

Niccolo Bolli

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136
papers

12,925
citations

37
h-index

113
g-index

154
ext. papers

16,340
ext. citations

7.3
avg, IF

5.16
L-index

#	Paper	IF	Citations
136	Signatures of mutational processes in human cancer. <i>Nature</i> , 2013 , 500, 415-21	50.4	5895
135	Genomic Classification and Prognosis in Acute Myeloid Leukemia. <i>New England Journal of Medicine</i> , 2016 , 374, 2209-2221	59.2	1999
134	Heterogeneity of genomic evolution and mutational profiles in multiple myeloma. <i>Nature Communications</i> , 2014 , 5, 2997	17.4	564
133	Mobile DNA in cancer. Extensive transduction of nonrepetitive DNA mediated by L1 retrotransposition in cancer genomes. <i>Science</i> , 2014 , 345, 1251343	33.3	250
132	Origins and functional consequences of somatic mitochondrial DNA mutations in human cancer. <i>ELife</i> , 2014 , 3,	8.9	229
131	Association of a germline copy number polymorphism of APOBEC3A and APOBEC3B with burden of putative APOBEC-dependent mutations in breast cancer. <i>Nature Genetics</i> , 2014 , 46, 487-91	36.3	208
130	Both carboxy-terminus NES motif and mutated tryptophan(s) are crucial for aberrant nuclear export of nucleophosmin leukemic mutants in NPMc+ AML. <i>Blood</i> , 2006 , 107, 4514-23	2.2	201
129	Identification of novel mutational drivers reveals oncogene dependencies in multiple myeloma. <i>Blood</i> , 2018 , 132, 587-597	2.2	196
128	Acute myeloid leukemia with mutated nucleophosmin (NPM1): is it a distinct entity?. <i>Blood</i> , 2011 , 117, 1109-20	2.2	178
127	A high-risk, Double-Hit, group of newly diagnosed myeloma identified by genomic analysis. <i>Leukemia</i> , 2019 , 33, 159-170	10.7	176
126	Altered nucleophosmin transport in acute myeloid leukaemia with mutated NPM1: molecular basis and clinical implications. <i>Leukemia</i> , 2009 , 23, 1731-43	10.7	156
125	Translocations and mutations involving the nucleophosmin (NPM1) gene in lymphomas and leukemias. <i>Haematologica</i> , 2007 , 92, 519-32	6.6	156
124	Precision oncology for acute myeloid leukemia using a knowledge bank approach. <i>Nature Genetics</i> , 2017 , 49, 332-340	36.3	155
123	Immunohistochemistry predicts nucleophosmin (NPM) mutations in acute myeloid leukemia. <i>Blood</i> , 2006 , 108, 1999-2005	2.2	146
122	Cell line OCI/AML3 bears exon-12 NPM gene mutation-A and cytoplasmic expression of nucleophosmin. <i>Leukemia</i> , 2005 , 19, 1760-7	10.7	124
121	Pan-cancer analysis of whole genomes identifies driver rearrangements promoted by LINE-1 retrotransposition. <i>Nature Genetics</i> , 2020 , 52, 306-319	36.3	122
120	KLF2 mutation is the most frequent somatic change in splenic marginal zone lymphoma and identifies a subset with distinct genotype. <i>Leukemia</i> , 2015 , 29, 1177-85	10.7	118

