

Cytotoxic Effect of *Rhopalurus junceus* Scorpion Venoms on the HeLa Cell Line

Georgenis García-Oliva

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Abstract

Introduction: Venom composition in both sexes of *Rhopalurus junceus* scorpion has the same major components but it is different in many compounds. However, it is unknown whether the sex of this scorpion influences in its biological activity. **Objective:** To compare the cytotoxic effect of this scorpion from both sexes on the HeLa tumor cells. **Materials and Methods:** Protein content and gel image were analyzed by SDS-PAGE and Image J 1.46 software, respectively. HeLa and Vero cells were treated with each sex and a mixture of both sexes of this scorpion venom. Percentage of cell viability and the morphological changes were determined by MTT assay and phase contrast microscopy, respectively.

Index terms— *Rhopalurus junceus* scorpion venom, female, male, mixture of two sexes, heLa.

1 Introduction

The scorpion venom is a highly complex and heterogeneous mixture of compounds, mainly proteins and peptides (Ahmadi et al., 2020). Around 2000 scorpion species have been described and only 30 species of the Buthidae family are considered dangerous to humans (Desales-Salazar et al., 2020). In a study carried out with two scorpions from Buthidae family: *Androctonus finitimus* and *Hottentota tumulus*, it was shown that the quantity and quality of extracted venom were associated with temperature, diet and the extraction method (Tobassum et al., 2018). Besides, venom composition can be influenced by different factors like sex, geographical location, age, time intervals of extraction and others (Pucca et al., 2014). For example, males and females of *Tityus nororientalis* scorpions produce venoms with different composition and activity (De Sousa et al., 2010). venom decreases the viability of tumor cells of epithelial origin and has no cytotoxic effect on normal cells (Díaz-García et al., 2013). In a proteomic comparative analysis of male and female *R. junceus* scorpion venom, from their 200 components just 63 were common and the most abundant component appeared in both sexes (Rodríguez-Ravelo et al., 2015). Previous studies by our group, using a mixture of venom of female and male scorpions in the same proportion and similar laboratory conditions have demonstrated their cytotoxic and apoptotic effect against cancer cells (Díaz-García et al., 2013; Díaz-García et al., 2015; Díaz-García et al., 2017; Yglesias-Rivera et al., 2019). Considering the described scenario, the objective of the present study was determine if there are differences among *R. junceus* scorpion venoms from female and males individually and mixture.

2 II.

3 Methods

4 a) Scorpion Venom source

Female and male adults of *Rhopalurus junceus* scorpions, collected in Isla de la Juventud (Cuba), were kept in captivity for at least one month before venom extraction by electrical stimulation. Scorpions were maintained under Bioterium conditions in individual ancer is among the leading causes of death worldwide and has a major impact in both developed and underdeveloped countries (Siegel et al., 2020). Specifically, cervical cancer is one of the main public health problem affecting middle-aged women, particularly in developing countries (Arbyn et al., 2020). Conventional antitumor therapies used in clinical practice are surgery, radiotherapy and chemotherapy

43 (Somayeh et al., 2017). These treatments are effective only to some extent, as they are not applicable in all
44 cases and the undesirable side effects often make them impractical (Topcul and Cetin, 2014) *Rhopalurus junceus*
45 (*R. junceus*) belongs to Buthidae family and is an endemic scorpion from Cuba. Preclinical studies have shown
46 that *R. junceus* scorpion Keywords: *Rhopalurus junceus* scorpion venom, female, male, mixture of two sexes,
47 HeLa. plastic containers at $23 \pm 1^\circ\text{C}$ temperature, $60 \pm 10\%$ relative humidity and 12:12 h light-dark cycle,
48 in the laboratories belonging to the Entrepreneurial Group of Biopharmaceuticals and Chemicals Productions
49 (LABIOFAM). Bioterium conditions about management of scorpion colonies and collection of venom have been
50 approved by the Ministry of Science, Technology and Environment of Cuba (CITMA 20/2016). Three groups
51 of scorpions containing 50 female (G1), 50 male (G2) or a mixture of 25 females with 25 males (G3) were used.
52 Venom was dissolved in distilled water and centrifuged at 15000xg for 15min. The supernatant was filtered by
53 using a 0.2 μm syringe filter and stored at -20°C until used. The protein concentration was calculated by Lowry
54 Modified Method (Herrera et al., 1999).

