# Robert J Klaassen

List of Publications by Year
in descending order



| 154 | 3,453 | 147801 | 168389 |
|---|---|---|---|
| papers | citations | 31 h-index | 53 g-index |

| 157 | 157 | 157 | 4227 |
|---|---|---|---|
| all docs | docs citations | times ranked | citing authors |

| # | Article | IF | Citations |
|---|---|---|---|
| 1 | Health-related quality of life and fatigue in children and adults with pyruvate kinase deficiency. Blood Advances, 2022, 6, 1844-1853. | 5.2 | 12 |
| 2 | Poor outcome after hematopoietic stem cell transplantation of patients with unclassified inherited bone marrow failure syndromes. European Journal of Haematology, 2022, 108, 278-287. | 2.2 | 1 |
| 3 | Novel cysteine substitution p.(Cys1084Tyr) causes variable expressivity of qualitative and quantitative VWF defects. Blood Advances, 2022, , . | 5.2 | 0 |
| 4 | Integration of Patient-reported Outcome Measures in Pediatric Hematology: A Qualitative Methods Study. Journal of Pediatric Hematology/Oncology, 2022, Publish Ahead of Print, . | 0.6 | 0 |
| 5 | Patient-Reported Outcomes Measurement in Radiation Oncology: Interpretation of Individual Scores and Change over Time in Clinical Practice. Current Oncology, 2022, 29, 3093-3103. | 2.2 | 5 |
| 6 | SARSâ€CoVâ€2 vaccination in pediatric patients with immune thrombocytopenia. Pediatric Blood and Cancer, 2022, 69, e29760. | 1.5 | 3 |
| 7 | Successful use of recombinant activated factor VII administered via automated bolus pump following emergency laparoscopic appendectomy in a patient with mild congenital FVII deficiency: Case report. Pediatric Blood and Cancer, 2021, 68, e28974. | 1.5 | 0 |
| 8 | Updating the Canadian Hemophilia Outcomesâ€"Kidsâ€™ Life Assessment Tool (CHOâ€KLAT) in the era of extended halfâ€life clotting factor concentrates. Research and Practice in Thrombosis and Haemostasis, 2021, 5, 403-411. | 2.3 | 5 |
| 9 | Quality of life is an important indication for secondâ€line treatment in children with immune thrombocytopenia. Pediatric Blood and Cancer, 2021, 68, e29023. | 1.5 | 4 |
| 10 | Measuring the impact of hemophilia on families: Development of the Hemophilia Family Impact Tool (Hâ€FIT). Research and Practice in Thrombosis and Haemostasis, 2021, 5, e12519. | 2.3 | 1 |
| 11 | Patterns of joint damage in severe haemophilia A treated with prophylaxis. Haemophilia, 2021, 27, 666-673. | 2.1 | 1 |
| 12 | Preferences and Health-Related Quality-of-Life Related to Disease and Treatment Features for Patients with Hemophilia A in a Canadian General Population Sample. Patient Preference and Adherence, 2021, Volume 15, 1407-1417. | 1.8 | 5 |
| 13 | Factor product utilization and health outcomes in patients with haemophilia A and B on extended halfâ€life concentrates: A Canadian observational study of realâ€world outcomes. Haemophilia, 2021, 27, 751-759. | 2.1 | 9 |
| 14 | Health services use by children identified as heterozygous hemoglobinopathy mutation carriers via newborn screening. BMC Pediatrics, 2021, 21, 296. | 1.7 | 1 |
| 15 | Successful treatment of pediatric primary hepatic Burkitt lymphoma using rituximab: A case report. Pediatric Blood and Cancer, 2021, 68, e29259. | 1.5 | 1 |
| 16 | Phase 3 randomised trial of eltrombopag versus standard first-line pharmacological management for newly diagnosed immune thrombocytopaenia (ITP) in children: study protocol. BMJ Open, 2021, 11, e044885. | 1.9 | 2 |
| 17 | Characterization of physical literacy in children with chronic medical conditions compared with healthy controls: a cross-sectional study. Applied Physiology, Nutrition and Metabolism, 2021, 46, 1073-1082. | 1.9 | 7 |
