Evaluation of the mobile phone electromagnetic radiation on serum iron parameters in rats

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Abstract

Background: Electromagnetic fields (EMF) created by mobile phones during communication have harmful effects on different organs.

Objectives: It was aimed to investigate the effects of an EMF created by a mobile phone on serum iron level, ferritin, unsaturated iron binding capacity and total iron binding capacity within a rat experiment model.

Methods: A total of 32 male Wistar albino rats were randomly divided into the control, sham, mobile phone speech (2h/day) and stand by (12 h/day) groups. The speech and stand by groups were subjected to the EMF for a total of 10 weeks.

Results: No statistically significant difference was observed between the serum iron and ferritin values of the rats in the speech and stand by groups than the control and sham groups (p>0.05). The unsaturated iron binding capacity and total iron capacity values of the rats in the speech and stand by groups were significantly lower in comparison to the control group (p<0.01).

Conclusion: It was found that exposure to EMF created by mobile phones affected unsaturated iron binding capacity and total iron binding capacity negatively.

Keywords: cellular phone, electromagnetic fields, ferritin, serum iron

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Introduction

Mobile phones have become our time's most popular communication devices with the help of the addition of new functions. Mobile phones work based on the transmission of radio frequency signals and they create an electromagnetic field (EMF) around them while doing so¹. These devices have become widespread increased the concerns about the potentially harmful biological effects of the EMF they create. Previously, negative effects of EMF created by mobile phones were reported on hormones² and parameters such as hemoglobin, hematocrit and red blood cells³. Additionally it was reported that exposure to EMF reduced the levels of blood serum elements such as magnesium, iron and copper⁴.

Iron is the one of serum element which has an important role in the active sites of a wide range of proteins involved in energy metabolism, respiration, and DNA synthesis⁵. While 65% of the iron in humans is bound to hemoglobin responsible for oxygen-carrying, 10% of it is found in the structures of myoglobin, cytochrome and enzymes that contain ions. 20-30% of the iron in the body is non-toxicly stored as bound to ferritin and hemosiderin molecules. The transportation of the iron is achieved by the transferrin protein⁶. While transferring is a protein with high affinity and low capacity (each transferrin molecule is bound to two iron atoms), there are receptors on its surface for each cell that requires iron⁷. Unsaturated iron binding capacity (UIBC) is the measurement of the fraction of the serum transferrin which is not bound with

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iron and it indicates the amount of iron that can be saturated with the free transferrin in the blood. Total iron binding capacity (TIBC) indicates the maximum amount of iron essential to saturate serum transferrin.

There are few studies in the literature on the effects of particularly the EMF created by mobile phones on serum iron parameters. Previously it was reported the EMF created by mobile phones affect the serum ferritin levels negatively. In another study, it was observed that the serum iron and ferritin levels of people living close to high voltage electric cables which create EMF around were relatively lower. Further experimental studies are needed in this field in order to clearly reveal the relationship between EMF and serum iron parameters. In this study, it was aimed to investigate the effects of exposure to the EMF created by mobile phones on serum iron level, ferritin, UIBC and TIBC.

Methods

The experimental protocol was approved by Animal Experiments Local Ethics Committee decision numbered 11.2011/54-4 and dated 28.11.2011. All animals were kept in wire mesh cages for 12 hours at night and 12 hours in daytime, at circadian rhythms, 24-26 °C ambient temperature and 45-60% humidity. A total of thirty-two (n=32) male Wistar Albino rats were used in the study. These rats were 8-10 weeks old with 200 ± 20 g weight. The rats were divided into four groups as the Control (C; n=8), Sham (Sh; n=8), mobile phone Speech (Sp; n=8) and mobile phone Stand by (S-by; n=8). The EMFs were generated by the mobile phones using GSM mobile phone systems. The properties of mobile phones were; the highest SAR value was 0,96 W/kg and each of them had an 890-915 MHz carrier frequency band, 217 Hz modulation frequency, 250 mW maximum average power and 2 W maximum peak power.

