Research Article

Quantitative Characterization of the Hemorrhagic, Necrotic, Coagulation-Altering Properties and Edema-Forming Effects of Zebra Snake (Naja nigricincta nigricincta) Venom

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This study was designed to investigate the cytotoxicity and haemotoxicity of the Western barred (zebra) spitting cobra (Naja nigricincta nigricincta) venom to help explain atypical and inconsistent reports on syndromes by Namibian physicians treating victims of human ophidian accidents. Freeze-dried venom milked from adult zebra snakes was dissolved in phosphate buffered saline (PBS) for use in this study. Haemorrhagic and necrotic activity of venom were studied in New Zealand albino rabbits. Oedema-forming activity was investigated in 10-day-old Cobb500 broiler chicks. Procoagulant and thrombolytic activity was investigated in adult Kalahari red goat blood in vitro. The rabbit skin minimum hemorrhagic dose (MHD) for N. n. nigricincta was 9.8 μg. The minimum necrotizing dose (MND) for N. n. nigricincta venom was 12.2 μg. The N. n. nigricincta venom showed linear dose-dependent procoagulant activity on goat blood (p < 0.05). Likewise, N. n. nigricincta venom showed linear dose-dependent thrombolytic activity on goat blood (p < 0.05, n = 6). Subplantar injection of N. n. nigricincta venom (25 μg, 50 μg, 75 μg, and 100 μg) into chick paw resulted in peak oedema of 35.5%, 38.5%, 42.9%, and 47.5%, respectively, two hours after injection. Paw oedema subsided within five hours to a mean volume ranging from 5% (25 μg venom) to 17.6% (100 μg venom). In conclusion, though N. n. nigricincta belongs to the genus Elapidae, the current study has shown its venom to possess potent hemorrhagic, necrotic (cytotoxic), and paradoxically, both procoagulant and thrombolytic activity. The authors propose further work to fractionate, isolate, and elucidate the structure of the various N. n. nigricincta venom toxins as a prelude to the development of an antivenom.

1. Introduction

Since time immemorial, man has always suffered from envenomation resulting from snakebites. Although accurate statistics have proved elusive, it is estimated that the global burden of snakebites stands at 1.2 to 5.5 million bites per year, 25,000-125,000 deaths per year, and about 400,000 victims left with permanent disability [1]. In 2009, snakebite was declared a neglected tropical disease by the WHO [2]. The World Health Organization declared snake envenomation as a significant Sub-Saharan disease problem [3].

In Namibia, like in most other developing countries, the majority of snake bites result from the overlap of human and snake habitats, domiciliation of rodents (main prey of most snakes), the nocturnal and heat seeking poikilothermic nature of snakes, and accidents during snake handling. Some of these snakebites lead to fatalities and wound complications culminating in debilitating physical deformities in victims [4, 5] and associated socioeconomic problems resulting from these disabilities [6, 7]. The vast size of Namibia as a country also poses a potential problem of bringing emergency health care to such snakebite victims.

Venomous snakes belong to five main families: Hydrophiinae, Elapidae, Viperidae, Crotalidae, and Colubridae [8, 9]. These snakes possess venom glands that can synthesize, store, and secrete up to 50-60 proteins/peptides of varying
structure but are capable of causing damage at the bite site and systematically [2, 10]. The venom components are usually fairly similar in snakes of the same family [1]. Venoms of snakes belonging to the families Elapidae (mainly cobras and mambas) and Hydrophiinae (mainly sea snakes) are highly neurotoxic and produce flaccid paralysis and respiratory paralysis in animals [2, 11–15]. Viperidae (vipers), Colubridae (back-fanged venomous snakes, e.g., Boomslang and the Twig snake), and Crotalidae (pit-vipers) venoms produce in addition to systemic/lethal effects, striking local effects, namely, hemorrhage, necrosis, and oedema [11, 16] as well as alterations in coagulability of blood [17–19]. The protein components of the spitting cobra of Naja naja sputatrix comprises the proteins, three-finger toxins (3FTXs), phospholipase A\(_2\) (PLA\(_2\)), nerve growth factors, and snake venom metalloproteinase in that order [15]. The Zebra snake (N. n. nigricincta) is a venomous spitting snake belonging to the Elapidae family and found only in Namibia and Southern Angola [20].

