

1 **Plant-Parasitic Nematodes (PPNs) Associated**
2 **with Sugarcane (*Saccharum officinarum*) in**
3 **Selected Areas of Northern Cebu, Philippines**

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ABSTRACT

Plant-parasitic nematodes (PPNs) are reported to be associated with sugarcane and are considered detrimental to the crop, resulting to low quality produce and low yielding ability. This study aimed at identifying the plant-parasitic nematode genera associated with sugarcane, as well as its prevalence in the selected sampling sites of Northern Cebu, Philippines. Root and soil samples were collected from Bogu City (Phil99-1793 & VMC84-947), Medellin (VMC84-947 & Mauritius), and San Remigio (VMC84-947 & Mauritius). Nematodes from roots and soil rhizosphere were extracted following the standard methods and were identified up to genus level based on morphological characteristics. A total of eleven (11) plant-parasitic nematode genera (*Criconema*, *Ditylenchus*, *Filenchus*, *Helicotylenchus*, *Hoplolaimus*, *Meloidogyne*, *Pratylenchus*, *Rotylenchulus*, *Rotylenchus*, *Tylenchus*, and *Xiphinema*) which belong to nine different families were identified. Among these identified nematode genera, *Meloidogyne* were the most abundant in sugarcane roots, and *Helicotylenchus* in soil rhizosphere. However, no nematode genera recovered from the roots were considered prevalent as their frequency and abundance were too low, which occurred only in some localities. In roots, *Helicotylenchus* was the predominant nematode genera due to its high frequency of occurrence in all sampling sites and high abundance in two sampling sites. Other identified nematode genera were less prevalent as observed in their relatively low abundance and frequency of occurrence. These results may advance farmers' knowledge on plant-parasitic nematodes associated with sugarcane in their locality and assess appropriate control measures in mitigating them.

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Keywords: Plant-parasitic nematodes, morphological characteristics, abundance, frequency, prevalence

27 1. INTRODUCTION

28

29 Sugarcane (*Saccharum officinarum* L.) is a tall, perennial grass that belongs to the
30 *Poaceae/Gramineae* family and is considered as one of the most important plantation crops
31 grown in tropical and subtropical regions of the world, including the Philippines. It is
32 propagated primarily by means of cuttings. Seed cane is spaced at 1.4 to 1.8 meters (4.5 to 6
33 feet) apart at densities of 10,000 to 25,000 per hectare (Yamane, 2023). Aside from sugar as
34 its main product, it also produces extensive by-products such as biofuels, ethanol, straw,
35 bagasse (cane fibers), molasses, and rum (Yamane, 2023).

36 According to Manzoor et al. (2023), approximately 28.3 million hectares are planted with
37 sugarcane in more than 90 countries with a total production of about 1.69 billion tons
38 worldwide. Brazil, India, and the European Union are considered as the top sugarcane-
39 producing countries which contribute to 40% of the total worldwide production (Anandh, 2018).
40 In the Philippines, out of the approximately 30M ha. of the country's total land area, sugarcane
41 production is planted to about 422, 500 ha. with average annual yield of 57 tons/ha.,
42 contributing an annual revenue of about PhP 70 billion (Tobias, 2020). Based on the Sugar
43 Regulatory Administration's (SRA) sugar policy for crop year 2020-2021, 93% of the total
44 output would go to domestic markets and 7% to the US (USDA, 2021).

45 With favorable conditions, growing cultivated areas, and increasing productivity rates amidst
46 the pandemic, infestations of pathogenic diseases in sugarcane have been a constant problem
47 which is prevalently increasing. Extreme consideration of its security is needed because it is
48 considered as a top export commodity. One major pathogen causing defects to sugarcane are
49 the nematodes that feed mostly on the roots of the plant. Nematodes are small worms that
50 cause serious damage to sugarcane (Blair et al., 1999). Symptoms of nematodes in sugarcane
51 depend on the genus and include reduced root and shoot mass, reduced tillering, stunted,
52 sparse and distorted roots, and less obvious symptoms such as necrosis and discoloration of
53 roots (Ramouthar & Bhuiyan, 2018). Tillering is reduced and the height of the canopy may
54 vary considerably because shoot elongation is affected. Below ground, swellings and galls are
55 relatively easy to see on primary roots and the fine root system (Stirling, 2023).

