









# Effect of antileukotriene treatment on capsular contracture: an experimental study

*Efeito do tratamento com antileucotrieno em contratura capsular: estudo experimental*

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## ■ ABSTRACT

**Introduction:** 40% of patients undergoing radiotherapy after breast reconstruction by silicone prosthesis implant may develop prosthesis encapsulation. Several strategies have already been tested to prevent capsule contracture with unsatisfactory results. This study analyzed the effect of topical antileukotriene (AL) on capsular contracture formation in rats with silicone implants associated with irradiation. **Methods:** Silicone blocks were implanted in the dorsal region in 20 female rats Wistar with weights ranging from 200-250g. The animals were divided into two groups: control (injection of 0.9% saline solution into the tissue around the implant) and intervention group (injection of 10mg of AL into the tissue around the implant). Immediately after surgery, the animals were irradiated with a single dose of 10Gy. After two months, we collected capsule samples for histological analysis and gene expression analysis of the following biomarkers: iNOS, VEGF-a and MMP-9. **Results:** Vascular density was lower in the AL group when compared to the control group ( $55.4 \pm 30.0$  vs.  $81.8 \pm 26.7$ ,  $p=0.05$ , respectively). Similarly, VEGF-a had the same behavior (control group -  $0.34 \pm 0.1$  vs. group Al -  $0.02 \pm 0.001$ ,  $p=0.04$ ). **Conclusion:** This study suggested that treatment with AL decreases angiogenesis in animals submitted to silicone implants and underwent radiotherapy.

**Keywords:** Radiotherapy; Mouse; Capsular contracture in implant; Vascular endothelium growth factor A; Matrix metalloproteinases inhibitors.

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## ■ RESUMO

**Introdução:** 40% dos pacientes submetidos à radioterapia após reconstrução de mama por implante de prótese de silicone podem desenvolver encapsulamento da prótese. Diversas estratégias já foram testadas para prevenir a contratura da cápsula com resultados insatisfatórios. Este estudo analisou o efeito do antileucotrieno (AL) tópico na formação de contratura capsular em ratos com implantes de silicone associados à irradiação. **Métodos:** Foram implantados blocos de silicone na região dorsal em 20 ratas fêmeas, espécie Wistar com peso variando de 200-250g. Os animais foram divididos em dois grupos: controle (injeção de solução fisiológica 0,9% no tecido ao redor do implante) e grupo intervenção (injeção de 10mg de AL no tecido ao redor do implante). Imediatamente após a cirurgia os animais foram irradiados com dose única de 10Gy. Após dois meses, coletamos amostras de cápsulas para análise histológica e análise da expressão gênica dos seguintes biomarcadores: iNOS, VEGF-a e MMP-9. **Resultados:** A densidade vascular foi menor no grupo AL quando comparado ao grupo controle ( $55,4 \pm 30,0$  vs.  $81,8 \pm 26,7$ ,  $p=0,05$ , respectivamente). Da mesma forma, o VEGF-a teve o mesmo comportamento (grupo controle -  $0,34 \pm 0,1$  vs. grupo Al -  $0,02 \pm 0,001$ ,  $p=0,04$ ). **Conclusão:** Este estudo sugeriu que o tratamento com AL diminui a angiogênese em animais submetidos a implantes de silicone e submetidos à radioterapia

**Descritores:** Radioterapia; Rato; Contratura capsular em implante; Fator A de crescimento do endotélio vascular; Inibidores de metaloproteinases de matriz.

## INTRODUCTION

Breast implants are routinely used in cosmetic and restorative breast plastic surgery; according to the American Society of Plastic Surgery, in 2016, 290,467 implants for aesthetic purposes and 109,256 implants for repair procedures<sup>1</sup>. Although widely used, there is a risk of capsular contracture in 10.6% of operated cases<sup>2</sup>. However, several patients should undergo radiotherapy as local breast cancer treatment when performing reconstructive surgery, such as breast reconstruction. In these cases, the prevalence of this complication is greater than 40%, and the need for removal of the prosthesis is 15%<sup>2</sup>.

