C Ellen Van Der Schoot

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/11087250/publications.pdf

Version: 2024-02-01

114 papers 7,044 citations

71102 41 h-index 80 g-index

119 all docs

119 docs citations

119 times ranked

11920 citing authors

#	Article	IF	CITATIONS
1	Large-scale genotyping identifies 41 new loci associated with breast cancer risk. Nature Genetics, 2013, 45, 353-361.	21.4	960
2	Multiple independent variants at the TERT locus are associated with telomere length and risks of breast and ovarian cancer. Nature Genetics, 2013, 45, 371-384.	21.4	493
3	New gene functions in megakaryopoiesis and platelet formation. Nature, 2011, 480, 201-208.	27.8	401
4	The Pi-linked receptor FcRIII is released on stimulation of neutrophils. Nature, 1988, 333, 667-669.	27.8	395
5	Afucosylated IgG characterizes enveloped viral responses and correlates with COVID-19 severity. Science, 2021, 371, .	12.6	244
6	Competition for FcRn-mediated transport gives rise to short half-life of human IgG3 and offers therapeutic potential. Nature Communications, 2011, 2, 599.	12.8	220
7	A HaemAtlas: characterizing gene expression in differentiated human blood cells. Blood, 2009, 113, e1-e9.	1.4	215
8	A prominent lack of IgG1-Fc fucosylation of platelet alloantibodies in pregnancy. Blood, 2014, 123, 471-480.	1.4	187
9	Comparative gene expression profiling of in vitro differentiated megakaryocytes and erythroblasts identifies novel activatory and inhibitory platelet membrane proteins. Blood, 2007, 109, 3260-3269.	1.4	153
10	Prenatal typing of Rh and kell blood group system antigens: The edge of a watershed. Transfusion Medicine Reviews, 2003, 17, 31-44.	2.0	119
11	Low antiâ€ <scp>R</scp> h <scp>D I</scp> g <scp>G</scp> â€ <scp>F</scp> câ€fucosylation in pregnancy: a new variable predicting severity in haemolytic disease of the fetus and newborn. British Journal of Haematology, 2014, 166, 936-945.	2.5	109
12	A proinflammatory monocyte response is associated with myocardial injury and impaired functional outcome in patients with ST-segment elevation myocardial infarction. American Heart Journal, 2012, 163, 57-65.e2.	2.7	103
13	Rapid genotyping of blood group antigens by multiplex polymerase chain reaction and DNA microarray hybridization. Transfusion, 2005, 45, 667-679.	1.6	100
14	SMIM1 underlies the Vel blood group and influences red blood cell traits. Nature Genetics, 2013, 45, 542-545.	21.4	96
15	Noninvasive Prenatal Diagnosis of Fetal Rhesus D. Obstetrics and Gynecology, 2005, 106, 841-844.	2.4	95
16	Migration of Human Hematopoietic Progenitor Cells Across Bone Marrow Endothelium Is Regulated by Vascular Endothelial Cadherin. Journal of Immunology, 2002, 168, 588-596.	0.8	93
17	The composition of the mesenchymal stromal cell compartment in human bone marrow changes during development and aging. Haematologica, 2012, 97, 179-183.	3. 5	89
18	One single dose of 200â€fμg of antenatal RhIG halves the risk of antiâ€D immunization and hemolytic disease of the fetus and newborn in the next pregnancy. Transfusion, 2008, 48, 1721-1729.	1.6	88

