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Significant Association of Caveolin-1 Single Nucleotide Polymorphisms with Childhood Leukemia in Taiwan

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Backgrounds: A growing body of evidence indicates that caveolin-1 (Cav-1) may influence the development of human cancers. However, the exact role of caveolin-1 in child leukemia is still controversial. We investigated six novel polymorphic variants of Cav-1, includes C521A (rs1997623), G14713A (rs3807987), G21985A (rs12672038). T28608A (rs3757733), T29107A (rs7804372), and G32124A (rs3807992), and analyzed the association of specific genotype with child leukemia susceptibility. Materials and Methods: In total, 266 patients with child leukemia and 266 age-matched healthy controls recruited from two major medical centers in Taiwan were genotyped investigating these polymorphisms' association with child leukemia. Results: We found that there were significant differences between child leukemia and control groups in the distributions of their genotypes (P=4.1*10° and 0.0167) and allelic frequencies (P=4.9*10⁻¹⁰ and 3.7*10⁻³) in the Cav-1 G14713A and T29107A polymorphisms, respectively. As for the haplotype analysis, those who had GG/AT or GG/AA at Cav-1 G14713A/T29107A showed a decreased risk of child leukemia compared to those with GG/TT, while those of any other combinations were of increased risk. Conclusion: The A allele of the Cav-1 G14713A is risky, the A allele of the Cav-1 T29107A is protective, for the development of child leukemia and may be novel useful genomic markers for early detection of child leukemia.

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