Nematodes as Sentinels of Heavy Metals and Organic Toxicants in the Soil

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Abstract: Field and laboratory research has repeatedly shown that free-living soil nematodes differ in their sensitivity to soil pollutants. In this paper, we analyze whether nematode genera proved sensitive or tolerant toward heavy metals and organic pollutants in six long-term field experiments. We discuss overlaps between nematode physiological responses to heavy metals and to organic pollutants, which may explain why nematodes can exhibit co-tolerance toward several contaminants. We propose a simple method for separating direct effects of soil contamination on nematode populations from indirect effects mediated through the food chain. Finally, we analyze the extent to which nematodes exhibited consistent responses across the experiments analyzed. Our results show that (a) indirect effects of pollution were generally strong; (b) fewer nematode genera were tolerant than sensitive; (c) many genera, including practically all Adenophorea, exhibited a common response pattern to contaminants; and (d) several genera of the Secernentea exhibited differential tolerance toward particular pollutants. We conclude that bioindication of soil contamination should preferentially be based on tolerant, and less on sensitive, nematodes. We provide a list of nematode genera that may potentially serve as differential bioindicators for specific soil contaminants.

Key words: Bioindicators, ecology, heavy metals, nematodes, organic toxicants, sentinels, soil pollution.

Among soil organisms, nematodes are seen as the most promising candidates for bioindication of soil status (Cortet et al., 1999; Achazi, 2002). Using the well-established classifications of nematode feeding types and cp-groups (Yeates et al., 1993; Bongers and Bongers, 1998), researchers have consistently exploited nematodes to investigate the propagation of broadly defined disturbance and fertilization effects through the soil ecosystem (Freckman and Ettema, 1993; Villeneuve et al., 2001). It has been shown repeatedly that nematodes respond differentially to xenobiotic substances (Bongers et al., 2001; De Nardo and Grewal, 2003; Jonker et al., 2004). However, a recognized concept enabling a nematode-based indication of more specific impacts than disturbance and fertilization, such as the indication of a specific heavy metal or a specific organic toxicant in the soil, is lacking.

In fact, many of the physiological responses of nematodes to toxic substances are highly unspecific (Table 1). As the very first and immediate reaction on sensing a toxic substance, nematodes can cease pharyngeal pumping and thereby avoid intake of the toxicant (Jones and Candido, 1999). Nematodes have elaborate sensorial equipment, including receptors for cadmium and copper ions (Sambongi et al., 1999), which enables them to avoid intake of a broad spectrum of substances by means of the same behavioral response mechanism. Once a toxicant has passed into the body, a plethora of molecular decontamination mechanisms is induced (Downs et al., 2001). Superoxide dismutases accumulate in response to oxidative stress and are one of the main anti-oxidant defense pathways (Fridovich, 1995). Cytochrome P450 has both physiologically relevant oxidative and reductive reactions and catalyzes many xenobiotic-based substrates (Menzel et al., 2001). Glutathiones and glutathione S-transferases are involved in the detoxification of organic xenobiotics and in the discharge of metal ions from the cell (Eaton and Bammler, 1999; Sies, 1999). Metallothioneins can act as scavengers for radicals, and mainly they protect against metal toxicity by sequestering Zn, Cu, Cd, and Hg (Klaassen et al., 1999). Phytochelatins also sequester Cd, as well as As, Ag, and Cu (Clemens et al., 2001; Vatamaniiuk et al., 2001). In this way, metal ions are neutralized within the nematode body if they cannot be excreted (Vijver et al., 2004). Storage of Pb can, for example, reach levels where visible lead particulates are formed in the oesophageal region of Panagrolaimus superbus (Williams and Seraphin, 1998). Similarly, organic pollutants can accumulate in the tissue if metabolism and excretion do not keep pace with intake rates. For example, Haitzer et al. (2000) observed accumulation of pyrene in specific, presumably lipid-rich body regions within Caenorhabditis elegans. The processes of avoidance, detoxification, and sequestration of pollutants are accompanied by a range of more general mechanisms for protein survey and repair (e.g., affected by ubiquitin or by heat shock proteins).