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19	Clinical Applications of Cell-Free Fetal DNA From Maternal Plasma. Obstetrics and Gynecology, 2004, 103, 157-164.	2.4	87
20	Reliability of Fetal Sex Determination Using Maternal Plasma. Obstetrics and Gynecology, 2010, 115, 117-126.	2.4	87
21	Leukocyte-Endothelium Interaction Promotes SDF-1-dependent Polarization of CXCR4. Journal of Biological Chemistry, 2003, 278, 30302-30310.	3.4	85
22	<i>PHOX2B</i> Is a Novel and Specific Marker for Minimal Residual Disease Testing in Neuroblastoma. Journal of Clinical Oncology, 2008, 26, 5443-5449.	1.6	83
23	Glycosylation pattern of antiâ€platelet IgG is stable during pregnancy and predicts clinical outcome in alloimmune thrombocytopenia. British Journal of Haematology, 2016, 174, 310-320.	2.5	83
24	Non-invasive prenatal diagnosis and determination of fetal Rh status. Seminars in Fetal and Neonatal Medicine, 2008, 13, 63-68.	2.3	81
25	Fetal Sex Determination From Maternal Plasma in Pregnancies at Risk for Congenital Adrenal Hyperplasia. Obstetrics and Gynecology, 2001, 98, 374-378.	2.4	80
26	Mesenchymal Stromal Cell Migration: Possibilities to Improve Cellular Therapy. Stem Cells and Development, 2012, 21, 19-29.	2.1	80
27	Comparison of the Fc glycosylation of fetal and maternal immunoglobulin G. Glycoconjugate Journal, 2013, 30, 147-157.	2.7	76
28	Fine-Scale Mapping of the 5q11.2 Breast Cancer Locus Reveals at Least Three Independent Risk Variants Regulating MAP3K1. American Journal of Human Genetics, 2015, 96, 5-20.	6.2	76
29	C-reactive protein enhances IgG-mediated phagocyte responses and thrombocytopenia. Blood, 2015, 125, 1793-1802.	1.4	74
30	Monoclonal antibodies against myeloperoxidase are valuable immunological reagents for the diagnosis of acute myeloid leukaemia. British Journal of Haematology, 1990, 74, 173-178.	2.5	68
31	Detecting Minimal Residual Disease in Neuroblastoma: The Superiority of a Panel of Real-Time Quantitative PCR Markers. Clinical Chemistry, 2009, 55, 1316-1326.	3.2	65
32	Sensitivity of fetal <i>RHD</i> screening for safe guidance of targeted anti-D immunoglobulin prophylaxis: prospective cohort study of a nationwide programme in the Netherlands. BMJ, The, 2016, 355, i5789.	6.0	63
33	B precursor acute lymphoblastic leukemia third complementarity-determining regions predominantly represent an unbiased recombination repertoire: Leukemic transformation frequently occurs in fetal life. European Journal of Immunology, 1994, 24, 900-908.	2.9	57
34	Nuclear Receptors <i>Nur77</i> Nur71Modulate Mesenchymal Stromal Cell Migration. Stem Cells and Development, 2012, 21, 228-238.	2.1	56
35	Fc Galactosylation Promotes Hexamerization of Human IgG1, Leading to Enhanced Classical Complement Activation. Journal of Immunology, 2021, 207, 1545-1554.	0.8	56
36	<i><scp>RHD</scp></i> and <i><scp>RHCE</scp></i> variant and zygosity genotyping via multiplex ligation–dependent probe amplification. Transfusion, 2013, 53, 1559-1574.	1.6	50

