Effect of hyperbaric oxygen therapy in rats with subtotal splenectomy preserving the inferior pole

Efeito da oxigenoterapia hiperbárica em ratos submetidos à esplenectomia subtotal com preservação do polo inferior

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ABSTRACT

\textbf{Purpose:} To evaluate the effect of hyperbaric oxygen therapy on the survival and weight of rats submitted to subtotal splenectomy and on the viability and growth of the inferior pole.

\textbf{Methods:} Forty Wistar rats underwent subtotal splenectomy with preservation of the inferior pole and distributed into two groups: Group A (n=20) - not treated with hyperbaric oxygen, Group B (n=20) – treated with hyperbaric oxygen. These groups were divided into two subgroups of 10 animals each, according to the time of euthanasia, 15\textsuperscript{th} and 45\textsuperscript{th} days. The survival and weight of the animals were recorded. The inferior pole was measured, weighed and morphologically analyzed.

\textbf{Results:} All animals survived. The weight of the animals increased in all subgroups, but decreased on the 10\textsuperscript{th} day in the subgroups treated with hyperbaric oxygen (p<0.001). The viability of the inferior pole was more evident in animals treated on the 15\textsuperscript{th} day, but did not differ on the 45\textsuperscript{th} day. The growth of the inferior pole has not occurred on the 15\textsuperscript{th} and sim on the 45\textsuperscript{th} day post-operatório, nos animais não tratados (p<0,01) e tratados (p<0,05). Vascular and cellular increase in treated animals was significantly higher than in untreated ones.

\textbf{Conclusion:} Hyperbaric oxygen therapy did not affect the survival of animals but reduced their weight. It improved the viability of the inferior splenic pole, but did not interfere with their growth.

\textbf{Key words:} Splenectomy/adverse effects. Splenectomy/methods. Hyperbaric Oxygenation. Animal Experimentation.

RESUMO

\textbf{Objetivo:} Avaliar o efeito da oxigenoterapia hiperbárica na sobrevida e peso de ratos submetidos à esplenectomia subtotal e na viabilidade e crescimento do polo inferior.

\textbf{Métodos:} Quarenta ratos \textit{Wistar} foram submetidos às esplenectomias subtotal com preservação do polo inferior e distribuídos em dois grupos: A (n=20) - não tratados com oxigênio hiperbárico, B (n=20) - tratados. Esses grupos foram divididos em dois subgrupos de 10 animais cada, de acordo com a época de eutanásia: 15º e 45º dias. A sobrevida e peso dos animais foram anotadas. O polo inferior foi medido, pesado e analisado morfologicamente.

\textbf{Resultados:} Todos os animais sobreviveram. O peso aumentou em todos os subgrupos, porém diminuiu no 10º dia nos subgrupos tratados com oxigênio hiperbárico (p<0,001). A viabilidade do polo inferior foi mais evidente nos animais tratados no 15º dia, porém não diferiu no 45º dia. O crescimento do polo inferior não ocorreu no 15º e sim no 45º dia pós-operatório, nos animais não tratados (p<0,01) e tratados (p<0,05). O aumento celular e vascular nos animais tratados foi mais significativo do que nos animais não tratados.

\textbf{Conclusão:} A oxigenoterapia hiperbárica não interferiu na sobrevida dos animais, porém diminuiu o peso. Melhorou a viabilidade do polo inferior do baço, mas não interferiu no seu crescimento.

Introduction

For many years, the spleen, an organ of the reticuloendothelial system was considered non-essential to life and, therefore, its removal should not cause serious damage to the patient. In 1952, King and Shumacker Jr. reported the association between total splenectomy and the occurrence of sepsis in children. Since then, the risks of infection after splenectomy have been observed in children and adults, especially in the first two years after surgery. Complicated infections after splenectomy were also observed in experimental animals. Absence of the spleen is related to alterations in lipid metabolism in humans and in laboratory animals thus causing atherosclerosis.

