

# Functional response of *Cydnodromus picanus* (Acari: Phytoseiidae) on two-spotted spider mite, *Tetranychus urticae* (Acari: Tetranychidae)

## Respuesta funcional de *Cydnodromus picanus* (Acari: Phytoseiidae) sobre la araña bimaculada, *Tetranychus urticae* (Acari: Tetranychidae)

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### Abstract

The functional response of adult females of predatory mite *Cydnodromus picanus* Ragusa (Acari: Phytoseiidae) was evaluated at different egg densities (5, 10, 20, 30, 40, 50, 60, 70, 80, 100 and 150 eggs per predator) of *Tetranychus urticae* Koch (Acari: Tetranychidae), which had 16, 42 and 65 h since oviposition. The experimental design was in a completely randomized blocks with five replicates per treatment. The environmental conditions of the trials were 25±2°C temperature, 50±2% of relative humidity and a photoperiod of 16:8 hours (light: dark). The average consumption rates for the three kinds of prey showed no significant differences ( $p>0.05$ ) being 20.56±1.02, 18.59±0.79 and 18.38±0.94 prey/predator for eggs of 16, 42 and 65 h age, respectively. Using a logistic regression, a type II functional response on *C. picanus* females for the three kinds of eggs was determined. The values of response parameters for *C. picanus* females were as follows: Attack rate ( $a$ ): 0.055±0.006, 0.076±0.009 and 0.073±0.016; Handling time ( $T_h$ ): 0.684±0.036, 0.894±0.034 and 0.898±0.062; for eggs of 16, 42 and 65 h age, respectively. These values are within the range of variation for different species of phytoseiids. These results suggest that *C. picanus* could effectively regulate populations of *T. urticae* in the field.

**Key words:** Biological control, predation, Holling's disc equation, attack rate, handling time.

### Resumen

La respuesta funcional de hembras adultas del ácaro depredador *Cydnodromus picanus* Ragusa (Acari: Phytoseiidae) se evaluó a diferentes densidades de huevos de *Tetranychus urticae* Koch (Acari: Tetranychidae) que tenían 16, 42 y 65 horas de edad. Las densidades evaluadas fueron: 5, 10, 20, 30, 40, 50, 60, 70, 80, 100 y 150 huevos por depredador. Las condiciones ambientales de los ensayos fueron 25±2 °C de temperatura, 50±2% de humedad relativa y un fotoperiodo de 16:8 horas (luz: oscuridad). Las tasas de consumo promedio para los tres tipos de presas no presentaron diferencias significativas ( $p>0,05$ ) siendo de 20,56±1,02; 18,59±0,79 y 18,38±0,94 presas/depredador para los huevos de 16, 42 y 64 h de edad, respectivamente. Utilizando una regresión logística, se determinó una respuesta funcional de tipo II para las hembras de *C. picanus* para los tres tipos de presas. Los valores de los parámetros de la respuesta funcional para las hembras de *C. picanus* fueron: Tasa de ataque ( $a$ ): 0.055±0.006; 0.076±0.009 y 0.073±0.016; Tiempo de manipulación ( $T_h$ ): 0.684±0.036; 0.894±0.034 y 0.898±0.062, para los huevos de 26, 42 y 65 h d edad, respectivamente. Estos valores están dentro del rango de variación para diferentes especies de fitoseidos. Estos resultados sugieren que *C. picanus* podría efectivamente regular poblaciones de *T. urticae* a nivel de campo.

**Palabras clave:** Control biológico, depredación, Ecuación del disco de Holling, tasa de ataque, tiempo de manipulación.

## Introduction

*Tetranychus urticae* Koch (Acari: Tetranychidae) is a cosmopolitan and polyphagous pest that attacks many economic importance crops in Chile, such as vegetables, extensive cultures (cotton, corn, etc.), citrus, vine and other fruit and ornamental trees. Although different strategies for control of *T. urticae* are recommended worldwide, the use of natural enemies has been highlighted as a tool of first order for biological control of this pest, especially the use of predator mite *Phytoseiulus persimilis* Athias-Henriot (Acari: Phytoseiidae) (De Moraes, 2004). However, this practice has been less common than the use of chemical acaricides, which improperly used, can cause resurgence, resistance and other problems of pests due to the elimination of natural enemies (Vitelli & Sato, 2016).