| 18 | Immune tolerance induction using Fcâ€fusionâ€protein recombinant factor IX in severe haemophilia B. Haemophilia, 2021, 27, e776-e779. | 2.1 | 0 |

| # | Article | IF | Citations |
|---|---|---|---|
| 19 | The impact of extended halfâ€life factor concentrates on patient reported health outcome measures in persons with hemophilia A and hemophilia B. Research and Practice in Thrombosis and Haemostasis, 2021, 5, e12601. | 2.3 | 5 |
| 20 | Magnetic resonance imaging in boys with severe hemophilia A: Serial and endâ€ofâ€study findings from the Canadian Hemophilia Primary Prophylaxis Study. Research and Practice in Thrombosis and Haemostasis, 2021, 5, e12565. | 2.3 | 4 |
| 21 | Patient-reported Outcome Measures in Pediatric Non-Malignant Hematology: A Systematic Review. Journal of Pediatric Hematology/Oncology, 2021, 43, 121-134. | 0.6 | 5 |
| 22 | Evaluating the Impact of Thrombopoietin Receptor Agonists Medications on Patient Outcomes and Quality of Life in Pediatric Immune Thrombocytopenia. Blood, 2021, 138, 4072-4072. | 1.4 | 0 |
| 23 | Pharmacist Integration into the Hemophilia Treatment Centre: A Canadian Pilot Project to Optimize Treatment and Improve Cost-Savings Using Pharmacokinetic Assessments of Hemophilia Patients. Blood, 2021, 138, 832-832. | 1.4 | 0 |
| 24 | Development of the pyruvate kinase deficiency diary and pyruvate kinase deficiency impact assessment: Diseaseâ€specific assessments. European Journal of Haematology, 2020, 104, 427-434. | 2.2 | 9 |
| 25 | Fatigue in children and adolescents with immune thrombocytopenia. British Journal of Haematology, 2020, 191, 98-106. | 2.5 | 18 |
| 26 | Androgen therapy in inherited bone marrow failure syndromes: analysis from the Canadian Inherited Marrow Failure Registry. British Journal of Haematology, 2020, 189, 976-981. | 2.5 | 8 |
| 27 | Hemophilia prophylaxis adherence and bleeding using a tailored, frequencyâ€escalated approach: The Canadian Hemophilia Primary Prophylaxis Study. Research and Practice in Thrombosis and Haemostasis, 2020, 4, 318-325. | 2.3 | 16 |
| 28 | Reduction of Extramedullary Complications in Patients With Acute Myeloid Leukemia/Myelodysplastic Syndrome Treated With Azacitidine. Journal of Pediatric Hematology/Oncology, 2020, 42, 170-174. | 0.6 | 1 |
| 29 | IVMP+IVIG raises platelet counts faster than IVIG alone: results of a randomized, blinded trial in childhood ITP. Blood Advances, 2020, 4, 1492-1500. | 5.2 | 16 |
| 30 | Syrian Refugees and Their Impact on Health Service Delivery in the Pediatric Hematology/Oncology Clinics Across Canada. Journal of Pediatric Hematology/Oncology, 2020, 42, e107-e109. | 0.6 | 5 |
| 31 | Cellular and molecular architecture of hematopoietic stem cells and progenitors in genetic models of bone marrow failure. JCI Insight, 2020, 5, . | 5.0 | 6 |
| 32 | Genotypic and Phenotypic Spectrum of Dyskeratosis Congenita: Results from the Canadian Inherited Marrow Failure Registry. Blood, 2020, 136, 8-9. | 1.4 | 0 |
| 33 | Affairs of the heart, does congenital heart disease make you happier?. Archives of Disease in Childhood, 2019, 104, 105-105. | 1.9 | 0 |
| 34 | Validation of the school age selfâ€administered pediatric bleeding questionnaire (Selfâ€PBQ) in children aged 8â€“12Âyears. Pediatric Blood and Cancer, 2019, 66, e27709. | 1.5 | 5 |
| 35 | Secondâ€line treatments in children with immune thrombocytopenia: Effect on platelet count and patientâ€centered outcomes. American Journal of Hematology, 2019, 94, 741-750. | 4.1 | 37 |
