INTRODUCTION

Ofloxacin is a quinolone that has shown to be an efficient bactericidal drug for the treatment against Gram-positive and Gram-negative microorganisms, as well as against anaerobic organisms (Hopkins et al., 2001; Meiland et al., 2001).

Ofloxacin acts on infections of the respiratory and urinary tract, pelvic inflammatory diseases, skin infections, osteomyelitis, colitis, and on sexually transmitted diseases that have chlamydia as an etiologic agent (Neu, 1992, 1997; Paschoal et al., 2003). In cases of genitourinary affections, it is used in the dose of 400mg/day, from 10 to 14 days, helping to avoid complications in future pregnancies and preventing reinfections. Researches with quinolones have revealed toxicity in rat and rabbit fetuses, whose dams received high doses of quinolones (810mg/kg in rats and 50mg/kg in rabbits), by relating decrease in body weight gain, difficulties in bone growth, and arthropathy in young animals. However, such effects haven’t been verified in rats who were administered doses of up to 360mg/kg (Watanabe et al., 1992; Linseman et al., 1995).

In pregnant women who used ciprofloxacin (400 mg/day), perfloxacin (800 mg/day), and ofloxacin (800mg/day), the drugs crossed the placenta, entered breast milk presenting considerably high levels (Giamarellou et al., 1989).

On the other hand, studies demonstrate that, because of their toxic effect, the use of ciprofloxacin, perfloxacin, and ofloxacin in young animals (rats and rabbits) can lead to

**SUMMARY:** Ofloxacin presents an ample spectrum of antimicrobial action, including combating *Mycobacterium leprae*, and is currently employed as a substitute when the use of rifampicin is impossible. The objective of this work was to study alterations in testicular cell nuclei of suckling rats, whose dams were submitted to oral application of ofloxacin, and respective control groups. The method utilized was morphometry by the karyometric technique. The main structures observed in histological preparations of the testicles were interstitial cells, spermatogonias, and sustentacular cells. 10 Wistar rats were utilized, four treated and five controls, in the period of the first 25 days of life, whose dams received ofloxacin 12 mg/ Kg of body weight / day orally, before being killed on the 25th day after birth. The karyometric study of interstitial cells and spermatogonias revealed that there were no changes in the form of their nuclei (p > 0.05). Since sustentacular cell nuclei presented increased major diameter, minor diameter, mean geometric diameter, volume, area, volume/area ratio and perimeter, as well as an augmented and statistically different eccentricity (p < 0.05) in suckling pups whose dams were administered ofloxacin, the nuclei presented larger size and more elongated form. It was concluded that the sustentacular cells were more sensitive to the ofloxacin effect at the administered dose.

**KEY WORDS:** Quinolones; Ofloxacin; Morphology; Karyometry; Nuclear volume, Testicle; Lactation; Sustentacular cells; Rat.

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**KEY WORDS:** Quinolones; Ofloxacin; Morphology; Karyometry; Nuclear volume, Testicle; Lactation; Sustentacular cells; Rat.
malformation, development delay with knee arthropathy, tenosynovitis, and tendon rupture. They also demonstrate that dogs have high-sensitivity to the drug (Paschoal et al.).

Some authors observed both functional testicular impairment, such as decrease in sperm count and motility, and structural testicular impairment, such as cell lesions in rats (Paschoal et al.; Abd-Allah et al., 2000; Paschoal, 2002). Though quinolones don’t provoke gene mutations (Mayer & Bruch, 1986), recent studies reveal that the use of this antibiotics during pregnancy in young females must be thoroughly researched (Stahlmann & Lode, 1999).

The aim of this study was to quantify, through karyometry, possible nuclear alterations in testicular cells of suckling rats, whose dams received doses of ofloxacin.

MATERIAL AND METHOD

After the approval by the Animal Experimentation Ethics Committee at FAMERP – Faculdade de Medicina de São José do Rio Preto – SP – Brazil, four rats (Rattus norvegicus, of the Wistar strain) were used, as well as five control animals, born after the mating of six virgin females and two males of the same pedigree.

