Host Suitability of 32 Common Weeds to *Meloidogyne* hapla in Organic Soils of Southwestern Quebec¹

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Abstract: Thirty-two weeds commonly found in the organic soils of southwestern Quebec were evaluated for host suitability to a local isolate of the northern root-knot nematode Meloidogyne hapla under greenhouse conditions. Galls were observed on the roots of 21 species. Sixteen of the 21 had a reproduction factor (Pf/Pi = final number of M. hapla eggs and juveniles per initial number of M. hapla juveniles per pot) higher than carrot (Pf/Pi = 0.37), the major host crop in this agricultural area. Tomato cv. Rutgers was also included as a susceptible host and had the highest Pf/Pi value of 13.7. Bidens cernua, B. frondosa, B. vulgata, Erysimum cheiranthoides, Eupatorium maculatum, Matricaria matricarioides, Polygonum scabrum, Thalictrum pubescens, Veronica agrestis, and Sium suave are new host records for M. hapla. Bidens cernua, B. frondosa, B. vulgata, D. carota, M. matricarioides, Pasticana sativa, P. scabrum, S. suave, and Thlaspi arvense sustained moderate to high galling by M. hapla and supported high M. hapla production ($12.4 \leq Pf/Pi \geq 2.9$). Capsella bursa-pastoris, Chrysanthemum leucanthemum, Gnaphalium uliginosum, Stellaria media, and Veronica agrestis sustained moderate galling and supported moderate M. hapla reproduction $(2.8 \leq Pf/Pi \geq 0.5)$. Chenopodium album, C. glaucum, E. cheiranthoides, P. convolvulus, Portulaca oleracea, and Rorippa islandica supported low reproduction (0.25 \leq Pf/Pi \geq 0.02) and sustained low galling. Galling was observed on Senecio vulgaris but no eggs or juveniles; thus, S. vulgaris may be useful as a trap plant. Eupatorium maculatum, and T. pubescens harbored no distinct galling but supported low to moderate M. hapla reproduction, respectively. Amaranthus retroflexus, Ambrosia artemisiifolia, Echinochloa crusgalli, Erigeron canadensis, Oenothera parviflora, Panicum capillare, Setaria glauca, S. viridis, and Solidago canadensis were nonhosts. Our results demonstrate the importance of adequate weed control in an integrated program for the management of M. hapla in organic soil.

Key words: Canada, host range, Meloidogyne hapla, nematode, northern root-knot nematode, organic soil, weeds.

The northern root-knot nematode *Meloidogyne hapla* is an important condition pest of vegetable crops. In organic soils in southwestern Quebec, *M. hapla* is a major constraint to production of carrot (*Daucus carota* var. *sativa*), the primary cash crop (12,13). Soil fumigation with high rates of 1,3-dichloropropene is the principal control method for nematodes. However, a 2-year crop rotation with onion and small grains is an effective method for reducing *M. hapla* nematode population densities (1,2) and is used by many growers.

Weeds can reduce the efficacy of crop rotation for the management of plantparasitic nematodes. Results of host range studies showed that many weed species are highly susceptible to M. hapla and can serve as alternate hosts in carrot fields (3, 5,6,10). Our study was undertaken to determine the host suitability of 32 common weeds to a local isolate of M. hapla in the organic soils of southwestern Quebec.

MATERIALS AND METHODS

This experiment was conducted from May to July 1993 under greenhouse conditions. Young plants of Amaranthus retroflexus (redroot pigweed), Ambrosia artemisiifolia (common ragweed), Bidens cernua (nodding beggarticks), Bidens frondosa (devil's beggarticks), Bidens vulgata (tall beggarticks), Capsella bursa-pastoris (shepherd's-purse), Chenopodium album (lamb'squarters), Chenopodium glaucum (oakleaved goosefoot), Chrysanthemum leucanthemum (ox-eye daisy), Daucus carota (wild carrot), Erigeron canadensis (Canada fleabane), Echinochloa crusgalli (barnyard grass), Erysimum cheiranthoides (wormseed

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mustard), Eupatorium maculatum (spotted Joe-Pye weed), Gnaphalium uliginosum (low cudweed), Matricaria matricarioides (pineappleweed), Oenothera parviflora (smallflowered evening-primrose), Panicum capillare (witch grass), Pasticana sativa (wild parsnip), Polygonum convolvulus (wild buckwheat), Polygonum scabrum (green smartweed), Portulaca oleracea (purslane), Rorippa islandica (marsh yellow cress), Senecio vulgaris (common groundsel), Setaria glauca (yellow foxtail), Setaria viridis (green foxtail), Sium suave (water-parsnip), Solidago canadensis (Canada goldenrod), Stellaria media (chickweed), Thalictrum pubescens (pubescent meadow-rue), Thlaspi arvense, Veronica agrestis (field speedwell) were recovered from single-species infested field plots in May 1993. At transplantation, species of the Gramineae family were between first and third true-leaf stage, annual species were between cotyledon and second true-leaf stage, winter annual species were at the rosette stage; perennial species were either at the rosette stage or between the second and third true-leaf stage. Carrot cv. Sixpak II grown from seed and tomato (Lycopersicon esculentum) cv. Rutgers as 3-week-old transplants were included as susceptible checks.

