

Namibian spitting cobra, *Naja nigricincta nigricincta* (zebra snake): Oral flora and antibiotic sensitivity, a cross-sectional study

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This was a cross-sectional study with the aim of characterising *Naja nigricincta nigricincta*'s oral bacterial flora as well as accompanying sensitivities and resistance towards antibiotics. *Naja nigricincta nigricincta* (zebra snake) is a spitting cobra indigenous to Namibia. Nasopharyngeal and venom swabs for bacterial culture and antibiotic sensitivity were taken from 37 native zebra snakes originating from the Khomas region that were captured for removal and relocation. *Enterococcus faecalis*, *Proteus* spp., *Morganella morganii* and *Pseudomonas* spp. were the organisms most often cultured. The antibiotic sensitivity profiles of these organisms suggest ciprofloxacin or a third-generation cephalosporin plus gentamicin or piperacillin-tazobactam as prophylactic antibiotics in case of *Naja nigricincta nigricincta* bites.

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African spitting cobras, such as *Naja nigricollis*, *Naja mossambica* and *Naja nigricincta nigricincta*, most often bite at night while the victims are asleep. Spitting cobra bites frequently result in local necrosis and secondary infection, often culminating in disfigurement, loss of function and amputation.^[1-6] Small children and babies are often bitten. Children are particularly vulnerable to snake envenomation, and suffer a high morbidity and mortality from these severely cytotoxic bites.

Naja nigricincta nigricincta (Western barred spitting cobra/zebra snake) (Fig. 1) is endemic to central and northern Namibia and southern Angola, and accounts for most of the venomous bites encountered in these areas of Namibia.^[1,2,7,8]

This cytotoxic venom typically results in a severe dermonecrosis. This resembles a type of venom-induced necrotising fasciitis, with

fast-spreading necrosis in the fascial planes between skin and deeper-lying muscles.^[1,2]

Prophylactic antibiotics are not recommended following snakebite in southern Africa.^[9] In Namibia, antibiotics are routinely part of the treatment of all cytotoxic bites (Namibian Medical Snakebite Management guidelines – Drs PJC Buys and EL Saaiman, unpublished). An increasing number of studies suggest that soft-tissue infection is one of the most substantial complications of cytotoxic snakebites, and that pre-emptive antibiotics should be considered in patients with severe local envenomation.^[4,10-15]

The extensive tissue destruction and devitalisation, caused by local cytotoxic envenomation, predispose the wound to bacterial infection. Inoculation of bacteria originating from the snake's indigenous oral flora, the environment or the victim's surrounding skin can occur during a bite. Venom-induced dermonecrosis may thus expand and exacerbate into an accompanying soft-tissue infection, and even progress to infective necrotising fasciitis.^[6,16-19]

Comprehensive identification of the microbiology of bite wounds and oral flora of culprit snakes is pertinent in selecting suitable empirical and prophylactic antibiotics, preventing secondary infection and reducing morbidity.^[15,19-21]

No data exist on *Naja nigricincta nigricincta*'s oral microbiome. Very few case reports of *Naja nigricincta nigricincta* snakebites have been recorded. Microscopy, culture and sensitivity (MC&S) results on 10 *Naja nigricincta nigricincta* bite wound swabs (2012 - 2020) were recovered from case files and analysed. The swabs were all taken at different times post bite. There were no data regarding the swabbing procedures, the specific wound areas swabbed or the indications for taking the wound swabs. *Enterococcus faecalis*, *Morganella morganii* and *Proteus* spp. were cultured most often. Other Gram-negative bacteria cultured included *Klebsiella pneumoniae*, *Escherichia coli*, *Salmonella enterica*, *Serratia marcescens*, *Schewanella algae* and



Fig. 1. *Naja nigricincta nigricincta* (Western barred spitting cobra/zebra snake).

Chryseobacterium indologenes. The Gram-negative organisms were sensitive to third- and fourth-generation cephalosporins, ciprofloxacin, gentamicin and piperacillin-tazobactam (Table 1).

