

JOINT TRAUMA SYSTEM



BITES, STINGS, AND ENVENOMATION

CLINICAL PRACTICE GUIDELINE (CPG) TRAINING

Joint Trauma System Trauma Care Educational Program



DISCLOSURE/DISCLAIMER



- ◆ No financial disclosures
- ◆ The view(s) expressed herein are those of the author(s) and do not reflect the official policy or position of Brooke Army Medical Center, the U.S. Army or Air Force Medical Department, the U.S. Army or Air Force Office of the Surgeon General, or the Department of Defense or the U.S. Government.

AGENDA



- ◆ Purpose
- ◆ Summary
- ◆ Evaluation of Bites
- ◆ Mammalian Bites Treatment
- ◆ Marine Bites Treatment
- ◆ Arthropod Bites Treatment
- ◆ Envenomation Treatment
- ◆ Performance Improvement (PI) Monitoring
- ◆ References
- ◆ Appendices
- ◆ Contributors

PURPOSE



- ◆ These slides are based on the JTS Global Spider and Scorpion Envenomation Management CPG which gives an overview of spider and scorpion envenomation and presents a standardized approach to providers in the evaluation and treatment of patients with spider or scorpion induced poisoning.
- ◆ Date of CPG publication: 09 Feb 2021
- ◆ JTS CPGs are evidence-based guidelines developed by subject matter experts in the military and civilian communities. CPGs are compiled from DoD Trauma Registry data, health data abstracted from patient records and after action reports.
- ◆ Information contained in this presentation is only a guideline and not a substitute for clinical judgment.

SUMMARY



- ◆ Differences exist in treatment of bites from different species.
- ◆ Envenomation requires aggressive medical care.
- ◆ Antivenom is available for some species; data regarding the benefits and risks of many of these antivenoms are significantly limited.

EVALUATION OF BITES



- ◆ Evaluate all bites and stings from vertebrate organisms by plain radiograph as barbs and teeth are frequently left in patients after attacks.
- ◆ Address tetanus and rabies status (only for mammalian bites).
- ◆ Assess type of environment.
 - ◆ Marine environment can have decompression sickness if deep enough.
 - ◆ Some infectious organisms are more common in certain conditions than others.

EVALUATION OF BITES



Timing and identification of organism is important, if possible.

- ◆ Unknown timing and source common for arthropod bites
- ◆ If arthropod bite, neurotoxic/allergic effects will be immediate, loxoscelism will be delayed presentation.

TREATMENT OF MAMMALIAN BITES



- ◆ Aggressive washout and debridement as indicated
 - ◆ Delayed primary closure or healing by secondary intention safest
 - ◆ Close follow-up for 1 to 2 days
- ◆ Consider rabies vaccine and rabies immunoglobulin.
- ◆ Antibiotic prophylaxis, typically amoxicillin-clavulanate, for three days is recommended for high-risk wounds.
 - ◆ Associated significant crush injury
 - ◆ Deep puncture
 - ◆ Cat bites
 - ◆ Bites near joints, hands, face, or genitalia
 - ◆ Wounds requiring closure

TREATMENT OF MARINE BITES



- ◆ Aggressive washout and debridement (removal of barbs and teeth) as indicated
- ◆ Treatment for decompression sickness if applicable (generally happens within 48 hrs of rapid ascent below 30 ft)
- ◆ Antibiotic prophylaxis includes generally trimethoprim-sulfamethoxazole, ciprofloxacin, or doxycycline for 3 days
- ◆ Antivenins for box jellyfish and stonefish potentially available if applicable

TREATMENT OF ARTHROPOD BITES



- ◆ Neurotoxic and anaphylaxis effects from various scorpions, spiders, and other insects can happen.
 - ◆ Transfer to facility with appropriate antivenin if signs of systemic illness if applicable.
- ◆ Loxoscelism is generally the only bite that needs addressing surgically.
 - ◆ Commonly confused with cellulitis when patient did not know he/she was bit.
 - ◆ Typically, does not heal and worsens with appropriate course of antibiotics.
 - ◆ Will demarcate in 1-2 weeks after which debridement; closure or possible skin grafting is required.
 - ◆ Administer antibiotics only for signs of infection.
 - ◆ Symptomatic support of systemic symptoms.

