

Jaroslav P Maciejewski

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.

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|--------------------|-------------------------|---------------|-----------------|
| 314 papers | 7,694 citations | 37 h-index | 85 g-index |
| 328 ext. papers | 9,848 ext. citations | 5 avg, IF | 5.59 L-index |

| # | Paper | IF | Citations |
|-----------------|---|------|-----------|
| 3 ¹⁴ | Eltrombopag inhibits TET dioxygenase to contribute to hematopoietic stem cell expansion in aplastic anemia.. <i>Journal of Clinical Investigation</i> , 2022 , | 15.9 | 1 |
| 3 ¹³ | Clonal dynamics of hematopoietic stem cell compartment in aplastic anemia.. <i>Seminars in Hematology</i> , 2022 , 59, 47-53 | 4 | 0 |
| 3 ¹² | A study of Telomerase Reverse Transcriptase rare variants in myeloid neoplasia.. <i>Hematological Oncology</i> , 2022 , | 1.3 | 0 |
| 3 ¹¹ | Recruitment of MLL1 complex is essential for SETBP1 to induce myeloid transformation.. <i>IScience</i> , 2022 , 25, 103679 | 6.1 | 0 |
| 3 ¹⁰ | Circulating microbial content in myeloid malignancy patients is associated with disease subtypes and patient outcomes.. <i>Nature Communications</i> , 2022 , 13, 1038 | 17.4 | 2 |
| 3 ⁰⁹ | Aplastic anemia: Quo vadis?. <i>Seminars in Hematology</i> , 2022 , 59, 54-55 | 4 | 0 |
| 3 ⁰⁸ | Single-cell characterization of leukemic and non-leukemic immune repertoires in CD8 T-cell large granular lymphocytic leukemia.. <i>Nature Communications</i> , 2022 , 13, 1981 | 17.4 | 2 |
| 3 ⁰⁷ | The Similarity of Class II HLA Genotypes Defines Patterns of Autoreactivity in Idiopathic Bone Marrow Failure Disorders. <i>Blood</i> , 2021 , | 2.2 | 2 |
| 3 ⁰⁶ | Immunogenetic, Molecular and Clinical Determinants of Clonal Evolution in Aplastic Anemia and Paroxysmal Nocturnal Hemoglobinuria. <i>Blood</i> , 2021 , 138, 602-602 | 2.2 | |
| 3 ⁰⁵ | Epigenetic Enzyme Mutations in Myeloid Malignancies Are Selected By Chromatin-Remodeling Requirements That Vary By Lineage- and Maturation-Stage. <i>Blood</i> , 2021 , 138, 1148-1148 | 2.2 | |
| 3 ⁰⁴ | A Novel Machine Learning-Derived Molecular Classification Scheme with Prognostic Significance. <i>Blood</i> , 2021 , 138, 3666-3666 | 2.2 | 0 |
| 3 ⁰³ | EPOR/JAK/STAT Signaling Pathway As Therapeutic Target of Acute Erythroid Leukemia. <i>Blood</i> , 2021 , 138, 610-610 | 2.2 | 1 |
| 3 ⁰² | A Novel Approach to Induce ATRA Mediated Differentiation in NPM1 Mutant Acute Myeloid Leukemia. <i>Blood</i> , 2021 , 138, 786-786 | 2.2 | |
| 3 ⁰¹ | Is nature truly healing itself? Spontaneous remissions in Paroxysmal Nocturnal Hemoglobinuria. <i>Blood Cancer Journal</i> , 2021 , 11, 187 | 7 | 2 |
| 3 ⁰⁰ | Is Nature Truly Healing Itself? Spontaneous Remissions and Clonal Replacement in Paroxysmal Nocturnal Hemoglobinuria. <i>Blood</i> , 2021 , 138, 4303-4303 | 2.2 | |
| 299 | Mutant TP53 prevents Telomere Shortening in Acute Myeloid Leukemia. <i>Blood</i> , 2021 , 138, 375-375 | 2.2 | 0 |
| 298 | A Systematic Review and Meta-Analysis Comparing Type I and II FLT3 Inhibitors in Relapsed/Refractory Acute Myeloid Leukemia and High-Risk Myelodysplastic Syndrome. <i>Blood</i> , 2021 , 138, 1249-1249 | 2.2 | 0 |

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| 297 | Spectrum of Molecular Modes of Immune Escape in Idiopathic Aplastic Anemia and Paroxysmal Nocturnal Hemoglobinuria. <i>Blood</i> , 2021 , 138, 603-603 | 2.2 | 0 |
| 296 | Genomic Data Improves Prognostic Stratification in Adult T-Cell Acute Lymphoblastic Leukemia Patients Enrolled in Measurable Residual Disease-Oriented Trials. <i>Blood</i> , 2021 , 138, 3486-3486 | 2.2 | 1 |
| 295 | Molecular Signatures of Immune Pressure and Immune Escape in Hematological Malignancies. <i>Blood</i> , 2021 , 138, 1093-1093 | 2.2 | |
| 294 | Transcriptomic Profile Identifies Early Signatures of Immunoediting and a Potential Role for VISTA As a Molecular Target in Acute Myeloid Leukemia. <i>Blood</i> , 2021 , 138, 4467-4467 | 2.2 | |
| 293 | Therapeutic Targeting of TET-Dioxygenase Deficiency in Myeloid Malignancies. <i>Blood</i> , 2021 , 138, 3985-3985 | 2.2 | 0 |
| 292 | Vacuolization of hematopoietic precursors: an enigma with multiple etiologies. <i>Blood</i> , 2021 , 137, 3685-3689 | 2.2 | 12 |
| 291 | Somatic mutations in lymphocytes in patients with immune-mediated aplastic anemia. <i>Leukemia</i> , 2021 , 35, 1365-1379 | 10.7 | 10 |
| 290 | How I manage acquired pure red cell aplasia in adults. <i>Blood</i> , 2021 , 137, 2001-2009 | 2.2 | 7 |
| 289 | Functional analyses of human LUC7-like proteins involved in splicing regulation and myeloid neoplasms. <i>Cell Reports</i> , 2021 , 35, 108989 | 10.6 | 4 |
| 288 | Therapeutic Targeting of Protein Disulfide Isomerase PDIA1 in Multiple Myeloma. <i>Cancers</i> , 2021 , 13, | 6.6 | 2 |
| 287 | Machine learning integrates genomic signatures for subclassification beyond primary and secondary acute myeloid leukemia. <i>Blood</i> , 2021 , 138, 1885-1895 | 2.2 | 3 |
| 286 | Complex landscape of alternative splicing in myeloid neoplasms. <i>Leukemia</i> , 2021 , 35, 1108-1120 | 10.7 | 12 |
| 285 | Decitabine- and 5-azacytidine resistance emerges from adaptive responses of the pyrimidine metabolism network. <i>Leukemia</i> , 2021 , 35, 1023-1036 | 10.7 | 19 |
| 284 | Frequency and perturbations of various peripheral blood cell populations before and after eculizumab treatment in paroxysmal nocturnal hemoglobinuria. <i>Blood Cells, Molecules, and Diseases</i> , 2021 , 87, 102528 | 2.1 | 3 |
| 283 | Dexrazoxane enhances efficacy of all- retinoic acid in acute myeloid leukemia patient blast cells and cell lines. <i>Leukemia and Lymphoma</i> , 2021 , 62, 473-477 | 1.9 | |
| 282 | Analysis of distinct hotspot mutations in relation to clinical phenotypes and response to therapy in myeloid neoplasia. <i>Leukemia and Lymphoma</i> , 2021 , 62, 735-738 | 1.9 | 2 |
| 281 | Reduced red blood cell surface level of Factor H as a mechanism underlying paroxysmal nocturnal hemoglobinuria. <i>Leukemia</i> , 2021 , 35, 1176-1187 | 10.7 | 1 |
| 280 | Novel invariant features of Good syndrome. <i>Leukemia</i> , 2021 , 35, 1792-1796 | 10.7 | 4 |

