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Histopathological analysis and *in situ* localisation of Australian tiger snake venom in two clinically envenomed domestic animals

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**Highlights**

- Venom-induced pathological changes were observed in the lungs, kidneys and muscle tissue of both patients
- Evidence, not previously noted, of procoagulant venom effects were apparent, with formed thrombi in the heart, lungs (small fibrillar aggregates and larger, discrete thrombi) and kidneys
- Work has shown pathological evidence of procoagulant venom activity supporting previous suggestions that an initial thrombotic state occurs in envenomed patients
- Illustrates venom localisation in the lung, kidney and muscle tissues of clinically envenomed animals
Histopathological analysis and \textit{in situ} localisation of Australian tiger snake venom in two clinically envenomed domestic animals

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Conflicts of Interest: Nil

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Abstract

Objective: To assess histopathological changes in clinically envenomed tiger snake patients and identify tissue specific localisation of venom toxins using immunohistochemistry.

Samples: One feline and one canine patient admitted to the Murdoch Pet Emergency Centre (MPEC), Murdoch University with tiger snake (Notechis sp.) envenoming. Both patients died as a result of envenomation. Non-envenomed tissue was also collected and used for comparison.

Methodology: Biopsy samples (heart, lung, kidney and skeletal muscle tissue) were retrieved 1-2 hours post death and processed for histopathological examination using Haemotoxylin and Eosin, Martius Scarlet Blue and Periodic Acid Schiff staining. Tissues were examined by light microscopy and tissue sections subjected to immunohistochemical staining using in-house generated monoclonal and polyclonal antibodies against Notechis venoms.

Results: Venom-induced pathological changes were observed in the lungs, kidneys and muscle tissue of both patients. Evidence, not previously noted, of procoagulant venom effects were apparent, with formed thrombi in the heart, lungs (small fibrillar aggregates and larger, discrete thrombi) and kidneys. Immunohistochemical assays revealed venom present in the pulmonary tissue, in and around the glomerular capsule and surrounding tubules in renal tissue and scattered throughout the Gastrocnemius muscle tissue.

Conclusion: This work has shown pathological evidence of procoagulant venom activity supporting previous suggestions that an initial thrombotic state occurs in envenomed patients. We have shown that venom toxins are able to be localised to specific tissues, in this case, venom was detected in the lung, kidney and muscle tissues of clinically envenomed animals. Future work will examine specific toxin localisation using monoclonal antibodies and identify if antivenom molecules are able to reach their target tissues.

Keywords: Tiger snake, Notechis scutatus, snake bite, envenoming, immunohistochemistry, histopathology.

Periodic Acid Schiff, AV: antivenom, OCT: Optimal cutting temperature, APTT: Activated partial thromboplastin time.
1. Introduction

There are up to 3,000 reported snakebites in humans (Sutherland and Leonard 1995; White 2000) and an estimated 6,000 snakebites in domestic animals each year in Australia (Mirtschin et al. 1998), many requiring antivenom (AV) treatment. The most common, potentially lethal clinical feature seen in envenomed humans is venom-induced consumptive coagulopathy (VICC); a result of procoagulant toxins in the venom. The severity and occurrence of major clinical effects seen in human and animal victims appears to differ. As shown by Isbister et al., (2008) (Isbister et al. 2008), 90% of human tiger snake bite patients display VICC, with few presenting with neurotoxic and myotoxic effects. In contrast, canine victims present more often with paralysis, though may also present with clinical signs referable to coagulopathy (Holloway and Parry 1989). This may be a reflection of differences in time since bite to discovery and treatment in the two groups and that VICC may go unnoticed in animals. It is not uncommon for domestic animal envenomings to be discovered hours, sometimes days after the bite or when symptoms are observed, whereas envenomed humans are almost always noticed at the time of first aid investigation, with treatment following shortly thereafter. Cats rarely have clinical signs of coagulopathy but show severe paralysis and myotoxicity. Notably, cats have been reported to be about three times less sensitive (on a weight basis) to both brown and tiger snake venoms than dogs (Moisidis et al. 1996).