With respect to bioindication, the multitude of detoxification mechanisms available and the remarkable overlap between the mechanisms involved in the detoxification of heavy metals and organic pollutants may mean that nematodes frequently acquire multiple resistance against various pollutants simultaneously. The resulting nonspecific population response to pollutants will, in turn, largely blur an individual pollutant’s signal in the nematode community. It seems, therefore, a relevant question whether nematodes exhibit sufficiently distinct responses to different toxic substances to enable a differential bioindication of these substances. We
re-analyze published data from six large field investigations on long-term soil contamination as to the responses of nematodes to the pollutants applied. First, we illustrate the importance of indirect contamination effects on nematode populations and propose a simple method for isolating direct effects from food-chain effects. Then, we compare across experiments the consistency and the specificity in the numerical responses of nematode genera to selected toxicants.

Materials and Methods

Data sources: Due to the destructive and risky nature of toxicological field experiments, few investigations are conducted in which soil contamination is deliberately introduced in a controlled way in the field. However, information can also be drawn from sites that have been contaminated by industrial waste or by accidental pollution. For our analysis, we selected six investigations from the literature that included at least 2 yr duration of contamination and provided quantitative documentation of soil pollution status and nematode community.

The copper field experiment (Korthals et al., 1996) was located near Wageningen, The Netherlands. The dominant soil type is a flemic anthrosol. A continuous crop rotation of maize, potato, and oats was maintained on the site. In 1982, 128 plots of 6 x 11 m were established in eight blocks. A full factorial design of four copper levels (0, 250, 500, and 750 kg Cu/ha) and four pH levels (4.0, 4.7, 5.4, and 6.1 pH-KCl) was implemented in eight replicates randomized in eight blocks. In 1992, 10 yr after the start of the experiment, 30 soil cores (17-mm-diam. x 10-cm-depth) were taken from each of the 128 plots, mixed, and analyzed for pH, total and available copper content, and nematodes.

The sewage sludge experiment (Weiss and Larink, 1998) was conducted at North Nottinghamshire, England. The sewage sludge used (12 t dry matter/ha/yr) was applied in one treatment and sewage sludge with added Ni, Pb, Cd, Cr, Cu, and Zn in a second treatment, while a third treatment was fertilized with 100 kg N/ha/yr. Soil concentrations of heavy metals were monitored regularly. In 1989, 9 yr after the start of the experiment, 10 soil cores (3.9-cm-diam. x 20-cm-depth) were taken in each of four replicate plots of the treatments, bulked, and analyzed for nematodes. The site was contaminated by surface runoff of Cu-Cr-As-based timber preserving liquor from an adjacent timber-treatment plant. In 1989, the contamination was analyzed. In 1991, 50 soil cores (2.5-cm-diam. x 5-cm-depth) were taken from 11 plots on areas with low, medium, high, and extreme contamination. Soil concentrations of Cu, Cr, and As were measured and nematodes enumerated and identified.

The heavy metals experiment (Nagy, 1999) was conducted at Nagyhórcsök, Hungary, on calcareous loamy chernozem with a medium-to-deep humus layer. Various crops, such as maize, carrot, potato, peas, beet root, spinach, and winter wheat, were planted during the experiment. In 1991, a split-plot design with two replicate blocks was established, in which 13 individual heavy metals and microelements were ploughed into the soil at three to four different contamination levels of up to 810 kg/ha. Soil element concentrations were annually monitored. In 1996 and 1997, a total of 56 soil cores (2-cm-diam. x 10-cm-depth) were taken from the controls and from the 15 treatments with the highest contaminations, and nematodes were extracted.

The multiple metal experiment (Georgieva et al., 2002) was conducted at North Nottinghamshire,
United Kingdom, on sandy loam. Barley, Italian ryegrass, sugar beet, white clover, and peas were grown on the site. In 1982, a randomized plot design was established with the following treatments: Application of uncontaminated sludge (100 t dry solids/ha); sewage sludge contaminated artificially with four concentrations of either Ni, Cu, Zn, or combinations of Zn and Ni or Zn and Cu; and a control without any sewage sludge application. Four replicates were established for the uncontaminated sludge and no sludge treatments and two replicates for each heavy metal concentration. In 1986, further additions of naturally contaminated sludge were made to some of the plots. The soils were analyzed regularly for heavy metal content. In 1994, 12 yr after the first and 8 yr after the second treatment, each plot was sampled for nematodes with a bulk sample consisting of 20 cores (1.9-cm-diam. × 20-cm-depth).