In addition to the functions assigned to the spleen, many others are certainly still unknown, which strengthens the need to preserve all or part of it by splenorrhaphies, vascular occlusions, partial splenectomies and autoimplants. The most recently described procedure is subtotal splenectomy with the preservation of the inferior pole (STPI). In this technique, the blood supply is maintained through the vessels of the gastrosplenic ligament. This operation was performed in dogs and in rats and an evaluation of the inferior pole in the immediate postoperative period, showed changes in the viability of the remaining tissue, in some cases accompanied by changes in the lipid function. In the late postoperative period there was marked follicular hyperplasia and increased cell population of the inferior pole in dogs on the 60th day. Paulo et al. observed the growth of the remaining tissue on the 80th day. STPI performed in other models than the one described in our work, showed that the histological pattern of this pole becomes similar to normal splenic tissue in 45 days. It’s unclear, however, to which factor these changes can be attributed. A recent study showed that, on the 11th day after surgery, the animals treated with hyperbaric oxygen (HBO) had improved the lipid function and the viability of the inferior splenic pole in comparison with untreated animals. On the 70th day after surgery, however, there was no influence of the hyperbaric oxygen therapy on the viability of that pole.

In experimental studies it was found that hyperbaric oxygen therapy (HBO) has an angiogenic effect, stimulates collateral circulation, decreases the deleterious effects on liver and spleen in rats submitted to hepatic veins ligature with increased survival and improves the outcome of transplantation of pancreatic islets. Considering these facts, it was questioned whether this therapeutic approach could actually interfere with the viability and hence with the growth of the remaining inferior pole, after the operation. Thus, it would be possible to restore some or all the functions of the spleen, avoiding the complications of splenic dysfunction.

Another relevant aspect to be studied concerns the analysis of the effects of hyperbaric oxygen therapy on the evolution and survival of animals. In a previous work, the rats that underwent STPI and were treated with hyperbaric oxygen for 10 days showed significant weight loss on the 11th day after surgery. Moreover, these rats had a shorter survival than those that were not treated, although in a non-significant way. In order to investigate these issues, we decided to study the effect of hyperbaric oxygen therapy in rats submitted to subtotal splenectomy taking into consideration the following aspects: animals’ weight and survival, and viability and growth of the inferior splenic pole.

Methods

This work was performed at the Laboratory of Animal Experimentation, Research Center of the School of Sciences of Santa Casa de Misericordia de Vitoria - ES (EMESCAM). Animal manipulation followed the recommendations of the Brazilian Society of Laboratory Animal Science (SBCAL/COBEA), and was approved by the Ethics Research Committee of the Federal University of Minas Gerais, according to ETIC document n° 004/2008.

Sample characteristics and animal care

Forty male Wistar rats weighing between 274 and 313g (294.15 ± 10.33g) were used. They were obtained at the Animal Production Vivarium of the Research Center of EMESCAM. The animals were kept in appropriate cages and properly identified and maintained in STD 5 Vivarium Cabinet (Vidy Group - Sao Paulo, Brazil) under appropriate conditions such as acclimatization, temperature (20 to 22 °C), ventilation and light (12 hours light and 12 hours dark) control. The animals received rat chow (CR-1 Nuvilab autoclavable - Nuvital) and water ad libitum at all stages of the experiment.

Animal group formation

The animals were distributed at random into two groups: those that were treated with hyperbaric oxygen and those who were not treated. Each group was divided into two subgroups according to the day of euthanasia in the postoperative period:

- Group A (n=20) - not submitted to hyperbaric oxygen therapy
- A15 subgroup (n=10) - euthanasia on 15th day after surgery;
- A45 subgroup (n=10) - euthanasia on 45th day after surgery;
• Group B (n=20) - submitted to hyperbaric oxygen therapy
  - B15 subgroup (n=10) - euthanasia on 15th day after surgery;
  - B45 subgroup (n=10) - euthanasia on 45th day after surgery.

The animals were housed in groups of four in each cage, identified according to the group and subgroup. The animals were individually monitored throughout the experiment and information on each animal was recorded.

Anesthesia and splenectomy with preservation of the inferior pole

The surgical procedure was the same for all 40 animals in the experiment and the surgeon did not know which animals were assigned to each group.