Early studies by Kellaway, (Kellaway 1929, 1932, 1934) and more recently work by Harris et al., (Harris et al. 1975, 1981, 2000; Preston et al. 1990) and others, have examined the histopathological effects of Notechis sp. in envenomed animal tissue. In most of these studies the focus has been on the neurotoxic and myotoxic effects of this family of venoms and most involved the examination of experimentally envenomed animal muscle tissue. Similarly, Lewis, (1994) (Lewis 1994), examined skeletal muscle tissue of experimentally envenomed dogs, but also extended his histopathological analysis to the kidney looking at direct nephrotoxic effects of N.scutatus venom. Although these studies have aided our understanding of the mechanisms of actions of some of these neuro- and myotoxins, more specifically, notoxin, none have defined their specific tissue localisation. Moreover, none thus far have investigated the in vivo procoagulant effects of or described the histopathological changes in the heart and lung after Notechis species clinical envenomings.

Numerous studies have looked into the direct effects of snake venom toxins based on clinical outcomes, histopathological changes and in vitro effects on cells. Such studies include the
measurement of various clinical indications, such as serum creatine kinase (CK), creatinine levels for myotoxic effects (Ponraj and Gopalakrishnakone 1997, 1996) and International Normalisation Ratio/ Prothrombin Time (INR/PT), Activated Partial Thromboplastin Time (APTT), D-Dimer levels and whole blood clotting time for coagulopathic envenomings (Ferguson et al. 2002; Isbister et al. 2002, 2006). Testing of venoms in vitro on various animal derived primary and cultured cells (Hodgson and Wickramaratna 2002; Kubo et al. 2002; Kuruppu et al. 2006; Marshall and Herrmann 1989; Rigoni et al. 2004) and on animal nerve-muscle preparations (Ramasamy et al. 2004; Zamuner et al. 2004) have aided our understanding of mechanisms of action and toxicities, while others observed histopathological changes in animal tissues after injection of venom or purified venom toxins (Mandal and Bhattacharyya 2007; Ponraj, 1997) to define sites and mechanisms of action. The majority of these studies have focused on snakes found outside Australia. Where these studies focused on the clinical outcomes and in vitro functions of various snake venoms and their toxins, others have measured crude venom levels in tissues and organs of experimental animals and human autopsy specimens (Brunda et al. 2006; Selvanayagam et al. 1999). No studies to date have detected or identified the specific tissue (cellular) localisation of Australian venom toxins in the tissues or organs of clinically envenomed patients.

Venom toxins thought to be responsible for the most severe clinical effects seen in humans and domestic animals are: (i) prothrombin activating enzymes acting on coagulation factors leading to venom induced consumptive coagulopathy, (ii) pre and post-synaptic neurotoxins that result in paralysis and (iii) myotoxic components that promote muscle cell destruction (or myonecrosis). Therefore we might expect to find venom toxins localised on the endothelium of the vasculature, on neuronal cells and throughout muscle tissue respectively in envenomed patients. In the case of prothrombin activators, demonstration of a substantial amount of the toxin bound to endothelial surfaces would cause us to question the relevance of serum concentrations of this toxin as most of the active toxin may be on the endothelial surface rather than in the blood. The aims of this study were to examine clinically envenomed animal tissue for evidence of procoagulant, neurotoxic and myotoxic venom activity and to use a panel of developed antibodies to identify tissue specific toxin localisation.
2. Materials and Methods

2.1 Materials

Antibodies: mouse anti-CD146 was purchased from Abcam, anti-mouse FITC and anti-rabbit AF488 from Millipore, Streptavidin-Cy3, Hoescht 3342 stain and Protein-G-Sepahrose were obtained from Amersham Biosciences. SuperFrost slides (Ecco) and OCT embedding medium were purchased from ProSciTech, Perth. The SOLV solution (Xylene substitute) was purchased from Scot Scientific, Perth and foetal bovine serum (FBS) was purchased from Gibco.