56 Sugarcane in South Africa hosts more than 90 species of 28 genera of plant-parasitic
57 nematodes (PPNs). The more common species include *Helicotylenchus dihystra*,
58 *Pratylenchus zaei*, *Xiphinema elongatum* and on sandy soils, *Meloidogyne javanica*, which
59 caused an estimated 9% loss of production (Berry et al., 2017). Several studies have
60 uncovered PPN diversity in sugarcane fields in other countries such as Mauritius (Spaull,
61 1990), India (Sundararaj & Mehta, 1993), Kenya (Chirchir et al., 2008), South Africa (Berry et
62 al., 2017), and Brazil (Noronha et al., 2017), revealing the most common PPN genera
63 associated with sugarcane as *Meloidogyne*, *Pratylenchus*, *Tylenchorhynchus*, *Rotylenchulus*,
64 and *Helicotylenchus* (Singh et al., 2020).

65 In the Philippines, there are few studies on plant-parasitic nematodes (PPNs) associated with
66 sugarcane. In Negros Occidental and Panay Island, *Tylenchorhynchus* was the most
67 predominant nematode genera associated with sugarcane (Husmillo et al., 1983). Knowledge
68 on the identification and prevalence of PPNS associated with sugarcane are important in order
69 to know how they cause reduction in yield and to assess which appropriate control measures
70 are effective in mitigating them. This study aimed to identify the PPN genera associated with
71 sugarcane in Northern Cebu, Philippines through morphological characteristics and to
72 determine the prevalent PPNS in the area based on population density, frequency, and
73 abundance.

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75 **2. MATERIALS AND METHODS**

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77 **2.1 Sampling Sites and Location of the Study**

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79 Root and soil samples were collected from three sugarcane production areas in Northern Cebu
 80 (Bogo, Medellin, and San Remigio) on February 2023 (Figure 1). In each municipality, three
 81 sugarcane fields were randomly sampled and soil characteristics (texture, pH, temperature,
 82 moisture), sugarcane varieties grown, and management practices employed were recorded
 83 (Table 1). In each field, root samples were taken from five sample plants that showed
 84 symptoms of nematode infection (i.e., stunted growth, leaf discoloration, chlorosis/yellowing,
 85 necrosis, galling, root lesions, and shortened roots) which were collected by excavation using
 86 a digging bar. The roots were then separated from the cane and were placed in plastic bags,
 87 labeled, and stored properly. Approximately 200 grams of soil samples were also collected
 88 from the soil rhizosphere of each plant, up to 15 cm depth using hand trowel and bolo. These
 89 soil samples were placed in plastic bags, labeled, and stored properly. The samples were
 90 brought to the Crop Protection laboratory of Cebu Technological University – Barili Campus
 91 for extraction and identification.
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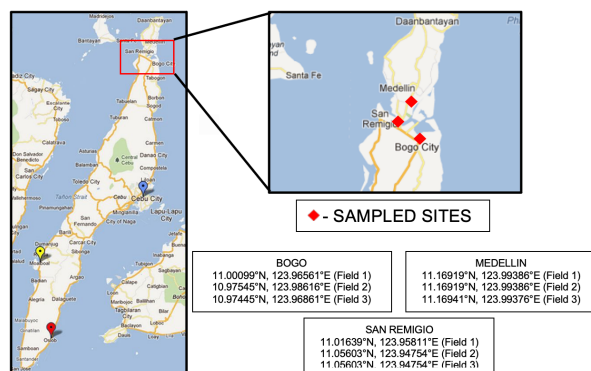


Fig 1. Sampling sites in Northern Cebu, Philippines

Table 1. Soil texture, pH, temperature, moisture, sugarcane varieties grown, and practices employed in the sampling sites

Sampling Sites	Soil Texture	Soil pH	Temperature	Soil Moisture	Variety	Practices after Harvest
Bogo	Clayey	6.8	23	Dry	Phil99-1793, VMC 84-947	Trash Burning / Mulching
Medellin	Loamy Clay	6.6	23	Normal	VMC 84-947, Mauritius	Trash Burning / Mulching
San Remigio	Sandy Loam	4.5	23.8	Normal	VMC 84-947, Mauritius	Trash Burning / Mulching

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104 **2.2 Extraction of Nematodes from Plant Roots**

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106 Nematode extraction was done at the Crop Protection laboratory of the Department of
107 Agriculture and Food Science, Cebu Technological University – Barili Campus. Plant root
108 samples were washed with running water and were cut into small pieces, approximately up to
109 1 cm using a pair of scissors. Then, 5g of cut roots were macerated for about six seconds
110 using a blender. The macerated roots were placed into a wire mesh lined with tissue paper
111 and were placed over a bowl following a modified Baermann tray method. Water was added
112 to the tray just enough to touch the bottom part of the mesh to get a clear suspension. Then,
113 the set-up was incubated for 48 hrs. and application of water mists were done whenever
114 necessary to avoid drying up.