The drug used was ofloxacin in the dose of 12mg/kg body weight, via a gastric tube. We opted for the use of the therapeutic dose, considering that the administered dose to treat an human adult (of up to 70 kg) is of 800mg/day, from 7 to 14 consecutive days.

From the first to the 25th day after birth, during lactation (pre-puberty period for the animal), 12mg/kg body weight of ofloxacin solved in 4ml of distilled water were administered directly in the stomach of the dams with the aid of a gastric tube. The same quantity of saline solution was given to dams of the animals selected for the control group. Around the 25th day after birth, five control males, and other four, whose mothers were treated, were chosen at random and, then, sacrificed by means of inhalation of sulfuric ether.

Histological technique. All the animals involved in this experiment were immersed in ALFAC fixing solution for 24 hours. After that, they were dried and weighed on a Mettler H64 balance, and immersed in alcohol 80%. Next, the testicles were removed through an inguinal incision and, then, fixed, cut transversely, dehydrated, cleaned, and embedded in paraffin. The paraffin inclusions were sectioned at 5 µm thickness. Nine sections of each block were selected and stained with hematoxilin-eosin.

The main structures observed were tunica albuginea, interstitial cells, germinative epithelium, sustentacular cells, and spermatogonias.

Morphometric Technique: Karyometry. Images from the nuclei of testicular cells were projected on white paper with a final magnification of 1.240x using a H500-Hund Wetzlar optical microscope with immersion objective and Leitz Wetzlar camera lucida. Fifty nuclear images of each of the three types of studied cells were considered, totaling the analysis of 150 nuclei for each animal. The camera lucida projected images that were, then, contoured with black pencil and had their major diameter (D) and minor diameter (d) calculated. From those diameters, the following nuclear parameters were estimated according to the formulas (Sala et al., 1981):

<table>
<thead>
<tr>
<th>Variables</th>
<th>Formulas</th>
</tr>
</thead>
<tbody>
<tr>
<td>Major diameter (µm)</td>
<td>M = (D•d) ½</td>
</tr>
<tr>
<td>Perimeter (µm)</td>
<td>P = (π 1/2) • [3/2 • (D + d) - M]</td>
</tr>
<tr>
<td>Mean diameter ratio</td>
<td>R = D / d</td>
</tr>
<tr>
<td>Volume (µm³)</td>
<td>V = π • 1/6 • M³</td>
</tr>
<tr>
<td>Area (µm²)</td>
<td>A = π • 1/4 • M²</td>
</tr>
<tr>
<td>Volume/area ratio</td>
<td>V/A = 2/3 • M</td>
</tr>
<tr>
<td>Shape coefficient</td>
<td>S = 4 • π • A • 1/P ²</td>
</tr>
<tr>
<td>Contour index</td>
<td>I = P / A ½</td>
</tr>
<tr>
<td>Eccentricity</td>
<td>E =[(D + d) ½ • (D - d) ½ ] • 1/D</td>
</tr>
</tbody>
</table>

Statistical analysis. To analyse the parameters obtained from karyometry, a software described by Maia Campos and Sala (Department of Stomatology, Faculdade de Odontologia de Ribeirão Preto – SP - Brazil, USP) was utilized. The Mann-Whitney test was used for the statistical analysis, with a 95% confidence interval.

RESULTS

Target organ. The testicles were intra-abdominal, similar, with blood vessels and tunica albuginea presenting normal characteristics and numerous spermatogenic cords. They still didn’t present lumen and had interstitial cells, sustentacular cells, spermatogonias and primary spermatocytes (undergoing cell division).

Morphometric results – Karyometry. The following results, presented in Tables I, II, III, refer to the data obtained through karyometry of testicular cells nuclei (interstitial, sustentacular and spermatogonias) from suckling rats, whose dams were submitted to ofloxacin.
Interstitial cells (Leydig). In Table I, the data distribution were similar for all the variables from both studied groups; thus, there was no significant statistical difference (p>0.05) when the Mann-Whitney test was used.

Spermatogonias. In Table II, when the values from the studied groups were distributed, they showed to be very similar for every analysed variable. The Mann-Whitney test did not present differences in data distribution (p>0.05).