ENVENOMATION OVERVIEW



◆ Local Manifestations

burning pain within minutes, edema, erythema, swelling, ecchymosis, hemorrhagic bullae, lymphangitis/lymphadenopathy, necrosis (late finding)

◆ Systemic Manifestations

nausea and vomiting (earliest findings) weakness, headache, tachycardia, paresthesias, bulbar symptoms, diplopia, twitching, consumptive coagulopathy, rhabdomyolysis, muscle paralysis, renal failure, capillary leakage, pulmonary edema, hypotension, and shock

ENVENOMATION EVALUATION (I)



- ◇ Presume all snake bites are venomous and send the patient to a location with antivenin and intensive care capabilities.
 - ◆ A significant minority are dry bites (no venom).
 - ◆ If no symptoms, admit the patient to a ward and watch for 24 hours. If still no symptoms, the patient can be safely discharged.
- ◇ Admit those with symptoms/signs to ICU.

ENVENOMATION EVALUATION (2)



- ◆ Initial evaluation should include trauma assessment.
- ◆ Additional labs/tests (when possible) should include:
 - ◆ Complete blood count
 - ◆ Prothrombin time/international normalized ratio (PT/INR)
 - ◆ Fibrinogen
 - ◆ EKG/electrocardiogram
 - ◆ Creatine kinase
 - ◆ Complete metabolic panel
 - ◆ Urine protein/blood/myoglobin
- ◆ Laboratory derangements will help guide supportive care.

ENVENOMATION TREATMENT



Initial treatment is supportive outside of antivenin

- ◆ Remove constricting clothing and clean wound,
- ◆ Mark site of the bite to demarcate initial erythema and swelling.
- ◆ Avoid FFP, cryoprecipitate and platelets as they may worsen consumptive coagulopathy.

PI MONITORING



◆ Intent (Expected Outcomes)

- ◆ Rapid evaluation and transfer to site with antivenin capability for envenomation
- ◆ Tetanus, rabies, and antibiotic prophylaxis when appropriate

◆ Performance/Adherence Metrics

- ◆ Transfer of patients with moderate to severe symptoms (grades 3 and 4) to antivenom if not available at site
- ◆ Aggressive use of benzodiazepines as indicated for agitation, neuromuscular stimulation, tachycardia, and hypertension
- ◆ Tetanus prophylaxis for all bites and stings.

◆ Data Source

- ◆ Patient Record
- ◆ Department of Defense Trauma Registry

REFERENCES (I)