| 36 | Reanalysing genomic data by normalized coverage values uncovers CNVs in bone marrow failure gene panels. Npj Genomic Medicine, 2019, 4, 30. | 3.8 | 3 |

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|---|---|---|---|
| 37 | The social impact of early psychological maturity in adolescents with cancer. Psycho-Oncology, 2019, 28, 586-592. | 2.3 | 11 |
| 38 | Development of Patient-Reported Outcome Measures (Symptoms and Impacts) in Adults with Pyruvate Kinase Deficiency. Blood, 2019, 134, 3447-3447. | 1.4 | 0 |
| 39 | Physician decision making in selection of secondâ€line treatments in immune thrombocytopenia in children. American Journal of Hematology, 2018, 93, 882-888. | 4.1 | 30 |
| 40 | Molecular analysis and genotypeâ€phenotype correlation of Diamondâ€Blackfan anemia. Clinical Genetics, 2018, 93, 320-328. | 2.0 | 26 |
| 41 | The burden of disease in pyruvate kinase deficiency: Patientsâ€™ perception of the impact on healthâ€related quality of life. European Journal of Haematology, 2018, 101, 758-765. | 2.2 | 25 |
| 42 | Tailored frequency-escalated primary prophylaxis for severe haemophilia A: results of the 16-year Canadian Hemophilia Prophylaxis Study longitudinal cohort. Lancet Haematology,the, 2018, 5, e252-e260. | 4.6 | 31 |
| 43 | Normalizing hepcidin predicts TMPRSS6 mutation status in patients with chronic iron deficiency. Blood, 2018, 132, 448-452. | 1.4 | 16 |
| 44 | Predictors of Hydroxyurea Use in Children with Sickle Cell Disease. Blood, 2018, 132, 4934-4934. | 1.4 | 0 |
| 45 | Observational Study of Real-World Factor Utilization and Health Outcomes in Patients with Hemophilia in Canada. Blood, 2018, 132, 4813-4813. | 1.4 | 1 |
| 46 | Increasing Incidence and Prevalence of Pathologic Hemoglobinopathies Among Children in Ontario, Canada from 1991-2013. Blood, 2018, 132, 4698-4698. | 1.4 | 3 |
| 47 | Generation and optimization of the selfâ€administered pediatric bleeding questionnaire and its validation as a screening tool for von Willebrand disease. Pediatric Blood and Cancer, 2017, 64, e26588. | 1.5 | 20 |
| 48 | Paediatric health-related quality of life: what is it and why should we measure it?. Archives of Disease in Childhood, 2017, 102, 393-400. | 1.9 | 77 |
| 49 | Patterns and influences in healthâ€related quality of life in children with immune thrombocytopenia: A study from the Dallas ITP Cohort. Pediatric Blood and Cancer, 2017, 64, e26405. | 1.5 | 19 |
| 50 | Evaluation of the Greek TranQol: a novel questionnaire for measuring quality of life in transfusion-dependent thalassemia patients. Annals of Hematology, 2017, 96, 1937-1944. | 1.8 | 11 |
| 51 | Severe Neutropenia and Anemia in a Child With Epilepsy and Copper Deficiency on a Ketogenic Diet. Pediatric Neurology, 2017, 76, 93-94. | 2.1 | 8 |
| 52 | Reply to comment on: Generation and optimization of the selfâ€administered pediatric bleeding questionnaire and its validation as a screening tool for von Willebrand disease. Pediatric Blood and Cancer, 2017, 64, e26763. | 1.5 | 0 |
| 53 | Pediatric Oncology Clinic Care Model: Achieving Better Continuity of Care for Patients in a Medium-sized Program. Journal of Pediatric Hematology/Oncology, 2017, 39, 476-480. | 0.6 | 2 |
| 54 | The clinical impact of copy number variants in inherited bone marrow failure syndromes. Npj Genomic Medicine, 2017, 2, . | 3.8 | 10 |

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|---|---|---|---|
