

# Clinical aspects and consequences of envenoming by a captive Rhinoceros viper (*Bitis nasicornis*) in Hungary

Tamás Malina<sup>a</sup>, László Krecsák<sup>b</sup>

<sup>a</sup> University of Szeged, Department of Systematic Zoology and Ecology, Dugonics tér 13, Szeged, Hungary

<sup>b</sup> Eötvös Loránd University, Department of Systematic Zoology and Ecology, Pázmány Péter s. 1/C, Budapest, Hungary

## Abstract

A case of Rhinoceros viper (*Bitis nasicornis*) bite is reported. The bitten hand distended to the wrist, which was tense and painful, with only mild local livid discolouration manifested around the fang mark. Slight hypertension with moderate tachycardia and temporary coagulopathy were observed. The patient received analgesic and intravenous fluids, antibiotics and anti-tetanus therapy. Use of antiserum was not necessary. The bitten person was treated in the main centre for snake-

bite first aid: in the Toxicological Ward of Erzsébet Hospital of Budapest. We attach importance to the implications of this case because envenoming by *B. nasicornis* being relatively rare in captivity all over the world (particularly in Europe and the USA), as well as in the wilderness in Africa.

*Key words:* snake; first aid; venom; coagulopathy; oedema

## Introduction

Snake-bite accidents caused by different venomous species deserve much more attention worldwide [1], especially since there is limited experience and a paucity of information available for bites (and their consequences) from certain species, e.g., *B. gabonica* or this case of *B. nasicornis*. These are medically important species, mainly due to their highly toxic venom and large venom yield. The Rhinoceros viper (*Bitis nasicornis*) is one of the largest vipers in Africa with its total length 100–150 cm, however the adult size of this species differs across its range [2]. Its distribution spreads from West Africa through Central Africa to western Kenya, thus its geographical range partly overlaps with that of the closely related *B. gabonica* [2, 3].

In Africa, the Puff adder (*Bitis arietans*) is responsible for the majority of bites from the *Bitis* genera [3] due to its large distribution and diverse habitats, while the other two puff adder species mentioned above have hardly ever inflicted accidents in their original habitats [3, 4]. Unfortunately, there is a lack of experience in *B. nasicornis* envenomations and the symptoms caused, as accidents are rare in its natural environment due to its sedentary and nocturnal lifestyle [2, 3]. Data of *B. nasicornis* bites in their natural habitats among

the local human population are often unreliable, as positive identification of the snake seldom occurs. Envenomings are not frequent in captivity either, probably due to the snake's calm behaviour [5]. Only four bites of *B. nasicornis* have been recorded, all in the USA [6–8]: one in Minnesota in 1965 [6], later another victim suffered two bites in St. Louis, Missouri in April then in May in 2002 [7], and one fatal accident occurred in Dayton, Ohio in 2003 [7, 8].

Nowadays the keeping of tropical venomous snakes is in fashion in Hungary [9]. The species of the genus *Bitis* are beloved and all three big-bodied *Bitis* species (*B. arietans*, *B. gabonica*, *B. nasicornis*) are present in private collections. *Bitis arietans* has already been involved in several severe or even life-threatening accidents in the last decades in Hungary [9, 10]. On account of the similar effects of the toxin of *B. nasicornis* compared to the venom of *B. gabonica*, toxicologists rely on experiences gathered during accidents caused by this species when regarding the consequences of *B. nasicornis* intoxication [11]. For this reason we attach particular importance to reporting the circumstances of a single *B. nasicornis* accident and its clinical outcomes, discovered during a nationwide survey of snake-bite incidents in Hungary.

## Case report

A 30-year-old healthy male was transported by car to the Toxicological Ward of Erzsébet Hospital of Budapest, the main centre for snake-bite first aid, on 9<sup>th</sup> November 2001. The person was bitten by an adult (90–100 cm long) male Rhinoceros viper (*B. nasicornis*) (fig. 1). He had never been bitten previously (patient's personal comment). The snake's last feeding date was unknown. The captive bred specimen's ancestral origin was Ghana, West Africa. The specimen was estimated to be 2–3 years old and its owner bought it, when it was still juvenile (patient's personal comment). According to the patient, the snake had always

been calm and handled easily and special caution was only necessary during the snake's feeding time. The accident happened when the owner pulled the viper from its cage with a snake-hook.