On 1 May 1993, approximately 1,000 liters of an organic soil (80% organic matter; pH 5.5) was collected from an experimental field plot infested with M. hapla. The bulked soil was thoroughly mixed in a large floor bin and stored in 200-liter barrels before use. Plants were transplanted into 20-cm-diam. plastic pots containing 4 liters of infested organic soil containing an initial population density (Pi) of approximately 18,000 second-stage juveniles (12) of M. hapla as determined by a modified Baermann pan method (10). No effort was made to maintain equal numbers of plants per pot per species. Depending on the size of each weed species, the number of plants per pot varied from 1 to 70. Plants were watered daily as needed. Greenhouse temperatures were 22 ± 3 °C with natural daylight only.

The experimental design was a random-

ized complete block with 34 treatments replicated six times. Roots were separated by sifting the contents of each pot on a coarse sieve (1.5 cm) layed over a plastic tray for recovering the soil. Fresh root weight, number of galls, and eggs per root system (7) were determined after 9 weeks. The growth stage of species was also recorded according to a standardized plant growth stage scale known as BBCH (8). Soil J2 population densities in each plot were assayed from a 100-cm³ soil subsample as described for inoculum preparation. Final population densities (Pf) were computed as the total number of J2 and eggs extracted from soil and roots for each pot, and reproduction factors (Pf/Pi) were calculated.

All data were transformed to $\log_{10}(x + 1)$ values before statistical analysis. Data were subjected to analysis of variance, and treatment means were separated by the Waller-Duncan k-ratio *t* test.

RESULTS AND DISCUSSION

The 32 species of weeds evaluated in this study belonged to 27 genera in 12 families. The presence of distinct galling was observed on the roots of 21 species, and reproduction was monitored on 22 species. Sixteen of the 22 host weeds had a Pf/Pi greater than carrot cv. Sixpak II (Pf/Pi = 0.37), the major host crop in this agricultural area (Table 1). Tomato cv. Rugers was included as a susceptible host and had the highest Pf/Pi value of 13.7. Although the Pf/Pi value allowed discrimination of the relative host status of weed species, it must be used with care because Pf is a function of the time at which it is measured. In this experiment, a relatively short 9-week growth period was used because many weed species had already reached their senescence stage. This, along with a high Pi value, could explain the generally low Pf/Pi values reached by M. hapla host plants, including carrot and tomato. Bidens cernua, B. frondosa, B. vulgata, E. cheiranthoides, E. maculatum, M. matricarioides, P. scabrum, T. pubescens, S. suave, and V. agres-