When inert materials, such as silicone are implanted in the human body, the body forms a connective tissue capsule around the implant to isolate it from adjacent tissues. In some cases, this envelope becomes fibrotic due to trauma, infection, and other unknown causes (capsular contracture). Depending on the degree of fibrosis may cause aesthetic deformity, pain and inflammation of the implant site<sup>3</sup>. Therefore, when the prosthesis is encapsulated, the treatment should be surgical with the removal of the prosthesis and replacement by another prosthesis in another anatomical plane, for example, if the prosthesis is placed above the pectoral muscle and there is a

contracture, it should be replaced by another prosthesis implanted below the muscle. This procedure aims to avoid recurrence of contracture<sup>4,5</sup>.

For this reason, several techniques have been studied to prevent the occurrence of capsular contractures, such as the use of textured implants, implantation of the prosthesis in a submuscular pocket<sup>4</sup>, use of medications such as corticosteroids, cyclosporine<sup>5</sup> and antileukotrienes<sup>6,7</sup>. All these therapeutic alternatives showed variable results without composing clinical evidence.

Bastos et al. (2012)<sup>8</sup> showed positive results of using Montelukast, an antagonist of leukotrienes receptors, to prevent capsular contracture.

## OBJECTIVE

This study will analyze the effect of Montelukast on animal models of radio-induced capsular contracture. Therefore, this study aimed to analyze the effect of the antileukotriene drug (Montelukast<sup>®</sup>) on capsular contracture formation in rats with silicone implants who underwent radiotherapy within two months.

## METHODS

Twenty Wistar rats were analyzed, with weights ranging from 250-300g. All animals were kept in a

bioterium with controlled temperature between 22-23°C, day/night cycle of 12.12h, water and food ad libitum. This study followed the national rules for the standardization of studies on laboratory animals (CONCEA) and was approved by the Ethics Committee on Animal Use of the Faculty of Medicine of the University of São Paulo (CEUA-FMUSP) under registration 067/2017.

### Surgical procedure

The animals were anesthetized by a solution composed of 100mg/kg of weight of ketamine hydrochloride (Ketamin®, Cristália, Brazil) and 10mg/kg of xylazine hydrochloride (Rompun®, Bayer, Brazil) by intraperitoneal route.

After intraperitoneal anesthesia, the animals were positioned in ventral decubitus. Trichotomy was then performed in an area of 6cm<sup>2</sup> in the region of the midline of the spine just below the scapulae. Topical chlorhexidine was used for antiseptis in this region.

A 2cm horizontal incision was performed using a 15-inch blade scalpel. Pocket dissection of 6 cm<sup>2</sup> below the plane of the panniculus carnosus of the animal and hemostasis with an electric scalpel. Before performing the closing of the pocket was implanted a 2x2 cm silicone block. And then the animals were randomized into two groups: control (n=10), cream injection around the silicone block using 1ml of 0.09% saline solution; antileukotriene group (n=10), tissue injection around the silicone block using 1ml of antileukotriene (Singulair®, Merck Sharp & Dohme, Campinas, Brazil) diluted in 0.9% saline solution.

### Radiotherapy

All animals were submitted in the immediate postoperative period to a radiotherapy session. The radiation was made a specific irradiator for small animals (Rad Source RS2000, Quastar™), with an X-ray of 160KV of 25mAs.

The animals were positioned in lateral decubitus contained by an immobilizer that allowed the emission of rays in the tangential plane as used in patients. A 5mm lead plate was also inserted that had a window for the irradiation of the animal, ensuring treatment only in the desired region. A single dose of 10Gy was used in two opposite fields (5Gy per field) with a 6Gy/min card.

### Histological analysis

After two months, the animals were euthanized by intraperitoneal injection of ketamine hydrochloride (Ketamin®, Cristália, Brazil) at a dose of 180mg/kg of weight, associated with xylazine (Rompun®, Bayer,

Brazil) at a dose of 15mg/kg and then placed in a 100% CO chamber.

Samples of the capsule were collected around the silicone block through an incision in the back of the animals. Part of the samples was fixed with formalin 4% for 24h and then colored with hematoxylin-eosin (HE) and picosirius for microscopy analysis (Nikon eclipse E600®, Japan) in magnification of 200 and 400 times.

