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Darine Villela ¹ , Silvia Souza da Costa ¹ , Angela M. Vianna-Morgante ¹ , Ana G	C. V. Krepischi ¹ ,
Carla Rosenberg ¹	

¹Department of Genetics and Evolutionary Biology, Institute of Biosciences, University of São Paulo, Rua do Matão 277, 05508-090, São Paulo, SP, Brazil

Running title: Targeted sequencing for detecting CNVs and SNVs

Keywords: exome sequencing, targeted sequencing, DNA copy number variations, CNVs, single nucleotide variants, SNVs, chromosomal imbalances

Address for correspondence: Darine Villela, PhD, Department of Genetics and Evolutionary Biology, Institute of Biosciences, University of São Paulo, Rua do Matão 277, 05508-090, São Paulo-SP, Brazil. Phone Number: +55 1130917573. E-mail: darinevillela@gmail.com.

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