THE EFFECTS OF VIBROACOUSTICALLY INDUCED MICROVIBRATIONS ON ARTERIAL BLOOD PRESSURE AND OXIDATIVE STRESS IN RATS

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ABSTRACT

Vibroacoustics, a scientific field that has been intensively studied for the last thirty years, uses the properties of sound waves (infrasound, ultrasound, noise and music) to induce vibrations that, like a sound wave, may have both useful and harmful effects. The aim of this study was to examine the effects of vibroacoustically induced microvibrations on arterial blood pressure and markers of oxidative stress in the blood. The experiments were performed on Wistar male rats that had a body mass of 180-200 g and were divided into control and experimental groups (6 rats in each). In the experimental group, microvibrations were induced using the Vitafon vibroacoustic apparatus (Vitafon, St. Petersburg, Russian Federation), which delivers sound waves of varying frequencies by a process called "phoning". Up to 60 minutes of phoning time was delivered to the kidney and liver using 4 different regimens that included a 5-minute stabilisation time; up to four 10-minute phoning regimens, with 5-minute breaks between each single regimen, at a 30 Hz-18000 kHz frequency range, and 2.8 μm-12.3 μm microwave amplitudes. After the completion of a phoning regimen, animals were sacrificed and the oxidative stress markers were measured in blood samples (O2 lipid peroxidation index, superoxide dismutase, catalase, and glutathione) and compared with the values of markers in the control group. Systolic arterial pressure was analysed after the acute application of up to four different regimens of vibroacoustic microvibrations. Systolic arterial pressure decreased significantly during the administration of the second regimen in comparison to the control group. Systolic arterial pressure returned, almost completely, to the initial value after the administration of the third and fourth regimens. There was no significant change in diastolic arterial pressure after the acute administration of up to four different regimens, although the pressure decreased slightly after the first and second regimens and returned to the initial value during the administration of the third and fourth regimens. Analysis of oxidative stress markers showed a statistically significant change in the catalase level. No statistically significant differences were found in the other oxidative stress harmful effects.
Rohracher found that the body surface of humans, or more generally homeotherms, produced constant vibrations, which he named microvibrations (1). The most significant observation of his intensive work is that the sources of these microvibrations are the heart rate and vascular and muscle activities, which produce microvibrations in the infrasound and sound range, respectively. Rohracher further showed that maintenance of microvibrations in an organism demands considerable muscle engagement, or energy consumption and that the amplitude of microvibrations is a sensitive psychophysical measure of muscle tension and total body activity. For example, in a healthy human (or other homeothermic animal), the amplitude of these microvibrations is 1-5 international units at rest, with a frequency range of 6-12 Hz/sec (vibrations per second). Rohracher demonstrated that it is possible to detect microvibrations that, originating in the striated muscle system, over the whole body; contractions of striated muscles have a manifold magnification of the amplitude of microvibrations but do not affect their frequency; and microvibrations are constant during the registration of frequency.

Several studies have been done performed that support Rohracher’s conclusions (2, 3, 4, 5, 6, 7). Based on these studies, the aim of this paper was to examine the acute effects of vibroacoustic microvibrations on arterial blood pressure and markers of oxidative stress in blood.

**MATERIAL AND METHODS**

**Experimental protocol**

The experiments were performed on Wistar albino rats, aged 8 weeks, with a body weight of 230-250 g. The rats were placed in experimental and control groups, with n=6 in each group, and. Experiments were carried out on each animal individually. All research procedures were carried out in accordance with the Declaration of Helsinki (last updated in 2005) and principles of Good Laboratory Practice (GLP) and were approved by the Ethical Committee for the Welfare of Experimental Animals, Faculty of Medical Sciences, University of Kragujevac. Baseline measurements were obtained for all of the parameters tested in rats without prior exposure to vibroacoustically induced microvibrations, which were used as controls. Rats were exposed to acute vibroacoustically induced microvibrations of defined amplitude and frequency using two vibroacoustic device emitters applied to the skin in the topographical area of the liver and kidney. The following settings on the vibroacoustic device (Vitaфон, St. Petersburg, Russian Federation) were used: regimen 1, lower frequency of the 1st subrange within the limits 30–60 Hz; regimen 2, upper frequency of the 1st subrange within the limits 1–3 kHz; regimen 3, lower frequency of the 2nd subrange within the limits 0.3–0.8 kHz; regimen 4, upper frequency of the 2nd subrange within the limits 9–18 kHz; number of microvibration frequency subranges, 2; length of a single cycle of microvibration frequency change, 80–160 sec; amplitude of microvibra-
tion at the lowest frequency for settings 1 and of 3, 2.8–5.4 μm and for settings 2 and 4, 6–12.3 μm; and period of impulse modulation, 0.5–1.2 sec. The duration of vibroacoustically induced microvibration stimulation was 60 min. divided into individual 10 min. regimens, with 5 min. breaks between each.