55 **5 b) SDS-PAGE and determination of molecular weight (MW)**

56 Electrophoretic analysis of each pooled venom was carried out according to the previous method (Díaz-García et
57 al., 2015) with 4% stacking gel and 16% separating gel under non-reduced and reduced (2mercaptoethanol, 95°C ,
58 10 min) conditions using an electrophoresis chamber (Biorad). All samples were dissolved in a sample buffer
59 (50mM Tris-HCl, pH 6.8, 0.1M DTT, 10% glycerol, 2% SDS, and 0.1% bromophenol blue). In each well, 50 μg of
60 venom was applied and a protein MW marker was used. The run conditions were 120 V to free current for two
61 hours. The gels were stained with Coomassie Brilliant Blue G-250 and were subsequently rinsed with Methanol:
62 Acetic acid: Water (45:10:45). The gels were photographed and analyzed using ImageJ 1.46 software.

63 **6 c) Cell line and culture**

64 HeLa (cervix adenocarcinoma ATCC CCL-2?) cell line was maintained in minimum essential medium (MEM).
65 Vero (normal African green monkey kidney ATCC CRL-1586?) cell line was maintained in Dulbecco's modified
66 Eagle's medium. The mediums of both cell lines were supplemented with 2 mM of glutamine and non-essential
67 amino acids, 10% of fetal bovine serum (SFB) and penicillin-streptomycin 100 UI/mL -100 $\mu\text{g}/\text{mL}$. The cells
68 were grown in a humidified atmosphere, 5% CO_2 at 37°C .

69 **7 d) In vitro cell viability assay (MTT assay)**

70 The effect of scorpion venom on cell viability was determined by the MTT Assay (Mosmann, 1983). HeLa
71 cells (1×10^4 /well) and Vero cells (1×10^4 /well) were plated in 50 μl of medium/well in 96-well culture plates
72 (Costar Corning, Rochester, NY) and incubated overnight in a humidified atmosphere of 5% (v/v) CO_2 at 37
73 $^\circ\text{C}$. After incubation, 50 μl of venom was dissolved in medium at final concentration of 0.0625, 0.125, 0.25, 0.5
74 and 1mg/mL and was added in five well for every concentration. Cells without scorpion venom were used as
75 untreated control. After 72h of incubation, 10 μl of 5mg/mL of sterile MTT was added per well and incubated for
76 another 3h. The supernatant was carefully removed, 150 μl DMSO was added per well and incubated for 15min
77 at 37°C . The absorbance was determined in a microplate reader (ELISA MRX Revelation Dynex Technologies
78 560nm with 630nm as reference). Absorbance from untreated cells was considered as 100% of growth and used
79 for viability calculation. The effect of scorpion venom on the viability for human cell lines panel was expressed as
80 the percentage of viability, using the formula: $\% \text{viability} = \frac{A_{560-630\text{nm}} \text{ of treated cells}}{A_{560-630\text{nm}} \text{ of control}}$
81 cells $\times 100\%$. The IC₅₀ values (venom concentration that causes 50% reduction of the cell) from cancer cells
82 were determined. The experiments were performed three times by triplicate.

83 **8 Phase-contrast microscopy**

84 After treatments, cells were washed with PBS and morphological changes in culture were then observed under
85 microscope IX-71 (Olympus Corporation, Tokyo, Japan). Images were captured using the camera DP-72
86 (Olympus Corporation, Tokyo, Japan) and 10X objectives.