The hematoxylin-eosin-stained slides analyzed the presence of inflammatory cells and vascular density (arteriole count) in 10 fields along the histological sections. The slides stained by the picosirius method graphically demonstrated collagen deposition along with the capsule around the silicone block.

### Gene expression analysis

#### Total RNA extraction

Panniculus carnosus samples were macerated using the Tissue Lyser LT device (Qiagen, Germantown, USA). 1.0ml Trizol® (Invitrogen-Life Technologies, Carlsbad, USA) and stainless-steel beads were added to microcentrifuge tubes with samples. Fragmentation was performed for 6 minutes at 50Hz.

After the removal of the spheres, 0.2 ml of chloroform (Merck) was added. The samples were centrifuged for 15 minutes at 12,000rpm at 4°C. After centrifugation, the aqueous phase was transferred to a new microcentrifuge tube, and 0.5ml of icy isopropyl alcohol (Merck) was added for RNA precipitation. The samples were incubated at room temperature for 10 minutes and then centrifuged at 12,000rpm for 10 minutes at 4°C. The supernatant was discarded, and the RNA precipitate was washed with 1.0ml of 75% ethanol. The RNA pellet was resuspended in 50 to 100µL of sterile ultrapure water free of DNase/RNase (Invitrogen-Life Technologies, Carlsbad, USA).

The concentration of the extracted RNAs was determined in a NanoDrop™ ND-1000 spectrophotometer (NanoDrop Technologies, Inc., Wilmington, USA). The degree of purity was evaluated using the 260/280nm ratio, using only RNAs whose ratio was ≥1.8. To analyze the integrity of the RNAs, agarose gel electrophoresis was performed to verify the 28S and 18S bands. The extracted RNAs were stored at -80°C until use.

#### cDNA Synthesis

For cDNA synthesis from the total RNA, the High Capacity RNA-to-cDNA (Applied Biosystems) kit was used in GeneAmp 2400 thermocycler (Applied Biosystems). In the final volume of 20µL: 1.0µL of enzyme mix; 10.0µL of RT buffer; 20µL q.s. of sterile ultrapure water free of DNase/RNase and total RNA (500ng). For reaction and reaction stoppage, the tubes were incubated at 37°C for

60 minutes and 95°C for 5 minutes, respectively. cDNA samples were stored at -20°C until use.

### Real-time polymerase chain reaction (qRT-PCR)

The analysis of the expression of the mRNA levels of the genes of interest was performed by qRT-PCR in the StepOnePlus™ (Applied Biosystems) thermocycler with the TaqMan® Gene Expression Assays (Applied Biosystems) system.

Probes and primers for the Genes C5AR1 (Rn02134203), ICAM1 (Rn 00564227), NOS2 (Rn 00561646), VEGF (Rn 01511602) and the endogenous control ACTB (Rn 00667869) were acquired from the list of inventoried assays of the company Applied Biosystems.

QRT-PCR was performed in duplicate for each sample using: 10.0µL TaqMan® Universal Master Mix II 2X; 1µL TaqMan® Gene Expression Assay 20X and 4µL of diluted cDNA (dilution 1:5) in the final volume of 20µL, in plates of 96 wells covered with an optical sealant.

The reaction conditions were 50°C for 2 minutes, 95 for 10 minutes, followed by 40 cycles at 95°C for 15 seconds and 60°C for 1 minute.

For the calculation of the expression level of each target gene, GenEx Standard 6.1 (MultiD Analyses AB) software was used, which uses the 2-delta delta Ct method for relative quantification, where Ct (threshold cycle) is the real-time PCR cycle, in which amplification reaches the logarithmic phase, where delta Ct is the difference of expression between the target gene and endogenous control of a given sample, and delta delta Ct corresponds to the difference between the delta Ct of the sample and the delta Ct of the control.

