The effect of adrenalectomy on neutrophil and monocyte migration into rat peritoneal and pleural cavities was investigated. Carrageenin- or thioglycollate-induced neutrophil emigration into both cavities was enhanced by adrenalectomy. In contrast, monocyte migration into peritoneal cavities induced by these two stimuli was significantly decreased. In pleural cavities, adrenalectomy enhanced the monocyte migration induced by carrageenin but had no effect on that induced by thioglycollate. Administration of physiological doses of glucocorticoids reversed the effect of adrenalectomy on monocyte migration by both stimuli into both cavities. The results support the hypothesis that endogenous glucocorticoids negatively control neutrophil migration independently of the site or type of stimulus. Their role in monocyte migration is, however, dependent on the site of injury and on the type of inflammatory stimulus. There is no obvious explanation for the divergent influence of endogenous glucocorticoids on the monocyte emigration into peritoneal and pleural cavities observed with different stimuli.

**Key words:** Adrenalectomy, Endogenous glucocorticoids, Inflammation, Monocyte migration, Neutrophil migration

**Introduction**

It is generally assumed that migration of phagocytes (neutrophils or monocytes) into the various body cavities or tissues is governed by factors released by inflammatory stimuli. This assumption is supported by a great number of studies on neutrophil migration, including those which show that during the initial stages of the inflammatory process an increase in glucocorticoid (GCC) secretion by the adrenal glands inhibits neutrophil migration, independently of the site of inflammation. In acute inflammation, GCC secretion is induced by the hypothalamic-pituitary-adrenal (HPA) axis through the influence of factors released in the inflamed area. Among these factors, cytokines such as interleukin-1 (IL-1), IL-6 or tumour necrosis factor (TNF) have been shown to be involved. IL-1, in addition to its effect on the HPA axis, also has an effect on the adrenal gland, to directly stimulate GCC secretion. The enhancement of neutrophil migration by inflammatory stimuli in adrenalectomized (ADX) or in animals pretreated with GCC antagonists such as RU 486 may be explained by the diminished capacity of the body cells to release lipocortins or by facilitation of the release of chemotactic cytokines. The lipocortins, which belong to the superfAMILY of calcium binding proteins termed annexins, have anti-phospholipase A activity. GCC induce the release of these proteins and therefore reduce the biosynthesis of leukotrienes thought to be involved in the control of neutrophil migration. In this context, it is interesting to note that in ADX animals there is a reduction in the amounts of both lipocortin-1 and lipocortin-1 mRNA in the body fluids, and that the physiological replacement of corticosterone restores the normal levels of both these components. Moreover, in ADX animals, Perret et al. demonstrated that macrophages release more IL-1 and prostaglandin E than do macrophages from sham-operated (SHO) ones. Conversely, GCC inhibited the in vivo neutrophil recruiting activity on pro-inflammatory cytokines such as interleukin IL-1, IL-6, IL-8 and TNF.

To date, the studies investigating the role of endogenous GCC on leukocyte migration into inflammatory sites have focused on neutrophil migration. In this paper, we have compared the effect of adrenalectomy on neutrophil and mononuclear phagocyte migration into two different sites of inflammation (the pleural and peritoneal cavities) induced by two different stimuli (carrageenin or thioglycollate). The results confirm the observations of other authors that adrenalectomy potentiates neutrophil migration into peritoneal or pleural cavities. However, with regard to monocyte migration, the response to adrenalectomy depended heavily on
the cavity tested. Adrenalectomy reduced the migration into peritoneal cavities induced by both carrageenin (Cg) and thioglycollate (Tg), but had no effect or enhanced the monocyte migration into pleural cavities induced by Tg and Cg, respectively.

Materials and Methods

Drugs and chemicals: Carrageenin and thioglycollate were purchased from Marine Colloids Inc. and Merck Laboratories, respectively. The steroid hormones oestr-a-1,3,5(10)-trien-3,17 diol (β-oestradiol) and corticosterone were obtained from Sigma Chemical Company, St Louis, MO. Dexamethasone 21-acetate (Decadronal) was from Merck Sharp & Dohme. All other reagents were purchased from Sigma Chemical Company (USA).

Animals: Adult male Wistar rats weighing 150–180 g obtained from the Central Animal House of the Faculty of Medicine (Ribeirão Preto) were used in all experiments. The animals received water and food ad libitum.