The organic pollution investigation (Blakely et al., 2002) was performed in Toledo, OH, in the vicinity of a creosote reservoir. The reservoir was established at the site about 50 yr earlier and since then had been leaking into adjacent soil and groundwater. The soils analyzed were vegetated by mixed deciduous forest with a dense understory of perennial and annual herbs and vines. Thirty intact soil cores (5.1-cm-diam. × 7.6-cm-depth) were taken on each of three sampling occasions in 1998. Phenanthrene, fluoranthene, pyrene, and benzo-a-pyrene concentrations were measured, and nematode community composition was analyzed.

**Data analysis:** The copper field experiment was analyzed separately to serve as a validation control (see Table 2). To illustrate indirect Cu effects mediated via the food chain, we compared the responses of nematode feeding groups to the Cu concentration gradient of the copper field experiment. In a second analysis, we compared nematode responses of the four experiments with multiple heavy metal pollution. To make it possible to compare data across the different experiments, data were normalized as follows: Counts of nematode genera were expressed as relative proportions of their feeding type (DoFT = Dominance within Feeding Type), thereby correcting both for indirect toxicity effects through changes in feeding sources and for differences across investigations in sampling and extraction. We see this as a natural extension from the widely adopted habit of expressing genus importance as dominance, i.e., as relative proportions of the total community. To correct for differences between investigations in heavy metal bioavailability, the concentrations of heavy metals were given a different weight for each investigation and for each feeding type, to provide the best regression coefficient between heavy metals and nematode responses in the combined data of the meta-analysis. The Excel Solver tool was applied to optimize

### Table 2. Data sources and analyses of the six studies used in this investigation, together with information on soil, vegetation, pollutants applied, organic amendments (if applied), years since contamination, and the contaminants considered. Data of the copper field experiment and of the organic pollution investigation were treated separately. Data of the four experiments with multiple heavy metal contamination were combined in a meta-analysis.

<table>
<thead>
<tr>
<th>Name of experiment and reference</th>
<th>Soil and vegetation</th>
<th>Pollutants and organic amendment</th>
<th>Yr</th>
<th>Analyzed contaminants</th>
</tr>
</thead>
<tbody>
<tr>
<td>Separate analysis: Cu effects, food-chain effects</td>
<td>Fimic anthrosol, continuous crop rotation of maize, potato, oat</td>
<td>Combinations of Cu and acidification</td>
<td>10</td>
<td>Cu</td>
</tr>
<tr>
<td>Copper field</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Meta-analysis: Cr, Cu, Ni, and Zn effects across experiments</td>
<td>Luvisol, crop rotation with maize, wheat, oat, sugar beet, potato</td>
<td>Mixture of Ni, Cd, Cr, Cu, Pb, Zn; sewage sludge</td>
<td>9</td>
<td>Cr, Cu, Zn</td>
</tr>
<tr>
<td>Sewage sludge</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Timber preservative</td>
<td>Typic Dystrochrept, pasture with ryegrass, clover</td>
<td>Mixture of As, Cr, Cu</td>
<td>2</td>
<td>Cr, Cu</td>
</tr>
<tr>
<td>Heavy metals</td>
<td>Chernozem, various crops: maize, carrot, potato, peas, beet, spinach, wheat</td>
<td>Al, As, Ba, Cd, Cr, Cu, Hg, Mo, Ni, Pb, Se, Sr, Zn, individually</td>
<td>5</td>
<td>Cr, Cu, Ni, Zn</td>
</tr>
<tr>
<td>Multiple metal</td>
<td>Sandy loam, various crops: barley, ryegrass, beet, clover, pea-se</td>
<td>Cu, Ni, Zn individually and mixtures Ni+Zn, Cu+Zn; sewage sludge</td>
<td>12</td>
<td>Cu, Ni, Zn</td>
</tr>
<tr>
<td>Separate analysis: Comparison of heavy metal effects and effects of organic pollutants</td>
<td>Soil not specified, mixed deciduous forest</td>
<td>Creosote (mixture of organic pollutants)</td>
<td>50</td>
<td>benzo-a-pyrene, five-ring-PAH, fluoranthene, phenanthrene, pyrene</td>
</tr>
<tr>
<td>Organic pollution</td>
<td></td>
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<td></td>
</tr>
</tbody>
</table>

1. (Korthals et al., 1996)
2. (Weiss and Larink, 1991)
3. (Yeates et al., 1995)
4. (Nagy, 1999)
5. (Georgieva et al., 2002)
6. (Blakely et al., 2002)
Table 3. Overview of nematode responses to selected heavy metals and organic toxicants comparing results from the copper field experiment, the joint analysis of experiments with multiple heavy metal contamination, and the organic pollution investigation. Nematode genera and toxic substances are arranged to group similar nematode responses in adjacent cells of the table (two-way joining).

