

Anne C Goodeve

List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

130
papers

4,796
citations

34
h-index

67
g-index

135
ext. papers

5,221
ext. citations

5.1
avg. IF

5.24
L-index

#	Paper	IF	Citations
130	Genotypes of European and Iranian patients with type 3 von Willebrand disease enrolled in 3WINTERS-IPS. <i>Blood Advances</i> , 2021 , 5, 2987-3001	7.8	0
129	Bleeding symptoms in patients diagnosed as type 3 von Willebrand disease: Results from 3WINTERS-IPS, an international and collaborative cross-sectional study. <i>Journal of Thrombosis and Haemostasis</i> , 2020 , 18, 2145-2154	15.4	9
128	Characterization of large in-frame von Willebrand factor deletions highlights differing pathogenic mechanisms. <i>Blood Advances</i> , 2020 , 4, 2979-2990	7.8	2
127	Curated disease-causing genes for bleeding, thrombotic, and platelet disorders: Communication from the SSC of the ISTH. <i>Journal of Thrombosis and Haemostasis</i> , 2019 , 17, 1253-1260	15.4	31
126	Next Generation Sequencing in Newborn Screening in the United Kingdom National Health Service. <i>International Journal of Neonatal Screening</i> , 2019 , 5, 40	2.6	14
125	Profile of Mutations Identified in the 3WINTERS-IPS Project on European & Iranian Patients with Previously Diagnosed Type 3 Von Willebrand Disease.. <i>Blood</i> , 2018 , 132, 1184-1184	2.2	
124	Clustering of Bleeding Symptoms in Patients Previously Diagnosed As Type 3 Von Willebrand Disease: Results from a Large Cohort of Type 3 Von Willebrand Disease (the 3Winters-Ips Project). <i>Blood</i> , 2018 , 132, 2465-2465	2.2	
123	Genetic and Laboratory Diagnosis 2018 , 153-166		
122	Evaluation of a semi-automated von Willebrand factor multimer assay, the Hydrigel 5 von Willebrand multimer, by two European Centers. <i>Research and Practice in Thrombosis and Haemostasis</i> , 2018 , 2, 790-799	5.1	23
121	The common single nucleotide variants c.2365A>G and c.2385T>C modify VWF biosynthesis and clearance. <i>Blood Advances</i> , 2018 , 2, 1585-1594	7.8	12
120	Genetic analysis of bleeding disorders. <i>Haemophilia</i> , 2016 , 22 Suppl 5, 79-83	3.3	1
119	Clinical and laboratory variability in a cohort of patients diagnosed with type 1 VWD in the United States. <i>Blood</i> , 2016 , 127, 2481-8	2.2	76
118	Diagnosing von Willebrand disease: genetic analysis. <i>Hematology American Society of Hematology Education Program</i> , 2016 , 2016, 678-682	3.1	15
117	A high-throughput sequencing test for diagnosing inherited bleeding, thrombotic, and platelet disorders. <i>Blood</i> , 2016 , 127, 2791-803	2.2	135
116	In silico analysis highlights the copy number variation mechanism responsible for the historically reported VWF exon 42 deletion. <i>Haemophilia</i> , 2016 , 22, e484-7	3.3	
115	ßIIB variants defined by next-generation sequencing: predicting variants likely to cause Glanzmann thrombasthenia. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2015 , 112, E1898-907	11.5	33
114	HemophiliaB: molecular pathogenesis and mutation analysis. <i>Journal of Thrombosis and Haemostasis</i> , 2015 , 13, 1184-95	15.4	56

