## 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>

#### 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 snakebite 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.

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

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

#### 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).





#### 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 wideawake 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 discolouration 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  $2\times300$  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.

## 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 particular 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 myotoxicity) [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 <b>09. 11. 2001</b> (Unit)	Patient's rates <b>10. 11. 2001</b> (Unit)	Normal rates [ <b>26, 27</b> ] (Unit)
Blood			
WBC	9.2 (M/µL)	12.5 (M/µL)	4.5–11.0 (M/µL)*
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/µL)	4.90 (M/μL)	4.5–5.8 (M/µL)*
Haemoglobin (Hb)	163 (g/l)	150 (g/l)	130–180 (g/l)*
Haematocrit	0.51 (l/l)	0.475 (1/1)	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.)	_	_
Urine			
RBC	3–4 (C/visual area)	_	>2*
WBC	5–6 (C/visual area)	_	>2*

anti-tetanus prophylaxis and antibiotic (clindamicin 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 samesized *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 87

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.

We are grateful to the Toxicological Ward of Erzsébet Hospital of Budapest, which permitted the application of this snake-bite and to Balázs Buzás and Zsolt Dernei for the photographs. Dr. Balázs Tóth (Medico Uno Pharma Ltd., Hungary) and the four anonymous reviewers are acknowledged for their valuable comments on the manuscript.

Correspondence: Tamás Malina University of Szeged Department of Systematic Zoology and Erology Dugonics tér 13 H-6722 Szeged E-Mail: dyspholidus@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.
- 2 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.
- 3 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.
- 4 Wildi SM, Gämperlic A, Beer G, Markwaldera K. Severe envenoming by a Gaboon viper (Bitis gabonica). Swiss Med Wkly. 2001;131:54–5.
- 5 Boyer DM. Notes on the natural history and husbandry of the Rhinoceros viper (Bitis nasicornis). Reptiles. 1995;3:9–12.
- 6 Parrish HM. Rarity of snakebites in Minnesota. Minnesota Med. 1965;48:1071–6.
- 7 Keyler DA. Exotic venomous snakebites in the USA at the milenia crossover. Newsl Minnesota Herpetol Soc. 2005;25:6–9.
- 8 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.
- 9 Turchányi B, Szalontay T, Zacher G. Snake-bite injuries. Orv Hetilap. 2000;141:1067–71 (In Hungarian).
- 10 Takács Z, Janisch M, Korsós Z. Contribution to the epidemiological and Clinical aspects of snake bites in Hungary. Toxicon. 1987;25:376.
- 11 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.
- 12 Marsh N, Glatston A. Venom of the Rhinoceros horned viper, Bitis nasicornis. Toxicon. 1974;12:621–8.
- 13 Rao GG. Risk factors for the spread of antibiotic-resistant bacteria. Drugs. 1998;55:323–30.
- 14 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.

- 15 Warrell DA. Treatment of bites by adders and exotic venomous Snakes. BMJ. 2005;331:1244–7.
- 16 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.
- 17 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.
- 18 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.
- 20 Mallow D, Ludwig D, Nilson G. True Vipers: Natural History and Toxinology of Old World Vipers. Florida: Krieger Publishing Co.; 2003; 359 pp.
- 21 Warrell DA, Ormerod LD, Davidson NM. Bites by puff adder (Bitis arietans) in Nigeria, and whole value of antivenom. BMJ. 1975;4:697–700.
- 22 Ghalayini R, Elliott WB. Cardiovascular effects of Horned viper (Bitis nasicornis) venom in the rat. Toxicon. 1985;23:567.
- 23 Mackay J, Ferguson JC, McNicol GP. Effects of the venom of the rhinoceros horned viper (Bitis nasicornis) on blood coagulation, platelet aggregation, and fibrinolisys. J Clin Path. 1970;23:789–96.
- 24 Phelps T. Poisonous Snakes. Dorset: Blandford Press; 1981; 237 pp.
- 25 Chippaux JP, Williams V, White J. Snake venom variability: methods of study, results and interpretation. Toxicon. 1991;29: 1279–1303.
- 26 Clinical Lab Reference Range Guide 1– 102. [home page on the Internet] Available from: http://www.hosp.uky.edu/Clinlab/report.pdf [update Tuesday, March 10, 2006]
- 27 Reference Ranges Common Pathology Tests [home page on the Internet] Available from: http://www.hoslink.com/LabResults/refranges.htm

Formerly: Schweizerische Medizinische Wochenschrift

# Swiss Medical Weekly

The European Journal of Medical Sciences

## The many reasons why you should choose SMW to publish your research

What Swiss Medical Weekly has to offer:

- SMW's impact factor has been steadily rising. The 2006 impact factor is 1.346.
- Open access to the publication via the Internet, therefore wide audience and impact
- Rapid listing in Medline
- LinkOut-button from PubMed with link to the full text website http://www.smw.ch (direct link from each SMW record in PubMed)
- No-nonsense submission you submit a single copy of your manuscript by e-mail attachment
- Peer review based on a broad spectrum of international academic referees
- Assistance of professional statisticians for every article with statistical analyses
- Fast peer review, by e-mail exchange with the referees
- Prompt decisions based on weekly conferences of the Editorial Board
- Prompt notification on the status of your manuscript by e-mail
- Professional English copy editing

Editorial Board

Prof. Jean-Michel Dayer, Geneva
Prof Paul Erne, Lucerne
Prof. Peter Gehr, Berne
Prof. André P. Perruchoud, Basel
Prof. Andreas Schaffner, Zurich (editor in chief)
Prof. Werner Straub, Berne (senior editor)
Prof. Ludwig von Segesser, Lausanne International Advisory Committee Prof. K. E. Juhani Airaksinen, Turku, Fin-

land Prof. Anthony Bayes de Luna, Barcelona, Spain

Prof. Hubert E. Blum, Freiburg, Germany Prof. Walter E. Haefeli, Heidelberg, Germany

- Prof. Nino Kuenzli, Los Angeles, USA Prof. René Lutter, Amsterdam,
  - The Netherlands
- Prof. Claude Martin, Marseille, France Prof. Josef Patsch, Innsbruck, Austria Prof. Luigi Tavazzi, Pavia, Italy
- We evaluate manuscripts of broad clinical interest from all specialities, including experimental medicine and clinical investigation.

We look forward to receiving your paper!

Guidelines for authors: http://www.smw.ch/set\_authors.html

All manuscripts should be sent in electronic form, to:

EMH Swiss Medical Publishers Ltd. SMW Editorial Secretariat Farnsburgerstrasse 8 CH-4132 Muttenz

Manuscripts:	submission@smw.ch
Letters to the editor:	letters@smw.ch
Editorial Board:	red@smw.ch
Internet:	http://www.smw.ch



Official journal of the Swiss Society of Infectious Diseases, the Swiss Society of Internal Medicine and the Swiss Respiratory Society

Supported by the FMH (Swiss Medical Association) and by Schwabe AG, the long-established scientific publishing house founded in 1488