119	Genomic patterns of progression in smoldering multiple myeloma. <i>Nature Communications</i> , 2018 , 9, 3363-7.4	99
118	Cytoplasmic mutated nucleophosmin (NPM) defines the molecular status of a significant fraction of myeloid sarcomas. <i>Leukemia</i> , 2007 , 21, 1566-70	10.7 97
117	Genomic landscape and chronological reconstruction of driver events in multiple myeloma. <i>Nature Communications</i> , 2019 , 10, 3835	17.4 94
116	Analysis of the genomic landscape of multiple myeloma highlights novel prognostic markers and disease subgroups. <i>Leukemia</i> , 2018 , 32, 2604-2616	10.7 90
115	Mutated nucleophosmin detects clonal multilineage involvement in acute myeloid leukemia: Impact on WHO classification. <i>Blood</i> , 2006 , 108, 4146-55	2.2 82
114	Born to be exported: COOH-terminal nuclear export signals of different strength ensure cytoplasmic accumulation of nucleophosmin leukemic mutants. <i>Cancer Research</i> , 2007 , 67, 6230-7	10.1 81
113	Hemopoietic-specific Sf3b1-K700E knock-in mice display the splicing defect seen in human MDS but develop anemia without ring sideroblasts. <i>Leukemia</i> , 2017 , 31, 720-727	10.7 76
112	A practical guide for mutational signature analysis in hematological malignancies. <i>Nature Communications</i> , 2019 , 10, 2969	17.4 73
111	Processed pseudogenes acquired somatically during cancer development. <i>Nature Communications</i> , 2014 , 5, 3644	17.4 68
110	Expression of the cytoplasmic NPM1 mutant (NPMc+) causes the expansion of hematopoietic cells in zebrafish. <i>Blood</i> , 2010 , 115, 3329-40	2.2 61
109	A dose-dependent tug of war involving the NPM1 leukaemic mutant, nucleophosmin, and ARF. <i>Leukemia</i> , 2009 , 23, 501-9	10.7 59
108	Biological and prognostic impact of APOBEC-induced mutations in the spectrum of plasma cell dyscrasias and multiple myeloma cell lines. <i>Leukemia</i> , 2018 , 32, 1044-1048	10.7 52
107	Multiple myeloma clonal evolution in homogeneously treated patients. <i>Leukemia</i> , 2018 , 32, 2636-2647	10.7 51
106	Zebrafish microRNA-126 determines hematopoietic cell fate through c-Myb. <i>Leukemia</i> , 2011 , 25, 506-14	10.7 49
105	A DNA target-enrichment approach to detect mutations, copy number changes and immunoglobulin translocations in multiple myeloma. <i>Blood Cancer Journal</i> , 2016 , 6, e467	7 47
104	Identification and functional characterization of a cytoplasmic nucleophosmin leukaemic mutant generated by a novel exon-11 NPM1 mutation. <i>Leukemia</i> , 2007 , 21, 1099-103	10.7 46
103	Differential and limited expression of mutant alleles in multiple myeloma. <i>Blood</i> , 2014 , 124, 3110-7	2.2 42
102	Cytoplasmic mutated nucleophosmin is stable in primary leukemic cells and in a xenotransplant model of NPMc+ acute myeloid leukemia in SCID mice. <i>Haematologica</i> , 2008 , 93, 775-9	6.6 42

101	Characterization of gene mutations and copy number changes in acute myeloid leukemia using a rapid target enrichment protocol. <i>Haematologica</i> , 2015 , 100, 214-22	6.6	38
100	Aberrant subcellular expression of nucleophosmin and NPM-MLF1 fusion protein in acute myeloid leukaemia carrying t(3;5): a comparison with NPMc+ AML. <i>Leukemia</i> , 2006 , 20, 368-71	10.7	37
99	Timing the initiation of multiple myeloma. <i>Nature Communications</i> , 2020 , 11, 1917	17.4	36
98	In human genome, generation of a nuclear export signal through duplication appears unique to nucleophosmin (NPM1) mutations and is restricted to AML. <i>Leukemia</i> , 2008 , 22, 1285-9	10.7	36
97	Development and validation of a comprehensive genomic diagnostic tool for myeloid malignancies. <i>Blood</i> , 2016 , 128, e1-9	2.2	36
96	Tumor protein D52 (TPD52): a novel B-cell/plasma-cell molecule with unique expression pattern and Ca(2+)-dependent association with annexin VI. <i>Blood</i> , 2005 , 105, 2812-20	2.2	34
95	Aberrant cytoplasmic expression of C-terminal-truncated NPM leukaemic mutant is dictated by tryptophans loss and a new NES motif. <i>Leukemia</i> , 2007 , 21, 2052-4; author reply 2054; discussion 2055-6	10.7	32
94	Ddx18 is essential for cell-cycle progression in zebrafish hematopoietic cells and is mutated in human AML. <i>Blood</i> , 2011 , 118, 903-15	2.2	29
93	Angioimmunoblastic T cell lymphoma: novel molecular insights by mutation profiling. <i>Oncotarget</i> , 2017 , 8, 17763-17770	3.3	29
92	Revealing the impact of structural variants in multiple myeloma. <i>Blood Cancer Discovery</i> , 2020 , 1, 258-273		28
91	Detailed molecular characterisation of acute myeloid leukaemia with a normal karyotype using targeted DNA capture. <i>Leukemia</i> , 2013 , 27, 1820-5	10.7	26
90	The human NPM1 mutation A perturbs megakaryopoiesis in a conditional mouse model. <i>Blood</i> , 2013 , 121, 3447-58	2.2	25
89	cpsf1 is required for definitive HSC survival in zebrafish. <i>Blood</i> , 2011 , 117, 3996-4007	2.2	25
88	Moving From Cancer Burden to Cancer Genomics for Smoldering Myeloma: A Review. <i>JAMA Oncology</i> , 2020 , 6, 425-432	13.4	25
87	Classification and Personalized Prognostic Assessment on the Basis of Clinical and Genomic Features in Myelodysplastic Syndromes. <i>Journal of Clinical Oncology</i> , 2021 , 39, 1223-1233	2.2	25
86	Cytoplasmic nucleophosmin in myeloid sarcoma occurring 20 years after diagnosis of acute myeloid leukaemia. <i>Lancet Oncology</i> , 2006 , 7, 350-2	21.7	24
85	Early Relapse Risk in Patients with Newly Diagnosed Multiple Myeloma Characterized by Next-generation Sequencing. <i>Clinical Cancer Research</i> , 2020 , 26, 4832-4841	12.9	22
84	Comprehensive detection of recurring genomic abnormalities: a targeted sequencing approach for multiple myeloma. <i>Blood Cancer Journal</i> , 2019 , 9, 101	7	22