<table>
<thead>
<tr>
<th>Genus</th>
<th>Family</th>
<th>Class</th>
<th>Feeding type</th>
<th>P-classification</th>
<th>Heavy metals and organic toxicants&lt;sup&gt;d&lt;/sup&gt;</th>
<th>Cu (Cu Field Exp.)</th>
<th>Zn (Meta-Analysis)</th>
<th>Cr (Meta-Analysis)</th>
<th>Ni (Meta-Analysis)</th>
<th>Benzpyrene</th>
<th>FreisingER</th>
</tr>
</thead>
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<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>Fluoranthene</td>
<td>Pyrene</td>
<td>Fluoranthene</td>
<td>Cu (Meta-Analysis)</td>
<td>Zn (Meta-Analysis)</td>
<td>Cr (Meta-Analysis)</td>
<td>Ni (Meta-Analysis)</td>
</tr>
<tr>
<td>Aporcelaimidae</td>
<td>Aporcelaimidae</td>
<td>A</td>
<td>PR</td>
<td>5</td>
<td>-0.33</td>
<td>-0.34</td>
<td>-0.34</td>
<td>n.s.</td>
<td>n.s.</td>
<td>n.s.</td>
<td>n.s.</td>
</tr>
<tr>
<td>Ditylenchus</td>
<td>Anguinidae</td>
<td>S</td>
<td>HF</td>
<td>2</td>
<td>n.s.</td>
<td>n.s.</td>
<td>-0.22</td>
<td>-0.39</td>
<td>n.s.</td>
<td>n.s.</td>
<td>n.s.</td>
</tr>
<tr>
<td>Bitylenchus</td>
<td>Dolichodoridae</td>
<td>S</td>
<td>PP</td>
<td>3</td>
<td>-0.25</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Bursia</td>
<td>Tylenchidae</td>
<td>S</td>
<td>PP</td>
<td>2</td>
<td>-0.35</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Meanhobaldites</td>
<td>Rhabditidae</td>
<td>S</td>
<td>BF</td>
<td>1</td>
<td>-0.35</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Acrobeloides</td>
<td>Cephalobidae</td>
<td>S</td>
<td>BF</td>
<td>2</td>
<td>-0.46</td>
<td>-0.25</td>
<td>-0.07</td>
<td>-0.93</td>
<td>+0.49</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Molinimus</td>
<td>Dolichodoridae</td>
<td>S</td>
<td>PP</td>
<td>3</td>
<td>-0.48</td>
<td>+0.85</td>
<td>+0.74</td>
<td>+0.96</td>
<td>+0.79</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Cordicithus</td>
<td>Cephalobidae</td>
<td>S</td>
<td>BF</td>
<td>2</td>
<td>-0.35</td>
<td>-0.49</td>
<td>-0.32</td>
<td>-0.93</td>
<td>+0.09</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Monhysterus</td>
<td>Monhysteridae</td>
<td>A</td>
<td>BF</td>
<td>1</td>
<td>-0.26</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Tylencholaimus</td>
<td>Tylencholaimidae</td>
<td>A</td>
<td>HF</td>
<td>4</td>
<td>-0.33</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>+1.00</td>
</tr>
<tr>
<td>Trichodorus</td>
<td>Trichodoridae</td>
<td>A</td>
<td>PP</td>
<td>4</td>
<td>-0.55</td>
<td>-0.67</td>
<td>-0.00</td>
<td>-0.31</td>
<td>+0.41</td>
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</tr>
<tr>
<td>Acrobelis</td>
<td>Cephalobidae</td>
<td>S</td>
<td>BF</td>
<td>2</td>
<td>-0.78</td>
<td>-0.64</td>
<td>-0.77</td>
<td>-0.85</td>
<td>+0.25</td>
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<td></td>
</tr>
<tr>
<td>Dipthirhopora</td>
<td>Dipthirhoporidae</td>
<td>A</td>
<td>HF</td>
<td>3</td>
<td>-0.65</td>
<td>-0.45</td>
<td>-0.69</td>
<td>-0.74</td>
<td>+0.04</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Aphelenchus</td>
<td>Aphelenchidae</td>
<td>S</td>
<td>HF</td>
<td>2</td>
<td>-0.42</td>
<td>-0.62</td>
<td>-0.63</td>
<td>-0.44</td>
<td>+0.74</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Alaimus</td>
<td>Alaimidae</td>
<td>A</td>
<td>BF</td>
<td>4</td>
<td>-0.73</td>
<td>-0.64</td>
<td>-0.04</td>
<td>+0.46</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Prismatolaimus</td>
<td>Prismatolaimidae</td>
<td>A</td>
<td>BF</td>
<td>3</td>
<td>n.s.</td>
<td>n.s.</td>
<td>n.s.</td>
<td>-0.24</td>
<td>+0.06</td>
<td>-0.67</td>
<td>-0.10</td>
</tr>
<tr>
<td>Bastiania</td>
<td>Bastianidae</td>
<td>A</td>
<td>BF</td>
<td>3</td>
<td>-0.53</td>
<td>-0.71</td>
<td>-0.25</td>
<td></td>
<td></td>
<td></td>
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</tr>
<tr>
<td>Divolaimus</td>
<td>Discolaimidae</td>
<td>A</td>
<td>PR</td>
<td>5</td>
<td>-0.53</td>
<td>-0.58</td>
<td>-1.00</td>
<td>-0.47</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Response type 1