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37	Antigen specificity determines antiâ€red blood cell IgGâ€Fc alloantibody glycosylation and thereby severity of haemolytic disease of the fetus and newborn. British Journal of Haematology, 2017, 176, 651-660.	2.5	50
38	Use of Biâ€Allelic Insertion/Deletion Polymorphisms as a Positive Control for Fetal Genotyping in Maternal Blood. Annals of the New York Academy of Sciences, 2006, 1075, 123-129.	3.8	48
39	Evaluation of prenatal RHD typing strategies on cell-free fetal DNA from maternal plasma. Transfusion, 2006, 46, 2142-2148.	1.6	46
40	SAFEâ€"The <i>S</i> pecial Nonâ€invasive <i>A</i> dvances in <i>F</i> etal and Neonatal <i>E</i> valuation Network: aims and achievements. Prenatal Diagnosis, 2008, 28, 83-88.	2.3	46
41	Functional platelet defects in children with severe chronic ITP as tested with 2 novel assays applicable for low platelet counts. Blood, 2014, 123, 1556-1563.	1.4	46
42	Prophylactic antiâ€ <scp>D</scp> preparations display variable decreases in <scp>F</scp> câ€fucosylation of antiâ€ <scp>D</scp> . Transfusion, 2015, 55, 553-562.	1.6	45
43	The highly variable RH locus in nonwhite persons hampers RHD zygosity determination but yields more insight into RH-related evolutionary events. Transfusion, 2005, 45, 327-337.	1.6	44
44	<scp>H</scp> 435â€containing immunoglobulin <scp>G</scp> 3 allotypes are transported efficiently across the human placenta: implications for alloantibodyâ€mediated diseases of the newborn. Transfusion, 2014, 54, 665-671.	1.6	43
45	Extensive Ethnic Variation and Linkage Disequilibrium at the FCGR2/3 Locus: Different Genetic Associations Revealed in Kawasaki Disease. Frontiers in Immunology, 2019, 10, 185.	4.8	43
46	RHC and RHc genotyping in different ethnic groups. Transfusion, 2002, 42, 634-644.	1.6	39
47	Characterization of platelet-specific alloantigens by immimoblotting: localization of Zw and Bak antigens. British Journal of Haematology, 1986, 64, 715-723.	2.5	38
48	The controversy about controls for fetal blood group genotyping by cell-free fetal DNA in maternal plasma. Current Opinion in Hematology, 2011, 18, 467-473.	2.5	36
49	Afucosylated Plasmodium falciparum-specific IgG is induced by infection but not by subunit vaccination. Nature Communications, 2021, 12, 5838.	12.8	36
50	The Majority of Human Memory B Cells Recognizing RhD and Tetanus Resides in IgM+ B Cells. Journal of Immunology, 2014, 193, 1071-1079.	0.8	34
51	On the Perplexingly Low Rate of Transport of IgG2 across the Human Placenta. PLoS ONE, 2014, 9, e108319.	2.5	32
52	Cell cycle and tissue of origin contribute to the migratory behaviour of human fetal and adult mesenchymal stromal cells. British Journal of Haematology, 2010, 148, 428-440.	2.5	30
53	Methylated RASSF1a Is the First Specific DNA Marker for Minimal Residual Disease Testing in Neuroblastoma. Clinical Cancer Research, 2012, 18, 808-814.	7.0	30
54	Genotyping to prevent Rh disease. Current Opinion in Hematology, 2017, 24, 544-550.	2.5	30

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55	Recovery and functional activity of mononuclear bone marrow and peripheral blood cells after different cell isolation protocols used in clinical trials for cell therapy after acute myocardial infarction. EuroIntervention, 2008, 4, 133-138.	3.2	29
56	Production of recombinant Ig molecules from antigen-selected single B cells and restricted usage of Ig-gene segments by anti-D antibodies. Journal of Immunological Methods, 2005, 298, 9-20.	1.4	28
57	MEIS1 regulates early erythroid and megakaryocytic cell fate. Haematologica, 2014, 99, 1555-1564.	3 . 5	28
58	Patients with IgG1-anti-red blood cell autoantibodies show aberrant Fc-glycosylation. Scientific Reports, 2017, 7, 8187.	3.3	27
59	BIGH3 modulates adhesion and migration of hematopoietic stem and progenitor cells. Cell Adhesion and Migration, 2013, 7, 434-449.	2.7	25
60	Frequency and characterization of known and novel <i>RHD</i> variant alleles in 37Â782 Dutch Dâ€negative pregnant women. British Journal of Haematology, 2016, 173, 469-479.	2 . 5	25
61	Cell-Free Fetal DNA Is Not Present in Plasma of Nonpregnant Mothers. Clinical Chemistry, 2004, 50, 679-681.	3.2	23
62	Molecular typing of human platelet and neutrophil antigens (HPA and HNA). Transfusion and Apheresis Science, 2014, 50, 189-199.	1.0	23
63	Fc-Glycosylation in Human IgG1 and IgG3 Is Similar for Both Total and Anti-Red-Blood Cell Anti-K Antibodies. Frontiers in Immunology, 2018, 9, 129.	4.8	23
64	Impact of genetic variation in the <i>SMIM1</i> gene on Vel expression levels. Transfusion, 2015, 55, 1457-1466.	1.6	22
65	Minimal residual disease detection in autologous stem cell grafts from patients with high risk neuroblastoma. Pediatric Blood and Cancer, 2015, 62, 1368-1373.	1.5	22
66	Reduced FcRn-mediated transcytosis of IgG2 due to a missing Glycine in its lower hinge. Scientific Reports, 2019, 9, 7363.	3.3	21
67	Whole-Genome Sequencing Identifies Patient-Specific DNA Minimal Residual Disease Markers in Neuroblastoma. Journal of Molecular Diagnostics, 2015, 17, 43-52.	2.8	19
68	Stability of PCR Targets for Monitoring Minimal Residual Disease in Neuroblastoma. Journal of Molecular Diagnostics, 2012, 14, 168-175.	2.8	18
69	Combining Hypermethylated RASSF1A Detection Using ddPCR with miR-371a-3p Testing: An Improved Panel of Liquid Biopsy Biomarkers for Testicular Germ Cell Tumor Patients. Cancers, 2021, 13, 5228.	3.7	18
70	Mesenchymal Neuroblastoma Cells Are Undetected by Current mRNA Marker Panels: The Development of a Specific Neuroblastoma Mesenchymal Minimal Residual Disease Panel. JCO Precision Oncology, 2019, 3, 1-11.	3.0	17
71	Multiplex blood group typing by cellular surface plasmon resonance imaging. Transfusion, 2019, 59, 754-761.	1.6	16
72	Human IgG lacking effector functions demonstrate lower FcRn-binding and reduced transplacental transport. Molecular Immunology, 2018, 95, 1-9.	2.2	15