113	Identification and Characterization of Novel Variations in Platelet G-Protein Coupled Receptor (GPCR) Genes in Patients Historically Diagnosed with Type 1 von Willebrand Disease. <i>PLoS ONE</i> , 2015 , 10, e0143913	3.7	5
112	Identification of Novel Mutations with Molecular Modelling of Missense Mutations of Congenital Afibrinogenemia Patients in Pakistan. <i>Blood</i> , 2015 , 126, 4664-4664	2.2	
111	von Willebrand Disease: Molecular Aspects 2014 , 353-361		
110	The UK National External Quality Assessment Scheme for heritable bleeding disorders. <i>Seminars in Thrombosis and Hemostasis</i> , 2014 , 40, 261-8	5.3	6
109	Genomics of bleeding disorders. <i>Haemophilia</i> , 2014 , 20 Suppl 4, 50-3	3.3	9
108	The diagnosis and management of von Willebrand disease: a United Kingdom Haemophilia Centre Doctors Organization guideline approved by the British Committee for Standards in Haematology. <i>British Journal of Haematology</i> , 2014 , 167, 453-65	4.5	222
107	p.P2063S: a neutral VWF variant masquerading as a mutation. <i>Annals of Hematology</i> , 2014 , 93, 505-6	3	3
106	Characterization and Expression of an In-Frame Exon 33-34 Deletion Causing Type 1 VWD. <i>Blood</i> , 2014 , 124, 1525-1525	2.2	
105	Assessment of the F9 genotype-specific FIX inhibitor risks and characterisation of 10 novel severe F9 defects in the first molecular series of Argentinian patients with haemophilia B. <i>Thrombosis and Haemostasis</i> , 2013 , 109, 24-33	7	26
104	Principles of care for the diagnosis and treatment of von Willebrand disease. <i>Haematologica</i> , 2013 , 98, 667-74	6.6	139
103	Specific and global coagulation assays in the diagnosis of discrepant mild hemophilia A. <i>Haematologica</i> , 2013 , 98, 1980-7	6.6	35
102	VWF propeptide and ratios between VWF, VWF propeptide, and FVIII in the characterization of type 1 von Willebrand disease. <i>Blood</i> , 2013 , 121, 2336-9	2.2	69
101	VWF sequence variants: a data goldmine. <i>Blood</i> , 2013 , 122, 471-3	2.2	3
100	Identification and characterisation of mutations associated with von Willebrand disease in a Turkish patient cohort. <i>Thrombosis and Haemostasis</i> , 2013 , 110, 264-74	7	12
99	The Complete Type I VWD Cohort Of The Zimmerman Program For The Molecular and Clinical Biology Of VWD - Phenotypic Assignment, Mutation Frequency, and Bleeding Assessment. <i>Blood</i> , 2013 , 122, 332-332	2.2	5
98	Genetics of haemostasis. <i>Haemophilia</i> , 2012 , 18 Suppl 4, 73-80	3.3	5
97	Quality in molecular biology testing for inherited thrombophilia disorders. <i>Seminars in Thrombosis and Hemostasis</i> , 2012 , 38, 600-12	5.3	16
96	Intracellular Retention, Enhanced Clearance, and Defective FVIII Binding Are Common Features of Von Willebrand Factor D'-D3 Domain Mutations in Patients with Von Willebrand Disease Type 1 From the European Mcmdm-1VWD Study. <i>Blood</i> , 2012 , 120, 99-99	2.2	1