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| 279 | A Phase II Trial of Imatinib Mesylate as Maintenance Therapy for Patients With Newly Diagnosed C-kit-positive Acute Myeloid Leukemia. <i>Clinical Lymphoma, Myeloma and Leukemia</i> , 2021 , 21, 113-118 | 2 | 2 |
| 278 | Clonal trajectories and cellular dynamics of myeloid neoplasms with SF3B1 mutations. <i>Leukemia</i> , 2021 , 35, 3324-3328 | 10.7 | 0 |
| 277 | Clinical and basic implications of dynamic T cell receptor clonotyping in hematopoietic cell transplantation. <i>JCI Insight</i> , 2021 , 6, | 9.9 | 2 |
| 276 | Phase 2 study of danicopan in patients with paroxysmal nocturnal hemoglobinuria with an inadequate response to eculizumab. <i>Blood</i> , 2021 , 138, 1928-1938 | 2.2 | 7 |
| 275 | Influence of Killer Immunoglobulin-Like Receptors and Somatic Mutations on Transplant Outcomes in Acute Myeloid Leukemia. <i>Transplantation and Cellular Therapy</i> , 2021 , 27, 917.e1-917.e9 | | 0 |
| 274 | Personalized Prediction Model to Risk Stratify Patients With Myelodysplastic Syndromes. <i>Journal of Clinical Oncology</i> , 2021 , 39, 3737-3746 | 2.2 | 14 |
| 273 | Treatment outcomes for patients with myelodysplastic syndrome/myeloproliferative neoplasms with ring sideroblasts and thrombocytosis. <i>Leukemia and Lymphoma</i> , 2021 , 1-6 | 1.9 | 0 |
| 272 | Germline DDX41 mutations cause ineffective hematopoiesis and myelodysplasia. <i>Cell Stem Cell</i> , 2021 , 28, 1966-1981.e6 | 18 | 6 |
| 271 | T-cell large granular lymphocytic leukemia associated with inclusion body myositis. <i>International Journal of Laboratory Hematology</i> , 2021 , | 2.5 | 0 |
| 270 | A geno-clinical decision model for the diagnosis of myelodysplastic syndromes. <i>Blood Advances</i> , 2021 , 5, 4361-4369 | 7.8 | 2 |
| 269 | Large Granular Lymphocytic Leukemia: From Immunopathogenesis to Treatment of Refractory Disease. <i>Cancers</i> , 2021 , 13, | 6.6 | 2 |
| 268 | Monoclonal IgM gammopathy in adult acquired pure red cell aplasia: culprit or innocent bystander?. <i>Blood Cells, Molecules, and Diseases</i> , 2021 , 91, 102595 | 2.1 | 1 |
| 267 | TET-dioxygenase deficiency in oncogenesis and its targeting for tumor-selective therapeutics. <i>Seminars in Hematology</i> , 2021 , 58, 27-34 | 4 | 4 |
| 266 | Implication of PIGA genotype on erythrocytes phenotype in Paroxysmal Nocturnal Hemoglobinuria. <i>Leukemia</i> , 2021 , 35, 2431-2434 | 10.7 | 3 |
| 265 | Baseline clinical characteristics and disease burden in patients with paroxysmal nocturnal hemoglobinuria (PNH): updated analysis from the International PNH Registry. <i>Annals of Hematology</i> , 2020 , 99, 1505-1514 | 3 | 27 |
| 264 | Human erythroleukemia genetics and transcriptomes identify master transcription factors as functional disease drivers. <i>Blood</i> , 2020 , 136, 698-714 | 2.2 | 16 |
| 263 | SF3B1-mutant MDS as a distinct disease subtype: a proposal from the International Working Group for the Prognosis of MDS. <i>Blood</i> , 2020 , 136, 157-170 | 2.2 | 72 |
| 262 | 5-formylcytosine and 5-hydroxymethyluracil as surrogate markers of TET2 and SF3B1 mutations in myelodysplastic syndrome, respectively. <i>Haematologica</i> , 2020 , 105, e213-e215 | 6.6 | 1 |

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| 261 | Distinct mutational pattern of myelodysplastic syndromes with and without 5q- treated with lenalidomide. <i>British Journal of Haematology</i> , 2020 , 189, e133-e137 | 4.5 | 1 |
| 260 | Molecular landscape and clonal architecture of adult myelodysplastic/myeloproliferative neoplasms. <i>Blood</i> , 2020 , 136, 1851-1862 | 2.2 | 34 |
| 259 | From Bench to Bedside and Beyond: Therapeutic Scenario in Acute Myeloid Leukemia. <i>Cancers</i> , 2020 , 12, | 6.6 | 7 |
| 258 | Large granular lymphocytic leukaemia after solid organ and haematopoietic stem cell transplantation. <i>British Journal of Haematology</i> , 2020 , 189, 318-322 | 4.5 | 3 |
| 257 | Distinctive and common features of moderate aplastic anaemia. <i>British Journal of Haematology</i> , 2020 , 189, 967-975 | 4.5 | 4 |
| 256 | A Novel Therapeutic Strategy for Preferential Elimination of Multiple Myeloma Cells By Targeting Protein Disulfide Isomerase. <i>Blood</i> , 2020 , 136, 32-33 | 2.2 | |
| 255 | Leveraging Whole Genome Sequencing to Define the Mutational Landscape in Paroxysmal Nocturnal Hemoglobinuria. <i>Blood</i> , 2020 , 136, 8-8 | 2.2 | |
| 254 | Type of TP53 Mutations Affects Subclonal Configuration and Selection Pressure for Acquisition of Additional Hits in Contralateral Alleles. <i>Blood</i> , 2020 , 136, 25-25 | 2.2 | |
| 253 | Multicenter Validation of a Personalized Model to Predict Hypomethylating Agent Response in Myelodysplastic Syndromes (MDS). <i>Blood</i> , 2020 , 136, 54-55 | 2.2 | |
| 252 | Immunogenomics of Paroxysmal Nocturnal Hemoglobinuria: A Model of Immune Escape. <i>Blood</i> , 2020 , 136, 21-22 | 2.2 | |
| 251 | Impact of HLA Evolutionary Divergence on Clinical Features of Patients with Aplastic Anemia and Paroxysmal Nocturnal Hemoglobinuria. <i>Blood</i> , 2020 , 136, 2-3 | 2.2 | |
| 250 | Inhibition of Critical DNA Dioxygenase Activity in IDH1/2 Mutant Myeloid Neoplasms. <i>Blood</i> , 2020 , 136, 28-28 | 2.2 | |
| 249 | Molecular and Clinical Aspects of Acute Myeloid Leukemia with Inv(3)(q21q26)/t(3;3)(q21;q26) Carrying Spliceosomal Mutations. <i>Blood</i> , 2020 , 136, 7-8 | 2.2 | 0 |
| 248 | The Genomic Landscape of WilmsTumor 1 (WT1) Mutant Acute Myeloid Leukemia. <i>Blood</i> , 2020 , 136, 28-28 | 2.2 | |
| 247 | Venetoclax Inhibition of Pyrimidine Synthesis Guides Methods for Integration with Decitabine or 5-Azacytidine That Are Non-Myelosuppressive. <i>Blood</i> , 2020 , 136, 26-27 | 2.2 | |
| 246 | Molecular and Expression Characterization of Monosomy 7 and Del(7q). <i>Blood</i> , 2020 , 136, 33-33 | 2.2 | |
| 245 | Implication of Piga Genotype on Clinical Features of PNH. <i>Blood</i> , 2020 , 136, 34-35 | 2.2 | |
| 244 | Genotype-Phenotype Relationships and Therapeutic Targets in Acute Erythroid Leukemia. <i>Blood</i> , 2020 , 136, 17-18 | 2.2 | 1 |

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| 243 | Double Genetic Hits and Subclonal Mosaicism in the Ras Signaling Pathway in Myeloid Neoplasia. <i>Blood</i> , 2020 , 136, 34-35 | 2.2 | |
| 242 | Immunogenomics of Aplastic Anemia: The Role of HLA Somatic Mutations and the HLA Evolutionary Divergence. <i>Blood</i> , 2020 , 136, 20-21 | 2.2 | |
| 241 | Rare Germline Alterations of Myeloperoxidase Predispose to Myeloid Neoplasms and Are Associated with Increased Circulating Burden of Microbial DNA. <i>Blood</i> , 2020 , 136, 2-3 | 2.2 | |
| 240 | Characterization of the Blood and Bone Marrow Microbiome of MDS Patients and Associations with Clinical Features. <i>Blood</i> , 2020 , 136, 34-35 | 2.2 | 0 |
| 239 | A Phase I/II Trial of CPX-351 + Palbociclib in Patients with Acute Myeloid Leukemia. <i>Blood</i> , 2020 , 136, 13-14 | 2.2 | 1 |
| 238 | Leukemia Relapse after Allogeneic Hematopoietic Stem Cell Transplantation: From Recapitulation/Acquisition of Leukemogenic Hits to Immune Escape Due to Somatic Class I/ II HLA Mutations. <i>Blood</i> , 2020 , 136, 21-21 | 2.2 | |
| 237 | Role of Oligoadenylate Synthetases in Myeloid Neoplasia. <i>Blood</i> , 2020 , 136, 29-30 | 2.2 | |
| 236 | Aberrant Telomere Length and Composition Are Recurrent Features of Myeloid Disorders. <i>Blood</i> , 2020 , 136, 29-30 | 2.2 | 1 |
| 235 | TET2 Inhibitory Effects of Eltrombopag Contribute Its Hematopoietic Activity. <i>Blood</i> , 2020 , 136, 2-3 | 2.2 | |
| 234 | Genomic Landscape of Splicing Factor Mutant Acute Myeloid Leukemia. <i>Blood</i> , 2020 , 136, 36-36 | 2.2 | |
| 233 | A Personalized Clinical-Decision Tool to Improve the Diagnostic Accuracy of Myelodysplastic Syndromes. <i>Blood</i> , 2020 , 136, 33-35 | 2.2 | 2 |
| 232 | The Clonal Trajectories of SF3B1 Mutations in Myeloid Neoplasia. <i>Blood</i> , 2020 , 136, 8-8 | 2.2 | 1 |
| 231 | Genotype-Phenotype Correlations in Patients with Myeloid Malignancies Using Explainable Artificial Intelligence. <i>Blood</i> , 2020 , 136, 31-32 | 2.2 | 1 |
| 230 | The Genomic Landscape of Myeloid Neoplasms Evolved from AA/PNH. <i>Blood</i> , 2020 , 136, 2-2 | 2.2 | 1 |
| 229 | Clinical Impacts of Germline DDX41 Mutations on Myeloid Neoplasms. <i>Blood</i> , 2020 , 136, 38-40 | 2.2 | 2 |
| 228 | Targeted Sequencing of 7 Genes Can Help Reduce Pathologic Misclassification of MDS. <i>Blood</i> , 2020 , 136, 32-33 | 2.2 | 1 |
| 227 | Impact of Pathogenic Germ Line Variants in Adults with Acquired Bone Marrow Failure Syndromes Vs. Myeloid Neoplasia. <i>Blood</i> , 2020 , 136, 1-1 | 2.2 | 1 |
| 226 | Genomics of therapy-related myeloid neoplasms. <i>Haematologica</i> , 2020 , 105, e98-e101 | 6.6 | 10 |

