



Variations in *WNT3* gene are associated with incidence of non-syndromic cleft lip with or without cleft palate in a northeast Chinese population

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ABSTRACT. Non-syndromic cleft lip with or without cleft palate (NSCL/P) is a common birth defect. Several *WNT* genes are involved in craniofacial embryogenesis, and therefore may play an important role in the etiology of NSCL/P. Two SNPs (rs3809857 and rs9890413) in the *WNT3* gene were subjected to case-control and case-parent analysis by polymerase chain reaction-restriction fragment length polymorphism (PCR-RFLP) in 236 unrelated patients with NSCL/P, including 128 elementary families (185 mothers and 154 fathers), and 400 control individuals from northeast China. The rs3809857 SNP, under the assumption of a dominant model, was found to induce a 2-fold lower risk of NSCL/P $OR_{GG \text{ vs } GT + TT} = 0.605$, 95%CI = 0.436-0.839, $P = 0.003$). Moreover, the family-based association test revealed an under-

transmission for the minor allele T. On the other hand, we observed a significant association in the case-control and case-parent analysis of the SNP rs9890413. In addition, the P values for the haplotype of rs3809857-rs9890413 were observed to be statistically significant ($P = 0.004$). In conclusion, our study confirmed the association between the *WNT3* variant and NSCL/P in the population tested.

Key words: Cleft lip; Cleft palate; Family-based association test; *WNT3*