Effects of long-term construction noise on health of adult female Wistar rats

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Abstract

The aim of this study was to investigate the influence of long-term building construction noise from refurbishment, which including vibration, on some physiological parameters and histopathological changes of organs of Wistar rats. Twenty 12 month old female rats were divided into two groups: rats group I (n = 10) were exposed to long-term construction noise and rats group II (n = 10) were kept under normal noise level. Study results revealed that long-term construction noise from building refurbishment has an influence on body weight, haematological and some serum biochemical parameters affects caecal microbiota, and causes histopathological changes in the organs of adult female Wistar rats. It was noticed that rats in group I exhibited significantly higher mean values for total protein, albumin and lower values for glucose, AST, ALT, blood urea nitrogen, haematological and caecal microbiota parameters than rats in group II. The most common pathologies were determined in the kidney, liver and lungs. Other observed pathologies were lymphadenopathy, catarrhal inflammation of the intestines, spleen hyperplasia and mammary gland adenofibroma. Single cases were subcutaneous fibroma in the thoracic region, abortus with uterine inflammation and thymus hyperplasia with formation of cysts were found.

Key words: Wistar rat, weight, blood, caecal microbiota, histopathology

Introduction

Human interaction and physical environmental factors are part of the stimuli presented to laboratory animals every day, influencing their behaviour and physiology and contributing to their welfare (Castelhano-Carlos and Baumans 2009). The concept of the five freedoms (freedom from thirst, hunger and malnutrition, freedom from discomfort, freedom from pain, injury and disease, freedom to express normal behavior and freedom from fear and distress) should be applied to achieve animal’s welfare.

Cage cleaning and design, bedding materials, nutrition, light intensity, noise, and veterinary care effect
laboratory rodent behaviours, physiology, health and welfare. The effects of sound on animal physiology and behavior depend not only on its intensity (or loudness), which is measured in decibels (dB), its frequency, which is measured in hertz (Hz), and its duration and pattern (including vibration potential), but also on the hearing ability of the animal species and strain, the age and physiological state of the animal at the time of exposure, to what sounds the animal has been exposed during its lifetime (noise exposure history of the animal) and to the predictability of the acoustic stimulus (Clough 1982, Gamble 1982, Voipio 1997, Turner et al. 2005, Burn 2008). Meaningful sounds at relatively low-intensity levels can have a considerable impact on animal physiology and behaviour by engaging limbic structures and higher centers involved in determining context and meaning (Turner et al. 2005).

Rodents have a different spectrum of audible sounds with maximum sensitivity at frequencies that are inaudible to humans. Humans can perceive frequencies from 20 Hz to 20 kHz, the frequencies from 400 Hz to 4.8 kHz being important for speech. Rodents not only produce sounds that we can hear, but also produce and hear frequencies that are inaudible to humans (above 20 kHz), perceiving sounds up to 80 kHz (Kelly and Masterton 1977, Bjork et al. 2000, Turner et al. 2005).

Exposure to high levels of noise can create physical and psychological stress and reduce productivity (Ng 2000). The negative impact of building construction noise from demolition and refurbishment on human health is well known (Suter 2002, Leensen et al. 2011). Most investigations have focused on human hearing, but the effects of sound do not stop with the ears. Non auditory effects of noise exposure are those effects that do not cause hearing loss but can still change the normal processes of other organ systems, such as the cardiovascular and digestive systems.

Regarding noise influences on laboratory animal physiology and behaviour, both auditory (hearing damage) and non auditory effects have been reported and reviewed in the literature (Voipio 1997, Burn 2008). Intense noise exposure can damage the cochlea and inner ear and lead to a cascade of auditory effects along the entire central auditory cascade (Turner et al. 2005). However, during a one year period of neighboring building refurbishment changes in rat behaviour (lethargy), coat appearance and litter size were noted. There is no information on low long-term construction noise exposure of *Wistar* rats may affect their health.

The aim of this study was to investigate the influence of long-term building construction noise from refurbishment, which includes vibration, on some physiological parameters and histopathological changes of organs of *Wistar* rats.

