



Revista de Biología Tropical

ISSN: 0034-7744

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Universidad de Costa Rica

Costa Rica

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Edema induced by *Bothrops asper* (Squamata: Viperidae) snake venom and its inhibition by Costa Rican plant extracts
Revista de Biología Tropical, vol. 54, núm. 2, junio, 2006, pp. 245-252
Universidad de Costa Rica
San Pedro de Montes de Oca, Costa Rica

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Inflammatory effects caused by *B. asper* are poorly neutralized by commercial antivenoms used for the treatment of these envenomations (Gutiérrez *et al.* 1981, Lomonte 1985, Gutiérrez and Lomonte 1989). Thus, there is a need to find new therapeutic alternatives to alleviate inflammation, which is an important feature in these envenomations.

Plants have been used successfully for the treatment of many diseases (Senior 1996, Ahlemeyer and Krieglsteim 2003). Ferreira *et al.* (1992) have shown, that extracts and fractions obtained from *Curcuma longa*, used in traditional medicine, have anti-inflammatory, and immuno-modulatory activities. Ethnomedically, many plants have been suggested to have anti-venom activity in several countries. A recent investigation described that at least 578 species of higher plants had some anti-venom action (Martz 1992), so plants constitute an extremely rich source of potential venom-inhibitory substances.

In the present work we investigated the characteristics of the edema induced by *B. asper* venom and the anti-inflammatory activity of the leaves extracts from *U. baccifera* (Urticaceae), *L. speciosa* (Urticaceae), *U. leptophylla* (Urticaceae), *C. nutans* (Asteraceae), and *S. viminea* (Lamiaceae), and the roots of *U. tomentosa* (Rubiaceae).

MATERIALS AND METHODS

Venom. It was obtained from adult specimens of *B. asper* collected in the Pacific Region of Costa Rica and corresponding to pools from more than 40 individuals. After freeze-drying, the venom was maintained at -20°C.

Plant extracts. The plants *U. baccifera*, *L. speciosa*, *U. leptophylla*, *C. nutans*, *S. viminea* and *U. tomentosa* were collected during January and March of 1999 in Cartago, Limón and San José (Costa Rica). The plants were identified at source by Luis Poveda and voucher specimens were deposited at the Herbarium of the Universidad Nacional in the

following numbers JVR 6997, JVR 7001, JVR 6998, JVR 6995. Specimens of *S. viminea* and *U. tomentosa* were deposited at the Herbarium of the Universidad de Costa Rica with the numbers USJ 0057229 and USJ 63390.

Collected material was washed, chopped and dried for 3 days at about 40° C, and was ground to a fine dust with a Wiley-type mill. An infusion (10% w/v) was made at 70°C for 30 minutes. The solution was filtered out and concentrated using a rotary evaporator at 40°C. Finally, all plant extracts were freeze-dried and kept at 5°C until used.

Experimental animals. Adult male Sprague-Dawley rats (*Rattus norvegicus*) with a body weight ranging from 180 to 220 g were used throughout the experiments. These animals were supplied by the Animal Care Unit (Universidad de Costa Rica). The animal protocol was approved by the Committee for the Use and Care of Animals of the University of Costa Rica.

Drugs and chemicals. Indomethacin (Merck), dexamethasone (Merck), diphenhydramine (Merck), sodium chloride (Merck) and sodium bicarbonate (Merck) were used throughout the study.

Evaluation of the of edema induced by *B. asper*. In order to evaluate the edema induced by *B. asper* and establish the challenge-dose, groups of six male rats were injected s.c. in the right hind paw with 100, 50, 25, 10 and 5 µg/50 µl of *B. asper* venom dissolved in 0.15M NaCl, whereas the left hind paw was injected with 50 µl of 0.15M NaCl. The paw volume was measured plethysmographically (Ugo Basile, model 7140, Italy) at 1, 2, 4, 6 and 24 hours after *B. asper* venom injection. In addition, groups of six male rats were injected i.p. with various doses of dexamethasone (1 and 0.5 mg/kg) or with diphenhydramine (50 and 25 mg/kg). One hour later, animals were injected s.c. in the right hind paw with 50 µg/50 µl of *B. asper* venom, whereas the left hind paw was injected

with 50 μ l of 0.15M NaCl. The paw volume was measured as described above.