Two other regional publications originating from KwaZulu-Natal (KZN), South Africa, examining the microbiology from infected cytotoxic snakebite wounds.^[15,22] Blaylock^[22] studied wound swabs taken from 14 cytotoxic snakebite victims, with associated necrosis, abscesses and haematomas. The snakes were mostly unidentified. In this study, *Morganella morganii*, *Proteus* spp., *Citrobacter* and *Serratia* spp. were cultured most often. No antibiotic profiles on the organisms were done (Table 2).

Wagener *et al.*^[4] analysed the microbiology results of 42 snakebite patients who required surgical debridement for extensive skin and soft-tissue necrosis after snakebite. The snakes responsible were not identified. Snakes most likely responsible for cytotoxic bites in this area of KZN are *Naja mossambica* (Mozambique spitting cobra), *Bitis arietans* (puff adder) and *Hemachatus haemachatus* (rinkhals).^[8] At the time of debridement, tissue samples of necrotic or infected tissue were sent for bacteriological analysis. *Morganella morganii*, *Enterococcus faecalis* and *Proteus* spp. were most often encountered. Other organisms cultured were *Salmonella enterica*, *Enterobacter cloacae*, *Escherichia coli*, *Klebsiella pneumoniae* and *Citrobacter freundii* (Table 2). The Gram-negative *Enterobacteriaceae* showed a resistance to penicillins and first- and second-generation cephalosporins, but were sensitive to third-generation cephalosporins, ciprofloxacin and aminoglycosides.^[4]

Both the above KZN study (Wagener *et al.*) and the *Naja nigricincta nigricincta* bite wound swab results are very similar to Taiwanese and Chinese publications on wound infections secondary to snakebite, where *Morganella morganii* and *Enterococcus* spp. were the most common pathogens found. All these Gram-negative organisms displayed a similar resistance to penicillins and first and second-generation cephalosporins (Table 2).^[4,14,17,23]

According to above bite wound microbiology, piperacillin-tazobactam, a quinolone, or second- or third-generation cephalosporin are proposed as empirical therapy following snakebite.^[4,14,17,23]

The possibility that primary infections are caused by the inoculation of the snake's oral flora during a bite is illustrated by a recent case report. A 2-and-a-half-year-old boy presented with an infective (*Proteus vulgaris*) necrotising fasciitis following *Naja nigricincta nigricincta* bite with rapid deterioration into multi-organ failure and death. A *Proteus vulgaris* with the same antibiotic profile was cultured from the mouth of the culprit snake (Table 3).^[6]

This case underscored the lack of current data on *Naja nigricincta nigricincta*'s oral biome, the limited data on post-*Naja nigricincta nigricincta* bite wound microbiology and the inadequacy of the post-snakebite antibiotic protocol then in place.

Study

This was a cross-sectional study. The aim was to characterise the patterns of oral bacterial flora in *Naja nigricincta nigricincta*, to determine the antibiotic sensitivity and resistance of above pathogens and to develop rational guidelines for antimicrobial prophylaxis after *Naja nigricincta nigricincta* snakebite injury in Namibia.

The study was conducted between 18 November 2020 and 15 April 2021. A total of 37 zebra snakes (*Naja nigricincta nigricincta*) that were caught for removal and relocation were used in the study. All snakes came from the Khomas region of Namibia, with the GPS location of where the snake was caught recorded in accordance with the Ministry of Environment, Forestry and Tourism's Human-Snake-Conflict-Mitigation programme (Fig. 2). The snakes were identified and measured, and the gender established, and milked by a local expert. Both an oropharyngeal and venom swab were taken from each snake using aseptic techniques by a medical professional. The swabs were sent for MC&S using conventional culture methods.

All the snakes were relocated into the wild.