Bilateral adrenalectomy: The animals were surgically prepared 7 days before the experiments. Following light ether anaesthesia, the rats of one group had both of their adrenal glands completely removed (ADX animals), while those of a second group underwent the same process to induce surgical stress, but were left with their glands intact (SHO animals). The ADX rats received an additional supply of saline solution (NaCl, 0.9% w/v) to drink. At the end of the experiments, the ADX animals in which there was macroscopic evidence of remnant adrenal tissue were rejected. The inflammatory stimuli were injected on the seventh day after surgery.

Experimental models of inflammation: The neutrophil and mononuclear cell migration into peritoneal and pleural cavities were investigated in SHO and ADX animals. In some experiments, the ADX animals were treated with steroid hormones.

(a) Peritoneal cavity. Cg resuspended in 3 ml of phosphate buffered saline (PBS) or 5 ml of thioglycollate (Tg, 3% w/v), was injected intraperitoneally. The migration of neutrophils was analysed 4 h after the injection of Cg (300 µg/rat). While the migration of mononuclear cells was analysed 48, 72 or 96 h after the injection of Cg (500 µg/rat) or Tg (150 mg/rat).

(b) Pleural cavity. In pleural cavities, neutrophil migration was also analysed 4 h after the intrapleural injection of Cg (300 µg/ml/rat) and mononuclear cell migration was analysed 48, 72 or 96 h after Cg (500 µg/ml/rat) or Tg (40 mg/ml/rat) injection.

(c) Collection of exudates. At selected intervals, the cells which had migrated were harvested from the peritoneal and pleural cavities. For this, the animals were anaesthetized with ether and exsanguinated. Both naïve and stimulated peritoneal and pleural cavities were washed with 10 ml and 5 ml of heparinized PBS (5 U/ml), respectively. In some experiments, the peritoneal cavities were washed with 10 ml of trypsin diluted in PBS (TRY, 1.25% w/v). Total and differential cell counts of the collected washings were estimated as previously described by Souza and Ferreira. The characterization of macrophages (esterase positive cells) performed using a nonspecific esterase staining technique described by Yam et al. The results were expressed as the number of mononuclear or esterase positive cells per cavity.

Leukogram count: Total leukocyte, neutrophil and mononuclear cell counts were performed on blood samples obtained from the tail vein of ADX or SHO rats which had or had not been treated with Tg (150 mg/5 ml/rat, i.p.) 48 h before. The results were recorded as the number of cells per ml of blood.

Administration of steroid hormones: Subcutaneous doses of 2, 20, 200, 2000 and 6000 µg/kg of oestradiol (ST) and corticosterone (CT) dissolved in peanut oil, and dexamethasone (DX) dissolved in sterile saline, were injected daily from the sixth day after surgery until the day mononuclear cell migration was measured. The control animals were injected with vehicle solution.

Statistical analysis: The results are expressed as the mean ± standard error of the mean (S.E.M.) and statistical comparisons between the SHO and ADX groups were done using Student's t-test. p values < 0.05 were considered to be statistically significant.

Results

Neutrophil migration into peritoneal cavities measured 4 h after Cg injection was significantly enhanced by adrenalectomy (Fig. 1A). In contrast, mononuclear phagocyte migration evaluated 48, 72 and 96 h after Cg injection was significantly reduced in ADX rats compared with SHO animals (Fig. 1B). A similar pattern of mononuclear phagocyte migration was observed when Tg instead of Cg was injected into the abdominal cavities (Fig. 2).

In the pleural cavities, both neutrophil and mononuclear phagocyte migration induced by Cg were enhanced in ADX animals (Fig. 3A and B). However, the mononuclear phagocyte migration into these cavities induced by Tg was not affected by adrenalectomy (Fig. 4). The number of resident mononuclear phagocytes present in the non-stimulated peritoneal or pleural cavities was not modified by adrenalectomy (cross-hatched bars in all figures).
Adrenalectomy and monocyte migration

FIG. 1. Adrenalectomy potentiates carrageenin induced neutrophil migration but inhibits mononuclear cell migration into peritoneal cavities. The bars show the number of neutrophils (panel A) and the number of mononuclear cells (panel B) in naïve cavities (cross-hatched bars) and in cavities pre-stimulated (open bars) with Cg (300 µg/cavity, panel A) or (500 µg/cavity, panel B), in sham-operated (SHO) or adrenalectomized (ADX) rats. The values are the means ± S.E.M. of the number of animals indicated above each bar. Asterisks indicate significant differences between the SHO and ADX groups (p < 0.05).