Though belonging to the family of Elapids, empirical evidence suggests that the Zebra snake has acquired highly potent cytotoxic, hemorrhagic, anticoagulant, and thrombolytic toxins whilst retaining their familial neurotoxins. Namibia has had a very high number of both human and animal victims of the Zebra snake (Buys, 2016; personal communication). Snake antivenom immunoglobulins are the only specific treatment for envenoming by snakebites [4, 21]. Clinically, administering antivenom to the affected patient within a very limited time frame (<2 hours) efficiently reverses many of the detrimental systemic effects caused by snake venom [22].

2. Materials and Methods

2.1. Snake Venom. Venom was carefully and humanely milked by an expert snake handler from snakes that were caught in suburban Windhoek and later relocated to the surrounding Savanna bushveld after replenishment of venom gland stores. Approximate age, sex, length, girth, location of capture, and location of release were geo-referenced and recorded on a database for surveillance of snakes in and around Windhoek. Venom was diluted with distilled water and freeze-dried overnight using a Vitis Freeze Dryer (United Scientific). The resultant venom powder from each snake was stored in a separate sealed and appropriately labelled glass vial at -30°C until time of use. For this study, serial dilutions of venom (250 to 1000 μg/ml) from a single snake were made by dissolving known quantities of venom powder in Phosphate Buffered Saline at pH 7.4 freshly prepared from tablets (Sigma-Aldrich). Sterile 0.5ml needles were used to administer intradermal and subplantar injections of solutions.

2.2. Animals. Male and female albino New Zealand rabbits (about 2.5 kg weight) were obtained through City Pets, Windhoek, and reared in the small stock section at Neudamm farm, University of Namibia. Day old broiler Cobb500 chicks were obtained from Namib Poultry and reared in the Poultry section at Neudamm farm. Blood for thrombolytic studies was obtained from Kalahari red stud goats reared at the Neudamm farm.

2.3. Minimum Hemorrhagic Dose (MHD). The MHD is defined as the least amount of venom (μg dry weight) which, when injected intradermally into rabbits, results in a hemorrhagic lesion of 10 mm diameter 24 hours later [25]. Aliquots of 0.1 ml PBS containing 75, 10, 25, 50, 75, and 100 μg of venom were injected into the shaved dorsal skin of each of six adult rabbits marked with grids of 25mm squares (n=6). Three replicates were performed for each dose on different randomly chosen squares of the grid on each rabbit and then mean values were determined for each concentration on each animal. The animals were sacrificed after 24 hours, the dorsal skin was removed, and the diameter of the lesions was measured on the inner surface of the skin in two directions at right angles using calipers and with the aid of background illumination. The MHD was calculated using the regression equations relating the doses of venom to the mean diameters of the haemorrhagic lesions.

2.4. Minimum Necrotic Dose (MND). The MND is defined as the least amount of venom (μg dry weight) which, when injected intradermally into rabbits, results in a necrotic lesion of 5mm diameter 72 hours later [25]. The method used was the same as that for the MHD, except that the skin was removed 72 hours after injection. The MND was calculated using the regression equations relating the doses of venom to the mean diameters of the necrotic lesions.

2.5. Percentage Thrombolysis. Venous blood was drawn from healthy adult male Kalahari red goats (n = 6) of which 500μl of blood was transferred to each of previously weighed microcentrifuge tubes to form clot. Phosphate buffered saline (PBS) at pH 7.4 was added to lyophilized heparin vial (1000 I.U.) and mixed properly to create a stock solution from which serial dilutions of 0.05, 0.5, 5, and 50 I.U. heparin were made for observation of thrombolytic activity of heparin using the in vitro method developed by Prasad et al., 2006.
This protocol was adapted to measure the thrombolytic activity of 100\(\mu\)L solutions containing 25, 50, 75, or 100 \(\mu\)g Zebra snake venom on 500\(\mu\)L goat blood. This experiment was repeated three times with blood from each animal and mean values were determined for each concentration for blood from each animal.