115

116 **2.3 Extraction of Nematodes from Soil**

117

118 Soil samples were mixed thoroughly and 200 mL representative were processed using a
119 combination of decanting and sieving (using a 212 μm and 53 μm aperture size) methods
120 followed by incubation in modified Baermann tray method. The soil was placed in a wire mesh
121 lined with tissue paper which were placed over into a dish water and left at room temperature
122 for 48 hrs.

123

124 **2.4 Preservation of Nematodes**

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126 After the specified extraction time, nematode suspensions were collected into test tubes and
127 were allowed to settle for 12 hrs. The supernatant was sucked using a pipette to reduce the
128 volume of suspension. Nematodes were killed and fixed by adding hot 4% formalin (65°C-
129 70°C) into the nematode suspensions. Preserved nematodes were placed in the refrigerator
130 until identification and assessment of population density.

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132 **2.4 Identification of Plant-Parasitic Nematodes**

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134 The extracted nematodes from soil and root samples were manually picked using a picking
135 needle from the clear suspension under a stereomicroscope and were mounted to glass slide
136 with a drop of water, covered with cover slip and sealed with colorless nail polish to prevent
137 drying up of mounts. Plant-parasitic nematodes were identified to genus level following keys
138 and references of Siddiqi (2006), Mekete et al. (2012), and others.

139

140 **2.4 Assessment of Nematode Population Density and Prevalence**

141

142 Population densities of plant-parasitic genera from root and soil samples per plant per
143 sampling site were determined in three 5 ml aliquots in a counting dish under a
144 stereomicroscope (Olympus, Germany). Nematode counts were expressed as 5g/roots and
145 per 200 ml soil.

146

147 The prevalence of plant-parasitic nematode genera in roots and soil were determined
148 using the relationship between abundance index (AI) and frequency (F) of occurrence
149 computed using the formula:

150

$$\text{Abundance (AI)} = \log \frac{\text{total number of nematodes per gram roots or}}{\text{L soil in all positive samples}} \frac{\text{number of positive samples}}{\text{number of positive samples}}$$

151

$$\text{Frequency (F)} = \frac{\text{number of samples positive with nematodes}}{\text{total number of samples}} \times 100$$

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154 Based on the limits established by Fortuner and Merny (1973), a nematode was regarded as
 155 abundant if abundance ≥ 1.3 (20 individuals/g of roots) and abundance ≥ 2.3 (200 individuals/L
 156 of soil) in roots and soil, respectively. A nematode was regarded as frequent in the soil or roots
 157 when observed in at least 30% of the samples.

158

159 **3. RESULTS AND DISCUSSION**

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161 **3.1 Plant-Parasitic Nematode Genera Associated with Sugarcane**

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163 Various plant-parasitic nematode (PPN) genera were found from the root and soil samples of
 164 sugarcane in the selected sites of Northern Cebu, Philippines (Table 2). A total of eleven (11)
 165 genera of PPNs belonging to seven (7) different families were found associated with
 166 sugarcane in the surveyed fields. PPNs found include *Criconema* (*Criconematidae*),
 167 *Ditylenchus* (*Anguinidae*), *Filenchus* (*Tylenchidae*), *Helicotylenchus* (*Hoplolaimidae*),
 168 *Hoplolaimus* (*Hoplolaimidae*), *Meloidogyne* (*Heteroderidae*), *Pratylenchus* (*Pratylenchidae*),
 169 *Rotylenchulus* (*Hoplolaimidae*), *Rotylenchus* (*Hoplolaimidae*), *Tylenchus* (*Tylenchidae*), and
 170 *Xiphinema* (*Longidoridae*). Of these genera, *Filenchus*, *Meloidogyne*, *Helicotylenchus*, and
 171 *Pratylenchus* were found associated with roots of sugarcane plants and all genera were
 172 present in the soil rhizosphere except *Meloidogyne* and *Filenchus*.
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175 The nematode genera identified in this study were also reported in previous works conducted
 176 in the Philippines as well as in other countries. Davide (1988) reported four (4) major nematode
 177 genera associated with sugarcane in the country. Davide (1988) observed that *Meloidogyne*
 178 *incognita* bannered the species that were present in the collected samples of sugarcane-
 179 growing areas in the country, followed by *Tylenchorhynchus martini*, *Rotylenchulus reniformis*,
 180 and *Pratylenchus zaeae*. In another study, *Tylenchorhynchus*, *Helicotylenchus*, *Pratylenchus*,
 181 and *Xiphinema* were observed to be predominant plant-parasitic nematode genera infecting
 182 sugarcane crops in Negros and Panay Island (Husmillo et al., 1983). These reported
 183 nematode genera were also found to occur in sugarcane crops in Northern Cebu with the
 184 exception of *Tylenchorhynchus*.