FIG. 2. Adrenalectomy inhibits thioglycollate induced mononuclear cell migration into rat peritoneal cavities. The bars show the number of mononuclear cells collected from naïve cavities (cross-hatched bars) or from cavities pre-stimulated with Tg (150 mg/cavity) 48, 72 and 96 h before (open bars) in sham-operated (SHO) or adrenalectomized (ADX) rats. The values are the means ± S.E.M. of the number of animals indicated above each bar. Asterisks indicate significant differences between the SHO and ADX groups (p < 0.05).

FIG. 3. Adrenalectomy potentiates carrageenin induced neutrophil and mononuclear cell migration into rat pleural cavities. The bars show the number of neutrophils (panel A) and the number of mononuclear cells (panel B) in naïve cavities (cross-hatched bars) and in cavities pre-stimulated (open bars) with Cg (300 µg/cavity, panel A) or (500 µg/cavity, panel B), in sham-operated (SHO) or adrenalectomized (ADX) rats. The values are the means ± S.E.M. of the number of animals indicated above each bar. Asterisks indicate significant differences between the SHO and ADX groups (p < 0.05).

FIG. 4. Adrenalectomy does not alter the thioglycollate induced mononuclear cell migration into rat pleural cavities. The bars show the number of mononuclear cells collected from naïve cavities (cross-hatched bars) and from cavities pre-stimulated with Tg (150 mg/cavity) 48, 72 and 96 h before (open bars) in sham-operated (SHO) and adrenalectomized (ADX) rats. The values are the means ± S.E.M. of the number of animals indicated above each bar. There were no significant differences in the migratory responses of mononuclear cells in the SHO and ADX groups.

Regardless of whether they were pre-stimulated or not with Tg, all ADX animals had higher leukocyte, neutrophil and mononuclear cell counts than did SHO rats (Table 1).

Daily treatment of ADX animals with DX (2 – 2000 µg/kg) resulted in a bell-shaped curve in which a low dose of DX (20 µg/kg) restored the Tg-induced mononuclear phagocyte migration into peritoneal cavities to levels similar to those observed in SHO animals, while a pharmacological dose (2 000 µg/kg) significantly enhanced the reduction in mononuclear phagocyte migration. Daily treatment with CT also dose dependently reverted the mononuclear cell migration into the peritoneal cavities of ADX animals. At none of the doses studied did ST have any effect on the mononuclear phagocyte migration in ADX rats (Fig. 5).

Discussion

The results presented above agree with previous experiments showing that adrenalectomy or pretreatment with RU 486 enhances the neutrophil migration induced by inflammatory stimuli such as Cg into pleural cavities and subcutaneous air pouches. In addition, a greater inflammatory exudation in ADX animals (data not shown) was observed. These responses are in line with the suggestion that during the acute inflammatory responses the release of GCC by the adrenal cortex partially down-regulates neutrophil migration and exudation via a reduced release of chemotactic cytokines or arachidonic acid metabolites.

Paradoxically, adrenalectomy reduced monocyte migration into peritoneal cavities by both stimuli utilized but did not affect or enhance migration into

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Table 1. Leukogram of sham-operated (SHO) and adrenalectomized (ADX) rats in the presence or absence of thioglycollate (Tg) pre-stimulation

| Cell type            | Cell countsa | Non-stimulated | Tg stimulated |
|----------------------|--------------|----------------|---------------|
|                      | SHO          | ADX            | SHO           | ADX           |
| Total leukocytes     | 9.7 ± 0.7    | 22.1 ± 2.3*    | 7.5 ± 0.9     | 19.1 ± 0.5*   |
| Neutrophils          | 3.8 ± 1.0    | 7.0 ± 1.1*     | 1.9 ± 0.3     | 6.6 ± 1.0*    |
| Mononuclear cells    | 5.7 ± 2.2    | 14.1 ± 2.2*    | 5.0 ± 0.5     | 12.5 ± 0.5*   |

* Thioglycollate (150 mg/rat) was injected intraperitoneally 48 h before.

Expressed as the number of cells x 10^6 ml^-1. The values are means ± S.E.M. of six observations.