*Cydnodromus picanus*, was described in 2000 on *Citrus aurantium* L. (Rutaceae) and reported only for Chile (Ragusa, 2000; De Moraes *et al.*, 2004). Recent researches suggest *C. picanus*, could be considered as junior synonymy of *Cydnodromus idaeus* (Denmark and Muma) (Tixier *et al.*, 2011). According to Ragusa *et al.* (2000), this species could be used in arid and warm agroecosystems, where is difficult to find phytoseiids that are adapted to these extreme conditions to control tetranychids. Based on classification of McMurtry & Croft (1997), *C. picanus* is a generalist predator belonging to type III lifestyle. In conditions of absence of prey can feed of pollen from different species (Ragusa *et al.* 2000), predated on *Oligonychus yothersi* (McGregor) in *Persea americana* Mill. (Lauraceae) (Rioja & Vargas, 2009) and on *Tetranychus cinnabarinus* Boisduval in *Dianthus caryophyllus* Linn. (Caryophyllaceae) (Tello *et al.*, 2009). The citrus are one of the more important crops in the zone of Pica, Chile (20° 29' 12.4" S; 69° 19' 33.9" O), where *C. picanus* had been found associated to *Panonychus citri* (McGregor) (Ragusa *et al.*, 2000).

The present research provides information about the predatory efficiency of *C. picanus* throughout functional response. The equations of the functional response are a key component of any mechanistic model, because they describe the interactions among consumers and its resources (Villemereuil & López-Sepulcre, 2011). This information is vital for the implementation of conservation programs for natural enemies and augmentation-type biological control (Ragusa *et al.*, 2010). The effectivity of a predator is directly related to the type of its functional response. Predatory arthropods unfold one of the three typical functional responses, but these responses may vary with the crop phenology, the habitat heterogeneity, the predator age and other biotic and abiotic factors. In type I functional

response, there is a lineal increase of predator attack rate respect to prey density (with a slope equal to the search efficiency), until reaching a point at which the maximum attack rate remains constant, even if the prey density increases. The response is estimated by a linear equation  $N_a = \alpha + \beta N_o$ ; where  $N_a$ =number of prey eaten,  $N_o$ =prey density (number of prey offered),  $\alpha$  and  $\beta$  are the y-intercept and the slope of prediction line, respectively. In type II functional response appears another parameter, in addition to the search efficiency, termed "handling time" ( $T_h$ ) defined as the time that invests the predator in chasing, capturing, consuming and digesting a prey. In type II functional response, the number of prey consumed hyperbolically approaches to an asymptote in which the maximum attack rate is expressed ( $K=T/T_h$ ). This type of response is estimated by a curved line function:  $N_a = aN_oT / (1 + aN_oT_h)$ , where  $N_a$ =number of prey consumed,  $a$ =constant attack rate or instantaneous search rate,  $N_o$ =initial prey density,  $T$ =total available time and  $T_h$ =handling time. The type III functional response is performed when the prey-consumed number approaches an asymptote as a sigmoid function (increases in the proportion of prey killed to an inflection point and then decreases similarly to type II response). The type III functional response may be derived using the model of Hassell (Hassell, 1978), where the attack rate ( $a$ ) is a hyperbolic function of prey density:  $a = (d + hN_o) / (1 + cN_o)$ , substituting the value of the Holling equation, obtaining the type III model:  $N_a = (dN_oT + hN_o^2T) / (1 + cN_o + dN_oT_h + bN_o^2T_h)$ , where  $N_a$ =number of prey consumed,  $N_o$ =initial prey density,  $T$ =total time,  $T_h$ =handling time and  $b$ ,  $c$ , and  $d$  are constants.

Tello *et al.* (2009), proved the life parameters of *C. picanus* when fed with eggs, the same happened with survival, achieving 100% when fed with this stadium. For those experiments, eggs less than 20 h old were performed. In addition, the functional response of this phytoseiid was affected by the maturity of the egg offered as food.

The aim of this research was to evaluate the consumption rates of *C. picanus* (= *C. idaeus*) in relation to eggs density of *T. urticae* and to determine the effect of egg maturity on functional response parameters, under laboratory conditions.