Anti-inflammatory activity. The anti-edematogenic properties of the extracts were quantified in the rat paw edema model (Di Rosa *et al.* 1971). Groups of six male rats were injected i.p. with either 250 or 500 mg/kg of each extract. One hour later, the animals were injected s.c. in the right hind paw with 50 μ g/50 μ l of *B. asper* venom, whereas the left hind paw was injected with 50 μ l of 0.15M NaCl. The paw volume was measured at 1, 2, 4, 6 and 24 hours after venom injection. The control group was injected only with venom (50 μ g/50 μ l) in the right hind paw and with saline solution in the left paw. A group of six rats were treated with indomethacin (100 mg/kg i.p.) as control for the anti-inflammatory activity. Edema was expressed as percentage of the difference between the left paw and the right paw volumes and compared with venom control.

Statistical analysis. Results are presented as mean \pm S.E.M. (n= 6) and the Student's *t* test was used to determine the significance of the differences between the mean values of two experimental groups. *p* values < 0.05 were considered significant.

RESULTS

Evaluation of the edema induced by *B. asper*. Animals treated with increasing doses of *B. asper* venom, showed a dose-related effect. Maximum effects were obtained with 100 μ g/rat. The dose of 50 μ g was selected as challenge dose for the determination of the anti-inflammatory activity of plant extracts (Fig. 1A). The results obtained with treatment with dexamethasone and diphenhydramine are shown in Fig. 1B and Fig. 1C. Only dexamethasone was able to significantly reduce venom-induced edema.

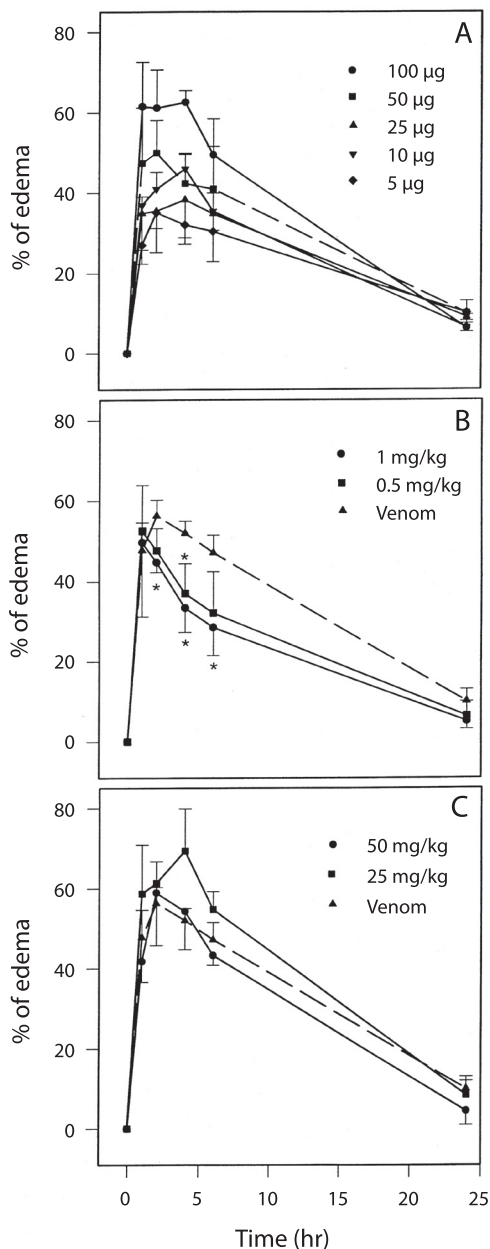


Fig. 1. *Bothrops asper* edema-forming activity in the rat hind paw model. A. Dose-response curve of *B. asper* venom. B. Inhibition of inflammatory effect of *B. asper* venom (50 μ g/50 μ l) by dexamethasone. C. Inhibition of inflammatory effect of *B. asper* venom (50 μ g/50 μ l) by diphenhydramine.