| 55 | Quality of life of pediatric oncology patients: Do patient-reported outcome instruments measure what matters to patients?. Quality of Life Research, 2017, 26, 273-281. | 3.1 | 25 |
| 56 | Health-Related Quality of Life. Pediatric Oncology, 2017, , 735-747. | 0.5 | 3 |
| 57 | A Prospective Study of the Association Between Clinically Significant Bleeding in PICU Patients and Thrombocytopenia or Prolonged Coagulation Times*. Pediatric Critical Care Medicine, 2017, 18, e455-e462. | 0.5 | 11 |
| 58 | Case 1: A newborn with pancytopenia. Paediatrics and Child Health, 2016, 21, 9-11. | 0.6 | 0 |
| 59 | Risk stratification in febrile neutropenic episodes in adolescent/young adult patients with cancer. European Journal of Cancer, 2016, 64, 101-106. | 2.8 | 15 |
| 60 | Thrombopoietin Receptor Agonist Use in Children: Data From the Pediatric ITP Consortium of North America ICON2 Study. Pediatric Blood and Cancer, 2016, 63, 1407-1413. | 1.5 | 70 |
| 61 | Reply to: Vitamin insufficiencies/deficiencies in relation to sickle cell disease severity and associated morbidity. Pediatric Blood and Cancer, 2016, 63, 2250-2251. | 1.5 | 0 |
| 62 | Response to treatment with azacitidine in children with advanced myelodysplastic syndrome prior to hematopoietic stem cell transplantation. Haematologica, 2016, 101, 1508-1515. | 3.5 | 27 |
| 63 | Azacitidine in Children with Advanced Myelodysplastic Syndrome as a Bridging Treatment Prior to Hematopoietic Stem Cell Transplantation. Biology of Blood and Marrow Transplantation, 2016, 22, S263-S264. | 2.0 | 0 |
| 64 | Prospective international validation of the Quality of Life in Myelodysplasia Scale (QUALMS). Haematologica, 2016, 101, 781-788. | 3.5 | 50 |
| 65 | Generic and disease-specific quality of life among youth and young men with Hemophilia in Canada. BMC Hematology, 2016, 16, 13. | 2.6 | 20 |
| 66 | Nutrient Insufficiencies/Deficiencies in Children With Sickle Cell Disease and Its Association With Increased Disease Severity. Pediatric Blood and Cancer, 2016, 63, 1060-1064. | 1.5 | 29 |
| 67 | Hospitalizations in pediatric patients with immune thrombocytopenia in the United States. Platelets, 2016, 27, 472-478. | 2.3 | 14 |
| 68 | Predicting microbiologically defined infection in febrile neutropenic episodes in children: global individual participant data multivariable meta-analysis. British Journal of Cancer, 2016, 114, 623-630. | 6.4 | 47 |
| 69 | Factors Impacting Quality of Life in Thalassemia Patients; Results from the Intercontinenthal Collaborative Study. Blood, 2016, 128, 3633-3633. | 1.4 | 15 |
| 70 | Clinical Characteristics and Quality of Life of Children with ITP Starting Second Line Treatments: Data from the ITP Consortium of North America ICON1 Study. Blood, 2016, 128, 249-249. | 1.4 | 7 |
| 71 | a Randomized, Double-Blinded, Placebo Controlled Study of IVIG Vs. IVIG with High Dose Methylprednisolone in Rapidly Augmenting Platelet Counts in Childhood ITP. Blood, 2016, 128, 868-868. | 1.4 | 0 |
| 72 | Physician Factors Determining Treatment Decisions in Selecting Second Line Agents for Pediatric ITP. Blood, 2016, 128, 1008-1008. | 1.4 | 0 |

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|---|---|---|---|
| 73 | Comparison of Bleeding Tools in a Cohort of Pediatric Patients with ITP: Data from the Pediatric ITP Consortium of North America ICON1 Study. Blood, 2016, 128, 4752-4752. | 1.4 | 0 |
| 74 | The impact of category, cytopathology and cytogenetics on development and progression of clonal and malignant myeloid transformation in inherited bone marrow failure syndromes. Haematologica, 2015, 100, 633-642. | 3.5 | 26 |