| Coslenchus   | Tylenchidae | S | PP           | 2                | +0.29       | -0.51   | +0.06       | -0.55              |                   |                   |                   |            |
| Necrophilus  | Cephalobidae | S | BF           | 2                | -0.03       | +0.36   | +0.39       | -0.89              | -0.58              |                   |                   |            |
| Semura       | Semuridae | S | PR           | 2                | +0.40       | -0.66   | -0.47       | -0.68              |                   |                   |                   |            |
| Pratylenchus | Pratylenchidae | S | PP           | 3                | +0.60       | +0.48   | -0.57       | -0.94              | -0.58              |                   |                   |            |

Response type 2

| Pseudhalenchus | Anguinidae | S | HF           | 2                | +0.35       | +0.32   | +0.23       | -0.19              |                   |                   |                   |            |
| Protohobaldites | Rhabditidae | S | BF           | 1                | +0.34       |         |             |                   |                   |                   |                   |            |
| Chloroporus   | Cephalobidae | S | BF           | 2                | +0.75       | +0.41   | +0.16       | -0.70              | -0.01              |                   |                   |            |
| Rhabditis     | Rhabditidae | S | BF           | 1                | n.s.       | n.s.    | n.s.       | +0.40              | +0.24              | n.s.               | n.s.               | n.s.       |
| Aphelenchoides | Aphelenchoididae | S | HF           | 2                | +0.35       | +0.52   | +0.61       | +0.59              | +0.02              | -0.25               | -0.23               | -0.25       |
| Paratylenchus | Paratylenchidae | S | PP           | 2                | n.s.       | n.s.    | n.s.       | +0.13              | +0.71              | -0.42               | +0.43               | -0.26       |
| Criconemoides | Criconemidae | S | PP           | 3                | -0.08       | +0.66   | -0.07       |                   |                   |                   |                   |            |
| Tylenchus     | Tylenchidae | S | PP           | 2                | -0.28       | -0.80   | -0.86       | -0.74              | -0.84              |                   |                   |            |
| Heterocephalobus | Cephalobidae | S | BF           | 2                | -0.96       | 1.00    | +0.81       | +0.88              |                   |                   |                   |            |
| Cephalobus    | Cephalobidae | S | BF           | 2                | -0.14       | +0.35   | -0.00       | +0.80              | -0.17              |                   |                   |            |
| Plectus       | Plectidae | A | BF           | 2                | +0.05       | -0.33   | -0.46       | +0.43              | +0.56              |                   |                   |            |
| Hopolaimus    | Alaimidae | S | PR           | 3                | n.s.       | n.s.    | n.s.       | +0.31              |                  |                   |                   |            |

Response type 3

| Cerceolaimus | Aporcelaimidae | A | PR           | 5                | -0.33       | -0.34   | -0.34       | n.s.               | n.s.               | n.s.               | n.s.               | n.s.       |

<sup>a</sup> Three major response types emerge from this grouping: Type 1: Cu/Zn sensitivity, Type 2: Ni sensitivity, and Type 3: specific tolerances.