1. Warrell DA. Venomous bites, stings, and poisoning. *Infectious Disease Clinic of North America*. June 2012; Vol 26(2):207-223.
2. Garb JE, Hayashi CY. Molecular evolution of α -latrotoxin, the exceptionally potent vertebrate neurotoxin in black widow spider venom. *Molecular Biology Evolution*. May 2013; Issue 30(5):999-1014.
3. Maretic Z. Latrodectism: variations in clinical manifestations provoked by *Latrodectus* species of spiders. *Toxicon*. 1983; Vol 21, Issue (4):457-66.
4. Halmo LS, Hurst IA, Ng PC, Wang GS. *Latrodectus Facies* after *Latrodectus Hesperus* Envenomation in a Pediatric Patient. *Journal of Emergency Medicine*. Oct 2019; Vol 57, Issue (4):523-526.
5. Ramialiharisoa A, de Haro L, Jouglard J, Goyffon M. Le latrodectisme à Madagascar [Latrodectism in Madagascar]. *Medecine Tropicale (Mars)*. Dec 1993; Vol 54, Issue (2):127-30.
6. Dart RC, Bush SP, Heard K, et al. The efficacy of antivenin *latrodectus* (black widow) equine immune F(ab')₂ versus placebo in the treatment of latrodectism: a randomized, double-blind, placebocontrolled, clinical trial. *Annals of Emergency Medicine*. Sep 2019; Vol 74, Issue (3):439-449.
7. Offerman SR, Daubert GP, Clark RF. The treatment of black widow spider envenomation with antivenin *latrodectus mactans*: a case series. *The Permanente Journal*. Summer 2011; Vol 15, Issue (3):76-81.
8. Nordt SP, Clark RF, Lee A, Berk K, Lee Cantrell F. Examination of adverse events following black widow antivenom use in California. *Clinical Toxicology (Phila)*. Jan 2012; Vol 50(1):70-73.
9. Sánchez EE, Migl C, Suntravat M, Rodriguez-Acosta A, Galan JA, Salazar E. The neutralization efficacy of expired polyvalent antivenoms: An alternative option. *Toxicon*. Oct 2019; Vol 168:32-39.
10. Isbister GK, O'Leary M, Miller M, Brown SG, Ramasamy S, James R, Schneider JS. A comparison of serum antivenom concentrations after intravenous and intramuscular administration of redback (widow) spider antivenom. *British Journal of Clinical Pharmacology*. Jan 2008; Vol 65(1):139-43.
11. Goddard J. *Physician's guide to arthropods of medical importance*. CRC press; 19 Apr 2016.
12. Chaim OM, Trevisan-Silva D, Chaves-Moreira D, et al. Brown spider (*loxosceles* genus) venom toxins: Tools for biological purposes. *Toxins (Basel)*. Mar 2011; Vol 3(3):309-344.
13. Kurpiewski G, Forrester LJ, Barrett JT, Campbell BJ. Platelet aggregation and sphingomyelinase D activity of a purified toxin from the venom of *Loxosceles reclusa*. *Biochimica et Biophysica Acta (BBA)*. 18 Dec 1981; Vol 678(3):467-476.

REFERENCES (2)



14. Robinson JR, Kennedy VE, Doss Y, Bastarache L, Denny J, Warner JL. Defining the complex phenotype of severe systemic loxoscelism using a large electronic health record cohort. *PLoS One*. 19 Apr 2017 ; Vol 12(4):e0174941.
15. Swanson DL, Vetter RS. Bites of brown recluse spiders and suspected necrotic arachnidism. *New England Journal of Medicine*. 17 Feb 2005; Vol 352 (7):700-707.
16. Said A, Hmiel P, Goldsmith M, Dietzen D, Hartman ME. Successful use of plasma exchange for profound hemolysis in a child with loxoscelism. *Pediatrics*. Nov 2014; Vol 134(5):e1464-7.
17. Kong EL, Hart KK. Tarantula Spider Toxicity. 31 May 2020. In: StatPearls [Internet]. Treasure Island (FL): StatPearls Publishing.
18. Nimorakiotakis B, Winkel KD. The funnel web and common spider bites. *Australian Family Physicians*. April 2004; Vol 33(4):244-251.
19. Hodgson WC. Pharmacological action of Australian animal venoms. *Clinical and Experimental Pharmacology and Physiology*. Jan 1997; Vol 24(1):10-17.
20. Isbister GK, Sellors KV, Beckmann U, Chiew AL, Downes MA, Berling I. Catecholamine-induced cardiomyopathy resulting from life-threatening funnel-web spider envenoming. *The Medical Journal of Australia*. 5 Oct 2015; Vol 203(7):302-304.
21. Braitberg G, Segal L. Spider bites - Assessment and management. *Australian Family Physicians*. Nov 2009; Vol 38(11):862-7.
22. Chippaux JP, Goyffon M. Epidemiology of scorpionism: a global appraisal. *Acta Tropica*. Aug 2008; Vol 107, Issue 2:71-79.
23. Goyffon M, Vachon M, Broglio N. Epidemiological and clinical characteristics of the scorpion envenomation in Tunisia, *Toxicon*. 1982; Vol 20:337-344.
24. O'Connor A, Ruha AM. Clinical course of bark scorpion envenomation managed without antivenom. *Journal of Medical Toxicology*. Sep 2012; Vol 8(3):258-262.
25. Bahloul M, Chaari A, Dammak H, et al. Pulmonary edema following scorpion envenomation: mechanisms, clinical manifestations, diagnosis and treatment. *International Journal of Cardiology*. 10 Jan 2013; Vol 162(2):86-91.
26. Dehghani R, Charkhloo E, Seyyedi-Bidgoli N, et al. A Review on Scorpionism in Iran. *Journal of Arthropod-Borne Disease*. 25 Dec 2018; Vol 12(4):325-333. eCollection 2018 Dec.