2.6. Coagulation-Altering Activity. Venous blood was drawn from healthy adult male Kalahari red goats (n = 6) of which 500\(\mu\)L of blood was transferred to each of previously weighed 1.5ml Eppendorf tubes containing 100\(\mu\)L of 250, 500, 750, or 1000\(\mu\)g/ml of venom, 0.05, 0.5, 5, or 50 I.U. heparin (positive control) or 100\(\mu\)L of PBS (negative control) and gently mixed to avoid haemolyzing the blood. The mixtures were incubated at 37\(^\circ\)C for 90 minutes to allow clotsto form. After the incubation period, filter paper strips were used to drain any liquid contents from the microcentrifuge tubes. This experiment was repeated three times and mean values were determined for each concentration for blood from each animal. The clot weight was then determined and compared with mean clot weight from tubes mixed with PBS. Percentage coagulation was calculated using the equation below:

\[
\text{\%Coagulation} = \frac{\text{clot weight in tube with venom or heparin}}{\text{clot weight in tube with PBS}} \times 100
\]

2.7. Edema-Forming Effects. Subplantar injection of known quantities (25, 50, 75, and 100\(\mu\)g in 0.1ml PBS solution) of snake venom was performed into the right paws of 10-day-old chicks (about 250 - 300g weight). Eighteen chicks were used for this protocol. The change in the paw volume was quantified using the chick paw edema method by Fereidoni and coworkers [26] and improved by Ainooson and others [27]. Formalin (2.5\%) was used as a standard edema-forming substance (positive control) and PBS was used as the negative control. The experiment was repeated three times using different chicks for each level of treatment and mean values of proportional change in paw size per concentration were determined.

2.8. Ethical Statement. All animals were used with the ethical approval from the University of Namibia Ethical Clearance Committee (Certificate: NCREC/01/2018/1). All procedures performed on animals and disposal of animals/animal tissues followed a protocol approved by the University of Namibia Ethical Clearance Committee. In the course of this study, the researchers strictly adhered to the WHO guidelines [24].

2.9. Statistical Analysis. Descriptive and inferential statistics were performed in SPSS version 25 using one way ANOVA with Tukey’s post-hoc test. P values \(\leq 0.05\) were considered statistical significant.

3. Results

As shown in Figure 1, intradermal injection of \(N. n. nigricincta\) venom produced significant haemorrhagic lesions within 24hrs of injection. Maximum average diameter (40mm, \(n = 6\)) was recorded with the highest amount of \(N. n. nigricincta\) venom (100\(\mu\)g) injected.

As shown in Figure 2, \(N. n. nigricincta\) venom showed a significant dose-dependent increase in the diameter of the hemorrhagic lesions with each increase in the amount of venom injected into each site (\(p < 0.05\), \(n = 6\)). Hemorrhagic lesion diameter showed a very strong logarithmic dependence on dose of venom injected (\(R^2 = 0.90\)). The MHD determined from this relationship for \(N. n. nigricincta\) was 9.8\(\mu\)g.

As shown in Figure 3, \(N. n. nigricincta\) venom showed a significant dose-dependent increase in the diameter of the necrotic lesions with each increase in the amount of venom injected into each site (\(p < 0.05\), \(n = 6\)). Necrotic lesion diameter showed a very strong logarithmic dependence on dose of venom injected (\(R^2 = 0.93\)). The MND was determined from this relationship for \(N. n. nigricincta\) (12.4 \(\mu\)g).