### Materials and Methods

#### Experiment conditions and experimental animals

*Wistar* strain rats were kept in two different premises (No. 1 and No. 2) under the same normal conditions: light-dark cycle of 12:12 hours, temperature maintained at 22 ± 2°C and relative humidity was 55 ± 5%. Rats were given *ad libitum* access to high-quality feed of the same batch and water throughout the whole study. The commercial standard diet (PA-11700000-171) for rodents contained crude protein 19.91%, fat 12.05%, and crude fiber 2.79% and 7.72% cellulose in 1 kg of feed. All procedures were approved by the local Animal Ethical Committee (No. license G2-16).

The rats were housed in standard polycarbonate cages (Techniplast, Italy), where minimum enclosure size was 800 cm², floor area per animal 350 cm² and minimum 18 cm height according to Directive 2010/63 EU of the European Parliament and of the Council of 22 September 2010 on the protection of animals used for scientific purposes.

The three-storey building under refurbishment was only 10 meters away from *Wistar* rat premises No. 1. Refurbishment works started early in the morning and continued until late afternoon five days a week and work lasted for about one year. Five days a week construction noise from electric drills, electric saws, grinders, cutting hammers and other devices ranged from 87 to 120 dB and the vibration level was $A(8) = 4m/s²$ for two years.

Immediately after the refurbishment works finished twenty randomly selected 12-months- old female rats were divided into two groups in accordance with their allocated premises. Rats group I (n = 10) were kept in premises No. which is located alongside refurbished building and rats of group II (n = 10) were held in premises No. 2 which is situated approximately 100 meters away from the construction site and the noise level did not exceed 60 dB at any time (control group).

#### Recordings of respiration and body weight

The respiratory rate was measured by calculating the frequency of respiratory movements per minute (cycles per minute, cpm) in a quiet but awake state (eyes opened, no motor activity). Body weight (g) was then measured using KERN PCB000-1 scales (Germany).
Blood sampling and analysis

Rats were killed using an overdose of carbon dioxide in a Quietek™ CO2 induction system chamber (USA). Trunk blood was collected in plastic tubes containing anticoagulant for haematological analysis and plastic serum tubes for biochemical analysis. Blood samples for haematological analyses were delivered to the laboratory within 1 h of collection and promptly assayed. Samples for biochemical analyses were centrifuged (4000 x g, 10 mm) in an EBA-200 centrifuge (Germany) and stored at -20°C until analysis.

The red blood cell count (RBC) was calculated using Neubauer’s counting chamber (Germany) under an OLYMPUS CX 22LED microscope (Germany). The white blood cell count (WBC) and haemoglobin content (Hb) were analyzed using a QBC Autoread Plus system (QBC Diagnostics Inc., USA).

Serum levels of total protein (T-Pro), albumin (Alb), glucose (Glu), triglycerides (TG), total cholesterol (T-Cholesterol), aspartate aminotransferase (AST), alanine aminotransferase (ALT), blood urea nitrogen (BUN) were analysed using a SPOTCHEM EZ SP-4430 automated biochemical analyser (Arkray Inc., Japan).

Intestinal content sampling and microbiological analysis

The caecum was removed aseptically from the abdominal cavity. The specimen was placed on a glass dish and cut open along its longitudinal incision. For microbiological enumeration, the intestinal contents were carefully collected without scraping the mucosa and delivered to the laboratory immediately after sampling.

Caecal contents (1 g) were homogenized in 9 ml saline peptone water by vortex mixing (IKA mini shaker MS2, USA). Bacteria growing and enumeration was using a standard method. Serial decimal dilutions carried out and as parallel triplicate cultures was inoculated onto Plate count agar (Liofilchem, Italy) for enumeration of aerobic and facultative anaerobic bacteria (LST EN ISO 4833:2003); de Man, Rogosa and Sharpe (MRS) agar (Biolife, Italy) for Lactobacillus spp. (LST ISO 15214:2009); Slanetz and Bartley agar+TTC (Liofilchem, Italy) for enterococci (ISO 7899-2:2000); Violet red bile glucose agar (Liofilchem, Italy) for Enterobacteriaceae (ISO 21528-2:2004).

The plates were incubated under aerobic conditions for growing of aerobic and facultative anaerobic bacteria at 30°C for 48 hours and for Enterobacteriaceae at 37°C for 24 hours. Plates for growing of mesophilic Lactobacillus spp. were incubated at 30°C in 2.5-liter jars (Oxoid) under microaerophilic conditions that were created using a CampyGen gas pack (Oxoid) for 72 hours.

