Martin L Olsson

List of Publications by Year
in descending order

156 papers	5,596 citations	50276 46 h-index	98798 67 g-index
164 all docs	164 docs citations	164 times ranked	4507 citing authors

#	Article	IF	Citations
1	Recommendation for validation and quality assurance of nonâ€invasive prenatal testing for foetal blood groups and implications for <scp>IVD</scp> risk classification according to <scp>EU</scp> regulations. Vox Sanguinis, 2022, 117, 157-165.	1.5	7
2	The EHA Research Roadmap: Transfusion Medicine. HemaSphere, 2022, 6, e670.	2.7	2
3	Novel or not? Reference alleles, genes, and genomes to unmask the true nature of the ABO*AW.10 allele associated with weak A phenotype. Transfusion, 2022, 62, 721-724.	1.6	2
4	Glycoproteomic and Phenotypic Elucidation of B4GALNT2 Expression Variants in the SID Histo-Blood Group System. International Journal of Molecular Sciences, 2022, 23, 3936.	4.1	4
5	Novel RHD variant causing <scp>RhD</scp> negative phenotype identified in a pregnant woman. Transfusion, 2022, 62, .	1.6	0
6	Human radical scavenger Î±1-microglobulin protects against hemolysis in vitro and Î±1-microglobulin knockout mice exhibit a macrocytic anemia phenotype. Free Radical Biology and Medicine, 2021, 162, 149-159.	2.9	19
7	Rh disease prevention: the European Perspective. ISBT Science Series, 2021, 16, 106-118.	1.1	6
8	An agnostic study of associations between ABO and RhD blood group and phenome-wide disease risk. ELife, 2021, 10, .	6.0	25
9	Platelets inhibit erythrocyte invasion by Plasmodium falciparum at physiological platelet:erythrocyte ratios. Transfusion Medicine, 2021, , .	1.1	0
10	Network pharmacology of triptolide in cancer cells: implications for transcription factor binding. Investigational New Drugs, 2021, 39, 1523-1537.	2.6	7
11	Dimerization of small integral membrane protein 1 promotes cell surface presentation of the Vel blood group epitope. FEBS Letters, 2020, 594, 1261-1270.	2.8	2
12	Disruption of the tumour-associated EMP3 enhances erythroid proliferation and causes the MAM-negative phenotype. Nature Communications, 2020, 11, 3569.	12.8	18
13	A novel ABO allele with a 21â€bp duplication identified in two unrelated European individuals with weak A expression. Transfusion Medicine, 2020, 30, 508-512.	1.1	0
14	Characterization of GYP*Mur and novel GYP*Bunâ€like hybrids in Thai blood donors reveals a qualitatively altered s antigen. Vox Sanguinis, 2020, 115, 472-477.	1.5	7
15	SMIM1, carrier of the Vel blood group, is a tail-anchored transmembrane protein and readily forms homodimers in a cell-free system. Bioscience Reports, 2020, 40, .	2.4	3
16	The Xg blood group system: no longer forgotten. Immunohematology, 2020, 36, 4-6.	0.2	0
17	May the FORS be with you: a system sequel. Immunohematology, 2020, 36, 14-18.	0.2	1
18	The P1PK blood group system: revisited and resolved. Immunohematology, 2020, 36, 99-103.	0.2	0

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19	Multiple miscarriages in two sisters of Thai origin with the rare P^k phenotype caused by a novel nonsense mutation at the B3GALNT1 locus. Transfusion Medicine, 2019, 29, 202-208.	1.1	6
20	Towards universally acceptable blood. Nature Microbiology, 2019, 4, 1426-1427.	13.3	2
21	A novel singleâ€nucleotide substitution in the proximal ABO promoter gives rise to the B₃ phenotype. Transfusion, 2019, 59, E1-E3.	1.6	6
22	Missense mutations in the C-terminal portion of the B4GALNT2-encoded glycosyltransferase underlying the Sd(aâ^') phenotype. Biochemistry and Biophysics Reports, 2019, 19, 100659.	1.3	19
