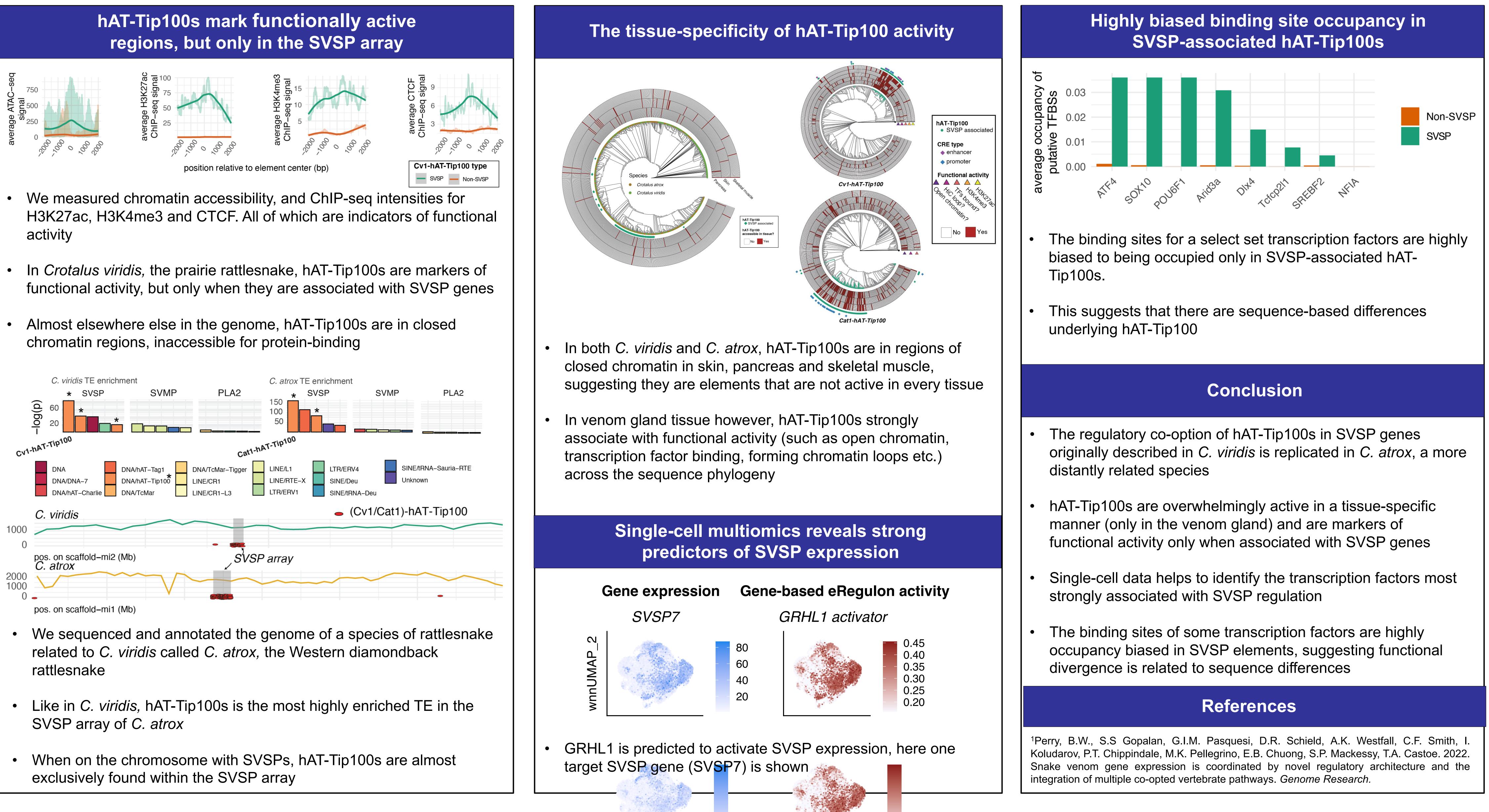
## The diversification of viperid venom genes highlights genomic mechanisms underlying transposon-mediated cis-regulatory element evolution

## and T.A. Castoe<sup>1</sup>

<sup>1</sup>University of Texas at Arlington, Arlington, TX; <sup>2</sup>University of California, Santa Cruz, CA; <sup>3</sup>Broad Institute of Massachusetts Institute of Technology, Cambridge, MA; <sup>4</sup>University of Northern Colorado, Greeley, CO; <sup>5</sup>AntlerA Therapeutics, San Carlos, CA

## Abstract

hAT-Tip100 transposable elements (TEs) are known to have been co-opted as regulatory loci (i.e., promoters and enhancers) for the snake venom serine protease (SVSP) gene family in a single viper species<sup>1</sup>. This is notable because TEs are sequences that are usually silenced after insertion to protect against their many deleterious effects (e.g., disruption of normal gene regulation or protein-coding potential, ectopic recombination etc.). However, the process by which a normally silenced element becomes one that has a defined regulatory role is not well understood. Here, we leverage functional and comparative genomics applied to the hAT-Tip100 SVSP gene regulatory system of rattlesnakes to understand the genomic drivers of TEdriven regulatory element evolution.



S.S. Gopalan<sup>1</sup>, S.N. Smith<sup>1</sup>, B.W. Perry<sup>2</sup>, K. Ballard<sup>1</sup>, Y.Z. Francioli<sup>1</sup>, C. Kim<sup>1</sup>, E. Betran<sup>1</sup>, J.P Demuth<sup>1</sup>, D.C Card<sup>3</sup>, S.R. Kerwin<sup>4</sup>, S.P. Mackessy<sup>4</sup>, S. Seshagiri<sup>5</sup>,



UNIVERSITY OF TEXAS ARLINGTON