87 **9 f) Statistical analysis**

88 Kruskal-Wallis non-parametric test and Dunn's multiple comparison tests was used to compare different assays.
89 Two-way Anova and Bonferroni posttest were performed to analyze differences in MW, protein band intensity
90 and cell viability. The IC₅₀ value was determined by interpolation of tendency line from linear regression curve.
91 GraphPad Prism version 5.01 for Windows, (GraphPad Software, San Diego California, USA) for $p < 0.05$ was
92 used for all analysis.

93 **10 III.**

94 **11 Results**

95 There was no significant difference among total protein concentration of female (G1), male (G2) and the mixture
96 of both sexes (G3) of *R. junceus* scorpion venom in our experimental conditions (Figure 1). The values of Mean

97 \pm SD for G1, G2 and G3 were 8.3 ± 1.1 mg/mL, 8.5 ± 2.9 mg/mL and 8.2 ± 1.6 mg/mL, respectively. However,
98 the electrophoretic analysis of protein content from G1, G2 and G3 under non-reduced and reduced conditions
99 demonstrated similarities and differences among experimental groups (Figure ??). The graphic represents the
100 mean \pm SD of total protein concentration values of three independent experiments with female (G1), male (G2)
101 and female + male (G3) *R. juncus* scorpion venom. Data were analyzed using Kruskal-Wallis followed by Dunn
102 test.

103 Figure ??: SDS-PAGE analysis of male, female and the mixture of both sexes *R. juncus* scorpion venom. A)
104 Nonreduced SDS-PAGE. B) Reduced SDS-PAGE. G1: female, G2: male, G3: female + male *R. juncus* scorpion
105 venom. Separating gel 16%, stacking gel 4%. Molecular weight protein marker from 10 kDa-175 kDa was used
106 Six bands were observed in non-reduced (Figure ??A) and reduced (Figure ??B) conditions of SDS-PAGE, in
107 the venom of female and male scorpions. While in the case of the venom obtained from the mixture of scorpions
108 of both sexes, six bands were observed under non-reduced conditions and seven under reduced conditions. The
109 comparison of molecular weight (MW) and protein band intensity among the groups of *R. juncus* scorpion are
110 presented in Table ?. Non-reduced electrophoresis conditions showed five similarities in the MW of (55, 45, 39,
111 28 and 19 kDa) from G1 and G2 scorpion groups. However, in these conditions was observed a difference between
112 them. A band of 11kDa was displayed only in G2 and 12kDa band only in G1 and G3. The molecular weights
113 observed in non-reduced conditions of the mixture of both sexes were: 52, 45, 39, 28, 19 and 12 kDa.

114 Electrophoresis under reduced conditions displayed two resemblances (at 44 and 11 kDa) and four differences
115 between the protein MW of G1 and G2 groups. An 11kDa coincident band was observed in all groups. There
116 were no statistically significant differences between the molecular weight of the bands in both electrophoretic
117 conditions for G1 and G2. The appearance of 47kDa band in G3 group was the only statistically significant
118 difference in MW respect to G1 and G2 groups.

119 Regarding to intensity values, statistically significant differences were found between both G1 and G2 groups
120 for the bands at 12 and 11 kDa ($p < 0.001$); for G3 group in the bands at 52 ($p < 0.05$) and 12 kDa ($p < 0.001$)
121 concerning to G1 and G2 groups. All these results were based in electrophoretic analysis under non-reduced
122 conditions. In reduced conditions, the Year 2020

123 12 Global

124 13 Table 1: Comparison of MW and intensity of protein band 125 in female (G1), male (G2) and female + male (G3) of *R.* 126 *juncus* scorpion venom

127 Legend: Data were analyzed using two-way ANOVA followed by Bonferroni test: a?: $p < 0.05$ (G1), a???: $p < 0.001$
128 (G1); b?: $p < 0.05$ (G2), b???: $p < 0.001$ (G2) and c?: $p < 0.05$ (G3), c???: $p < 0.001$ (G3).

