Comparison of Fumigant and Nonfumigant Nematicides for Control of *Meloidogyne chitwoodi* on Potato¹

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Abstract: The fumigant 1,3-dichloropropene (1,3-D) effectively controlled Meloidogyne chitwoodi on Russet Burbank potato, Solanum tuberosum. There was a maximum of 4% infected and galled tubers from the 1,3-D treatment after 2,028 degree-days with a base temperature of 5 C (DD5). This compared to 66% infected and galled tubers in aldicarb at-plant treated plots. Soil temperature, as determined by DD5, and timing of chemical applications affected the nematicidal activity on *M. chitwoodi* (P < 0.05). Aldicarb was most effective when applied postplant (PP) during the nematode reproductive cycle. After 1,684 DD5 of growth, there were 59, 26, 22, and 6% infected and galled tubers from untreated control plots and aldicarb treatments of 2.1 g/m row at 600 DD5, 2.1 g/m row at 1,228 DD5, and 1.3 g/m row at 600 DD5 plus 2.1 g/m row at 1,228 DD5, respectively. No aldicarb treatments were effective over a growing period of 2,028 DD5; 34% of the tubers were infected and galled following the most effective aldicarb treatment (1.3 g/m row at 504 DD5 plus 2.1 g/m row at 996 DD5).

Key words: aldicarb, chemical control, Columbia root-knot nematode, degree-day, 1,3-dichloropropene, Meloidogyne chitwoodi, potato, reproduction, soil temperature, Solanum tuberosum.

The Columbia root-knot nematode, *Meloidogyne chitwoodi* Golden, O'Bannon, Santo & Finley, is one of the most important pathogens attacking potato, *Solanum tuberosum* L. (4). It is found in most potatoproducing regions of the western United States (2,5,8,13). It invades and galls potato tubers, thus suppressing both tuber quantity and quality (4,5,10,11).

Soil temperature affects M. chitwoodi hatching, development, and reproduction and thereby influences the severity of nematode invasion and tuber galling (5). Approximately 1,000 and 500 degree-days at a base temperature of 5 C (DD5) are required for the reproduction of the first and future nematode generation, respectively (5,11). The greater the number of nematode generations during a growing season, the greater the degree of infection and galling of potato tubers (5,6,9,11,14). Because of the large population increase of M. chitwoodi during the growing season, only fumigant nematicides provide acceptable chemical control (2,5,10,12).

The objective of this study was to determine 1) whether the nematicidal efficacy of aldicarb was comparable to 1,3-dichloropropene (1, 3-D) if application timing was correlated to nematode reproduction and 2) the effect of soil temperature, as determined by DD5, on aldicarb control of M. *chitwoodi*.

MATERIALS AND METHODS

Three separate experiments were conducted over a 3-year period at Logan, Utah, in microplots fabricated from redwood. Microplot boxes were 3.0 m wide \times 4.3 m $\log \times 0.6$ m deep and were open at the bottom. Boxes were buried 0.46 m deep, and soil was excavated from the microplots and replaced with a 1,3-D fumigated sandy loam soil (80% sand, 11% silt, 9% clay, 0.7% organic carbon; pH 7.7). Meloidogyne chitwoodi inoculum came originally from potato at Ft. Hall, Idaho, and was cultured in a greenhouse on wheat, Triticum aestivum L. cv. Nugaines. Inoculum was introduced into the microplots on Nugaines transplants and maintained on that host for 2 years.

Spring mean nematode population densities (Pi) of second-stage juveniles (J2) were determined from 10 soil samples per microplot, composited from 14 soil cores (30cm centers \times 30 cm deep) per sample, collected with a 1.9-cm-d probe. Elutriation (3) and alcohol flotation (7) were used to process 100-cm³ aliquants of each soil sam-

Received for publication 25 March 1988.

¹ Cooperative investigation by USDA ARS and the Utah Agricultural Experiment Station. Journal Paper No. 3615. ² Nematologist, USDA ARS, Forage and Range Research Laboratory, Utah State University, Logan, UT 84322.