| 75 | Is the <scp>G</scp>ive <scp>Y</scp>outh a <scp>V</scp>oice questionnaire an appropriate measure of teenâ€centred care in paediatric oncology: a <scp>R</scp>asch measurement theory analysis. Health Expectations, 2015, 18, 1686-1697. | 2.6 | 3 |
| 76 | Sports Participation in Children and Adolescents with Immune Thrombocytopenia (ITP). Pediatric Blood and Cancer, 2015, 62, 2223-2225. | 1.5 | 18 |
| 77 | Experience with central venous access devices (<scp>CVAD</scp>s) in the Canadian hemophilia primary prophylaxis study (<scp>CHPS</scp>). Haemophilia, 2015, 21, 469-476. | 2.1 | 13 |
| 78 | Improving diagnostic precision, care and syndrome definitions using comprehensive next-generation sequencing for the inherited bone marrow failure syndromes. Journal of Medical Genetics, 2015, 52, 575-584. | 3.2 | 78 |
| 79 | Thalassaemia in children: from quality of care to quality of life. Archives of Disease in Childhood, 2015, 100, 1051-1057. | 1.9 | 22 |
| 80 | Genes Influencing the Development and Severity of Chronic ITP Identified through Whole Exome Sequencing. Blood, 2015, 126, 73-73. | 1.4 | 6 |
| 81 | Second Line ITP Therapy in the UK: Results from the UK Paediatric ITP Registry. Blood, 2015, 126, 3478-3478. | 1.4 | 0 |
| 82 | Copy Number Variants Underlying Inherited Bone Marrow Failure Syndromes. Blood, 2015, 126, 2416-2416. | 1.4 | 0 |
| 83 | Molecular Analysis of Diamond Blackfan Anemia and Genotype-Phenotype Correlation: Experience from the Canadian Inherited Marrow Failure Registry. Blood, 2015, 126, 3621-3621. | 1.4 | 1 |
| 84 | Validation and reliability of a diseaseâ€specific quality of life measure (the) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 307 Td (<scp>T</scp>r Haematology, 2014, 164, 431-437. | 2.5 | 36 |
| 85 | Caregiving, single parents and cumulative stresses when caring for a child with cancer. Child: Care, Health and Development, 2014, 40, 184-194. | 1.7 | 51 |
| 86 | Understanding the healthcare experiences of teenaged cancer patients and survivors. Child: Care, Health and Development, 2014, 40, 723-730. | 1.7 | 8 |
| 87 | Considering quality of life for children with cancer: a systematic review of patient-reported outcome measures and the development of a conceptual model. Quality of Life Research, 2014, 23, 771-789. | 3.1 | 67 |
| 88 | Patient-reported outcomes for the myelodysplastic syndromes: a new MDS-specific measure of quality of life. Blood, 2014, 123, 451-452. | 1.4 | 27 |
| 89 | Mutations in TRNT1 cause congenital sideroblastic anemia with immunodeficiency, fevers, and developmental delay (SIFD). Blood, 2014, 124, 2867-2871. | 1.4 | 162 |
| 90 | Pediatric Oncology Clinic Care Model: How to Care for Patients to Achieve Better Continuity of Care in a Medium Sized Program. Blood, 2014, 124, 6000-6000. | 1.4 | 0 |

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|---|---|---|---|
| 91 | Preliminary International Validation of the Quality of Life in Myelodysplasia Scale (QUALMS). Blood, 2014, 124, 1328-1328. | 1.4 | 0 |
| 92 | Risk Factors for Poor Survival after Hematopoietic Stem Cell Transplantation in Inherited Bone Marrow Failure Syndromes. Blood, 2014, 124, 2592-2592. | 1.4 | 0 |
| 93 | Application of Novel Next Generation Sequencing Gene Panel Assay to Genetic and Clinical Diagnosis of Inherited Bone Marrow Failure Syndromes. Blood, 2014, 124, 257-257. | 1.4 | 0 |
