

# Barbara Tazon-Vega

## 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.

18  
papers

3,433  
citations

14  
h-index

19  
g-index

19  
ext. papers

4,086  
ext. citations

8.9  
avg, IF

4.69  
L-index

#	Paper	IF	Citations
18	Variant t(11;22)(q13;q11.2) with involvement in mantle cell lymphoma.. <i>Leukemia and Lymphoma</i> , <b>2022</b> , 1-4	1.9	
17	Is acute lymphoblastic leukemia with mature B-cell phenotype and rearrangements a new entity? A systematic review and meta-analysis. <i>Leukemia and Lymphoma</i> , <b>2021</b> , 62, 2202-2210	1.9	0
16	Immunological and genetic kinetics from diagnosis to clinical progression in chronic lymphocytic leukemia. <i>Biomarker Research</i> , <b>2021</b> , 9, 37	8	2
15	Cell free circulating tumor DNA in cerebrospinal fluid detects and monitors central nervous system involvement of B-cell lymphomas. <i>Haematologica</i> , <b>2021</b> , 106, 513-521	6.6	25
14	Usefulness of NGS for Diagnosis of Dominant Beta-Thalassemia and Unstable Hemoglobinopathies in Five Clinical Cases. <i>Frontiers in Physiology</i> , <b>2021</b> , 12, 628236	4.6	2
13	Prognostic impact of micromegakaryocytes in primary myelodysplastic syndromes.. <i>Leukemia and Lymphoma</i> , <b>2021</b> , 1-9	1.9	
12	Spanish Guidelines for the use of targeted deep sequencing in myelodysplastic syndromes and chronic myelomonocytic leukaemia. <i>British Journal of Haematology</i> , <b>2020</b> , 188, 605-622	4.5	14
11	A CLK3-HMGA2 Alternative Splicing Axis Impacts Human Hematopoietic Stem Cell Molecular Identity throughout Development. <i>Cell Stem Cell</i> , <b>2018</b> , 22, 575-588.e7	18	24
10	Clinical Sequencing Uncovers Origins and Evolution of Lassa Virus. <i>Cell</i> , <b>2015</b> , 162, 738-50	56.2	176
9	Long noncoding RNAs regulate adipogenesis. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , <b>2013</b> , 110, 3387-92	11.5	315
8	Integrative annotation of human large intergenic noncoding RNAs reveals global properties and specific subclasses. <i>Genes and Development</i> , <b>2011</b> , 25, 1915-27	12.6	2492
7	Clinical value of NPHS2 analysis in early- and adult-onset steroid-resistant nephrotic syndrome. <i>Clinical Journal of the American Society of Nephrology: CJASN</i> , <b>2011</b> , 6, 344-54	6.9	58
6	Clinical utility of genetic testing in children and adults with steroid-resistant nephrotic syndrome. <i>Clinical Journal of the American Society of Nephrology: CJASN</i> , <b>2011</b> , 6, 1139-48	6.9	157
5	Evaluation of genome coverage and fidelity of multiple displacement amplification from single cells by SNP array. <i>Molecular Human Reproduction</i> , <b>2009</b> , 15, 739-47	4.4	34
4	Genetic testing for X-linked Alport syndrome by direct sequencing of COL4A5 cDNA from hair root RNA samples. <i>American Journal of Kidney Diseases</i> , <b>2007</b> , 50, 257.e1-14	7.4	25
3	Study of candidate genes affecting the progression of renal disease in autosomal dominant polycystic kidney disease type 1. <i>Nephrology Dialysis Transplantation</i> , <b>2007</b> , 22, 1567-77	4.3	19
2	Male-to-male transmission of X-linked Alport syndrome in a boy with a 47,XXY karyotype. <i>European Journal of Human Genetics</i> , <b>2005</b> , 13, 1040-6	5.3	17

- 1 Autosomal recessive Alport's syndrome and benign familial hematuria are collagen type IV diseases. *American Journal of Kidney Diseases*, **2003**, 42, 952-9 7.4 41