<sup>b</sup> A: Adenophorea, S: Secernentea.

<sup>c</sup> BF: bacterial feeders, HF: hyphal feeders, PP: plant feeders, PR: predators.

<sup>d</sup> Spearman rank correlation coefficients. Negative correlations are in italics. Significant correlations are marked in bold; p-values were corrected for multiple testing. n.s.: not significant. Empty cells denote absence or very low abundance of a taxon.
regression coefficients. The values of the relative weights were handled conservatively and were kept within 0.1 and 10. Spearman rank correlations were calculated for each nematode genus and each of the four heavy metals, Cu, Zn, Ni, and Cr, that were shared between at least two experiments. The truncated probability method (TPM; Zaykin et al., 2002; Neuhäuser, 2004) was applied to account for replicate testing and to keep cumulative error below 0.05. Finally, the regression coefficients obtained for heavy metals were compared to those reported from the organic pollution investigation. Regression coefficients were rearranged in a table by means of two-way joining to reflect similarities of the effects of toxicants and similarities of the responses of nematodes (Table 3). The Statistica software package (StatSoft Inc., Tulsa, OK) was used for all statistical analyses.

**Results**

**Differences between trophic channels:** In the copper field experiment, there was a general and strong decrease of the nematode population density at higher Cu concentrations (Fig. 1). However, there were marked differences between feeding groups. Plant feeders proved the most strongly affected group, exhibiting a marked decline at Cu concentrations higher than 1 ppm. Bacterial feeders remained comparatively less affected up to a Cu concentration of around 2 ppm, while rapidly degrading at higher concentrations. In contrast, fungal feeders increased in abundance with increasing Cu concentrations up to 4 ppm and declined moderately at higher concentrations. We consider it unlikely that nematodes with a hyphal-feeding habit are generally resistant to Cu while nematodes with a plant-feeding habit are generally susceptible to Cu. Instead, we assume that the abundances of nematode trophic groups followed the responses of their feeding sources to the contamination. Specifically, plants showed a dramatic inhibition of growth at higher Cu concentrations, while fungal growth may have been partly fostered by the soil acidification brought about by the combined pH and Cu treatment in the experiment.

**Comparison across investigations:** Of the 70 nematode taxa registered in total, 34 exhibited significant correlations with at least one of the toxic substances analyzed (Table 3). Correlations of nematode genera with Cu were coherent between the copper field experiment and the other heavy metal experiments. None of the significant correlations with Cu was contradictory between experiments; however, the ensemble of multiple heavy metal experiments showed less significant correlations, as is to be expected by the higher variation of data encompassed. The overall comparison of results from the copper field experiment, the four experiments with multiple heavy metals combined, and the organic pollution experiment reveals three major response types of nematodes: (i) Cu/Zn sensitivity, (ii) Ni sensitivity, and (iii) specific tolerances. Type I genera exhibited negative correlations with Cu and in many cases also with Zn. Possibly, genera of this response type also tend to be susceptible to organic pollutants, as indicated to some extent by *Ditylenchus*. The Aporcelaimidae were allocated to response type 1 due to their susceptibility to fluoranthene, pyrene, and phenanthrene. At the same time, type 1 nematodes were less affected by Cr and tended to be correlated positively with Ni, albeit not significantly in most cases. Practically all Adenophorea belonged to this response type. In direct contrast, response type 2 encompassed Secernentea showing negative correlations with Ni. The genera of response type 3 were characterized by positive correlations with pollutants. They showed a stronger differentiation of responses. Four genera were particularly tolerant to Cu, three to Zn, and three to Cr; *Plectus* was tolerant to Ni and the Hoplolaimidae were tolerant to benzo-a-pyrene and the five-ring-PAH. This response type was represented almost exclusively by Secernentea, the only exception being *Plectus*. The error probability for the positive correlation between *Plectus* and Ni is at the margin of the cut-off criterion for pairwise correlations ($P_{\text{pairwise}} < 0.005$), which is required to keep the cumulative error below the significance threshold. The result for *Plectus* is therefore subject to the cumulative error probability, which amounted to 0.045.