After being bitten the victim was transported to hospital within an hour and did not receive any first aid until admission. The bite occurred with one fang only and was clearly visible on the pulp of the patient's right index-finger (fig. 2). At admission he had clear consciousness, was wide-awake and in a good mood considering the circumstances. Local symptoms were moderate: the bitten hand was oedematous (oedema extended to the wrist), the fingers were moderately swollen and tense, and slight livid discoloration developed around the fang mark (fig. 2). No other local symptoms occurred. He experienced slight hypertension, with systolic pressure of 155 mmHg, and diastolic of 80 mmHg. The ECG did not show any abnormal changes despite the fact that the venom of *B. nasicornis* can affect heart function [11, 12]. Only moderate tachycardia (105/min) was observed at the time of admission. Of the laboratory findings some rates, especially those of the cardiovascular system, diverged from the normal ones (table 1). The envenomed person had mild temporary coagulopathy (table 1), represented by the following laboratory results: INR 1.30; PTT 32 sec; and IVY 1.15 min.

Infusion (lactated Ringer solution 500 ml) and strong antiphlogistic (Tramadolium chloratum 50 mg i.v.) were administered at admission. Infusion is administrated routinely following mild to moderate snake-bite envenomation in Hungary [9]. Furthermore the patient received tetanus antitoxin injection (1 ampoule – 5 ml) and antibiotic therapy (Clindamicin 2×300 mg). The analgesic was necessary due to the tension and intense pain of the hand. Antivenom therapy was not considered. Hospital observation lasted for 4 days only. The patient complained of pain – presumably due to arthralgia of the bitten hand – for two weeks after the incident.

**Figure 1**

The Rhinoceros viper (*Bitis nasicornis*), which caused the accident (photograph by Balázs Buzás).



**Figure 2**

Local consequences of the envenoming (photograph by Zsolt Dernei).



## Discussion

There are differences of opinion regarding routine antibiotic use in cases of snake-bite. On the one hand the risk of routine antibiotic therapy leading to increases in resistant bacterial phyla [13] is well known, on the other, antibiotic use is not essential in many snake-bite incidents [14, 15]. Many authors [16–18] assume that routine antibiotic therapy is necessary due to the bacterial flora of snakes' saliva, especially in cases of accidents caused by a species with haemo-cytotoxic venom such as the family of Viperidae [18].

Justifying the use and explaining the benefit of anti-tetanus therapy is problematic, in particu-

lar since tetanus after snake-bite has not been documented [19]. Necrosis, blistering and abscess formation are often observed following *Bitis* bites [4, 8, 20, 21] but are caused in part by actions of the haemo- and cytotoxicity of the venom [18]. Local necrosis evolves on account of the properties of the venom (i.e., haemo- cyto- and myotox- icity) [1, 18], possibly interacting with the bacterial flora of snake's saliva, and the natural flora of human skin [18].

A similar case was recorded in 2002, when severe necrosis necessitated amputation of the patient's finger following snake-bite [7]. In our case

**Table 1**

Laboratory findings (\*normal rates in males).

Laboratory analysis	Patient's rates 09. 11. 2001 (Unit)	Patient's rates 10. 11. 2001 (Unit)	Normal rates [26, 27] (Unit)
<b>Blood</b>			
WBC	9.2 (M/ $\mu$ L)	12.5 (M/ $\mu$ L)	4.5–11.0 (M/ $\mu$ L)*
Lymphocytes	22.9 (%)	12.6 (%)	17–45 (%)
Granulocytes	73.4 (%)	83.0 (%)	42–74 (%)
Lymphocytes abs.	2.1 (G/l)	1.6 (G/l)	1.5–4.0 (G/l)
Granulocytes abs.	6.8 (G/l)	10.4 (G/l)	2.0–7.5 (G/l)
RBC	5.38 (M/ $\mu$ L)	4.90 (M/ $\mu$ L)	4.5–5.8 (M/ $\mu$ L)*
Haemoglobin (Hb)	163 (g/l)	150 (g/l)	130–180 (g/l)*
Haematocrit	0.51 (l/l)	0.475 (l/l)	40–50 (%)*
Mean Cell Volume (MCV)	95 (fl)	97 (fl)	80–100 (fl)
Mean Cell Haemoglobin (MCH)	30.3 (pg)	30.6 (pg)	27–32 (pg)
Mean Cell Haemoglobin Concentration (MCHC)	318.9 (g/l)	315.6 (g/l)	300–350 (g/l)
Thrombocyte count	271 (G/l)	250 (G/l)	150–400 (G/l)
Thrombin time	20 (sec)	19 (sec)	16–21 (sec)
Fibrinogen	2.0 (g/l)	1.8 (g/l)	1.5–4.0 (g/l)
Activated Partial Thromboplastin Time (PTT)	32 (sec)	33 (sec)	<29 (sec)
International Normalised Ratio (INR)	1.30	1.70	–
Coagulation time	4.10 (min.)	4.25 (min)	<9 (min)
SGOT	43 (U/l)	–	<50 (U/L)*
Alkaline phosphatase	143 (U/l)	–	25–100 (U/l)
Carbamid	8.0 (mmol/l)	–	3.0–8.0 mmol/l
IVY	1.15 (min.)	–	–
<b>Urine</b>			
RBC	3–4 (C/visual area)	–	>2*
WBC	5–6 (C/visual area)	–	>2*

