

Project Title: Rearing and evaluating *Rhinusa linariae* and *Mecinus janthinus*, biological controls of yellow toadflax.

PI:

Andrew Norton

Professor

Colorado State University 1177 Campus Delivery, Fort Collins, CO 80523-1177. Voice: (970) 491-7421. FAX: (970) 491-3865. andrew.norton@colostate.edu

Contact:

Carol Bell Randall

Entomologist

USDA Forest Service

Northern and Intermountain Regions 2502 E Sherman Ave Coeur d'Alene, ID 83814 Phone: 208-769-3051 Fax: 208-769-3062 Email: crandall@fs.fed.us

Amount Requested (yearly and total), Project Leveraging :

Source	Year 1	Year 2	Year 3	Total
USFS-FHP-FHTET	\$24,800	\$24,250	\$25,672	\$74,722
CSU Match	\$21,297	\$26,732	\$28,077	\$76,106

Project Goals and Supporting Objectives

Linaria vulgaris, or yellow toadflax is an invasive weed, now found in 49 US States and 9 Canadian provinces. The species is on the noxious weed lists for at least 8 Western states. Like its congener Dalmatian toadflax, yellow toadflax was originally introduced as an ornamental plant. It now invades range, forest and park areas where it displaces native vegetation. The plant's chemical defenses (iridoid glycosides and alkaloids) are distasteful and toxic to livestock and other animals (Polunin 1969).

Toadflax species (*Linaria dalmatica* and *L. vulgaris*) have been the targets of classical biological control programs since the 1960's. Classical biological control has successfully been implemented to control Dalmatian toadflax in many areas. To date, several insect species have been introduced for toadflax control, including two seed-feeding beetles (which were also introduced adventively), the defoliating moth *Calophasia lunula* and the stem-boring weevil *Mecinus janthiniformis*. *Mecinus janthiniformis* has become the most effective weed biological control agent for Dalmatian toadflax in North America. Research into the genetics of these weevils (Tosevski et al. 2011) and the locations of their initial collection in Europe has now revealed that two sibling biotypes (cryptic species) were included in the original importation and release of stem borers: *Mecinus janthinus* and *M. janthiniformis*. *Mecinus janthiniformis* is the species/biotype that has proven successful for control of Dalmatian toadflax (*Linaria dalmatica*), while *M. janthinus* is the species currently found on *L. vulgaris* in Montana. The yellow toadflax weevil has subsequently been collected, reared in captivity, and redistributed at various sites in

Colorado. We have evidence of establishment, but populations have grown slowly, perhaps the result of several dry years following the releases or some other, unknown, factor. *Mecinus janthinus* populations grow more slowly than *M. janthiniformis* (pers. obs. Dan Bean, pers. communication, Sharlene Sing, pers. communication).

Rhinusa linariae (previously named *Gymnetron linariae*) (Curculionidae) is a univoltine root-galling weevil specializing on yellow toadflax. Adults begin feeding on toadflax in early May, and after mating females descend to the roots and lay eggs. Larvae develop in galls on the root system. Field studies have found up to 20 galls per plant, and in caged test studies plants with more than 100 galls were recorded (Jordan, 1994). Work at the CABI biological control center in Switzerland resulted in its permitting for release into the United States and Canada in 1994.

Initial releases of the weevil failed to establish in the US. However, 2 populations have established in British Columbia and have recently reached a density high enough to support collection and re-distribution efforts. In 2009 the Colorado Department of Agriculture biological control insectary imported 200 *R. linariae* and released them in a field cage in the White River National Forest. These weevils did not establish (Dan Bean, personal communication). In 2014 we imported root galls from the British Columbia site, overwintered these in the lab, and in spring of 2015 released the resulting adults onto yellow toadflax populations in Colorado. We do not know yet if these weevils have established.

Both *M. janthinus* and *R. linariae* have potential as biological control agents for yellow toadflax. However, unlike *M. janthiniformis* on Dalmatian toadflax, populations of these species grow more slowly. While reports from Montana for *M. janthinus* (Sharlene Sing, pers. communication) and British Columbia for *R. linariae* (Susan Turner, pers. communication) suggest that both weevils are capable of substantially reducing weed population densities, the few sources of the weevils for re-distribution has limited their implementation in yellow toadflax biocontrol. I propose to address this by creating a mass-rearing program at CSU's experimental farm, and to use these weevils for impact studies on yellow toadflax and for re-distribution efforts throughout the US. Further, we will disseminate the mass rearing techniques we develop to stakeholders and other rearing facilities such as the Nez Perce Bio-Control Center in Lapwai, Idaho and the Colorado Department of Agriculture's Biological Control insectary in Palisade, Colorado.