After fasting for six hours, the animals were weighed (electronic scale Filizola® model MF-6 - sensitivity to 1g) and anesthetized with ketamine hydrochloride (Vetaset®, Fort Dodge-Iowa, USA) at a dose of 50 mg/kg associated with xylazine hydrochloride (Kensol®, König-Avellaneda, Argentina) at a dose of 5 mg/kg applied intraperitoneally. The rats were immobilized on the surgical table, had their chest and abdominal wall shaved and underwent antiseptics with topical Polvidine.

The surgical procedure consisted of: a) midline laparotomy approximately 2.5 cm in length, which started 0.5 cm below the xiphoid process; b) exploration of the abdominal cavity and mobilization of the spleen to the surface of the abdominal cavity; c) ligation and sectioning of vessels that irrigate the upper and middle portion of the spleen, close to the splenic surface, with mononylon 6.0 (Shalom®, Shalom Wire Surgical Ltda, Goias, Brazil); d) sectioning of the spleen below the lacquered vessels, keeping the wound unsutured and without omentum protection and inferior pole irrigated by vessels of the gastrosplenic ligament17 (Figure 1); e) measuring the spleen’s inferior pole in its central part, as to length, width and thickness, using plastic caliper; f) suturing of the abdominal wall in plans, using mononylon 6.0 (Shalom®, Shalom Wire Surgical Ltda, Goias, Brazil).

At surgery, 5 mL of 0.9% saline were administered subcutaneously and repeated every 24 hours for two days, aiming at fluid replacement.

The upper portion of the excised spleen was fixed in neutral 10% buffered formalin (potential of Hydrogen - pH 7) 24 to 48 hours and processed in paraffin for histological analysis. This splenic tissue was used to control the inferior splenic pole for microscopic examination.

Postoperatory

After the surgical procedures, the animals, still anesthetized, were identified with ear tags and returned to their cages of origin. The animals received water and food ad libitum and 200 mg/kg paracetamol (Tylenol®, Janssen-Cilag, Sao Paulo, Brazil) orally, dissolved in their drinking water, for 72 hours. The evolution of the animals was monitored throughout the postoperative period.

Hyperbaric oxygen therapy (HBO)

Hyperbaric oxygen therapy was performed only in group B, according to the protocol16. Immediately after recovery from anesthesia, the rats were placed in the hyperbaric chamber; where oxygen pressure was progressively adjusted for 15 minutes to up to 2.5 atmospheres. Animals were exposed to this therapy for 90 minutes, followed by exposition to gradual decompression chamber for 15 minutes. This procedure was performed twice daily during the first three days and once daily for seven days.
Euthanasia and removal of the remaining splenic tissue

At the 15th and 45th days of the postoperative period, the rats of group A and group B were weighed (electronic scale Filizola® model MF-6 - sensitivity of 1g) and intraperitoneally anesthetized with sodium thiopental (Thiopentax®, Cristalia, Sao Paulo, Brazil) at a dose of 50 mg/kg. The animals were then euthanized with a lethal dose of sodium thiopental and 10% potassium chloride 10% (Farmace, Ceara, Brazil), intracardiac (300 mg/kg). The animals underwent laparotomy through an inverted “U” incision in the abdominal wall. In the exploration of the cavity, the abdominal viscera were inspected as well as the presence or absence of adhesions and the aspect of the inferior splenic pole. The remaining splenic tissue was excised for examination.

Measurement and weighing of the inferior splenic pole

Before its removal, the central part of the inferior pole was measured (length, width, thickness) with a plastic caliper, the same way it was done during subtotal splenectomy. After the removal of the inferior pole it was weighed to 0.001g precision on an Adventurer OHAUS model AR 3130 balance (Sao Paulo, Brazil). Information on each animal (measure and weight of the inferior pole) was recorded.

Macroscopic examination

The inferior splenic pole of each animal was macroscopically examined for color, consistency and presence or absence of necrosis. The specimen was photographed and fixed in 10% neutral formaldehyde solution (pH 7) 24 to 48 hours and processed in paraffin for morphological analysis.