## Material and methods

### Establishment of colonies

The bioassays were performed in the Regional Research Center INIA La Cruz (Instituto de Investigaciones Agropecuarias (INIA), La Cruz, Chile), at a temperature of 25±2°C, 50±2% of relative humidity (RH) and a photoperiod of 16:8 h (L:D). *Cydnodromus picanus*, was collected from trees of *C. aurantium* in Pica, Chile (20°29' S; 69°19'

O) and bred in acrylic plates of 12x12 cm according to the methodology described by Swirski, Amitai & Dorzia (1970), in breeding rooms and fed only with pollen from *Oxalis* sp (Oxidales) to avoid habituation to prey. *Tetranychus urticae* obtained from breeding rooms located in INIA La Cruz. The colonies of this tetranychid had a time of about four months since they were collected in the field. *Tetranychus urticae* was reproduced on bean plants (*Phaseolus vulgaris* L.), that were infested with the mite when they had two true leaves. To maintain available plants, 100 pots weekly seeded with 20 bean seeds.

### Consumption rate and functional response

The functional response of *C. picanus* on *T. urticae* eggs of different maturity time studied in separated bioassays. For each maturity time, one phytoseiid adult female, previously subjected to 24 h fasting, was confined in an artificial arena (black acrylic plate of 4 cm diameter) (Tello *et al.*, 2009). These *C. picanus* females came from a cohort of eggs of the same age, using newly emerged females. Glue (Point Sticken Glue, Point Chile S.A.) was applied in the edges plates to prevent the mites escape. Therefore, *T. urticae* eggs of different maturity times (16, 42 and 65 h) were introduced as preys in the plates at densities of 5, 10, 20, 30, 40, 50, 60, 70, 80, 100 and 150 eggs per plate. *T. urticae* eggs were obtained by depositing on a bean leaf about 100 females to oviposit and at 16h were transferred into trial arenas, replicating this for eggs of 42 and 65 h. The number of *T. urticae* eggs consumed by the predator recorded at 24 h (experimental time). The observations were made using a stereomicroscope with 40X magnification (Zeiss Stemi 6V 6, Alemania).

### Statistical analysis

The consumption among eggs with different maturity (16, 42 and 65 h) were compared. In addition, for each age, among different densities. Because consumption data is not normally distributed, even after applying transformations (e.g. logarithmic and square root). The non-parametric Kruskal-Wallis test with post-hoc comparison were performed to determine the type of functional response, data of each density were fitted to maximum likelihood logistic regression (CATMOD procedure, SAS Institute, Cary, NC, USA). (Equation 1).

$$\frac{N_a}{N_0} = \text{Prob}(Y = 1) = \frac{\exp(P_0 + P_1 N_0 + P_2 N_0^2 + P_3 N_0^3)}{1 + \exp(P_0 + P_1 N_0 + P_2 N_0^2 + P_3 N_0^3)}$$

Equation 1

Where Y, is the dichotomous dependent variable, representing a non-consumed (Y=0) or consumed (Y=1) prey at the end of the trial,  $N_a$  is the number of prey consumed,  $N_0$  is the number of prey available; and  $P_0$ ,  $P_1$ ,  $P_2$  and  $P_3$  are the intercept, linear, quadratic and cubic coefficient, respectively. Negative or positive significant linear coefficients (e.g.,  $P_1$ ) indicate a type II or III functional response, respectively. The slope of type II response decreases (e.g.,  $P_1$  is negative), while that slope of type III increases (e.g.,  $P_1$  is positive). After determining the type of functional response, its parameters (attack rate and handling time) are estimated through linearization of Holling's disc equation depending on the type of functional response (Xiao & Fadamiro, 2010). Type II equation was used:  $a$  and  $T_h$  parameters were calculated using a nonlinear regression by least squares based on Gauss-Newton method (SAS Institute, Cary, NC, USA). The  $a/T_h$  value indicates the effectiveness of depredation. The maximum depredation rate ( $K$ ) was calculated as  $T/T_h$ . To perform comparisons between  $a$  and  $T_h$  parameters, standard deviation was estimated with 95% confidence interval using Jackknife statistical technique available in SAS (SAS Institute, Cary, NC, USA). Later, the parameters were compared using non-parametric Kruskal-Wallis test ( $\alpha=0.05$ ) due to heterogeneity of variances and non-normally distributed residuals.

## Results and discussion

### Consumption rate

The average consumption rate of *C. picanus* did not show significant differences among the three types of eggs offered [ $KW=4.04$ ,  $gl=2$ ;  $P=0.131$ ]. This indicates that the maturity time of eggs offered as prey have no influence in the preference by *C. picanus*. The predator consumed 54, 52 and 51% of initial egg density at 16, 42 and 65 h, respectively, not showing significant differences [ $KW=1.166$ ;  $gl=2$ ;  $P=0.556$ ] (Table 1).