Permit

In line with regulations imposed by the Namibian Ministry of Environment, Forestry and Tourism regarding handling of animals for scientific purposes, the study protocol was submitted for study permission from the National Commission on Research Science and Technology

Table 1. Organisms cultured from wound swabs of 10 patients bitten by *Naja nigricincta nigricincta* and antibiotic sensitivities

Patient number	Days post bite	Organism	Penicillin			Cephalo-sporin		Carba-penem	Amino-glycoside		Other	
			A	AC	PT	3	4	M	AK	G	CF	ST
4	1	<i>Proteus vulgaris</i> (G-)		R		S			S			
13	2	<i>Morganella morganii</i> (G-)					S	S		S		
		<i>Enterococcus faecalis</i> (G+)	S									
15	2	<i>Morganella morganii</i> (G-)			S		S	S		S	S	
		<i>Enterococcus faecalis</i> (G+)	S	S								
11	4	<i>Chryseobacterium indol</i> (G-)			S		S				S	
		<i>Enterococcus faecalis</i> (G+)	S									
5	6	<i>Serratia marcescens</i> (G-)	R	S							S	S
16	7	<i>Morganella morganii</i> (G-)								S	S	S
		<i>Enterococcus faecalis</i> (G+)	S									
7	10	<i>Morganella morganii</i> (G-)				S	S	S		S	S	S
		<i>Enterococcus faecalis</i> (G+)	S									
8	10	<i>Proteus</i> spp. (G-)		S		S				S		
9	14	<i>Klebsiella pneumoniae</i> (G-)				S		S		S		
		<i>Morganella morganii</i> (G-)				S		S		S		
14	11	<i>Schewanella algae</i> (G-)			S		S			S		

A = ampicillin; AC = amoxicillin-clavulanate; PT = piperacillin-tazobactam; 3 = third-generation cephalosporin; 4 = fourth-generation cephalosporin; M = meropenem; AK = amikacin; G = gentamicin; CF = ciprofloxacin; ST = sulfa-trimethoprim; G- = Gram negative; G+ = Gram positive; R = resistant; S = sensitive.

(NCRST) (Section 4 of Research Science and Technology Act, Act No. 23 of 2004).

Permit AN20200222 was granted in accordance with Section 45 of the Nature Conservation Ordinance 4 of 1975 of the Republic of Namibia, regarding animals for scientific purposes.

Catching and swabbing

Expert snake handlers caught the snakes. The GPS location of where the snake was caught was recorded using the Epicollect app (Fig. 2).

Cloacal probing with a blunt sexting probe^[24] to establish gender, as well as snout-to-vent measurements using the tube restraint method, were done by a local snake expert.^[24]

The snakes were milked using the acknowledged method of voluntary injection of the venom into a receptacle through a rubber or para-film membrane (Fig. 3).^[25] The para-films were cleaned with 90% ethanol spray. A first milking was followed immediately by a second milking through another ethanol-cleaned para-film. This was

done to achieve a more representative sample of uncontaminated venom – rather like a midstream urine sample. A swab was taken from the second sample.

After milking, the snakes' mouths were opened by a small sterile speculum and an oropharyngeal swab taken (Fig. 4). All swabs were taken by a medical professional and were kept refrigerated before being sent to NAMPath laboratory.

Bacterial identification and antibiotic susceptibility

A Gram smear was made aseptically onto a slide, prior to inoculation, which was then stained and viewed under 100× objective lenses. The swab was inoculated onto 5% blood agar, McConkey agar and chocolate agar plates. After inoculating the plates, the swab was transferred into a tube containing thio-glycolate enrichment medium. Both the inoculated plates and thio-glycolate medium with the swab inside were then incubated

Table 2. Comparison of the frequency of organisms cultured from the venom and oropharynx of 21 adult *Naja nigricincta nigricincta* compared with wound swabs of 10 patients bitten by *Naja nigricincta nigricincta*, as well as wound swabs from 42 and 14 patients with secondary wound infection, post snakebite in KZN, SA^[4,22]