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| 225 | Rare germline variant contributions to myeloid malignancy susceptibility. <i>Leukemia</i> , 2020 , 34, 1675-1678 | 10.7 | 3 |
| 224 | Clonal dynamics of aplastic anemia/paroxysmal nocturnal hemoglobinuria. <i>Leukemia and Lymphoma</i> , 2020 , 61, 1242-1245 | 1.9 | 1 |
| 223 | Extended experience with a non-cytotoxic DNMT1-targeting regimen of decitabine to treat myeloid malignancies. <i>British Journal of Haematology</i> , 2020 , 188, 924-929 | 4.5 | 8 |
| 222 | Machine learning demonstrates that somatic mutations imprint invariant morphologic features in myelodysplastic syndromes. <i>Blood</i> , 2020 , 136, 2249-2262 | 2.2 | 21 |
| 221 | Context dependent effects of ascorbic acid treatment in TET2 mutant myeloid neoplasia. <i>Communications Biology</i> , 2020 , 3, 493 | 6.7 | 13 |
| 220 | Large granular lymphocytic leukemia coexists with myeloid clones and myelodysplastic syndrome. <i>Leukemia</i> , 2020 , 34, 957-962 | 10.7 | 16 |
| 219 | Leukemia evolving from paroxysmal nocturnal hemoglobinuria. <i>Leukemia</i> , 2020 , 34, 327-330 | 10.7 | 1 |
| 218 | Targeting of CD38 by the Tumor Suppressor miR-26a Serves as a Novel Potential Therapeutic Agent in Multiple Myeloma. <i>Cancer Research</i> , 2020 , 80, 2031-2044 | 10.1 | 19 |
| 217 | Promoter Methylation Is Linked to Defective Homologous Recombination Repair and Elevated to Disrupt Myeloid Differentiation in Myeloid Malignancies. <i>Clinical Cancer Research</i> , 2019 , 25, 2513-2522 | 12.9 | 6 |
| 216 | The mutational burden of therapy-related myeloid neoplasms is similar to primary myelodysplastic syndrome but has a distinctive distribution. <i>Leukemia</i> , 2019 , 33, 2842-2853 | 10.7 | 19 |
| 215 | Impact of germline CTC1 alterations on telomere length in acquired bone marrow failure. <i>British Journal of Haematology</i> , 2019 , 185, 935-939 | 4.5 | 5 |
| 214 | Chronic myeloid leukemia: Two mysteries. <i>Leukemia Research</i> , 2019 , 79, 3-5 | 2.7 | 1 |
| 213 | The functional mechanisms of mutations in myelodysplastic syndrome. <i>Leukemia</i> , 2019 , 33, 2779-2794 | 10.7 | 11 |
| 212 | Effects of the Therapeutic Armamentarium on Survival and Time to Next Treatment in CMML Subtypes: An International Analysis of 950 Cases Coordinated By the AGMT Study Group. <i>Blood</i> , 2019 , 134, 844-844 | 2.2 | 3 |
| 211 | RORA Is a Potential Prognostic Biomarker and Therapeutic Target for Patients with Acute Myeloid Leukemia. <i>Blood</i> , 2019 , 134, 2696-2696 | 2.2 | 1 |
| 210 | Geno-Clinical Model for the Diagnosis of Bone Marrow Myeloid Neoplasms. <i>Blood</i> , 2019 , 134, 4238-4238 | 2.2 | 2 |
| 209 | Combined Treatment with Lenalidomide and Epoetin Alfa Leads to Durable Responses in Patients with Epo-Refractory, Lower Risk Non-Deletion 5q [Del(5q)] MDS: Final Results of the E2905 Intergroup Phase III Study - an ECOG-ACRIN Cancer Research Group Study, Grant CA180820, and the National Cancer Institute of the National Institutes of Health. <i>Blood</i> , 2019 , 134, 842-842 | 2.2 | 3 |
| 208 | A Personalized Prediction Model to Risk Stratify Patients with Acute Myeloid Leukemia (AML) Using Artificial Intelligence. <i>Blood</i> , 2019 , 134, 2091-2091 | 2.2 | 8 |

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| 207 | Novel Molecular Pathogenesis and Therapeutic Target in Acute Erythroid Leukemia. <i>Blood</i> , 2019 , 134, 914-914 | 2.2 | 1 |
| 206 | TET Dioxygenase Inhibition As a Therapeutic Strategy in TET2 Mutant Myeloid Neoplasia. <i>Blood</i> , 2019 , 134, 880-880 | 2.2 | 2 |
| 205 | Idiopathic aplastic anemia vs hypocellular myelodysplastic syndrome. <i>Hematology American Society of Hematology Education Program</i> , 2019 , 2019, 97-104 | 3.1 | 9 |
| 204 | Gene-centric functional dissection of human genetic variation uncovers regulators of hematopoiesis. <i>ELife</i> , 2019 , 8, | 8.9 | 7 |
| 203 | T-cell large granular lymphocytic leukemia evolution post-transplant: The Cleveland Clinic experience.. <i>Journal of Clinical Oncology</i> , 2019 , 37, e19072-e19072 | 2.2 | |
| 202 | CUL1: Novel Therapeutic Target in Myeloid Neoplasms Harboring -7/Del(7q). <i>Blood</i> , 2019 , 134, 1281-1281.2 | | |
| 201 | The Biological and Clinical Implications of the Alternative Splicing Landscape of 1,258 Myeloid Neoplasm Cases. <i>Blood</i> , 2019 , 134, 769-769 | 2.2 | |
| 200 | Extended Experience with a Very Low Dose, Metronomic, Subcutaneous Decitabine Regimen Intended to Deplete DNMT1 without Cytotoxicity. <i>Blood</i> , 2019 , 134, 1279-1279 | 2.2 | |
| 199 | Pharmacologic Normalization of Altered Transcriptome of SF3B1 Mutant Myeloid Neoplasia. <i>Blood</i> , 2019 , 134, 564-564 | 2.2 | |
| 198 | Molecular Characterization of EP300 Mutant Myeloid Neoplasia. <i>Blood</i> , 2019 , 134, 5043-5043 | 2.2 | |
| 197 | TET2 Loss Accelerates Leukemogenesis By Disrupting Mismatch Repair Proteins. <i>Blood</i> , 2019 , 134, 1200-1200 | | |
| 196 | Long-Term Experience with Large Granular Lymphocytic Leukemia Evolving after Solid Organ and Hematopoietic Stem Cell Transplantation. <i>Blood</i> , 2019 , 134, 1226-1226 | 2.2 | |
| 195 | MPO as a Novel Susceptibility Gene in Myeloid Malignancies. <i>Blood</i> , 2019 , 134, 5402-5402 | 2.2 | 0 |
| 194 | Angioimmunoblastic T-Cell Lymphoma: Molecular Characterization of Clonal T and B-Cells and a Patient Derived Xenograft Model of Coexisting T and B-Cell Proliferations. <i>Blood</i> , 2019 , 134, 1572-1572 | 2.2 | |
| 193 | Large Granular Lymphocytic Leukemia Coexists with Clonal Hematopoiesis of Indeterminate Potential. <i>Blood</i> , 2019 , 134, 3743-3743 | 2.2 | |
| 192 | Predicting Response to Hypomethylating Agents in Patients with Myelodysplastic Syndromes (MDS) Using Artificial Intelligence (AI). <i>Blood</i> , 2019 , 134, 2089-2089 | 2.2 | |
| 191 | A Single Arm, Phase II Study of Eltrombopag to Enhance Platelet Count Recovery in Older Patients with Acute Myeloid Leukemia (AML) Undergoing Remission Induction Therapy. <i>Blood</i> , 2019 , 134, 2595-2595 | 2.2 | |
| 190 | Genetics of Monosomy 7 and Del(7q) in MDS Informs Potential Therapeutic Targets. <i>Blood</i> , 2019 , 134, 1703-1703 | 2.2 | 1 |