| 94 | Familyâ€centred care: a qualitative study of Chinese and South Asian immigrant parents' experiences of care in paediatric oncology. Child: Care, Health and Development, 2013, 39, 185-193. | 1.7 | 26 |
| 95 | Quality of life in immune thrombocytopenia following treatment. Archives of Disease in Childhood, 2013, 98, 895-897. | 1.9 | 17 |
| 96 | Evaluation of health related quality of life in children with immune thrombocytopenia with the PedsQLâ„¢ 4.0 Generic Core Scales: a study on behalf of the pays de Loire pediatric hematology network. Health and Quality of Life Outcomes, 2013, 11, 193. | 2.4 | 11 |
| 97 | Quality of life in childhood immune thrombocytopenia: International validation of the kids' ITP tools. Pediatric Blood and Cancer, 2013, 60, 95-100. | 1.5 | 33 |
| 98 | Understanding the Importance of Using Patient-Reported Outcome Measures in Patients With Immune Thrombocytopenia. Seminars in Hematology, 2013, 50, S39-S42. | 3.4 | 10 |
| 99 | Single-Parent Caregivers of Children with Cancer. Journal of Pediatric Oncology Nursing, 2013, 30, 45-55. | 1.5 | 26 |
| 100 | Standardization of bleeding assessment in immune thrombocytopenia: report from the International Working Group. Blood, 2013, 121, 2596-2606. | 1.4 | 179 |
| 101 | A novel syndrome of congenital sideroblastic anemia, B-cell immunodeficiency, periodic fevers, and developmental delay (SIFD). Blood, 2013, 122, 112-123. | 1.4 | 101 |
| 102 | A survey of the management of newborns with severe hemophilia in Canada. Paediatrics and Child Health, 2013, 18, 189-193. | 0.6 | 3 |
| 103 | Utility Of The Immature Platelet Fraction In Predicting Recovery With Or Without Treatment In Pediatric Immune Thrombocytopenia. Blood, 2013, 122, 3546-3546. | 1.4 | 1 |
| 104 | Introducing the Tran Qol: A New Disease-Specific Quality of Life Measure for Children and Adults with Thalassemia Major. Journal of Blood Disorders & Transfusion, 2013, 04, . | 0.1 | 9 |
| 105 | Characterization Of The Immature Platelet Fraction (IPF) Parameters In Children With Immune Thrombocytopenia. Blood, 2013, 122, 3547-3547. | 1.4 | 1 |
| 106 | An evidence-based threshold for thrombocytopenia associated with clinically significant bleeding in pediatric intensive care unit patients*. Pediatric Critical Care Medicine, 2012, 13, e316-e322. | 0.5 | 6 |
| 107 | Understanding the health impact of caregiving: a qualitative study of immigrant parents and single parents of children with cancer. Quality of Life Research, 2012, 21, 1595-1605. | 3.1 | 48 |
| 108 | A <scp>C</scp>anadian survey of selfâ€infusion practices in persons with haemophilia A. Haemophilia, 2012, 18, e403-5. | 2.1 | 2 |

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|---|---|---|---|
| 109 | Rasch analysis of the PedsQL: an increased understanding of the properties of a rating scale. Journal of Clinical Epidemiology, 2012, 65, 1117-1123. | 5.0 | 40 |
| 110 | Impact of caring for a child with cancer on single parents compared with parents from twoâ€parent families. Pediatric Blood and Cancer, 2012, 58, 74-79. | 1.5 | 19 |
| 111 | Response to steroids predicts response to rituximab in pediatric chronic immune thrombocytopenia. Pediatric Blood and Cancer, 2012, 58, 221-225. | 1.5 | 29 |
| 112 | Pilot study of the effect of romiplostim on child healthâ€related quality of life (HRQoL) and parental burden in immune thrombocytopenia (ITP). Pediatric Blood and Cancer, 2012, 58, 395-398. | 1.5 | 41 |
| 113 | Immigrant to Canada, newcomer to childhood cancer: a qualitative study of challenges faced by immigrant parents. Psycho-Oncology, 2012, 21, 558-562. | 2.3 | 34 |