**Table 1.** Average consumption rate ( $\pm$ SE) and proportion of prey consumed ( $\pm$ SE) for *Cydnodromus picanus* fed with *Tetranychus urticae* eggs of different maturity times

Prey	Average consumption rate	Na/No
Eggs of 16 h	20.56 $\pm$ 1.02 a	0.54 $\pm$ 0.03 a
Eggs of 42 h	18.59 $\pm$ 0.79 a	0.52 $\pm$ 0.04 a
Eggs of 65 h	18.38 $\pm$ 0.94 a	0.51 $\pm$ 0.04 a

Equal letters in the same column indicate no significant differences according to Kruskal-Wallis test ( $P<0.05$ );  $N_a$ =number of prey consumed;  $N_0$ =initial prey density; SE=standard error.

From a density of 50 eggs.predator<sup>-1</sup>, the age of eggs did not affect the predation by *C. picanus*; 5 ( $KW=0.32$ ,  $gl=2$ ,  $P=0,3679$ ), 10 ( $KW=2.33$ ,  $gl=2$ ,

$P < 0.1503$ ), 20 ( $KW = 0.43$ ,  $gl = 2$ ,  $P = 0.7884$ ), 30 ( $KW = 4.76$ ,  $gl = 2$ ,  $P < 0.0913$ ), 40 ( $KW = 0.04$ ,  $gl = 2$ ,  $P = 0.9807$ ) and 50 ( $KW = 1.35$ ,  $gl = 2$ ,  $P = 0.5053$ ). The Kruskal-Wallis test did not show significant differences ( $P > 0.05$ ) for those prey densities (Table 2). Differences showed densities of 60 ( $KW = 12.49$ ,  $gl = 2$ ,  $P < 0.05$ ), 70 ( $KW = 8.75$ ,  $gl = 2$ ,  $P < 0.05$ ) and 100 ( $KW = 8.54$ ,  $gl = 2$ ;  $P < 0.05$ ) eggs.predator<sup>-1</sup>. The number of attacked eggs per capita was significantly affected by prey density for each egg maturity (16h:  $KW = 54.87$ ,  $gl = 10$ ,  $P < 0.0001$ ), 42h:  $KW = 43.23$ ,  $gl = 10$ ,  $P < 0.0001$  and 65h:  $KW = 46.31$ ,  $gl = 10$ ,  $P < 0.0001$ ).

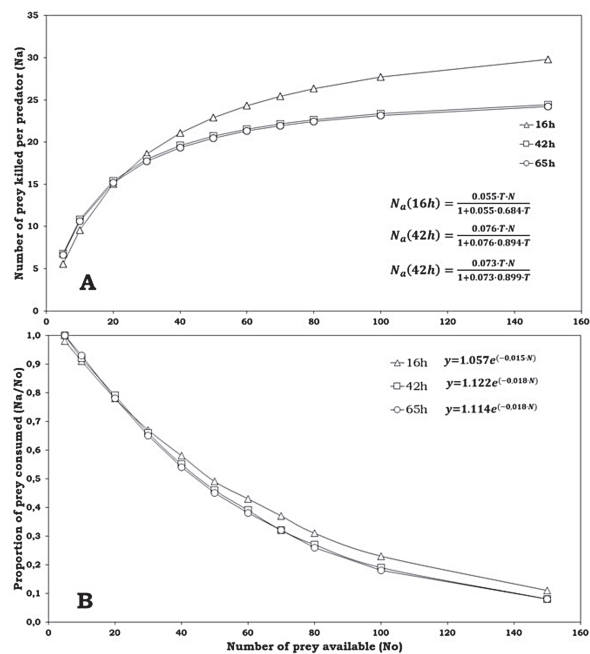
**Table 2.** Prey consumption (mean ± SE) for *Cydnodromus picanus* when fed with *Tetranychus urticae* eggs of three maturity times at different prey densities