anti-tetanus prophylaxis and antibiotic (clindamycin 2x300 mg/d) administration were deemed necessary to prevent secondary infections. Antibiotics and tetanus antitoxin are often applied for snake-bite therapy in Hungary [9]. The venom of *B. nasicornis* is haemo-cytotoxic, potently destructive to cellular elements of blood, degrading tissues and causing coagulation disorders [3, 12, 22, 23]. It also contains myocardiotoxins [11] affecting cardiac function and blood pressure, thus it can readily provoke arrhythmia, cardiac-mediated hypotension and myocardial damage.

Local and systemic bleeding can occur owing to the haemotoxic components of the venom. The venom yield of an adult specimen is high (200 mg – dry weight) [2], similar to the same-sized *B. arietans* (150–250 mg – dry weight) [24]. With its long fangs (250–300 mm) the species usually injects the venom directly into the muscles. The lethal dose of venom is only 8.6 mg/kg i.m., thus *B. nasicornis* produces the most toxic

venom intramuscularly of the big-bodied *Bitis* species (i.e., *B. gabonica* and *B. arietans*) [20]. An envenoming is potentially as dangerous as one arising from one of the two taxa mentioned above and can be fatal [7]. In Europe, two polyvalent antivenins (Ipser Africa® Pasteur Vaccins, France; and Anti-Bitis-Echis-Naja Africa®, Institut Pasteur, France) were produced against the bite of *B. nasicornis*, while another polyvalent serum (Saimr Polyvalent Snake Antivenom®, South African Vaccine Producers, Johannesburg, South Africa) is made in Africa. The gender of the snake is important as sexual venom variation is present in this species [12]. The female's venom contains one component absent from the venom of males [12], which might influence the outcome of a poisoning. Geographical venom variation might also influence the development of symptoms, a fact already proved in many other species [25]. The body size and age of the species affect the ophidism as well. Juveniles of variant species of Viperidae have more toxic venom, which has higher coagulant activity and lower proteolytic effects [25]. Experience shows that *B. nasicornis* bites cause intense pain and moderate to severe oedema of the bitten extremity. Compartment syndrome may also develop in severe cases, as has been observed among victims of *B. arietans* and/or *B. gabonica*. In our case, the local effects were similar to one of the US cases, where also the finger was bitten. Following this latter event the hand and the wrist became oedematous and intensive pain developed after injury [7]. Local necrosis also occurred [7], although this is often less serious and extensive than in the case of *B. arietans* bites.

In this case the patient was lucky not to apply a tourniquet as first aid; since the application of pressure dressings contributes to necrosis at the various *Bitis* bites [17]. Besides the typical local symptoms of puffadder-like bites, systemic symptoms also developed in our patient involving moderate tachycardia, slight hypertension and temporary coagulopathy. Temporary coagulopathy included shortened prothrombin time (table 1), and disappeared spontaneously without any medical intervention. It is likely that different coagulopathies will develop in moderate and severe poisonings by *B. nasicornis* owing to the haemostatic activity of their venom [23]; however these may also appear in mild cases. Fortunately, *B. nasicornis* are seldom encountered in Hungarian private collections, probably owing to difficulties in their procurement. The accident in question may have been contributed to by human negligence, as occurs in almost every snake-bite incident, mainly with specimens in captivity [15]. Most probably this was “only” a warning bite, which occurred with one fang only and the snake injected a small amount of venom. Thus our patient had very lucky.

