



ORAL PRESENTATION

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A NKG2D Thr72Ala polymorphism is not associated with HAM/TSP and proviral load values in Peruvian HTLV-1 infected patients with HAM/TSP and asymptomatic carriers

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Background

NKG2D is an activating receptor mainly expressed on Natural Killer (NK) cells, its ligands has an abnormal expression in virus infected cells and tumoral cells. The SNP rs2255336 G/A generates a substitution of alanine to threonine (NKG2D Thr72Ala) associated with a low and high cytotoxic activity of NK cells respectively. This study investigated the association between the NKG2D Thr72Ala polymorphism with HAM/TSP and PVL values in Peruvian HTLV-1 infected patients with HAM/TSP and asymptomatic carriers.

Methods

215 Peruvian HTLV-1 infected patients (142 asymptomatic carriers (AC) and 73 HAM/TSP) were evaluated. NKG2D Thr72Ala polymorphism were genotyped by polymerase chain reaction using allele-specific primers, 37 ancestry informative markers (AIM) were genotyped to correct by population stratification using the first three principal components. Proviral load (PVL) was measured using the endogenous retrovirus 3 (ERV-3). Association of the NKG2D Thr72Ala polymorphism with HAM/TSP and PVL were performed by univariate analysis using Pearson's chi-square test and Mann-Whitney U-test and by multivariate analysis using logistic regression and linear regression analysis. STATA software was used for all analysis.

Results

The NKG2D Thr72Ala polymorphism with G/G genotype showed a higher prevalence in asymptomatic carrier

(64.13%) respect to HAM/TSP patients (34.87%), however these differences were not statistically significant ($P > 0.05$). No association was found between the different genotypes of NKG2D Thr72Ala with PVL ($P > 0.05$), or with the presence of HAM/TSP ($P > 0.05$). The assessment of the association between allele G (OR = 2.3760; CI = 0.7744 to 7.2901) or the allele A (OR = 0.4209; CI = 0.1372 to 1.2914) with the presence of HAM/TSP were not statistically significant ($P > 0.05$).

Conclusions

The results of this study indicate the need to evaluate the association of polymorphisms in NKG2D gene with HAM/TSP and PVL in other independent populations of HTLV-1 infected individuals in order to verify an association of disease state with host genetic.

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