185

186 In Kilimanjaro region of Tanzania, Singh et al. (2020) revealed the presence of seven plant-
 187 parasitic nematode (PPN) genera; *Helicotylenchus*, *Hemicyliophora*, *Pratylenchus*,
 188 *Rotylenchulus*, *Scutellonema*, *Meloidogyne*, and *Tylenchorhynchus* after they performed
 189 morphological and molecular analyses of PPNs from 12 sugarcane plantation sites of
 190 Tanganyika Planting Company (TPC). Four of these nematode genera (*Helicotylenchus*,
 191 *Meloidogyne*, *Pratylenchus*, and *Rotylenchulus*) were also found to occur in the sugarcane
 192 fields of Northern Cebu, Philippines.

193

194 In Paraná, Brazil, Martinha et al. (2022) reported that *Pratylenchus*, *Meloidogyne*,
 195 *Helicotylenchus*, *Xiphinema*, *Hoplolaimus*, *Tylenchus*, and *Ditylenchus* were some of
 196 the nematode genera that caused damages to the sugarcane-growing areas. These nematode
 197 genera were also observed in the present study, suggesting that these nematodes are
 198 commonly associated with sugarcane production.

199

200 **Table 2. Soil texture, pH, temperature, moisture, sugarcane varieties grown, and**
 201 **practices employed in the sampling sites**

Roots		Rhizosphere	
Family	Genus	Family	Genus
<i>Heteroderidae</i>	<i>Meloidogyne</i>	<i>Anguinidae</i>	<i>Ditylenchus</i>
<i>Hoplolaimidae</i>	<i>Helicotylenchus</i>	<i>Criconimatidae</i>	<i>Criconema</i>

<i>Pratylenchidae</i>	<i>Pratylenchus</i>	<i>Hoplolaimidae</i>	<i>Hoplolaimus</i> <i>Helicotylenchus</i> <i>Rotylenchus</i> <i>Rotylenchulus</i>
<i>Tylenchide</i>	<i>Filenchus</i>	<i>Longidoridae</i> <i>Pratylenchidae</i> <i>Tylenchidae</i>	<i>Xiphinema</i> <i>Pratylenchus</i> <i>Tylenchus</i>

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3.2 Genus Identification

205 Morphological identification yielded 11 genera of plant-parasitic nematodes (PPNs)
206 associated with sugarcane in Northern Cebu, Philippines. The important diagnostic characters
207 that aid in the genus identification are herein presented:

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Genus: *Criconema* (Mekete et al., 2012) (Figure 2)

211

Annuli smooth or variously ornamented, annuli of labial region smooth; labial region usually
212 with six pseudolips wide and clearly set off the next body annulus; tail conoid-pointed to
213 bluntly-rounded.



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Fig 2. Genus: *Criconema* (A, 10x) whole body; (B, 100x) anterior; and (C, 100x) posterior

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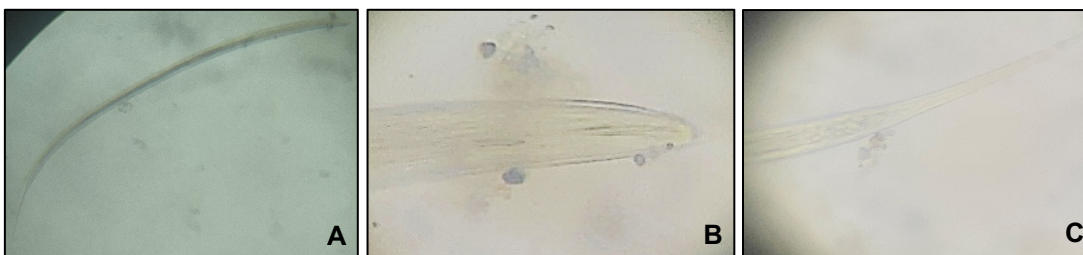
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Genus: *Ditylenchus* (Siddiqi, 2000) (Figure 3)

219

Body usually under 1.5 mm long, not curving strongly when relaxed; mature adults slender;
220 tails elongate-conoid to subcylindrical or filiform.

220



221

Fig 3. Genus: *Ditylenchus* (A, 10x) whole body; (B, 100x) anterior; and (C, 100x) posterior

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Genus: *Filenchus* (Siddiqi, 2000) (Figure 4)

228

Body small to moderately large (0.3-1.3 mm), straight to arcuate when relaxed. Cephalic

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region broadly-rounded or conoid-rounded; tails generally filiform and straight, may be

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elongate-conoid and slightly ventrally arcuate but not hooked.