Organism	Venom and oropharynx (21 snakes), %		Wounds (64 patients), %	
	Adult <i>Naja nigricincta nigricincta</i> (Namibia)	Namibia (10 patients), <i>Naja nigricincta nigricincta</i> bites	KZN (42 patients), wound infections post bite, 2017	KZN (14 patients), wound infections post bite, 1999
<i>Enterococcus faecalis</i> (G+)	71.4	50	31	-
<i>Morganella morganii</i> (G-)	19	50	40.5	28.5
<i>Proteus</i> spp. (G-)	71.4	20	23.8	28.5
<i>Salmonella enterica</i> (G-)	9.6	10	7.1	7.1
<i>Klebsiella pneumoniae</i> (G-)	4.8	10	2.4	-
<i>Citrobacter freundii</i> (G-)	9.6	-	2.4	21.4
<i>Acinetobacter</i> (G-)	9.6	-	-	-
<i>Yersinia</i> spp. (G-)	4.8	-	-	-
<i>Pseudomonas</i> spp. (G-)	19	-	-	-
<i>Elizabethkingia</i> spp. (G-)	4.8	-	-	-
<i>Vibrio</i> spp. (G-)	4.8	-	-	-
<i>Serratia marcescens</i> (G-)	-	10	-	21.4
<i>Schewanella algae</i> (G-)	-	10	-	-
<i>Chryseobacterium indologenes</i> (G-)	-	10	-	-
<i>Escherichia coli</i> (G-)	-	-	2.4	14.2
<i>Enterobacter complex</i> (G-)	-	-	4.8	7.1
<i>Staphylococcus</i> spp. (G+)	-	-	-	7.1
<i>Streptococcus</i> spp. (G-)	-	-	-	7.1

KZN = KwaZulu-Natal Province; SA = South Africa; G+ = Gram positive; G- = Gram negative.

Table 3. Organisms cultured from the patient's wound swab and the culprit *Naja nigricincta nigricincta*, with antibiotic sensitivities

Organism	Location		Penicillins			Cephalo- sporins			Carba- penems		Amino- glycosides		Other	
	Patient	Snake	AC	A	PT	2	3	4	IM	M	AK	G	CF	ST
<i>Proteus vulgaris</i>	Wound	Mouth, pharynx, cloaca	R	R	S	R	R	S	R	S	S	S	S	S
<i>Morganella morganii</i>	-	Mouth, venom, cloaca	R	R	R	R	R	R	R	S	S	S	S	S
<i>Salmonella</i> spp.	-	Pharynx		S	S		S		S	S			S	S
<i>Enterococcus faecalis</i>	-	Venom, skin	S	S								S		
<i>Pseudomonas aeruginosa</i>	-	Cloaca			S		S	S	I	S	S	S	S	

AC = amoxicillin-clavulanate; A = ampicillin; PT = piperacillin-tazobactam; 2 = second-generation cephalosporin; 3 = third-generation cephalosporin; 4 = fourth-generation cephalosporin; IM = imipenem; M = meropenem; AK = amikacin; G = gentamicin; CF = ciprofloxacin; ST = sulfa-trimethoprim; R = resistant; S = sensitive; I = intermediate sensitivity.



Fig. 2. The red dots represent the capture locations of 22 of the 37 snakes milked for this study. All snakes represented here were in direct conflict with humans in an urban setting. The map was constructed using QGIS 3.24.1.



Fig. 3. Milking of snake.



Fig. 4. Oropharyngeal swab.

at 37°C. The 5% blood agar and chocolate agar were incubated under CO₂ conditions.

Plates were read after 24 hours of incubation, and identification was by use of the manual API 10S method for lactose fermenters. For non-lactose fermenters and Gram-positive cocci, the automated Vitek 2

system was utilised. Antibiotic susceptibilities were determined either by the manual Kirby-Bauer method or the automated Vitek system (minimum inhibitory concentration assay).

After 24 hours, the swabs that were inoculated into the thioglycolate enrichment medium were re-inoculated onto 5% blood agar, chocolate agar and McConkey agar and incubated at 37°C.