Haemodynamic measurements

All animals were anaesthetised (35 mg/kg sodium pentobarbital; i.p.). The mean arterial pressure (MAP) was determined directly through the femoral artery catheter (PE-50, Clay-Adams, Parsippany, NY, USA) using a low-volume displacement transducer (P23 Db, Statham, Oxnard, CA, USA) and was recorded on a direct writing recorder.

Measurement of oxidative stress parameters in rat blood

Rats were anesthetized with ether and sacrificed using cervical dislocation. For both control and experimental groups, n1/2=12. Blood was collected in tubes (12x100), with 50 I.J. heparin/ml of blood, and kept frozen at -20°C until used for biochemical measurements. The following parameters of redox status were determined spectrophotometrically from the blood samples: index of lipid peroxidation (measured as TBARS), SOD, CAT and GSH. The presence of thiobarbituric acid reactive substances (TBARS) was used to estimate the degree of lipid peroxidation in plasma by adding 1% thiobarbituric acid (TBA) in 0.05 M NaOH to an aliquot of plasma followed by a 15 min. incubation at 100°C and reading at 530 nm. Distilled water was used as a blank probe. A TBA extract was obtained by combining 0.8 mL of plasma and 0.4 mL of trichloroacetic acid (TCA), incubating the sample on ice for 10 minutes and centrifuging the sample for 15 min. at 6000 rpm, as described previously (8). To calculate the activity of endogenous antioxidants, haemoglobin was measured according to the Drabkin method (9). Isolated red blood cells (RBCs) were washed three times with 3 volumes of ice-cold 0.9 mmol/L NaCl and hemolysate containing approximately 50 g Hb/L prepared. Superoxide dismutase (SOD) activity was determined by the epinephrine method. A 100 μL sample of lysate was mixed with 1 mL of carbonate buffer followed by addition of 100 μL of epinephrine. Detection of SOD was performed at 470 nm. (10). Catalase (CAT) activity was determined according to Beutler (11). Lysates were diluted with distilled water (1:7 v/v) and treated with chloroform-ethanol (0.6:1 v/v) to remove haemoglobin. The sample (100 μL) was mixed with 50 μL of catalase buffer and 1 mL of 10 mM H2O2. Detection of CAT was performed at 360 nm. Distilled water was used as a blank probe. The level of reduced glutathione (GSH) was determined by the oxidation of GSH with 5, 5-dithiobis-6, 2-nitrobenzoic acid using the Beutler method (12). The concentration of oxidative stress parameters is expressed as nanomoles per millilitre of red blood cells (RBCs).

Statistical analysis

Statistical analysis of experimental data included the following basic descriptive statistics: the mean value (X), standard deviation (SD) and standard error mean (SEM). For testing the normality of distribution parameters, the Kolmogorov-Smirnov test was used. To test the statistical significance of the results and to confirm the hypothesis, the following statistical tests were used: Student’s t-test (parametric test), for dependent and independent variables and the Mann Whitney U test, for differences between the parameters. A database analysis of the results was performed using software package SPSS 10th 0 (SPSS Inc., Chicago, IL, USA). Ap < 0.05 was considered statistically significant.

RESULTS

Acute effects of vibroacoustic microvibrations of a specified amplitude and frequency on the arterial blood pressure of rats in vivo

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An analysis of systolic arterial pressure after the acute application of vibroacoustic microvibrations using different regimens over the rat’s body (5-minute single phonation time, with 5-minute breaks between single regimens) showed a statistically significant decrease of systolic arterial pressure during the administration of the second regimen compared to control (p < 0.05). A visible decrease in pressure was noted during the administration of the first regimen, but this was not statistically significant. Systolic arterial pressure returned to a nearly baseline level after the administration of the third and fourth regimens of vibroacoustic microvibration (Figure 1).

Diastolic arterial pressure did not decrease significantly after the acute administration of vibroacoustic microvibra-
tions using different regimens over the rat’s body (5-minute single phoning time, with 5-minute breaks between single regimens). There was a trend, however, towards decreased diastolic arterial pressure during the first and second regimens, which returned to nearly baseline levels during the administration of the third and fourth regimens (Figure 2).