| 114 | Communication and language challenges experienced by Chinese and South Asian immigrant parents of children with cancer in Canada: Implications for health services delivery. Pediatric Blood and Cancer, 2012, 58, 572-578. | 1.5 | 47 |
| 115 | Outcome and Clinical Characteristics of Clonal and Malignant Myeloid Transformation in Inherited Bone Marrow Failure Syndromes. Blood, 2012, 120, 1266-1266. | 1.4 | 0 |
| 116 | Vitamin D and Zinc Deficiency Associated with Increased Pain Episodes in Children with Sickle Cell Disease. Blood, 2012, 120, 3237-3237. | 1.4 | 3 |
| 117 | Comparative analysis of Shwachman-Diamond syndrome to other inherited bone marrow failure syndromes and genotype-phenotype correlation. Clinical Genetics, 2011, 79, 448-458. | 2.0 | 50 |
| 118 | Identifying determinants of quality of life of children with cancer and childhood cancer survivors: a systematic review. Supportive Care in Cancer, 2011, 19, 1275-1287. | 2.2 | 105 |
| 119 | Exploring predictors of optimism among parents of children with cancer. Psycho-Oncology, 2011, 20, 411-418. | 2.3 | 29 |
| 120 | Quality of life during active treatment for pediatric acute lymphoblastic leukemia. International Journal of Cancer, 2011, 128, 1213-1220. | 5.1 | 87 |
| 121 | Parents of children with cancer: Which factors explain differences in healthâ€related quality of life. International Journal of Cancer, 2011, 129, 1190-1198. | 5.1 | 68 |
| 122 | Genetic analysis of inherited bone marrow failure syndromes from one prospective, comprehensive and population-based cohort and identification of novel mutations. Journal of Medical Genetics, 2011, 48, 618-628. | 3.2 | 55 |
| 123 | Fatal Neurotoxicity in a Patient With Down Syndrome Treated With Chemotherapy, Irradiation, Stem Cell Transplant, and Clofarabine. Journal of Pediatric Hematology/Oncology, 2010, 32, e111-e113. | 0.6 | 3 |
| 124 | Health-related quality of life outcomes for patients with immune thrombocytopenic purpura. Annals of Hematology, 2010, 89, 51-54. | 1.8 | 7 |
| 125 | Evaluating the ability to detect change of healthâ€related quality of life in children with Hodgkin disease. Cancer, 2010, 116, 1608-1614. | 4.1 | 29 |
| 126 | Nurses provide valuable proxy assessment of the healthâ€related quality of life of children with Hodgkin disease. Cancer, 2010, 116, 1602-1607. | 4.1 | 20 |

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|---|---|---|---|
| 127 | Caregiving Demands in Parents of Children With Cancer: Psychometric Validation of the Care of My Child With Cancer Questionnaire. Journal of Pediatric Nursing, 2010, 25, 258-263. | 1.5 | 11 |
| 128 | Response to Steroids Predicts Response to Rituximab In Pediatric Chronic Immune Thrombocytopenia.. Blood, 2010, 116, 3681-3681. | 1.4 | 0 |
| 129 | The North American Chronic Immune Thrombocytopenia Registry (NACIR): Demographics and Treatment Responses. Blood, 2010, 116, 2509-2509. | 1.4 | 0 |
| 130 | A case of moyamoya syndrome and hemoglobin E/betaâ€thalassemia. Pediatric Blood and Cancer, 2009, 52, 422-424. | 1.5 | 15 |
| 131 | Relationships among bleeding severity, healthâ€related quality of life, and platelet count in children with immune thrombocytopenic purpura. Pediatric Blood and Cancer, 2009, 53, 652-654. | 1.5 | 51 |
| 132 | Factors affecting the delivery of familyâ€centered care in pediatric oncology. Pediatric Blood and Cancer, 2009, 53, 1079-1085. | 1.5 | 24 |
| 133 | Parental optimism in poor prognosis pediatric cancers. Psycho-Oncology, 2009, 18, 783-788. | 2.3 | 11 |