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| 189 | Fatty Acid Binding Protein FABP5: A Novel Therapeutic Target in Acute Myeloid Leukemia. <i>Blood</i> , 2019 , 134, 2553-2553 | 2.2 | 1 |
| 188 | Clonal Cytopenias of Undetermined Significance Are Common in Cytopenic Adults Evaluated for MDS in the National MDS Study. <i>Blood</i> , 2019 , 134, 4271-4271 | 2.2 | |
| 187 | Invariant phenotype and molecular association of biallelic mutant myeloid neoplasia. <i>Blood Advances</i> , 2019 , 3, 339-349 | 7.8 | 18 |
| 186 | mutations define a specific subgroup of MDS and MDS/MPN patients with favorable outcomes with intensive chemotherapy. <i>Blood Advances</i> , 2019 , 3, 922-933 | 7.8 | 39 |
| 185 | Subclonal STAT3 mutations solidify clonal dominance. <i>Blood Advances</i> , 2019 , 3, 917-921 | 7.8 | 12 |
| 184 | Distinct clinical and biological implications of in myeloid neoplasms. <i>Blood Advances</i> , 2019 , 3, 2164-2178 | 7.8 | 12 |
| 183 | Therapy-related acute lymphoblastic leukemia is a distinct entity with adverse genetic features and clinical outcomes. <i>Blood Advances</i> , 2019 , 3, 4228-4237 | 7.8 | 16 |
| 182 | Invariant patterns of clonal succession determine specific clinical features of myelodysplastic syndromes. <i>Nature Communications</i> , 2019 , 10, 5386 | 17.4 | 29 |
| 181 | Molecular pathogenesis of disease progression in MLL-rearranged AML. <i>Leukemia</i> , 2019 , 33, 612-624 | 10.7 | 18 |
| 180 | Mutation clonal burden and allogeneic hematopoietic cell transplantation outcomes in acute myeloid leukemia and myelodysplastic syndromes. <i>Bone Marrow Transplantation</i> , 2019 , 54, 1281-1286 | 4.4 | 17 |
| 179 | New drugs for pharmacological extension of replicative life span in normal and progeroid cells. <i>Npj Aging and Mechanisms of Disease</i> , 2019 , 5, 2 | 5.5 | 7 |
| 178 | and mutations in myelodysplastic syndromes (MDS): clonal architecture and impact on outcomes. <i>Leukemia and Lymphoma</i> , 2019 , 60, 1587-1590 | 1.9 | 9 |
| 177 | Effectiveness of eculizumab in patients with paroxysmal nocturnal hemoglobinuria (PNH) with or without aplastic anemia in the International PNH Registry. <i>American Journal of Hematology</i> , 2019 , 94, E37-E41 | 7.1 | 12 |
| 176 | Mutations Sensitize Acute Myeloid Leukemia to PARP Inhibition and This Is Reversed by IDH1/2-Mutant Inhibitors. <i>Clinical Cancer Research</i> , 2018 , 24, 1705-1715 | 12.9 | 53 |
| 175 | Wild-type and mutated IDH1/2 enzymes and therapy responses. <i>Oncogene</i> , 2018 , 37, 1949-1960 | 9.2 | 127 |
| 174 | Mutations in DNMT3A, U2AF1, and EZH2 identify intermediate-risk acute myeloid leukemia patients with poor outcome after CR1. <i>Blood Cancer Journal</i> , 2018 , 8, 4 | 7 | 21 |
| 173 | Targeting the MALAT1/PARP1/LIG3 complex induces DNA damage and apoptosis in multiple myeloma. <i>Leukemia</i> , 2018 , 32, 2250-2262 | 10.7 | 70 |
| 172 | Clinical features and treatment outcomes in large granular lymphocytic leukemia (LGLL). <i>Leukemia and Lymphoma</i> , 2018 , 59, 416-422 | 1.9 | 42 |

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|-----|---|------|----|
| 171 | Therapeutic outcomes using subcutaneous low dose alemtuzumab for acquired bone marrow failure conditions. <i>British Journal of Haematology</i> , 2018 , 183, 133-136 | 4.5 | 6 |
| 170 | Clonal PIGA mosaicism and dynamics in paroxysmal nocturnal hemoglobinuria. <i>Leukemia</i> , 2018 , 32, 2507-2511 | 2.2 | 6 |
| 169 | Tet2 Regulates Osteoclast Differentiation by Interacting with Runx1 and Maintaining Genomic 5-Hydroxymethylcytosine (5hmC). <i>Genomics, Proteomics and Bioinformatics</i> , 2018 , 16, 172-186 | 6.5 | 18 |
| 168 | Leukemogenic nucleophosmin mutation disrupts the transcription factor hub that regulates granulomonocytic fates. <i>Journal of Clinical Investigation</i> , 2018 , 128, 4260-4279 | 15.9 | 61 |
| 167 | Fanconi Anemia germline variants as susceptibility factors in aplastic anemia, MDS and AML. <i>Oncotarget</i> , 2018 , 9, 2050-2057 | 3.3 | 12 |
| 166 | Heterozygous CTC1 Variants in Acquired Bone Marrow Failure. <i>Blood</i> , 2018 , 132, 3866-3866 | 2.2 | |
| 165 | BRCA1 & BRCA2 Germline Variants Are Enriched in MDS/AML and Portend Higher Average Mutational Burden. <i>Blood</i> , 2018 , 132, 4352-4352 | 2.2 | |
| 164 | Association of MHC Class I Chain-Related Gene a (MICA) Polymorphisms with Allogeneic Hematopoietic Cell Transplantation Outcomes in Acute Myeloid Leukemia. <i>Blood</i> , 2018 , 132, 2075-2075 | 2.2 | |
| 163 | Novel Small Molecule Stimulants of Hematopoietic Stem Cells and Their Mode of Action. <i>Blood</i> , 2018 , 132, 1302-1302 | 2.2 | |
| 162 | Analysis of Even a Limited Number of Genes Indicates a Strong Inherited Component in Otherwise Typical Sporadic MDS. <i>Blood</i> , 2018 , 132, 3074-3074 | 2.2 | |
| 161 | Differences in Genomic Patterns between African Americans and Whites with Acute Myeloid Leukemia. <i>Blood</i> , 2018 , 132, 1527-1527 | 2.2 | |
| 160 | Survival Outcomes of Patients with Therapy-Related Myelodysplastic Syndromes in the United States. <i>Blood</i> , 2018 , 132, 371-371 | 2.2 | |
| 159 | Rational management approach to pure red cell aplasia. <i>Haematologica</i> , 2018 , 103, 221-230 | 6.6 | 29 |
| 158 | The evolution of paroxysmal nocturnal haemoglobinuria depends on intensity of immunosuppressive therapy. <i>British Journal of Haematology</i> , 2018 , 182, 730-733 | 4.5 | 9 |
| 157 | Mutational landscape of myelodysplastic/myeloproliferative neoplasm-unclassifiable. <i>Blood</i> , 2018 , 132, 2100-2103 | 2.2 | 26 |
| 156 | Germline loss-of-function and alterations in adult myelodysplastic syndromes. <i>Blood</i> , 2018 , 132, 2309-2313 | 2.2 | 23 |
| 155 | Consequences of mutant TET2 on clonality and subclonal hierarchy. <i>Leukemia</i> , 2018 , 32, 1751-1761 | 10.7 | 30 |
| 154 | Transcriptomic rationale for synthetic lethality-targeting ERCC1 and CDKN1A in chronic myelomonocytic leukaemia. <i>British Journal of Haematology</i> , 2018 , 182, 373-383 | 4.5 | 4 |