95	von Willebrand disease. <i>Genetics in Medicine</i> , 2011 , 13, 365-76	8.1	60
94	p.Tyr365Cys change in factor VIII: haemophilia A, but not as we know it. <i>British Journal of Haematology</i> , 2011 , 154, 618-25	4.5	16
93	Nomenclature of genetic variants in hemostasis. <i>Journal of Thrombosis and Haemostasis</i> , 2011 , 9, 852-5	15.4	29
92	The molecular basis of von Willebrand disease: the under investigated, the unexpected and the overlooked. <i>Haematologica</i> , 2011 , 96, 798-800	6.6	8
91	Diagnosis and management of von Willebrand disease in the United Kingdom. <i>Seminars in Thrombosis and Hemostasis</i> , 2011 , 37, 488-94	5.3	11
90	The international society on thrombosis and haemostasis von Willebrand disease database: an update. <i>Seminars in Thrombosis and Hemostasis</i> , 2011 , 37, 470-9	5.3	61
89	Haemophilia A and von Willebrand's disease. <i>Haemophilia</i> , 2010 , 16 Suppl 5, 79-84	3.3	10
88	The impact of bleeding history, von Willebrand factor and PFA-100(®) on the diagnosis of type 1 von Willebrand disease: results from the European study MCMDM-1VWD. <i>British Journal of Haematology</i> , 2010 , 151, 245-51	4.5	32
87	von Willebrand Disease: Molecular Aspects 2010 , 278-285		1
86	Polymorphic variation within the VWF gene contributes to the failure to detect mutations in patients historically diagnosed with type 1 von Willebrand disease from the MCMDM-1VWD cohort. <i>Haematologica</i> , 2010 , 95, 2163-5	6.6	13
85	The genetic basis of von Willebrand disease. <i>Blood Reviews</i> , 2010 , 24, 123-34	11.1	114
84	Tribbles-1 and -2 are tumour suppressors, down-regulated in human acute myeloid leukaemia. <i>Immunology Letters</i> , 2010 , 130, 115-24	4.1	32
83	Genetic testing for von Willebrand disease: the case for. <i>Journal of Thrombosis and Haemostasis</i> , 2010 , 8, 13-6	15.4	24
82	Vicenza deciphered: modeling the von Willebrand disease enigma: commentary on accelerated clearance alone explains ultralarge multimers in VWD Vicenza. <i>Journal of Thrombosis and Haemostasis</i> , 2010 , 8, 1271-2	15.4	2
81	Another step towards understanding hemophilia A molecular pathogenesis. <i>Journal of Thrombosis and Haemostasis</i> , 2010 , 8, 2693-5	15.4	
80	Investigation of the Role of Copy Number Variation In the Pathogenesis of Type 1 Von Willebrand Disease. <i>Blood</i> , 2010 , 116, 2218-2218	2.2	
79	Identification and characterization of a novel P2Y ₁₂ variant in a patient diagnosed with type 1 von Willebrand disease in the European MCMDM-1VWD study. <i>Blood</i> , 2009 , 113, 4110-3	2.2	59
78	When 1 plus 1 equals 3 in VWD. <i>Blood</i> , 2009 , 114, 933-4	2.2	2

77	Re-evaluation of three Israeli families initially diagnosed with type 1 von Willebrand disease in light of the ISTH update on von Willebrand factor pathophysiology and classification. <i>Haemophilia</i> , 2008 , 14, 621-4	3.3	2
76	Type 1 von Willebrand disease: application of emerging data to clinical practice. <i>Haemophilia</i> , 2008 , 14, 685-96	3.3	19
75	Authors' response to Dong-Zhi Li's letter: Routine molecular screening for common β -Thalassaemia deletions is necessary as part of an antenatal screening programme. <i>Journal of Medical Screening</i> , 2008 , 15, 47-47	1.4	0
74	Molecular genetic testing of hemophilia A. <i>Seminars in Thrombosis and Hemostasis</i> , 2008 , 34, 491-501	5.3	18
73	Methylation of the suppressor of cytokine signaling 3 gene (SOCS3) in myeloproliferative disorders. <i>Haematologica</i> , 2008 , 93, 1635-44	6.6	68
72	Response to desmopressin is influenced by the genotype and phenotype in type 1 von Willebrand disease (VWD): results from the European Study MCMDM-1VWD. <i>Blood</i> , 2008 , 111, 3531-9	2.2	162
71	Identification of type 1 von Willebrand disease patients with reduced von Willebrand factor survival by assay of the VWF propeptide in the European study: molecular and clinical markers for the diagnosis and management of type 1 VWD (MCMDM-1VWD). <i>Blood</i> , 2008 , 111, 4979-85	2.2	123
70	Gene structure, expression profiling and mutation analysis of the tumour suppressor SHP1 in Caucasian acute myeloid leukaemia. <i>Leukemia</i> , 2007 , 21, 2390-3	10.7	24
69	Sixteen novel hemophilia A causative mutations in the first Argentinian series of severe molecular defects. <i>Haematologica</i> , 2007 , 92, 842-5	6.6	24
68	Is routine molecular screening for common alpha-thalassaemia deletions necessary as part of an antenatal screening programme?. <i>Journal of Medical Screening</i> , 2007 , 14, 60-1	1.4	6
67	Phenotype and genotype of a cohort of families historically diagnosed with type 1 von Willebrand disease in the European study, Molecular and Clinical Markers for the Diagnosis and Management of Type 1 von Willebrand Disease (MCMDM-1VWD). <i>Blood</i> , 2007 , 109, 112-21	2.2	309
66	Genetics of type 1 von Willebrand disease. <i>Current Opinion in Hematology</i> , 2007 , 14, 444-9	3.3	9
65	Enhanced carrier and prenatal diagnosis in the Italian haemophilia B population. <i>Blood Transfusion</i> , 2007 , 5, 117-9	3.6	
64	Mutational analysis of the von Willebrand factor gene in type 1 von Willebrand disease using conformation sensitive gel electrophoresis: a comparison of fluorescent and manual techniques. <i>Haematologica</i> , 2007 , 92, 550-3	6.6	13
63	The UK National External Quality Assessment Scheme (UK NEQAS) for molecular genetic testing in haemophilia. <i>Thrombosis and Haemostasis</i> , 2006 , 96, 597-601	7	19
62	Investigation of Underlying Reasons of Factor VIII Deficiency in Haemophilia A Patients with Undetectable Mutations in the Factor VIII Gene.. <i>Blood</i> , 2006 , 108, 1042-1042	2.2	
61	The UK National External Quality Assessment Scheme (UK NEQAS) for molecular genetic testing in haemophilia. <i>Thrombosis and Haemostasis</i> , 2006 , 96, 597-601	7	5
60	Aberrant methylation of the negative regulators RASSF1A, SHP-1 and SOCS-1 in myelodysplastic syndromes and acute myeloid leukaemia. <i>British Journal of Haematology</i> , 2005 , 129, 60-5	4.5	71