Microscopic examination

The fragments of the superior portion (upper and middle) and the inferior splenic pole were processed in 3µm thick paraffin blocks and stained with hematoxylin-eosin. The microscopy was performed with binocular microscopes by two pathologists who did not know to which subgroup the animals belonged. 10 slides were analyzed in subgroup A15, 10 in subgroup A45, 10 in subgroup B15 and 10 in subgroup B45 and 10 slides with the tissue of the superior portion of the spleen at random selection (control). In each slide pathologists examined 10 random fields (100X) and investigated the following parameters: lymphatic follicles (germinal centers), number of lymphocytes, number of sinusoids, cellular proliferation and blood vessels.

Variables studied and statistical tests

Descriptive statistics was used to calculate the arithmetic mean and standard deviation of the rats’ weight as well as the weight, length, width and thickness of the inferior splenic pole.

Student’s t-test for related samples was used to compare the weight of animals in groups A and B, preoperatively to postoperatively (10th, 15th and 45th day). This test was also used to compare the length, width and thickness of the inferior splenic pole of animals in groups treated or not treated with hyperbaric oxygen, from the beginning to the end of the experiment.

Student’s t-test for independent samples was used to compare the weight of the inferior pole of the animals treated with hyperbaric oxygen with those that were untreated.

Fisher’s exact test was utilized to compare the frequency of the feasibility of the inferior pole of the animals treated with hyperbaric oxygen with that of untreated animals.

All tests were two-tailed and P values were considered significant when equal or less than 5%.

Results

Survival of animals and examination of the abdominal cavity

There were no deaths in both groups. The abdominal viscera showed no changes from the first surgery (intraoperative). Adhesions of the inferior splenic pole to the neighboring structures (stomach, greater omentum) were noted in all animals treated or not with hyperbaric oxygen.

Weight of animals

The weight of the animals in the group that was not treated with hyperbaric oxygen increased significantly at the end of the experiment compared to its beginning (Subgroups A15 - p<0.0001 and A45 p=0.0001). In animals treated with hyperbaric oxygen, B15 and B45 subgroups, there was weight loss on the 1st to the 10th day postoperatively (p<0.01) and subsequently the animals gained weight (p<0.0001) (Table 1).

Acta Cirúrgica Brasileira - Vol. 26 (3) 2011 - 159
### TABLE 1 - Rat weight (g) in groups treated and not treated with hyperbaric oxygen in the preoperative period, at the 10th day and at the postoperative period, submitted to subtotal splenectomy with the preservation of the inferior pole.

<table>
<thead>
<tr>
<th>Group</th>
<th>Subgroup</th>
<th>Preoperative AM (SD)</th>
<th>10th day AM (SD)</th>
<th>Postoperative AM (SD)</th>
<th>P</th>
<th>P1</th>
<th>P2</th>
</tr>
</thead>
<tbody>
<tr>
<td>A</td>
<td>A15</td>
<td>302.80 ± 8.54</td>
<td>319.90 ± 9.80</td>
<td>339.40 ± 8.26</td>
<td>&lt;0.0001</td>
<td>&lt;0.0001</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td></td>
<td>A45</td>
<td>294.40 ± 4.52</td>
<td>314.30 ± 6.05</td>
<td>382.50 ± 18.43</td>
<td>&lt;0.0001</td>
<td>&lt;0.0001</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>B</td>
<td>B15</td>
<td>293.33 ± 11.73</td>
<td>288.22 ± 9.99</td>
<td>309.44 ± 14.96</td>
<td>&lt;0.0001</td>
<td>&lt;0.0001</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td></td>
<td>B45</td>
<td>286.00 ± 8.76</td>
<td>281.10 ± 10.50</td>
<td>412.10 ± 27.40</td>
<td>&lt;0.0001</td>
<td>&lt;0.0001</td>
<td>&lt;0.0001</td>
</tr>
</tbody>
</table>

**AM -** Arithmetic Mean; **SD -** Standard Deviation. T test for related samples. p<0.05 significant; **P** - comparison between the weight of animals of the same subgroup, between the preoperative period and the 10th day of the experiment; **P1** - comparison between the weight of the animals of the same subgroup, between the preoperative and postoperative periods of the experiment; **P2** - comparison between the weight of animals of the same subgroup, between the 10th day and postoperative period of the experiment.