Anti-inflammatory activity. Pretreatment with the extract of *U. tomentosa* diminished considerably the edema-forming activity of *B. asper* venom. The doses of 250 mg/kg and 500 mg/kg showed an antiinflammatory effect at all times, except for the dose of 250 mg/kg at one hour (Fig. 2A). The extract of *C. nutans* showed an important dose-dependent reduction of the edematogenic activity of *B. asper* venom at all times measured. Maximum inhibition was achieved with the highest dose tested (500 mg/kg) (Fig. 2B). Figure 2C shows the time-course of the edema after pretreatment with the extract of *L. speciosa*. In this case, animals evidenced a sedative effect. Those treated with 500 mg/kg died after 4 hours, therefore two more groups were treated with 125 and 62.5 mg/kg. The higher inhibitory effect was observed at a dose of 250 mg/kg (Fig. 2C). Furthermore, a not significant inhibition of edema was observed in animals pretreated with extracts at doses of 125 and 62.5 mg/kg (Fig. 2C). *U. baccifera* induced an important reduction of the edema induced by *B. asper* venom at all times measured (Fig. 3A). A partial inhibition was observed in the animals treated with extracts of *U. leptophylla* and *S. viminea*. Animals pretreated with *U. leptophylla* (250 mg/kg) presented a pro-inflammatory effect at 1 and 2 hours. This effect was not observed in animals treated with the dose of 500 mg/kg (Fig. 3B). In the case of *S. viminea*, a pro-inflammatory effect was observed at 1, 2 and 4 hours with 250 mg/kg. At the other times of measurement the results were similar to those of *U. leptophylla* (Fig. 3C). Indomethacin inhibited the edema at all times after venom injection (results not shown).

DISCUSSION

In this work we demonstrated the characteristics of the edema induced by *B. asper* venom in rats, showing the classical two phases that were described in mice (Chaves *et al.* 1995). We also demonstrated that extracts

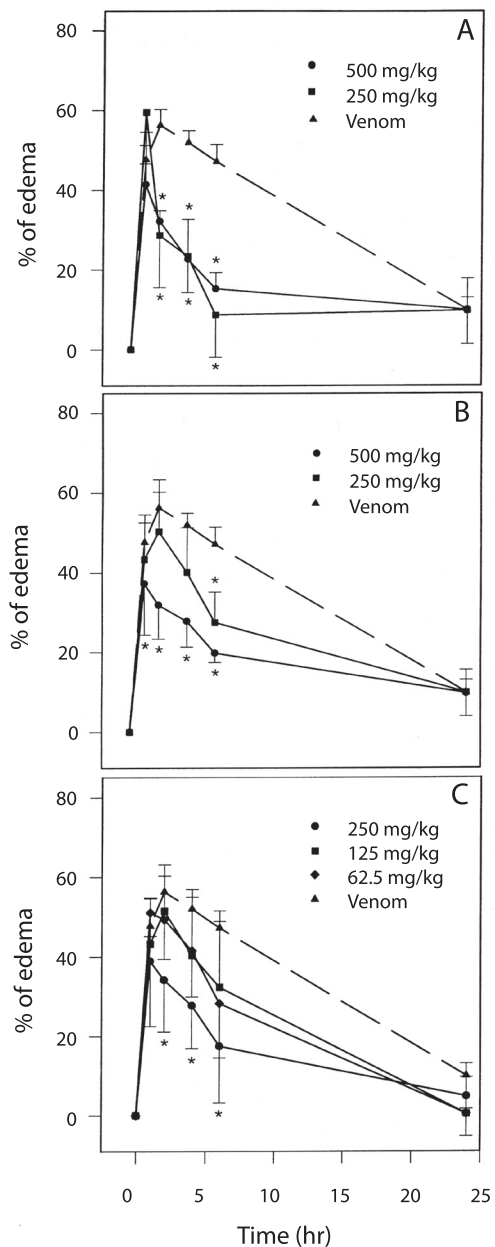


Fig. 2. Anti-inflammatory activity of three plant extracts in the rat hind paw model induced by *B. asper* venom. Groups of rats ($n=6$) were injected i.p with different doses of *U. tomentosa* (A), *C. nutans* (B) and *L. speciosa* (C). Indomethacin (100 mg/kg) was used as anti-inflammatory drug in all experiments. Results are shown as mean \pm SD (* $p < 0.05$).