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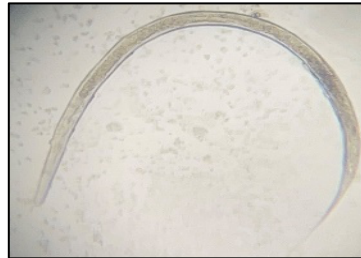


Fig 4. Genus: *Filenchus* whole body (40x)

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Genus: *Helicotylenchus* (Steiner, 1949) (Figure 5)

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Spirally-colloid or rarely arcuate; stylet robust; tail asymmetrical or hemispherical, short,

247

dorsally-convex conoid; vulva located at 70% of the total body length.



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Fig 5. Genus: *Helicotylenchus* (A, 40x) whole body; (B, 100x) female mid-body region showing the vulva (v); and (C, 100x) anterior

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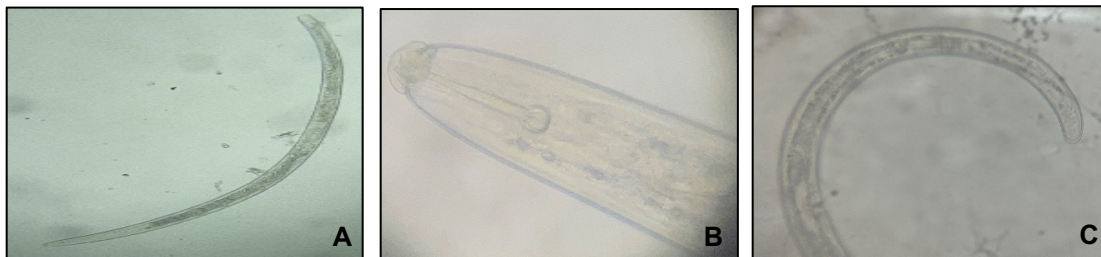
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Genus: *Hoplolaimus* (Daday, 1905; Siddiqi, 2000) (Figure 6)

254

Body arcuate or spiral-shaped; rounded stylet knobs, devoid of tooth-like projections; labial disc lemon-shaped or spindle-shaped; annulated female tail.

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Fig 6. Genus: *Hoplolaimus* (A, 10x) whole body; (B, 100x) anterior; and (C, 40x) posterior

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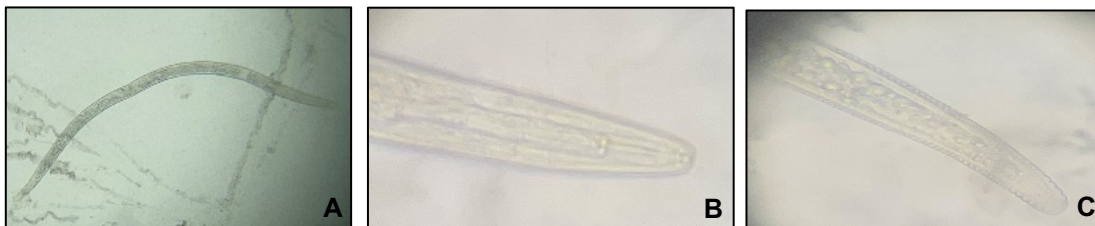
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262 **Genus: *Meloidogyne*** (Eisenback, J. & Triantaphyllou H., 1991) (Figure 7)
 263 Vermiform (male); rounded cephalic region, few annules; rounded tail, blunt tail tip (first stage
 264 juvenile) and vermiform, migratory and infective (second stage juvenile).



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275 **Fig 7. Genus: *Meloidogyne* whole body (40x)**
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278 **Genus: *Pratylenchus*** (Siddiqi, 2000) (Figure 8)
 279 Rounded stylet; anteriorly flat or indented basal knobs; terminus rounded (rarely pointed) tail,
 280 sub-cylindrical to conoid (female); terminus smooth or annulated.



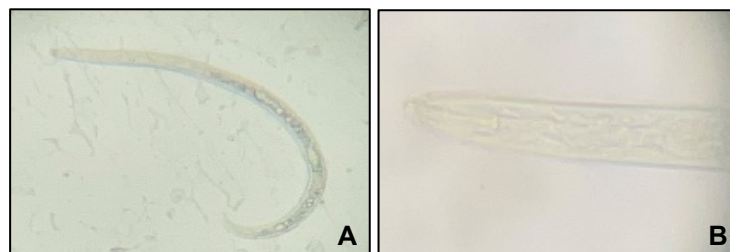
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282 **Fig 8. Genus: *Pratylenchus* (A, 10x) whole body; (B, 100x) anterior; and (C, 100x)**
 283 **posterior**
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286 **Genus: *Rotylenchulus*** (Siddiqi, 2000) (Figure 9)
 287 Small-sized (usually 0.5 mm or under); cephalic region continuous, with or without distinct
 288 annules; male stylet weaker than that of females; tail elongate-conoid; slender spicule in
 289 males.