23	Automatic for the people: a rapidly evolving movement for the future of genotyping. Transfusion, 2019, 59, 3545-3547.	1.6	1
24	ABO blood type and risk of porcine bioprosthetic aortic valve degeneration: SWEDEHEART observational cohort study. BMJ Open, 2019, 9, e029109.	1.9	4
25	Validated Reference Panel from Renewable Source of Genomic DNA Available for Standardization of Blood Group Genotyping. Journal of Molecular Diagnostics, 2019, 21, 525-537.	2.8	7
26	Noninvasive fetal <scp>RHD</scp> genotyping to guide targeted antiâ€D prophylaxisâ€“an external quality assessment workshop. Vox Sanguinis, 2019, 114, 386-393.	1.5	14
27	A large deletion spanning XG and GYG2 constitutes a genetic basis of the Xg_{null} phenotype, underlying antiâ€Xg^a production. Transfusion, 2019, 59, 1843-1849.	1.6	4
28	The P1 histoâ€blood group antigen is present on human red blood cell glycoproteins. Transfusion, 2019, 59, 1108-1117.	1.6	10
29	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.	1.5	75
30	Matrixâ€assisted laser desorption/ionization timeâ€ofâ€flight mass spectrometry analysis of 36 blood group alleles among 396 T hai samples reveals regionâ€specific variants. Transfusion, 2018, 58, 1752-1762.	1.6	12
31	Identification of human glycosyltransferase genes expressed in erythroid cells predicts potential carbohydrate blood group loci. Scientific Reports, 2018, 8, 6040.	3.3	13
32	Allele-selective RUNX1 binding regulates P1 blood group status by transcriptional control of A4GALT. Blood, 2018, 131, 1611-1616.	1.4	24
33	Vox Sanguinis International Forum on application of fetal blood grouping. Vox Sanguinis, 2018, 113, e26-e35.	1.5	8
34	ABO zygosity, but not secretor or Fc receptor status, is a significant risk factor for IVIG-associated hemolysis. Blood, 2018, 131, 830-835.	1.4	19
35	Vox Sanguinis International Forum on application of fetal blood grouping: summary. Vox Sanguinis, 2018, 113, 198-201.	1.5	10
36	Thorough analysis of unorthodox <scp>ABO</scp> deletions called by the 1000 Genomes project. Vox Sanguinis, 2018, 113, 185-197.	1.5	13

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37	GBGT1 is allelically diverse but dispensable in humans and naturally occurring antiâ€FORS1 shows an ABOâ€restricted pattern. Transfusion, 2018, 58, 2036-2045.	1.6	9
38	Disruption of a GATA1-binding motif upstream of XG/PBDX abolishes Xga expression and resolves the Xg blood group system. Blood, 2018, 132, 334-338.	1.4	26
39	An update on the GLOB blood group system (and former GLOB collection). Immunohematology, 2018, 34, 161-163.	0.2	5
40	FUT1 mutations responsible for the H-deficient phenotype in the Polish population, including the first example of an abolished start codon. Blood Transfusion, 2018, 16, 101-104.	0.4	3
41	An update on the GLOB blood group system (and former GLOB collection). Immunohematology, 2018, 34, 161-163.	0.2	2
42	SMIM1 variants rs1175550 and rs143702418 independently modulate Vel blood group antigen expression. Scientific Reports, 2017, 7, 40451.	3.3	16
43	A and B antigen levels acquired by group O donorâ€derived erythrocytes following <scp>ABO</scp>â€nonâ€identical transfusion or minor <scp>ABO</scp>â€incompatible haematopoietic stem cell transplantation. Transfusion Medicine, 2017, 27, 181-191.	1.1	16
44	The FORS awakens: review of a blood group system reborn. Immunohematology, 2017, 33, 64-72.	0.2	9
45	The FORS awakens: review of a blood group system reborn. Immunohematology, 2017, 33, 64-72.	0.2	5
