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List of Publications by Year
in descending order

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61 papers	2,185 citations	279487 23 h-index	233125 45 g-index
62 all docs	62 docs citations	62 times ranked	1581 citing authors

#	Article	IF	Citations
1	Accurate long-read sequencing allows assembly of the duplicated RHD and RHCE genes harboring variants relevant to blood transfusion. American Journal of Human Genetics, 2022, 109, 180-191.	2.6	11
2	An intron c.149â€2632T>A change in <scp>RHD</scp> is associated with aberrant transcription and very weak D phenotype. Transfusion, 2022, 62, .	0.8	1
3	First report of a null allele on a GYPB*s background: GYPB*s(37â€‰+â€‰4_8delAGTGA). Transfusion, 2022, 62, .	0.8	0
4	Impact of <scp>RHD</scp> genotyping on transfusion practice in <scp>Denmark</scp> and the <scp>United States</scp> and identification of novel <scp>RHD</scp> alleles. Transfusion, 2021, 61, 256-265.	0.8	9
5	ABO maternalâ€child discordance: Evidence of variable allelic expression and considerations for investigation. Transfusion, 2021, 61, 979-985.	0.8	0
6	A pair of <scp>S</scp>â€silencing single nucleotide variants cisâ€linked on <scp>GYPB</scp>. Transfusion, 2021, 61, E34-E36.	0.8	1
7	Screening of blood donors for sickle cell trait using a DNA â€based approach: Frequency in a multiethnic donor population. Transfusion, 2021, 61, 2008-2013.	0.8	5
8	Three new <scp>XK</scp> alleles; two associated with a <scp>McLeod RBC</scp> phenotype. Transfusion, 2021, 61, E69-E70.	0.8	3
9	Five novel silenced <scp>KEL</scp>*02 alleles. Transfusion, 2021, 61, E77-E79.	0.8	0
10	Use of an inâ€house trypsinâ€based method to resolve the interference of daratumumab. Transfusion, 2021, 61, 3000-3007.	0.8	1
11	An insertion/deletion polymorphism in the <scp>KLF1</scp> gene resulting in an In(Lu) phenotype. Transfusion, 2021, 61, E73-E74.	0.8	0
12	Five novel <scp>FY</scp> null alleles associated with typing discrepancies. Transfusion, 2021, 61, E80-E82.	0.8	0
13	A novel <scp>P1PK</scp> allele in two Bangladeshi sisters with a history of spontaneous abortion: <scp>A4GALT</scp>*<scp>02N</scp>(<scp>951C</scp>). Transfusion, 2021, 61, E71-E72.	0.8	1
14	How do I incorporate red cell genotyping to improve chronic transfusion therapy?. Transfusion, 2020, 60, 16-25.	0.8	7
15	A novel <scp>RHD*DAU</scp> allele with <scp>c.1136C>T (p.Thr379Met) and c.17C>T (p.) Tj ETQq1 1 0.784314 rgBT /Ov	0.8	0
16	Development and validation of a universal blood donor genotyping platform: a multinational prospective study. Blood Advances, 2020, 4, 3495-3506.	2.5	31
17	Multiple GYPB gene deletions associated with the Uâ^’ phenotype in those of African ancestry. Transfusion, 2020, 60, 1294-1307.	0.8	12
18	Overcoming the challenges of interpreting complex and uncommon RH alleles from whole genomes. Vox Sanguinis, 2020, 115, 790-801.	0.7	5

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19	It's time to phase out â€œserologic weak D phenotypeâ€•and resolve D types with RHD genotyping including weak D type 4. Transfusion, 2020, 60, 855-859.	0.8	27
20	Reliability of labeling red cell units with minor antigen historical results and process considerations. Transfusion, 2020, 60, 822-830.	0.8	3
21	<scp>RHCE*02</scp> (<scp>c.148G>A</scp>, <scp>p.Val50Ile</scp>) allele with silenced <scp>RHCE*Ce</scp> expression. Transfusion, 2020, 60, E23-E24.	0.8	0
22	American Society of Hematology 2020 guidelines for sickle cell disease: transfusion support. Blood Advances, 2020, 4, 327-355.	2.5	241
23	Automated typing of red blood cell and platelet antigens from whole exome sequences. Transfusion, 2019, 59, 3253-3263.	0.8	32
24	Blood group genotyping. Blood, 2019, 133, 1814-1820.	0.6	63
25	Molecular DNAâ€“Based Blood Group Typing. , 2019, , 131-138.		1
26	International Society of Blood Transfusion Working Party on Red Cell Immunogenetics and Blood Group Terminology: Report of the Dubai, Copenhagen and Toronto meetings. Vox Sanguinis, 2019, 114, 95-102.	0.7	75
27	Development and evaluation of a transfusion medicine genome wide genotyping array. Transfusion, 2019, 59, 101-111.	0.8	30
28	Monoclonal antiâ€CD47 interference in red cell and platelet testing. Transfusion, 2019, 59, 730-737.	0.8	65
29	Experience with RHD*weak D type 4.0 in the USA. Blood Transfusion, 2019, 17, 91-93.	0.3	10
30	The â€œnext generationâ€•reference laboratory?. Transfusion, 2018, 58, 277-279.	0.8	10
31	Novel mutations in KLF1 encoding the In(Lu) phenotype reflect a diversity of clinical presentations. Transfusion, 2018, 58, 196-199.	0.8	9
32	Automated typing of red blood cell and platelet antigens: a whole-genome sequencing study. Lancet Haematology,the, 2018, 5, e241-e251.	2.2	70
33	RH genotype matching for transfusion support in sickle cell disease. Blood, 2018, 132, 1198-1207.	0.6	79
34	Application of genomics for transfusion therapy in sickle cell anemia. Blood Cells, Molecules, and Diseases, 2017, 67, 148-154.	0.6	12
35	Whole-exome sequencing for RH genotyping and alloimmunization risk in children with sickle cell anemia. Blood Advances, 2017, 1, 1414-1422.	2.5	64
36	Rh and LW blood group antigens. , 2016, , 176-184.		2

