N Lannoy

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/7757319/publications.pdf

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15 papers	184 citations	1163065 8 h-index	1125717 13 g-index
15 all docs	15 docs citations	15 times ranked	238 citing authors

#	ARTICLE	IF	CITATIONS
1	CF Gene and Cystic Fibrosis Transmembrane Conductance Regulator Expression in Autosomal Dominant Polycystic Kidney Disease. Journal of the American Society of Nephrology: JASN, 2000, 11, 2285-2296.	6.1	41
2	Principles of genetic variations and molecular diseases: applications in hemophilia A. Critical Reviews in Oncology/Hematology, 2016, 104, 1-8.	4.4	24
3	Intron 22 homologous regions are implicated in exons 1â€“22 duplications of the F8 gene. European Journal of Human Genetics, 2013, 21, 970-976.	2.8	20
4	Genetic mosaicism in haemophilia: A practical review to help evaluate the risk of transmitting the disease. Haemophilia, 2020, 26, 375-383.	2.1	18
5	Identification of de novo deletion in the factor VIII gene by MLPA technique in two girls with isolated factor VIII deficiency. Haemophilia, 2009, 15, 797-801.	2.1	15
6	The â€˜royal diseaseâ€™â€“ haemophilia A or B? A haematological mystery is finally solved. Haemophilia, 2010, 16, 843-847.	2.1	14
7	Computational and molecular approaches for predicting unreported causal missense mutations in Belgian patients with haemophilia A. Haemophilia, 2012, 18, e331-9.	2.1	11
8	Comparative study of the prevalence of clotting factor deficiency in carriers of haemophilia A and haemophilia B. Haemophilia, 2017, 23, e471-e473.	2.1	8
9	Review of molecular mechanisms at distal Xq28 leading to balanced or unbalanced genomic rearrangements and their phenotypic impacts on hemophilia. Haemophilia, 2018, 24, 711-719.	2.1	7
10	Overrepresentation of missense mutations in mild hemophilia A patients from Belgium: founder effect or independent occurrence?. Thrombosis Research, 2015, 135, 1057-1063.	1.7	6
11	Tandem inversion duplication within F8 Intron 1 associated with mild haemophilia A. Haemophilia, 2015, 21, 516-522.	2.1	5
12	Usual and unusual mutations in a cohort of Belgian patients with hemophilia B. Thrombosis Research, 2017, 149, 25-28.	1.7	5
13	Incidental finding of unreported large duplication in F8 gene during prenatal analysis: Which management for genetic counselling?. Thrombosis Research, 2019, 182, 39-42.	1.7	4
14	Five int22h homologous copies at the Xq28 locus identified in intron22 inversion type 3 of the Factor VIII gene. Thrombosis Research, 2016, 137, 224-227.	1.7	3
15	Inhibitor epidemiology and geneticâ€related risk factors in people with haemophilia from CÃ´te dâ€™Ivoire. Haemophilia, 2020, 26, 79-85.	2.1	3