	Growth stage (BBCH)	Number per root system			
Host		Galls	Eggs	Eggs + juveniles	Pf/Pi ^a
Amaranthaceae		4649 ⁻¹¹¹			
Amaranthus retroflexus	65	0	0	0	
Carvophyllaceae					
Stellaria media	75	4 hi	271 kl	1,312 i	2.85 dc
Chenopodiaceae					
Chenopodium album	77	3 ijk	71 kl	154 lm	$0.08~{ m g}$
C. glaucum	77	6 ghi	112 jk	226 kl	0.25 f
Compositae		0	5		
Ambrosia artemisiifolia	51	0	0	0	
Bidens cernua	33	531 b	58,545 b	111,278 abc	8.59 abc
B. frondosa	59	500 Ь	31,744 b	69,183 bcd	12.37 ab
B. vulgata	45	274 с	11,042 bc	30,064 de	7.11 abcd
Chrysanthemum leucanthemum	66	255 cd	5.650 efg	6.725 i	0.74 f
Erigeron canadensis	65	0	0	0	
Fubatorium maculatum	56	Õ	304 ii	3.080 i	0.27 f
Gnaphalium uliginosum	66	48 f	2 190 fo	6.256 hi	0.83 ef
Matricaria matricarioides	80	80 e	3.787 def	55.610 cde	5.05 abcd
Senecio mulgaris	69	98 f	0	0	0.00 4004
Salidago conodensis	51	0	õ	õ	
Cruciferze	51	v	0	0	
Capsella bursa-bastoris	95	10 gh	376 ik	4 076 i	047 f
Emisimum chairanthaidas	64	4 hiik	570 JK 59 1/1	78 m	0.171
Poribbo islandisa	77	ե հվել 11-ի	28 I	28 m	0.05 g
Thesti amiana	11 99	10 ~	540 h;	7 022 ch	0.02 g
Thiaspi arvense	65	rog	540 m	1,955 gi	4.55 btu
Fahimeae	00	0	0	0	
Echinochioa crusgalli	02 50	0	0	0	
Panicum capillare	52	0	0	0	
Setaria giauca	4/	0	0	0	
S. viridis	11	0	0	0	
Onagraceae	=0	0	0	0	
Oenothera parviflora	78	0	0	0	
Polygonaceae		0.111	100 11	M 4 1 11	0 F0 C
Polygonum convolvulus	93	2 jki	168 jkl	741 jk	0.53 ef
P. scabrum	73	133 de	6,969 bc	27,936 def	3.08 d
Portulacaceae					
Portulaca oleracea	79	3 ijk	53 kl	55 m	$0.04 \mathrm{g}$
Ranunculaceae					
Thalictrum pubescens	65	0	5,425 cd	15,201 etg	0.81 e
Scrophulariaceae					
Veronica agrestis	77	4 hij	94 kl	1,372 kl	1.24 ef
Solanaceae					
Lycopersicon esculentum	45	778 ab	143,952 a	258,352 a	13.71 a
Umbelliferae					
Daucus carota	66	235 cd	5,705 cde	9.821 fgh	2.92 d
D. carota var. sativa cv. Sixpak II	45	38 f	873 gh	1,096 ij	0.37 ef
Pasticana sativa	68	883 a	81,247 a	128,247 abc	6.81 abcd
Sium suave	66	574 ab	64,076 ab	193,828 ab	10.29 ab

TABLE 1. Host suitability of common weeds in southwestern Quebec to Meloidogyne hapla.

Values are means of six replications for each weed species. Mean values followed with the same letter are not significantly different according to the Waller-Duncan k-ratio t test (k = 100).

* Pf/Pi = final M. hapla population/initial M. hapla per pot.

tis are herein reported as new host records for *M. hapla*.

Bidens cernua, B. frondosa, B. vulgata, D. carota, P. scabrum, S. suave, and T. arvense

sustained moderate to high galling and supported the highest *M. hapla* reproduction (12.4 \leq Pf/Pi \geq 2.9) (Table 1). Capsella bursa-pastoris, *C. leucanthemum*, *G. uligino*-

Species	Percentage of fields infested ^a					
	Car	rrot	Onion			
	1982	1993	1982	1993		
Amaranthus retroflexus	44	46	90	100		
Ambrosia artemisiifolia	11	38	40	57		
Bidens cernua	56	31	40	43		
B. vulgata	0	8	10	0		
Capsella bursa-pastoris	_	_	0	14		
Chenopodium album	11	38	70	57		
C. glaucum	0	31	20	43		
Echinochloa crusgalli	89	31	100	86		
Erigeron canadensis	56	46	0	71		
Erysimum cheranthoides	22	8	10	57		
Eupatorium perfoliatum		_	10	14		
Matricaria matricarioides	22	31	10	29		
Panicum capillare	44	0	20	14		
Pastinaca sativa	11	23	10	14		
Polygonum convolvulus	0	8	30	29		
P. scabrum	33	31	40	57		
Portulaca oleracea	22	31	20	14		
Rorippa islandica	11	38	30	100		
Senecio vulgaris	11	31	0	43		
Setaria viridis	11	0	10	0		
Sium suave	0	8	_			
Solidago canadensis	11	8	0	14		
Stellaria media	11	54	10	71		
Thlaspi arvense	0	15	0	43		
Veronica sp.	0	15				

TABLE 2. List of weed species from carrot and onion fields in organic soils in Quebec surveyed in 1982 and 1993 within carrot and onion production fields grown in organic soil.

^aTotal number of fields surveyed: carrot, 9 in 1982 and 13 in 1993; onion, 10 in 1982 and 17 in 1993.

sum, S. media, and V. agrestis supported moderate galling and sustained moderate *M. hapla* reproduction ($2.8 \le Pf/Pi \le 0.5$). Chenopodium album, C. glaucum, E. cheiranthoides, P. convolvulus, P. oleracea, and R. islandica supported low reproduction and sustained low galling $(0.25 \le Pf/Pi \ge 0.02)$. Senecio vulgaris, on which galls were found but no eggs or J2, may serve as a trap crop for M. hapla, as previously reported (10). This weed was reported to be present in more carrot and onion fields in organic soil in 1993 than previously reported (4). Amaranthus retroflexus, A. artemisiifolia, E. crusgalli, E. canadensis, O. parviflora, P. capillare, S. glauca, S. viridis, and S. canadensis were nonhosts. Although no distinct galling was observed on E. maculatum and T. pubescens, they supported low to moderate M. hapla reproduction, respectively. The economic threshold of M. hapla on organic-grown carrots is 9 juveniles/100 cm³

soil (13); therefore, low nematode population densities recorded on some weed species represent a threat to this highly susceptible cash crop.

