Induction of Micronuclei in Mice Lymphocytes Exposed to Microwave and Toluene

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Abstract: Increasing applications of microwave radiation are of great concern with regard to public health. Several studies have been conducted detect effects of microwave exposure genetic material leading to negative or questionable results. The Micronucleus (MN) assay which is proved to be a useful method for detection of radiation exposure-induced cytogenetic damage was used in the present study to investigate the genotoxic effect of microwave and toluene alone and in combination in Balb/c lymphocytes. The electromagnetic field with two frequencies (980, 950 MHz, 200 KHz Mod), 5 w and 500 ppm Toluene applied for two weeks. Microwave irradiation had no significant effect on the frequency of micronucleus induced, but exposure of animals to toluene alone and in combination with microwave have significantly increased the induced micronucleus (p<0.05). Indeed combination exposure of microwave and toluene showed higher rates of micronucleus in comparison with toluene alone. This study indicated that microwave radiation cannot induce any significant cytogenetic effects but, in combination with toluene could show synergistic effect.

Key words: Microwave, toluene, lymphocytes, micronucleus, mice

INTRODUCTION

Applications of microwave have been increased in recent years due to radars and police communication systems[1], high power satellite and TV transmitters, mobile phones, microwave ovens and medical devices[2]. The biological effects of this type of exposure on living organisms have been studied by different investigations[3,4]. The absorption of electromagnetic energy can cause biological effects in tissues. The Specific Absorption Rate (SAR) has been introduced for the assessment of absorbed energy from electromagnetic fields with biological systems. SAR measurement requires complicated technical procedures[5]. Several biological effects of microwave exposure such as effects on eyes, gonads and circulatory system can be thermal effects. It has shown that endogenous biological electrical activities of can be categorized as nonthermal effects of microwave exposure which occurs in low powers[6]. Occupational and non occupational exposure to low power microwave and its biological effects have increased in recent years.

Toluene as a chemical material have various industrial applications such as Benzene production, inks, adhesives and leather industry[7] Occupational exposure of 1.278000 American workers to toluene during 1981-1982 has been reported by NIOSH[8]. Vehicles and their fueling procedures are the main sources of toluene environmental pollution and therefore transportation employees are exposed to toluene occupationally[9] Toluene introduced as an alternative for Benzene but there are controversies about its harmful biological effects of toluene on lymphocytes of workers is reported[12,13]. By talking to accounting the probable harmful biological effects of microwave exposure and toluene alone and together, we planned this study to investigate the synergistic effects of two these agents.

MATERIALS AND METHODS

Animals: We used the micronucleus assay for quantification of cytotoxic effects of microwave and toluene on balbc lymphocytes[14]. The micronucleus assay was carried by lymphocyte cells of male bulb/c mice aged 2 months weighting 18-25g. Five equal populations[6] experimental groups were used in various conditions. There was a control group for each experimental group.
Table 1: Micronucleus in lymphocytes after microwave and toluene exposure

| Groups | power [w] | frequency [MHz] | Exposure time [week] | Modulation | Toluene [ppm] | MN*/1000 |
|--------|-----------|-----------------|----------------------|------------|---------------|-----------|
| 1      | 5         | 950             | 2                    | -          | -             | 14.16±1.47 |
| control | -         | -               | -                    | -          | -             | 11.66±1.03 |
| 2      | 5         | 890             | 2                    | -          | -             | 13.83±1.47 |
| control | -         | -               | -                    | -          | -             | 12±1.26   |
| 3      | 5         | 950             | 2                    | 200        | -             | 13.5±1.37 |
| control | -         | -               | -                    | -          | -             | 12.08±0.89 |
| 4      | -         | -               | 2                    | 500        | -             | 17.66±3.66** |
| control | -         | -               | -                    | -          | -             | 11.33±1.21 |
| 5      | 5         | 950             | 2                    | -          | 500           | 25.5±4.63** |
| control | -         | -               | -                    | -          | -             | 11.83±1.16 |

*Frequency of Micro Nucleus (MN) expressed as number of micronucleus in binucleated cells (BN) on 1000

**P<0.05

Micronuclei in cultured lymphocytes: All animals were sacrificed and their splenic cells were isolated by injection of RPMI 1640 medium (Gibco) to their spleen followed by grinding true cell mesh and focal density gradient centrifugation (700g, 15min), 200 μl of each cell suspension was cultured at 37 °C and 5% CO₂ in RPMI medium containing 10% fetal serum (FCS, Gibco) and 1.5% Phytohemagglutinin (PHA, Gibco). After 17.5 hours of incubation cytochalasin B (Cyt-B, sigma, 6μ/ml final concentration) was added to cell cultures. The cells were harvested after another 24 hours of incubation and laid slide using cytospin. The frequency of binucleate cells after methanol fixation and Gisma staining using light microscope.

RESULTS

Frequency of induced micronucleus observed in 10 groups including the control and exposed groups and corresponding statically analysis for each group are given in Table 1.

Mean frequency of induced micronucleus in 5 groups were 14.16, 13.83, 13.5, 17.66 and 25.5 respectively which with their corresponding control groups are shown in Fig. 2.

ANOVA analysis showed that the differences between groups of 4 and 5 and their controls were statistically significant, but no significant difference
was observed in the other three groups (groups 1, 2 and 3). The difference between groups 5 and 6 was significant as well (p<0.5).

DISCUSSION

Microwave telecommunication systems as an electromagnetic field used in our environment leading raised the natural electromagnetic noise by a factor of about $10^{12}$. Exposure of large groups of the population to this type of electromagnetic field and the probable biological effects on living organisms have revived considerable research attention. Biological effects at a cellular and molecular level such as DNA damage, chromosomal aberration probable genotoxic effects. Microwave telecommunication systems have been reported by several investigations. Toluene as a chemical volatile agent has been used widely in various industries and as an air Pollutant could exposure population. The cytogenetic effects of toluene have been reported in recent years. We considered the genotoxic effects of microwave and toluene alone and together by micronucleus assay. MN formation is correlated with the loss of either chromosome fragments or whole chromosomes. Due to their relatively small size, the microwave-induced MN here observed are likely hypothesized to arise via a clastogenic. The present work indicates that exposure to non-ionizing radiation in the range of microwave with toluene is able to cause, increase of lymphocytes MN frequency. This finding adds further evidence in favor of the use of cytogenetic assays as a dosimetry biomarker in epidemiological studies of workers occupationally exposed to a combination of non-ionizing radiation and chemical agents. The microwave exposure of 890, 950 MHz and power of 5w has no significant effect on the frequency of micronucleus observed. This result is in agreement with an investigation that used 935.2 MHz and 4.5w power density and also agrees with other study which used RF frequencies of 380, 900 and 1800 MHz with Amplitude modulation of 0.73-3.1 MHz on incidence of Sister Chromatid Exchange (SCE) of human lymphocytes. We also showed that exposure to 500 ppm toluene increased induced frequency of MN that agrees with investigation that reported occupational exposure to 40ppm toluene simultaneous exposure of microwave and toluene has higher micronucleus induction in compression with microwave and toluene alone which is in agreement with study that investigated the simultaneous effects of microwave and mitomycin on human blood cells in vitro it must be considered that really there are many chemical agents and an electromagnetic field which is genotoxic in our living and working environment, hence so, the synergistic effects of these chemical and physical agents can be considered as future investigations.

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