2.2 Samples

Feline and Canine samples were obtained from two patients with Tiger snake envenoming whom presented to Murdoch Pet Emergency Centre (MPEC), Murdoch University between October 2008 and March 2009. Ethics approval was obtained from Murdoch University (Ref: R2202/08) and The University of Western Australia (UWA) (Ref: F15811). Samples were collected with the owner’s consent. The species of snake responsible was, in both cases, determined by a venom detection kit (VDK), CSL Ltd from patient serum. Control canine and feline tissue was obtained from non-envenomed MPEC patients with ethical approval. Mice were maintained at the Biomedical Research Facility, UWA (Ref: RA/100/593).

2.3 Tissue collection and processing

Within one hour of death, multiple biopsy samples were collected from the envenomed canine and feline patients. Tissues collected were lung, heart, kidney and Gastrocnemius muscle. One sample of each tissue was immediately frozen in OCT medium stepwise on liquid nitrogen and stored at -80°C. The second biopsy sample of each tissue was placed in 10% formalin and maintained at room temperature. Post mortem urine and serum samples were collected from the canine and feline patients as part of their routine treatment.

Frozen canine tissue was trimmed and sectioned (10μm) on a Leica Microtome (Model CM30505) and sections fixed to slides for 5 minutes in ice cold acetone. Fixed slides were then stored at -80°C. Segments of formalin fixed tissue (canine and feline) were oriented to obtain transverse sections and immersed in 50% alcohol. Tissues were processed and stained by the Histology Department (Technologists Michael Slaven and Gerard Spoelstra), School of Veterinary and Biomedical Science, Faculty of Health Science, Murdoch University. In brief, tissues were
processed on a Tissue Tek VIP tissue processor and dehydrated with changes in various concentrations of alcohol. Tissues were then embedded in paraffin using a Leica EG 1150C and 4μm sections cut on a Leica microtome. All paraffin embedded tissue slides were stained with Haematoxylin and Eosin (H&E), Martius Scarlet Blue (MSB) and Periodic Acid Schiff (PAS). All tissue sections were examined by light microscopy.

2.4 Immunohistochemistry
Monoclonal and polyclonal (pAb) antibodies raised to detect *Notechis scutatus* venom (Nsv) and Notexin, both unconjugated and biotinylated, were optimised for use in Immunohistochemistry (IHC) using normal and frozen envenomed mouse tissue sections. Each of the antibodies was tested at varying concentrations (1:50, 1:100 and 1:200). Similarly, secondary antibodies (anti-mouse FITC, anti-rabbit AF488 and Streptavidin-Cy3) were titrated to determine optimal working dilutions. Prior to staining with antibodies, paraffin embedded tissue sections were deparaffinised and subjected to an antigen retrieval. This was accomplished by an initial immersion of the tissue sections in three changes of SOLV (Xylene substitute) solution for 10 minutes. Slides were subsequently transferred into 100% alcohol (3 changes of 3 mins each) followed by 95%, 70% and 50% ethanol respectively for 3 minutes each. Sections were rinsed twice with PBS (5 mins) and processed for antigen retrieval. Tissue sections were immersed in citrate buffer (10mM, pH 6.0) and incubated at 95ºC, 10 mins and allowed to cool for 20 minutes. Sections were washed in PBS and blocked with 10% FBS in PBS. Primary antibodies were then added and incubated overnight at 4ºC in a humidified chamber. Relevant antibody controls were prepared for each tissue type. Tissue sections were washed with 2 changes of PBS (5 mins) followed by the application of secondary antibodies; anti-mouse FITC and Streptavidin-Cy3 required a further 1 hour incubation and two additional washes with PBS. The final wash contained a 1:5,000 dilution of Hoescht stain (Amersham) to enable the visualisation of the cell’s DNA. Slides were viewed on an inverted fluorescent microscope (Olympus, IX71) and images captured with an Olympus (DP70) camera.
3. Results