83	Absence of nucleophosmin leukaemic mutants in B and T cells from AML with NPM1 mutations: implications for the cell of origin of NPMc+ AML. <i>Leukemia</i> , 2008 , 22, 195-8	10.7	21
82	Integrative analysis of the genomic and transcriptomic landscape of double-refractory multiple myeloma. <i>Blood Advances</i> , 2020 , 4, 830-844	7.8	21
81	Next-Generation Sequencing for Clinical Management of Multiple Myeloma: Ready for Prime Time?. <i>Frontiers in Oncology</i> , 2020 , 10, 189	5.3	20
80	A western blot assay for detecting mutant nucleophosmin (NPM1) proteins in acute myeloid leukaemia. <i>Leukemia</i> , 2008 , 22, 2285-8	10.7	20
79	A one-mutation mathematical model can explain the age incidence of acute myeloid leukemia with mutated nucleophosmin (NPM1). <i>Haematologica</i> , 2008 , 93, 1219-26	6.6	20
78	Prognostic impact of genetic characterization in the GIMEMA LAM99P multicenter study for newly diagnosed acute myeloid leukemia. <i>Haematologica</i> , 2008 , 93, 1017-24	6.6	16
77	Whole-genome sequencing reveals progressive versus stable myeloma precursor conditions as two distinct entities. <i>Nature Communications</i> , 2021 , 12, 1861	17.4	16
76	Sustained ventricular tachycardia in a thalidomide-treated patient with primary plasma-cell leukemia. <i>Nature Clinical Practice Oncology</i> , 2007 , 4, 722-5		11
75	Pan-cancer analysis of whole genomes reveals driver rearrangements promoted by LINE-1 retrotransposition in human tumours		10
74	Analysis of the genomic landscape of multiple myeloma highlights novel prognostic markers and disease subgroups. <i>Leukemia</i> , 2017 ,	10.7	9
73	Treating two concurrent B-cell and T-cell lymphoid neoplasms with alemtuzumab monotherapy. <i>Lancet Oncology</i> , 2004 , 5, 64-5	21.7	9
72	Biology of peripheral T cell lymphomas [Not otherwise specified: Is something finally happening? 2016 , 3, 9-18		9
71	Next-generation sequencing of a family with a high penetrance of monoclonal gammopathies for the identification of candidate risk alleles. <i>Cancer</i> , 2017 , 123, 3701-3708	6.4	8
70	Integration of transcriptional and mutational data simplifies the stratification of peripheral T-cell lymphoma. <i>American Journal of Hematology</i> , 2019 , 94, 628-634	7.1	7
69	Genomic landscape and chronological reconstruction of driver events in multiple myeloma		6
68	Clinical relevance of clonal hematopoiesis in persons aged 80 years. <i>Blood</i> , 2021 , 138, 2093-2105	2.2	6
67	Recurrent histone mutations in T-cell acute lymphoblastic leukaemia. <i>British Journal of Haematology</i> , 2019 , 184, 676-679	4.5	6
66	Limits and Applications of Genomic Analysis of Circulating Tumor DNA as a Liquid Biopsy in Asymptomatic Forms of Multiple Myeloma. <i>HemaSphere</i> , 2020 , 4, e402	0.3	5