Sustentacular cells (Sertoli). Table III presents the median values of the major diameter of Sertoli cells nuclei equal to 13.02 µm for the control group, and 14.62 µm for the treated group, as well as the median values of the nuclei minor diameter, 11.02 µm for the control group and 12.29 µm for the treated group. The median values of the mean nuclear diameter in 11.98 µm for the control group and 13.37 µm for the treated group. The index for the control group nuclear volume medians was of 887.77 µm³, and of 1182.00 µm³ for the treated group.

Table I. Values from the kariometry of interstitial cells nuclei taken from suckling pups whose dams were submitted to application of ofloxacin (T), and control groups (C).

<table>
<thead>
<tr>
<th>Variables</th>
<th>Control group n = 250</th>
<th>Treated group n = 200</th>
</tr>
</thead>
<tbody>
<tr>
<td>Major diameter (µm)</td>
<td>13.35 0.54 13.60</td>
<td>14.33 0.98 14.25</td>
</tr>
<tr>
<td>Minor diameter (µm)</td>
<td>8.87 0.37 8.80</td>
<td>9.47 0.31 9.51</td>
</tr>
<tr>
<td>Mean geometric diameter</td>
<td>10.82 0.41 10.74</td>
<td>11.56 0.53 11.45</td>
</tr>
<tr>
<td>Major/Minor</td>
<td>1.55 0.05 1.54</td>
<td>1.58 0.07 1.60</td>
</tr>
<tr>
<td>Volume (µm³)</td>
<td>589.06 68.07 599.16</td>
<td>715.45 750.16 704.03</td>
</tr>
<tr>
<td>Area (µm²)</td>
<td>93.87 7.40 93.02</td>
<td>107.50 9.37 105.18</td>
</tr>
<tr>
<td>Volume/ area</td>
<td>5.91 0.24 5.87</td>
<td>6.31 0.20 6.34</td>
</tr>
<tr>
<td>Perimeter</td>
<td>38.44 1.49 38.91</td>
<td>41.20 2.48 40.91</td>
</tr>
<tr>
<td>Eccentricity</td>
<td>0.70 0.01 0.70</td>
<td>0.70 0.03 0.71</td>
</tr>
<tr>
<td>Contour index</td>
<td>4.01 0.04 4.01</td>
<td>4.04 0.06 4.05</td>
</tr>
<tr>
<td>Shape coefficient</td>
<td>0.79 0.01 0.79</td>
<td>0.78 0.02 0.78</td>
</tr>
</tbody>
</table>

Table II. Values from the kariometry of spermatogonias nuclei taken from suckling pups whose dams were submitted to application of ofloxacin, and control groups.

<table>
<thead>
<tr>
<th>Variables</th>
<th>Control group n = 250</th>
<th>Treated group n = 200</th>
</tr>
</thead>
<tbody>
<tr>
<td>Major diameter (µm)</td>
<td>11.20 0.58 11.40</td>
<td>12.83 0.72 13.41</td>
</tr>
<tr>
<td>Minor diameter (µm)</td>
<td>9.11 1.33 9.32</td>
<td>10.51 1.13 11.00</td>
</tr>
<tr>
<td>Mean geometric diameter</td>
<td>10.08 1.32 10.33</td>
<td>11.59 1.25 12.20</td>
</tr>
<tr>
<td>Major/Minor</td>
<td>1.25 0.05 1.26</td>
<td>1.23 0.03 1.23</td>
</tr>
<tr>
<td>Volume (µm³)</td>
<td>553.81 197.45 544.66</td>
<td>796.62 223.89 903.42</td>
</tr>
<tr>
<td>Area (µm²)</td>
<td>82.31 20.49 84.96</td>
<td>108.24 21.91 118.42</td>
</tr>
<tr>
<td>Volume/area</td>
<td>6.07 0.88 6.21</td>
<td>7.00 7.75 7.35</td>
</tr>
<tr>
<td>Perimeter</td>
<td>33.56 4.09 34.37</td>
<td>38.48 2.26 40.33</td>
</tr>
<tr>
<td>Eccentricity</td>
<td>0.56 0.03 0.56</td>
<td>0.55 0.02 0.56</td>
</tr>
<tr>
<td>Contour index</td>
<td>3.76 0.04 3.77</td>
<td>2.90 1.68 3.72</td>
</tr>
<tr>
<td>Shape coefficient</td>
<td>0.88 0.02 0.89</td>
<td>0.89 0.01 0.89</td>
</tr>
</tbody>
</table>