129 Viability was significantly reduced on the HeLa tumor cell line treated respect to untreated control after 72h
130 for 0.25 mg/mL ($p < 0.05$), 0.5 mg/mL ($p < 0.001$) and 1 mg/mL ($p < 0.001$) for G2 group. The same results were
131 observed for G1 and G3 at 0.5 mg/mL ($p < 0.01$) and 1 mg/mL ($p < 0.001$) (Figure 3A). The effect on the Vero
132 cell line resulted in no significant difference in cell viability between treated and untreated cells for all groups
133 and concentrations studied (Figure 3B). However, no statistically significant differences were observed for the
134 percentages of cells viability among all groups and concentrations of *R. juncus* scorpion venom evaluated in
135 HeLa and Vero cell lines. The IC 50 values found for HeLa cells were not significantly different among G1
136 (1.13mg/mL), G2 (1.034 mg/mL) and G3 (1.175 mg/mL) groups. In Vero cell line, no cytotoxic effect was
137 observed in all concentration tested and the theoretical IC 50 values were higher than 1 mg/mL: 3.1, 3.5 and 2.6
138 mg/mL for G1, G2 and G3, respectively (Table ??).

139 14 Table 2:

140 The IC 50 values on the HeLa and the Vero cell lines exposed to G1, G2 and G3 of *R. juncus* scorpion venom.
141 Values represent the mean \pm SD derived from three independent experiments. Data showed as >1 means No
142 effect.

143 The morphological changes induced at 1 mg/mL of G1, G2 and G3 of *R. juncus* scorpion venom are shown
144 in the Figure 4. All of them induced a loss of membrane integrity on HeLa cells; meanwhile Vero cells were not
145 affected by scorpion venom from all studied group. Studies about venom recovery in *Androctonus ferox* and
146 *Hottentota tumulus* scorpions kept in the laboratory, demonstrated better yield and quantity by electrical method
147 than manual method. Also, it was revealed the influence of diet and temperature on venom production (Tobassum
148 et al., 2018). However, there are others parameters affecting venom consistency such as sex, geographical location,
149 age and time intervals for extraction (Pucca et al., 2014). Regarding this last parameters, it has been reported
150 that extended periods of Cuban *R. juncus* venom collection was positively correlated with the regeneration of
151 venom composition and the increase of cytotoxic effect against A549 lung cancer cells (Díaz-García et al., 2019).
152 On the other hand, 200 individual molecular masses were identified in male and female *R. juncus* scorpion venom
153 from which 63 are identical in both sexes (Rodríguez-Ravelo et al., 2015).