* Significant difference between SHO and ADX groups (p < 0.05, Student’s t-test).

the pleural cavities. The reduction of monocyte migration into the peritoneal cavities of ADX animals resulted from the absence of circulating GCC since it was reverted by daily treatment of the ADX animals with low doses of dexamethasone or corticosterone, but not by oestradiol. The reduction of mononuclear phagocyte migration into the peritoneal cavity of ADX animals was not caused by monocytopenia since adrenalectomy produced intense leukocytosis with monocytosis in animals which had or had not been stimulated with Tg. The reduction in the number of migrating monocytes in ADX animals could be due to an increased adherence of these cells to the abdominal wall. However, this possibility can be discarded since the addition of trypsin to the washing fluid did not increase the number of migrating monocytes in ADX animals (data not shown). Pharmacological (200 and 2,000 µg/kg) doses of dexamethasone which are able to inhibit neutrophil migration29,30 inhibited monocyte migration into the peritoneal cavities of SHO rats (data not shown) and enhanced the reduction in monocyte migration already observed in ADX animals (Fig. 5).

The differential effect of the stimuli in both cavities excludes a possible action of GCC on the monocytes. A simple explanation for the effect of adrenalectomy on monocyte migration into peritoneal cavities is that GCC, by a permissive effect, at physiological concentrations facilitates the release of chemotactic factors for monocytes by resident peritoneal cells. Should this be the case, this permissive effect would be under the influence of the microenvironment, since adrenalectomy had either no effect or enhanced the migration into pleural cavities induced by inflammatory stimuli. We have no explanation for the selective action of Tg or Cg on monocyte migration into the pleural cavities of ADX animals. This enhancement of monocyte migration could be abolished by daily treatment of the ADX animals with small doses of dexamethasone. Similarly, the treatment of SHO or ADX animals with pharmacological doses of dexamethasone also diminished the migration induced by Cg. These results indicate that pharmacological concentrations of GCC inhibit the release of a chemotactic factor for monocytes by pleural or peritoneal cells.

In conclusion, the results confirmed that endogenous GCC play an important role in modulating the acute events of the inflammation, by exerting a negative control on neutrophil migration and exudation which is independent of the site of injury. On the other hand, the effect of endogenously released corticoids on monocyte migration is dependent on the site of injury and on the type of inflammatory stimuli. Furthermore, the results clearly indicate that observations regarding monocyte migration in one tissue or cavity cannot be generalized to other tissues.

References

1. Wilkinson PC. Cellular accumulation and inflammation. In: Dale M, Foreman JC, eds. Textbook of Immunopharmacology (3rd edition). Oxford: Blackwell Scientific Publications, 1995; 209-222.

2. Faccioli LH, Souza GEP, Cunha FQ, Poole S, Pereira SH. Recombinant interleukin-1 and tumor necrosis factor induce neutrophil migration in vitro by indirect mechanisms. Agents Actions 1990; 36: 344-349.

3. Ribeiro RA, Flores CA, Cunha FQ, Pereira SH. IL-8 causes in vivo neutrophil migration by a cell-dependent mechanism. Immunology 1991; 73: 472-477.

4. Flower RJ, Pernet L, Persico P, Salome JA. A comparison of the acute inflammatory response in adrenalectomized and sham-operated rats. J R Soc Med 1996; 87: 57-62.
5. Lase L, Kowai S, Brandon DD, et al. Receptor-mediated effects of glucocorticoids on inflammation: enhancement of the inflammatory response with a glucocorticoid antagonist. J Steroid Biochem 1988; 29: 591-598.

6. Garcia-Leme J, Schapoval ES. Stimulation of hypothalmo-pituitary-adrenal axis by compounds formed in inflamed tissue. Br J Pharmacol 1975; 53: 75-83.

7. Besedovsky H, Del Rey A, Sorkin E, Dinarello CA. Immunoregulatory feedback between interleukin-1 and glucocorticoid hormones. Science 1986; 235: 652-654.

8. Sapienza R, Rivier C, Yamamoto G, Pletscher V, Vale W. Interleukin-1 stimulates secretion of hypothalamic corticotropin-releasing factor. Science 1987; 238: 522-524.

9. Naitech V, Pukata J, Tominga T, et al. Interleukin-6 stimulates the secretion of adrenocorticotropin hormone in conscious, freely moving rats. Biochem Biophys Res Commun 1988; 155: 1459-1465.