Treat- ment†	Rate a.i.	Time of application‡	Yield (quintal/ha)	Infected and galled tubers (%)§	
1,3-D	9.4 g/m (168 kg/ha)	Preplant	328 a	0 a	
1,3-D	7.6 g/m (135 kg/ha)	Preplant	336 a	2 ab	
Aldicarb	2.1 g/m (5.6 kg/ha)	At plant	307 a	25 d	
Aldicarb	2.1 g/m (5.6 kg/ha)	615 DD5	317 a	21 d	
Aldicarb	2.1 g/m (5.6 kg/ha)	916 DD5	308 a	13 c	
Aldicarb	2.1 g/m (5.6 kg/ha)	1,221 DD5	312 a	6b	
Untreated			316 a	39 e	

TABLE 1. Effect of time of application on chemical control of *Meloidogyne chitwoodi* on Russet Burbank potato after 1,344 DD5.

Means within a column with same letter are not different according to Duncan's new multiple-range test (P < 0.05).

 \pm 1,3-D applied broadcast 16 days preplant at 11 C. Aldicarb 15G applied at plant in a 30-cm band directly over the row and rototilled into the soil 15 cm deep. Aldicarb 15G applied postplant in 15-cm band on each side of row, and raked into soil 5-8 cm deep. All treatments were followed immediately with 2.5-3.0 cm water through sprinkler irrigation system. \pm DD5 = degree-days with a base of 5 C.

\$ Tubers stored at 7 C for 64 days (128 DD5) after harvest before tuber infection and galling percentages were determined.

ple. Mean numbers of J2 were 1.8 (range: 1.6-2.1)/cm³ soil.

with two thermographs located in two selected microplots.

The nematicide treatments, rates, and time of applications for each experiment are listed in Tables 1-3. The 1,3-D was applied broadcast with a fumigant injectorgun by injecting it 30 cm deep on 15-cm intervals. The fumigant was applied 16-18 days preplant at soil temperatures of 9-11 C. At-plant (AP) aldicarb was applied with a shaker jar in a 30-cm band directly over the row and rototilled into the soil 15 cm deep immediately before planting. For postplant (PP) treatments, equal amounts of aldicarb were applied in a 15-cm band on each side of the row and hand raked into the soil 5-8 cm deep. All treatments received 2.5-3.0 cm water through a sprinkler irrigation system immediately after each aldicarb application.

Soil was fertilized immediately before planting at rates used in commercial potato production in southeastern Idaho (5). Certified Russet Burbank seed potato was planted at 15-cm spacing on 100-cm rows, three rows per microplot. No herbicide was used, and the plots were hand weeded. Soil temperatures were recorded at 15 cm deep Fifty-six soil subsamples were collected with a 1.9-cm-d probe 30 cm deep at 15cm intervals 15 and 30 cm from each side of the center of each row immediately before harvest from each microplot. Subsamples were composited (eight subsamples per sample), thoroughly mixed, and 100-cm³ aliquants of each sample were processed to determine the numbers of *M. chitwoodi* J2. Tubers were harvested after 1,344 DD5, 1,684 DD5, and 2,028 DD5 (1) in experiments 1, 2, and 3, respectively, and fresh tuber weights were determined.

Tubers invaded by *M. chitwoodi* J2 were harvested before development of nematode-induced disease symptoms; i.e., necrotic tissue encircling embedded females preceding tuber galling. Hence, tubers were stored at 7 C for 64 days (128 DD5) to allow for symptom development. This simulated commercial operations where tubers are stored at approximately 7 C before being graded and processed. Data on tuber yields and infection and galling were subjected to analysis of variance and linear regression analysis. Treatment means were

Treat- ment†	Rate a.i.	Time of application	J2/cm³ soil 1,684 DD5‡	Infected and galled tubers§ (%)	Yield (quintal/ha)
1,3-D	7.6 g/m (135 kg/ha)	Preplant	0.6 a	2 a	312 a
Aldicarb	2.1 g/m (5.6 kg/ha)	At plant	8.3 g	36 e	305 a
Aldicarb	2.1 g/m (5.6 kg/ha)	600 DD5	5.8 f	26 d	313 a
Aldicarb	2.1 g/m (5.6 kg/ha)	1,228 DD5	4.2 def	22 cd	290 a
Aldicarb	1.3 + 1.3 g/m (3.4 + 3.4 kg/ha)	AP + 600 DD5	5.4 ef	24 cd	297 a
Aldicarb	1.3 + 1.3 g/m (3.4 + 3.4 kg/ha)	AP + 1,228 DD5	3.3 cd	18 c	302 a
Aldicarb	1.3 + 1.3 g/m (3.4 + 3.4 kg/ha)	600 DD5 + 1,228 DD5	3.0 cd	19 c	298 a
Aldicarb	1.3 + 2.1 g/m (3.4 + 5.6 kg/ha)	AP + 600 DD5	4.1 de	18 c	317 a
Aldicarb	1.3 + 2.1 g/m (3.4 + 5.6 kg/ha)	AP + 1,228 DD5	1.9 abc	10 b	310 a
Aldicarb	1.3 + 2.1 g/m (3.4 + 5.6 kg/ha)	600 DD5 + 1,228 DD5	1.0 ab	6 ab	322 a
Untreated	. 0, 7		10.6 h	59 f	314 a

TABLE 2. Effect of time of application on chemical control of *Meloidogyne chitwoodi* on Russet Burbank potato after 1,684 DD5.