46	Erythrogene: a database for in-depth analysis of the extensive variation in 36 blood group systems in the 1000 Genomes Project. Blood Advances, 2016, 1, 240-249.	5.2	122
47	International society of blood transfusion working party on red cell immunogenetics and terminology: report of the Seoul and London meetings. ISBT Science Series, 2016, 11, 118-122.	1.1	56
48	Sex-Discordant Blood Transfusions and Survival After Cardiac Surgery. Circulation, 2016, 134, 1692-1694.	1.6	12
49	An ageâ€dependent <scp>ABO</scp> discrepancy between mother and baby reveals a novel <scp>A</scp>^{weak} allele. Transfusion, 2015, 55, 422-426.	1.6	9
50	RIFINs are adhesins implicated in severe Plasmodium falciparum malaria. Nature Medicine, 2015, 21, 314-317.	30.7	166
51	Shiga Toxinâ€"Induced Complement-Mediated Hemolysis and Release of Complement-Coated Red Blood Cellâ€"Derived Microvesicles in Hemolytic Uremic Syndrome. Journal of Immunology, 2015, 194, 2309-2318.	0.8	65
52	Identification of the Molecular and Genetic Basis of PX2, a Glycosphingolipid Blood Group Antigen Lacking on Globoside-deficient Erythrocytes. Journal of Biological Chemistry, 2015, 290, 18505-18518.	3.4	20
53	Do ABO Blood Group Antigens Hamper the Therapeutic Efficacy of Mesenchymal Stromal Cells?. PLoS ONE, 2014, 9, e85040.	2.5	61
54	Allelic variants of <scp>PRDM</scp>9 associated with high hyperdiploid childhood acute lymphoblastic leukaemia. British Journal of Haematology, 2014, 166, 947-949.	2.5	12

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55	Large deletions involving the regulatory upstream regions of <scp>A</scp>4<scp>GALT</scp> give rise to principally novel <scp>P</scp>1<scp>PK</scp>â€null alleles. Transfusion, 2014, 54, 1831-1835.	1.6	8
56	International Society of Blood Transfusion Working Party on red cell immunogenetics and blood group terminology: Cancun report (2012). Vox Sanguinis, 2014, 107, 90-96.	1.5	69
57	Mixed field reactions in ABO and Rh typing chimerism likely resulting from twin haematopoiesis. Blood Transfusion, 2014, 12, 608-10.	0.4	8
58	<scp>P</scp>¹/<scp>P</scp>² genotyping of known and novel null alleles in the <scp>P</scp>1<scp>PK</scp> and <scp>GLOB</scp> histoâ€blood group systems. Transfusion, 2013, 53, 2928-2939.	1.6	12
59	Two cases of hemolytic uremic syndrome caused by Streptococcus pneumoniae serotype 3, one being a vaccine failure. Scandinavian Journal of Infectious Diseases, 2013, 45, 411-414.	1.5	11
60	Forssman expression on human erythrocytes: biochemical and genetic evidence of a new histo-blood group system. Blood, 2013, 121, 1459-1468.	1.4	60
61	Homozygosity for a null allele of SMIM1 defines the Vel-negative blood group phenotype. Nature Genetics, 2013, 45, 537-541.	21.4	75
62	P1PK: The blood group system that changed its name and expanded. Immunohematology, 2013, 29, 25-33.	0.2	31
63	Allelic Forms Of PRDM9 Associated With High Hyperdiploid Childhood Acute Lymphoblastic Leukaemia. Blood, 2013, 122, 1351-1351.	1.4	0
64	An update on the GLOB blood group system and collection. Immunohematology, 2013, 29, 19-24.	0.2	8
65	An update on the GLOB blood group system and collection. Immunohematology, 2013, 29, 19-24.	0.2	4
66	P1PK: the blood group system that changed its name and expanded. Immunohematology, 2013, 29, 25-33.	0.2	20
67	ABO Blood Groups Influence Macrophage-mediated Phagocytosis of Plasmodium falciparum-infected Erythrocytes. PLoS Pathogens, 2012, 8, e1002942.	4.7	39
68	H Blood Group System. , 2012, , 489-498.		32
69	Globoside Blood Group System. , 2012, , 609-613.		1