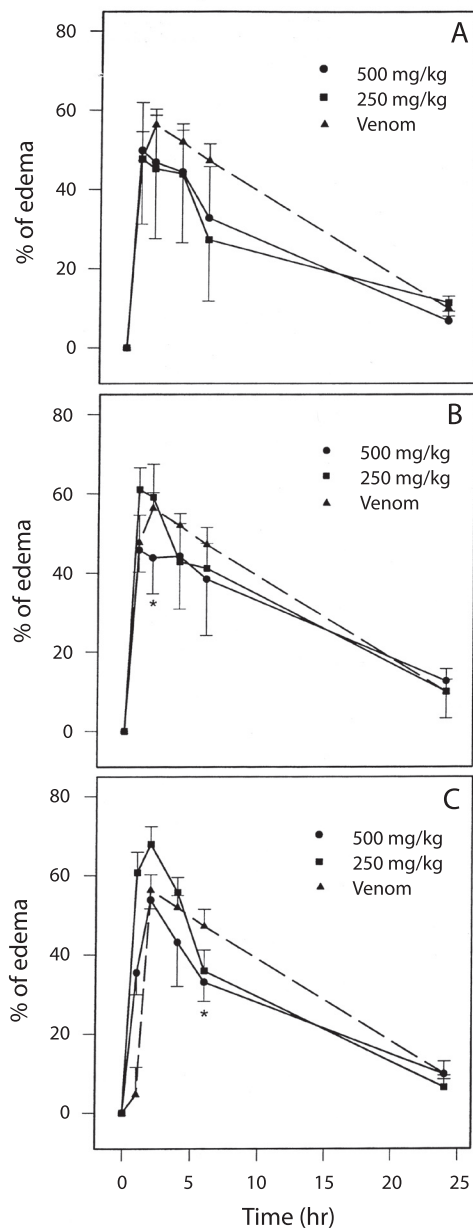


Fig. 3. Anti-inflammatory activity of three plant extracts in the rat hind paw model induced by *B. asper* venom. Groups of rats (n=6) were injected i.p with different doses of *U. baccifera* (A), *U. leptophylla* (B) and *S. viminea* (C). Indomethacin (100 mg/kg) was used as anti-inflammatory drug in all experiments. Results are shown as mean \pm SD (* $p < 0.05$).

of *U. tomentosa*, *C. nutans* and *L. speciosa* were able to neutralize the edema induced by *B. asper* venom. These two last species have been previously found to have anti-inflammatory activity in the carrageenan induced model (Badilla *et al.* 1999a).

One of the consequences of snakebites is local inflammation, especially in the croto-line species (Rosenfeld 1971, Ownby 1982, Gutiérrez *et al.* 1986; Chaves *et al.* 1995). *B. asper*, the most important snake in Central America (Campbell and Lamar 1989, Hardy 1994), induces a striking immediate dose-dependent edema in mice (Lomonte *et al.* 1993, Chaves *et al.* 1995). This snakebite may lead to shock, because of the loss of fluid, and to tissue compression (Garfin *et al.* 1985) which would contribute to the development of cardiovascular disturbances (Carlson *et al.* 1975). There are many inflammatory mediators which participate in the production of edema in a variety of inflammatory conditions (Posadas *et al.* 2000). Among others, histamine, prostaglandins, kinins and leukotrienes, could be implicated in the resulting edema in the case of snake venoms (Trebien and Calixto 1989, Chaves *et al.* 1995).

Edema induced by *B. asper* venom in rats follows the same pattern to the one induced in mice, characterized by a rapid initial first phase produced by mediators such as histamine and serotonin, and a delayed second phase mediated by prostaglandins (Chaves *et al.* 1995). On the basis of dose- response experiments, a dose of 50 μ g/rat was chosen as the challenge-dose, due to its effective inflammatory result without damaging the animal's overall physical integrity, and allowing to show the inhibitory effect of the plant extracts.

Edema seems to be clearly related with prostaglandin production, because an important reduction of the inflammatory effect is induced by dexamethasone, which is a PLA₂ inhibitor and by indomethacin a known inhibitor of cyclooxygenase. No histamine effect was shown in this animal model, because the

doses of diphenhydramine used (50 and 25 mg/kg), were not able to inhibit the first phase of the model.