### Statistical analysis

The variables were submitted to descriptive analysis. After checking the distribution and variances of the data, the two groups were compared using the Wilcoxon rank-sum test (non-normal distribution), considering an alpha p of 0.05. Stata software version 14 (StataCorp., 2015) was used. Stata Statistical Software: Release 14. College Station, TX: StataCorp LP, USA).

## RESULTS

Three animals in the control group and one in the AL group presented dehiscence and extrusion of the silicone block after four weeks of irradiation.

### Histological analysis

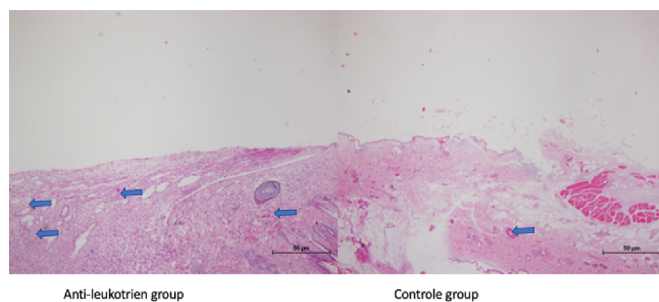
The analysis by the histomorphometry method showed no difference in the number of inflammatory cells. However, there was a decrease in vascular density

in the group treated with AL when compared to the control group (55.4±30.0 (AL) vs. 81.8±26.7 (control); p=0.05) (Table 1 and Figure 1).

**Table 1.** Analysis by histomorphometry refers to inflammatory cells and vascular density in the control and AL groups.

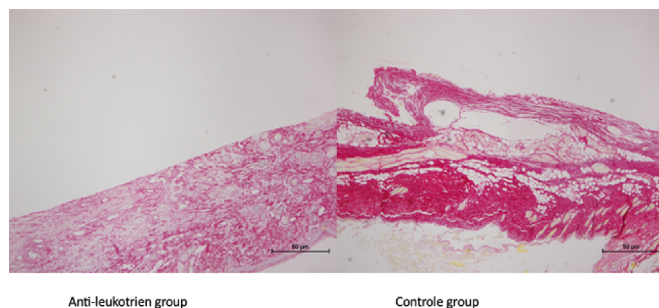
	Control (n=7) (average±SD)	Antileukotriene (n=9) (average±SD)
Inflammatory cells	8.25±5.9	4.4±4.0
Vascular density	81.8±26.7	55.4±30.0

SD = Standard deviation.



**Figure 1.** Slides colored by HE method, samples by groups. The blue arrows correspond to arterioles along with the capsule.

The slides colored by picosirius showed a higher amount of collagen in the control group (Figure 2).



**Figure 2.** Picosirius staining showed the density of collagen fibers in the control group when compared to the AL group.

Analysis of gene expression of biomarkers (iNOS, VEGF-a, MMP-9)

The AL group showed a lower amount of VEGF-a when compared to the control group (0.02±0.001 vs. 0.34±0.1, respectively, p=0.04) (Table 2).

## DISCUSSION

Leukotrienes are lipids (eicosanoid) belonging to the family of inflammation-mediating substances. Act on smooth muscle contraction (bronchoconstriction) and increasing microvascular permeability perpetuating the inflammatory reaction<sup>9</sup>.

**Table 2.** Gene expression of iNOS, VEGF-a, ICAM-1 and MMP-9 in the control and AL groups.

	Control (n=7) (average±SD)	Antileukotriene (n=9) (average±SD)
iNOS	0.013±0.01	0.005±0.001
VEGF-a	0.34±0.1	0.02±0.001
MMP-9	0.36±0.2	0.30±0.1

SD = Standard deviation.

Antileukotrienes, therefore, have the function of reducing and preventing smooth muscle contraction and reducing vascular permeability. Although the mechanism of action has not yet been elucidated, several studies have demonstrated the beneficial effect of antileukotrienes in inhibiting or minimizing capsular contracture after breast alloplastic implants<sup>6-8,10</sup>.

Our hypothesis was to analyze the effect of AL on the local inflammatory reaction (decrease in late neoangiogenesis and decrease in inflammatory cells) and inhibition of fibrosis formation. To enhance the results of this study and to pet the clinical situation of treatment adjunct to breast cancer reconstruction, all animals received a dose of irradiation immediately after surgery.