70	Noninvasive Single-Exon Fetal RHD Determination in a Routine Screening Program in Early Pregnancy. Obstetrics and Gynecology, 2012, 120, 227-234.	2.4	91
71	Pathological Conditions Involving Extracellular Hemoglobin: Molecular Mechanisms, Clinical Significance, and Novel Therapeutic Opportunities for Î±₁-Microglobulin. Antioxidants and Redox Signaling, 2012, 17, 813-846.	5.4	87
72	Flow cytometry evaluation of red blood cells mimicking naturally occurring ABO subgroups after modification with variable amounts of functionâ€spacerâ€lipid A and B constructs. Transfusion, 2012, 52, 247-251.	1.6	16

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73	A novel B^{weak} hybrid allele lacks three enhancer repeats but generates normal ABO transcript levels. Vox Sanguinis, 2012, 102, 55-64.	1.5	13
74	Identification of a novel A4GALT exon reveals the genetic basis of the P1/P2 histo-blood groups. Blood, 2011, 117, 678-687.	1.4	56
75	International Society of Blood Transfusion Working Party on red cell immunogenetics and blood group terminology: Berlin report. Vox Sanguinis, 2011, 101, 77-82.	1.5	75
76	The IgG-specific endoglycosidase EndoS inhibits both cellular and complement-mediated autoimmune hemolysis. Blood, 2010, 115, 5080-5088.	1.4	66
77	Increased levels of cell-free hemoglobin, oxidation markers, and the antioxidative heme scavenger Î±1-microglobulin in preeclampsia. Free Radical Biology and Medicine, 2010, 48, 284-291.	2.9	87
78	Many genetically defined ABO subgroups exhibit characteristic flow cytometric patterns. Transfusion, 2010, 50, 308-323.	1.6	47
79	Weak A phenotypes associated with novel ABO alleles carrying the A²â€related 1061C deletion and various missense substitutions. Transfusion, 2010, 50, 1471-1486.	1.6	38
80	KEL*02 alleles with alterations in and around exon 8 in individuals with apparent KEL:1,â˜2 phenotypes. Vox Sanguinis, 2010, 99, 150-157.	1.5	5
81	Fondaparinux or enoxaparin: A comparative study of postoperative bleeding in coronary artery bypass grafting surgery. Scandinavian Cardiovascular Journal, 2010, 44, 100-106.	1.2	18
82	The Bloodgen Project of the European Union, 2003–2009. Transfusion Medicine and Hemotherapy, 2009, 36, 162-167.	1.6	48
83	International Society of Blood Transfusion Committee on Terminology for Red Blood Cell Surface Antigens: Macao report. Vox Sanguinis, 2009, 96, 153-156.	1.5	65
84	A novel mutation in the complement regulator clusterin in recurrent hemolytic uremic syndrome. Molecular Immunology, 2009, 46, 2236-2243.	2.2	39
85	Will Genotyping Replace Serology in Future Routine Blood Grouping? – Opinion 4. Transfusion Medicine and Hemotherapy, 2009, 36, 232-233.	1.6	8
86	Transfusion medicine: a target on the move. Current Opinion in Hematology, 2009, 16, 466.	2.5	0
87	The human Pk histo-blood group antigen provides protection against HIV-1 infection. Blood, 2009, 113, 4980-4991.	1.4	69
88	The ABO blood group system revisited: a review and update. Immunohematology, 2009, 25, 48-59.	0.2	221
89	The ABO blood group system revisited: a review and update. Immunohematology, 2009, 25, 48-59.	0.2	156
90	A simple screening assay for the most common JK*0 alleles revealed compound heterozygosity in Jk(a-b-) probands from Guam. Immunohematology, 2009, 25, 165-9.	0.2	2

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91	Modifying the red cell surface: towards an ABOâ€universal blood supply. British Journal of Haematology, 2008, 140, 3-12.	2.5	46
92	Expression of a novel missense mutation found in the A4GALT gene of Amish individuals with the pâ€ƒphenotype. Transfusion, 2008, 48, 479-487.	1.6	23