The numbers of colony forming units were expressed as log_{10}CFU per gram of caecal content (log_{10}CFU/g).

Histopathological examination

During necropsy, after macroscopical evaluation, specimens of the following organs were collected: liver, pancreas, spleen, thymus, heart, lungs, stomach, colon, caecum, and kidney.

Tissue samples were fixed with 10% formalin solution. Paraffin blocks were made using Shandon Pathcentre (UK) and TES99 Medite Medizintechnik (Germany) equipment. Four micrometer sections were obtained using Shandon Pathcentre (UK) microscope. Sections were stained with hematoxylin-eosin (HE) using Sakura Accu-Cut SRM (Japan) equipment.

Statistical analysis

Statistical analysis of the results was carried out using the SPSS version 15 (licence No. 9900457; SPSS Inc., Chicago, IL). All data are presented as mean ± SEM (standard error of the mean). Differences in body weight, respiratory rate, haematological and biochemical parameters among groups were determined using Student’s paired t test. Differences in microbiological parameters among groups were compared by one-factor ANOVA. A p ≤ 0.05 was indicative of a statistically significant difference between groups.

Results

Body weight of Wistar rats group I, which were exposed to construction noise for one year, was less by 7.5% (p<0.05) in compare with rats group II, where the noise level was normal. Respiratory rate did not differ between the groups (p>0.05). The results of body weight and respiratory rate are shown in Table 1.

The results of haematological and biochemical parameters of group I and II rats from are presented in Table 2. The haematological parameters of rats exposed to long-term construction noise (group I) had significantly lower mean values than rats kept under a normal noise level (group II). The number of RBC
Table 1. Body weight and respiratory rate of Wistar rats exposed to different noise levels.

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Group I (n = 10)</th>
<th>Group II (n = 10)</th>
<th>p*</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>mean</td>
<td>SEM</td>
<td>mean</td>
</tr>
<tr>
<td>Body weight, g</td>
<td>265.97</td>
<td>2.13</td>
<td>287.52</td>
</tr>
<tr>
<td>Respiratory rate, cpm</td>
<td>58</td>
<td>1.64</td>
<td>60</td>
</tr>
</tbody>
</table>

SEM – standard error of the mean  
* Student’s paired t test

Table 2. Haematological and biochemical parameters of Wistar rats exposed to different noise levels.

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Group I (n = 10)</th>
<th>Group II (n = 10)</th>
<th>p*</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>mean</td>
<td>SEM</td>
<td>mean</td>
</tr>
<tr>
<td>RBC, × 10¹²/L</td>
<td>3.90</td>
<td>0.08</td>
<td>4.20</td>
</tr>
<tr>
<td>WBC, × 10⁹/L</td>
<td>5.60</td>
<td>0.13</td>
<td>8.10</td>
</tr>
<tr>
<td>Hb, g/L</td>
<td>111.00</td>
<td>0.11</td>
<td>117.00</td>
</tr>
<tr>
<td>T-Pro, g/L</td>
<td>76.20</td>
<td>2.31</td>
<td>55.90</td>
</tr>
<tr>
<td>Alb, g/L</td>
<td>41.20</td>
<td>1.67</td>
<td>36.00</td>
</tr>
<tr>
<td>Glu, mmol/L</td>
<td>3.31</td>
<td>0.74</td>
<td>7.52</td>
</tr>
<tr>
<td>TG, mmol/L</td>
<td>2.27</td>
<td>0.45</td>
<td>1.40</td>
</tr>
<tr>
<td>T-Cho, mmol/L</td>
<td>1.75</td>
<td>0.17</td>
<td>1.90</td>
</tr>
<tr>
<td>AST, IU/L</td>
<td>140.10</td>
<td>9.14</td>
<td>187.60</td>
</tr>
<tr>
<td>ALT, IU/L</td>
<td>19.60</td>
<td>2.34</td>
<td>69.10</td>
</tr>
<tr>
<td>BUN, mmol/L</td>
<td>6.20</td>
<td>0.29</td>
<td>8.56</td>
</tr>
</tbody>
</table>

* Student’s paired $t$ test

Table 3. Microbiological parameters of caecum content in Wistar rats exposed to different noise levels.