* Mann-Whitney Test; ns: non-significant at a 5% level.
The nuclear area was estimated in 114.89 µm² and in 147.68 µm² for the group undergoing treatment. The median for the volume/area ratio was 7.35 µm for the control and 8.19 µm for the treated. The median values for the perimeters data of sustentacular cells nuclei was of 39.33 µm, for the control group, and of 44.01 µm for the treated group. For the eccentricity, the indexes were 0.47 and 0.51, respectively for the control and treated groups. The difference in data distribution from one group to another is significant, indicating an increase in all the analysed variables of the treated group (p < 0.05). The median values of the difference between major and minor nuclear diameters of sustentacular cells were of 1.16 mm for the control group and 1.19 mm for the treated group. The values for the nuclei contour index were of 3.68, control, and 3.41, treatment; and for the shape coefficient, the values were 0.93 and 0.91 with p > 0.05.

### DISCUSSION

**Lactation, toxicity and rat development.** Lactation is a very important period for the growth and development of mammals and one of its main consequences is the reduction in mortality rates. For rats, these two first weeks of life are, as well, crucial (Lopes et al., 1974). An obstacle to the suckling behavior of pups is the use of chemical substances. In general, it is possible to say that most drugs ingested by the dam pass through the milk, generating minimal or potentially harmful effects (Zucolotto & Marino, 1991).

Because of its own peculiarities, newborn rat pups require constant precautions and vigilance when using drugs. During this special period, when the pup is independent of the dam’s metabolism, adaptation to a different environment depends on its own ability to absorb, conjugate, inactivate and excrete substances. The ingestion of chemical elements by the dam during prenatal and suckling periods, along with the factors previously cited, can considerably prolong the half-life of the drug in young organisms (King, 1991).

The 15th day of postnatal life, for the rat, represents a critical moment in testicular development (Rugh, 1968). Studies about drug effects during this period demonstrated alterations indicating toxicity (Azoubel, 1964, Lopes, 1973; Souza et al., 1975; Kempinas et al., 1985).

Despite the capacity of ofloxacin to reach the newborn animal, through the ingestion of dam’s milk, this research has only presented alterations at the cellular level. The karyometric study of Interstitial cells, whose rats were fed by dams exposed to ofloxacin, did not reveal changes in shape and volume. Every variable presented similar data (p > 0.05) whenever the treated group and the control group were compared, demonstrating that drug, despite passing from the dam to the pup through suckling, did not cause toxicity, in the utilized dose.
Insensitivity to the drug may have occurred because interstitial cells were mature. The previous period to this phase was of constant transformation, but, around the 21st day of life, the establishment of interstitial cell populations becomes more mature or differentiated.

Karyometry of spermatogonias from the pups whose dams were treated, revealed similar data between the treated and control groups, (p > 0.05), for every studied parameter, demonstrating that ofloxacin, when administrated to the suckling dam, in the right dosage, does not cause toxicity to these cell nuclei.

On the other hand, different data were found for the karyometric study of sustentacular cells. The major diameter, minor diameter, mean geometric diameter, volume, area, volume/area ratio, and perimeter were augmented (p < 0.05). As for eccentricity, it was significantly augmented and statistically different (p < 0.05) in suckling pups whose dams were administered ofloxacin. Because of the increased major and minor diameters, such alterations resulted in nuclei of increased sizes and more elongated shapes.

The results found in this study suggest the following conclusions: Ofloxacin, during lactation period expands the volume of sustentacular cells nuclei, increasing major diameter by the minor diameter, the area, and the volume; and changes the shape, making eccentricity unequal and diminishing the perimeter.

**Key Words:** Quinolonas; Ofloxacin; Morfología; Cáncer de Piel; Células Sustentaculares; Rata.

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