Density	16 h	42 h	65 h
5	5.00 ± 0.00 A d	4.83 ± 0.17 A e	5.00 ± 0.00 A e
10	8.83 ± 0.79 A d	9.67 ± 0.21 A de	10.00 ± 0.00 A de
20	19.00 ± 0.52 A bcd	18.50 ± 0.56 A cd	15.00 ± 2.91 A cde
30	16.33 ± 0.84 A d	18.83 ± 1.17 A bcd	20.67 ± 1.82 A abc
40	18.33 ± 0.92 A cd	18.67 ± 1.84 Abcd	18.17 ± 0.65 A bcd
50	23.67 ± 1.96 A abc	19.50 ± 1.73 A abc	21.00 ± 2.29 A abc
60	27.83 ± 1.56 A a	22.83 ± 1.42 A abc	16.00 ± 0.93 B cde
70	25.67 ± 0.84 AB ab	23.33 ± 0.99 B ab	32.17 ± 2.09 A a
80	23.50 ± 1.18 A abc	21.67 ± 0.56 A abc	20.17 ± 1.05 A bc
100	27.00 ± 1.41 A a	21.50 ± 0.89 B abc	20.83 ± 1.08 B bc
150	31.00 ± 2.37 A a	25.17 ± 1.35 A a	23.17 ± 1.30 A ab

Means in the same row followed by different uppercase letters are significantly different; means in the same column followed by different lowercase letters are significantly different ( $P < 0.05$ ; Kruskal-Wallis Test). SE=standard error.

### Functional Response

The logistic regression delivered a significantly negative linear parameter ( $P_1 < 0$ ) for the three classes of prey offered (eggs of 16, 42 and 65 h), suggesting that *C. picanus* performs a type II functional response for the three types of prey (Table 3). The functional response curves (Figure 1A), showed a high proportion of prey consumed at lower densities in the three kinds of prey. The proportion of eggs consumed by predator ( $N_a/N_0$ ) declined with the increase of prey density (Figure 1B); this implies that there is an inverse density-dependence, being confirmed by logistic regression (Table 3) that, for all ages of eggs, presented a negative linear coefficient ( $P_1 < 0$ ) and a positive quadratic coefficient ( $P_2 > 0$ ), both significant ( $P < 0.001$ ; Table 3).



**Figure 1.** Consumption of *Tetranychus urticae* eggs (A) and proportion of prey consumed (B) per *Cydnodromus picanus* females according to provided density (fitted equations inside plots)

**Table 3.** Selection of functional response model based on maximum likelihood analysis of parameter estimations (±SE) using the proportion of eggs consumed as a polynomial function of offered eggs density

Egg maturity	Linear ( $P_1$ )	Quadratic ( $P_2$ )	Cubic ( $P_3$ )	Best fitted model
16 h	-0.1082±0.0152 <sup>a</sup>	0.00098±0.000201 <sup>a</sup>	-3.07E±7.71E <sup>-7a</sup>	Tipo II
42 h	-0.1409±0.0169 <sup>a</sup>	0.00132±0.000220 <sup>a</sup>	-4.21E±8.41E <sup>-7a</sup>	Tipo II
65 h	-0.1149±0.0156 <sup>a</sup>	0.00103±0.000206 <sup>a</sup>	-3.26E±7.95E <sup>-7a</sup>	Tipo II

<sup>a</sup> Significant at  $P < 0.001$ , SE=standard error.

The comparison of functional response curves revealed significant differences among the three kinds of eggs offered ( $\alpha$ :  $KW = 11.94$ ,  $gl = 2$ ,  $P < 0.05$ ;  $T_h$ :  $KW = 11.51$ ,  $gl = 2$ ,  $P < 0.05$ ). The maximum number of prey consumed was significantly higher for eggs of maturity of 16 h (Table 4). The estimated parameters of functional response for *C. picanus*, for the three types of prey, showed that the attack rate was higher for eggs of 42 h and 65 h than for eggs of 16 h. On the other hand, *C. picanus* showed the lowest handling time ( $T_h$ ) for eggs of 16 h (Table 4). The slightly higher  $\alpha/T_h$  values recorded for *C. picanus* on eggs of 42 and 65 h suggest that this predator was more effective on consuming that sort of *T. urticae* eggs.



**Table 4.** Values of  $R^2$  and parameters (mean  $\pm$  SE) of functional response of *Cydnodromus picanus* adult females on *Tetranychus urticae* eggs of different maturity times estimated by disc equation.

Eggs Maturity	Disc Equation			$a/T_h$	K
	$a$	$T_h$	$R^2$		
16 h	0.055 $\pm$ 0.006 b	0.684 $\pm$ 0.036 b	0.97	0.080 $\pm$ 0.0 b	35.12 $\pm$ 0.37 a
42 h	0.076 $\pm$ 0.009 a	0.894 $\pm$ 0.034 a	0.98	0.085 $\pm$ 0.0 a	26.85 $\pm$ 0.19 b
65 h	0.073 $\pm$ 0.016 a	0.898 $\pm$ 0.062 a	0.93	0.081 $\pm$ 0.0 ab	26.70 $\pm$ 0.16 b

Values with equal letters in the same column are significantly not different according to Kruskal-Wallis test ( $P < 0.05$ );  $a$ =search or attack rate;  $T_h$ =handling time (day);  $R^2$ =determination coefficient;  $a/T_h$ =depredation efficiency; K=maximum depredation rate.