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37	Comprehensive red blood cell and platelet antigen prediction from whole genome sequencing: proof of principle. Transfusion, 2016, 56, 743-754.	0.8	81
38	RHD Zygosity Determination from Whole Genome Sequencing Data. Journal of Blood Disorders & Transfusion, 2016, 7, .	0.1	13
39	RHCE*ceAG (254C>G, Ala85Gly) is prevalent in blacks, encodes a partial ceâ€phenotype, and is associated with discordant RHD zygosity. Transfusion, 2015, 55, 2624-2632.	0.8	17
40	Financial implications of RHD genotyping of pregnant women with a serologic weak D phenotype. Transfusion, 2015, 55, 2095-2103.	0.8	47
41	How do I manage Rh typing in obstetric patients?. Transfusion, 2015, 55, 470-474.	0.8	21
42	It's time to phase in <scp>RHD</scp> genotyping for patients with a serologic weak <scp>D</scp> phenotype. Transfusion, 2015, 55, 680-689.	0.8	157
43	RHCE*ceTI encodes partial c and partial e and is often in cis to RHD*DIVa. Transfusion, 2013, 53, 741-746.	0.8	19
44	Erythroid transcription factor EKLF/KLF1 mutation causing congenital dyserythropoietic anemia type IV in a patient of Taiwanese origin: Review of all reported cases and development of a clinical diagnostic paradigm. Blood Cells, Molecules, and Diseases, 2013, 51, 71-75.	0.6	68
45	<scp>RHCE*ceMO</scp> is frequently in cis to <scp>RHD*DAU0</scp> and encodes a <scp>hr^S</scp>â€“, <scp>hr^B</scp>â€“, <scp>RH</scp>:â€“61 phenotype in black persons: clinical significance. Transfusion, 2013, 53, 2983-2989.	0.8	23
46	High prevalence of red blood cell alloimmunization in sickle cell disease despite transfusion from Rh-matched minority donors. Blood, 2013, 122, 1062-1071.	0.6	394
47	Nucleotide deletion in RHCE*cE (907delC) is responsible for a D-â€ƒ- haplotype in Hispanics. Transfusion, 2011, 51, 2142-2147.	0.8	12
48	DIIIa and DIII Type 5 are encoded by the same allele and are associated with altered RHCE*ce alleles: clinical implications. Transfusion, 2010, 50, 1303-1311.	0.8	53
49	The Rh and RhAG blood group systems. Immunohematology, 2010, 26, 178-186.	0.2	29
50	Molecular biology of the Rh system: clinical considerations for transfusion in sickle cell disease. Hematology American Society of Hematology Education Program, 2009, 2009, 178-184.	0.9	37
51	Molecular Genotyping in Transfusion Medicine. Clinical Chemistry, 2008, 54, 1948-1950.	1.5	17
52	The Structure and Function of the Rh Antigen Complex. Seminars in Hematology, 2007, 44, 42-50.	1.8	127
53	Rh complexities: serology and DNA genotyping. Transfusion, 2007, 47, 17S-22S.	0.8	35
54	Deciphering the function of the Rh family of proteins. Transfusion, 2005, 45, 117S-121S.	0.8	7

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55	Review: the Rh blood group D antigen . . . dominant, diverse, and difficult. Immunohematology, 2005, 21, 155-163.	0.2	22
56	A new hybrid RHCE gene (CeNR) is responsible for expression of a novel antigen. Transfusion, 2004, 44, 1047-1051.	0.8	13
57	Protein 4.2 is critical to CD47-membrane skeleton attachment in human red cells. Blood, 2004, 103, 1131-1136.	0.6	44
58	Ammonium Transport by the Erythrocyte Rh-Associated Protein.. Blood, 2004, 104, 1594-1594.	0.6	0
59	Fractional attachment of CD47 (IAP) to the erythrocyte cytoskeleton and visual colocalization with Rh protein complexes. Blood, 2003, 101, 1194-1199.	0.6	49
60	DNA analysis for donor screening of Dombrock blood group antigens. Immunohematology, 2003, 19, 73-76.	0.2	16
61	Rh and LW Blood Group Antigens. , 0, , 109-120.		3