Circular plastic experimental boxes were used in the Sh, Sp and S-by groups. One mobile phone was placed in the special sections located 1 cm below and in the middle of the experimental boxes (Figure 1). The rats in group Sh were placed in the box with their testes directed towards the mobile phone while the mobile phones were switched off. The rats were kept in this experimental box for 12 hours/day. The rats in the Sp and S-by groups were placed in experimental boxes in the same way as those in group Sh. Speech mode was obtained by the mobile phone in speech mode for 2 hours/day. The duration of standby mode was 12 hours/day. Experimental practices were carried out between 07:00-19:00 for groups Sh and S-by and 07:00-09:00 for the Sp group seven days a week. The rats in the C group were kept in their normal life cages in a separate room without being exposed to any type of electromagnetic field. The duration of the experiment was 10 weeks for all groups.

Evaluation of the serum iron parameters

At the end of the experiment (on the last day of 10th week) all the rats were anesthetized by Ketamine-Hydrochloride (50 mg/kg) and Xylazine-Hydrochloride (5 mg/kg) intraperitoneally. The anesthetization of rats in groups was made randomly. By thoracotomy, the thoracic cages of the rats were opened and blood samples were taken intracardially. Blood samples were drawn into common biochemical tubes and samples were centrifuged at 400 RPM for 10 minutes and blood serums were obtained. Serum samples were stored at -80 °C before the evaluations of serum iron parameters. The evaluated serum iron parameters were serum iron level, ferritin and UIBC. The total
iron binding capacity was calculated by the sum of serum iron level and UIBC\textsuperscript{14}. Serum iron parameters evaluated with electrochemiluminescence method on the Roche Cobas 8000 modular analyzer.

**Statistical analyses**

Kolmogorov Smirnov test was used to check whether the continuous variables fit normal distribution. In the comparisons of more than two groups of normally distributed variables, ANOVA and LSD multiple comparison tests were conducted whereas Kruskal Wallis and Dunn multiple comparison test were used for the variables that did not fit normal distribution. Differences between repeated measurements were tested using paired samples t test. Mean ± std. deviation values were given as the descriptive statistics. SPSS for Windows version 22.0 package program was used for statistical analyses and P<0.05 was accepted as the statistically significant value.

**Results**

Ferritin, serum iron level, UIBC and TIBC values of the rats in the Sp and S-by groups were compared to those of the rats in the Sh and C groups. It was observed that ferritin and serum iron levels did not change significantly in the S-by and Sp groups relative to the Sh and C groups (p>0.05). The statistically significant difference was seen UIBC values in the S-by and Sp groups than to the C group (p<0.01). It was observed that the TIBC levels of S-by and Sp groups decreased significantly in comparison to the C group (p<0.01) (Table 1).

**Table 1:**

The comparison of ferritin, serum iron level, UIBC and TIBC values in the experiment groups.

| Groups   | Ferritin (ng/mL) | Serum iron level (µg/dL) | UIBC (µg/dL) | TIBC (µg/dL) |
|----------|------------------|--------------------------|--------------|--------------|
| C (n:8)  | 0.666±0.078      | 223.3±46.4               | 539.4±176.5  | 762.6±202.4  |
| Sh (n:8) | 0.633±0.075      | 219.1±51.1               | 373.0±105.3  | 592.1±124.4  |
| Sp (n:8) | 0.589±0.072      | 195.4±22.2               | 350.0±36.0\textsuperscript{*} | 545.3±46.2\textsuperscript{*} |
| S-by (n:8) | 0.610±0.030   | 186.1±25.1               | 361.5±66.5\textsuperscript{*} | 547.7±67.8\textsuperscript{*} |

* C: Control group (n=8), Sh/Sham group (n=8), Sp: Speech group (n=8), S-by: Stand by group (n=8), UIBC: Unsaturated iron binding capacity, TIBC: Total iron binding capacity, (Mean±SD)

*\textsuperscript{p}<0.01: Compared to C group, \textsuperscript{**p}<0.05: Compared to C group.