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291 **Fig 9. Genus: *Rotylenchulus* (A, 10x) whole body; (B, 100x) posterior showing the**
 292 **male spicule; and (C, 100x) anterior**
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298 **Genus: *Rotylenchus*** (Siddiqi, 2000) (Figure 10)
 299 Cephalic region elevated, offset or continuous, round or truncate anteriorly, finely annulated;
 300 labial disc flat; stylet well-developed in both sexes; female tail cylindroid to dorsally convex-
 301 conoid.



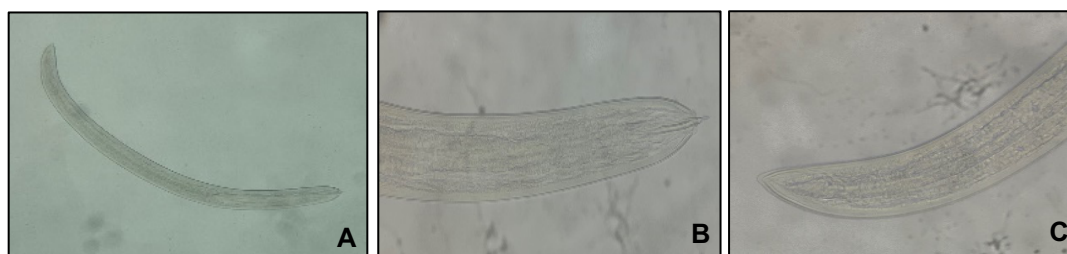
314
 315 **Fig 10. Genus: *Rotylenchus* (A, 10x) whole body; (B, 100x) anterior**

317
 318 **Genus: *Tylenchus*** (Siddiqi, 2000) (Figure 11)
 319 Small to medium-sized (0.4-1.3 mm), ventrally-curved upon relaxation; cephalic region
 320 continuous, annulated; tail ventrally-arcuate; spicules cephalated, arcuate; cloacal lips slightly
 321 raised.



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 323 **Fig 11. Genus: *Tylenchus* (A, 10x) whole body; (B, 100x) posterior; and (C, 100x)**
 324 **anterior**

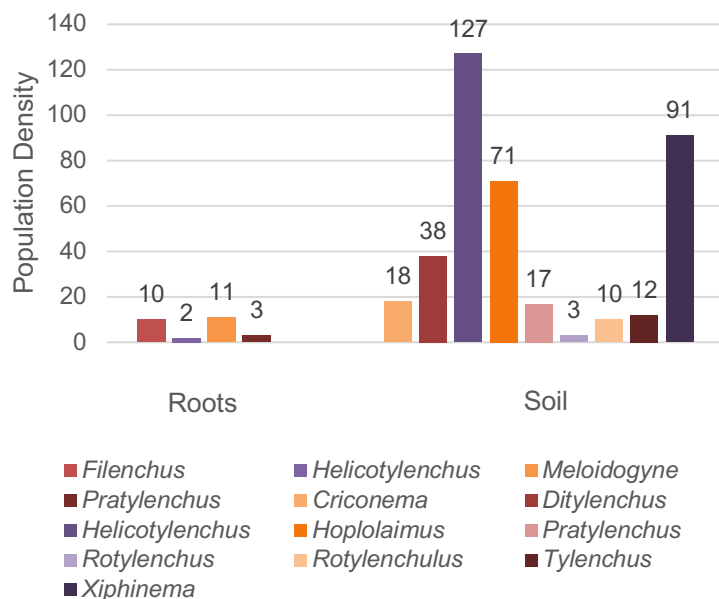
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 327 **Genus: *Xiphinema*** (Mekete et al., 2012; Tarjan, 1964) (Figure 12)
 328 Stylet long, straight, tapering to a long slender point with long extensions; body long, stylet
 329 extensions with sclerotized basal flanges; shorter, conical, and digitate tail.



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 331 **Fig 12. Genus: *Xiphinema* (A, 10x) whole body; (B, 100x) posterior; and (C, 100x)**
 332 **anterior**