Acute effects of vibroacoustic microvibrations on oxidative stress parameters in the rat blood

Analysis of oxidative stress parameters after the acute application of vibroacoustic microvibrations using 4 different regimens (5-minute single phoning time, with 5-minute breaks, total phoning time 60 minutes) showed a statistically significant increase in the levels of the antioxidant enzyme catalase (CAT). In contrast, TBARs, SOD and GSH levels decreased slightly however this was not statistically significant (Table 1).

DISCUSSION

Research into the effects of vibrations and sound on human physiology has been the focus of vibroacoustic science for several years because of its significance in the environment (i.e., the ecology of labour) and in medicine. This research is especially important given the knowledge of the potentially harmful effects of high intensity vibrations on both labourers who work with vibrating machines and devices and the cardiovascular and nervous systems, as well as the harmful effects of high intensity sound on the environment. On the other hand, there are many beneficial effects of vibrations and/or low frequency sound waves, as has been shown recently, with important research being done conducted to study the vibratory sensitivity of people and potential applications of infrasound, ultrasound, noise and music.

Based on his research, Olav Skille (13) pointed to three universal effects of sound and/or music induced vibrations on humans: a. low frequency vibrations can have a relaxing effect, while high frequencies can increase tension; b. “rhythmic” music can excite while non-rhythmic music can have a sedating effect; and c. loud music can lead to aggression, and quiet music can act to sedate.

Vibroacoustic devices, which produce sound-induced vibrations and stimulate humans, were developed over the last 30 years. Vibroacoustic therapy can be used in clinical settings. For example, music that causes sedation and/or pulse sinusoidal waves with low frequencies (between 20 Hz and 70 Hz) can be applied through a bed or armchair. Vibroacoustic therapy is currently used in the treatment of decubiti, to decrease arterial blood pressure, to reduce the heart rate, to improve circulation in post-operative treatment, and for stress-induced diseases. Reports on the improvement of circulatory beds in lower limbs and positive change of skin colour in patients treated with vibroacoustic therapy are anecdotal evidence of the benefits of this therapy (14). A hydrodynamic pump has been shown by Russian authors, to cause muscle fibres to tremble with sound oscillations, i.e., “a muscle sings at sound frequencies”. According to this concept, microvibrations are a physical agent that helps organisms by reducing peripheral resistance in capillary networks and increasing venous blood flow. It has been shown that the role of microvibrations in the pump-like functioning of vascular vessels of the venous and lymphatic systems leads to the unidirectional flow of both blood and lymph. The frequency of smooth muscle trembling in the vascular walls improves the efficiency of venous and lymphatic pumping as well as the amplitude of movement, i.e., oscillations of muscle tissue appears to align with the diameter of the lumen in venous and lymphatic vessels. By applying different regimens of a vibroacoustic stimulator, vibroacoustic waves of various shapes, frequencies, amplitudes and time length can be used to synchronise their energetic stimulation on vessels that can have many different diameters. Each blood or lymphatic vessel will have its own optimum frequency and characteristic energy wave based on its unique diameter. Another important characteristic is a reduction of resistance due to blood circulation. It is assumed that at certain frequencies, vibroacoustic microvibrations decrease the friction between blood layers, thus reducing viscosity and vascular resistance, leading to an increase of “shear stress”, which is the main physiological stimulus for the production of nitric oxide (NO).

As discussed, there are certain effects of low frequency sound applications on the human cardiovascular system. In a study performed on an Apollo mission, astronauts using infrasound treatment found no electrocardiographic disturbances when 21 male subjects aged 21 and 23 were stimulated by sounds ranging betweenfrom 2 Hz-12 Hz, with an intensity of 119-144 decibels, in thea simulation chamber. The heart rate increased in 6 subjects by more than 6 beats per minute during maximum stimulation, but in 5 subjects, the heart rate decreased (15). Respiratory function was evaluated by pneumographic impedance and was normal in all subjects exposed to low frequency stimu-
An acute application of vibroacoustic microvibrations in rats *in vivo* at low frequencies and amplitudes leads to a significant decrease in systolic arterial pressure (12%) and a trend towards decreased diastolic arterial pressure.

An acute application of vibroacoustic microvibrations in rats *in vivo* using 4 different regimens (total time 60 minutes, 5 minutes per single regime with 5-minute breaks after each) showed a slight decrease in certain markers of oxidative stress; however, this was not statistically significant. In contrast, the there was as statistically significant increase in the catalase (CAT) level.

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