Percentage thrombosis of goat blood showed an almost perfect negative logarithmic dependence on the dose of heparin (\(R^2 = 0.99991\)) (Figure 4). Each increase in amount of heparin (0.05 I.U., 0.5 I.U., 5 I.U., and 50 I.U.) incubated with goat blood showed a significant decrease in percentage thrombosis (\(p < 0.05\), \(n = 6\)). Incubation of 500\(\mu\)L goat blood
Figure 3: The dose-dependent necrotic activity of *N. n. nigricincta* venom on adult rabbit skin.

Figure 4: The dose-dependent anticoagulative activity of heparin on Kalahari red goat blood.

with 50 I.U. resulted in only 4.4% thrombosis whilst incubation of same volume of blood with 0.05 I.U. (a 1000 times less heparin) resulted in 99.5% thrombosis. These results show that heparin has significantly potent anticoagulant properties.

As shown in Figure 5, percentage thrombosis of Kalahari red goat blood showed a very strong linear dependence on the dose of *N. n. nigricincta* venom ($R^2 = 0.9892$). An increase in the amount of venom (25, 50, 75, and 100 μg) resulted in significantly higher levels of thrombosis (33.2%, 48.7%, 75.9%, and 93.6%, respectively). These results show that *N. n. nigricincta* venom has significantly potent procoagulant properties ($p<0.05$).

As shown in Figure 6, thrombolysis of Kalahari red goat blood clots showed very strong linear relationship with the doses of heparin ($R^2 = 0.98$) and *N. n. nigricincta* ($R^2 = 0.99$). At 100 μg *N. n. nigricincta* venom showed 60% thrombolysis which was significantly higher than the 45.6% thrombolysis in the presence of 50 I.U. heparin ($p<0.05, n=6$). At 75 μg *N. n. nigricincta* venom showed 50.9% thrombolysis which was significantly higher than the 43.9% thrombolysis in the presence of 5 I.U. heparin ($p<0.05, n=6$). At 50 μg *N. n. nigricincta* venom showed 50.1% thrombolysis which was significantly higher than the 40% thrombolysis in the presence of 0.5 I.U. heparin ($n<0.05$). At 25 μg *N. n. nigricincta* venom showed 44.4% thrombolysis which was significantly higher than the 36.4% thrombolysis in the presence of 0.05 I.U. heparin ($p<0.05$).

Subplantar injection of *N. n. nigricincta* venom into chick paw resulted in peak oedema 2 hrs after injection and subsided within 5 hrs to a mean volume ranging from 5% larger than the original volume (due to 25 μg venom) to 17.6% larger than the original volume (due to 100 μg venom) (Figure 7). The peak oedema was 35.5%, 38.5%, 42.9%, 47.5%, and 16.3% due to 25 μg, 50 μg, 75 μg, and 100 μg of venom and 2.5% formalin, respectively. Significant declines in oedema were noticed with smaller quantities of venom (100 μg > 75 μg > 50 μg, respectively) ($p<0.05; n=18$).