3.1 Case Histories

3.1.1 Case 1
Collie cross Kelpie canine, 2 years old, Female; presented to MPEC in cardiac and respiratory arrest within 2 hours of suspected envenoming. Upon presentation, the patient was intubated, ventilated, followed by CPR. Despite drug administration (adrenaline, atropine and IV Hartman’s at shock rates) and 4 attempts at defibrillation, there was no response. The owners decided to halt CPR given the poor prognosis. A venom detection kit (VDK) was performed on post mortem serum which was strongly and immediately positive for Tiger snake venom. The circulating serum crude venom (Nsv) concentration (post mortem) was determined to be 22.5ng/ml and notexin level 11.1ng/ml by immunoassay.

3.1.2 Case 2
Domestic Shorthair feline, 1 year old, Male; presented to MPEC early in the morning after having appeared normal the night before. On initial physical examination, the patient was in lateral recumbency with flaccid paralysis of all four limbs. There was generalised Lower Motor Neuron (LMN) paralysis, weak palpebral reflexes (eyelid/eyebrow), dilated pupils and absent pupillary light response (PLR). VDK on urine was strongly positive to Tiger snake venom. APPT was 75 seconds (normal range of 60 to 115) and serum Creatine kinase (CK) was 93,800 U/L (normal range of 50 to 100). Symptoms progressed to severe LMN paralysis and the feline became hypotensive, hypothermic and hypoxic. Urine showed signs of haemoglobinuria and/or myoglobinuria. Treatment included Chlorpheniramine (1mg/kg) IM and 7 vials of antivenom (AV) which included 2 vials of Summerland Serums Tiger/Multibrown AV administered within 15 minutes of the initial exam, a third vial 3 hours later, another vial 3 hours later and a further 3 vials 7 hours after the first dose administered. Each Summerland Serums Tiger/Multibrown AV contains 3000 units Tiger snake AV plus 1500 units of Brown snake AV. At this stage the feline was hypoventilating (PvCO$_2$ = 73mmHg, and ETCO$_2$ = 60mmHg) with decreased respiratory effort and rate, consistent with severe flaccid paralysis progressing to respiratory paralysis. At this point ventilation commenced.

Flaccid paralysis remained through the evening and the patient was maintained under light anaesthesia with alfaxalone by continuous rate infusion (CRI). Mild improvement was seen the following day, with the feline able to move and objecting to the endotracheal (ET) tube but displayed ongoing hypoventilation when attempting to wean from the ventilator. On day 3, CK
levels of 805,100 U/L, AST = 6,440 (normal range 26 to 43), ALT = 1,520 (normal range 0 to 83) with urea and creatine normal. Complete blood count (CBC) showed mild anaemia and a mild thrombocytosis/thrombocythaemia. Mild improvement in neurological status was observed and although the patient could maintain ETCO$_2$ levels within normal ranges for up to 30 minutes, the patient quickly developed respiratory fatigue without ventilator support. On day 4, CK serum concentrations had decreased to 226,720 U/L, AST = 1,520 and ALT = 1,360. The feline went into cardiac arrest during an ET tube change, failed to respond to CPR/adrenaline and subsequently passed away. Serum crude venom (Nsv) concentration (pre treatment) was found to be 28.7ng/ml.

3.2 Histopathological Changes

A summary of the tissue specific histopathological findings are presented in Table 1. Evidence of procoagulant venom effects were observed in the epicardium of the feline heart (Figure 1A and B) and throughout the feline pulmonary tissue (Figure 1C and E) as well as in the vasculature of the renal cortex in the canine patient (Figure 1F), apparent by visibly formed thrombi. The presence of fibrin was confirmed by MSB staining. The lungs of both animals showed multifocal to coalescing alveolar collapse, (Figure 2A and C), hence the lungs were unable to fully inflate, some of which may have been in part due to post-mortem changes. The presence of eosinophilic (proteinaceous) material within the alveoli is consistent with the presence of oedema and small aggregate of fibrillar deposits possibly suggestive of further procoagulant venom activity, indicated by arrows in Figure 2C.