Project justification

Yellow toadflax is an aggressive perennial noxious weed. Both it and its congener, Dalmatian toadflax, are unpalatable to wildlife and domesticated livestock. Both species will out compete native species, and both species are difficult to control using herbicides. Further, the collateral damage from herbicide applications often presents an additional management challenge via the appearance of secondary invaders and/or the need to actively re-vegetate treated areas. While Dalmatian toadflax is limited to lower and usually drier sites, yellow toadflax prefers more mesic and higher elevation portions of the landscape. As these areas often contain a more diverse plant community than lower drier sites (Stohlgren et al. 1999), yellow toadflax may have the potential for greater impact than Dalmatian toadflax.

Approach

The funding requested will be used for field collection of *R. linariae* and *M. janthinus* and the establishment of mass rearing colonies at CSU's experimental farm. Several years ago we secured 2.5 acres of land at the farm that is now dedicated to research on invasive plants. The site is equipped with an on-demand drip irrigation system. Following establishment of these colonies, we will perform a series of experiments to determine the impact of each agent on potted yellow toadflax plants grown under a gradient of resource availability.

Field collection of *R. linariae* and *M. janthinus*

In Spring and summer, 2016 we will travel to established sites to collect *M. janthinus* and *R. linariae*. *Mecinus janthinus* exists in small populations in western Montana. In early spring we will collect the previous years toadflax stems, return these to the lab and dissect out overwintered adults. Adults will be used to initiate a rearing colony at the CSU field station.

There are no known established populations of *Rhinusa linariae* in the United States. However, there is a site in British Columbia that has collectable populations. I have contacted the cooperators that manage the site, and depending upon next year's population size we will be able to collect the weevil in early August 2016. Current permit restrictions require that the weevil be reared through a 'critical life stage' in containment. In 2011 I developed such a containment facility. This facility has been inspected and approved by USDA-APHIS and I have the relevant permits for importation, contained rearing and transfer to field release for *R. linariae*.

Galls will be returned to the CSU campus and we will collect adults as they emerge in our containment facility. These adults will be used to initiate a rearing colony at the CSU field station.

Field collection of yellow toadflax

In early spring 2016, we will collect yellow toadflax plants from 2 sites in each of Colorado, Montana, Idaho and North Dakota. From each site, we will collect 50 individual plants for a total of 400 plants. Plants will be transported to CSU, washed of all soil material and individually potted in 1-gallon pots in the greenhouse. After 6 weeks of growth inside the greenhouse, we will move them to a shade house located on the CSU campus. Plants will be transplanted into the field insectary as needed.

Establishment of rearing site at CSU Agricultural Research, Education and Demonstration Center.

Mecinus janthinus rearing: In each of 2, 3m x 3m x 2.5m field cages we will transplant 8 individual plants from each of the 4 (state) x 2 (population) collection sites, for a total of 64 plants per cage. Plants will be planted 0.25 m on center, and will be labeled with the population from which it was collected. Plants will be fertilized and watered to maintain a vigorous and

healthy plant. Previous lab work by my group determined that *Mecinus janthinus* performs better on larger and more vigorous plants.

In late May we will release 300 adult *Mecinus janthinus* into each cage. Adults will be allowed to oviposit at will, and we will census the population every 2 weeks by shaking the foliage of 10 replicate plants into a 0.25 m x 0.30 m Rubbermaid™ tub and counting all adults. After counting, all adults will be returned to the cage.

In addition to estimating adult survival through the season, we will estimate oviposition through time by removing 4 sets of 5 (M:F) pairs of weevils from the field cage and placing them into 4 separate lab cages each containing 1 yellow toadflax plant. Weevils will be allowed to oviposit for 5 days and then will be removed from the plants. Plants will then be placed outside in the CSU shade house (away from natural sources of *Mecinus*) and watered and fertilized as needed. Stems from these plants will be dissected in the fall of each year to determine oviposition rates.