Oedema due to 50 μg of venom was significantly greater than that due to 25 μg of venom which in turn was greater than oedema caused by 2.5% formalin ($p<0.05$). Injection of PBS resulted in a peak increase in paw size of 4.6% within the first 30 mins. Paw size resolved back to normal (0% increase) within 2 hrs of injection. The change in paw size due to venom injection was significantly greater than that due to PBS injection throughout the 5 hrs of observation ($p<0.05$). The change in paw size due to 2.5% formalin was significantly greater than that due to PBS for the first 3 hrs after injection.
of haemorrhagic activity in studies on the venoms from these elapids. The assessment [29], though there is no documentation of MHD and MND of the Pakistan Naja naja was also recently documented of the Malayan blue coral snake [28]. The proteomic profile of the venom at 28.2 𝜇g (Nigeria), 47.15 𝜇g (Nigeria), and N. kauaihina venom at 28.2 𝜇g (Nigeria), and N. mossambica. The venoms from spitting elapids contains 67-73% three finger toxins (3FTXs), 22-30% phospholipases A2 (PLA2), 2.1% snake venom metalloproteinases (SVMPs), and minor quantities of nucleotidases and cysteine-rich secretory proteins (CRISSPs) (Hus et al. [32]). Up to five cytotoxins (cytotoxin 1, 2, 4, 5, and 11) have been isolated and strongly implicated in the cytotoxicity of N. mossambica, N annulifer, and N. pallida which are all close relatives of the N. n. nigricinta [32]. Similar findings with another spitting cobra species, Naja sputatrix, revealed its venom to contain 3FTXs (64.2%), PLA2 (31.2%), nerve growth factor (1.82%), and SVMPs (1.33%). About 48.08% of these 3FTXs were cytotoxins [15]. In addition, PLA2s (acidic PLA2 CM I, basic PLA2 I, and basic PLA2 CMIII) have also been implicated in the cytotoxicity of N. mossambica. SVMPs (cobilin, atragin, and atrase) have been identified and implicated as minor contributors to cytotoxic activity in N. atra and N. kaouthia venoms [33, 34]. Isolation and identification of specific cytotoxic SVMPs in Southern African spitting elapids is, however, not yet reported. It is therefore, logical to speculate that the Southern African spitting cobras also have a smaller contribution from SVMP’s towards their venoms’ cytotoxicity. CRISPs (annuliferin, nawafarin, and natrin I) have been isolated and confirmed to contribute towards the cytotoxic activity of N. nigricollis and N. annulifer both of which are close relatives of the N. n. nigricinta [32].

The cytotoxic mechanisms of 3FTXs, PLA2s, and SVMPs mainly involve the disruption of microvascular basement membranes [35] as well as endothelial cell membranes to result in the observed haemorrhage, oedema, and myonecrosis (resulting from disruption of plasma membranes of skeletal muscle cells) [32]. These mechanisms provide possible objectives in any further investigation of the cytotoxicity of N. n. nigricinta venom. In vitro exposure of whole goat blood to N. n. nigricinta venom resulted in enhancement of coagulation but the exposure of pre-formed goat blood clots resulted in profound thrombolytic activity. It is not unusual to find one venom containing both fibrinolytic (anticoagulant) and fibrinogenolytic (coagulant) activities [36, 37]. Snake venom serine proteinases (SVSPs) have been found in elapid, viperid, and colubrid venoms. These have been implicated in the interference with platelet aggregation, blood coagulation, fibrinolysis, complement system, and immune system [38]. Thrombin-like SVSPs (TLEs), however, have been implicated in procoagulation through activation of factor V, VIII, XIII, possibly VII and XI. TLEs have also been known to stimulate fibrinolysis and also activation of platelet aggregation [39]. Future studies with N. n. nigricinta venom can be guided towards proving or disproving involvement of these mechanisms.

An L-amino acid oxidase with human platelet aggregation activity from Ophiophagus hannah (king cobra) venom was isolated and characterized [40]. Cardiotoxin was isolated from Naja naja atra (Chinese cobra) venom; this toxin was able to potentiate platelet aggregation induced by ADP, thrombin, collagen, and venom phospholipase A2 [41]. Cobra venom phospholipase A2 showed conflicting effects on washed rabbit platelets, an initial reversible calcium-dependent aggregation followed by an inhibition of platelet

Figure 7: Dose-dependent oedema-forming effect of N. n. nigricincta venom on 10-day-old chick paw.

(p<0.05). There was no significant difference in change in paw size between 2.5% formalin and PBS at 4hrs and 5hrs after injection (p>0.05%).