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|-----|--|------|-----|
| 153 | Genetic abnormalities in myelodysplasia and secondary acute myeloid leukemia: impact on outcome of stem cell transplantation. <i>Blood</i> , 2017 , 129, 2347-2358 | 2.2 | 184 |
| 152 | Molecular features of early onset adult myelodysplastic syndrome. <i>Haematologica</i> , 2017 , 102, 1028-1034 | 4.6 | 13 |
| 151 | DDX41-related myeloid neoplasia. <i>Seminars in Hematology</i> , 2017 , 54, 94-97 | 4 | 28 |
| 150 | Tet2 loss leads to hypermutagenicity in haematopoietic stem/progenitor cells. <i>Nature Communications</i> , 2017 , 8, 15102 | 17.4 | 61 |
| 149 | Novel therapeutic strategies to target leukemic cells that hijack compartmentalized continuous hematopoietic stem cell niches. <i>Biochimica Et Biophysica Acta: Reviews on Cancer</i> , 2017 , 1868, 183-198 | 11.2 | 25 |
| 148 | Ubiquitination of hnRNPA1 by TRAF6 links chronic innate immune signaling with myelodysplasia. <i>Nature Immunology</i> , 2017 , 18, 236-245 | 19.1 | 60 |
| 147 | Dynamics of clonal evolution in myelodysplastic syndromes. <i>Nature Genetics</i> , 2017 , 49, 204-212 | 36.3 | 228 |
| 146 | Origins of myelodysplastic syndromes after aplastic anemia. <i>Blood</i> , 2017 , 130, 1953-1957 | 2.2 | 32 |
| 145 | Clinical implications of somatic mutations in aplastic anemia and myelodysplastic syndrome in genomic age. <i>Hematology American Society of Hematology Education Program</i> , 2017 , 2017, 66-72 | 3.1 | 10 |
| 144 | GATA4 loss of function in liver cancer impedes precursor to hepatocyte transition. <i>Journal of Clinical Investigation</i> , 2017 , 127, 3527-3542 | 15.9 | 26 |
| 143 | Recurrent genetic defects on chromosome 5q in myeloid neoplasms. <i>Oncotarget</i> , 2017 , 8, 6483-6495 | 3.3 | 24 |
| 142 | Extent and Clinical Implications of Subclonal Diversity in Paroxysmal Nocturnal Hemoglobinuria. <i>Blood</i> , 2017 , 130, 779-779 | 2.2 | 2 |
| 141 | The Mechanisms By Which Mutant-NPM1 Uncouples Differentiation from Proliferation Are Reversed By Several Drugs, Enabling Rational Multi-Component Non-Cytotoxic Differentiation Therapy. <i>Blood</i> , 2017 , 130, 878-878 | 2.2 | |
| 140 | The efficacy of current prognostic models in predicting outcome of patients with myelodysplastic syndromes at the time of hypomethylating agent failure. <i>Haematologica</i> , 2016 , 101, e224-7 | 6.6 | 30 |
| 139 | Subcutaneous Low Dose Alemtuzumab: Role As a Salvage Therapy in Immune -Mediated Marrow Failure Conditions. <i>Blood</i> , 2016 , 128, 1505-1505 | 2.2 | 0 |
| 138 | Molecular and Immunophenotypic Characteristics of Adult Acute Leukemias of Ambiguous Lineage. <i>Blood</i> , 2016 , 128, 1659-1659 | 2.2 | 1 |
| 137 | Genetic and Epigenetic Defects in the Autophagy Machinery in Myelodysplastic Syndromes. <i>Blood</i> , 2016 , 128, 4301-4301 | 2.2 | 2 |
| 136 | TP53 Mutations and Outcome in Patients with Myelodysplastic Syndromes (MDS). <i>Blood</i> , 2016 , 128, 4336-4336 | 2.2 | 2 |

| | | | |
|-----|---|------|-----|
| 135 | The Mechanism By Which Mutant Nucleophosmin (NPM1) Creates Leukemic Self-Renewal Is Readily Reversed. <i>Blood</i> , 2016 , 128, 444-444 | 2.2 | 4 |
| 134 | The Role of LUC7L2 in Splicing and MDS. <i>Blood</i> , 2016 , 128, 5504-5504 | 2.2 | 3 |
| 133 | Myb expression is critical for myeloid leukemia development induced by Setbp1 activation. <i>Oncotarget</i> , 2016 , 7, 86300-86312 | 3.3 | 16 |
| 132 | BCOR and BCORL1 mutations in Myelodysplastic Syndromes (MDS): Clonal Architecture and Impact on Outcomes. <i>Blood</i> , 2016 , 128, 4293-4293 | 2.2 | |
| 131 | Impact of SNP array karyotyping on the diagnosis and the outcome of chronic myelomonocytic leukemia with low risk cytogenetic features or no metaphases. <i>American Journal of Hematology</i> , 2016 , 91, 185-92 | 7.1 | 15 |
| 130 | Impact of allogeneic hematopoietic cell transplant in patients with myeloid neoplasms carrying spliceosomal mutations. <i>American Journal of Hematology</i> , 2016 , 91, 406-9 | 7.1 | 11 |
| 129 | High incidence of activating STAT5B mutations in CD4-positive T-cell large granular lymphocyte leukemia. <i>Blood</i> , 2016 , 128, 2465-2468 | 2.2 | 51 |
| 128 | Connect MDS/AML: design of the myelodysplastic syndromes and acute myeloid leukemia disease registry, a prospective observational cohort study. <i>BMC Cancer</i> , 2016 , 16, 652 | 4.8 | 10 |
| 127 | Somatic Mutations and Clonal Hematopoiesis in Aplastic Anemia. <i>New England Journal of Medicine</i> , 2015 , 373, 35-47 | 59.2 | 361 |
| 126 | Inherited and Somatic Defects in DDX41 in Myeloid Neoplasms. <i>Cancer Cell</i> , 2015 , 27, 658-70 | 24.3 | 228 |
| 125 | The Revised International Prognostic Scoring System (IPSS-R) is not predictive of survival in patients with secondary myelodysplastic syndromes. <i>Leukemia and Lymphoma</i> , 2015 , 56, 3437-9 | 1.9 | 19 |
| 124 | Loss of Tifab, a del(5q) MDS gene, alters hematopoiesis through derepression of Toll-like receptor-TRAF6 signaling. <i>Journal of Experimental Medicine</i> , 2015 , 212, 1967-85 | 16.6 | 65 |
| 123 | Dose-dependent role of the cohesin complex in normal and malignant hematopoiesis. <i>Journal of Experimental Medicine</i> , 2015 , 212, 1819-32 | 16.6 | 106 |
| 122 | Whole-exome sequencing enhances prognostic classification of myeloid malignancies. <i>Journal of Biomedical Informatics</i> , 2015 , 58, 104-113 | 10.2 | 9 |
| 121 | Radioprotection of IDH1-Mutated Cancer Cells by the IDH1-Mutant Inhibitor AGI-5198. <i>Cancer Research</i> , 2015 , 75, 4790-802 | 10.1 | 108 |
| 120 | The analysis of clonal diversity and therapy responses using STAT3 mutations as a molecular marker in large granular lymphocytic leukemia. <i>Haematologica</i> , 2015 , 100, 91-9 | 6.6 | 61 |
| 119 | Mutations in G protein β subunits promote transformation and kinase inhibitor resistance. <i>Nature Medicine</i> , 2015 , 21, 71-5 | 50.5 | 60 |
| 118 | Genomic patterns associated with hypoplastic compared to hyperplastic myelodysplastic syndromes. <i>Haematologica</i> , 2015 , 100, e434-7 | 6.6 | 18 |