333 **3.3 Population Density of Plant-Parasitic Nematode (PPN) Genera Associated with**
 334 **Sugarcane**
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336 Population densities of plant-parasitic nematodes associated with sugarcane in roots and soil
 337 differed (Figure 13). It can be noted that population densities of plant-parasitic nematodes
 338 from root samples were generally low that ranged only from 2 to 11 per 5g roots. *Meloidogyne*
 339 had the highest population density (11/5g roots) followed by *Filenchus* (10/5g roots), while
 340 *Pratylenchus* and *Helicotylenchus* had a population density of 3 per 5g roots and 2 per 5g
 341 roots, respectively. In soil, *Helicotylenchus* (127/200 ml soil) was the predominant nematode
 342 genera followed by *Xiphinema* (91/200 ml soil), and *Hoplolaimus* (71/200 ml soil). The rest of
 343 the nematode genera (*Criconema*, *Ditylenchus*, *Pratylenchus*, *Rotylenchulus*, *Rotylenchus*,
 344 and *Tylenchus*) were recovered at lesser densities that ranged from 3 to 38 per 200 ml soil.
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346
 347 **Fig 13. Population density of plant-parasitic nematode (PPN) genera associated with**
 348 **roots and soil of sugarcane grown in Northern Cebu, Philippines**
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350
 351 Population densities of nematode genera recovered from sugarcane roots were generally low
 352 which can be attributed to several factors. It was noted in this study that the sample plants
 353 were already ratooned twice or thrice and mostly aged at 11-12 months. In Queensland,
 354 Australia, Blair et al. (1999) found out that nematode population densities were significantly
 355 higher in the roots of plant crops than ratoon crops, specifying the *Pratylenchus* genera.
 356 Similarly, in Australia and Burkina Faso, Berry et al. (2007) observed that ratoon crops were
 357 less susceptible than plant crops as its root systems became stronger and more rigid as it
 358 aged. In soil, higher population densities were observed. McSorley (2019) stated that soil is
 359 an excellent and is one of the natural habitats for plant-parasitic nematodes, and 100 cc of soil
 360 may contain several thousands of them. In addition, Ramadhani et al. (2022) validated that
 361 *Helicotylenchus*, a migratory ectoparasite (Rathore, 2017), was one of the important nematode
 362 genera associated with sugarcane. In his research on the phytonematodes associated with
 363 sugarcane, *Helicotylenchus* was one of the major nematode genera recovered based on the

364 highest absolute population, which was also the frontrunner along with *Pratylenchus* in
365 Brathwaite's (1980) study.

366
367 Plant-parasitic nematodes that were found at low densities might be due to the management
368 practices of the farmers from Northern Cebu as they employ the "kaingin" system/trash burning
369 and trash mulching. "Kaingin" was also referred to as 'slash and burn' by farmers in Matalom,
370 Leyte (Lawrence et al., 1997). On the other hand, trash mulching was a common practice in
371 sugarcane production that helped protect the soil as it minimized and/or prevented soil erosion
372 (Samson et al., 2001). After harvest, 90% of the sugarcane farmers in Northern Cebu usually
373 employ trash burning and the remaining 10% employ trash mulching as their means of land
374 clearing and preparation for the next cropping season. In addition, once sugarcane has been
375 established, they do not employ any form of weed management or pest control until harvest.
376 According to Song (2023), a 4-year study of Bastow (2020) revealed that fire reduces soil
377 nematode population by 76% in a semiarid grassland. As a response, Song (2023)
378 investigated the effects of short-term prescribed fire on soil nematode communities and soil
379 properties in an old-field grassland in Northern China which resulted to an increased soil
380 nematode population by 77% and genus richness by 49%. For a more precise control of
381 nematode population, Stirling (2023) recommended crop rotation, use of resistant and tolerant
382 cultivars, nematicide application, and use of natural enemies or biocontrol agents like
383 nematode-trapping fungi, soil microarthropods (nematophagous mites - *Protogamasellus* sp.),
384 and bacteria in the genus *Pasteuria*.

385
386 The predominant nematode genera recovered in this study (*Helicotylenchus*) were considered
387 as ectoparasites (Marmonier, 2019) which live outside the roots of sugarcane. Contributing to
388 higher population densities, eggs of these plant-parasitic nematodes may remain dormant in
389 the soil for a few months until conditions are favorable.