59	JAK2 V617F Mutation is uncommon in chronic myelomonocytic leukaemia. <i>British Journal of Haematology</i> , 2005 , 130, 968	4.5	25
58	Mutations in PTPN11 are uncommon in adult myelodysplastic syndromes and acute myeloid leukaemia. <i>British Journal of Haematology</i> , 2004 , 124, 843-4	4.5	19
57	Activating loop mutations in the PDGFR alpha and beta genes are rare in core binding factor acute myeloid leukaemia. <i>British Journal of Haematology</i> , 2004 , 127, 123-4	4.5	5
56	Homeologous recombination between AluSx-sequences as a cause of hemophilia. <i>Human Mutation</i> , 2004 , 24, 440	4.7	32
55	Comparison of Phenotypic Assessment and Mutation Detection in the Diagnosis of Carrier State in Hemophilia: Identification of 10 Novel Mutations.. <i>Blood</i> , 2004 , 104, 4020-4020	2.2	
54	Promoter Methylation of RASSF1A, SHP1 and SOCS1 Genes in Acute Myeloid Leukaemia (AML) and Myelodysplastic Syndromes (MDS).. <i>Blood</i> , 2004 , 104, 2999-2999	2.2	
53	Analysis of factor VIII gene intron 1 inversion in Argentinian families with severe haemophilia A and a review of the literature. <i>Blood Coagulation and Fibrinolysis</i> , 2004 , 15, 569-572	1	15
52	Second hit mutations in the RTK/RAS signaling pathway in acute myeloid leukemia with inv(16). <i>Haematologica</i> , 2004 , 89, 106	6.6	34
51	The molecular basis of hemophilia A: genotype-phenotype relationships and inhibitor development. <i>Seminars in Thrombosis and Hemostasis</i> , 2003 , 29, 23-30	5.3	100
50	The incidence of inhibitor development according to specific mutations--and treatment?. <i>Blood Coagulation and Fibrinolysis</i> , 2003 , 14 Suppl 1, S17-21	1	27
49	Analysis of intracellular storage and regulated secretion of 3 von Willebrand disease-causing variants of von Willebrand factor. <i>Blood</i> , 2003 , 102, 2452-8	2.2	67
48	Mutational analysis of class III receptor tyrosine kinases (C-KIT, C-FMS, FLT3) in idiopathic myelofibrosis. <i>British Journal of Haematology</i> , 2003 , 120, 464-70	4.5	27
47	Incidence and prognosis of c-KIT and FLT3 mutations in core binding factor (CBF) acute myeloid leukaemias. <i>British Journal of Haematology</i> , 2003 , 121, 775-7	4.5	253
46	c-FMS mutational analysis in acute myeloid leukaemia. <i>British Journal of Haematology</i> , 2003 , 123, 749-50	4.5	14
45	FLT3 internal tandem duplication mutations are rare in agnogenic myeloid metaplasia. <i>Blood</i> , 2002 , 100, 364	2.2	9
44	Genomic structure of human FLT3: implications for mutational analysis. <i>British Journal of Haematology</i> , 2001 , 113, 1076-7	4.5	60
43	Identification of novel FLT-3 Asp835 mutations in adult acute myeloid leukaemia. <i>British Journal of Haematology</i> , 2001 , 113, 983-8	4.5	303
42	A standard nomenclature for von Willebrand factor gene mutations and polymorphisms. <i>Best Practice and Research in Clinical Haematology</i> , 2001 , 14, 235-40	4.2	5

