

# Cytogenetic Studies in 174 Consecutive Patients With Prader-Willi Syndromes

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Citation Report

#	ARTICLE	IF	CITATIONS
1	Chromosome Abnormalities in Malignant Hematologic Disorders. Mayo Clinic Proceedings, 1985, 60, 675-689.	3.0	38
2	Refined chromosome analysis as an independent prognostic indicator in de novo myelodysplastic syndromes. Blood, 1986, 67, 1721-1730.	1.4	223
3	Myelodysplasia Progressing to Acute Myeloblastic Leukemia in an HTLV-III Virus-Positive Homosexual Man with AIDS-Related Complex. American Journal of Clinical Pathology, 1986, 86, 788-791.	0.7	40
4	PRIMARY ACQUIRED SIDEROBLASTIC ANAEMIA. British Journal of Haematology, 1986, 64, 415-418.	2.5	17
5	Myelodysplastic Syndromes. Acta Haematologica, 1987, 78, 91-93.	1.4	4
6	Isodicentric X chromosome in a patient with myelodysplastic syndrome. Cancer Genetics and Cytogenetics, 1987, 27, 215-218.	1.0	9
7	Cytogenetic findings in siblings with a myelodysplastic syndrome. Cancer Genetics and Cytogenetics, 1987, 27, 241-249.	1.0	9
8	Trisomy 9 in hematologic disorders: Possible association with primary thrombocytosis. Cancer Genetics and Cytogenetics, 1987, 27, 73-78.	1.0	13
9	Associations between morphology, karyotype, and clinical features in myeloid leukemias. Human Pathology, 1987, 18, 211-225.	2.0	142
10	Establishment and characterization of a human myeloid cell line from Philadelphia chromosome-negative myeloblastic leukemia arising in a patient with myelodysplastic syndrome. Blood, 1987, 70, 1665-1672.	1.4	16
11	Myelodysplastic syndromes. A multiparametric study of prognostic factors in 336 patients. Cancer, 1987, 60, 3029-3032.	4.1	39
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13	5 Myelodysplastic syndromes. Best Practice and Research: Clinical Haematology, 1987, 1, 389-426.	1.1	44
14	Chromosome abnormalities and karyotypic evolution in 83 patients with myelodysplastic syndrome and predictive value for prognosis. Cancer, 1988, 62, 1129-1138.	4.1	125
15	Translocation t(11;21)(q24;q11.2) is a new nonrandomly occurring chromosome change in myelodysplastic syndromes. Cancer Genetics and Cytogenetics, 1988, 34, 33-40.	1.0	7
16	Is trisomy 11 another nonrandom chromosomal anomaly in acute nonlymphocytic leukemia and myelodysplastic syndromes?. Cancer Genetics and Cytogenetics, 1988, 35, 205-211.	1.0	14
17	Chronic myelomonocytic leukemia with trisomy 8 and a related clone with trisomy 8 and t(15;17). Cancer Genetics and Cytogenetics, 1988, 32, 287-292.	1.0	2
18	The Bone Marrow in Myeloproliferative and Dysmyelopoietic Syndromes. Hematology/Oncology Clinics of North America, 1988, 2, 669-694.	2.2	13

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20	Myelofibrosis in primary myelodysplastic syndromes: a clinico-morphological study of 10 cases. <i>British Journal of Haematology</i> , 1989, 71, 499-504.	2.5	94
21	Refractory anemia with excess of blasts in transformation hematologic and clinical study of 52 patients. <i>Cancer</i> , 1989, 64, 2340-2346.	4.1	67
22	Chronic Myelomonocytic Leukemia: Natural History and Prognostic Determinants. <i>Mayo Clinic Proceedings</i> , 1989, 64, 1246-1254.	3.0	79
23	Refractory anemia with excess of blasts in transformation. <i>Cancer Genetics and Cytogenetics</i> , 1989, 42, 55-65.	1.0	14
24	Trisomy 13 in a case of myelodysplastic syndrome. <i>Cancer Genetics and Cytogenetics</i> , 1989, 37, 153-156.	1.0	12
25	A t(3;5) in blastic phase of a Philadelphia chromosome-negative chronic myeloid leukemia. <i>Cancer Genetics and Cytogenetics</i> , 1989, 37, 163-168.	1.0	6
26	Twenty-six patients with hematologic disorders and X chromosome abnormalities. <i>Cancer Genetics and Cytogenetics</i> , 1989, 42, 173-185.	1.0	54
27	Acute leukemia with abnormal thrombopoiesis and inversions of chromosome 3. <i>Cancer Genetics and Cytogenetics</i> , 1989, 39, 167-179.	1.0	62
28	Trisomy 20 in acute myelogenous leukemia. <i>Cancer Genetics and Cytogenetics</i> , 1989, 39, 25-28.	1.0	3
29	Involvement of bands 9q21-q22 in five cases of acute nonlymphocytic leukemia. <i>Cancer Genetics and Cytogenetics</i> , 1989, 39, 55-64.	1.0	19
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38	Results of chromosome studies and their relation to morphology, course, and prognosis in 120 patients with de novo myelodysplastic syndrome. <i>Cancer Genetics and Cytogenetics</i> , 1990, 44, 15-26.	1.0	112
39	Translocation t(12;13)(p13.3;q12.2) is Not Restricted to Lymphoid Malignancies; Report of a Further Case with Hypereosinophilia. <i>Leukemia and Lymphoma</i> , 1991, 4, 211-213.	1.3	1
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51	Predominance of chromosome 5 deletions in myeloid neoplasia associated with solid tumors managed by surgical excision. <i>Cancer Genetics and Cytogenetics</i> , 1992, 58, 125-129.	1.0	0
52	Translocation (8;21) in two cases of refractory anemia with excess of blasts in transformation. <i>Cancer Genetics and Cytogenetics</i> , 1992, 58, 76-78.	1.0	18
53	Cytogenetic studies in 112 cases of untreated myelodysplastic syndromes. <i>Cancer Genetics and Cytogenetics</i> , 1992, 64, 12-20.	1.0	74
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64	Analysis of karyotype, SCE, and point mutation of RAS oncogene in Indian MDS patients. <i>Cancer Genetics and Cytogenetics</i> , 1993, 65, 12-20.	1.0	14
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