# Barbara Tazon-Vega

List of Publications by Year
in descending order



| 19 | 4,413 | 623188 | 839053 |
|---|---|---|---|
| papers | citations | 14 h-index | 18 g-index |

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

| # | Article | IF | Citations |
|---|---|---|---|
| 1 | Variant t(11;22)(q13;q11.2) withIGLinvolvement in mantle cell lymphoma. Leukemia and Lymphoma, 2022, 63, 1746-1749. | 0.6 | 1 |
| 2 | Cell free circulating tumor DNA in cerebrospinal fluid detects and monitors central nervous system involvement of B-cell lymphomas. Haematologica, 2021, 106, 513-521. | 1.7 | 75 |
| 3 | Usefulness of NGS for Diagnosis of Dominant Beta-Thalassemia and Unstable Hemoglobinopathies in Five Clinical Cases. Frontiers in Physiology, 2021, 12, 628236. | 1.3 | 7 |
| 4 | 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. | 0.6 | 2 |
| 5 | Immunological and genetic kinetics from diagnosis to clinical progression in chronic lymphocytic leukemia. Biomarker Research, 2021, 9, 37. | 2.8 | 5 |
| 6 | Prognostic impact of micromegakaryocytes in primary myelodysplastic syndromes. Leukemia and Lymphoma, 2021, , 1-9. | 0.6 | 0 |
| 7 | Spanish Guidelines for the use of targeted deep sequencing in myelodysplastic syndromes and chronic myelomonocytic leukaemia. British Journal of Haematology, 2020, 188, 605-622. | 1.2 | 25 |
| 8 | A CLK3-HMGA2 Alternative Splicing Axis Impacts Human Hematopoietic Stem Cell Molecular Identity throughout Development. Cell Stem Cell, 2018, 22, 575-588.e7. | 5.2 | 40 |
| 9 | Clinical Sequencing Uncovers Origins and Evolution of Lassa Virus. Cell, 2015, 162, 738-750. | 13.5 | 230 |
| 10 | Long noncoding RNAs regulate adipogenesis. Proceedings of the National Academy of Sciences of the United States of America, 2013, 110, 3387-3392. | 3.3 | 371 |
| 11 | Integrative annotation of human large intergenic noncoding RNAs reveals global properties and specific subclasses. Genes and Development, 2011, 25, 1915-1927. | 2.7 | 3,208 |
| 12 | 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. | 2.2 | 65 |
| 13 | 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. | 2.2 | 189 |
| 14 | Evaluation of genome coverage and fidelity of multiple displacement amplification from single cells by SNP array. Molecular Human Reproduction, 2009, 15, 739-747. | 1.3 | 36 |
| 15 | 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. | 0.4 | 25 |
| 16 | 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. | 2.1 | 27 |
| 17 | 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. | 1.4 | 21 |
| 18 | Collagen type IV (Â3-Â4) nephropathy: from isolated haematuria to renal failure. Nephrology Dialysis Transplantation, 2004, 19, 2429-2432. | 0.4 | 39 |

| # | Article | IF | Citations |
|---|---|---|---|
| 19 | Autosomal recessive Alportâ€™s syndrome and benign familial hematuria are collagen type IV diseases. American Journal of Kidney Diseases, 2003, 42, 952-959. | 2.1 | 47 |