

# Syed Khizer Hasan

## 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.

43  
papers

560  
citations

13  
h-index

22  
g-index

47  
ext. papers

657  
ext. citations

4.3  
avg, IF

3.18  
L-index

#	Paper	IF	Citations
43	Genetic ablation of pregnancy zone protein promotes breast cancer progression by activating TGF- $\beta$ /SMAD signaling. <i>Breast Cancer Research and Treatment</i> , <b>2021</b> , 185, 317-330	4.4	1
42	Clinical impact of panel-based error-corrected next generation sequencing versus flow cytometry to detect measurable residual disease (MRD) in acute myeloid leukemia (AML). <i>Leukemia</i> , <b>2021</b> , 35, 1392-1404	10.7	10
41	miRNA-mRNA Profiling Reveals Prognostic Impact of Expression in Acute Myeloid Leukemia. <i>Oncology Research</i> , <b>2020</b> , 28, 321-330	4.8	3
40	Characterization of therapy-related acute leukemia in hereditary breast-ovarian carcinoma patients: role of BRCA1 mutation and topoisomerase II-directed therapy. <i>Medical Oncology</i> , <b>2020</b> , 37, 48	3.7	0
39	Exome sequencing study of Russian breast cancer patients suggests a predisposing role for USP39. <i>Breast Cancer Research and Treatment</i> , <b>2020</b> , 179, 731-742	4.4	5
38	Over expression of brain and acute leukemia, cytoplasmic and ETS-related gene is associated with poor outcome in acute myeloid leukemia. <i>Hematological Oncology</i> , <b>2020</b> , 38, 808-816	1.3	1
37	A novel machine-learning-derived genetic score correlates with measurable residual disease and is highly predictive of outcome in acute myeloid leukemia with mutated NPM1. <i>Blood Cancer Journal</i> , <b>2019</b> , 9, 79	7	11
36	Utility of Immunophenotypic Measurable Residual Disease in Adult Acute Myeloid Leukemia-Real-World Context. <i>Frontiers in Oncology</i> , <b>2019</b> , 9, 450	5.3	6
35	Early and sensitive detection of PML-A216V mutation by droplet digital PCR in ATO-resistant acute promyelocytic leukemia. <i>Leukemia</i> , <b>2019</b> , 33, 1527-1530	10.7	10
34	Molecular Heterogeneity in Acute Promyelocytic Leukemia - a Single Center Experience from India. <i>Mediterranean Journal of Hematology and Infectious Diseases</i> , <b>2018</b> , 10, e2018002	3.2	5
33	Longitudinal detection of DNMT3A transcripts in patients with acute myeloid leukemia. <i>American Journal of Hematology</i> , <b>2018</b> , 93, E120-E123	7.1	6
32	Inhibition of novel GCN5-ATM axis restricts the onset of acquired drug resistance in leukemia. <i>International Journal of Cancer</i> , <b>2018</b> , 142, 2175-2185	7.5	13
31	Comparative genomic analysis of PML and RARA breakpoints in paired diagnosis/relapse samples of patients with acute promyelocytic leukemia treated with all-trans retinoic acid and chemotherapy. <i>Leukemia and Lymphoma</i> , <b>2018</b> , 59, 1268-1270	1.9	1
30	Long term clinical outcomes of adult hematolymphoid malignancies treated at Tata Memorial Hospital: An institutional audit. <i>Indian Journal of Cancer</i> , <b>2018</b> , 55, 9-15	0.9	5
29	Studies of protein-protein interactions in Fanconi anemia pathway to unravel the DNA interstrand crosslink repair mechanism. <i>International Journal of Biological Macromolecules</i> , <b>2017</b> , 104, 1338-1344	7.9	3
28	Structural and biophysical properties of h-FANCI ARM repeat protein. <i>Journal of Biomolecular Structure and Dynamics</i> , <b>2017</b> , 35, 3032-3042	3.6	0
27	The hidden genomic landscape of acute myeloid leukemia: subclonal structure revealed by undetected mutations. <i>Blood</i> , <b>2015</b> , 125, 600-5	2.2	14

