Effects of topical 2% cyclosporine A on the corneas of dogs subjected to lamellar keratoplasty with a graft of equine pericardium preserved in glycerin.

Clinical and morphological evaluation

Efeitos da aplicação tópica da ciclosporina A a 2% sobre a córnea de cães submetidos à ceratoplastia lamelar com implante de pericárdio de equino preservado em glicerina.

Avaliação clínica e morfológica

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ABSTRACT

The effects of topical 2% cyclosporine A on the cornea of dogs subjected to experimental lamellar keratoplasty with an equine pericardial graft were evaluated. Ten dogs were grouped to be evaluated 3, 7, 15, 30 and 60 days after surgery. Animals received bilateral grafts followed by the application of a 2% cyclosporine A ointment on the left eye (treated eye) and the ointment base on the right eye (control eye) twice a day. The ophthalmic evaluation showed profound bilateral blepharospasm, photophobia and a mucous secretion until the 7th day after surgery; corneal vascularization starting in the limbus was observed as early as the 3rd day in both eyes; opacification and vascularization were more intense in the treated cornea on days 15 and 30 after surgery. Vascularization was still evident on the 60th day, and looked similar in treated eyes and control eyes. The histologic evaluation showed a complete bilateral reepithelization and corneal vascularization three days after the surgery; intense vascularization in both eyes on days 15 and 30, that was much more pronounced in the treated cornea; and a bilateral predominance of polymorphonuclear cells until day 15, and mononuclear cells on day 30. Intact epithelium and stroma with new vessels, as well as graft absorption, in both eyes, were seen on the 60th day. The lamellar keratoplasty with equine pericardial graft was an effective model to study the inflammatory kinetics and corneal vascularization phenomenon. In this study, cyclosporine A did not inhibit corneal vascularization and it did not interfere in the corneal cicatrical process.

Key words: cyclosporine, cornea, pericardium, vascularization, keratoplasty, dogs.

RESUMO

Foram avaliados os efeitos da ciclosporina A a 2% sobre a córnea de cães submetidos à ceratoplastia lamelar experimental com implante de pericárdio de equino. Dez cães foram divididos em grupos para estudo aos três, sete, 15, 30 e 60 dias de pós-operatório, recebendo implantes bilaterais e em seguida aplicação da pomada com ciclosporina A a 2% no olho esquerdo (olho tratado) e somente a base da mesma pomada no olho direito (olho controle), duas vezes ao dia. A avaliação oftalmológica observou-se, bilateralmente, blefarospasma, fotofobia e secreção mucosa evidentes até 7 dias de pós-operatório; início de vascularização a partir do limbo aos três dias, bilateralmente; opacidade e vascularização mais intensos em córnea esquerda aos 15 e 30 dias; vascularização ainda evidente, bilateralmente, com aspecto semelhante aos 60 dias. À avaliação histopatológica observou-se, bilateralmente, completa reepitelização e vascularização aos três dias de pós-operatório; vascularização intensa aos 15 e 30 dias, bilateralmente, porém acentuada na córnea esquerda; predominio de polimorfonucleares até os 15 dias e mononucleares aos 30 dias, bilateralmente; epitélio e estroma íntegros com vasos ainda evidentes e implante incorporado ao estroma, bilateralmente, aos 60 dias. A utilização do implante de pericárdio de equino em ceratoplastias lamelares em cães constitui uma técnica eficaz, aplicável ao estudo da cinética inflamatória e fenômeno de vascularização. A ciclosporina A a 2% não inibiu ou diminuiu a formação vascular na córnea e também não interferiu no processo cicatricial neste estudo.

Palavras-chave: ciclosporina, córnea, pericárdio, vascularização, ceratoplastia, cães.

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INTRODUCTION

Corneal ulceration is very frequent and represents one of the most important eye diseases in veterinary ophthalmology. It requires quick intervention, even in the most superficial lesions, in order to avoid serious complications and preserve the vision (BROOKS, 2000). Many therapeutic attempts have been made over the years to repair the cornea in order to maintain its structure and, if possible, its transparency and function. Surgical therapeutic methods are very important in corneal ulcer cases with important corneal tissue loss, in which the only alternative is tissue substitution (BARROS et al., 1995).

