy?. Haemophilia, 2018, 24, 843-844.	1.0	5
32	Hepatitis C and bleeding disorders in Europe. The Journal of Haemophilia Practice, 2018, 5, 50-65.	0.2	0
33	CD1: A Singed Cat of the Three Antigen Presentation Systems. Archivum Immunologiae Et Therapiae Experimentalis, 2017, 65, 201-214.	1.0	9
34	Plasmodium reichenowi EBA-140 merozoite ligand binds to glycophorin D on chimpanzee red blood cells, shedding new light on origins of Plasmodium falciparum. Parasites and Vectors, 2017, 10, 554.	1.0	6
35	Evaluation of an amino acid residue critical for the specificity and activity of human Gb3/CD77 synthase. Glycoconjugate Journal, 2016, 33, 963-973.	1.4	11
36	Baculovirus-expressed Plasmodium reichenowi EBA-140 merozoite ligand is host specific. Parasitology International, 2016, 65, 708-714.	0.6	5

#	Article	IF	CITATIONS
37	Human Gb3/CD77 synthase reveals specificity toward two or four different acceptors depending on amino acid at position 211, creating Pk, P1 and NOR blood group antigens. Biochemical and Biophysical Research Communications, 2016, 470, 168-174.	1.0	20
38	Can mutations in the gene encoding transcription factor EKLF (Erythroid Krüppel-Like Factor) protect us against infectious and parasitic diseases?. Postepy Higieny I Medycyny Doswiadczalnej, 2016, 70, 1068-1086.	0.1	1
39	The Baculovirus-Expressed Binding Region of Plasmodium falciparum EBA-140 Ligand and Its Glycophorin C Binding Specificity. PLoS ONE, 2015, 10, e0115437.	1.1	19
40	Studies of a Murine Monoclonal Antibody Directed against DARC: Reappraisal of Its Specificity. PLoS ONE, 2015, 10, e0116472.	1.1	6
41	P1PK, GLOB, and FORS Blood Group Systems and GLOB Collection: Biochemical and Clinical Aspects. Do We Understand It All Yet?. Transfusion Medicine Reviews, 2014, 28, 126-136.	0.9	33
42	Genetyczne podstawy syntezy cukrowych antygenów grupowych krwi. Acta Haematologica Polonica, 2013, 44, 251-259.	0.1	2
43	A Single Point Mutation in the Gene Encoding Gb3/CD77 Synthase Causes a Rare Inherited Polyagglutination Syndrome. Journal of Biological Chemistry, 2012, 287, 38220-38230.	1.6	40
44	Bacterially expressed truncated F2 domain of Plasmodium falciparum EBA-140 antigen can bind to human erythrocytes Acta Biochimica Polonica, 2012, 59, .	0.3	5
45	Bacterially expressed truncated F2 domain of Plasmodium falciparum EBA-140 antigen can bind to human erythrocytes. Acta Biochimica Polonica, 2012, 59, 685-91.	0.3	2