After 48 hours of incubation, plates were read for identification and antibiotic susceptibility (as described above). Previously inoculated plates from the thioglycolate medium were checked for new bacterial growth not obtained from the initial culture plates. If new bacteria had grown, identification and antibiotic susceptibility were done as a follow-up.

Results from the Gram stain, identification and antibiotic susceptibility were entered into the laboratory information system, and laboratory reports generated.

Results

Sex and length

Twenty-one adult snakes, of which 15 were male (adult length ≥80 cm),^[26] measuring 90 - 165 cm, and 16 juvenile snakes, ranging from 28 - 35 cm, were caught.

Bacteria cultured

Organisms were cultured from all venom and oral samples. No anaerobic organisms were cultured. In clinical practice, juvenile *Naja nigricincta nigricincta* snakes are very seldom responsible for bites. The results from adult and juvenile snakes were differentiated and recorded into separate tables (Tables 4, 5 and 6).

Antibiotic profile

The antibiotic profiles of organisms cultured from adult and juvenile *Naja nigricincta nigricincta* oropharynges and venom are detailed in Table 7.

Discussion

Although venom is thought to be sterile with strong antimicrobial properties,^[22] archaea, algae, bacteria, fungi, protozoa and viruses have all been found to be present in certain venom microenvironments.^[27,28] The anatomy of the envenomation apparatus, i.e. an open duct attached to a liquid vessel with intermittent flow, may allow for the colonisation and facilitation of bacterial persistence and adaptation within antimicrobial venom.^[29] All the venom samples yielded organisms, and eight of the adult venom samples cultured different organisms than those from the same snake's oropharynx, suggesting venom colonisation. Further research in this field is needed.

The aim of this study was to identify oral snake pathogens with their specific antibiotic profile that can be inoculated during snakebite. Whether these organisms originate from the venom or the oropharynx is not of clinical relevance. In analysing the findings of the study, the results of each snake's venom and oropharyngeal swabs were thus combined.

No anaerobic organisms were cultured. The most frequent organisms cultured from *Naja nigricincta nigricincta* adult venom and oropharynx were *Enterococcus faecalis* and *Proteus* spp., both in 71.4% of cases, with *Morganella* spp. and *Pseudomonas* spp. in 19%. Also cultured were *Salmonella*, *Acinetobacter* and *Citrobacter* spp. (Tables 4 and 6). Pathogens were present in the oral flora (venom and/or oropharynx) of even the smallest snake. *Enterococcus faecalis* (43.7%) and *Proteus* spp. (87.5%) were most often cultured from the juvenile snakes (Tables 5 and 6).

In line with other publications, the similarity of the organisms cultured between individual *Naja nigricincta nigricincta*'s hints at the

Table 4. Bacteria cultured from adult *Naja nigricincta nigricincta* venom and oropharynx