In November of each year, after all weevil development has completed, we will harvest all stems from each plant by severing them at ground level. Stems will be placed, by plant, into 0.15 x 0.20 m clear plastic sweater boxes and held in a growth chamber set at 2°C (day) -4°C (night) with a 8:16 D:N photoperiod. This combination of temperatures and photoperiod has worked well for *M. janthinus* in the past. In late April we will gradually increase the temperature profile to 20°C (day) and 14°C (night) and a 14:10 D:N photoperiod. The increase in temperatures and day lengths will occur over a 4-week period.

Sweater boxes will be monitored twice weekly for emerged adults. Adults will be removed from the boxes and placed in groups of 200 (50:50 sex ratio) in cardboard containers with sprigs of yellow toadflax in moist florist foam. Containers will be stored at 10° C until they are used for experiments on impact (see below), addition to the rearing colonies, release within Colorado or shipment to Cooperators in other states.

***Rhinusa linariae* rearing**

We will use a similar design to mass rear *R. linariae*. We will use 2, 3m x 3m x 2.5m field cages each containing 64 yellow toadflax plants on 0.25 m centers to rearing the root-galler. Up to 200 overwintered *R. linariae* adults will be introduced into each cage in early May. Weevil abundance will be measured in the same manner as with *M. janthinus*, above.

As weevils emerge from galls in late summer, we will transfer these to a new 3m x 3m x 2.5m cage containing a new population of 64 toadflax plants. The first 400 weevils produced will be used for continuation of the mass rearing population (200 for each of 2 cages). We will transfer additional weevils to sweater boxes containing blocks of florist foam with 5 mm holes drilled through. After all weevils have emerged in late summer, we will excavate the plants and record the number of empty galls on each plant.

Boxes containing weevils will be overwintered as with *M. janthinus*, above. These weevils will be used for experiments on impact (see below) release within Colorado or shipment to Cooperators in other states.

Measurement of the impact of *M. janthinus* on potted plants under low – med – high resource conditions and low – med – high attack rates

We will use a factorial design to measure the impact of *M. janthinus* on potted yellow toadflax plants grown under low, medium and high resource conditions. Plants, started from seed the summer before, will be grown in 1 gallon pots in the shade house. In June, individually caged plants will receive 0, 2 pairs (M:F) or 8 pairs of weevils. Weevils will be allowed to oviposit for 4 weeks and then we will remove the weevils and the cages. Each weevil treatment level will be crossed with 3 resource levels grown in 50:50 low N mineral soil:sand. Low resource = no additional fertilizer, medium resource = ¼ strength Hoagland's solution every 2 weeks, high = ½ strength Hoagland's solution every 2 weeks. Each treatment combination will be replicated 8 times for a total of 3 (weevil) x 3 (resource) x 8 (replicates) = 72 pots. Plants will be maintained outside until the end of the growing season. At this time, plants will be harvested and we will measure the total above and below-ground biomass and the number of adult weevils produced for each. This experiment will be performed in 2017 and again in 2018.

Measure of the impact of *R. linariae* under low – med – high resource conditions and low – med – high attack rates

We will use the same design as above to measure the impact of *R. linariae* on yellow toadflax performance, substituting *R. linariae* for *M. janthinus*. Weevil treatment sizes will be 0, 4 pairs or 8 pairs per plant. Resource treatments, number of replicates and data collection will be the same as for *M. janthinus*, above. This experiment will be performed in 2017 and 2018.

Data Analysis

Mass rearing results: We will use mixed model ANOVA to evaluate the effect of yellow toadflax source population on weevil production. We will also record data on the persistence of the weevils through time, and for *M. janthinus*, we will also record the insect's oviposition schedule.

We will use mixed model ANOVA on appropriately transformed data to evaluate the effects of the biocontrol agents on toadflax performance as resource concentrations change. We will also examine the effects of resource additions on weevil performance using a similarly structured model as above.

Expected Products and Outcomes

In years 2 and 3, we will share our rearing methodologies, experience and data on weevil survival and oviposition through time with both the Nez Perce and the Palisade, CO biocontrol programs. These data and our experiences will assist in transferring the results of this project to other key stakeholders, and it is our hope that these cooperators will be able to continue mass rearing of these agents after this project has ended.

Dissemination of *R. linariae* and *M. janthinus* to CO and other stakeholders.

