DNA hazard in Furnace Operating Workers from a Power Plant

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Abstract

The aim of this study was to investigate the possible DNA damage in workers occupationally exposed to coal combustion products in furnace section of Afsin-Elbistan A power plant (Turkey). With this purpose, venous blood samples collected from 36 male power plant workers were analysed by comet assay for determining the level of DNA tail intensity. The obtained results were compared with those of control group consisting of 34 healthy male individuals. The data comparison showed that mean frequency of tail intensity was significantly higher in workers as compared to control group (P<0.05), respectively 9.94±2.51 and 8.48±2.31. Present study indicated the DNA damage in the peripheral lymphocytes of furnace operators, possibly due to the several chemical compounds present in the coal ash and gaseous emissions.

Keywords: Coal fired power plant; Afsin-Elbistan A; Power plant workers; Genotoxic risk; Comet assay

Introduction

Thermal power plants leads to the emission of significant amounts of SO2 and fine particulate matter, volatile organic compounds (VOCs) and toxic organic micro pollutants, such as polycyclic aromatic hydrocarbons (PAHs) into the environment [1,2]. These products of combustion represent a risk both to environment [3,4] and human [5,6]. Considering increased coal production and utilization, several researchers concerned with the safety and health of the workers who utilize coal and expose to coal combustion products in various workplace. Epidemiological studies have shown that exposure to PAHs is associated with time- and dose-dependent increases in risk of cancer namely lung, colon, and bladder, and adverse birth outcomes [7-9]. Beside, other concents of coal combustion products such as coal ash, quartz, carbon, nitrogen, and sulfur oxides, and trace elements have been reported to lead to various types of cancer [10,11] and also have been shown to be mutagenic and genotoxic by several studies [12-15].

A few studies concerning the genotoxic risks of workers occupationally exposed to coal combustion products in power plants, reported a significant increase in the level of chromosomal aberrations (CA), in acentric chromosome fragments and dicentric chromosome as well as the number of abnormal cells, sister chromatid exchanges (SCE), micronucleus (MN), and polyploid cells in workers as compared to control group [16,17].

In the present study, possible genotoxic risk of workers exposed to coal combustion products in the furnace section of Afsin-Elbistan A power plant, located in South-eastern Turkey, were investigated by comet assay. Afsin-Elbistan an established in 1983 is an old power plant where the pollution control is insufficient. So, the workers were exposed to the combustion products of coal-mixture of volatile substances and also coal ash, especially in the furnace.

Materials and Methods

Subjects and sampling

A total of 36 furnace operators from Afsin-Elbistan A power plant (south-eastern Turkey) and 34 healthy male from Kahramanmaras city (160 km far from the power plant) consented to participate in the study. The workers and control group were selected after questionnaire administration for obtaining information regarding age, occupation, and years of employment, life style, and health problems, if any. The years of exposure of the workers range between 9-28 years, the mean value was 21±3.22. The mean age of the workers was 46.46±5.78 years (range 37-53), it was 46.06±5.81 (range 32-55) for the control group. Venous blood samples of 5 ml were taken in heparinized tubes at the working site, codified and immediately transported at 4°C to the Toxicology Laboratory of Pharmacy Faculty (Gazi University, Ankara) for Comet assay.

Comet assay (Single-cell gel electrophoresis-SCGE)

The comet assay was performed under alkaline conditions using the method described previously by Singh et al. (18) with slight modification. The isolated lymphocytes (with Histopaste 1077 in phosphate buffer salin on ice) from heparinized blood samples were suspended (at ~2×10^5 cells/mL) were mixed with 100 μl of 0.65% low-melting-point agarose in PBS at 37°C and rapidly pipetted onto a frosted glass microscope slide precoated with 100 ml of 1% agarose, spread out with a coverslip and maintained at 4°C for 30 min to solidify. After removal of the coverslip, the slides were immerced in lysis solution (2.5 M NaCl, 100 mM Na2EDTA, 10 mM Tris, NaOH to pH 10.0, and 1% Triton X-100) for overnight at 4°C, to remove cellular proteins. Slides were initially placed in an electrophoresis tank containing 1 mM Na2EDTA and 300 mM NaOH, (pH 13) for 20 min. Afterward, the tank was set at 25 V (1.6 V/cm, 300 mA) for 20 min at an ambient temperature of 4°C. The slides were then washed three times for 5 min each, with Tris buffer (0.4 M Tris, pH 7.5), at 4°C before staining them with 65 μl ethidium bromide (EtBr 20 μg/ml) for analyzing. Analysis was carried

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out immediately after sample collection without freezing or storing. Cell viability, using trypan blue, was found to be over 95% at each time point of the study.

After the staining process, hundred cells were analysed using double slides selected randomly for examination at x200 magnification under a fluorescent microscope (Zeiss-Axioskop, Oberkochen, Germany) equipped with an excitation filter 515–560 nm and a 100 W Hg lamp. DNA migration (tail intensity) was measured, using Comet Assay III image analysis system (Perceptive Instruments, UK). All slides were coded and scored blindly.

Statistical analysis

The data were analysed with SPSS 15.0 for Windows statistical programme (SPSS, Chicago, IL). Differences between control and exposed groups were evaluated by the Mann-Whitney U test. Correlations between different variables were analysed by Spearman’s rho correlation test. For all statistical analyses, a level of at least 0.05 was used to determine significance.

Results and Discussion

The effects of exposing to coal combustion products in the furnace section of Afsin-Elbistan A power plant on the tail intensity selected as a marker of the genotoxic damage were presented in Table 1. Workers showed a significantly higher mean tail intensity (9.94±2.51) compared to the unexposed control subjects (8.48±2.31) (P<0.05).

We also analysed the data with respect to years of exposure and age of the individuals, to investigate the association between the marker and independent variables (Table 2). The results obtained from Spearman’s rho correlation analysis revealed no significant correlation between the years of exposure and tail intensity in workers (P>0.05). In neither workers nor controls subjects, the incidence of markers, show a significant correlation with age (P>0.05).

As compared to studies done to performed the environmental impact of energy production by coal, a few researches are available on the health risk of the workers occupationally exposed to coal combustion products in the power plants. In a study from Italy [19], malignant mesothelioma has been reported in power plant workers. In a study [20], the lung function impairment and respiratory symptoms of workers occupationally exposed to chemical products of burning coal and ashes. Elevated level of DNA damage might originate from the many chemical compounds present in the coal ash and also in the gaseous emissions. The mixtures of these compounds may exhibit additive interactions that can not be predicted by the any single substance.

Concurrently, to genotoxic risk of power plant workers, Bauman and Horvat [23] found significantly higher CA level in workers of a power station burning coal, than in control group. In another study, Leonard et al. [17], reported a significant increase in acentric chromosome fragments and dicentric chromosome as well as the number of abnormal cells in workers from coal fueled power plant as compared to control group. Beside, Celik et al. [24], reported considerably elevated CA, SCE, and MN levels in the peripheral lymphocytes of workers exposed to waste coal ash in a power plant. Moreover, the frequency of the polyploid cells was found to be significantly higher in the workers than in control subjects.

In the present study, the observed increase of DNA damage level in workers in terms of tail intensity were consistent with the previous cytogenetic studies [17,23,24] reported the genotoxic risks in the workers occupationally exposed to chemical products of burning coal and ashes. Elevated level of DNA damage might originate from the many chemical compounds present in the coal ash and also in the gaseous emissions. The mixtures of these compounds may exhibit additive interactions that can not be predicted by the any single substance.

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