**Discussion**

Deficiency or surplus of iron in the body leads to various diseases\textsuperscript{15}. Iron deficiency negatively affects the immune system\textsuperscript{16}, thyroid metabolism\textsuperscript{17} and cognitive functions\textsuperscript{18}. On the other hand, iron surplus is related to various pathological conditions such as cancer\textsuperscript{19}, neurodegenerative diseases\textsuperscript{20}, and damage of liver\textsuperscript{21} and testicles\textsuperscript{22}. Therefore, while keeping the serum iron levels and other iron parameters within a certain interval is important for body function, various complicated mechanisms take part in the establishment of this balance.

There are studies in the literature which show that EMF in different doses affect serum iron parameters. The implementation durations of EMF exposure and total experiment durations vary in these studies. Elferchichi et al.\textsuperscript{23} observed that a short 5-day exposure to a static magnetic field (SMF) of 128 mT negatively affected serum iron levels. It was also seen that exposure to the SMF of the same dosage for three times longer (15 days) decreased the serum iron levels\textsuperscript{24}. Fattahi et al.\textsuperscript{11}, in their in vitro study, found that an exposure of 30 minutes to EMFs created by mobile phones decreased the levels of serum ferritin. They explained this decrease with the oxidative stress occurring because of EMF.

In addition to the studies where EMF is applied for short durations, there are also other studies where applications are made for longer durations. It was shown that serum ferritin and iron values were affected negatively by long exposure to EMF\textsuperscript{12,25}. On the other hand, contrary to knowledge on the negative effects of EMF, there are also studies which showed that very low frequency EMF did not have an impact on serum iron levels\textsuperscript{4,26}. In this study, serum iron and ferritin levels in experiment groups had a tendency to decrease, while there were not significant differences found. The insignificance of the difference between experiment groups and the control group may have been caused by the compensation of iron by iron storage areas in the body such as liver, spleen and bone marrow.

Transferrin is primarily formed by the bonding of apo-transferrin synthesized in the liver with (Fe\textsuperscript{3+})\textsuperscript{27}. UIBC represents the fraction of the transferrin protein that is not bonded with iron. There are few studies in the
literature on the effects of EMF on transferrin. Djordjevich et al. found that the transferrin levels of rats exposed to SMF (16 mT) for 28 days were increased. Likewise, there are studies which reported that short (5 to 15 days) exposure to 128 mT SMF increased the serum transferrin levels, while these studies found a negative relationship between transferrin and serum iron levels. Contrary to previous results, in this study, it was observed that exposure to mobile phones working on EMF with 900MHz frequency, both in speech and stand by modes, significantly decreased UIBC which represents serum transferrin levels. It was seen that the difference between sham and control was also statistically significant. The limitations of the this study were that the parameters evaluated on the blood serum samples, the duration of experiment performed in a long period (10 week) and the occurring stress depending on the immobilization of rats for a long period. In connection to the significant decrease in UIBC, TIBC also decreased. The reason for the decrease in UIBC may be damage in the liver where transferrin is synthesized caused by the oxidative stress created by the EMF. Histopathologic investigation of the liver in further studies will be helpful in clearly revealing the activation mechanism between EMF and transferrin.

As a conclusion, mobile phone usage is becoming more and more widespread in the society. It is a necessity to investigate the potentially harmful effects of EMF created by mobile phones on different systems. In this study, it was determined that the EMF created by mobile phones on speech and standby did not affect serum iron and ferritin levels. On the other hand, it was observed that the EMF in both modes negatively affects UIBC. We suggest that these results will contribute to studies investigating the effects of EMF on hematologic parameters.

Conflicts of interest:
The authors of this paper have no conflicts of interest, including specific financial interests, relationships, and/or affiliations relevant to the subject matter or materials included.

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