### Weight of the remaining inferior pole

The weight of the inferior splenic pole increased significantly only in animals not treated with hyperbaric oxygen (A15 and A45 subgroups), (p<0.01). In the other comparisons that variable did not change significantly (Table 2).

### TABLE 2 - Final weight of the inferior splenic pole in rats treated and untreated with hyperbaric oxygen in the mediate preoperative and mediate postoperative periods (15th day) and in the late preoperative and late postoperative periods (45th day) who underwent subtotal splenectomy, with the preservation of the inferior pole.

<table>
<thead>
<tr>
<th>Group</th>
<th>Subgroup</th>
<th>Initial weight</th>
<th>Final weight</th>
<th>P1</th>
<th>P2</th>
</tr>
</thead>
<tbody>
<tr>
<td>A</td>
<td>A15</td>
<td>9.90 ± 2.07</td>
<td>8.60 ± 1.50</td>
<td>&gt;0.05</td>
<td></td>
</tr>
<tr>
<td></td>
<td>A45</td>
<td>10.10 ± 1.28</td>
<td>11.70 ± 1.70</td>
<td>&lt;0.01</td>
<td>&lt;0.05</td>
</tr>
<tr>
<td>B</td>
<td>B15</td>
<td>8.44 ± 1.57</td>
<td>8.22 ± 1.78</td>
<td>&gt;0.05</td>
<td></td>
</tr>
<tr>
<td></td>
<td>B45</td>
<td>8.90 ± 1.66</td>
<td>10.35 ± 2.22</td>
<td>&gt;0.05</td>
<td></td>
</tr>
</tbody>
</table>

**AM -** Arithmetic Mean; **SD -** Standard Deviation. **P** - comparison between the length of the inferior pole of the animals of the same subgroup, from the beginning to the end of the experiment. T test for independent samples. p<0.05 significant.

### Measures of the remaining inferior pole

The length of the inferior splenic pole increased significantly only in animals not treated with hyperbaric oxygen and which were euthanized at the 45th day (A45) (p<0.01). In other subgroups, this variable did not change (p>0.05) (Table 3).

### TABLE 3 - Length of the inferior splenic pole in rats treated and not treated with hyperbaric oxygen in the mediate preoperative and mediate postoperative periods (15th day) and in the late preoperative and late postoperative periods (45th day) that underwent subtotal splenectomy with the preservation of the inferior pole.

<table>
<thead>
<tr>
<th>Groups</th>
<th>Subgroups</th>
<th>Initial length</th>
<th>Final length</th>
<th>P</th>
</tr>
</thead>
<tbody>
<tr>
<td>A</td>
<td>A15</td>
<td>8.50 ± 1.17</td>
<td>7.50 ± 1.95</td>
<td>&gt;0.05</td>
</tr>
<tr>
<td></td>
<td>A45</td>
<td>8.50 ± 0.52</td>
<td>9.30 ± 1.05</td>
<td>&gt;0.05</td>
</tr>
<tr>
<td>B</td>
<td>B15</td>
<td>7.66 ± 0.66</td>
<td>7.50 ± 1.93</td>
<td>&gt;0.05</td>
</tr>
<tr>
<td></td>
<td>B45</td>
<td>7.90 ± 0.99</td>
<td>9.05 ± 1.69</td>
<td>&gt;0.05</td>
</tr>
</tbody>
</table>

**AM -** Arithmetic Mean; **DP -** Standard Deviation. **P** - comparison between the width of the inferior pole of the animals of the same subgroup, from the beginning to the end of the experiment. t-Test for related samples p<0.05 significant.

### TABLE 4 - Width of the inferior splenic pole in rats treated or not with hyperbaric oxygen in the mediate preoperative period and the mediate postoperative period (15th day) and in the late preoperative period and late postoperative period (45th day) who underwent subtotal splenectomy with the preservation of the inferior pole.