Snake number, sex	Venom bacteria	Oropharynx bacteria
1 male	<i>Klebsiella pneumoniae</i> , <i>Enterococcus faecalis</i>	<i>Citrobacter braakii</i> , <i>Enterococcus faecalis</i>
2 male	<i>Morganella morganii</i>	<i>Acinetobacter baumannii</i>
3 male	<i>Proteus mirabilis</i>	<i>Salmonella enterica</i> , <i>Enterococcus faecalis</i>
4 male	<i>Proteus mirabilis</i>	<i>Yersinia enterocolytica</i>
5 male	<i>Proteus vulgaris</i>	<i>Proteus vulgaris</i> , <i>Enterococcus faecalis</i>
6 male	<i>Proteus mirabilis</i>	<i>Proteus penneri</i>
7 male	<i>Enterococcus faecalis</i> , <i>Proteus penneri</i>	<i>Pseudomonas aeruginosa</i> , <i>Citrobacter freundii</i>
8 male	<i>Morganella morganii</i>	<i>Morganella morganii</i> , <i>Elizabethkingia meningoseptica</i> , <i>Enterococcus faecalis</i>
9 male	<i>Pseudomonas aeruginosa</i> , <i>Enterococcus faecalis</i>	No sample
10 female	<i>Enterococcus faecalis</i> , <i>Proteus vulgaris</i>	<i>Enterococcus faecalis</i> , <i>Proteus vulgaris</i>
11 female	<i>Enterococcus faecalis</i>	<i>Enterococcus faecalis</i> , <i>Vibrio alginolyticus</i>
13 female	<i>Enterococcus faecalis</i> , <i>Proteus vulgaris</i>	<i>Enterococcus faecalis</i> , <i>Acinetobacter baumannii</i> , <i>Pseudomonas aeruginosa</i>
22 female	<i>Enterococcus faecalis</i>	<i>Proteus mirabilis</i> , <i>Enterococcus faecalis</i>
23 female	<i>Proteus mirabilis</i>	<i>Proteus vulgaris</i>
24 male	<i>Enterococcus faecalis</i> , <i>Proteus vulgaris</i>	<i>Enterococcus</i> spp., <i>Proteus mirabilis</i>
25 male	<i>Proteus morganii</i>	<i>Proteus mirabilis</i>
26 male	<i>Enterococcus faecalis</i> , <i>Morganella morganii</i>	<i>Enterococcus faecalis</i> , <i>Proteus vulgaris</i> , <i>Salmonella enterica</i>
30 female	<i>Enterococcus faecalis</i> , <i>Proteus vulgaris</i>	<i>Enterococcus faecalis</i> , <i>Pseudomonas aeruginosa</i>
31 male	<i>Enterococcus faecalis</i>	<i>Enterococcus faecalis</i> <i>Morganella morganii</i>
33 male	<i>Enterococcus faecalis</i> , <i>Proteus penneri</i>	<i>Enterococcus faecalis</i> , <i>Proteus mirabilis</i>
34 male	<i>Enterococcus faecalis</i> , <i>Proteus penneri</i>	<i>Enterococcus faecalis</i> , <i>Proteus hauseri</i>

Table 5. Bacteria cultured from juvenile *Naja nigricincta nigricincta* venom and oropharynx

Snake number	Venom bacteria	Oropharynx bacteria
14, not sexed	<i>Providencia rettgeri</i>	<i>Proteus morganii</i>
15, not sexed	-	<i>Proteus mirabilis</i>
16 female	<i>Enterococcus faecalis</i> <i>Proteus mirabilis</i>	<i>Enterococcus faecalis</i> , <i>Proteus mirabilis</i>
17 female	<i>Enterococcus faecalis</i>	<i>Enterococcus faecalis</i> , <i>Proteus mirabilis</i>
18 female	<i>Streptococcus</i> spp. <i>Proteus mirabilis</i>	<i>Enterococcus faecalis</i> , <i>Proteus mirabilis</i>
19 female	<i>Proteus mirabilis</i>	<i>Proteus mirabilis</i>
20 female		<i>Proteus mirabilis</i>
21 female	<i>Enterococcus faecalis</i> <i>Proteus mirabilis</i>	<i>Staphylococcus epidermidis</i> , <i>Proteus mirabilis</i>
27 female	<i>Providencia</i> spp. <i>Enterococcus faecalis</i>	<i>Salmonella enterica</i> , <i>Pseudomonas aeruginosa</i> , <i>Enterococcus faecalis</i>
28 male	<i>Enterococcus faecalis</i> , <i>Sphingomonas paucimobilis</i> , <i>Proteus vulgaris</i>	<i>Enterococcus faecalis</i> , <i>Salmonella enterica</i>
29 male	<i>Proteus hauseri</i>	<i>Proteus hauseri</i> , <i>Kocuria varians</i>
32 male	No growth	No growth
35 male	<i>Proteus penneri</i> , <i>Enterococcus faecalis</i>	<i>Salmonella enterica</i>
36, not sexed	<i>Proteus hauseri</i>	<i>Proteus penneri</i>
37, not sexed	<i>Proteus vulgaris</i>	<i>Pseudomonas stutzeri</i>
38 female	<i>Proteus vulgaris</i>	<i>Proteus vulgaris</i>
39 female	<i>Salmonella enterica</i>	<i>Salmonella enterica</i>

possibility of a species-specific or host-specific oral microbiome.^[30] Further studies comparing the oral microbiome of different species will clarify this issue.