Equine pericardium was successfully used in the correction of entropion and to fill the orbital cavity of dog following enucleation (BARROS et al., 1985), as well as in the repair of the sclera and third eyelid in dogs (BARROS et al., 1990). In cases of penetrating sclerokeratectomy due to limbal neoplasia (BARROS et al., 1995); keratectomy with partial iridectomy due to a perforating lesion (BARROS et al., 1995) and in lamellar and penetrating keratoplasty in dogs, it was also effective because of the perfect integration between the pericardium and the cornea and the absence of rejection phenomena (KAVINSKI, 1980; BARROS et al., 1995; BARROS et al., 1997). Although the results are good with the use of pericardium for the anatomic preservation of frail or perforated cornea, the occurrence of cicatrical reaction with leucoma and corneal vascularization may be a limiting factor to its use, especially in axial or extensive wounds, for vision may be affected.

The present trial was performed in order to evaluate the lamellar keratoplasty with an equine pericardial graft preserved in glycerin and the effects of topical cyclosporine A on these corneas of dogs, because it does not interfere in the cicatrical process of the cornea (READ, 1996); because of its potential for decreasing corneal vascularization (EPSTEIN et al., 1987; LIPMAN et al., 1992; NORRBY, 1992; BENELLI et al., 1997) and because of the beneficial results observed using topical cyclosporine A on several corneal inflammatory affections (KASWAN et al., 1989; JACKSON et al., 1991; MORGAN & ABRAMS, 1991; READ, 1996; STEVENSON et al., 2000).

MATERIALS AND METHODS

All animals involved in this study were maintained and handled in accordance with the ARVO Statement for the Use of Animals in Ophthalmic and Vision Research. Ten mixed-breed male dogs, weighing around 15kg were divided in five groups of two dogs each to be studied after 3, 7, 15, 30 and 60 days of the surgery. Preoperative ophthalmic examination included slit-lamp biomicroscopy, fluorescein strip test, Schirmer tear test, TonoPen applanation tonometry and direct ophthalmoscopy. The anesthetic protocol included premedication using intramuscular 0,2mg.kg⁻¹ of 0,2% acepromazine; induction using intravenous 12,5mg.kg⁻¹ of Tiopental; and maintenance using 1,0-2,5% of halothane in semiclosed circle rebreathing system.

As an immediate post-surgical procedure, a 2% cyclosporine A ointment* was applied to the left eye (treated eye) and only the base of the ointment* to the right eye (control eye), twice a day, until the end of the evaluation period. No other topical or sistemic drug was used during these periods.

The animals were submitted to ophthalmic examination with slit-lamp biomicroscopy daily, in which parameters such as blepharospasm, photophobia, ocular secretion, corneal opacification, corneal vascularization and pigmentation were analyzed. Viability, permanence and transparency of the grafts were also observed. These parameters were classified as absent, minor, moderate, intense and major. At the end of the period of study, animals were euthanized and their eyes enucleated and fixed in a 10% buffered formaldehyde solution. Sections were stained by H & E, PAS and Masson’s trichrome stain, and examined by light microscopy. Aspects such as reepithelization, edema, corneal vascularization and polymorphonuclear and mononuclear leukocyte infiltration were analyzed.

RESULTS

Clinical aspects

Blepharospasm and photophobia were observed in all animals, and were intense up to the 7th postoperative day; they were minor on day 15 and bilaterally absent from the 30th day on. In relation to the 10% buffered formaldehyde solution. Sections were stained by H & E, PAS and Masson’s trichrome stain, and examined by light microscopy. Aspects such as reepithelization, edema, corneal vascularization and polymorphonuclear and mononuclear leukocyte infiltration were analyzed.

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type, evident from the 3rd to 15th postoperative day, and absent in the other observation periods, being moderate on the 7th day.

In relation to corneal opacification, it was minor adjacent to the graft on the 7th postoperative day;

moderate in the control cornea (Fig. 1A), and intense in the treated cornea on day 15 (Fig. 1B). It was still present bilaterally on day 30 (Figs. 1C and D), and it was reduced on the 60th postoperative day. A leucoma in the graft area was similar in both corneas.