4. Discussion

MHD and MND have been extensively used in the preclinical assessment of viperid and crotalid venoms as important WHO approved protocols for haemorrhagic and necrotizing venom toxins. Though longstanding knowledge of cytotoxins in spitting elapids exists, the presence of powerful cytotoxins has also been well investigated and documented in nonspitting elapids. Tan and coworkers characterized a significant cytotoxin contribution to the proteomic profile of the Malayan blue coral snake [28]. The proteomic profile of the Pakistan Naja naja was also recently documented [29], though there is no documentation of MHD and MND studies on the venoms from these elapids. The assessment of haemorrhagic activity in Micrurus pyrrhocryptus (a Latin American elapid) venom using these protocols in mice and rats produced negative results [30]. The MHD (rabbit) of N. n. nigricincta venom at 9.8 𝜇g was, however, almost similar to that of Bothrops atrox, a viper, determined using a similar assay on mice by researchers in Colombia (Otero et al., 2000). At 12.4 𝜇g, the MND of N. n. nigricincta venom was less than the 39.3 𝜇g of Echis ocellatus (Nigeria), 47.15 𝜇g of Echis leucogaster (Mali), the 24.9 𝜇g of Echis pyramidum (Kenya), the 64.8 𝜇g of Bitis arietans (Nigeria), and the 28.2 𝜇g of Bitis gabonica (Nigeria) from a study by Segura et al. [31]. These findings show that N. n. nigricincta venom has probably successfully acquired haemorrhagic activity equal to or even surpassing those of Viperidae. In this study, an attempt to reduce the number of animals used by using rabbits in place of rats and mice, however, resulted in a major limitation when comparing the findings from this study to those of other earlier studies. The findings from this pioneering study with Zebra snake venom, however, provide a basis way for the use of WHO protocols involving large numbers of mice and rat for future work to determine and compare the toxicity of this venom to other
aggregation with longer incubation times [42]. Two three-finger toxins, hemexin A and hemexin B, were isolated and purified from *Hemachatus haemachatus* (rinkhals) venom. Individually, hemexin A prolongs blood coagulation, but hemexin B does not show any effect on blood clotting. However, hemexin AB complex inhibits coagulation by noncompetitively inhibiting the Tissue Factor–Factor VIIa (extrinsic tenase) complex [43]. Studies to profile the proteomics of *N. n. nigricincta* venom would provide significant and relevant information that can be applied in combating envenomation from this species.

Oedema-causing toxins in snake venom have not been extensively studied. However, one study concluded that oedema induced by Bothrops snake venoms was multifactorial [44]. Other workers suggested that haemorrhagic toxins, through disruption of the microvasculature, resulted in extravasation which characterizes observed oedema in some envenomations [45, 46]. These authors also suggested the involvement of other toxins which acted directly on the endothelial cells of capillaries and venules thus increasing their permeability. Histamine release from mast cells as a result of phospholipases and cytotoxins was also a possible mechanism [47]. Another suggested mechanism was the release of prostaglandins resulting from phospholipase A2-induced liberation of arachidonic acid from plasma membranes [44]. In one study it was illustrated that *Bothrops jararaca* venom proteases activated plasma kininogens to bradykinin [44]. Kallikrein released after vascular damage was also suggested in the activation of kininogens to bradykinin [44]. Due to the major differences between viperid and elapids, suggestion of dilation (and thus oedema) at the site of inflammation [48]. bradykinin, an inflammatory mediator responsible for vascular dilatation of snake venom metalloproteinases: A journey of discovery and understanding," *Toxins*, vol. 8, no. 4, article no. 93, 2016.

5. Conclusion

In conclusion, fractionation of *N. n. nigricincta* venom and further investigations with cytotoxic, oedema-forming, procoagulant, and thrombolytic fractions separately may reveal the toxins responsible for observed activity of the venom in this study. Further work will then be required with *N. n. nigricincta* venom to unravel the mechanisms of action of any discovered toxins. The severity of the sequelae of the local cytotoxicity of this venom warrants a separate investigation into the formulation of effective intervention measures (both antivenom and other emergency on-site measures including phytotherapy) to reduce the fatalities and bodily deformities resulting from envenomation by this snake.

Data Availability

The data used to support the findings of this study are available from the corresponding author upon request.

Conflicts of Interest

The authors declare that they have no conflicts of interest.

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