65	mmsig: a fitting approach to accurately identify somatic mutational signatures in hematological malignancies. <i>Communications Biology</i> , 2021 , 4, 424	6.7	5
64	Application of Next-Generation Sequencing for the Genomic Characterization of Patients with Smoldering Myeloma. <i>Cancers</i> , 2020 , 12,	6.6	4
63	High-dose chemotherapy followed by autologous transplantation may overcome the poor prognosis of diffuse large B-cell lymphoma patients with MYC/BCL2 co-expression. <i>Blood Cancer Journal</i> , 2016 , 6, e491	7	4
62	Sf3b1 K700E Mutation Impairs Pre-mRNA Splicing and Definitive Hematopoiesis in a Conditional Knock-in Mouse Model. <i>Blood</i> , 2015 , 126, 140-140	2.2	4
61	A Journey Through Myeloma Evolution: From the Normal Plasma Cell to Disease Complexity. <i>HemaSphere</i> , 2020 , 4, e502	0.3	4
60	Cereblon enhancer methylation and IMiD resistance in multiple myeloma. <i>Blood</i> , 2021 , 138, 1721-1726	2.2	4
59	2021 European Myeloma Network review and consensus statement on smoldering multiple myeloma: how to distinguish (and manage) Dr. Jekyll and Mr. Hyde. <i>Haematologica</i> , 2021 , 106, 2799-2812	6.6	4
58	Rapid parallel acquisition of somatic mutations after NPM1 in acute myeloid leukaemia evolution. <i>British Journal of Haematology</i> , 2017 , 176, 825-829	4.5	3
57	Timing the Initiation of Multiple Myeloma. <i>SSRN Electronic Journal</i> ,	1	3
56	CDKN2A deletion is a frequent event associated with poor outcome in patients with peripheral T-cell lymphoma not otherwise specified (PTCL-NOS). <i>Haematologica</i> , 2021 , 106, 2918-2926	6.6	3
55	Specific targeting of the KRAS mutational landscape in myeloma as a tool to unveil the elicited antitumor activity. <i>Blood</i> , 2021 , 138, 1705-1720	2.2	3
54	Single- and double-hit events in genes encoding immune targets before and after T cell-engaging antibody therapy in MM. <i>Blood Advances</i> , 2021 , 5, 3794-3798	7.8	3
53	Reply to Pitiot et al. <i>Leukemia</i> , 2007 , 21, 2055-2056	10.7	2
52	Whole-Genome Sequencing Reveals Evidence of Two Biologically and Clinically Distinct Entities: Progressive Versus Stable Myeloma Precursor Disease. <i>Blood</i> , 2020 , 136, 47-48	2.2	2
51	A Next Generation Sequencing-Based Approach to Detect Gene Mutations, Copy Number Changes and IGH Translocations in Multiple Myeloma. <i>Blood</i> , 2014 , 124, 3364-3364	2.2	2
50	Dissecting Genetic and Phenotypic Heterogeneity to Map Molecular Phylogenies and Deliver Personalized Outcome and Treatment Predictions in AML. <i>Blood</i> , 2015 , 126, 803-803	2.2	2
49	Analysis of Mutational Signatures Suggest That Aid Has an Early and Driver Role in Multiple Myeloma. <i>Blood</i> , 2016 , 128, 116-116	2.2	2
48	Peripheral T-Cell Lymphomas Not Otherwise Specified: Potential Novel Molecular Entities Based on Both Tumor and Microenvironment Cellular Components. <i>Blood</i> , 2016 , 128, 4098-4098	2.2	2