Corneal vascular formation began at the limbus of the upper temporal region bilaterally on the 3rd postoperative day; it was more evident bilaterally on day 7, intense in the control cornea (Fig. 1A), and major, reaching the graft, in the treated cornea on day 15 (Fig. 1B); it was also intense bilaterally on day 30 (Figs. 1C and D); however, it was still prominent in the treated cornea (Fig. 1D). The appearance of both corneas was similar on the 60th postoperative day, when corneal vessels, some suture granulomas, and excellent transparency of the cornea and partial transparency of the implant were observed. None of the corneas presented pigmentation.

Histopathologic findings

Complete bilateral reepithelization was observed on the 3rd postoperative day, with the presence of regenerative phase epithelium, and acanthosis in the transition cornea/graft. On day 7, the basal layer of the epithelium was starting to be defined, and on day 15 it was stratified, presenting almost normal width. The stroma had fibroblastic hypertrophy with an increase in the collagenous matrix and bilateral inflammatory infiltrate with predominance of polymorphonuclear cells. On day 30, the epithelium presented bilaterally several polyedrical and cylindrical layers (Figs. 2A and B), and complete

Figure 1 – A) Control cornea at day 15: mild vascularization and light edema close to the implant; B) Treated cornea at day 15: heavy vascularization and strong edema at the implant site.

C) Control cornea at day 30: intense vascularization surrounding the implant; D) Treated cornea at day 30: intense vascularization over and surrounding the implant, discreet opacification close to the implant, exuberant granulation.
regeneration occurred by the 60th postoperative day when remains of the graft were observed among the reconstituted stroma suggesting an absorption process.

Edema adjacent to the graft was observed bilaterally on the 7th postoperative day, and was more evident on day 15 when disarrangement of the lamellae of the stroma with presence of fibroblasts and inflammatory cells could be observed. There was an increase in the width of the cornea in the keratoplasty site when compared to the intact cornea. In this phase, epithelial edema was also observed, and it was characterized by vacuolization which was also evident on the 30th postoperative day. These events disappeared on day 60.

Corneal vascular formation occurred associated with the inflammatory events, and vessels were observed bilaterally from the limbus towards the keratoplasty site on the 3rd postoperative day. On day 7, these vessels took up most of the stroma, bilaterally. On day 15, the phenomenon was intense in both corneas, with vessels in the interface graft/stroma, inside the graft and in the adjacent stroma. On day 30, vascularization was still intense bilaterally with vessels inside the graft and in the adjacent stroma (Figs. 2A and B). The vessels, however, took a greater part of the stroma of the treated corneas of the animals in this group (Fig.2B). On the 60th postoperative day, vessels were still present in similar amount bilaterally.

Inflammatory cell infiltrate was characterized, bilaterally, by the early infiltration of polymorphonuclear cells in the adjacent stroma, in the interface and inside the graft. It was present up to the 15th postoperative day associated with the intense vascularization in this phase. A major inflammatory setting was observed, with intra-epithelial inflammation in the treated cornea of the animals of the 15th day group. On day 30, a chronic inflammatory reaction was observed with the bilateral presence of mononuclear cells and rare polymorphonuclear cells in the adjacent stroma and inside the graft (Figs. 2A and B). On day 60, a discrete mononuclear cell infiltrate was observed in the corneal stroma bilaterally.

**DISCUSSION**

Photophobia and blepharospasm were evident immediately after the surgery and occurred due to stimulation of the sensors epithelium and stroma nerve terminations (WARING, 1984) produced by the surgical procedure and the presence of the graft, as well as the contact of the sutures with the eyelid conjunctiva. This is a frequent finding in keratoplasties with lamellar and penetrating grafts, in which there is no burying of the suture knots. Secretion was predominantly mucous, which is present in conjunctival and corneal processes, due to the stimulation of caliciform cells; it was, therefore, expected in such a surgical procedure (STARTUP, 1984). The characteristic of the ocular secretion was not influenced by the topical use of cyclosporine A, which presents specific immunosupressing action on T lymphocytes, and did not interfere in the innate immune response by phagocytic cells (DIASIO & LOBUGLIO, 1996), what explains its uncommon association with opportunistic infections (RYFFEL, 1989).
In relation to the cicatricial process, epithelial repair was quick, and characterized by sliding and mitosis of epithelial cells in a centripetal direction (PEIFFER JR. et al., 1999). Acanthosis observed in the extremities of the implant in this study confirms the mitotic capacity of these cells (PEIFFER JR. et al., 1999).