Male zebra snakes have a much larger home range with possibly more prey diversity than their female counterparts (unpublished data, Mr F Theart, 2022). This may account for the larger number of

adult males caught in this study. These differences could potentially influence variations between male and female oral microbacteria. The sample size from this study was too small to identify any substantial differences between the oral flora of the different sexes.

Whether the bacteria cultured can benefit the host by playing a role in enhancing venom effects or digestion is unclear.^[30,31] *Enterococcus*

faecalis and the Gram-negative bacteria cultured in our study have all been implicated in necrotising fasciitis.^[32] Inoculation of these bacteria into an area of extensive venom-induced destruction and tissue devitalisation, with a resultant soft-tissue infection, will further expand and exacerbate tissue damage, and may even culminate in an infective necrotising fasciitis.^[16-18] In the 'weaponised bacteria theory', Auffenberg^[33] postulated that bacteria in Komodo dragon (*Varanus komodoensis*) venom are a mechanism for prey debilitation and mortality, because of the absence of said bacteria in the reptile's oral microbiome. This raises the question of whether the 'weaponized bacteria theory' may be true for the *Naja nigricincta nigricincta* oral microbiome.^[34]

All the *Enterococcus faecalis* cultured were sensitive to penicillins, all cephalosporins, ciprofloxacin and gentamicin. The Gram-negative bacteria from the adult snakes displayed a resistance of 35.8% against amoxy-clavulanic acid, 78.5% against cephalothin (first-generation cephalosporin) and 60% against cefuroxime (second-generation

cephalosporin). Sensitivity of 92% was displayed towards ceftriaxone (third-generation cephalosporin), 96.3% to piperacillin-tazobactam, and 100% sensitivity to ciprofloxacin and gentamicin (Table 7). The pathogens cultured from the juvenile snakes showed less overall resistance and greater sensitivity towards penicillin and second-generation cephalosporins than their counterparts from adult snakes (Table 7).

Juvenile *Naja nigricincta nigricincta* snakes are very seldom responsible for bites (authors' own clinical experience). Only the data from the adult snake samples were taken into consideration for antibiotic selection suggestions. In view of above results, recommended antibiotic prophylaxis after *Naja nigricincta nigricincta* bites is ciprofloxacin or a third-generation cephalosporin plus gentamicin or piperacillin-tazobactam. Since controversy surrounds the use of ciprofloxacin in children, a third-generation cephalosporin plus gentamicin may be a safer option for the paediatric population.^[35] Piperacillin-tazobactam should be reserved for severely ill patients.

Table 6. Bacterial differentiation: A comparison of the frequency of individual bacteria as cultured from the different samples