<table>
<thead>
<tr>
<th>Grupos</th>
<th>Subgrupos</th>
<th>Largura inicial</th>
<th>Largura final</th>
<th>P</th>
</tr>
</thead>
<tbody>
<tr>
<td>A</td>
<td>A15</td>
<td>8.50 ± 1.17</td>
<td>7.50 ± 1.95</td>
<td>&gt;0.05</td>
</tr>
<tr>
<td></td>
<td>A45</td>
<td>8.50 ± 0.52</td>
<td>9.30 ± 1.05</td>
<td>&gt;0.05</td>
</tr>
<tr>
<td>B</td>
<td>B15</td>
<td>7.66 ± 0.66</td>
<td>7.50 ± 1.93</td>
<td>&gt;0.05</td>
</tr>
<tr>
<td></td>
<td>B45</td>
<td>7.90 ± 0.99</td>
<td>9.05 ± 1.69</td>
<td>&gt;0.05</td>
</tr>
</tbody>
</table>

**AM -** Arithmetic Mean; **DP -** Standard Deviation. **P** - comparison between the width of the inferior pole of the animals of the same subgroup, from the beginning and the end of experiment. t-Test for related samples p<0.05 significant.
The thickness of the inferior splenic pole increased significantly only in animals treated with hyperbaric oxygen and euthanized on 45th day (B45). In the other subgroups, that variable did not change significantly (p>0.05).

**TABLE 5** - Thickness of the inferior splenic pole in rats treated and not treated with hyperbaric oxygen in the mediate preoperative and mediate postoperative periods (15th day) and late preoperative and late postoperative periods (45th day) who underwent subtotal splenectomy with the preservation of the inferior pole.

<table>
<thead>
<tr>
<th>Groups</th>
<th>Subgroups</th>
<th>Initial thickness</th>
<th>Final thickness</th>
<th>p</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>AM ± SD</td>
<td>AM ± SD</td>
<td></td>
</tr>
<tr>
<td>A</td>
<td>A15</td>
<td>4.20 ± 0.91</td>
<td>3.35 ± 1.05</td>
<td>&gt;0.05</td>
</tr>
<tr>
<td></td>
<td>A45</td>
<td>4.00 ± 0.81</td>
<td>4.45 ± 0.89</td>
<td>&gt;0.05</td>
</tr>
<tr>
<td>B</td>
<td>B15</td>
<td>3.44 ± 0.46</td>
<td>3.61 ± 0.74</td>
<td>&gt;0.05</td>
</tr>
<tr>
<td></td>
<td>B45</td>
<td>3.15 ± 0.57</td>
<td>4.35 ± 1.13</td>
<td>&lt;0.05</td>
</tr>
</tbody>
</table>

AM - Arithmetic Mean. SD - Standard Deviation. 
*p* - comparison between the thickness of the inferior pole of the animals of the same subgroup, between the beginning and end of the experiment. T-test for related samples; *p*≤0.05 significant.

**Macroscopic evaluation of the inferior splenic pole**

In eight animals that underwent hyperbaric oxygen therapy (subgroup A15) the macroscopic appearance of the inferior pole was normal and in 60% of the cases adhesions were found at the hepatic lobe and the intestinal loops. Vessels were observed at the periphery of the inferior splenic pole adhered to the omentum. In two animals of this subgroup (20%) necrosis was observed at the end and central portion of the inferior pole. In the rats of subgroup A45, the macroscopic appearance of the inferior pole was normal in all cases, with vascularized adhesions on the greater omentum, intestinal loops and liver. Only one case was described as showing few adhesions.

In animals submitted to hyperbaric oxygen therapy (subgroup B15), the macroscopic appearance of the inferior pole was considered viable, with vascularized adhesions by the greater omentum, intestinal loops and liver. In the animals of subgroup B45, the macroscopic aspect of the inferior pole was considered feasible in all cases, with vascularized adhesions by the greater omentum, intestinal loops and liver. There was no macroscopic difference between subgroups A45 and B45.