U. tomentosa, *C. nutans* and *L. speciosa* extracts significantly reduced venom-induced edema. Since inhibition was observed from 1 to 6 hours, it is likely that the compounds in these extracts are acting both the first and second phases of this inflammatory response (Gutiérrez *et al.* 1986, Trebien and Calixto 1989, Chaves *et al.* 1995, Badilla *et al.* 1999b, Posadas *et al.* 2000). *U. tomentosa* has already been reported as a plant that inhibits the carrageenan-induced edema (Kiuchi *et al.* 1983, Aquino *et al.* 1991, Aguilar *et al.* 2002); our studies confirm its effect with *B. asper* venom as inflammation inductor.

The effects found in the first 2 hours with 250 mg/kg of *S. viminea* extracts could suggest an effect on serotonin and histamine liberation, although the effects at 4 to 6 hours suggest an action on prostaglandin production. These effects were not observed at 500 mg/kg. The *U. leptophylla* had a similar behavior.

Although this study was not designed to investigate the mechanism of inhibition, it might be said that aqueous extracts from *U. tomentosa*, *C. nutans* and *L. speciosa* are capable of inhibiting the production of mediators involved in the inflammation process induced by *B. asper* venom, effects that had been found in studies made with plant extracts (Kiuchi *et al.* 1983).

Currently, the mainstream scientific treatment for snakebite envenomations is the parenteral administration of antivenoms (Warrell 1992). Polyvalent (Crotalinae) antivenom is highly effective in the neutralization of the systemic effects in the case of *B. asper* bites, while local injurious effects are neutralized only to a partial extent (Gutiérrez *et al.* 1981, 1985). This is attributed to the quick development of local hemorrhage, edema and myonecrosis after venom injection (Gutiérrez *et al.* 1981, Moreira *et al.* 1992; Lomonte *et al.* 1994, Chaves *et al.* 1995). Therefore, it is necessary to find out complementary treatments aimed at the inhibition of toxins responsible of local tissue

damage. Plant extracts constitute a rich source of novel compounds of potential therapeutic interest in the inhibition of venom toxins. In this regard, previous studies performed with crude extracts and fractions of various plants have shown that they possess anti-snake venom activity (Martz 1992). Recently, Borges *et al.* (2000) demonstrated the ability of extract from *Casearia sylvestris* to inhibit phospholipase A₂, myotoxic, anticoagulant, and edema-forming activities of some snake and bee venoms and of various phospholipases A₂ isolated from these venoms. Our results suggest that the plant extracts investigated contain anti-inflammatory agents that reduce the extent of *B. asper* venom-induced edema.

ACKNOWLEDGMENTS

The authors thank Lorena Chavarría, Alexis Torres, members of the staff of the Department of Pharmacology and Toxicology, School of Pharmacy, Universidad de Costa Rica and Juan Carlos Brenes, of CIPRONA, for their technical assistance. This work was supported by CONICIT, Ministerio de Ciencia y Tecnología, Costa Rica and Vicerrectoría de Investigación (No 817-98-514), Universidad de Costa Rica. We also thank Norman Farnsworth and NAPRALERT, of the University of Illinois in Chicago, for kindly allowing our access to the database.

RESUMEN

Se investigó la capacidad de los extractos de las hojas de *Urera baccifera*, *Loasa speciosa*, *Urtica leptophylla*, *Chaptalia nutans*, *Satureja viminea* y de la raíz de *Uncaria tomentosa* para inhibir el edema inducido por el veneno de *Bothrops asper* por métodos pletismométricos. Los grupos de ratas fueron inyectados intraperitonealmente con varias dosis de cada extracto y una hora más tarde se inyectó veneno por vía subcutánea en la pata trasera derecha de la rata. Se evaluó el edema en distintos intervalos de tiempo. Los resultados muestran que la actividad edematogénica fue inhibida en los animales que recibieron los extractos de raíz de *U. tomentosa*, hojas de *C. nutans* y *L. speciosa*. Los extractos de hojas de *U. baccifera* mostraron leve

inhibición del efecto del veneno. El extracto de hojas de *S. viminea* y en menor grado el de *U. leptophylla* indujeron un efecto pro inflamatorio.

Palabras clave: actividad anti-edematogénica, extractos de plantas, venenos, serpientes, *Bothrops asper*.

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