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Joint Meeting on

4th World Congress on

HUMAN GENETICS & GENETIC DISEASES and **3**rd International Conference on

MOLECULAR MEDICINE & DIAGNOSTICS

April 19-20, 2018 Dubai, UAE

Genetic polymorphism of apolipoprotein E in hemorrhagic stroke: Case-control study

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Aim: It is a case-control study realized at the Hospital of Constantine. It discusses the relationship between polymorphism of apolipoprotein E and hemorrhagic stroke.

Method: The determination of the polymorphism of apolipoprotein E was carried out by PCR-digestion (polymerase chain reaction) using the enzyme of restriction HhaI. The study population consisted of 81 Algerian patients with hemorrhagic stroke and 509 control subjects.

Results: Three isoforms of apolipoprotein E have been identified. The allelic distribution of apoE in the general population showed a predominance of the allele ε 3 (84.3%) followed distantly by allele ε 4 (10.7%) and ε 2 (5%) respectively. In hemorrhagic stroke patients, allele frequencies of £4 and £2 are respectively 10.5% and 3.3%. These frequencies are not statistically different as reported in the control group. The assessment of the odds ratio of patient subjects with allele $\varepsilon 4$, $\varepsilon 2$, $\varepsilon 3/\varepsilon 4$ and $\varepsilon 2/\varepsilon 3$ compared to control subjects with genotype $\varepsilon_3/\varepsilon_3$ did not show any statistical association between the polymorphism of the apoE and the set of hemorrhagic stroke.

Conclusion: The distribution of apolipoprotein E allele frequencies in the population of Constantine is similar to that of Southern Europe. The £2, £4 alleles do not appear to be implied in the occurrence of this affection; nevertheless, large additional studies are necessary to confirm these results.

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