This study demonstrated a 50% decrease in inflammatory cell count in the AL group ( $4.4 \pm 4.0$ ) compared to the control group ( $8.25 \pm 5.9$ ). Although it did not reach significance, there is a tendency to observe less inflammatory reaction in the AL group. In the histological analysis regarding vascular density, this study demonstrated a lower count of neoformed vessels in the AL group when purchasing with the control group ( $55.4 \pm 30.0$  vs.  $81.8 \pm 26.7$ , respectively;  $p=0.05$ ). These two findings suggest decreased chronic inflammation in the silicone implant capsule. Kang et al. (2015)<sup>11</sup> analyzed the use of Zafirlukast in the formation of silicone implant capsules in rabbits. These authors demonstrated a similar result to our study; animals treated with AL showed decreased capsular contracture after two months of follow-up. Still, Kang et al. (2015)<sup>11</sup> also analyzed the inflammatory reaction around the silicone implant. Again, this group demonstrated a decrease in macrophages, as presented in our study.

Still, Kang et al. (2015)<sup>11</sup> studied an intermediate outcome measuring the gene expression of myofibroblasts. The authors showed a decrease in the expression of the myofibroblast gene in the AL-treated group compared to the control group. Based on these literature findings, our group explored other pathways responsible for the mechanism of action of AL.

The expression of the iNOS gene was higher in the control group than in the AL group ( $0.013 \pm 0.01$  and  $0.005 \pm 0.001$ , respectively) ( $p=0.08$ ), but in this study,

this difference showed a tendency to significance. The action of nitric oxide inducible synthetase (iNOS) refers to an enzyme in the inflammatory response mediated by macrophages. And, consequently, the increase in nitric oxide (NO) produced by macrophages induces a pro-inflammatory state. Therefore, the decrease in gene expression in the AL group suggests an anti-inflammatory effect on the silicone implant capsule. As demonstrated by McNeill et al. (2015)<sup>12</sup>. According to this study, the dissection of macrophages, according to the free radical pathway, demonstrated a decrease in the inflammatory process due to the inhibition of iNOS<sup>11,12</sup>.

Another biomarker studied by McNeill et al. (2015)<sup>12</sup> was VEGF-a as a possible factor contributing to the mechanism of action to the anti-inflammatory effect of AL. In our study, the gene expression of VEGF-a decreased in the group treated with AL compared to the control group. Reflecting on this outcome, the decrease in the long-term neoangiogenesis marker (2 months) may demonstrate the transition of phenotype 1 macrophages to phenotype 2; there is the resolution of the inflammatory process<sup>13</sup>. Another intermediate outcome analyzed in this study was the gene expression of MMP-9, which corresponds to gelatinase metabolism (extracellular matrix component). In addition, several authors have analyzed other metalloproteinases, intending to study collagen metabolism in tissue regeneration<sup>14,15</sup>.

This study found no difference in gelatinase (MMP-9) between groups. However, this study presented some limitations: the anatomical structure of the mine model adopted in this study presented some differences (panniculus carnosus) concerning humans.

## CONCLUSION

This study suggests that the treatment with AL decreases angiogenesis in animals submitted to silicone implants and underwent radiotherapy

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## COLLABORATIONS

CPC Analysis and/or data interpretation, Conception and design study, Conceptualization, Final manuscript approval, Formal Analysis, Methodology, Writing - Review & Editing.

<b>HAC</b>	Analysis and/or data interpretation, Conception and design study, Investigation, Methodology, Realization of operations and/or trials, Writing - Review & Editing.
<b>MSK</b>	Data Curation, Investigation, Methodology, Realization of operations and/or trials, Writing - Original Draft Preparation.
<b>SAM</b>	Data Curation, Realization of operations and/or trials, Writing - Original Draft Preparation.
<b>JMB</b>	Analysis and/or data interpretation, Conception and design study, Methodology, Writing - Review & Editing.
<b>RG</b>	Conception and design study, Final manuscript approval, Supervision, Visualization, Writing - Review & Editing.

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