**Discussion**

Our results suggest that a broad spectrum of nematode genera show a common response pattern to the
toxicants analyzed, while some genera acquired specific and differential tolerances toward particular substances. It must be emphasized that dominance within feeding type (DoFT), used to quantify nematode importance in this investigation, does not offer an absolute measure of a nematode’s potential to cope with a given impact. Instead, DoFT provides a relative measure of sensitivity as compared to the responses of the other nematodes from the same feeding type. We interpret the abundances of nematode feeding groups in the copper field experiment to be strongly influenced by responses of feeding sources to the copper contamination. Such food-chain effects distort the proportions of the individual genera within the nematode community, depending on the feeding type of each genus. Therefore, both nematode abundances and dominances reflect the superimposition of direct effects of toxicants on nematodes and indirect effects through the responses of feeding sources to the toxicants. In contrast, DoFT isolates the direct effects from the superimposition by treating the food chains separately. Analyses that we performed using nematode abundances and dominances did not yield consistent results, and we found DoFT by far the better choice.

Interpretation of the results is partially impeded by the fact that a number of nematode genera were present in only one of the datasets, which prevents a reliable generalization of their behavior. The sparse data available on long-term effects of organic toxicants means that only the most reserved of conclusions can be drawn. Generally, pollution can induce tolerance in nematodes through selection of tolerant strains within the population (Millward and Grant, 2000). Thus, our long-term analysis reflects the potential of nematode genera to adapt themselves to pollution in the long term, and this potential may not be representative of other investigations performed shortly after a contamination event. Given these limitations, our analysis revealed several consistent patterns. Consistency was explicitly visible by the lack of contradiction between the two separate analyses for Cu. Consistency was implicitly ensured by the meta-analysis of the four datasets with multiple heavy metals. If a nematode genus responded inconsistently between datasets, then no significant correlation would emerge across datasets.

Contrary to our expectations derived from the overlap of molecular mechanisms involved in the detoxification of heavy metals and organic toxicants, nematodes generally did not exhibit multiple resistance against various pollutants simultaneously. Instead, the majority of nematode taxa exhibited multiple sensitivity. Overall, tolerance of nematodes toward toxicants (16 significant positive correlations) was half as frequent as sensitivity (35 significant negative correlations), which illustrates that tolerance, not sensitivity, represented the special case. As a consequence, bioindication of specific toxicants often will be more powerful if based on tolerant nematodes rather than on sensitive nematodes. The relative persistence of tolerant nematodes, in turn, qualifies them better as sentinels (Beeby, 2001). Tolerance to heavy metals and to organic toxicants was found almost exclusively in Secernentea. However, not all Secernentea exhibited such a behavior. Some Secernentea were particularly sensitive to Ni. Many other Secernentea resembled the largely unspecific Adenophorea. This differentiation within the Secernentea will certainly obscure any relationship between the Secernentea/Adenophorea ratio and a toxicant. No coherent response pattern was visible on the family level. The Cephalobidae presented an extreme example where all kinds of different behaviors were shown by its genera. It is known that, even within the same genus, different species can exhibit contrasting responses to heavy metals (Sturhan, 1986). Therefore, it seems possible that tolerance to toxic substances, or the potential for selective adaptation, evolved repeatedly, independently, and differentially within the Secernentea based on a general pre-adaptation of this group. This implies that relatively little information can be drawn from the taxonomic relationships between nematodes about their potential as bioindicators of toxicants. Empirical knowledge of their responses to toxic substances is indispensable. From our analysis, we conclude that Chilophaeus and Pratylenchus are good candidates for substance-specific bioindication of Cu, Pratylenchus and Criconemoides of Cr, and Tylenchus and Cephalobus of Zn, as seen from their exclusive and positive correlations with these metals. Good candidates for bioindication of organic toxicants possibly can be found in the family Hoplolaimidae.

**Literature Cited**


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