Many weeds are hosts on which population densities of M. hapla can be maintained or increased to higher densities than those on carrot. The presence of weeds after herbicide application in crop production fields represents an important source of nematode inoculum for the following crop. A recent weed survey in organic soils in Quebec revealed that the number of weed species did not increase in carrot fields from 1982 to 1993 (34 vs. 35) but increased dramatically in onions (38 vs. 46) (4). The percentage of fields infested with monocotyledonous weed species either has remained stable or has decreased (Table 2). Weeds from the family Cruciferae were present in more onion production fields than previously reported.

Infestations of Matricaria matricarioides, R. islandica, S. media, and S. vulgaris are increasing in both carrot and onion production fields. Weed species with the highest M. hapla reproduction factor are found in small isolated groups in fields (B. cernua, P. scabrum, S. suave), on the perimeter of fields (B. cernua, B. vulgata, S. suave), (4), or along ditches (B. cernua, B. frondosa, B. vulgata, S. suave, D. carota, and P. scabrum). These infested areas act as important reservoirs of M. hapla and may aid dissemination of the nematode during flooding of fields caused by snowmelt in the spring, or by soil deposition during winter wind erosion of fields (9).

Of the 22 weed species reported in this study as hosts for M. hapla, 12 and 10 weed species had increased their presence in carrot and onion fields, respectively, between 1982 and 1993 (Table 2). Stellaria media, C. glaucum, M. matricarioides, R. islandica, T. arvense, P. scabrum, and P. sativa were common to both carrot and onion crops in 1993. Our results emphasize the importance of adequate weed control in an integrated program for management of M. hapla in organic soils where rotation with onion and carrot is commonly practiced. Crop rotation with a nonhost crop such as cereal will not be successful unless weeds are controlled.

LITERATURE CITED

1. Bélair, G. 1992. Effects of cropping sequences on population densities of *Meloidogyne hapla* and carrot yield in organic soil. Journal of Nematology 24: 450–456. 2. Bélair, G., and L. E. Parent. 1996. Using crop rotation to control *Meloidogyne hapla* Chitwood and improve marketable carrot yield. HortScience 31: 106–108.

3. Bendixen, L. E. 1986. Weed hosts of *Meloido-gyne*, the root-knot nematodes. Pp. 101–167 *in* K. Noda and B. L. Mercado, eds. Weeds and the environment in the tropics. Chiang Mai, Thailand: Asian-Pacific Weed Society.

4. Benoit, D. L., and M. Bélanger. 1994. Inventaire des mauvaises herbes dans les cultures de carottes et d'oignons en sol organique. Agriculture and Agri-Food Canada. Horticultural Research and Development Centre Research Summary 23:24-28.

5. Edwards, W. H. and R. K. Jones. 1984. Additions to the weed host range of *Meloidogyne hapla*. Plant Disease 68:811-812.

6. Gaskin, A. G., and H. W. Crittenden. 1956. Studies of the host range of *Meloidogyne hapla*. Plant Disease Reporter 40:265-270.

7. Hussey, R. S., and K. R. Barker. 1973. A comparison of methods of collecting inocula from *Meloidogyne* spp., including a new technique. Plant Disease Reporter 57:1025–1028.

8. Lancashire, P. D., H. Bleiholder, T. Van Den Boom, P. Langelüddeke, R. Stauss, E. Weber, and A. Witzenberger. 1991. A uniform decimal code for growth stages of crops and weeds. Annals of Applied Biology 119:561–601.

9. Orr, C. C., and O. H. Newton. 1971. Distribution of nematodes by wind. Plant Disease Reporter 55:61-63.

10. Townshend, J. L. 1963. A modification and evaluation of the apparatus for the Oostenbrink direct cotton-wool filter extraction method. Nematologica 9:106–110.

11. Townshend, J. L., and T. R. Davidson. 1962. Some weed hosts of the northern root-knot nematode, *Meloidogyne hapla* Chitwood, 1949, Ontario. Canadian Journal of Botany 40:543–548.

12. Vrain, T. C. 1978. Dissémination et importance des nématodes phytoparasites dans les sols organiques du Québec. Phytoprotection 59:186 (Abstr.).

13. Vrain, T. C. 1982. Relationship between *Meloidogyne hapla* density and damage to carrots in organic soils. Journal of Nematology 14:50–57.