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73	TGFBI Expressed by Bone Marrow Niche Cells and Hematopoietic Stem and Progenitor Cells Regulates Hematopoiesis. Stem Cells and Development, 2018, 27, 1494-1506.	2.1	15
74	Different Balance of Wnt Signaling in Adult and Fetal Bone Marrow-Derived Mesenchymal Stromal Cells. Stem Cells and Development, 2016, 25, 934-947.	2.1	14
7 5	Transient and chronic childhood immune thrombocytopenia are distinctly affected by Fc- \hat{l}^3 receptor polymorphisms. Blood Advances, 2019, 3, 2003-2012.	5.2	14
76	The Metastatic Bone Marrow Niche in Neuroblastoma: Altered Phenotype and Function of Mesenchymal Stromal Cells. Cancers, 2020, 12, 3231.	3.7	14
77	Hypermethylated <i>RASSF1A</i> as Circulating Tumor DNA Marker for Disease Monitoring in Neuroblastoma. JCO Precision Oncology, 2020, 4, 291-306.	3.0	14
78	Antiâ€platelet antibodies in childhood immune thrombocytopenia: Prevalence and prognostic implications. Journal of Thrombosis and Haemostasis, 2020, 18, 1210-1220.	3.8	13
79	Specific and Sensitive Detection of Neuroblastoma mRNA Markers by Multiplex RT-qPCR. Cancers, 2021, 13, 150.	3.7	13
80	Improving Risk Stratification for Pediatric Patients with Rhabdomyosarcoma by Molecular Detection of Disseminated Disease. Clinical Cancer Research, 2021, 27, 5576-5585.	7.0	13
81	Novel Circulating Hypermethylated RASSF1A ddPCR for Liquid Biopsies in Patients With Pediatric Solid Tumors. JCO Precision Oncology, 2021, 5, 1738-1748.	3.0	13
82	RhIg-prophylaxis is not influenced by FCGR2/3 polymorphisms involved in red blood cell clearance. Blood, 2017, 129, 1045-1048.	1.4	12
83	lgG-Fc glycosylation before and after rituximab treatment in immune thrombocytopenia. Scientific Reports, 2020, 10, 3051.	3.3	12
84	Potential Diagnostic Approaches for Prediction of Therapeutic Responses in Immune Thrombocytopenia. Journal of Clinical Medicine, 2021, 10, 3403.	2.4	12
85	Multiplex ligation-dependent probe amplification (MLPA) assay for blood group genotyping, copy number quantification, and analysis of $\langle i \rangle$ RH $\langle i \rangle$ variants. Immunohematology, 2015, 31, 58-61.	0.2	11
86	The restricted use of IGHV3 superspecies genes in anti-Rh is not limited to hyperimmunized anti-D donors. Transfusion, 2006, 46, 2162-2168.	1.6	10
87	Neuroblastoma messenger RNA is frequently detected in bone marrow at diagnosis of localised neuroblastoma patients. European Journal of Cancer, 2016, 54, 149-158.	2.8	10
88	Noninvasive Fetal Blood Group Typing. , 2018, , 125-156.		10
89	Associations between single nucleotide polymorphisms and erythrocyte parameters in humans: A systematic literature review. Mutation Research - Reviews in Mutation Research, 2019, 779, 58-67.	5.5	10
90	Noninvasive prenatal blood group and <scp>HPA</scp> â€la genotyping: the current <scp>E</scp> uropean experience. Transfusion, 2013, 53, 2834-2836.	1.6	9