**Microscopic evaluation of the inferior splenic pole**

Lymph follicles were larger and more frequent in the inferior pole of the animals in subgroup A15 than in the superior portion of the spleen (control). These follicles were larger in the inferior pole of animals of the subgroups A45, B15, B45 than in the animals of subgroup A15 and in the superior splenic portion (control). The presence of lymphocytes was more frequent in the inferior splenic pole of the animals in subgroup A15 than in the superior portion (control).

There was a greater amount of lymphocytes in subgroups A45, B15 and B45 than in the superior portion (control). There was no marked difference in the amount of sinusoids between the superior splenic portion (control) and the inferior pole of the four subgroups in a conventional morphological study. However, the cellular and vascular increase in subgroups B15 and B45 is noteworthy when compared to the inferior pole of the A15 and A45 subgroups. Cell proliferation was more pronounced in the inferior pole of the A15 subgroup than in the superior portion (control) and was less intense in the superior portion (control) than in other subgroups.

There was an increase of lymphocyte cell proliferation in subgroup A45, B15 and B45. The presence of cells in the wall and inside the vessels was more intense in the inferior pole of the B15 and B45 subgroups and occurred with less intensity in the superior splenic portion, which was excised in partial splenectomy.

**Discussion**

Preservation of splenic tissue is important to avoid immunodeficiency caused by splenectomy, especially in children. Bradshaw and Thomas reported that the larger the remaining spleen, the most significant the protection against sepsis and they calculated that 25% of the critical mass of residual splenic tissue would restore normal phagocytic function. This suggestion is accepted by most authors. Van Wyck et al. in an experimental work with rats considered that it was necessary to maintain a third of the total splenic tissue to restore splenic function. In this work, subtotal splenectomy preserving the inferior pole kept about a third of the total splenic tissue.

When performing subtotal splenectomy, it is important to follow the technical care suggested by Paulo et al. The present study followed two other recommendations made by the author of the subtotal splenectomy technique. The first one was to preserve the thin and transparent peritoneal fold extending from the anterior face of the greater curvature of the antrum to the anterior surface.
of the inferior splenic pole. This approach was adopted to keep the remaining fixed, i.e., with its cut surface facing anteriorly. The second one was not to fix the inferior pole below the greater curvature of the stomach, with the intention of preventing the twist of this remnant. Previous experience with the technique of subtotal splenectomy in rats showed that fixing the inferior pole in the stomach resulted in the high percentage of necrosis of this remnant. This phenomenon possibly occurred because fixing the inferior pole could compromise the blood supply of the inferior splenic pole due to the twist or traction of the small vessels of the pedicle. Thus, the inferior pole should be placed in the abdominal cavity with its cut surface facing toward the skull. After this maneuver, the greater omentum adheres naturally to the pole, which can be beneficial.

Hyperbaric oxygen therapy was performed according to the protocol suggested by Paulo et al.17. These authors suggested two sessions of HBO in the first three days and a daily session within seven days after STPI to improve the viability of the inferior pole or even accelerate its growth, as this treatment has an angiogenic effect.

In 1999, Paulo et al.17 reported signs of impairment of the viability and function of the inferior pole in dogs in the early postoperative period. Hence, the use of HBO in rats subjected to STPI could improve the viability of this remnant.

In this study there were no surgical complications. All animals which underwent STPI presented satisfactory recovery. Only one animal from subgroup B15 was excluded from the study because of the hyperbaric oxygen treatment because it presented, in the third postoperative day, after the session of HBO, signs of dehydration: bristling hair, malaise and high upper respiratory frequency. Nevertheless, all animals survived.

HBO can help the animal lose weight, as demonstrated in a later study24,29. In this work, the animals subjected to this treatment lost weight during the first ten days and the group which was not submitted to the treatment gained weight during the same period. Possible causes of weight loss were the daily manipulation of the rats, their stay in the hyperbaric chamber during the period stipulated by the protocol and the stress resulting from the procedure. In this study, some animals treated with HBO showed chromodacryorrhea (tear red), which is indicative of stress or pain in rats.