41	Endoplasmic reticulum retention and prolonged association of a von Willebrand's disease-causing von Willebrand factor variant with ERp57 and calnexin. <i>Biochemical and Biophysical Research Communications</i> , 2001 , 280, 448-53	3.4	24
40	Somatic mosaicism in hemophilia A: a fairly common event. <i>American Journal of Human Genetics</i> , 2001 , 69, 75-87	11	125
39	A Standard Nomenclature for von Willebrand Factor Gene Mutations and Polymorphisms. <i>Thrombosis and Haemostasis</i> , 2001 , 85, 929-931	7	43
38	A second case of Hb Renert [β 133(H11)Val \rightarrow Ala]. <i>Hemoglobin</i> , 2001 , 25, 337-40	0.6	3
37	A standard nomenclature for von Willebrand factor gene mutations and polymorphisms. On behalf of the ISTH SSC Subcommittee on von Willebrand factor. <i>Thrombosis and Haemostasis</i> , 2001 , 85, 929-31	7	7
36	FLT3 internal tandem duplication mutations in adult acute myeloid leukaemia define a high-risk group. <i>British Journal of Haematology</i> , 2000 , 111, 190-195	4.5	7
35	Are aberrant BCR/ABL transcripts more common than previously thought?. <i>British Journal of Haematology</i> , 2000 , 111, 1109-1111	4.5	0
34	Two novel type 2N von Willebrand disease-causing mutations that result in defective factor VIII binding, multimerization, and secretion of von Willebrand factor. <i>Blood</i> , 2000 , 95, 2000-2007	2.2	57
33	A novel von Willebrand disease-causing mutation (Arg273Trp) in the von Willebrand factor propeptide that results in defective multimerization and secretion. <i>Blood</i> , 2000 , 96, 560-568	2.2	49
32	Relationship between Factor VIII Mutation Type and Inhibitor Development in a Cohort of Previously Untreated Patients Treated with Recombinant Factor VIII (Recombinate). <i>Thrombosis and Haemostasis</i> , 2000 , 83, 844-848	7	51
31	A novel von Willebrand disease-causing mutation (Arg273Trp) in the von Willebrand factor propeptide that results in defective multimerization and secretion. <i>Blood</i> , 2000 , 96, 560-568	2.2	15
30	FLT3 internal tandem duplication mutations in adult acute myeloid leukaemia define a high-risk group. <i>British Journal of Haematology</i> , 2000 , 111, 190-5	4.5	227
29	Are aberrant BCR/ABL transcripts more common than previously thought?. <i>British Journal of Haematology</i> , 2000 , 111, 1109-11	4.5	28
28	Two novel type 2N von Willebrand disease-causing mutations that result in defective factor VIII binding, multimerization, and secretion of von Willebrand factor. <i>Blood</i> , 2000 , 95, 2000-7	2.2	8
27	A novel von Willebrand disease-causing mutation (Arg273Trp) in the von Willebrand factor propeptide that results in defective multimerization and secretion. <i>Blood</i> , 2000 , 96, 560-8	2.2	18
26	Relationship between factor VIII mutation type and inhibitor development in a cohort of previously untreated patients treated with recombinant factor VIII (Recombinate). Recombinate PUP Study Group. <i>Thrombosis and Haemostasis</i> , 2000 , 83, 844-8	7	9
25	Null Alleles Are not a Common Cause of Type 1 von Willebrand Disease in the British Population. <i>Thrombosis and Haemostasis</i> , 1999 , 82, 1373-1375	7	6
24	A rapid method for haemophilia B mutation detection using conformation sensitive gel electrophoresis. <i>British Journal of Haematology</i> , 1999 , 104, 915-8	4.5	34

