Comparative proteomics and transcriptomics of *Nephilingis cruentata* spider venom and glands

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Introduction: The Nephilidae family comprises 61 spiders, including the highly synantropic genre Nephilingis, commonly characterized by large webs and sexual size dimorfism. remarkable Despite some few studies about characterization of acylpolyamines in venom, there is a lack of studies of Nephilingis cruentata venom proteome and transcriptome. Objectives: Investigate the N. cruentata venom and gland proteome by data dependent LC-MS/MS acquisition and transcriptomics comparing protein matches among biological samples. Material and Methods: Venom extracted by a low voltage electrical stimulator and gland supernatant pools were separated, centrifuged and stored at -80°C until digestion. Disulfide bonds were reduced and alkylated with dithiothreitol and iodoacetamide, respectively and pools were digested with trypsin at 37°C overnight. Peptides were separated in 75 µm x 100 mm capillary columns by a linear gradient of 5-35% of acetonitrile and 0.1% of formic acid, ionized by ESI and analyzed in data dependent acquisition mode in a LTQ Orbitrap Velos. Proteins were identified in PEAKS Studio against the compiled transcriptome database. **Results and Discussion:** A total of 33.108 predicted proteins were translated from the venom gland transcriptome and 406 proteins were found in N. cruentata venom and gland proteomes. 276 proteins were unique for glands, as arginine kinase, while 105 proteins were exclusively found in venoms, such as astacin-like metalloprotease toxin. Only 25 out of 406 proteins were common to both groups, showing a wide diversity of proteins in pools of venom and glands of *N.cruentata*. **Conclusions:** The findings in *N.cruentata* venom show a great diversity of proteins in gland and venom. That diversity may be related to proteins with metabolism functions in glands but also with roles in the maturation of venom toxins.