The repair of the stroma occurs more slowly, through fibroblastic proliferation. In the final phase, fibroblastic tissue decreases and collagenous fibers are reorganized in a parallel fashion, determining the intensity of the scar and the subsequent transparency of the cornea. This was observed in the later periods of evolution, with the bilateral incorporation of equine pericardium to the corneal stroma. Complete and bilateral repithelization on the 3rd postoperative day confirms that cyclosporine A does not present adverse effects on the epithelium, during the cicatricial process in the cornea (BEHRENS-BAUMAN et al., 1986; BELIN et al., 1990).

Edema is the main responsible factor for the loss of corneal transparency (PEIFFER JR. et al., 1999) and thin vesicles are found when it affects the epithelium. When affecting the stroma, it produces diffuse mist, and is characterized by an increase in cornea thickness (SPENCER, 1996); this was bilaterally observed on day 15 and 30.

The degree of opacification is not only associated with the edema, but also to the number and concentration of inflammatory cells, to the presence or absence of neovessels and to enzyme-induced stroma necrosis (SPENCER, 1996). This may explain the higher intensity of cornea opacification on days 15 and 30, when intense vascularization and important inflammatory infiltrates occurred, mainly in the treated cornea.

In relation to vascular phenomena, vascularization was observed bilaterally on the 3rd postoperative day. This may be associated with the hypothesis that angiogenic factors or cytokines are released in the first hours after the surgical procedure (NAUMANN & SAUTTER, 1988; KOCH, 1992) with vessels in the cornea 2 to 4 days after the loss of epithelium and stroma (PEIFFER JR. et al., 1999).

Conical vascularization favors the cicatricial process, including tissue remodeling (EPSTEIN et al., 1987). On the other hand, it is not desirable due to the loss of corneal transparency (PEIFFER JR. et al., 1999). Several trials have been performed in order to evaluate aspects related to corneal vascularization (SUNDERKOTTTER et al., 1991a; REHANY & WAISMAN, 1994; KENYON et al., 1996; BENELLI et al., 1997; BARROS et al., 2000) and several therapeutic schemes were attempted in order to decrease or inhibit it, mainly the topic or systemic administration of cyclosporine A at different concentrations (LIPMAN et al., 1992; REHANY & WAISMAN, 1994; BENELLI et al., 1997).

Cyclosporine A is a cyclic polypeptide from the fungus Tolypocladium inflatum Gans, and presents the ability to inhibit the activation and proliferation of T-helper lymphocytes, by blocking the release of lymphokines (DIASTO & LOBUGLIO, 1996). It was successfully used in topical treatment, decreasing the neovascularization in the cornea of dogs presenting keratoconjunctivitis sicca (KASLAN et al., 1989). It also reduced corneal vascularization that developed in mice after an intracorneal injection of interleukin-2, when intramuscularly administered (LIPMAN et al., 1992). Cyclosporine A given subcutaneously also has been found to suppress new vessel formation induced in the mesenteric-window assay in rats (NORRBY, 1992). It also is reported to produce clinical improvement in psoriasis and certain epithelial neoplasms in which angiogenesis is prominent (NORRBY, 1992). It decreased the growth of vessels in the cornea of rats after xenotransplantation and chemical cautery using silver nitrate, when a 4% solution was topically and systematically administered. However, it was not effective in the blockage of new vessels growth induced by the insertion of an angiogenic disc in the subcutaneous tissue, when systematically or intradisc administered (BENELLI et al., 1997).

Topical administration of cyclosporine A does not produce a measurable systemic concentration or contralateral ocular immunosupression (FOETS et al., 1985; BELIN et al., 1990; JACKSON et al., 1991; BENELLI et al., 1997; STEVENSON et al., 2000) enabling each animal to be its own control (JACKSON et al., 1991).