Means within a column with same letter are not different according to Duncan's new multiple-range test (P < 0.05).

† 1,3-D applied broadcast 16 days preplant at 9 C. Aldicarb 15G applied at plant (AP) in a 30-cm band directly over the row and rototilled into the soil 15 cm deep. Aldicarb 15G applied postplant in 15-cm band on each side of row, and raked into soil 5-8 cm deep. All aldicarb treatments were followed immediately with 2.5-3.0 cm water through sprinkler irrigation system.

 $\ddagger DD5 = degree-days$ with a base of 5 C.

\$ Tubers stored at 7 C for 60 days (120 DD5) after harvest before tuber infection and galling percentages were determined.

compared by Duncan's new multiple-range test.

All experiments were similar, except for differences of accumulated DD5, and time of aldicarb applications.

RESULTS AND DISCUSSION

Experiment 1: Infection and galling of potato tubers by M. chitwoodi were less (P < 0.05) in all treated than untreated plots (Table 1). Only the aldicarb PP treatment at 1,221 DD5 was as effective as 1,3-D in controlling M. chitwoodi. There were no differences in tuber yields among treatments.

Experiment 2: All aldicarb treatments reduced (P < 0.05) the nematode population density below that of the untreated control (Table 2). The single AP treatment of aldicarb provided the poorest nematode control of the chemical treatments. The PP applications of aldicarb were consistently better than AP in controlling *M. chitwoodi.* The most effective aldicarb treatment was 1.3 g/m at 600 DD5 PP plus 2.1 g/m at 1,228 DD5 PP. This treatment was equivalent to the 1,3-D treatment. There was a positive correlation (r = 0.94) between the final nematode population density (Pf) and the percentage of infected and galled potato tubers.

Experiment 3: The 1,3-D was more effective than aldicarb in reducing nematode population densities and infection and galling of potato tubers (Table 3). There was a strong correlation between the percentage of infected and galled tubers and the nematode Pf (r = 0.90). None of the aldicarb treatments reduced *M. chitwoodi* infection and galling of tubers below that allowed by potato processors (5). The

Treat- ment†	Rate a.i.	Time of application	J2/cm³ soil 2,028 DD5‡	Infected and galled tubers (%)§	Yield (quintal/ha)
1,3 - D	7.6 g/m (135 kg/ha)	Preplant	0.8 a	4 a	326 a
Aldicarb	2.1 g/m (5.6 kg/ha)	At plant	12.5 f	66 e	312 a
Aldicarb	2.1 g/m (5.6 kg/ha)	996 DD5	10.2 e	56 de	319 a
Aldicarb	1.3 + 1.3 g/m (3.4 + 3.4 kg/ha)	AP + 996 DD5	8.3 d	49 cd	297 a
Aldicarb	1.3 + 1.3 g/m (3.4 + 3.4 kg/ha)	504 DD5 + 996 DD5	7.8 cd	45 c	332 a
Aldicarb	1.3 + 2.1 g/m (3.4 + 5.6 kg/ha)	AP + 996 DD5	6.5 bc	39 bc	321 a
Aldicarb	1.3 + 2.1 g/m (3.4 + 5.6 kg/ha)	504 DD5 + 996 DD5	5.9 b	34 b	318 a
Untreated	, G, ,		14.9 g	94 f	303 a

TABLE 3. Effect of time of application on chemical control of *Meloidogyne chitwoodi* on Russet Burbank potato after 2,028 DD5.

Means within a column with same letter are not different according to Duncan's new multiple-range test (P < 0.05).

† 1,3-D applied broadcast 16 days preplant at 9 C. Aldicarb 15G applied at plant (AP) in a 30-cm band directly over the row and rototilled into the soil 15 cm deep. Aldicarb 15G applied postplant in 15-cm band on each side of row, raked into the soil 5-8 cm deep. All aldicarb treatments followed immediately with 2.5-3.0 cm water through sprinkler irrigation system.