The average eggs consumption rate is similar to those obtained for adult females of this phytoseiid species. Ragusa *et al.* (2000) indicates a daily consumption of about 16 eggs of *T. urticae* at 26 $\pm$ 1°C

and 70 $\pm$ 5% HR. Tello *et al.* (2009) registered the same consumption value (16 eggs.day<sup>-1</sup>) with *T. cinabarinus* eggs at 29.44 $\pm$ 1.47°C and 42.35 $\pm$ 5.01% RH.

*Cydnodromus picanus*, showed a type II functional response for each maturity time of eggs. This type of functional response is registered for different phytoseiid species, including: *Amblyseius californicus* (Mc-Gregor) (Gotoh, Nozawa & Yamaguchi, 2004); *Neoseiulus longispinosus* (Evans) (Rahman, Babu, Roobakkumar & Perumalsamy, 2012); *Phytoseiulus persimilis* Athias-Henriot, *Galendromus occidentalis* (Nesbitt) (Xiao & Fadamiro, 2010); *Neoseiulus cucumeris* (Oudemans) (Madadi *et al.*, 2007). In general, the results obtained in this study for  $a$  and  $T_h$  parameters of functional response for *C. picanus* are within the range of variation for different phytoseiid species (Table 5).

**Table 5.** Functional response parameters ( $a$  and  $T_h$ ) for different phytoseiid species depredating on *Tetranychus urticae* eggs in a range between 15°C and 35°C

Phytoseiid	Prey	Temp (°C)	Attack rate ( $a$ )	Handling time ( $T_h$ )	References
<i>Neoseiulus californicus</i>	<i>Tetranychus urticae</i>	25	1,676	0,041	Kuştutan and Çakmak (2009)
<i>Neoseiulus californicus</i>	<i>Tetranychus urticae</i>	27	0,935	0,012	Landeros <i>et al.</i> (2013)
<i>Neoseiulus californicus</i>	<i>Tetranychus urticae</i>	26	0,490	1,733	Xiao, Osborne, Chen and McKenzie (2013)
<i>Amblyseius swirskii</i>	<i>Tetranychus urticae</i>	26	0,421	0,518	Xiao <i>et al.</i> (2013)
<i>Neoseiulus californicus</i>	<i>Tetranychus urticae</i>	35	0,209	0,951	Ahn, Kim and Lee (2010)
<i>Typhlodromus athiasae</i>	<i>Tetranychus urticae</i>	25	0,115	0,130	Saker, Dahiah, Mofleh and Basheer (2015)
<i>Phytoseiulus persimilis</i>	<i>Tetranychus urticae</i>	25	0,114	3,514	Seiedy, Saboori, Allhyari, Talaei-Hassanloui and Tork (2012)
<i>Neoseiulus californicus</i>	<i>Tetranychus urticae</i>	25	0,094	1,645	Farazmand, Fathipou and Kamali (2012)
<i>Cydnodromus picanus</i>	<i>Tetranychus urticae</i> (huevos 42h)	25	0,076	0,894	Present study
<i>Cydnodromus picanus</i>	<i>Tetranychus urticae</i> (huevos 65h)	25	0,073	0,899	Present study
<i>Neoseiulus californicus</i>	<i>Tetranychus urticae</i>	30	0,071	1,331	Ahn <i>et al.</i> (2010)
<i>Neoseiulus californicus</i>	<i>Tetranychus urticae</i>	25	0,064	1,655	Ahn <i>et al.</i> (2010)
<i>Neoseiulus californicus</i>	<i>Tetranychus urticae</i>	20	0,061	1,998	Ahn <i>et al.</i> (2010)
<i>Cydnodromus picanus</i>	<i>Tetranychus urticae</i> (huevos 16h)	25	0,055	0,684	Present study
<i>Neoseiulus californicus</i>	<i>Tetranychus urticae</i>	15	0,045	4,459	Ahn <i>et al.</i> (2010)
<i>Iphiseius degenerans</i>	<i>Tetranychus urticae</i>	25	0,017	3,441	Fantinou <i>et al.</i> (2012)