| 134 | Evaluating familyâ€centred service in paediatric oncology with the measure of processes of care (MPOCâ€20). Child: Care, Health and Development, 2009, 35, 16-22. | 1.7 | 26 |
| 135 | Identification of paediatric cancer patients with poor quality of life. British Journal of Cancer, 2009, 100, 82-88. | 6.4 | 75 |
| 136 | Measuring disease-specific quality of life in rare populations: a practical approach to cross-cultural translation. Health and Quality of Life Outcomes, 2009, 7, 92. | 2.4 | 46 |
| 137 | Iliopsoas hematoma following a bone marrow aspirate. Pediatric Blood and Cancer, 2008, 51, 146-146. | 1.5 | 0 |
| 138 | Spontaneous remission of myelodysplastic syndrome with monosomy 7 in a young boy. Cancer Genetics and Cytogenetics, 2008, 182, 122-125. | 1.0 | 22 |
| 139 | Impact of Caring for a Child With Cancer on Parentsâ€™ Health-Related Quality of Life. Journal of Clinical Oncology, 2008, 26, 5884-5889. | 1.6 | 177 |
| 140 | Clinical and Genetic Analysis of Unclassifiable Inherited Bone Marrow Failure Syndromes. Pediatrics, 2008, 122, e139-e148. | 2.1 | 49 |
| 141 | Health-related Quality of Life: Changes in Children Undergoing Chemotherapy. Journal of Pediatric Hematology/Oncology, 2008, 30, 292-297. | 0.6 | 31 |
| 142 | Validity, Reliability, and Responsiveness of a New Measure of Health-Related Quality of Life in Children with Immune Thrombocytopenic Purpura: The Kidsâ€™ ITP Tools. Journal of Pediatrics, 2007, 150, 510-515.e1. | 1.8 | 82 |
| 143 | Central venous catheter thrombosis as a cause of SVC obstruction and cardiac tamponade in a patient with Diamond-Blackfan anemia and iron overload. Pediatric Blood and Cancer, 2006, 46, 112-114. | 1.5 | 5 |
| 144 | Disease progression in recently diagnosed patients with inherited marrow failure syndromes: A Canadian inherited marrow failure registry (CIMFR) report. Pediatric Blood and Cancer, 2006, 47, 918-925. | 1.5 | 18 |

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|---|---|---|---|
| 145 | Quantitative Assessment of Drug Treatment Side Effects in Children with Idiopathic Thrombocytopenic Purpura (ITP).. Blood, 2004, 104, 3030-3030. | 1.4 | 3 |
| 146 | High-Risk Neuroblastoma in Ontario: A Report of Experience From 1989 to 1995. Journal of Pediatric Hematology/Oncology, 2003, 25, 8-13. | 0.6 | 9 |
| 147 | Using Decision Analysis Techniques to Deal With ???Unanswerable??? Questions in Idiopathic Thrombocytopenic Purpura. Journal of Pediatric Hematology/Oncology, 2003, 25, S62-S63. | 0.6 | 2 |
| 148 | Acute immune thrombocytopenic purpura: What do we do when things go well?. Paediatrics and Child Health, 2002, 7, 384-385. | 0.6 | 1 |
| 149 | The diagnosis and treatment of von Willebrand disease in children. Paediatrics and Child Health, 2002, 7, 245-249. | 0.6 | 14 |
| 150 | Initial Bone Marrow Aspiration in Childhood Idiopathic Thrombocytopenia: Decision Analysis. The American Journal of Pediatric Hematology/oncology, 2001, 23, 511-518. | 1.3 | 26 |
| 151 | â€œLow-Riskâ€•Prediction Rule for Pediatric Oncology Patients Presenting With Fever and Neutropenia. Journal of Clinical Oncology, 2000, 18, 1012-1012. | 1.6 | 207 |
| 152 | Randomized Placebo-Controlled Trial of Oral Antibiotics in Pediatric Oncology Patients at Low-Risk With Fever and Neutropenia. The American Journal of Pediatric Hematology/oncology, 2000, 22, 405-411. | 1.3 | 60 |
| 153 | Neuroblastoma in Down syndrome. , 1999, 33, 125-125. | | 1 |
| 154 | Clinical Measures. , 0, , 89-98. | | 0 |