23	c-kit proto-oncogene exon 8 in-frame deletion plus insertion mutations in acute myeloid leukaemia. <i>British Journal of Haematology</i> , 1999 , 105, 894-900	4.5	201
22	A study of Wilson disease mutations in Britain. <i>Human Mutation</i> , 1999 , 14, 304-11	4.7	61
21	A common splice site mutation is shared by two families with different type 2N von Willebrand disease mutations. <i>Thrombosis and Haemostasis</i> , 1999 , 82, 1061-4	7	2
20	Advances in carrier detection in haemophilia. <i>Haemophilia</i> , 1998 , 4, 358-64	3.3	12
19	Laboratory methods for the genetic diagnosis of bleeding disorders. <i>International Journal of Laboratory Hematology</i> , 1998 , 20, 3-19		21
18	Precise Carrier Diagnosis in Families with Haemophilia A: Use of Conformation Sensitive Gel Electrophoresis for Mutation Screening and Polymorphism Analysis. <i>Thrombosis and Haemostasis</i> , 1998 , 79, 723-726	7	98
17	Late Relapsing Childhood Lymphoblastic Leukemia. <i>Blood</i> , 1998 , 92, 2334-2337	2.2	51
16	Late Relapsing Childhood Lymphoblastic Leukemia. <i>Blood</i> , 1998 , 92, 2334-2337	2.2	
15	Precise carrier diagnosis in families with haemophilia A: use of conformation sensitive gel electrophoresis for mutation screening and polymorphism analysis. <i>Thrombosis and Haemostasis</i> , 1998 , 79, 723-6	7	13
14	BCR-ABL Transcript With an e19a2 (c3a2) Junction in Classical Chronic Myeloid Leukemia. <i>Blood</i> , 1997 , 89, 3064-3064	2.2	31
13	Clonal stability in late-relapsing childhood lymphoblastic leukaemia. <i>British Journal of Haematology</i> , 1997 , 98, 992-4	4.5	15
12	HLA Class II Profile: A Weak Determinant of Factor VIII Inhibitor Development in Severe Haemophilia A. <i>Thrombosis and Haemostasis</i> , 1997 , 77, 234-237	7	129
11	HLA class II profile: a weak determinant of factor VIII inhibitor development in severe haemophilia A. UKHCDO Inhibitor Working Party. <i>Thrombosis and Haemostasis</i> , 1997 , 77, 234-7	7	34
10	A rapid and cost effective method for analysis of dinucleotide repeat polymorphisms in the factor VIII gene. <i>Blood Coagulation and Fibrinolysis</i> , 1996 , 7, 672-7	1	13
9	Characterisation of Type 2N von Willebrand Disease Using Phenotypic and Molecular Techniques. <i>Thrombosis and Haemostasis</i> , 1996 , 75, 959-964	7	26
8	von Willebrand Factor/Factor VIII Binding Is not Affected by the Arg89Gln Polymorphism in von Willebrand Factor. <i>Thrombosis and Haemostasis</i> , 1996 , 76, 820-821	7	
7	Characterisation of type 2N von Willebrand disease using phenotypic and molecular techniques. <i>Thrombosis and Haemostasis</i> , 1996 , 75, 959-64	7	1
6	Factor VIII gene rearrangements in patients with severe haemophilia A. <i>Lancet, The</i> , 1994 , 343, 329-30	40	51

- 5 A comparison of the allelic frequencies of ten DNA polymorphisms associated with factor VIII and factor IX genes in Thai and Western European populations. *Blood Coagulation and Fibrinolysis*, **1994**, 5, 29-35 1 9
- 4 Reassortants of influenza B viruses for use in vaccines: an evaluation. *Archives of Virology*, **1985**, 83, 169-176 10
- 3 Laboratory Analysis of Von Willebrand Disease: Molecular Analysis 204-215
- 2 Molecular Diagnosis of von Willebrand Disease: The Genotype 114-126 2
- 1 Phenotypic and Genotypic Signatures of VWF Exon 18 in Eastern Saudi Patients Previously Diagnosed with Type 1 von Willebrand Disease. *International Journal of General Medicine*, Volume 15, 5385-5394 2.3