The search (or attack) rate and the handling time are parameters used to determine the magnitude of the functional responses. This parameter could be affected by different factors, such as the predator velocity, the prey movements and the spent time in subduing each prey (Hassell, 1978). In this research, the handling time for eggs of 16 h was significantly lower than for eggs of 42 and 65 h, which could be explained for the presence of a higher amount of vitellus and the easiness to extract it. When eggs mature, the embryonic

development makes diminish the amount of vitellus, making difficult its extraction and making to take longer time to predator in feeding. Cabrera, Donohuea, Khalil, Sonenshinec & Roa (2009), pointed that there is no available studies describing the characteristics of proteins in the yolk of mite eggs. According to these authors, in newly oviposited tetranychids eggs (<3 h), the embryo development would be minimal and the proteins would be more available for predators. Therefore, we can deduce that as time progress

and egg maturity with it; the eggs of 42 and 65 h present higher embryo development, so the amount of vitellus must be minimal and thus of lower nutritional value.

## Conclusion

This research compared the average consumption rates and the functional response of predator mite *C. picanus* fed with *T. urticae* eggs of different maturity times. The results show that adult females of *C. picanus* exhibit a type II functional response and a proportion of prey consumed that ranged between 51% and 54% for the three types of prey offered. Studies of predation in the field, with different crops and preys (tetranychid species), are necessary to consider this predator as a potential biological control agent in arid zones of Chile.