26	Two novel methods for rapid detection and quantification of DNMT3A R882 mutations in acute myeloid leukemia. <i>Journal of Molecular Diagnostics</i> , <b>2015</b> , 17, 179-84	5.1	7
25	Understanding the molecular pathogenesis of acute promyelocytic leukemia. <i>Best Practice and Research in Clinical Haematology</i> , <b>2014</b> , 27, 3-9	4.2	55
24	Genomic analysis of therapy-related acute promyelocytic leukemias arising after malignant and non-malignant disorders. <i>American Journal of Hematology</i> , <b>2014</b> , 89, 346-7	7.1	2
23	Clustering of genomic breakpoints at the MLL locus in therapy-related acute leukemia with t(4;11)(q21;q23). <i>Genes Chromosomes and Cancer</i> , <b>2014</b> , 53, 248-54	5	5
22	Minimal Residual Disease by Multiparametric Flow Cytometry Predicts Relapse Free Survival better than Over-Expression of WT1 and BAALC in Acute Myeloid Leukemia. <i>Blood</i> , <b>2014</b> , 124, 1064-1064	2.2	
21	Identification of emerging FLT3 ITD-positive clones during clinical remission and kinetics of disease relapse in acute myeloid leukaemia with mutated nucleophosmin. <i>British Journal of Haematology</i> , <b>2013</b> , 161, 533-40	4.5	30
20	Rapid detection of IDH2 (R140Q and R172K) mutations in acute myeloid leukemia. <i>Annals of Hematology</i> , <b>2013</b> , 92, 1319-23	3	7
19	Biology and management of therapy-related acute promyelocytic leukemia. <i>Current Opinion in Oncology</i> , <b>2013</b> , 25, 695-700	4.2	12
18	Comparative molecular analysis of therapy-related and de novo acute promyelocytic leukemia. <i>Leukemia Research</i> , <b>2012</b> , 36, 474-8	2.7	12
17	Acute myeloid leukemia developing in patients with autoimmune diseases. <i>Haematologica</i> , <b>2012</b> , 97, 805-17	6.6	50
16	Presenting features and treatment outcome of acute promyelocytic leukemia arising after multiple sclerosis. <i>Haematologica</i> , <b>2011</b> , 96, 621-5	6.6	21
15	Molecular pathogenesis of secondary acute promyelocytic leukemia. <i>Mediterranean Journal of Hematology and Infectious Diseases</i> , <b>2011</b> , 3, e2011045	3.2	23
14	Risk of acute promyelocytic leukemia in multiple sclerosis: coding variants of DNA repair genes. <i>Neurology</i> , <b>2011</b> , 76, 1059-65	6.5	32
13	Utilization of molecular phenotypes to detect relapse and optimize the management of acute promyelocytic leukemia. <i>Clinical Lymphoma, Myeloma and Leukemia</i> , <b>2010</b> , 10 Suppl 3, S139-43	2	
12	Evidence for direct involvement of epirubicin in the formation of chromosomal translocations in t(15;17) therapy-related acute promyelocytic leukemia. <i>Blood</i> , <b>2010</b> , 115, 326-30	2.2	55
11	Analysis of t(15;17) chromosomal breakpoint sequences in therapy-related versus de novo acute promyelocytic leukemia: association of DNA breaks with specific DNA motifs at PML and RARA loci. <i>Genes Chromosomes and Cancer</i> , <b>2010</b> , 49, 726-32	5	29
10	Identification of a potential "hotspot" DNA region in the RUNX1 gene targeted by mitoxantrone in therapy-related acute myeloid leukemia with t(16;21) translocation. <i>Genes Chromosomes and Cancer</i> , <b>2009</b> , 48, 213-21	5	20
9	Molecular analysis of t(15;17) genomic breakpoints in secondary acute promyelocytic leukemia arising after treatment of multiple sclerosis. <i>Blood</i> , <b>2008</b> , 112, 3383-90	2.2	68

8	Molecular Characterization of the t(15;17)(q22;21) in Epirubicin-Related Acute Promyelocytic Leukaemia. <i>Blood</i> , <b>2008</b> , 112, 791-791	2.2	
7	Timing of allogeneic BMT for CML patients in India: does it affect response?. <i>American Journal of Hematology</i> , <b>2007</b> , 82, 329	7.1	
6	Impact of FLT3 internal tandem duplications on Indian acute promyelocytic leukemia patients: prognostic implications. <i>Hematology</i> , <b>2007</b> , 12, 99-101	2.2	9
5	Childhood CML in India: b2a2 transcript is more common than b3a2. <i>Cancer Genetics and Cytogenetics</i> , <b>2006</b> , 169, 76-7		19
4	Acute promyelocytic leukemia with secondary myelofibrosis -- case report and review of the literature. <i>American Journal of Hematology</i> , <b>2006</b> , 81, 476-7	7.1	7
3	Over-representation of bcr3 subtype of PML/RARalpha fusion gene in APL in Indian patients. <i>Annals of Hematology</i> , <b>2005</b> , 84, 781-4	3	2
2	The Pathophysiology of Acute Promyelocytic Leukemia161-168		
1	Acute promyelocytic leukemia: pathophysiology and clinical results update131-140		