93	Investigation into Aâ€ƒantigen expression on O² heterozygous groupâ€ƒOâ€“labeled red blood cell units. Transfusion, 2008, 48, 1650-1657.	1.6	11
94	Identification of a GH110 Subfamily of Î±1,3-Galactosidases. Journal of Biological Chemistry, 2008, 283, 8545-8554.	3.4	52
95	Blood grouping discrepancies between ABO genotype and phenotype caused by O alleles. Current Opinion in Hematology, 2008, 15, 618-624.	2.5	26
96	The O² allele: questioning the phenotypic definition of an ABO allele. Immunohematology, 2008, 24, 138-147.	0.2	6
97	The O2 allele: questioning the phenotypic definition of an ABO allele. Immunohematology, 2008, 24, 138-47.	0.2	4
98	Structural Effects of Naturally Occurring Human Blood Group B Galactosyltransferase Mutations Adjacent to the DXD Motif. Journal of Biological Chemistry, 2007, 282, 9564-9570.	3.4	19
99	Quantitative and qualitative evaluation of plasma and urine Î±1-microglobulin in healthy donors and patients with different haemolytic disorders and haemochromatosis. Clinica Chimica Acta, 2007, 386, 31-37.	1.1	16
100	Bacterial glycosidases for the production of universal red blood cells. Nature Biotechnology, 2007, 25, 454-464.	17.5	259
101	International Society of Blood Transfusion Committee on Terminology for Red Cell Surface Antigens: Cape Town report. Vox Sanguinis, 2007, 92, 250-253.	1.5	56
102	Structural basis for red cell phenotypic changes in newly identified, naturally occurring subgroup mutants of the human blood group B glycosyltransferase. Transfusion, 2007, 47, 864-875.	1.6	28
103	An extensive polymerase chain reactionâ€“alleleâ€specific polymorphism strategy for clinical ABO blood group genotyping that avoids potential errors caused by null, subgroup, and hybrid alleles. Transfusion, 2007, 47, 2110-2125.	1.6	53
104	The Development of Severe Anti-HPA 1a-Related Neonatal Alloimmune Thrombocytopenia Is Influenced by the Maternal ABO Type.. Blood, 2007, 110, 2093-2093.	1.4	2
105	SI14 Stripped Red Cells?Efforts to Eliminate A and B Antigens from the Blood Supply. Transfusion Medicine, 2006, 16, 9-9.	1.1	0
106	The cis-AB Blood Group Phenotype: Fundamental Lessons in Glycobiology. Transfusion Medicine Reviews, 2006, 20, 207-217.	2.0	28
107	Heterozygosity for the Non-Deletional O2 Allele Does Not Cause Discrepancies in Automated Blood Donor ABO Grouping.. Blood, 2006, 108, 957-957.	1.4	1
108	ADP Acting on P2Y₁₃Receptors Is a Negative Feedback Pathway for ATP Release From Human Red Blood Cells. Circulation Research, 2005, 96, 189-196.	4.5	122

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109	New and unusual O alleles at the ABO locus are implicated in unexpected blood group phenotypes. Transfusion, 2005, 45, 70-81.	1.6	47
110	Blood group genotype analysis for the quality improvement of reagent test red blood cells. Vox Sanguinis, 2005, 88, 265-270.	1.5	24
111	The Abantu phenotype in the ABO blood group system is due to a splice-site mutation in a hybrid between a new O1-like allelic lineage and the A2 allele. Vox Sanguinis, 2005, 88, 256-264.	1.5	19
112	Novel glycolipid variations revealed by monoclonal antibody immunochemical analysis of weak ABO subgroups of A. Vox Sanguinis, 2005, 89, 27-38.	1.5	16
113	A clue to the basis of allelic enhancement: occurrence of the Ax subgroup in the offspring of blood group O parents. Transfusion Medicine, 2005, 15, 435-442.	1.1	21
114	Two previously proposed P1/P2-differentiating and nine novel polymorphisms at the A4GALT (Pk) locus do not correlate with the presence of the P1 blood group antigen. BMC Genetics, 2005, 6, 49.	2.7	19