15 CONCLUSIONS

154 The present study, demonstrated that total protein concentration was similar among female, male and the
155 mixture of both sexes of *R. junceus* scorpion venom. Moreover, the numbers of bands in all sex studied groups
156 were identical from both electrophoretic conditions, with no statistically significant differences in the molecular
157 weight in female and male scorpions. Nevertheless, unique bands were observed in each one of the sexes. with
158 poisonous animals how *Cerastes cerastes* snake, where specific bands were found in male (42 and 39 kDa) and
159 female (46 and 44kDa) venoms ??Sarhan et al, 2017). However, in the study with *Cerastes cerastes* snake
160 only was shown the MW in the electrophoretic analysis. While we also determined, the intensity of each one
161 of the bands obtained under both electrophoretic conditions. Statistically significant differences were observed
162 between both sexes (G1 and G2) for the intensity of the bands to 12 kDa (under non-reduced conditions) and
163 11 kDa (under non-reduced and reduced conditions). Previous proteomic studies with *R. junceus* scorpion venom
164 disagree to current study where the scorpion venom was kept in captivity. That study evaluated by HPLC
165 and mass spectrometry, the venom of females and males scorpions kept in its natural medium. As results, the
166 relative abundance of identical components was different among the genders (Rodríguez-Ravelo et al., 2015).
167 We have already reported the majority band below 14 kDa in the electrophoretic profile (Díaz-García et al.,
168 2015). Several authors that work with scorpion venom have reported the presence of the mixture peptides in a
169 diffuse area conformed by small molecules lower than 14 kDa, due to the little resolution power of SDS-PAGE
170 technique, these peptides cannot be observed separated (Hernández-Betancourt et al., 2009). This is the first
171 study which compares the electrophoretic profiles of this venom from different sexes and the combination of
172 both sexes (same proportion) from scorpion maintained under conditions of captivity. Previous studies with no
173 same proportion have been done (Díaz-García et al., 2013; Díaz-García et al., 2017; Díaz-García et al., 2019).
174 One of the important contributions of this study was the demonstration that morphological changes and loss of
175 viability on the HeLa tumoral cell line; and the lack of cytotoxicity on the Vero cell line induced by *R. junceus*
176 scorpion venom was independently of gender. This biological effect could be possible because the most abundant
177 components are present in both sexes (Rodríguez-Ravelo et al., 2015). Also, the bands of the most intensity
178 in all sex studied groups corresponded to proteins with low MW that they are the main responsible for their
179 therapeutic potentialities. However, next studies are needed to compare this biological effect of venom from
180 scorpions with different sexes with other tumor cells.

181 V.

15 Conclusions

182 In our experimental conditions, there were similarities in the protein concentration and some differences in the
183 electrophoretic profile of female, male and the combination of both sexes of *Rhopalurus junceus* scorpion venom.
184 However, the cytotoxic effect of *Rhopalurus junceus* scorpion venom is maintained regardless of the sex of the
185 scorpions on the HeLa tumor cell line.
186