| | | | |
|-----|--|------|-----|
| 117 | TET 2 Alterations in Myeloid Malignancies, Impact on Clinical Characteristics, Outcome, and Disease Predisposition. <i>Blood</i> , 2015 , 126, 1645-1645 | 2.2 | 1 |
| 116 | Clonal Mutational Landscape of Childhood Myelodysplastic Syndromes. <i>Blood</i> , 2015 , 126, 1662-1662 | 2.2 | 5 |
| 115 | Survival Outcomes of Leukemias and Myelodysplastic Syndromes Occurring As Second Cancers in the United States: A SEER Registry-Based Population Analysis. <i>Blood</i> , 2015 , 126, 2507-2507 | 2.2 | 2 |
| 114 | Somatic Mutations in MDS Patients Are Associated with Clinical Features and Predict Prognosis Independent of the IPSS-R: Analysis of Combined Datasets from the International Working Group for Prognosis in MDS-Molecular Committee. <i>Blood</i> , 2015 , 126, 907-907 | 2.2 | 73 |
| 113 | Aberrant DNA Methylation Is Associated with a Poor Outcome in Juvenile Myelomonocytic Leukemia. <i>PLoS ONE</i> , 2015 , 10, e0145394 | 3.7 | 18 |
| 112 | APC mutations in myeloid malignancies: Incidence and impact on leukemogenesis.. <i>Journal of Clinical Oncology</i> , 2015 , 33, 11047-11047 | 2.2 | 1 |
| 111 | Activation of the Unfolded Protein Response with the First-in-Class P97 Inhibitor CB-5083 Induces Stable Disease Regression and Overcomes Ara-C Resistance in AML. <i>Blood</i> , 2015 , 126, 1350-1350 | 2.2 | 0 |
| 110 | Impact of STAT3 Mutations on Clinical Features and Treatment Outcomes in Large Granular Lymphocyte Leukemia. <i>Blood</i> , 2015 , 126, 2216-2216 | 2.2 | |
| 109 | Dose-Dependent Role of the Cohesin Complex in Normal and Malignant Hematopoiesis. <i>Blood</i> , 2015 , 126, 435-435 | 2.2 | |
| 108 | Determinants of Phenotypic Commitment and Clonal Progression--Conclusions from the Study of Clonal Architecture in CMML. <i>Blood</i> , 2015 , 126, 2848-2848 | 2.2 | |
| 107 | Radioactive Iodine Treatment of Thyroid Cancer and Risk of Myelodysplastic Syndromes. <i>Blood</i> , 2015 , 126, 612-612 | 2.2 | |
| 106 | Real World Outcomes of Less Well-Characterized Acute Leukemias: A Population-Based Survival Analysis Using SEER Registry (1973-2012). <i>Blood</i> , 2015 , 126, 4491-4491 | 2.2 | |
| 105 | Network-Based Analysis of Exome Sequencing Mutations Identifies Molecular Subtypes of Myelodysplastic Syndromes. <i>Blood</i> , 2015 , 126, 611-611 | 2.2 | |
| 104 | Synergistic effect of major histocompatibility complex class I-related chain a and human leukocyte antigen-DPB1 mismatches in association with acute graft-versus-host disease after unrelated donor hematopoietic stem cell transplantation. <i>Biology of Blood and Marrow Transplantation</i> , 2014 , 20, 1835-40 | 4.7 | 14 |
| 103 | The driver and passenger effects of isocitrate dehydrogenase 1 and 2 mutations in oncogenesis and survival prolongation. <i>Biochimica Et Biophysica Acta: Reviews on Cancer</i> , 2014 , 1846, 326-41 | 11.2 | 93 |
| 102 | ETV6 and signaling gene mutations are associated with secondary transformation of myelodysplastic syndromes to chronic myelomonocytic leukemia. <i>Blood</i> , 2014 , 123, 3675-7 | 2.2 | 16 |
| 101 | Genetic alterations of the cohesin complex genes in myeloid malignancies. <i>Blood</i> , 2014 , 124, 1790-8 | 2.2 | 151 |
| 100 | Myeloid malignancies with chromosome 5q deletions acquire a dependency on an intrachromosomal NF- κ B gene network. <i>Cell Reports</i> , 2014 , 8, 1328-38 | 10.6 | 46 |

| | | | |
|----|---|------|------|
| 99 | Deep sequencing reveals stepwise mutation acquisition in paroxysmal nocturnal hemoglobinuria. <i>Journal of Clinical Investigation</i> , 2014 , 124, 4529-38 | 15.9 | 87 |
| 98 | Molecular pathogenesis of myelodysplastic syndromes. <i>Translational Medicine @ UniSa</i> , 2014 , 8, 19-30 | 0.5 | 10 |
| 97 | Impact and Function of Somatic PHF6 Mutations in Myeloid Neoplasms. <i>Blood</i> , 2014 , 124, 3581-3581 | 2.2 | |
| 96 | Haploinsufficiency and Deletions of G3BP1 on Chromosome 5q Result in Induction of TP53. <i>Blood</i> , 2014 , 124, 784-784 | 2.2 | |
| 95 | Deep sequencing of the T-cell receptor repertoire in CD8+ T-large granular lymphocyte leukemia identifies signature landscapes. <i>Blood</i> , 2013 , 122, 4077-85 | 2.2 | 50 |
| 94 | STAT3 mutations indicate the presence of subclinical T-cell clones in a subset of aplastic anemia and myelodysplastic syndrome patients. <i>Blood</i> , 2013 , 122, 2453-9 | 2.2 | 87 |
| 93 | A case of mistaken identity: When lupus masquerades as primary myelofibrosis. <i>SAGE Open Medical Case Reports</i> , 2013 , 1, 2050313X13498709 | 0.7 | 2 |
| 92 | Whole Exome Sequencing Detecting Kinesin Family Gene Defects In Myeloid Neoplasm. <i>Blood</i> , 2013 , 122, 2762-2762 | 2.2 | |
| 91 | Molecular Predictors Of Response To Lenalidomide In Myeloid Malignancies. <i>Blood</i> , 2013 , 122, 2807-2807 | 2.2 | |
| 90 | Introduction: molecular pathogenesis of hematologic malignancies. <i>Seminars in Oncology</i> , 2012 , 39, 9-12 | 5.5 | 2 |
| 89 | STAT3 mutations unify the pathogenesis of chronic lymphoproliferative disorders of NK cells and T-cell large granular lymphocyte leukemia. <i>Blood</i> , 2012 , 120, 3048-57 | 2.2 | 285 |
| 88 | Revised international prognostic scoring system for myelodysplastic syndromes. <i>Blood</i> , 2012 , 120, 2454-65 | 6.5 | 1799 |
| 87 | Somatic STAT3 mutations in large granular lymphocytic leukemia. <i>New England Journal of Medicine</i> , 2012 , 366, 1905-13 | 59.2 | 535 |
| 86 | Long-Term Safety of Sustained Eculizumab Treatment in Patients with Paroxysmal Nocturnal Hemoglobinuria. <i>Blood</i> , 2012 , 120, 1260-1260 | 2.2 | 1 |
| 85 | Non-Cytotoxic Differentiation Therapy Based On Mechanism of Disease Produces Complete Remission in Myelodysplastic Syndromes (MDS) with High Risk Cytogenetics. <i>Blood</i> , 2012 , 120, 1696-1696 | 2.2 | 3 |
| 84 | Impact of Mutations in the Spliceosome Machinery and Ring Sideroblasts in Patients with Myeloid Malignancies Who Received Conventional Chemotherapy or Allogeneic Hematopoietic Cell Transplantation. <i>Blood</i> , 2012 , 120, 1973-1973 | 2.2 | 1 |
| 83 | Whole Exome Analysis Reveals Spectrum of Gene Mutations in Juvenile Myelomonocytic Leukemia. <i>Blood</i> , 2012 , 120, 170-170 | 2.2 | |
| 82 | Haploinsufficiency of Mir-146a in High-Risk Del(5q) MDS/AML Requires an Intrachromosomal Gene Network Involving p62/TRAF6/NF- κ B. <i>Blood</i> , 2012 , 120, 557-557 | 2.2 | |