115	The Levels of Blood Group ABO Transcripts in Peripheral Blood Are Allele-Dependent but Not Correlated to Variations in the 5â€²-Regulatory Region.. Blood, 2005, 106, 1897-1897.	1.4	1
116	Novel Mechansims for Pk (Gb3) Inhibition of HIV Infection.. Blood, 2005, 106, 1430-1430.	1.4	0
117	Transcriptional Profile of Carbohydrate Blood Group Genes in Erythroid Versus Neutrophil Differentiation of Human CD34+ Cells In Vitro.. Blood, 2005, 106, 4242-4242.	1.4	0
118	Low Antibody Levels against Cell Wallâ€“Attached Proteins ofStreptococcus pyogenesPredispose for Severe Invasive Disease. Journal of Infectious Diseases, 2004, 189, 797-804.	4.0	53
119	Genetic heterogeneity at the glycosyltransferase loci underlying the GLOB blood group system and collection*. British Journal of Haematology, 2004, 125, 528-536.	2.5	25
120	Genetic basis of blood group diversity. British Journal of Haematology, 2004, 126, 759-771.	2.5	48
121	Clonogenicity, gene expression and phenotype during neutrophil versus erythroid differentiation of cytokine-stimulated CD34+ human marrow cells in vitro. British Journal of Haematology, 2004, 127, 451-463.	2.5	32
122	A novel blood group B subgroup: serological and genetic studies. Transfusion Medicine, 2004, 14, 173-180.	1.1	8
123	Autologous del(20q)-positive erythroid progenitor cells, re-emerging after DLI treatment of an MDS patient relapsing after allo-SCT, can provide a normal peripheral red blood cell count. Bone Marrow Transplantation, 2004, 33, 559-563.	2.4	2
124	Universal red blood cellsâ€”enzymatic conversion of blood group A and B antigens. Transfusion Clinique Et Biologique, 2004, 11, 33-39.	0.4	54
125	ATP Release from Red Blood Cells Is Regulated by a Negative Feedback Pathway where ADP Acts on P2Y13 Receptors.. Blood, 2004, 104, 1576-1576.	1.4	0
126	Additional molecular bases of the clinically important p blood group phenotype. Transfusion, 2003, 43, 899-907.	1.6	33

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127	ABO exon and intron analysis in individuals with the AweakB phenotype reveals a novel O1v-A2 hybrid allele that causes four missense mutations in the A transferase. BMC Genetics, 2003, 4, 17.	2.7	49
128	Processing of the lipocalin Î±1-microglobulin by hemoglobin induces heme-binding and heme-degradation properties. Blood, 2002, 99, 1894-1901.	1.4	116
129	Molecular Basis of the Globoside-deficient Pk Blood Group Phenotype. Journal of Biological Chemistry, 2002, 277, 29455-29459.	3.4	65
130	Novel alleles at the JK blood group locus explain the absence of the erythrocyte urea transporter in European families*. British Journal of Haematology, 2002, 116, 445-453.	2.5	38
131	Phenotype prediction by DNA-based typing of clinically significant blood group systems in Jordanian blood donors. Vox Sanguinis, 2002, 83, 55-62.	1.5	10
132	Platelet concentrate increases bone ingrowth into porous hydroxyapatite. Orthopedics, 2002, 25, 169-72.	1.1	31
133	Intercellular adhesion molecule-4 binds Î±4Î²1 and Î±V-family integrins through novel integrin-binding mechanisms. Blood, 2001, 98, 458-466.	1.4	96
134	Genomic analysis of clinical samples with serologic ABO blood grouping discrepancies: identification of 15 novel A and B subgroup alleles. Blood, 2001, 98, 1585-1593.	1.4	176
135	Polymorphism and recombination events at the ABO locus: a major challenge for genomic ABO blood grouping strategies. Transfusion Medicine, 2001, 11, 295-313.	1.1	112
136	The ABO blood group gene: A locus of considerable genetic diversity. Transfusion Medicine Reviews, 2001, 15, 177-200.	2.0	85