Hydropic changes were observed in kidney tissue of both patients’, a degenerative change where cells swell with fluid was observed, indicative of cells becoming vacuolated. No signs of proteinuria or necrosis were observed in the canine tissue; however, distinct changes indicating increased glomerular permeability were noted in the feline kidney. In the feline occasional fibrillar deposits were seen in the glomeruli; glomerular capillary thrombus formation may be indicative of procoagulant toxin activity. Expanded Bowman’s spaces containing an amorphous pale eosinophilic material, confirmed by PAS staining, was suggestive of proteinuria (Figure 3B and C) in the feline tissue. The appearance of hypereosinophilic epithelial cells without the presence of an accompanying inflammatory response was indicative of mild post-mortem autolysis. No pathological changes were observed in the canine GM tissue, which was not surprising as the time since envenoming to presentation and death was within 3 hours, thereby not allowing the toxins sufficient time to reach the muscle fibres and cause myonecrosis. In contrast, significant changes
were observed in the feline tissue including selective degeneration of myofibres with an
accompanying inflammatory response (Figure 4A and B); and as indicated by arrows in Figures
4C and D, macrophages are seen actively phagocytosing individual necrotic myofibres. Non-
envenomed feline muscle tissue is presented for comparison (Figure 4E and F)

3.3 Tissue localisation of venom toxins

Although minor histopathological observations were made in both cardiac tissues, modest
fluorescence was apparent when staining the tissue with an anti-Nsv IgG pAb. Limited staining of
frozen sections of the entire canine myocardium along with a speckled staining throughout the
cardiac muscle tissue was seen when compared to the secondary alone control (not shown) and the
non-envenomed tissue control. However, when probing the canine pulmonary tissue with the same
pAb, multifocal to coalescing intense staining was observed, indicating the presence of a
significant amount of venom in this tissue (Figure 5A and B). A commercially available anti-
CD146 antibody, reactive with endothelial cells and smooth muscle, also showed limited staining
on paraffin embedded tissue and so it was difficult to determine exactly where the venom
localised. As illustrated in Figure 5A and B, it appeared as though the toxins were not confined to
one specific location (i.e. blood vessels, bronchi or bronchioles), but scattered throughout the
tissue. This is consistent with the appearance of nerve fibres, to be confirmed with further
investigation. All fluorescence observed was absent in secondary alone controls and non-
envenomed tissue sections (Figure 5C and D).

Correlating to histopathological changes observed in the feline kidney, IHC experiments carried
out on frozen canine kidney tissue resulted in anti-Nsv staining in and around the glomerular
capsule in the renal cortex, as well as in surrounding tubules (Figure 6A). No distinct staining
above background fluorescence was observed in the feline kidney, which may be the result of a
combination of the administration of antivenom, the time since envenoming and collection of the
tissue (>4 days), as opposed to the canine tissue, which was collected within 3 hours post
envenomation. Although it may have been too early following envenomation to observe
histopathological changes in the canine GM tissue, IHC assays revealed that venom was present.
Illustrated in Figure 6D and E, using a biotinylated anti-Nsv (then Strep-Cy3), punctate staining
pattern was observed in both patients through the Gastrocnemius muscle tissue itself as well as a
staining of toxins within the vasculature (centre bottom bundle). All staining observed was greater
than the fluorescence intensity of the secondary alone and non-envenomed control tissue (Figure
6C and F) controls at the same exposure.
4. Discussion

The purpose of this work was to initially assess the histopathological changes in several organs of clinically envenomed tiger snake patients, confirmed by VDK, and relevant controls and secondly to conduct preliminary experiments to localise venom toxins in vivo. What become very clear through this work is that histopathological change in tissues in response to Australian elapid envenoming is an area poorly represented in the literature.