Cyclosporine A 2% did not inhibit or decrease the formation of vessels in this trial, although it may have showed anti-angiogenic or angiostatic activity in other studies (LIPMAN et al., 1992; NORRBY, 1992; BENELLI et al., 1997). The mechanisms by which cyclosporine A reduces or inhibits angiogenesis in some, but not in all models, remain unrecognized (BENELLI et al., 1997). One of the hypothesis is that it may be related to the nature and the permanence of the injury; to the quality and time of the surgical procedure and to the kind of suture used (NASISSE, 1985; KERN, 1990). Therefore, the angiogenic response may vary significantly according to these characteristics, to the species involved, to the type of biological membrane used and to its particular angiogenic ability (BARROS et al., 2000).
Cyclosporine A inhibits only the transcription of the genes for interleukin-2 (IL-2) and interleukin-4 (IL-4) (RIZZO, personal communication, 2000). Thus, the other lymphokines (DIASTO & LOBUGLIO, 1996; ROOK & BALKWILL, 1999; GOLDSBY, 2000a) may produce angiogenic effects, what could explain the similarity between the control and treated corneas on the 3rd, 7th and 60th postoperative day. Cyclosporine A also increases the expression of TGF-β, (PRASHAR et al., 1995; KHANNA et al., 1997; SHIN et al., 1998) which presents evident angiogenic effects (ROBERTS et al., 1986; SUnderkotter et al., 1991b).

It is important to note that, during the cicatricial process, epithelial cells, keratocytes and stromal fibroblasts may also synthesize and release growth factors with angiogenic activity (SWANK & Hosgood, 1996).

As it is know that several cells take part in the angiogenesis phenomenon, and although lymphocytes are relevant in this process, macrophages are also very important due to their intense secretion of factors that may be directly or indirectly angiogenic, contributing with more than one factor at each stage of the angiogenesis (POLVERINI et al., 1977), what may compensate for the angiogenic effects of the lymphokines blocked by cyclosporine A.

The occurrence of a more intense vascularization process on days 15 and 30 in the corneas treated with cyclosporine A 2% may be due to the combined action of the angiogenic cytokines secreted by macrophages and those released by the lymphocytes and not affected by cyclosporine, together with the increase of TGF-β expression (PRASHAR et al., 1995; KHANNA et al., 1997; SHIN et al., 1998). Another possibility is that, because of the inhibition of the synthesis and release of IL-2 and IL-4, and of the increase of TGF-β expression, both caused by cyclosporine A, there would be an unbalance between angiogenic and anti-angiogenic factors. This balance is extremely important for the control of angiogenesis, which is dose-dependent (STRIETER et al., 1992). Therefore, it may affect the secretory pattern of lymphocytes and macrophages, what would result in the increase in angiogenic cytokine release by these cells.

In relation to the inflammatory reaction, cell infiltrate was characterized by the invasion of the site with polymorphonuclear and mononuclear lymphocytes. Neutrophils were the most prevalent in the beginning of the inflammatory process, followed by monocytes and lymphocytes that also migrated to the site of injury. The early inflammatory cell infiltrated is considered a pre-requisite for angiogenesis induction (SUNDERKOTTER et al., 1991a), for it acts in the degradation of proteins from the extracellular matrix and would justify the beginning of the vascularization process as early as the 3rd postoperative day, bilaterally.

An important stage in leukocyte migration is the fixation of the cells to the vascular endothelium, as a consequence of the interaction between the molecules on the membranes of the leukocytes with the corresponding molecules on the activated endothelium (SPENCER, 1996), controled by the expression of the adhesion molecule E-selectin on the surface of the endothelium (KOCH et al., 1995). It has recently been suggested that this molecule presents direct angiogenic effect on the endothelial cells (KOCH et al., 1995) and that cyclosporine A may promote the low regulation of the expression of this molecule by the vascular endothelium, what would suppress the growth of vessels (BENELLI et al., 1997). In the present trial, however, inflammatory cell infiltrate was observed as early as the 3rd postoperative day, and was bilaterally evident up to day 60. This would confirm that cyclosporine A did not present any effect on this adhesion molecule. It is important to emphasize that cytokines TNF-α and IL-1, which stimulate the secretion of E-selectin by the endothelium (KOCH et al., 1995; GOLDSBY, 2000b) are not affected by cyclosporine A.

CONCLUSIONS

Lamellar keratectomy with equine pericardium graft preserved in 98% glycerin was a feasible model for the study of inflammation kinetics and angiogenesis phenomenon in the cicatricial reaction of the cornea. There was intense vascularization and an important fibroblast response, with the permanence of cicatricial leucoma and corneal vessels on the 60th postoperative day.

Cyclosporine A 2% did not interfere in the corneal cicatricial process generated by keratoplasty with lamellar equine pericardium graft. It also did not decrease the corneal vascularization process in this study.

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ACQUISITION

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