Barbara Tazon-Vega

List of Publications by Year
in descending order

19	4,413	623699	839512
papers	citations	14 h-index	18 g-index

19	19	19	8826
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Integrative annotation of human large intergenic noncoding RNAs reveals global properties and specific subclasses. Genes and Development, 2011, 25, 1915-1927.	5.9	3,208
2	Long noncoding RNAs regulate adipogenesis. Proceedings of the National Academy of Sciences of the United States of America, 2013, 110, 3387-3392.	7.1	371
3	Clinical Sequencing Uncovers Origins and Evolution of Lassa Virus. Cell, 2015, 162, 738-750.	28.9	230
4	Clinical Utility of Genetic Testing in Children and Adults with Steroid-Resistant Nephrotic Syndrome. Clinical Journal of the American Society of Nephrology: CJASN, 2011, 6, 1139-1148.	4.5	189
5	Cell free circulating tumor DNA in cerebrospinal fluid detects and monitors central nervous system involvement of B-cell lymphomas. Haematologica, 2021, 106, 513-521.	3.5	75
6	Clinical Value ofNPHS2Analysis in Early- and Adult-Onset Steroid-Resistant Nephrotic Syndrome. Clinical Journal of the American Society of Nephrology: CJASN, 2011, 6, 344-354.	4.5	65
7	Autosomal recessive Alportâ€™s syndrome and benign familial hematuria are collagen type IV diseases. American Journal of Kidney Diseases, 2003, 42, 952-959.	1.9	47
8	A CLK3-HMGA2 Alternative Splicing Axis Impacts Human Hematopoietic Stem Cell Molecular Identity throughout Development. Cell Stem Cell, 2018, 22, 575-588.e7.	11.1	40
9	Collagen type IV (Â3-Â4) nephropathy: from isolated haematuria to renal failure. Nephrology Dialysis Transplantation, 2004, 19, 2429-2432.	0.7	39
10	Evaluation of genome coverage and fidelity of multiple displacement amplification from single cells by SNP array. Molecular Human Reproduction, 2009, 15, 739-747.	2.8	36
11	Genetic Testing for X-Linked Alport Syndrome by Direct Sequencing of COL4A5 cDNA From Hair Root RNA Samples. American Journal of Kidney Diseases, 2007, 50, 257.e1-257.e14.	1.9	27
12	Study of candidate genes affecting the progression of renal disease in autosomal dominant polycystic kidney disease type 1. Nephrology Dialysis Transplantation, 2007, 22, 1567-1577.	0.7	25
13	Spanish Guidelines for the use of targeted deep sequencing in myelodysplastic syndromes and chronic myelomonocytic leukaemia. British Journal of Haematology, 2020, 188, 605-622.	2.5	25
14	Male-to-male transmission of X-linked Alport syndrome in a boy with a 47,XXY karyotype. European Journal of Human Genetics, 2005, 13, 1040-1046.	2.8	21
15	Usefulness of NGS for Diagnosis of Dominant Beta-Thalassemia and Unstable Hemoglobinopathies in Five Clinical Cases. Frontiers in Physiology, 2021, 12, 628236.	2.8	7
16	Immunological and genetic kinetics from diagnosis to clinical progression in chronic lymphocytic leukemia. Biomarker Research, 2021, 9, 37.	6.8	5
17	Is acute lymphoblastic leukemia with mature B-cell phenotype and KMT2A rearrangements a new entity? A systematic review and meta-analysis. Leukemia and Lymphoma, 2021, 62, 2202-2210.	1.3	2
18	Variant t(11;22)(q13;q11.2) withIGLinvolvement in mantle cell lymphoma. Leukemia and Lymphoma, 2022, 63, 1746-1749.	1.3	1

#	Article	IF	Citations
19	Prognostic impact of micromegakaryocytes in primary myelodysplastic syndromes. Leukemia and Lymphoma, 2021, , 1-9.	1.3	0