This work was conducted in consultation with an experienced Veterinary Specialist in Emergency Medicine and Critical Care plus a Pathologist which has ensured that no signs commonly presenting in normal domestic animals have been misinterpreted as venom induced effects.

4.1 Heart

In the canine tissue, mild, peracute, multifocal haemorrhage was observed (not shown) which is consistent with findings reported by Tibballs et al. (Tibballs et al. 2002). The canine heart tissue showed changes within a short time period (2 hours), therefore suggestive that the coagulopathic venom toxins are fast acting toxins, exerting their effects soon after envenoming. Although there were no signs of cardiac myocyte degeneration (indicative of myotoxic venom activity) in either patient’s heart tissue, a small epicardial thrombus was detected in the feline heart sections, indicative of procoagulant toxin action.

4.2 Lung

The histopathological changes in the lungs of both pulmonary tissues are consistent with the clinical outcomes seen in both patients. Several thrombi were observed within the feline vasculature (Figure 1C-E) which would more than likely manifest clinically as dyspnoea. The feline had been ventilated for several days post envenoming and went into cardiac arrest when being weaned off the ventilator. It is likely that the venous thrombosis was directly caused by the original snake envenoming, but time spent on the ventilator may have contributed to the fibrinous alveolar exudates observed. The pulmonary thrombosis evident was sufficiently significant to be functionally catastrophic. Direct pathological thrombosis, more specifically pulmonary embolism as a result of snake envenoming has only previously been described in cases involving the Martinique vipers and related species from the West Indies (Dong-Zong et al. 2002; Thomas et al. 1995).
It has been suggested that in true coagulopathy a brief window of thrombus formation as the toxin enters the circulation is expected (White 2005), which prior to the activation of fibrinolysis gives rise to the consumption of coagulation factors, seen clinically in patients. Thrombus formation has previously been documented in experimental dogs in response to brown snake envenoming (Tibballs et al. 1991, 1992). This work has also shown pulmonary thrombus formation as well as emboli observed in heart and kidney tissues. It has been suggested that even a few minutes of such thrombotic complications can be devastating for the victim, as the occlusion of critical vessels by thrombi can result in cardiac arrest and death (White 2005).

4.3 Kidney

Although Tiger snake envenoming has been widely reported to cause kidney damage in clinically envenomed patients (Cheng and Currie 2004; Currie 2000; Hood 1975; Jolles et al. 1998; White 1995), it is unclear whether renal failure is a direct result of the venom’s nephrotoxicity or a secondary effect of myoglobinuria. Different types of kidney damage induced by snake venoms and purified venom toxins (from snakes found outside Australia) has been thoroughly reviewed by Sitprija et al., (1982) (Sitprija et al. 1982), allowing us to utilise this work when examining kidney damage caused by Australian snakes.

Interestingly, within the vasculature of the canine renal cortex, a small thrombus/embolus was observed (Figure 1F). Usually, thrombi are attached to the vascular endothelium, however, in this case it is possible that the thrombus detached from the endothelium as an embolus and was freely circulating through the patient’s vasculature. One particular study of the renal effects of tiger snake envenoming in experimental dogs showed the presence of lesions, indicating acute tubular necrosis (Lewis 1994). The author also noted the presence of variable amounts of proteinaceous material within the renal tubules. Our work is consistent with these findings where mild proteinuria was observed within the proximal loops of Henle and the Bowmans’s space of the feline kidney. This is indicative of increased glomerular permeability and is suggestive of direct toxin activity. In the canine kidney, anti-Nsv staining showed a punctuate fluorescence pattern in and around the glomerulus and surrounding tubules suggesting large amounts of venom are being filtered through the kidneys most likely leading to the renal damage identified.
4.4 Muscle