| | | | |
|----|---|-----|----|
| 81 | SNP array-based karyotyping: differences and similarities between aplastic anemia and hypocellular myelodysplastic syndromes. <i>Blood</i> , 2011 , 117, 6876-84 | 2.2 | 88 |
| 80 | Clonal drift demonstrates unexpected dynamics of the T-cell repertoire in T-large granular lymphocyte leukemia. <i>Blood</i> , 2011 , 118, 4384-93 | 2.2 | 50 |
| 79 | Genetic and Epigenetic Defects in DNA Repair Lead to Synthetic Lethality of Poly (ADP-Ribose) Polymerase (PARP) Inhibitors in Aggressive Myeloproliferative Disorders. <i>Blood</i> , 2011 , 118, 400-400 | 2.2 | 4 |
| 78 | Association of SF3B1 with Ring Sideroblasts in patients, In Vivo, and In Vitro models of Spliceosomal Dysfunction. <i>Blood</i> , 2011 , 118, 457-457 | 2.2 | 2 |
| 77 | Clinical and Genomic Characterization of Chromosome 7 Lesions in Myeloid Malignancies,. <i>Blood</i> , 2011 , 118, 3549-3549 | 2.2 | |
| 76 | Defining the Topography of Deletion 5q Using SNP-A Identifies Patients with More Aggressive Disease and Correlates with Additional Lesions. <i>Blood</i> , 2011 , 118, 2795-2795 | 2.2 | |
| 75 | A Novel View of Paroxysmal Nocturnal Hemoglobinuria (PNH) Pathogenesis: Do Pathologic PNH Hematopoietic Stem/Progenitor Cells (HSPCs) Displace Normal HSPCs From Their Niches in Bone Marrow Because They Are More Motile Due to Defective Adhesion and Enhanced Migratory Properties?. <i>Blood</i> , 2011 , 118, 732-732 | 2.2 | |
| 74 | Prognostic Factors of Response and Survival in CMML Patients Treated with Azacitidine (AZA). <i>Blood</i> , 2011 , 118, 1726-1726 | 2.2 | |
| 73 | Increased Group B Killer Cell Immunoglobulin-Like Receptor (KIR) Haplotypes with Mismatched MHC Class I and Altered NK Repertoire Distribution in Bone Marrow Failure Syndromes. <i>Blood</i> , 2011 , 118, 2412-2412 | 2.2 | |
| 72 | Human Telomerase Reverse Transcriptase (hTERT) Deficiency in Myelodysplastic Syndrome (MDS) Demonstrates Mechanistic Linkage to Aplastic Anemia Pathophysiology. <i>Blood</i> , 2011 , 118, 791-791 | 2.2 | |
| 71 | Acquired Molecular Defects in Spliceosome Machinery: Novel Pathogenetic Pathways in Myeloid Leukemogenesis. <i>Blood</i> , 2011 , 118, 271-271 | 2.2 | |
| 70 | Prognostic Factors for Post-Transplant Outcomes in Patients with Myelodysplastic Syndromes (MDS). <i>Blood</i> , 2011 , 118, 2015-2015 | 2.2 | |
| 69 | A Proof of Principle Clinical Trial in Myelodysplastic Syndromes of Non-Cytotoxic Differentiation Therapy with Decitabine,. <i>Blood</i> , 2011 , 118, 3830-3830 | 2.2 | |
| 68 | Distinction of Early and Late Molecular Events In Patients with Myelodysplastic Syndromes (MDS) Who Progressed to Acute Myeloid Leukemia (AML),. <i>Blood</i> , 2011 , 118, 3566-3566 | 2.2 | |
| 67 | Pathogenesis of MONOSOMY 7 In BONE MARROW FAILURE SYNDROMES. <i>Blood</i> , 2011 , 118, 2411-2411 | 2.2 | |
| 66 | The Impact of Molecular Lesions in Post-Transplant Acute Myeloid Leukemia (AML) in Correlation with Cytogenetic Abnormalities,. <i>Blood</i> , 2011 , 118, 4137-4137 | 2.2 | |
| 65 | Cytogenetic Predictors of Response to Lenalidomide In Myeloid Malignancies without Del(5q). <i>Blood</i> , 2010 , 116, 4016-4016 | 2.2 | 1 |
| 64 | New TET2, ASXL1 and CBL Mutations Have Poor Prognostic Impact In Systemic Mastocytosis and Related Disorders. <i>Blood</i> , 2010 , 116, 3076-3076 | 2.2 | |

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|----|--|-----|-----|
| 63 | SNP-A Karyotyping Provides Clinically Relevant Results In Myeloid Hematologic Disorders with Unsuccessful Routine Cytogenetic Testing.. <i>Blood</i> , 2010 , 116, 3374-3374 | 2.2 | |
| 62 | HTLV-1 Epitope (BA21) Reactivity in Rare Bone Marrow Failure Diseases. <i>Blood</i> , 2010 , 116, 1724-1724 | 2.2 | |
| 61 | Single Nucleotide Polymorphism Array (SNP-A) Genomic Profiling of Mantle Cell Lymphoma (MCL) Against a Large Control Database Reveals Recurring Copy Number Alterations (CNAs) and Copy Neutral Loss of Heterozygosity (CN-LOH). <i>Blood</i> , 2010 , 116, 2001-2001 | 2.2 | |
| 60 | CBL, CBLB, TET2, ASXL1, and IDH1/2 Mutations as Well as Additional Chromosomal Aberrations Constitute Molecular Events Contributing to Malignant Progression In Advanced Philadelphia Chromosome-Positive Disorders.. <i>Blood</i> , 2010 , 116, 3396-3396 | 2.2 | |
| 59 | A High Resolution Analysis of Chromosome 21 Amplification In Myeloid Malignancies Reveals An Association with a Specific Cytogenetic Subgroup and Enhanced ERG Gene Expression.. <i>Blood</i> , 2010 , 116, 1687-1687 | 2.2 | |
| 58 | Clonotype Switching Indicates Propensity for Clonal Outgrowth From Diverse Components of the T Cell Repertoire In T Cell Large Granular Lymphocyte Leukemia.. <i>Blood</i> , 2010 , 116, 1171-1171 | 2.2 | |
| 57 | Next Generation Exome Sequencing for Identification of the Gene Mutations Associated with Loss of Heterozygosity on Chromosome 7 In Myeloid Malignancies. <i>Blood</i> , 2010 , 116, 297-297 | 2.2 | |
| 56 | Gene Alterations In Acute Megakaryoblastic Leukemia (AMKL): a Comparison of AMKL with and without Down Syndrome. <i>Blood</i> , 2010 , 116, 875-875 | 2.2 | |
| 55 | Therapeutic implications of variable expression of CD52 on clonal cytotoxic T cells in CD8+ large granular lymphocyte leukemia. <i>Haematologica</i> , 2009 , 94, 1407-14 | 6.6 | 32 |
| 54 | Application of array-based whole genome scanning technologies as a cytogenetic tool in haematological malignancies. <i>British Journal of Haematology</i> , 2009 , 146, 479-88 | 4.5 | 118 |
| 53 | A Study Comparing Dosing Regimens and Efficacy of Subcutaneous to Intravenous Azacitidine (AZA) for the Treatment of Myelodysplastic Syndromes (MDS).. <i>Blood</i> , 2009 , 114, 3797-3797 | 2.2 | 5 |
| 52 | C-Cbl but Not TET2 Mutations Are Present in Patients with Juvenile Myelomonocytic Leukemia.. <i>Blood</i> , 2009 , 114, 420-420 | 2.2 | 1 |
| 51 | Identification of Genetic Polymorphisms Contributing to Risk in MDS Using Innovative GWAS Approaches.. <i>Blood</i> , 2009 , 114, 734-734 | 2.2 | |
| 50 | Reconciling Phenotype with Genotype in the MPN: Impact of SNP Array-Based Chromosomal Analysis.. <i>Blood</i> , 2009 , 114, 1893-1893 | 2.2 | |
| 49 | Whole genome scanning as a cytogenetic tool in hematologic malignancies. <i>Blood</i> , 2008 , 112, 965-74 | 2.2 | 116 |
| 48 | Phenotypic differences between healthy effector CTL and leukemic LGL cells support the notion of antigen-triggered clonal transformation in T-LGL leukemia. <i>Journal of Leukocyte Biology</i> , 2008 , 83, 589-601 | 6.5 | 25 |
| 47 | Circulating Cytokine Profiles of Patients with Acquired Aplastic Anemia and Myelodysplastic Syndrome. <i>Blood</i> , 2008 , 112, 1038-1038 | 2.2 | 1 |
| 46 | SNP Array-Based Analysis of Chromosome 17 Reveals Biallelic TP53 Mutations Due to Uniparental Disomy 17p in Advanced MDS and AML with Cooperating Deletions of Chromosomes 5 and 7. <i>Blood</i> , 2008 , 112, 2521-2521 | 2.2 | 2 |