Thus, it is interesting to quantify catecholamines and cortisol to compare the stress level of animals subjected to HBO treatment with those animals which were not treated with HBO. The viability of the inferior pole was detected macroscopically in 80% of the untreated animals (subgroup A15) and in 100% of the animals treated with hyperbaric oxygen (subgroup B15). This difference, however, was not significant. Microscopic analysis revealed that the animals in subgroup B15 (treated) compared to subgroup A15 (untreated) exhibited larger lymph follicles, a greater number of cells and vessels and higher lymphocytic proliferation. These data suggest a better viability of the inferior splenic pole in animals treated with hyperbaric oxygen than in those which were not treated (B15).

It is known that HBO has an angiogenic effect, which may have contributed to the results of this work. The protective effect of HBO on the inferior pole was already reported24. On 45th day there was no difference in the viability of this pole between the subgroups treated and not treated with hyperbaric oxygen from the macroscopic point of view. However, there was a greater amount of vessels and cells in subgroup A45 compared to B45 in the microscopic evaluation, in relation to lymphoid follicles, lymphocytes and lymphocyte proliferation. This seems to indicate that HBO in the 45 day subgroup had an effect as striking as that in the subgroup of 15 days. It is important to remember that HBO was administered during the first ten days postoperatively and that the B45 subgroup received no such treatment for 35 days. Thus, during this period without treatment, the phenomena which occurred in subgroup A45 could have helped to avoid the macroscopic and microscopic differences between the inferior pole of the groups treated and not treated with hyperbaric oxygen. It should be emphasized that the morphological aspect does not necessarily have functional correspondence. Therefore, the tissue may be viable and appear normal even if it does not work properly35.

The growth of the inferior splenic pole has already been mentioned. Torres et al.23 in a model of subtotal splenectomy, different from the one in STPI, described the regeneration of this remnant in rats on the 45th day. It was not known, however, if HBO could interfere with the growth of this pole. The present work is an attempt to settle this doubt. In this work, on the 15th day, when the length, width and thickness of the pole were measured from the beginning to the end of the experiment, it was verified that the inferior pole did not grow in the animals treated or not with hyperbaric oxygen. However, in subgroups of 45 days this remainder grew. A significant increase in the length of the inferior splenic pole in subgroup A45 was obtained as well as an increase in the thickness of this pole in subgroup B45. It should be noted that the growth of this remnant is suggested by the fact that their average weight in animals of 45 days was markedly higher than in animals of 15 days not treated with hyperbaric oxygen.

In the treated animals, this pole was heavier in the group

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Nunes TA et al.
of 45 days, but not significantly. In the microscopic analysis, subgroups A45, B15 and B45, when compared to subgroup A15 showed increased lymphoid follicles, a great number of lymphocytes, higher cell proliferation and a great number of vessels. This suggests a further growth of the inferior pole of these subgroups. Although this growth has not occurred macroscopically in subgroup B15, microscopy shows signs of growth.

These results support the view that the morphological changes of the inferior splenic pole occur with the passing of time. Paulo et al.23 emphasized the improvement of lipid metabolism in the spleen. A recent study showed that the inferior pole grew significantly in length, width and thickness, from the 1st to the 80th day postoperatively, after STPI. The microscopic sign of this phenomenon was cell hyperplasia23. The inferior splenic pole grows for different reasons: the animal’s growth, inflammation, and unknown factors. In this study, there was an increase in animal weight between the preoperative and postoperative periods and unknown factors. In this study, there was an increase in animal weight between the preoperative and postoperative periods, but not significantly. In the microscopic analysis, animal weight between the preoperative and postoperative periods, and 45th day. It is possible that on the 45th day after surgery the initial inflammatory process has been minimized, and thus did not contribute to the growth of the inferior pole.

Studies on immunohistochemistry and molecular genetics are being developed to complement this work in order to clarify these issues.

Conclusion

Hyperbaric oxygen therapy did not affect the survival of animals but reduced their weight. It improved the viability of the inferior splenic pole, but did not interfere with their growth.

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