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Group I (n = 10)</th>
<th>Group II (n = 10)</th>
<th>p*</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>mean</td>
<td>SEM</td>
<td>mean</td>
</tr>
<tr>
<td>Total count aerobic and facultative anaerobic bacteria, log$_{10}$CFU/g</td>
<td>5.52</td>
<td>0.12</td>
<td>5.87</td>
</tr>
<tr>
<td>Mesophilic Lactobacillus spp., log$_{10}$CFU/g</td>
<td>4.67</td>
<td>0.16</td>
<td>5.26</td>
</tr>
<tr>
<td>Enterobacteriaceae, log$_{10}$CFU/g</td>
<td>3.75</td>
<td>0.11</td>
<td>4.11</td>
</tr>
<tr>
<td>Enterococci, log$_{10}$CFU/g</td>
<td>4.03</td>
<td>0.29</td>
<td>4.63</td>
</tr>
</tbody>
</table>

SEM – standard error of the mean, CFU – colony forming units  
* one-factor ANOVA

in blood of rats group II was higher by 7.7% (p<0.05), WBC by 44.6% (p<0.05) and the concentration of Hb by 5.4% (p<0.05) in compare to the rats group I.
As seen in Table 2, the group I rats exhibited significantly higher mean values for T-Pro, Alb and significantly lower values for Glu, AST, ALT and BUN than rats group II. The mean of serum T-Pro and Alb in rats exposed to excessive noise (group I) were higher by 36.31% (p<0.001) and 14.44% (p<0.05), respectively, compared to the rats group II. The mean of Glu content in blood of rats group I was lower by 55.9%, BUN by 27.57%, AST by 25.3% and ALT by 71.64% than rats kept under normal noise level (group II). Differences in mean values of TG and T-Cho concentration in serum between groups were not observed.
Some variation of caecal microbiota are shown in Table 3. The total count of aerobic and facultative anaerobic bacteria, Lactobacillus spp. and enterococci in caecal content of rats exposed to long-term noise were statistically significantly lower by 5.96%
During histopathological examination the most common pathologies were determined in the kidney, liver, and lungs. They are presented in Fig. 1.

In addition to these pathologies common pathologies were lymphadenopathy, catarrhal inflammation of the intestines (Fig. 1), spleen hyperplasia (Fig. 2) and mammary gland adenofibroma (Fig. 3). Only one case was of subcutaneous fibroma in the thoracic region (Fig. 4), one of uterine inflammation (Fig. 5), and one case of thymus hyperplasia and cyst formation (Fig. 6) were determined in histopathological examination.

In Table 4 the data show that only one rat did not have any pathology. One rat had only one pathology, swelling of the liver. Other rats had different pathologies (from 5 to 9).

The most common rat lung pathologies were pneumonia and emphysema (each 36.4%), congestion (18.2%), and edema (9.1%).

Histologically (Fig. 7) there were damaging of lung tissue, infiltration of neutrophils and macrophages and, as a consequence of inflammation we determined bronchiectasis (Fig. 8), cholesterol crystals (Fig. 9), granulation tissue formation (Fig. 10) and emphysema (Fig. 11).

Frequently, rat kidney pathologies were vacuolation of podocytes (58.3%) and kidney swelling (p<0.05), 11.22% (p<0.01) and 12.96% (p<0.05), respectively, compared to the rats kept under normal noise level.
Fig 1. Enlarged white pulp (lymphoid follicles) in rat spleen (arrows). HE, x40.

Fig 2. Small intestine mucosa -infiltration of lymphocytes, plasmocytes and eosinophils. Edema of intestinal villi. HE, x40.

Fig 3. Adenofibroma of mammary gland of rat. Abundant connective tissue (thin arrow) and glandular part (thick arrow) with milk stones, HE, x40.

Fig 4. Subdermal fibroma in rat – at left HE, x40; at right – gross view.

Fig 5. Purulent inflammation of rat uterus (endometritis, miometritis and perimetritis) with fetus putrification, HE, x40.

Fig 6. Hyperplasia and cysts (arrow) formation in rat thymus. HE, x200.
Fig. 7. Purulent exudate in rat bronchiole: abundant neutrophils (upper arrow) and macrophages (lower arrows) are visible. HE, x200.

Fig. 8. Bronchiectasis in rat lung. Bronchiol diameter is enlarged, filled with mucoid – purulent exudate. Lung tissue around bronchiole is atelectatic and hyperemic. HE, x200.