47	The molecular pathogenesis of multiple myeloma. <i>Hematology Reports</i> , 2020 , 12, 9054	0.9	2
46	DIS3 mutations in multiple myeloma impact the transcriptional signature and clinical outcome. <i>Haematologica</i> , 2021 ,	6.6	2
45	What Is New in the Treatment of Smoldering Multiple Myeloma?. <i>Journal of Clinical Medicine</i> , 2021 , 10,	5.1	2
44	Copy number signatures predict chromothripsis and clinical outcomes in newly diagnosed multiple myeloma. <i>Nature Communications</i> , 2021 , 12, 5172	17.4	2
43	Bowel perforation from occult ileal involvement after diagnosis in a case of primary mediastinal large B-cell lymphoma. <i>BMJ Case Reports</i> , 2016 , 2016,	0.9	1
42	Reply to Aberrant cytoplasmic expression of C-terminal truncated NPM leukaemic mutant is dictated by tryptophans loss and a new NES motif by Falini et al.. <i>Leukemia</i> , 2007 , 21, 2054-2054	10.7	1
41	The Genomic and Transcriptomic Landscape of Double-Refractory Multiple Myeloma. <i>Blood</i> , 2019 , 134, 3056-3056	2.2	1
40	Specific Targeting of KRAS Using a Novel High-Affinity KRAS Antisense Oligonucleotide in Multiple Myeloma. <i>Blood</i> , 2019 , 134, 3104-3104	2.2	1
39	Genomic Landscape and Its Prognostic Implications in Multiple Myeloma Using a Targeted Sequencing Approach. <i>Blood</i> , 2015 , 126, 370-370	2.2	1
38	GATA-3 Expression in Peripheral T-Cell Lymphomas (PTCL): Identification of a Cut-Off and Prognostic Value in PTCL-NOS Versus Others Hystotypes. <i>Blood</i> , 2015 , 126, 3889-3889	2.2	1
37	Personally Tailored Risk Prediction of AML Based on Comprehensive Genomic and Clinical Data. <i>Blood</i> , 2015 , 126, 85-85	2.2	1
36	Chemotherapy-Related Mutational Signatures Reveal the Origins of Therapy-Related Myeloid Neoplasms. <i>Blood</i> , 2021 , 138, 3271-3271	2.2	1
35	Revealing the impact of recurrent and rare structural variants in multiple myeloma		1
34	Whole genome sequencing provides evidence of two biologically and clinically distinct entities of asymptomatic monoclonal gammopathies: progressive versus stable myeloma precursor condition		1
33	A rare case of atypical chronic lymphocytic leukaemia presenting as nephrotic syndrome. <i>BMJ Case Reports</i> , 2017 , 2017,	0.9	1
32	ROBO1 Promotes Homing, Dissemination, and Survival of Multiple Myeloma within the Bone Marrow Microenvironment. <i>Blood Cancer Discovery</i> , 2021 , 2, 338-353	7	1
31	Functional Impact of Genomic Complexity on the Transcriptome of Multiple Myeloma. <i>Clinical Cancer Research</i> , 2021 , 27, 6479-6490	12.9	1
30	The 2020 BMT CTN Myeloma Intergroup Workshop on Immune Profiling and Minimal Residual Disease Testing in Multiple Myeloma. <i>Transplantation and Cellular Therapy</i> , 2021 , 27, 807-816		1