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91	A Conceptual Framework for Optimizing Blood Matching Strategies: Balancing Patient Complications Against Total Costs Incurred. Frontiers in Medicine, 2018, 5, 199.	2.6	9
92	C-Reactive Protein Enhances IgG-Mediated Cellular Destruction Through IgG-Fc Receptors in vitro. Frontiers in Immunology, 2021, 12, 594773.	4.8	9
93	Clinical characteristics of human platelet antigen (HPA)â€1a and HPAâ€5b alloimmunised pregnancies and the association between platelet HPAâ€5b antibodies and symptomatic fetal neonatal alloimmune thrombocytopenia. British Journal of Haematology, 2021, 195, 595-603.	2.5	9
94	Biological and structural characterization of murine TRALI antibody reveals increased Fc-mediated complement activation. Blood Advances, 2020, 4, 3875-3885.	5.2	8
95	Will Genotyping Replace Serology in Future Routine Blood Grouping? & Samp; ndash; Opinion 5. Transfusion Medicine and Hemotherapy, 2009, 36, 234-235.	1.6	7
96	The Elements Steering Pathogenesis in IgG-Mediated Alloimmune Diseases. Journal of Clinical Immunology, 2016, 36, 76-81.	3.8	7
97	Peripheral Stem Cell Apheresis is Feasible Post 131Iodine-Metaiodobenzylguanidine-Therapy in High-Risk Neuroblastoma, but Results in Delayed Platelet Reconstitution. Clinical Cancer Research, 2019, 25, 1012-1021.	7.0	7
98	A variant RhAG protein encoded by theRHAG*572Aallele causes serological weak D expression while maintaining normal RhCE phenotypes. Transfusion, 2019, 59, 405-411.	1.6	7
99	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
100	Immunoassay for quantification of antigen-specific lgG fucosylation. EBioMedicine, 2022, 81, 104109.	6.1	7
101	Development of a recombinant antiâ€Vel immunoglobulin M to identify Velâ€negative donors. Transfusion, 2019, 59, 1359-1366.	1.6	6
102	HIP (HPA-screening in pregnancy) study: protocol of a nationwide, prospective and observational study to assess incidence and natural history of fetal/neonatal alloimmune thrombocytopenia and identifying pregnancies at risk. BMJ Open, 2020, 10, e034071.	1.9	6
103	Rh disease prevention: the European Perspective. ISBT Science Series, 2021, 16, 106-118.	1.1	6
104	Neutrophil Antigens, from Bench to Bedside. Immunological Investigations, 1995, 24, 245-272.	2.0	4
105	Fast and lowâ€cost direct ELISA for highâ€throughput serological HPAâ€1 a typing. Transfusion, 2019, 59, 2989-2996.	1.6	4
106	A functional spleen contributes to afucosylated IgG in humans. Scientific Reports, 2021, 11, 24045.	3.3	4
107	SMIM1 missense mutations exert their effect on wild type Vel expression late in erythroid differentiation. Transfusion, 2021, 61, 236-245.	1.6	3
108	Identification of a novel frequentRHCE*ce308Tvariant allele in Chinese D– individuals, resulting in a C+c– phenotype. Transfusion, 2016, 56, 2314-2321.	1.6	2

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109	Identification of a novel singleâ€nucleotide mutation in <i>SMIM1</i> gene that results in low Vel antigen expression. Transfusion, 2019, 59, E8-E10.	1.6	2
110	Fetal <i>RHD</i> genotyping after bone marrow transplantation. Transfusion, 2016, 56, 2122-2126.	1.6	1
111	Predicting anti-RhD titers in donors: Boostering response and decline rates are personal. PLoS ONE, 2018, 13, e0196382.	2.5	1
112	Pediatric Neuroblastoma: Molecular Detection of Minimal Residual Disease. Pediatric Cancer, 2012, , 47-63.	0.0	0
113	Factors contributing to the pathogenesis of lgGâ€mediated alloimmune disease. ISBT Science Series, 2016, 11, 126-132.	1.1	O
114	Gene-expression-based monocyte-specific clustering of acute myeloid leukemias reveals novel associations. Leukemia and Lymphoma, 2017, 58, 1721-1725.	1.3	0