Fig. 9. Cholesterol crystal (thin arrow) formation in rat lung between macrophages (thick arrow). HE, x200.

Fig. 10. Granulation tissue formation in periphery of abscess. x200.

Fig. 11. Emphysema in rat lung. HE, x40.

Fig. 12. Vacuolization of podocytes in rat kidney (black arrow). HE, x200.
Histopathological lesions are shown in Fig. 12.

The main changes in the liver were edema (66.7%) and lipidosis (25%). Histopathological changes are presented in Fig. 13.

Discussion

Many laboratories have found that nearby construction or building maintenance has produced a downturn in breeding success or noticeable disturbance of their animals. Chronic sources of sound in the laboratory may include ventilation, building noise, animal noise, and human noise (Patterson-Kane and Farnworth 2006). Long-term or short-term exposure to sound pollution disrupts many physiological functions in different systems of the body in humans and animals (Stansfeld and Matheson 2003, Kivimäki et al. 2006, Yildirim et al. 2007).

The different kinds of stress can cause reduction of body weight in experimental animals (Gamaro et al. 2003, Bekris et al. 2005, Zardooz et al. 2006, Lucca et al. 2009). In the present study, the weight of Wistar rats exposed to construction noise for one year was less in comparison to the rats kept under normal noise level. This could be related to insufficient digestion and absorption of nutrients because of inflammatory processes in the intestinal mucosa found during histopathological examination.

Several studies reported higher haemoglobin and RBC values (Ola-Davies et al. 2014, Vinodini et al. 2015), and lower WBC values (Osonuga et al. 2012) in rats of Wistar strain than we found. Ashaolu et al. (2011) determined similar results to ours for RBC counts in the blood of Wistar rats. The present study demonstrated that the WBC count decreased in rats exposed to excessive noise (group I) compared to the rats in the control (group II). Erken et al. (2008) claimed that a 95-dB machine sound changes the haematological parameters of rats such as red blood cell deformability and aggregation. Moreover, long-term noise exposure could lead to an increase in the number of red blood cells, and serum concentration of haemoglobin (Monsefi et al. 2006, Wankhar et al. 2014).

It was noted that total protein content and albumin value of blood were higher, but blood urea nitrogen was lower in group I compared to the group II. These results are in line with Kadi et al. (2014); however, several studies detected a higher content of total protein (Ola-Davies et al. 2014, Vinodini et al. 2015). The increased protein and albumin levels indicate impairment in the normal function of the liver, as recently revealed (Imafidon and Okunrobo 2012); it seems that long-term construction noise exposure changes the normal functions of the rat liver.

The enzymatic activity of ALT and AST were studied to evaluate liver malfunctions. Liver enzyme levels are usually raised in acute hepatotoxicity, but tend to decrease with prolonged intoxication due to damage to the liver (Obi et al. 2004). In the present study AST enzyme activity of blood was three times higher compared to the results of Ola-Davies et al. (2014). Moreover, similar results of ALT activity were reported by Kadi et al. (2014) and Ola-Davies et al. (2014).

Rahma (2011) reported that noise of 80-100 dBA caused a significant increase in plasma glucose of rats but the cholesterol and triglycerides have no significant difference compared to control. Rostamkhani et al. (2012) observed a non-significant reduction of plasma triglyceride, and unchanged concentration of glucose and plasma cholesterol following chronic stress exposure compared to the control. No changes in the serum level of glucose in rats exposed to chronic noise were noticed by Armario et al. (1985). We found that rats exposed to long-term construction noise had a significantly lower content of Glu compared to the rats kept under normal noise level. However, no differences in the serum concentration of TG and T-Chol were noticed. If the stress continues for a long period it may be possible that the stress response of the animals become “distress”, in which apparent precipitation or exacerbation of metabolic impairments may occur (Dhabhar and McEwen 1997).

In our study we observed pathological changes in the rat intestine and lungs. Mainly inflammation was noted. In association with this there were changes in immune system organs such as lymph nodes, but less in the thymus. This may be related to disbacteriosis due to stress. The changes in biological function occurring during a stress response may suppress immune competence, rendering the animal susceptible
to pathogens that may be present in the environment (Moberg, 2000). All experiments of Baldwin (2007) demonstrated that 90 dB white noise reduced stimulation of the parasympathetic nervous system, and also induced an inflammatory response in the intestinal mucosa, resulting in structural damage.