10. Sharp BM, Matao SG, Peterton PI, Newton R, Chag C, MacGillen R. Tumor necrosis factor-α is a potent ACTH secretagogue: comparison to interleukin-1. Endocrinology 1989; 124: 3131-3135.

11. Rohl M, Drazenovich KA, Barbooe J, Dinarello CA, Gobb GF. Direct stimulation of adrenal corticotropin by interleukin-1. Surgery 1987; 102: 140-146.

12. Hirsta F. The role of lipocortin in cellular function as a second messenger of glucocorticoids. In: Schleimer RP, Claman HN, Oronsky A, eds. Antimflammatory Steroid Action—Basic and Clinical Aspects. San Diego: Academic Press Inc., 1989; 67-95.

13. Peers SH, Flower RJ. The role of lipocortin in corticosteroid actions. Ann Rev Res 1990; 141: 518-521.

14. Dural D, Prease-Begian M. Glucocorticoids and prostaglandin synthesis: we cannot see the wood for the trees. Prostagland Leukot Essent Fatty Acids 1982; 45: 85-112.

15. Blackwell CJ, Camuccio R, Di Rosa M, et al. Glucocorticoids induce the formation and release of anti-inflammatory and anti-phospholipase proteins into the peritoneal cavity of the rat. Br J Pharmacol 1982; 76: 185-194.

16. Erasla M, Russo Marie F. A purified lipocortin protein blocks the anti-inflammatory effect of glucocorticosteroids in vitro and in vivo. Br J Pharmacol 1989; 97: 1051-1058.

17. Vaidyanathan RN, Frey RJ, Inbaraby M, Gillman MF, Frey SM. Adrenalectomy decreases lipocortin-1 messenger ribonucleic acid and tissue protein content in rats. Biochemistry 1992; 120: 565-591.

18. Bieleski RP, Ezzedine AJ, Flowers RJ. Glucocorticoid and non-glucocorticoid induction of lipocortin (anetra) 1 and 2 in rat peritoneal leucocytes in vitro. Br J Pharmacol 1993; 108: 66-72.

19. Perretti M, Bocheneczi C, Scapigliati G, Parente L. The effect of adrenalectomy on interleukin-1 release in vitro and in vivo. Br J Pharmacol 1989; 98: 1137-1142.

20. Snyder DS, Unanue ER. Corticosterone inhibits murine macrophage l-arginine metabolism and interleukin-1 production. J Immunol 1982; 129: 1803-1805.

21. Staruch MJ, Wood DD. Reduction of serum interleukin-1-like activity after treatment with dexamethasone. J Leukocyte Biol 1985; 37: 193-207.

22. Wargke A, Shyamshang G, Shalby R. Glucocorticoids inhibit the production of IL-6 from monocytes, endothelial cells and fibroblasts. Br J Immunol 1990; 20: 2439-2444.

23. Antilla HSI, Reitamo S, Cenka M, Hurme M. Signal transduction pathways leading to the production of IL-8 by human monocytes are differentially regulated by dexamethasone. Cll Exp Immunol 1992; 89: 509-512.

24. Waga A, Balske O. Glucocorticoids suppress the production of tumor necrosis factor by lipopolysaccharide-stimulated human monocytes. Immunology 1988; 63: 299-302.

25. Barber AE, Coyle SM, Marano MA, et al. Glucocorticoid therapy alters hormonal and cytokine responses to endotoxin in man. J Immunol 1993; 150: 1999-2006.

26. Garcia-Leme J, Farnisky SJ. Hormonal control of inflammatory responses. Mediat Inflamm 1993; 2: 181-198.

27. Souza GE, Ferrelies S. Blockade of interleukin-1 activity by dexamethasone induces the in vivo migration of peritoneal macrophages into the peritoneal cavity. Agents Actions 1995; 47: 97-103.

28. Yam LT, Li CY, Crossby WH. Cytotoxic identification of monocytes and granulocytes. Am J Clin Path 1971; 55: 283-290.

29. Kuribara A, Ouchi K, Tsuchufji S. Reduction by dexamethasone of chemotactic activity in inflammatory exudates. J Clin Invest 1985; 76: 11-16.

30. Tsuchufji S, Kuribara A, Ojima F. Mechanisms of antiinflammatory action of dexamethasone blockade by hydrocortisone methylate and actinomycin D of the inhibitory effect of dexamethasone on leukocyte infiltration in inflammatory sites. J Pharmacol Exp Ther 1984; 229: 257-263.

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