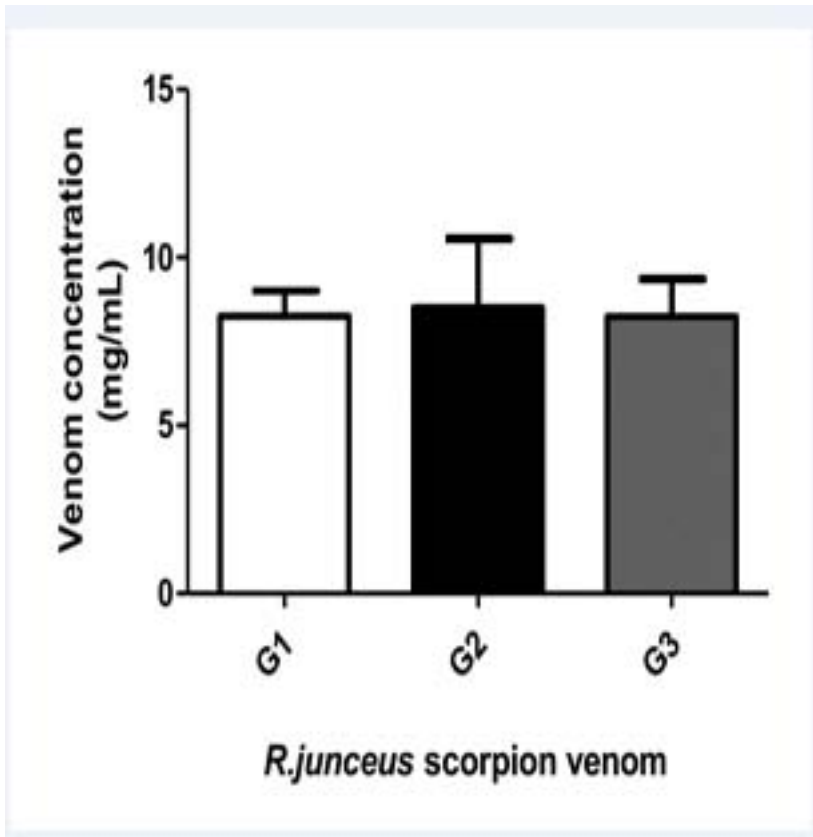
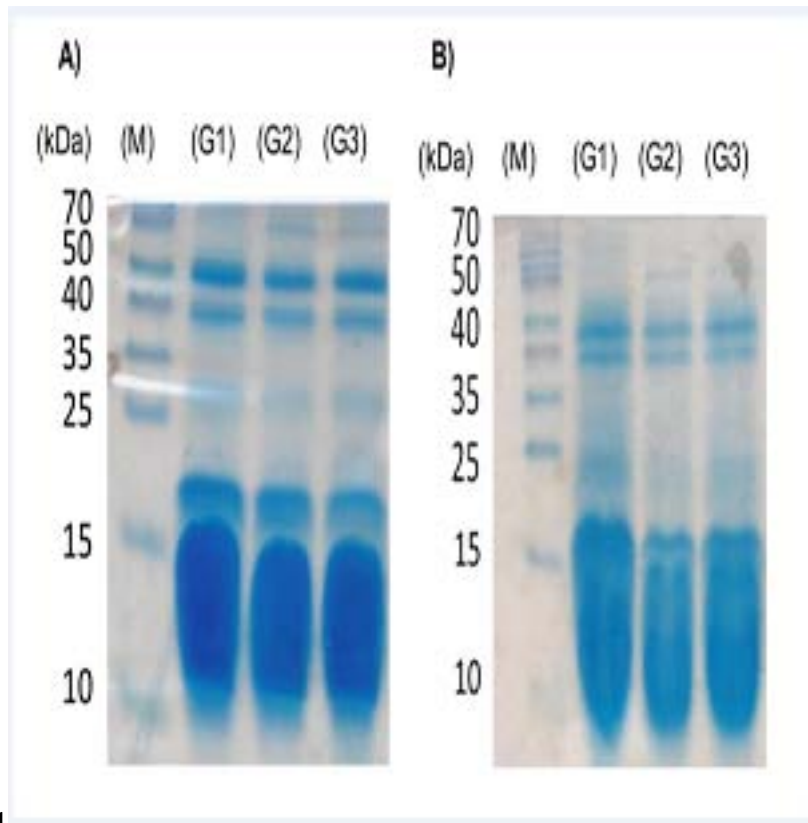


Figure 1:



1

Figure 2: Figure 1 :