Kleessen et al. (1999) noticed that rats are sensitive to multifactorial disorders. Microbial dysbiosis in the caecum leads to changes in the digestive system which could promote the development of potentially pathogenic microflora, which could cause digestive troubles, which might have similar effects on the health status: intestinal health, caecal characteristics and microflora population. Rats are coprophagic and ingest a large number of bacteria along the whole gastrointestinal tract. The large intestine contains a complex bacteria flora, with the principal groups being the Enterobacteria, enterococci, lactobacilli, bacteroides species and bifidobacteria. The strictly anaerobic bacteria (bacteroides, bifidobacteria), which outnumber the aerobes (enterobacteria, enterococci) by a factor 100 to 1000, are present in large numbers in rats. In the intestine the most numerous organisms are lactobacilli, bacteroides species and bifidobacteria (10^9-10^10 CFU/ml which outnumber the enterobacteria by a factor of more than 100). Stress causes hypothalamic-pituitary-adrenal (HPA) axis activation and gut microbiota dysbiosis (Galley and Bailey, 2014). The activation of the HPA axis produces changes in certain populations of bacteria (Pullinger et al. 2010).

Fewer changes were determined in the reproductive system. Only one rat had abortus with putrificated fetuses and inflammation of the uterus. Stress can suppress reproduction. Nakamura et al. (1996) noticed that mechanically induced whole-body vibration of 9-11 day gestation female Wistar rats resulted in decreased uterine blood flow, increases in plasma corticosterone, and decreases in plasma progesterone and prostaglandin E2, but no changes were found in plasma estradiol and prostaglandin F- compared to control rats. In contrast our studied rats were 18 months old, and according Hung et al. (2003) reproductive aging in female rats results from both aging and estrogen exposure, and may be rooted in the oxidative stress theory of aging.

Our study of rats showed different levels of degeneration (swelling, lipoidosis) in the liver and kidney. These pathologies can be in association with inflammation processes in other organs and while the cells of the hepatic parenchyma have the flexibility to adapt to changing physiological demands, with reversible functional and morphological alterations, sufficient stress or noxious stimuli may lead to inability to maintain homeostasis and adverse cellular adaptations. The morphological response to injurious stimuli depends on the nature of the injury and its severity and duration (Thoolen et al. 2010). Helal et al. (2011) detected that noise exposure showed some vacuolated hepatocytes with enlarged nuclei of the endothelial lining compared with the control group. Xue et al. (2014) indicated that noise could lead to pathological changes in the liver because the exposure to noise increased the level of oxygen free radicals and lipid peroxidation. They consider that the possible mechanism was that the rats were in a state of stress when they were exposed to noise, and therefore under this condition the oxygen free radicals in the body and malondialdehyde (a product of oxygen free radicals) content would be higher than normal.

We diagnosed spontaneous cases of neoplasms in rat subcutis (fibroma) and mammary gland (adenofibroma). In many studies, a major cause of death of rats is neoplasms. In some rat studies, benign and malignant cutaneous and subcutaneous neoplasms are a common cause of death (Snyder et al. 2016). Fibromas are the most common tumor of the subcutis at about 5% incidence in male Fischer 344 and Wistar rats (Poteracki and Walsh 1998, Suckow et al. 2006). According to Suckow et al. (2006) primary tumors of the mammary gland are some of the most common tumors in rats occurring in up to a third of females by 2 years of age. In our study we found mammary gland tumors by 18 months of age. For Wistar rats and Fischer 344, the majority of tumours are fibroadenomas. Boorman et al. (1990), Poteracki and Walsh (1998) reported that fibroadenomas were the most common mammary tumor (43% of all integumentary tumors, 78% of all mammary tumors).

**Conclusion**

From the results of this study it can be concluded that long-term construction noise from building refurbishment has an influence on body weight, haematological and some serum biochemical parameters, caecum microbiota, and histopathological changes in the organs of adult female Wistar rats. The most common pathologies where determined in the kidney, liver and lungs. Other pathologies were lymphadenopathy, catarrhal inflammation of the intestines, spleen hyperplasia and mammary gland adenofibroma. There were single cases of subcutaneous fibroma in the thoracic region, abortus with uterine inflammation, and thymus hyperplasia with cyst formation.
References