The effects of tiger snake phospholipase A$_2$ myotoxins on muscle fibres have been well documented and shown to produce skeletal muscle paralysis by inhibiting synaptosomal membrane choline transport from nerve endings (Mollier et al. 1990). Harris et al. (Harris et al. 1975, 2003; Harris and MacDonell 1981; White 1987) showed that the purified notexin caused destruction of the myofibres in a mosaic pattern and Preston et al., (1990) (Preston et al. 1990) found a dose-dependent relationship between the extent of muscle loss and the amount of venom injected in experimental animals. The feline muscle tissue in this study also clearly showed myotoxicity with an inflammatory response as defined by Mebs and Ownby, (1990) (Mebs and Ownby 1990) and Harris and Maltin (Harris and Maltin 1982), respectively.

No signs of necrosis were observed in the canine muscle which may be due to the short time frame between envenoming and presentation, as opposed to the feline. Lewis, (1994) (Lewis, 1994) found that after the injection of $N$.scutatus venom into the muscle tissue of dogs, segmental necrosis of the underlying tissue within 1 hr of envenomation was present and the degeneration of myofibres at 3 hrs post bite. A major difference here is that the toxin was delivered locally and directly to the musculature rather than through the circulation as in the clinical patients described here.

In the canine GM muscle, although no histopathological changes were observed, a clear staining pattern was observed. This staining was consistent with the histopathological changes observed in the feline muscle. Venom staining in the canine appeared the brightest in several locations throughout the connective tissue between muscle epimysium with many less bright areas appearing in the endomysium (shown to contain capillaries) and within the muscle fibres themselves. Although not as prominent as in the canine, a mild scattered staining pattern was observed in the feline muscle (not shown). Considering the myonecrosis observed in the feline muscle by standard histopathological examination, it is not surprising that venom was detected within the tissue.
Concluding remarks

In this work it was not anticipated that venom would be detected in the feline tissues as this patient had been treated with 7 vials of antivenom and therefore venom toxins would more than likely already be bound by antibody molecules. In the canine patient however, envenoming and presentation was within 2 hours, with no antivenom treatment and we therefore anticipated venom toxins to be abundant in the canine tissue, which coincides with our results.

This work has described the examination of post mortem heart, lung, kidney and Gastrocnemius muscle tissue from two tiger snake envenomed animal patients admitted to MPEC. For the first time, pathological evidence of procoagulant venom activity has been presented, with numerous thrombi observed in the lungs, heart and kidney of the clinically envenomed patients. This supports the proposal that an initial thrombotic state occurs in snakebite patients that leads to the depletion of vital clotting factors, manifesting in VICC, clinically seen in human and domestic animal victims. A panel of antibodies has been developed against various Australian snake venoms and individual toxins for use in localising venom toxins \textit{in vivo} in envenomed patient tissue. Results presented here suggest the presence of venom toxins in each of the tissues examined and offer a valuable tool in further clarifying toxin localisation.