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|----|--|-----|---|
| 45 | Differences Between Normal and Leukemic Stem Cell-Specific Methylome Indicates Aberrantly Silenced Genes Involved in the Pathogenesis of Malignant Evolution. <i>Blood</i> , 2008 , 112, 599-599 | 2.2 | 1 |
| 44 | FISH and SNP-Array Karyotyping Improve the Detection of Recurrent Chromosomal Defects Including Del(5q), Monosomy 7, Del(7q), Trisomy 8, and Del(20q) in Myelodysplastic Syndromes.. <i>Blood</i> , 2008 , 112, 1483-1483 | 2.2 | |
| 43 | Progressive Chromatin Repression and Promoter Methylation of a-catenin Correlates with AML Transformation in Patients with and without 5q Deletions.. <i>Blood</i> , 2008 , 112, 3369-3369 | 2.2 | |
| 42 | Investigations of Genetic Risk Factors in MDS and AML Using High- Density 6.0 Affymetrix Arrays. <i>Blood</i> , 2008 , 112, 638-638 | 2.2 | 1 |
| 41 | BACH2 Directly Regulates Expression of Foxp3 in UCB CD4+ T-Cells. <i>Blood</i> , 2008 , 112, 4760-4760 | 2.2 | |
| 40 | Expression of MICA by Granulocytes in Neutropenia Due to Large Granular Lymphocyte Leukemia Points towards Cytotoxicity Exerted Via NKG2D on Clonal Cytotoxic T Cells.. <i>Blood</i> , 2008 , 112, 1262-1262 | 2.2 | 2 |
| 39 | Permissive Conditions for Evolution of PNH Clones Are Characterized by Overproduction of IFN- γ by Clonal CD4 and CD8 T Cells, Fas-L by CTLs, and Promoted by Immunogenetic Background. <i>Blood</i> , 2008 , 112, 4116-4116 | 2.2 | |
| 38 | SNP-Array Based Karyotyping Complements Routine Cytogenetics in Diagnosis and Risk Stratification Schemes of MDS. <i>Blood</i> , 2008 , 112, 639-639 | 2.2 | |
| 37 | Alemtuzumab Shows Significant Efficacy in T-LGL Leukemia and Refractory Cases Are Due to GPI-Deficiency of LGL Clones.. <i>Blood</i> , 2008 , 112, 2038-2038 | 2.2 | |
| 36 | SNP-A Based Karyotyping Facilitates Improved Mapping of Deletions and Uniparental Disomy within the Long Arm of Chromosome 5 in Myeloid Disorders.. <i>Blood</i> , 2007 , 110, 2435-2435 | 2.2 | 1 |
| 35 | Phospho-IkappaB Is Abnormally Expressed in Bone Marrow of CMML Patients.. <i>Blood</i> , 2007 , 110, 2450-2450 | 2.2 | 1 |
| 34 | SNP-Array Karyotyping Reveals the Presence of Previously Cryptic Clonal Chromosomal Aberrations Including Segmental UPD in Patients with Fanconi Anemia.. <i>Blood</i> , 2007 , 110, 1678-1678 | 2.2 | |
| 33 | SNP-A Karyotyping Demonstrates a High Incidence of Segmental Uniparental Disomy in Patients with CMML and Shows Impact of Newly Identified Chromosomal Aberrations on Clinical Course.. <i>Blood</i> , 2007 , 110, 2428-2428 | 2.2 | |
| 32 | SNP-Array-Based Karyotyping Has Impact on Cytogenetic Diagnosis and Prognosis of Non-Core Binding Factor Primary and Secondary AML.. <i>Blood</i> , 2007 , 110, 597-597 | 2.2 | |
| 31 | Can Genomic Copy Number Variants Be a Part of Complex Genetic Traits Predisposing to Marrow Failure?.. <i>Blood</i> , 2007 , 110, 106-106 | 2.2 | |
| 30 | SNP Arrays Facilitate Genotyping of Non-Synonymous SNP in MDS To Identify Disease Susceptibility Loci.. <i>Blood</i> , 2007 , 110, 2421-2421 | 2.2 | |
| 29 | Decreased Expression of Membrane-Bound Proteinase 3 by a GPI-Deficient Granulocytes May Contribute to Thrombophilic Propensity in PNH.. <i>Blood</i> , 2007 , 110, 3673-3673 | 2.2 | |
| 28 | SNP Array Karyotyping Improves Detection Rate of Clonal Chromosomal Abnormalities in Refractory Anemia with Ringed Sideroblasts.. <i>Blood</i> , 2007 , 110, 4132-4132 | 2.2 | |

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|----|---|----------------|-----|
| 27 | Immune-mediated bone marrow failure syndromes of progenitor and stem cells: molecular analysis of cytotoxic T cell clones. <i>Folia Histochemica Et Cytobiologica</i> , 2007 , 45, 5-14 | 1.4 | 34 |
| 26 | High Density SNP Arrays Reveal That Distinct Clonal Lesions Including Uniparental Disomy Can Be Detected in a Proportion of Patients with Aplastic Anemia with Normal Metaphase Cytogenetics.. <i>Blood</i> , 2006 , 108, 125-125 | 2.2 | 3 |
| 25 | Identification of Chromosomal Abnormalities in Healthy Bone Marrow Using 250K SNP Arrays.. <i>Blood</i> , 2006 , 108, 2076-2076 | 2.2 | 1 |
| 24 | Differential Expression of SLAM Family Receptor Markers in Normal Human Hematopoietic Stem Cells and Their Malignant Counterpart in MDS and AML.. <i>Blood</i> , 2006 , 108, 1897-1897 | 2.2 | |
| 23 | Influence of Killer Immunoglobulin-Like Receptor (KIR) Matching on Achieving T Cell (CD3+) Complete Donor Chimerism (CDC) in Related Donor Nonmyeloablative Allogeneic Hematopoietic Stem Cell Transplantation (NMHSCT).. <i>Blood</i> , 2006 , 108, 3012-3012 | 2.2 | 1 |
| 22 | A Decision Analysis To Determine the Appropriate Treatment for Low-Risk Myelodysplastic Syndromes.. <i>Blood</i> , 2005 , 106, 2533-2533 | 2.2 | 2 |
| 21 | Phosphatidylinositol-3-Phosphate Kinase Pathway Activation Protects Leukemic Large Granular Lymphocytes from Undergoing Homeostatic Apoptosis.. <i>Blood</i> , 2005 , 106, 739-739 | 2.2 | 1 |
| 20 | Presence of JAK2 Mutations in MDS/MPD-u WHO Classified Patients and Not Other Forms of MDS Suggests Their Derivation from Classical Myeloproliferative Syndrome.. <i>Blood</i> , 2005 , 106, 369-369 | 2.2 | 1 |
| 19 | Molecular Identification of Alloreactive CTL Precursors in Hematopoietic Stem Cell Transplantation.. <i>Blood</i> , 2005 , 106, 597-597 | 2.2 | |
| 18 | High-Resolution Genomic Arrays Facilitate Detection of Novel Cryptic Chromosomal Lesions in MDS.. <i>Blood</i> , 2005 , 106, 370-370 | 2.2 | |
| 17 | Immunogenetic Factors Determining Evolution of T-Cell Large Granular Lymphocyte Leukemia and Associated Cytopenias.. <i>Blood</i> , 2005 , 106, 2211-2211 | 2.2 | |
| 16 | Evolution of clonal cytogenetic abnormalities in aplastic anemia. <i>Leukemia and Lymphoma</i> , 2004 , 45, 433-40 | 1.9 | 63 |
| 15 | In-vivo dominant immune responses in aplastic anaemia: molecular tracking of putatively pathogenetic T-cell clones by TCR beta-CDR3 sequencing. <i>Lancet, The</i> , 2004 , 364, 355-64 | 4 ⁰ | 192 |
| 14 | Transfer of glycosylphosphatidylinositol-anchored proteins to deficient cells after erythrocyte transfusion in paroxysmal nocturnal hemoglobinuria. <i>Blood</i> , 2004 , 104, 3782-8 | 2.2 | 34 |
| 13 | Prospective Molecular Identification of Alloreactive CTL Clones in Allogeneic Hematopoietic Stem Cell Transplantation.. <i>Blood</i> , 2004 , 104, 4972-4972 | 2.2 | |
| 12 | Differential Comparative Genomic Hybridization Analysis of Normal and Glycosyl Phosphatidyl Inositol Deficient Clones in Paroxysmal Nocturnal Hemoglobinuria.. <i>Blood</i> , 2004 , 104, 2831-2831 | 2.2 | |
| 11 | Pathologic Clonal CTL Responses - Non Random Nature of the TCR Restriction in LGL Leukemia.. <i>Blood</i> , 2004 , 104, 3241-3241 | 2.2 | |
| 10 | A Pilot Application of SELDI Serum Proteomics in Bone Marrow Failure Syndromes.. <i>Blood</i> , 2004 , 104, 2822-2822 | 2.2 | |

| | | | |
|---|--|------|-----|
| 9 | KIR Gene Distribution in Hematologic Disorders.. <i>Blood</i> , 2004 , 104, 1624-1624 | 2.2 | |
| 8 | Polarized CTL Responses Detected in Patients with Autoimmune Neutropenia.. <i>Blood</i> , 2004 , 104, 1459-1459 | 1 | |
| 7 | In Search for the Specificity of Clonal CTL in T-LGL Leukemia - Generation of Soluble LGL-Derived T Cell Receptor.. <i>Blood</i> , 2004 , 104, 4645-4645 | 2.2 | |
| 6 | High-Resolution Genomic Scan for Cryptic Chromosomal Lesions in MDS and AML.. <i>Blood</i> , 2004 , 104, 3427-3427 | 2.2 | |
| 5 | Efficient Identification of T-Cell Clones Associated with Graft-Versus-Host Disease (GvHD) in Target Tissue for Subsequent Detection in Peripheral Blood.. <i>Blood</i> , 2004 , 104, 2243-2243 | 2.2 | |
| 4 | Hematopoietic stem cells in aplastic anemia. <i>Archives of Medical Research</i> , 2003 , 34, 520-7 | 6.6 | 34 |
| 3 | Immune pathophysiology of aplastic anemia. <i>International Journal of Hematology</i> , 2002 , 76 Suppl 1, 207-14 | 20 | |
| 2 | Genetic and environmental effects in paroxysmal nocturnal hemoglobinuria: this little PIG-A goes "Why? Why? Why?". <i>Journal of Clinical Investigation</i> , 2000 , 106, 637-41 | 15.9 | 63 |
| 1 | The pathophysiology of acquired aplastic anemia. <i>New England Journal of Medicine</i> , 1997 , 336, 1365-72 | 59.2 | 417 |