Organism	Adult			Juvenile		
	Venom	Oropharynx	Venom and oropharynx	Venom	Oro-pharynx	Venom and oropharynx
	Samples, <i>n</i>		Snakes, <i>n</i>	Samples, <i>n</i>		Snakes, <i>n</i>
	21 <i>n</i> (%)	20 <i>n</i> (%)	21 <i>n</i> (%)	14 <i>n</i> (%)	16 <i>n</i> (%)	16 <i>n</i> (%)
<i>Enterococcus faecalis</i> (G+)	13 (61.9)	13 (65)	15 (71.4)	5 (35.7)	5 (31.3)	7 (43.8)
<i>Morganella morganii</i> (G-)	3 (14.2)	2 (10)	4 (19)			
<i>Proteus species</i> (G-)	13 (61.9)	10 (50)	15 (71.4)	10 (71.4)	12 (75)	14 (87.5)
<i>Pseudomonas</i> spp. (G-)	1 (4.8)	3 (15)	4 (19)			2 (12.5)
<i>Citrobacter freundii</i> (G-)		2 (10)	2 (9.6)			
<i>Acinetobacter</i> (G-)		2 (10)	2 (9.6)			
<i>Yersinia</i> spp. (G-)		1 (5)	1 (4.8)			
<i>Elizabethkingia</i> spp. (G-)		1 (5)	1 (4.8)			
<i>Salmonella enterica</i> (G-)		2 (10)	2 (9.6)	1 (7.1)	4 (25)	4 (25)
<i>Klebsiella pneumoniae</i> (G-)	1 (4.8)		1 (4.8)			
<i>Vibrio</i> spp. (G-)		1 (5)	1 (4.8)			
<i>Providencia</i> spp. (G-)				1 (7.1)		1 (6.3)
<i>Spingomonas</i> spp. (G-)				1 (7.1)		1 (6.3)
<i>Kocuria</i> spp. (G+)					1 (6.3)	1 (6.3)
<i>Staphylococcus epidermidis</i> (G+)					1 (6.3)	1 (6.3)
<i>Streptococcus</i> spp. (G+)				1 (7.1)		1 (6.3)

G+ = Gram positive; G- = Gram negative.

Table 7. Antibiotic profile of organisms cultured from adult and juvenile *Naja nigricincta nigricincta* oropharynx and venom

Antibiotic	All Gram-negative organisms cultured				Enterococcus faecalis (A and J)
	Adult (A)		Juvenile (J)		Sensitive, %
	Sensitive,%	Resistant, %	Sensitive, %	Resistant,%	
Ampicillin	21	79	54.5	45.5	100
Amoxy-clavulanate	56.4	35.8	93	7	100
Cephalothin (1st generation)	21.5	78.5	61	39	
Cefuroxime (2nd generation)	34.3	60	50	50	
Ceftazidime (3rd generation)	89.5	10.5	100		
Cefotaxime (3rd generation)	80	20	100		
Ceftriaxone (3rd generation)	92	8	95		100
Ciprofloxacin	100	-	100		100
Gentamicin	98	-	100		94.5
Piperacillin-tazobactam	96.3	3.7	100		

Limitations

The snakes that were swabbed were not snakes responsible for bites.

This study was conducted from mid-November to mid-April. Seasonal variations in snake oral flora are unknown, and current results may not reflect other time periods. The sample size of 37 snakes is not large, and all the snakes originated from the Khomas region in Namibia. The results may therefore not be representative of the oral flora of all *Naja nigricincta nigricincta* from all geographical areas, or of other spitting cobras.

Practical issues and financial constraints necessitated conventional culture methods, although sequencing techniques would have been able to identify more bacterial species.

Conclusion

Although subject to several limitations, this study has provided an initial baseline database on the oral microbiology of *Naja nigricincta nigricincta* with concomitant antibiotic sensitivities and resistance. This was an important first step in the quest to identify suitable prophylactic and empirical antimicrobial therapy secondary to *Naja nigricincta nigricincta* snakebite. The next step will be to perform a comprehensive standardised study on the microbiology of *Naja nigricincta nigricincta* snakebite wounds.

The antibiotic profile from *Naja nigricincta nigricincta*'s oral microbiome, as cultured in this study, is very similar to the profile of the Gram-negative *Enterobacteriaceae* cultured from both the KZN study and the *Naja nigricincta nigricincta* bite wounds (Tables 1 and 7).^[4] Based on these findings, prophylactic antimicrobial therapy after *Naja nigricincta nigricincta* bites should comprise of ciprofloxacin or a third-generation cephalosporin plus gentamicin or piperacillin-tazobactam.

Declaration. None.

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Author contributions. ES: literature review, protocol, supervise milking procedure, swab venom and snake's oropharynx, data analysis, article compilation. PB: supervisor, oversee data analysis and article compilation. FT: identification, catching, handling, milking and releasing snakes.

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