 $\ddagger DD5 = degree-days$ with a base of 5 C.

§ Tubers stored at 7 C for 64 days (128 DD5) after harvest before tuber infection and galling percentages were determined.

greatest Pf values were in the 2.1 g/m aldicarb AP treatment.

The relationship between soil temperature and nematode reproduction was evident from this study. The greater the DD5, the greater the nematode activity, resulting in increased nematode reproduction and tuber infection and galling. There were no differences in plant growth and tuber yields among treatments. This is consistent with the results from a previous study made in eastern Idaho (5). These findings, however, differ from those reported from Washington and western Idaho (10,11). In these states, the reduction in tuber yields from M. chitwoodi was due to increased stress of the nematode on plant growth. Because of the greater Pi levels and DD5, there are more nematode generations and increased parasitism (5,10). It is the effect of M. chitwoodi on the quality, not quantity, of tuber yields that makes it such an important pathogen to potato in eastern Idaho.

Nonfumigant nematicides are easy to apply and are not generally phytotoxic. To control *M. chitwoodi* effectively, however, a nonfumigant nematicide must have a longer residual period than aldicarb or its application must be timed accurately. Continued investigations are needed for better methodology of nonfumigant nematicide applications for increased efficacy.

LITERATURE CITED

1. Arnold, C. Y. 1960. Maximum-minimum temperatures as a basis for computing heat units. Proceedings of the American Society for Horticultural Science 76:682-692.

2. Bahme, J. B., S. D. Van Gundy, and M. N. Schroth. 1987. Population dynamics and control of *Meloidogyne chitwoodi* on potato in northern California. Journal of Nematology 19:510-511 (Abstr.).

3. Byrd, D. W., Jr., K. R. Barker, H. Ferris, C. J. Nusbaum, W. E. Griffin, R. H. Small, and C. A. Stone. 1976. Two semi-automatic elutriators for extracting nematodes and certain fungi from soil. Journal of Nematology 8:206-212.

4. Finley, A. M. 1981. Histopathology of *Meloi*dogyne chitwoodi (Golden et al.) on Russet Burbank potato. Journal of Nematology 13:486-491.

5. Griffin, G. D. 1985. Host-parasite relationship of *Meloidogyne chitwoodi* on potato. Journal of Nematology 17:395-399.

6. Inserra, R. N., G. D. Griffin, and D. V. Sisson.

1983. Effect of temperature and root leachates on embryonic development and hatching of *Meloidogyne chitwoodi* and *M. hapla.* Journal of Nematology 15: 123-127.

7. Jenkins, W. R. 1964. A rapid centrifugal-flotation technique for separating nematodes from soil. Plant Disease Reporter 48:692.

8. Nyczepir, A. P., J. H. O'Bannon, G. S. Santo, and A. M. Finley. 1982. Incidence and distribution of *Meloidogyne chitwoodi* and *M. hapla* in potato from the northwestern United States. Journal of Nematology 14:347-353.

9. O'Bannon, J. H., and G. S. Santo. 1984. Effect of temperature on reproduction of *Meloidogyne chitwoodi* on *M. hapla* alone and in combination on potato and *M. chitwoodi* on rotation plants. Journal of Nematology 16:309-312.

10. Pinkerton, J. N., and G. S. Santo. 1986. Con-

trol of *Meloidogyne chitwoodi* in commercially grown Russet Burbank potatoes. Plant Disease 70:860-863.

11. Pinkerton, J. N., G. S. Santo, and H. Mojtahedi. 1986. Population dynamics of *Meloidogyme chitwoodi* in relation to Russet Burbank potato tuber penetration. Journal of Nematology 18:627 (Abstr.).

12. Santo, G. S., and M. Qualls. 1984. Control of *Meloidogyne* spp. on Russet Burbank potato by applying metham sodium through center pivot irrigation systems. Journal of Nematology 16:159–161.

13. Santo, G. S., J. H. O'Bannon, A. M. Finley, and A. M. Golden. 1980. Occurrence and host range of a new root-knot nematode (*Meloidogyne chitwoodi*) in the Pacific Northwest. Plant Disease 64:951-952.

14. Santo, G. S., and J. H. O'Bannon. 1981. Effect of soil temperature on the pathogenicity and reproduction of *Meloidogyne chitwoodi* and *M. hapla* on potato. Journal of Nematology 13:483-486.