Currently, demand for *M. janthinus* and *R. linariae* exceeds supply. We will work with USFS to identify cooperators with appropriate release sites. Priority will be given to those cooperators with a history of or commitment to collecting data on plant densities and weevil establishment through time. I will obtain the necessary permits for interstate movement of the weevils.

Publish a paper on the impact of *R. linariae* and *M. janthinus* under varying resource conditions.

Resource availability commonly alters a plant's ability to tolerate herbivory, but research indicates that resource addition can either *increase* or *decrease* plant tolerance. Ecological theory (Wise and Abrahamson 2005) suggests that the impact of the root galler will *decrease* with increasing below ground resources. This is because in conditions where soil nutrients are not limiting, root herbivory should have little impact on plant performance. In contrast, the impact of the stem borer should be insensitive to resource conditions. Above ground herbivory should not impact the ability of the plant to acquire the limiting resource (soil nutrients) and thus plants should be equally affected by *M. janthinus* across resource levels.

In addition to addressing an important theoretical aspect of weed biological control impacts, these results will also provide an estimate of biocontrol agent impacts, data that has not been previously published for either species.

These results will also be presented at the W3185 annual meeting and the Annual meeting of the Entomological Society of America.

Train a graduate student in weed biological control

The majority of project funds requested will go toward the training of a graduate student. This (to be determined) master's student will gain essential skills in the collection, rearing and implementation of weed biological control. Sufficient travel funds are requested to send this student to regional and National meetings of biological control scientists and practitioners to complete this student's training.

Budgets**Sponsor Contribution**

		Year 1	Year 2	Year 3	Total
PERSONNEL SALARIES					
	Academic Faculty:	\$0	\$0	\$0	\$0
	Fringe Rate	\$0	\$0	\$0	\$0
	Administrative Prof	\$0	\$0	\$0	\$0
	Fringe	\$0	\$0	\$0	\$0
	State Classified:	\$0	\$0	\$0	\$0
	Fringe	\$0	\$0	\$0	\$0
	Post Docs, Interns, (\$0	\$0	\$0	\$0
	Fringe	\$0	\$0	\$0	\$0
	Other Staff	\$0	\$0	\$0	\$0
	Fringe	\$0	\$0	\$0	\$0
	Student Hourly:	\$0	\$0	\$0	\$0
	Fringe	\$0	\$0	\$0	\$0
	GRA's:	\$18,750	\$19,500	\$20,280	\$58,530
	Fringe	\$1,350	\$1,422	\$1,498	\$4,270
TOTAL SALARY:		\$18,750	\$19,500	\$20,280	\$58,530
TOTAL FRINGE:		\$1,350	\$1,422	\$1,498	\$4,270
TOTAL PERSONNEL:		\$20,100	\$20,922	\$21,778	\$62,800
DOMESTIC TRAVEL:		\$2,000	\$2,288	\$3,894	\$8,182
INTERNATIONAL TRAVEL:		\$1,200	\$0	\$0	\$1,200
MATERIALS AND SUPPLIES		\$1,500	\$1,040	\$0	\$2,540
TOTAL OTHER DIRECT:		\$0	\$0	\$0	\$0
SUBCONTRACTS:		\$0	\$0	\$0	\$0
EQUIPMENT:		\$0	\$0	\$0	\$0
TOTAL DIRECT COSTS:		\$24,800	\$24,250	\$25,672	\$74,722
Facilities & Administrative:		\$0	\$0	\$0	\$0
Consortium F&A		\$0	\$0	\$0	\$0
TOTAL:		\$24,800	\$24,250	\$25,672	\$74,722

