Yanli Ji

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/857635/publications.pdf

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		1163117	1125743
22	189	8	13
papers	citations	h-index	g-index
22	22	22	139
all docs	docs citations	times ranked	citing authors

#	Article	IF	CITATIONS
1	Hemolytic disease of the fetus and newborn due to alloantiâ€M: three Chinese case reports and a review of the literature. Transfusion, 2019, 59, 385-395.	1.6	24
2	The distribution of <scp>MNS</scp> hybrid glycophorins with Mur antigen expression in Chinese donors including identification of a novel <i><scp>GYP</scp>.Bun</i> allele. Vox Sanguinis, 2016, 111, 308-314.	1.5	19
3	The summary of <i><scp>FUT</scp>1</i> and <i><scp>FUT</scp>2</i> genotyping analysis in <scp>C</scp> hinese paraâ€ <scp>B</scp> ombay individuals including additional nine probands from <scp>G</scp> uangzhou in <scp>C</scp> hina. Transfusion, 2013, 53, 3224-3229.	1.6	18
4	<i>RHD</i> genotype and zygosity analysis in the Chinese Southern Han D+, Dâ^ and D variant donors using the multiplex ligationâ€dependent probe amplification assay. Vox Sanguinis, 2017, 112, 660-670.	1.5	18
5	Genotyping for Glycophorin GYP(B-A-B) Hybrid Genes Using a Single Nucleotide Polymorphism-Based Algorithm by Matrix-Assisted Laser Desorption/Ionisation, Time-of-Flight Mass Spectrometry. Molecular Biotechnology, 2016, 58, 665-671.	2.4	13
6	Distribution of maternal red cell antibodies and the risk of severe alloimmune haemolytic disease of the foetus in a Chinese population: a cohort study on prenatal management. BMC Pregnancy and Childbirth, 2020, 20, 539.	2.4	13
7	Novel alleles at the <scp>K</scp> ell blood group locus that lead to <scp>K</scp> ell variant phenotype in the <scp>D</scp> utch population. Transfusion, 2015, 55, 413-421.	1.6	12
8	Genotyping analysis of MNS blood group GP(Bâ€Aâ€B) hybrid glycophorins in the Chinese Southern Han population using a highâ€resolution melting assay. Transfusion, 2018, 58, 1763-1771.	1.6	12
9	Validation of the multiplex ligationâ€dependent probe amplification assay and its application on the distribution study of the major alleles of 17 blood group systems in <scp>C</scp> hinese donors from <scp>G</scp> uangzhou. Transfusion, 2017, 57, 423-432.	1.6	8
10	Hyporegenerative anemia in antiâ€Mâ€associated hemolytic disease of the fetus. Transfusion, 2021, 61, 1908-1915.	1.6	8
11	Incidence of <scp>antiâ€D</scp> alloimmunization in Dâ€negative individuals receiving Dâ€positive red blood cell transfusion: A systematic review and metaâ€analysis. Vox Sanguinis, 2022, 117, 633-640.	1.5	8
12	A novel 519_525dup mutation of <i><scp>KLF</scp>1</i> gene identified in a <scp>C</scp> hinese blood donor with <scp>L</scp> u(a–b–) phenotype. Transfusion, 2013, 53, 1619-1620.	1.6	7
13	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
14	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
15	Red blood cell genotyping in China. ISBT Science Series, 2016, 11, 55-68.	1.1	6
16	Secondary alloanti-D immunization post transfusion of "Asia type―DEL red blood cells. Transfusion and Apheresis Science, 2022, 61, 103458.	1.0	4
17	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
18	Serological screening and genetic analysis of RhCE variants in the Chinese Southern Han donors. Transfusion Medicine, 2021, 31, 271-276.	1.1	2

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19	Successful prenatal management of two foetuses affected by antibodies against GP.Mur with prenatal genotyping analysis and a literature review. Blood Transfusion, 2021, 19, 135-143.	0.4	1
20	Major applications and limitations of blood group genotyping in China. ISBT Science Series, 2018, 13, 365-370.	1.1	0
21	Molecular genetic analysis of Mi a â€positive hybrid glycophorins revealed two novel alleles of GP .Vw and multiple variant transcripts of GYPB existing in both the homozygous GP .Mur and wildâ€type GPB individuals. Transfusion, 2021, 61, 2477-2486.	1.6	O
22	Psychological impact of the COVID â€19 pandemic on young professionals in blood banks and transfusion services: A global crossâ€sectional survey. Vox Sanguinis, 2022, , .	1.5	0