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Effect of polymorphisms of selected DNA repair genes on lung cancer development with respect to chromium exposure.

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Objective: Chromium is a well-known mutagen and carcinogen involved in lung cancer development. DNA repair genes play an important role in eliminating of genetic changes caused by chromium exposure. In the present study, we investigated the polymorphisms of following DNA repair genes as XRCC3 participating in homologous recombination, hOOG1 involved in base excision repair as well as MLH1 and MSH2 functioning in mismatch repair and the risk they present towards the development of lung cancer with emphasis to effect of chromium exposure.

Material and methods: We analysed 106 individuals; 45 patients exposed to chromium with diagnosed lung cancer and 61 healthy controls. Genotypes were determined by PCR-RFLP method.

Results: We found out increased risk of lung cancer development in hOOG1 (rs1052133) CC genotype in dominat model (OR = 1.862), and in MLH1 (rs1800734) AA genotype in recessive model (OR = 4.28).

Conclusions: We conclude that gene polymorphisms in DNA repair genes may underscore the risk of lung cancer development in chromium exposed individuals.

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