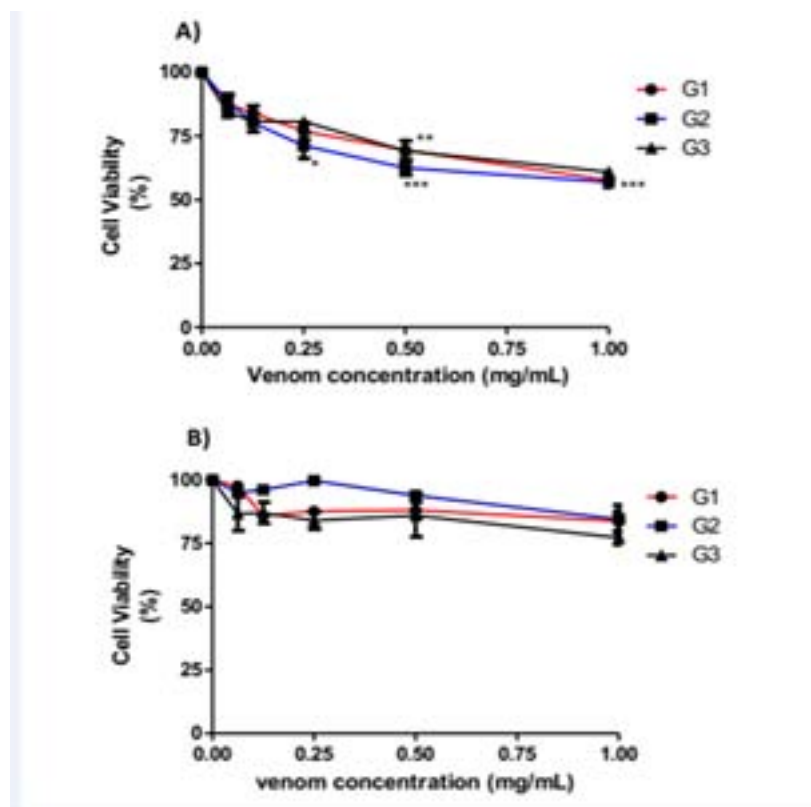
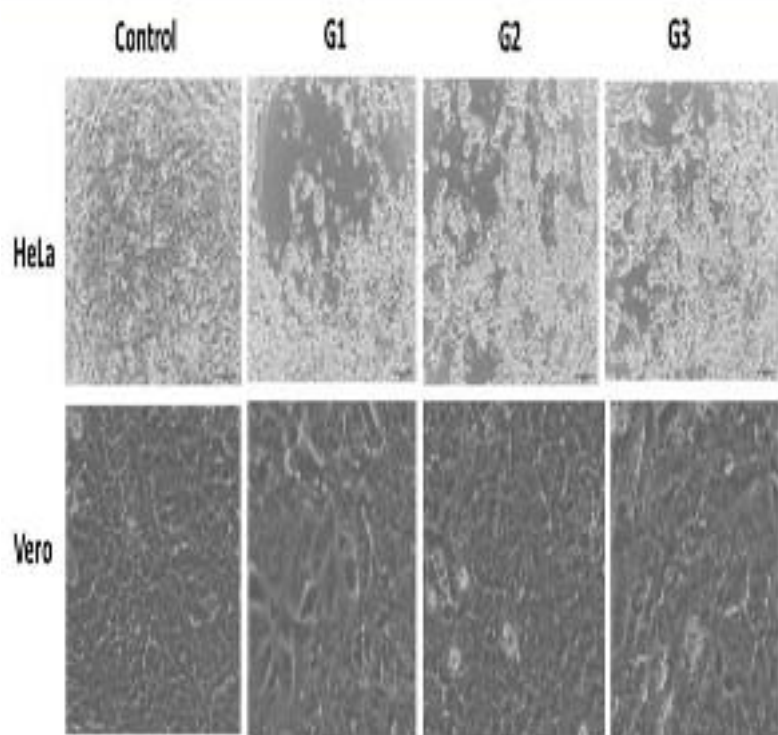


Figure 3:

SDS-PAGE	G1		G2		G3	
	MW (kDa)	Band Intensity	MW (kDa)	Band Intensity	MW (kDa)	Band Intensity
non-reduced conditions	55	585	55	1105	52	1762 ^{a,b}
	45	1813	45	2060	45	1642
	39	1090	39	1201	39	991
	28	759	28	1178	28	695
	19	2055	19	2714	19	2883
	12	21540 ^{b,c}	11	26950 ^{a,c}	12	24290 ^{a,b}
reduced conditions	66	656	65	729	64	1011
	50	922	49	1295	52	816
	44	412	44	363	47 ^{a,b}	448
	23	1374	25	1465	42	674
	17	2858	18	923	24	1266
	11	10926 ^{b,c}	11	15010 ^{a,c}	19	957
					11	7479 ^{a,b}

3

Figure 4: Figure 3 :



4

Figure 5: Figure 4 :

Samples of <i>R.juncus</i> scorpion venom	IC ₅₀ (mg/mL)	
	HeLa	Vero
G1	1.13 ± 0.004	>1
G2	1.034 ± 0.262	>1
G3	1.175 ± 0.128	>1

Figure 6:

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190 scorpion bioterium facilities LABIOFAM-Isla de la Juventud.

191 .2 Conflict of Interest

192 The authors declare no conflict of interest.

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195 commercial, or not-for-profit sectors.

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