CSU Contribution

CSU Contribution		Year 1	Year 2	Year 3	Total
PERSONNEL SALARIES					
	Academic Faculty:	\$4,568	\$7,390	\$7,686	\$19,644
	Fringe Rate	\$1,160	\$1,901	\$2,003	\$5,064
	Administrative Prof	\$0	\$0	\$0	\$0
	Fringe	\$0	\$0	\$0	\$0
	State Classified:	\$0	\$0	\$0	\$0
	Fringe	\$0	\$0	\$0	\$0
	Post Docs, Interns, (\$0	\$0	\$0	\$0
	Fringe	\$0	\$0	\$0	\$0
	Other Staff	\$0	\$0	\$0	\$0
	Fringe	\$0	\$0	\$0	\$0
	Student Hourly:	\$0	\$0	\$0	\$0
	Fringe	\$0	\$0	\$0	\$0
	GRA's:	\$0	\$0	\$0	\$0
	Fringe	\$0	\$0	\$0	\$0
TOTAL SALARY:		\$4,568	\$7,390	\$7,686	\$19,644
TOTAL FRINGE:		\$1,160	\$1,901	\$2,003	\$5,064
TOTAL PERSONNEL:		\$5,728	\$9,291	\$9,689	\$24,708
DOMESTIC TRAVEL:		\$0	\$0	\$0	\$0
INTERNATIONAL TRAVEL:		\$0	\$0	\$0	\$0
MATERIALS AND SUPPLIES		\$0	\$0	\$0	\$0
INDIRECT:		\$12,466	\$0	\$38,607	\$51,073
TOTAL DIRECT COSTS:		\$18,194	\$9,291	\$48,296	\$75,781
INDIRECT (F&A) COSTS:		\$0	\$0	\$0	\$0
TOTAL:		\$18,194	\$9,291	\$48,296	\$75,781

Budget narrative

Personnel

I request funding for 1 Graduate Research Associate for 7.5 months (1 semester and 3 summer months) per year at an initial salary of \$2,500 per month in the first year, with a 4% inflationary increase in years 2 and 3. The CSU fringe rate for the GRA is calculated at 7.2% in year 1, 7.29% in year 2, and 7.39% in year 3. Fringe will be charged at the rate in effect when the expense is incurred. The total salary and fringe requested comes to **\$62,800** for the 3 years of the project. The GRA will take the lead in executing all field and greenhouse rearing and experiments.

Travel

For domestic travel, I request \$2,000 in year 1 for travel to collection sites in Colorado and Montana for *M. janthinus* and yellow toadflax. In year 2 I request \$1,000 to send the GRA to the annual meeting of the Western Biological Control coordinating committee (W3185) and \$1,200 to travel release sites in Colorado. In year 3, I request \$1,000 for travel to the annual meeting of

the Western Biological Control coordinating committee (W3185), \$1,400 to travel to the annual meeting of the Entomological Society of America and \$1,200 to travel to release sites in Colorado.

For international travel, I request \$1,200 in year 1 to collect *R. linariae* in British Columbia, Canada.

Travel costs include mileage or airfare, plus lodging, and per diem if overnight stays are necessary. Total travel funds request come to **\$9,382** for the 3 years of the project.

Materials and Supplies

In year 1 I request \$500 for irrigation supplies for the insectary, \$600 for greenhouse bench space to establish the rearing colony and \$400 for materials to repair field cages. In year 2, I request \$624 for bench space and \$416 for cage materials. The total request for materials and supplies is **\$2,540**

The total funds requested are **\$74,722**.

CSU Matching funds:

Personnel

PI Norton will contribute 0.40, 0.70 and 0.70 months in years 1, 2, and 3 to project design, hiring and mentoring the GRA and reporting. Dr. Norton's salary is \$10,151 per month, with a 4% inflationary increase in years 2 and 3. CSU fringe for faculty is calculated at 25.4% in year 1, 25.73% in year 2 and 26.12% in year 3. Fringe will be charged at the rate in effect when the expense is incurred. The total salary and fringe contributed equals **\$24,708** for the 3 years of the project.

Indirect

Indirect costs on CSU's cost share amount, plus unrecovered indirect costs on the federal request, which are calculated at CSU's federally negotiated rate of 51% MTDC in Year 1 and 52% MTDC in years 2 and 3 of modified total direct costs: $\$12,466 + \$38,607 = \$51,073$

The total matching funds are **\$75,144**

Timeline

	2016				2017				2018				2019	
	W	Sp	Su	F	W	Sp	Su	F	W	Sp	Su	F	W	Sp
Project starts		xx												
Field collect YT, <i>M. janthinus</i>.		xx												
Plant rearing populations.		xx												
Field collect <i>R. linariae</i>.			xx											
Collect 1st year Mj stems, evaluate.				xx										
Overwinter Mj, RI				xx										
Release Mj, RI to mass rearing cages.						xx								
Start Mj impact exp.						xx								
Evaluate Mj impact, Mj and RI performance in rearing cages. Data Analysis.							xx							
Overwinter Mj, RI.							xx	xx						
Release Mj, RI to mass rearing cages.										xx				
Start Mj impact exp.										xx				
Start RI impact exp.										xx				
Evaluate Mj impact, Mj and RI performance in rearing cages.												xx		
Data analysis, write paper.												xx	xx	xx
Mj available for re-distribution.						xx					xx			xx
RI available for re-distribution											xx			xx