29	Redefining Mutational Profiling Using RNA-Seq: Insight into the Functional Mutational Landscape of Multiple Myeloma. <i>Blood</i> , 2015 , 126, 837-837	2.2	○
28	Whole Genome Sequencing of Unique Paired SMM/MGUS Progressing to MM Samples Reveals a Genomic Landscape with Diverse Evolutionary Pattern. <i>Blood</i> , 2016 , 128, 2088-2088	2.2	○
27	CD40 Activity on Mesenchymal Cells Negatively Regulates OX40L to Maintain Bone Marrow Immune Homeostasis Under Stress Conditions. <i>Frontiers in Immunology</i> , 2021 , 12, 662048	8.4	○
26	Clinical, Morphological and Clonal Progression of VEXAS Syndrome in the Context of Myelodysplasia Treated with Azacytidine. <i>Clinical Hematology International</i> , 1	1.8	○
25	The Molecular Basis of Haematological Malignancies 2015 , 314-331		
24	Revealing Transcriptome Deregulation upon Genomic Complexity in Multiple Myeloma. <i>Blood</i> , 2020 , 136, 3-4	2.2	
23	Dissection of Bone Marrow Microenvironment By Single Cell RNA Sequencing in Warm AIHA Patients: A Proof-of-Concept Analysis. <i>Blood</i> , 2021 , 138, 931-931	2.2	
22	MGUS and Chip: Two Faces, but Not of the Same Medal. <i>Blood</i> , 2021 , 138, 3800-3800	2.2	
21	Tumor Protein D52 (TPD52): A Novel B Cell/Plasma Cell Molecule Identified through a Proteomic Approach and Characterized by Unique Expression Pattern and Ca ²⁺ -Dependent Association with Annexin VI.. <i>Blood</i> , 2004 , 104, 3652-3652	2.2	
20	Exon-12 Nucleophosmin (NPM) Mutation and Aberrant Cytoplasmic Expression of NPM Protein in Leukemia Cell Line OCI-AML3.. <i>Blood</i> , 2005 , 106, 2376-2376	2.2	
19	Prognostic Impact of Genetic Characterization in the GIMEMA LAM99P Study for Newly Diagnosed Adult AML. Relevance of Combined Analysis of Conventional Karyotyping, FLT3 and NPM Mutational Status.. <i>Blood</i> , 2005 , 106, 226-226	2.2	
18	Mechanism of Altered Nucleo-Cytoplasmic Traffic of Nucleophosmin in Acute Myelogenous Leukemia Carrying Exon-12 NPM Mutations (NPMc+ AML).. <i>Blood</i> , 2005 , 106, 4396-4396	2.2	
17	Reciprocal Interaction between NPM Leukemic Mutants and Arf: Structural Basis and Functional Consequences.. <i>Blood</i> , 2006 , 108, 1939-1939	2.2	
16	Function of Nucleophosmin in Zebrafish Hematopoiesis.. <i>Blood</i> , 2007 , 110, 2644-2644	2.2	
15	The Landscape of Structural Variant Signatures in Multiple Myeloma Identifies Distinct Disease Subgroups with Implications for Pathogenesis. <i>Blood</i> , 2018 , 132, 109-109	2.2	
14	Mytype: A Capture Based Sequencing Approach to Detect Somatic Mutations, Copy Number Changes and IGH Translocations in Multiple Myeloma. <i>Blood</i> , 2018 , 132, 5588-5588	2.2	
13	Whole Genome Sequencing Reveals Recurrent Structural Driver Events in Peripheral T-Cell Lymphomas Not Otherwise Specified. <i>Blood</i> , 2018 , 132, 4115-4115	2.2	
12	Clinical Relevance of Clonal Hematopoiesis in the Oldest-Old Population: Analysis of the "Health and Anemia" Study. <i>Blood</i> , 2018 , 132, 750-750	2.2	

- 11 Lack of Significant Differences in Somatic Alterations between MGUS, SMM and Symptomatic Multiple Myeloma: A Result from Comprehensive Genomic Profiling Study. *Blood*, **2019**, 134, 3089-3089 2.2
- 10 Timing the Initiation of Multiple Myeloma. *Blood*, **2019**, 134, 573-573 2.2
- 9 Differential and Limited Expression of Mutant Alleles in Multiple Myeloma. *Blood*, **2014**, 124, 2007-2007 2.2
- 8 The Complex Landscape of Rearrangements in Smoldering and Symptomatic Multiple Myeloma Revealed By Whole-Genome Sequencing. *Blood*, **2016**, 128, 236-236 2.2
- 7 RNA-Seq De Novo Assembly of Clonal Immunoglobulin Rearrangements Identifies Interesting Biology and Uncovers Prognostic Features in Multiple Myeloma. *Blood*, **2016**, 128, 195-195 2.2
- 6 The Role of RNA Helicase Dead Box 18 in Zebrafish Hematopoiesis and Human MDS. *Blood*, **2008**, 112, 500-500 2.2
- 5 Cleavage and Polyadenylation Specificity Factor 1 Is Required for Definitive Hematopoietic Stem Cell Survival In Zebrafish.. *Blood*, **2010**, 116, 1606-1606 2.2
- 4 Upregulation of eIF4E in Nucleophosmin 1 (NPM1) Haploinsufficient Cells Alters CCAAT Enhancer Binding Protein Alpha (C/EBP β) Activity: Implications for MDS and AML. *Blood*, **2011**, 118, 2432-2432 2.2
- 3 Whole Exome Sequencing Of Multiple Myeloma Reveals An Heterogeneous Clonal Architecture and Genomic Evolution. *Blood*, **2013**, 122, 399-399 2.2
- 2 A new case of myelodysplastic syndrome associated with t(3;3)(q21;q26) and inv(11)(p15q22). *Tumori*, **2020**, 106, NP18-NP22 1.7
- 1 PS1345 CIRCULATING TUMOR DNA AS A LIQUID BIOPSY IN SMOLDERING MULTIPLE MYELOMA TO IDENTIFY BIOMARKERS OF PROGRESSION. *HemaSphere*, **2019**, 3, 614-615 0.3