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## References

- Ahn, J.J., Kim, K.W. & Lee, J.H. (2010). Functional response of *Neoseiulus californicus* (Acari: Phytoseiidae) to *Tetranychus urticae* (Acari: Tetranychidae) on strawberry leaves. *J Appl Entomol*, 134(2), 98–104. <http://dx.doi.org/10.1111/j.1439-0418.2009.01440.x>
- Cabrera, A.R., Donohuea, K.V., Khalil, S.M.S., Sonenshine, D.E., & Roa, R.M. (2009). Characterization of vitellin protein in the two-spotted spider mite, *Tetranychus urticae* (Acari: Tetranychidae). *J Insect Physiol*, 55(7), 655–661. <http://dx.doi.org/10.1016/j.jinsphys.2009.04.006>
- De Moraes, G.J., McMurtry, J.A., Denmark, H.A. & Campos, C.B. (2004). A revised catalog of the mite family Phytoseiidae. *Zootaxa*, 1(434), 1–494. <http://dx.doi.org/10.11646/zootaxa.434.1.1>
- Fantinou, A.A., Baxevani, A., Drizou, F., Labropoulos, P., Perdakis, D.,... (2012). Consumption rate, functional response and preference of the predaceous mite *Iphiseius degenerans* to *Tetranychus urticae* and *Eutetranychus orientalis*. *Exp Appl Acarol*, 58(2), 133–144. <http://dx.doi.org/10.1007/s10493-012-9557-6>
- Farazmand, A., Fathipour, Y. & Kamali, K. (2012). Functional response and mutual interference of *Neoseiulus californicus* and *Typhlodromus bagdasarjani* (Acari: Phytoseiidae) on *Tetranychus urticae* (Acari: Tetranychidae). *Int J Acarol*, 38(5), 369–376. <http://dx.doi.org/10.1080/01647954.2012.655310>
- Gotoh, T., Nozawa, M. & Yamaguchi, K. (2004). Prey consumption and functional response of three acarophagous species to eggs of the two-spotted spider mite in the laboratory. *Appl Entomol Zool*, 39(1), 97–104. <http://dx.doi.org/10.1303/aez.2004.97>
- Hassell, M.P. (1978). The dynamics of arthropod predator-prey systems. *Monogr Popul Biol*, 13(3-7), 1-237.
- Kuřtutan, O. & Çakmak, I. (2009). Development, fecundity, and prey consumption of *Neoseiulus californicus* (McGregor) fed *Tetranychus cinnabarinus* Boisduval. *Turk J Agric For*, 33, 19–28. <http://dx.doi.org/10.3906/tar-0806-39>
- Landeros, J., Cerda, P., Badii, M.H., Aguirre, L.A., Cerna, E.,... (2013). Functional response of *Neoseiulus californicus* on *Tetranychus urticae* on apple leaves. *Southwest Entomol*, 38(1), 79–84. <http://dx.doi.org/10.3958/059.038.0108>
- Madadi, H., Enkegaard, A., Brodsgaard, H. F., Kharrazi-Pakdel, A., Mohaghegh, J.,... (2007). Host plant effects on the functional response of *Neoseiulus cucumeris* to onion thrips larvae. *J Appl Entomol*, 131(9–10), 728–733. <http://dx.doi.org/10.1111/j.1439-0418.2007.01206.x>
- Ragusa, S., Vargas, R., Tsolakis, H. & Ashbach, R. (2000). Laboratory studies on the influence of various food substances on some biological and life-table parameters of *Cydnodromus picanus* Ragusa (Parasitiformes, Phytoseiidae) associated with citrus trees in the Chilean desert. *Phytophaga*, 10, 1–23.
- Rahman, V., Babu, A., Roobakkumar, A. & Perumalsamy, K. (2012). Functional and numerical responses of the predatory mite, *Neoseiulus longispinosus*, to the red spider mite, *Oligonychus coffeae*, infesting tea. *J Insect Sci*, 12, 125. <http://dx.doi.org/10.1673/031.012.12501>
- Rioja, T.S. & Vargas, R.M. (2009). Life table parameters and consumption rate of *Cydnodromus picanus* Ragusa, *Amblyseius graminis* Chant, and *Galendromus occidentalis* (Nesbitt) on avocado red mite *Oligonychus yothersi* (McGregor) (Acari: Phytoseiidae, Tetranychidae). *Chilean J Agric Res*, 69(2), 160–170. <http://dx.doi.org/10.4067/S0718-58392009000200005>
- Saker, I., Dahiah, H., Mofleh, M. & Basheer, A. (2015). Functional response of the predatory mite, *Typhlodromus athiasae* Porath and Swirski (Acari: Phytoseiidae) to the two spotted spider mite, *Tetranychus urticae* Koch (Acari: Tetranychidae) infesting bean. *Egypt J Biol Pest Co*, 25(1), 1–5.
- Seiedy, M., Saboori, A., Allahyari, H., Talaei-Hassanloui, R. & Tork M. (2012). Functional response of *Phytoseiulus persimilis* (Acari: Phytoseiidae) on untreated and *Beauveria bassiana*—treated adults of *Tetranychus urticae* (Acari: Tetranychidae). *J Insect Behav*, 25(6), 543–553. <http://dx.doi.org/10.1007/s10905-012-9322-z>
- Tello, V., Vargas, R., Araya, J., & Cardemil, A. (2009). Biological parameters of *Cydnodromus picanus* and *Phytoseiulus persimilis* raised on the carmine spider mite, *Tetranychus cinnabarinus* (Acari: Phytoseiidae, Tetranychidae). *Cienc Inv Agr*, 36(2), 277–290. <http://dx.doi.org/10.4067/S0718-16202009000200012>
- Tixier, M-S., Tsolakis, H., Ragusa, S., Poinso, A., Ferrero, M., Okassa, M.,... (2011). Integrative taxonomy demonstrates the unexpected synonymy between two predatory mite species: *Cydnodromus idaeus* and *C. picanus* (Acari: Phytoseiidae). *Invertebr Syst*, 25, 273–281. <http://dx.doi.org/10.1071/IS11025>
- Villemerueil, P.B. & López-Sepulcre, A. (2011). Consumer functional responses under intra- and in-

- ter-specific interference competition. *Ecol Model*, 222(3), 419-426. <http://dx.doi.org/10.1016/j.ecolmodel.2010.10.011>
- Vitelli, M. & Sato, M. (2016). Pyrethroid resistance in *Phytoseiulus macropilis* (Acari: Phytoseiidae): cross-resistance, stability and effect of synergists. *Exp Appl Acarol*, 68, 71-82. <http://dx.doi.org/10.1007/s10493-015-9984-2>
- Xiao, Y. & Fadamiro, H. (2010). Functional responses and prey-stage preferences of three species of predacious mites (Acari: Phytoseiidae) on citrus red mite, *Panonychus citri* (Acari: Tetranychidae). *Biol Control*, 53(3), 345-352. <http://dx.doi.org/10.1016/j.biocontrol.2010.03.001>
- Xiao, Y., Osborne, L.S., Chen, J., & McKenzie, C.L. (2013). Functional responses and prey-stage preferences of a predatory gall midge and two predacious mites with two-spotted spider mites, *Tetranychus Urticae*, as host. *J Insect Sci*, 13(8), 1-12. <http://dx.doi.org/10.1673/031.013.0801>