References

- Jordan, K. 1994. *Gymnetron linariae* Panzer (Col.: Curculionidae). A candidate for biological control of Dalmatian and yellow toadflax in North America. International Institute of Biological Control.
- Polunin, 1969. Flowers of Europe. Oxford University Press.
- Stohlgren T.J., Binkley D., Chong G.W., Kalkhan M.A., Schell L.D., Bull K.A., Otsuki Y., Newman G., Bashkin M., Son Y. 1999. Exotic plant species invade hot spots of native plant diversity. Ecol. Mono. 69(1): 25-46.
- Tosevski, I., R. Caldara, J. Jovic, G. Hernandez-Vera, C. Baviera, A. Gassmann and B.C. Emerson. 2011. Morphological, molecular and biological evidence reveal two cryptic species in *Mecinus janthinus* Germar (Coleoptera, Curculionidae), a successful biological control agent of Dalmatian toadflax, *Linaria dalmatica* (Lamiales, Plantaginaceae). Syst. Entomol. 36: 741-753.
- Ward SM, Reid SD, Harrington J. 2008. Genetic variation in invasive populations of yellow toadflax (*Linaria vulgaris*) in the western United States. Weed Science 56(3) 394-399.
- Wise, M.J. and Abrahamson, W.G. 2005. Beyond the compensatory continuum: Environmental resource levels and plant tolerance of herbivory. Oikos 109: 417-428.

Andrew P Norton**EDUCATION**

1985 B.A. Biology, University of California, Santa Cruz
 1995 Ph. D. Entomology, University of California, Berkeley

ACADEMIC POSITIONS

Years (2013 -) Professor, BSPM, Colorado State University
 Years (2008 -2013) Associate Professor, BSPM, Colorado State University
 Years (2002-2008) Assistant Professor, BSPM, Colorado State University,
 Years (2000-2002) Research Associate / Assistant Professor, Special Appointment,

Pest ecology and management, with an emphasis on plant insect interactions and weed biological control**Recent relevant publications**

Bright, D.E., Kondratieff B.C., and **Norton A.P.** (2014) First record of the “splendid tamarisk weevil”, *Coniatus splendidulus* F. (Coleoptera: Curculionidae: Hyperinae), in Colorado, USA. *The Coleopterists Bulliten* 67(3): 1-2.

Cariveau, D. P., **Norton, A.P.** 2014. Direct effects of a biocontrol agent are greater than indirect effects through flower visitors for the alien plant Dalmatian toadflax (*Linaria dalmatica*: Scrophulariaceae). *Biological Invasions*, 16(9), 1951-1960.

Williams, W. I., Friedman, J. M., Gaskin, J. F., **Norton, A.P.** 2014. Hybridization of an invasive shrub affects tolerance and resistance to defoliation by a biological control agent. *Evolutionary Applications*, 7(3), 381-393.

Blumenthal, D. **Norton, A.P.**, Cox, S.E., Hardy, E.M., Liston, G.E., Kennaway, L., Booth, D.T., Derner, J.D. 2012. *Linaria dalmatica* invades south-facing slopes and less grazed areas in grazing tolerant mixed-grass prairie. *Biological Invasions* 14(2) 395-404

Cariveau, D.† and **Norton, A.P.**. 2009. Spatially contingent interactions between an exotic and native plant mediated through flower visitors. *Oikos*: 118: 107-114.

Bean, D., **Norton, A.P.**, Jashenko, R., Cristofaro M., and Schaffner, U. 2008 Status of Russian olive biological control in North America. *Ecological Restoration* 26:105-107.

Norton, A.P., Blair, A.C.†, Hardin, J.G., Nissen, S.J. and Brunk, G.R. 2008. Herbivory and novel weapons: No evidence for enhanced competitive ability or allelopathy induction of *Centaurea diffusa* by biological controls. *Biological Invasions* 10(1): 79-88.