137	Allele-related variation in minisatellite repeats involved in the transcription of the blood group ABO gene. Transfusion Medicine, 1999, 9, 219-226.	1.1	43
138	Heterogeneity of the blood group Axallele: genetic recombination of common alleles can result in the Axphenotype. Transfusion Medicine, 1998, 8, 231-238.	1.1	55
139	Heterogeneity of the O Alleles at the Blood Group ABO Locus in Amerindians. Vox Sanguinis, 1998, 74, 46-50.	1.5	45
140	Different Genotypes Causing Indiscernible Patterns of A Expression on A el Red Blood Cells as Visualized by Scanning Immunogold Electron Microscopy. Vox Sanguinis, 1998, 75, 47-51.	1.5	14
141	A clinically applicable method for determining the three major alleles at the Duffy (FY) blood group locus using polymerase chain reaction with allele-specific primers. Transfusion, 1998, 38, 168-173.	1.6	82
142	Genomic typing of the Kidd blood group locus by a singleâ€tube alleleâ€specific primer PCR technique. British Journal of Haematology, 1998, 102, 1010-1014.	2.5	36
143	The Fyxphenotype is associated with a missense mutation in theFyballele predicting Arg89Cys in the Duffy glycoprotein. British Journal of Haematology, 1998, 103, 1184-1191.	2.5	87
144	Heterogeneity of the O Alleles at the Blood Group ABO Locus in Amerindians. Vox Sanguinis, 1998, 74, 46-50.	1.5	3

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145	Heterogeneity of the O alleles at the blood group ABO locus in Amerindians. Vox Sanguinis, 1998, 74, 46-50.	1.5	5
146	Different genotypes causing indiscernible patterns of A expression on A(el) red blood cells as visualized by scanning immunogold electron microscopy. Vox Sanguinis, 1998, 75, 47-51.	1.5	1
147	Molecular Analysis of theOAlleles at the Blood Group ABO Locus in Populations of Different Ethnic Origin Reveals Novel Crossing-Over Events and Point Mutations. Biochemical and Biophysical Research Communications, 1997, 234, 779-782.	2.1	50
148	Evidence for a New Type of O Allele at the ABO Locus, due to a Combination of the A^2 Nucleotide Deletion and the A^el Nucleotide Insertion. Vox Sanguinis, 1996, 71, 113-117.	1.5	27
149	Evidence for a New Type of O Allele at the ABO Locus, due to a Combination of the A2Nucleotide Deletion and the AelNucleotide Insertion. Vox Sanguinis, 1996, 71, 113-117.	1.5	37
150	Frequent Occurrence of a Variant O¹ Gene at the Blood Group ABO Locus. Vox Sanguinis, 1996, 70, 26-30.	1.5	64
151	Polymorphisms at the ABO locus in subgroup A individuals. Transfusion, 1996, 36, 309-313.	1.6	51
152	A Rapid and Simple ABO Genotype Screening Method Using a Novel B/O2versus A/O2Discriminating Nucleotide Substitution at the ABO Locus. Vox Sanguinis, 1995, 69, 242-247.	1.5	74
153	A Rapid and Simple ABO Genotype Screening Method Using a Novel B/O^2 versus A/O^2 Discriminating Nucleotide Substitution at the ABO Locus. Vox Sanguinis, 1995, 69, 242-247.	1.5	71
154	An Ael Allele-Specific Nucleotide Insertion at the Blood Group ABO Locus and Its Detection Using a Sequence-Specific Polymerase Chain Reaction. Biochemical and Biophysical Research Communications, 1995, 216, 642-647.	2.1	57
155	Characterization of Monoclonal Anti-alpha1-Microglobulin Antibodies: Binding Strength, Binding Sites, and Inhibition of Lymphocyte Stimulation. Scandinavian Journal of Immunology, 1991, 34, 655-666.	2.7	16
156	Alpha1-Microglobulin is Mitogenic to Human Peripheral Blood Lymphocytes. Regulation by both Enhancing and Suppressive Serum Factors. Immunobiology, 1990, 180, 221-234.	1.9	21