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Table 1: Histopathological findings of analysed tiger snake envenomed domestic animal tissue
Figure 1 Histopathological evidence of procoagulant venom activity in clinically envenomed animal tissue. A. Feline cardiac tissue, x200, MSB stained. Myocardium (Myo) on the left and Subepicardial adipose tissue (Ep) to the right containing coronary arteries (CA) and vein (CV) surrounded by muscular connective tissue (collagen stains blue). The vein (top right) contains an early thrombus, B. Feline cardiac tissue, x 400, MSB stained. A higher magnification of the thrombus inside the vascular lumen (VL), C. Feline pulmonary tissue, x100, H&E stained. Notice the thickening of the alveolar septa (a) and the thrombi (Thr) in the smaller blood vessels (BV), D. Feline pulmonary tissue, x 200, MSB stained. Large bronchus (Br) filled with eosinophilic material surrounded by Hyaline cartilage (Hc). Above bronchus is a large blood vessel lumen (BVL) containing a large early thrombus, E. x400, MSB stained. Large thrombus in large muscular artery magnified further and composed of erythrocytes (yellow), leukocytes (nuclei brown/black) and fibrin (red/pink) and clusters of platelets appearing amphophilic (pink/purple), F. Canine kidney, x100, MSB stained. Observe the renal cortex blood vessel containing a small thrombus formation (red).
Figure 2 Histopathological changes in envenomed and non-envenomed pulmonary tissue. A. Canine tissue, x100, MSB stained. Notice the general congestion and a blood vessel (BV) filled with erythrocytes (yellow) and two bronchi (BR) filled with eosinophilic (proteinaceous) material and the presence of extravascular erythrocytes, indicative of haemorrhage. B. Canine tissue, x200, MSB stained. Notice hyperaemia (pooling of erythrocytes within the vasculature) within the alveolar septa and alveolar atelectasis (collapse). Also apparent is the proteinaceous, lacy material present in the bronchioles (Br). C. Feline tissue, x 200, MSB stained. Observe multifocal small aggregates of red fibrillar deposits throughout the alveolar septa and occasionally in the alveolar lumen, indicated by arrows. D. x100 and E. x200 Non-envenomed Canine tissue, MSB stained and F. x100, Non-envenomed Feline tissue, MSB stained.
Figure 3 Histopathological observations of tiger snake envenomed feline kidney tissue. A. x400, MSB stained. Notice the irregularly shaped glomeruli (GL) with expanded Bowman’s space (Bs) filled with eosinophilic (proteinaceous) material. Arrows indicate fibrillar deposits (bright red). Proximal convoluted tubules (PT), distal convoluted tubules (DT) and collecting tubules (CT) indicated, B. x200, PAS stained and C. x400, PAS stained. Notice occasional minor proteinuria indicated by lacy material in the proximal loops of Henle (PL) and Bowman’s space (arrows), D. Non-envenomed feline glomerulus, MSB stained, x 200.
Figure 4 Histopathological observations of tiger snake envenomed Feline Gastrocnemius muscle tissue. A. x100, MSB stained. Notice the scattering of necrotic muscle fibres highlighted in blue/purple, B. x200, H&E stained. Muscle fibres with moderate inflammatory infiltrate, C. x200 and D. x400, MSB stained, both showing individual and grouped necrotic fibres and inflammatory infiltrate. All arrows indicate the presence of individual macrophages phagocytosing necrotic muscle fibres. E. x 200, H&E stained, non-envenomed feline muscle and F. x 200, MSB stained non-envenomed feline muscle.
Figure 5 Anti-Nsv staining of tiger snake envenomed and non-envenomed canine lung tissue. Formalin fixed post-mortem pulmonary tissue was processed and embedded in paraffin. Sections were stained with an anti-Notechis scutatus pAb (left) and Hoechst stain, a merged image is presented (centre). A. and B. offer two distinct fields of view at x 400 magnification. Notice the antibody staining (red) and cell nuclei (blue). C. Non-envenomed canine lung, x 200, stained with anti-NsPA/Strep-Cy3 and D. Hoechst stained non-envenomed canine lung, x 200.
Figure 6 Anti-Nsv staining of tiger snake envenomed canine kidney and muscle tissue. x400. A. Renal tissue stained with anti-Nsv pAb followed by anti-rabbit AF488. Notice areas of bright staining in and around the glomerulus (GL) and surrounding tubules. Bowman’s space (Bs) is also indicated. B. Hoechst stain and C. anti-rabbit AF488 secondary alone control. D. GM tissue, x200, stained with anti-Nsv pAb Biotin followed by Streptavidin-Cy3. Notice the bundle of cellular material within the vascular endothelium as well as punctate staining throughout the muscle tissue. E. Hoechst stain and F. x100, Non-envenomed canine muscle tissue stained with anti-Nsv pAb Biotin followed by Streptavidin-Cy3.
Figure 1
Click here to download high resolution image
Figure 2
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Figure 3

A

B

C

D