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*Correspondence:*

*Tamás Malina*

*University of Szeged*

*Department of Systematic Zoology and Erology*

*Dugonics tér 13*

*H-6722 Szeged*

*E-Mail: dyspolidus@gmail.com*

## References

- Gutiérrez JM, Theakston RDG, Warrell DA. Confronting the neglected problem of snakebite envenoming: the need for a global partnership. *Plos Med.* 2006;3:1–5.
- Spawls S, Branch B. *The Dangerous Snakes of Africa. Natural History: Species directory: Venoms and snakebite.* South Africa: South African Book Publishers; 1995; 192 pp.
- Dorandeu F. The big vipers of Africa of the genus *Bitis* (Gray, 1842) and their venom. *Zoological, biochemical and clinical data. Med Trop. (Mars)* 1991;51:293–306.
- Wildi SM, Gämperlic A, Beer G, Markwaldera K. Severe envenoming by a Gaboon viper (*Bitis gabonica*). *Swiss Med Wkly.* 2001;131:54–5.
- Boyer DM. Notes on the natural history and husbandry of the Rhinoceros viper (*Bitis nasicornis*). *Reptiles.* 1995;3:9–12.
- Parrish HM. Rarity of snakebites in Minnesota. *Minnesota Med.* 1965;48:1071–6.
- Keyler DA. Exotic venomous snakebites in the USA at the millennia crossover. *Newsl Minnesota Herpetol Soc.* 2005;25:6–9.
- Watson WA, Litovitz TL, Rogers GC, Klein-Schwartz W, Youniss J, Rose SR, Borys, D, Mary E. 2002 Annual Association of Poison Control Centers toxic exposure surveillance system. *Am J Emerg Med.* 2003;21:353–421.
- Turchányi B, Szalontay T, Zacher G. Snake-bite injuries. *Orv Hetilap.* 2000;141:1067–71 (In Hungarian).
- Takács Z, Janisch M, Korsós Z. Contribution to the epidemiological and Clinical aspects of snake bites in Hungary. *Toxicon.* 1987;25:376.
- Alloatti G, Gattullo D, Losano G, Marsh NA, Pagliaro P, Vono P. The mechanical effects of Rhinoceros horned viper (*Bitis nasicornis*) venom on the isolated perfused guinea-pig heart. *Exp Physiol.* 1991;76:611–4.
- Marsh N, Glatston A. Venom of the Rhinoceros horned viper, *Bitis nasicornis*. *Toxicon.* 1974;12:621–8.
- Rao GG. Risk factors for the spread of antibiotic-resistant bacteria. *Drugs.* 1998;55:323–30.
- Bernheim A, Lorenzetti E, Licht A, Markwalder K, Schneemann M. Three cases of severe neurotoxicity after cobra bite (*Naja kaouthia*). *Swiss Med Wkly.* 2001;131:227–8.
- Warrell DA. Treatment of bites by adders and exotic venomous Snakes. *BMJ.* 2005;331:1244–7.
- Jorge MT, Riberio LA, Silva MLR, Kusano EJU, Mendonça JS. Bacteriology of abscess complicating Bothrops snake bites in humans: a prospective study. *Toxicon.* 1994;6:743–8.
- Jorge MT, Nishioka SA, Oliveira RB, Ribeiro LA, Silveira PVP. *Aeromonas hydrophila* soft-tissue infection as a complication of snakebite: report of three cases. *Ann Trop Med Parasitol.* 1998;92:213–7.
- Otero R, Johnayro G, Mesa MB, Duque E, Rodríguez O, Arango JL, et al. Complications of Bothrops, Prothidium, and Bothriechis snakebites in Columbia. A clinical study of 39 cases attended in a university hospital. *Toxicon.* 2002;40:1107–14.
- Bubalo P, Curi I, Fišter K. Characteristics of Venomous Snakebites in Herzegovina. *CMJ.* 2004;45:50–3.
- Mallow D, Ludwig D, Nilson G. *True Vipers: Natural History and Toxinology of Old World Vipers.* Florida: Krieger Publishing Co.; 2003; 359 pp.
- Warrell DA, Ormerod LD, Davidson NM. Bites by puff adder (*Bitis arietans*) in Nigeria, and whole value of antivenom. *BMJ.* 1975;4:697–700.
- Ghalayini R, Elliott WB. Cardiovascular effects of Horned viper (*Bitis nasicornis*) venom in the rat. *Toxicon.* 1985;23:567.
- Mackay J, Ferguson JC, McNicol GP. Effects of the venom of the rhinoceros horned viper (*Bitis nasicornis*) on blood coagulation, platelet aggregation, and fibrinolysis. *J Clin Path.* 1970;23:789–96.
- Phelps T. *Poisonous Snakes.* Dorset: Blandford Press; 1981; 237 pp.
- Chippaux JP, Williams V, White J. Snake venom variability: methods of study, results and interpretation. *Toxicon.* 1991;29:1279–1303.
- Clinical Lab Reference Range Guide 1– 102. [home page on the Internet] Available from: <http://www.hosp.uky.edu/Clin-lab/report.pdf> [update Tuesday, March 10, 2006]
- Reference Ranges – Common Pathology Tests [home page on the Internet] Available from: <http://www.hoslink.com/LabResults/refranges.htm>