REFERENCES (3)



27. Cupo P. Clinical update on scorpion envenoming. *Revista da Sociedade Brasileira Medicina Tropical*. Nov-Dec 2015; Vol 48(6):642-649.
28. Dizaji R, Sharafi A, Pourahmad J, et al. The effects of *Hemiscorpius lepturus* induced-acute kidney injury on PGC-1 α gene expression: From induction to suppression in mice. *Toxicon*. 30 Jan 2020; Vol 174:57-63.
29. Chakroun-Walha O, Karray R, Jerbi M, et al. Value of troponin levels in the diagnosis of cardiac dysfunction in moderate scorpion envenomation. *Human and Experimental Toxicology*. Jun 2018; Vol 37(6):580-586.
30. Sagheb MM, Sharifian M, Moini M, Sharifian AH. Scorpion bite prevalence and complications: report from a referral centre in southern Iran. *Tropical Doctor*. Apr 2012; Vol 42(2):90-91.
31. Rodrigo C, Gnanathanan A. Management of scorpion envenoming: a systematic review and metaanalysis of controlled clinical trials. *Systematic Reviews*. 8 Apr 2017; 6(1): Article 74.
32. Prasad R, Mishra OP, Pandey N, Singh TB. Scorpion sting envenomation in children: factors affecting the outcome. *Indian Journal of Pediatrics*. May 2011; Vol 78(5):544-548.
33. Abroug F, ElAtrous S, Nouira S, et al. Serotherapy in scorpion envenomation: a randomised controlled trial. *Clinical Trial Lancet*. 11 Sep 1999; Vol 354, Issue (9182):906-909.
34. Boyer L, Degan J, Ruha A-M, et al. Safety of intravenous equine F(ab')₂: Insights following clinical trials involving 1534 recipients of scorpion antivenom. *Toxicon*. Dec 2013; Vol 76:386-393.
35. León G, Herrera M, Segura Á, Villalta M, Vargas M, Gutiérrez JM. Pathogenic mechanisms underlying adverse reactions induced by intravenous administration of snake antivenoms. *Toxicon*. 15 Dec 2013; Vol 76:63-76.
36. Dhimi S, Panesar SS, Roberts G, et al. Management of anaphylaxis: a systematic review. *Allergy*. Feb 2014; Vol 69(2):168-175.
37. Simons FER, Ebisawa M, Sanchez-Borges M, et al. 2015 update of the evidence base: World Allergy Organization anaphylaxis guidelines. *World Allergy Organization Journal J*. 2015; Vol 8(1):32.
38. Simons FER, Arduoso LRF, Bilò MB, et al. 2012 Update: World Allergy Organization Guidelines for the assessment and management of anaphylaxis. *Current Opinion Allergy Clinical Immunology*. Aug 2012; Vol 12(4):389-399.
39. Isbister GK, Bawaskar HS. Scorpion envenomation. *New England Journal of Medicine*. 31 July 2014; Vol 371(5):457-63



LIST OF APPENDICES IN CPG

- ◆ **Appendix A: Medical Facilities and Stocked Antivenins**
- ◆ **Appendix B: CROFAB Treatment Algorithm**
- ◆ **Appendix C: Additional Information Regarding Off-label Uses in CPGs**

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