

Int J Hum Genet, 17(2): 64-71 (2017) DOI:10.31901/24566330.2017/17.02.03

Determination of Genotoxic Damage by Comet Assay in Smokers

E. G. Karahan^{1#}, A. G. Tomatir^{1*}, I. Açikba⁰¹⁺, A. B. Er^{1\$}, F. Evyapan², B. Akdag³ and P. E. Arslan^{1^}

 ¹Pamukkale University, Department of Medical Biology, Faculty of Medicine, Denizli, Turkey
²Pamukkale University, Department of Chest Diseases, Faculty of Medicine, Denizli, Turkey
³Pamukkale University, Department of Biostatistics, Faculty of Medicine, Denizli, Turkey
E-mail: ^{1#}<elif.g.turkecan@gmail.com>, ^{1*}< tomatir@pau.edu.tr>, ¹⁺<iacikbas@pau.edu.tr>, ^{1\$}<buketer@gmail.com>, ²<fevyapan@pau.edu.tr>, ³<bakdag@pau.edu.tr>, ^{1^}<elvanars@gmail.com>

KEYWORDS Genomic Instability. Genotoxicity. Peripheral Blood. Single Cell Gel Electrophoresis. Smoking

ABSTRACT The clinical course of most diseases related to smoking has a strong relationship with genotoxicity. In this study, the researchers aimed to compare DNA damage of smokers and non-smokers to determine the genotoxic risk. In total, 50 volunteers were included in this study; 30 of them smokers and 20 of them forming the non-smoker control group. Peripheral blood samples taken from the volunteers were determined with comet assay. The researchers determined the DNA damage ratio as 12, 75 (\pm 7. 14) from smokers 10, 41 (\pm 3.41) from non-smokers (p> 0.05), and also higher DNA damage in male smokers than female ones (p<0. 05). There was no correlation between age and DNA damage. In conclusion, there was no significant difference between smokers and genomic instability was adversely affected.

Abbreviations: SCGE: Single Cell Gel Electrophoresis; DNA: Deoxyribonucleic Acid; WHO: World Health Organization; HMA: High Melting Agarose; LMA: Low Melting Agarose