Presented at the ACMT Annual Scientific Meeting 2021 – Virtual

Published in J Med Tox 2021; 17:94

003. Late Hemotoxicity following North American Rattlesnake Envenomation Treated with Crotalidae Immune F(ab')2 (Equine) Antivenom (Anavip®)

Meghan B Spyres^{1,2}, Gabriel Padilla³, Richard D Gerkin^{1,2}, Anne- Michelle Ruha^{1,2}, On Behalf of the ToxIC Investigators Consortium (ToxIC)

Background: Late hemotoxicity is common following rattlesnake envenomation treated with Fab antivenom (CroFab®). Clinical trials showed Anavip® to be superior to Crofab® in preventing late hemotoxicity, but this effect has not been demonstrated in broader populations. Hypothesis: Patients with rattlesnake envenomation treated with Anavip® will have lower incidence of late hemotoxicity than those treat- ed with Crofab®.

Methods: This is a multicenter analysis of prospectively collected data from patients with snakebite reported to the ToxIC NASBR spanning 2019. Inclusion criteria included administration of antivenom for rattlesnake envenomation. Cases were excluded for incomplete data and if less than one set of follow-up laboratory values was obtained. Data collected included demographics, envenomation characteristics, laboratory values, and treatment ad- ministered. Statistics: t-test and Fisher's exact test. Hemotoxicity was defined as platelets <120 K/mm³ or fibrinogen < 170 mg/dL; hemotoxicity was considered late on follow-up if values met threshold and were decreased from discharge value.

Results: One hundred thirteen rattlesnake bites receiving antivenom were reported to the NASBR in 2019, 111 were included. Twenty-seven received Anavip® alone, 50 received Crofab® alone, and 34 received both antivenoms. Baseline patient and envenomation characteristics were similar between the groups. For single antivenom cases, Anavip® cases received more vials than Crofab® cases (17 vs. 11; p < 0.001), but discrete dosing occurrences were similar (2.6 vs. 2.9 events; p = 0.11). Late hemotoxicity occurred in 13 cases in the Crofab® group (35.1%) and in 0 cases in the Anavip® group (0.0%, 95% CI 0.00–0.16; p = 0.001); 2 (5.4%) Crofab® cases required readmission and retreatment; no Anavip® cases were readmitted or retreated for hemotoxicity. One case (3.2%) of late hemotoxicity was reported in patients receiving both Anavip® and Crofab®.

Conclusion: Late hemotoxicity was not detected in NASBR Registry patients treated with Anavip® alone after rattlesnake envenomation; Crofab®-associated late hemotoxicity was similar to previous years.

¹Banner University Medical Center Phoenix, Phoenix, AZ, USA.

²University of Arizona College of Medicine Phoenix, Phoenix, AZ, USA.

³University of Southern California Keck Medical School, Los Angeles, CA, USA