

Book chapters and article reviews:

Bartolo C, and Viswanatha DS (March 2000) Molecular diagnostics in pediatric acute leukemias. Clinics in Laboratory Medicine: Acute leukemias 20 (1) 139-182.

Bartolo C and Viswanatha DS (2001) Molecular methods for minimal residual disease detection in hematolymphoid diseases: Principles and Applications. Journal of Clinical Ligand Assay 24 (2): 76-93.

Journal Articles:

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Beaubier NT, Hart AP, **Bartolo C**, Willman CL, Viswanatha D (2000) Comparison of capillary electrophoresis and polyacrylamide gel electrophoresis for the evaluation of T and B cell clonality by polymerase chain reaction. Diagnostic Molecular Pathology 9 (3):121-131

Bartolo C, Mendell JR, Prior TW (1998) Identification of a missense mutation in a Friedreich's Ataxia patient: Implications for diagnosis and carrier studies. American Journal of Medical Genetics 79: 396-399.

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Prior TW, **Bartolo C**, Papp AC, Snyder PJ, Sedra MS, Burghes AHM, Kissel JT, Tsao CY, Mendell JR (1997) Dystrophin expression in a Duchenne muscular dystrophy patient with a frameshift deletion. Neurology 48: 486-488.

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Prior TW, **Bartolo C**, Papp AC, Snyder PJ, Sedra MS, Burghes ARM, Mendell JR (1994) Identification of a Missense Mutation, Single Base Deletion, and a Polymorphism in the Dystrophin Exon 16. Human Molecular Genetics 3: 1173-1174.

Prior TW, Papp AC, Snyder PJ, Burghes AHM, **Bartolo C**, Sedra MS, Western LM, Mendell JR (1993) A Missense Mutation in the Dystrophin Gene in a Duchenne Muscular Dystrophy Patient. *Nature Genetics* 4: 357-360.

Prior TW, Papp AC, Snyder PJ, Burghes AHM, Sedra MS, Western LM, **Bartolo C**, Mendell JR (1993) Exon 44 Nonsense Mutation in Two Duchenne Muscular Dystrophy Brothers Detected by Heteroduplex Analysis. *Human Mutation* 2: 192-195.

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