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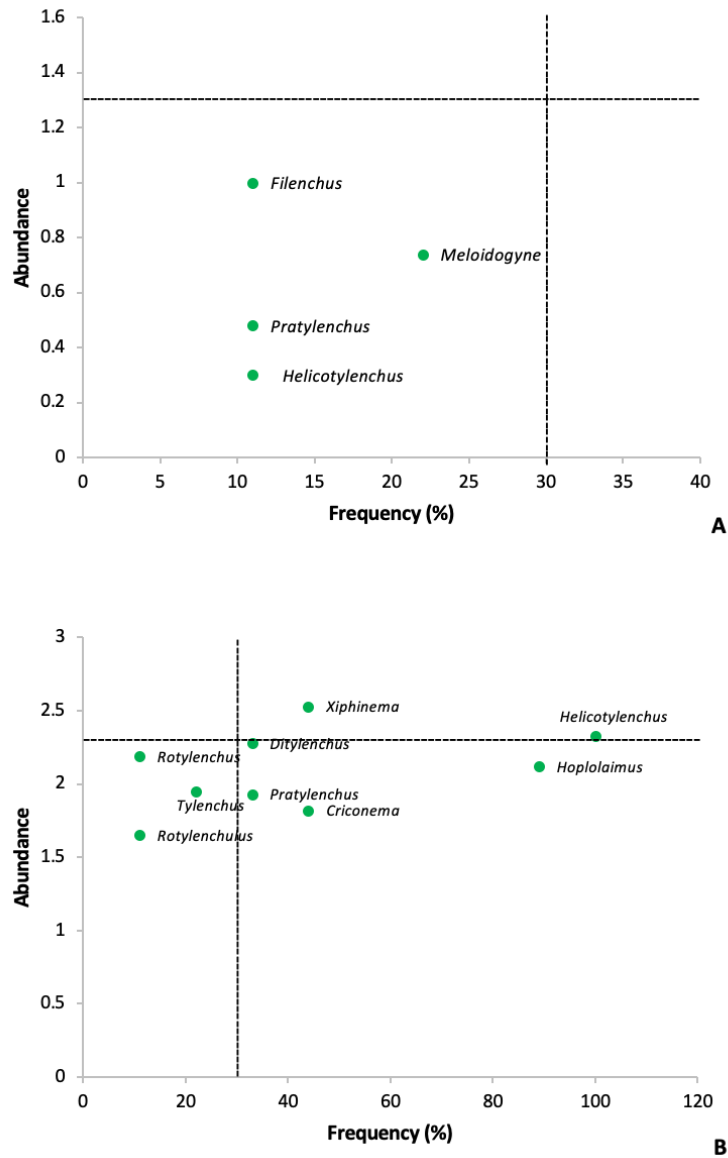
391 **3.4 Prevalence of Plant-Parasitic Nematode (PPN) Genera Associated with Sugarcane**

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393 Based on frequency and abundance, prevalence of plant-parasitic nematodes in roots and soil
394 rhizosphere of sugarcane varied (Figure 14). Based on the limits by Fortuner and Merny
395 (1973), no nematode genera were considered as prevalent in roots as none reached the limit
396 (AI, 1.3 and F,30%). Abundance index of the four nematode genera ranged only from 0.3 to
397 1, while the frequency of occurrence was 11% (*Filenchus*, *Meloidogyne*, and *Pratylenchus*)
398 and 22% (*Helicotylenchus*) which were both below the threshold limits. In soil, *Helicotylenchus*
399 and *Xiphinema* were the most prevalent nematode genera (AI, 2.33 and 2.53, respectively; F,
400 100% and 44%, respectively) as they surpassed the limits (AI,2.3 and F, 30%) set by Fortuner
401 and Merny (1973). Similarly, Ramadhani et al. (2022) also observed that *Helicotylenchus* was
402 the most prevalent nematode genera in Rajawali II, West Java. Román (1961) also revealed
403 that *Helicotylenchus* was the most prevalent nematode genera in the selected fields of
404 sugarcane in Central Soller and San Sebastián, Puerto Rico. *Xiphinema* was also noted to be
405 prevalent in South Africa (Berry et al., 2008) and in Mauritius (Lamberti et al., 1987).
406 *Criconema*, *Ditylenchus*, *Hoplolaimus*, and *Pratylenchus* were also regarded as frequent
407 nematodes of sugarcane soil rhizosphere as their frequencies ranged from 33% to 89%, which
408 were beyond the frequency limit (30%). However, these nematode genera were considered
409 as omnipresent pathogens (Pascual et al., 2017; Fortuner and Merny, 1973) because of their
410 abundance that ranged only from 1.65 to 2.28, which were below the threshold limit (2.3) in
411 soil.

412

413 Among the nematode genera identified in this study, *Filenchus*, *Meloidogyne*, *Rotylenchulus*,
414 *Rotylenchus*, and *Tylenchus* were considered as fortuitous parasites (Pascual et al., 2017;
415 Fortuner and Merny, 1973) which do not cause economic loss due to low frequency and
416 abundance as described by Fortuner and Merny (1973).



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Figure 14. Frequency and abundance of plant-parasitic nematodes (PPNs) in sugarcane roots (A) and soil rhizosphere (B)

3.5 Occurrence and Abundance of Plant-Parasitic Nematode (PPN) Genera Associated with Sugarcane in the Localities

Nematode genera recovered from the different municipalities varied in frequency of occurrence and abundance (Table 3). In Bogó, the observed nematode genera were present in more than 30% of the fields surveyed but their abundance were generally low. In Medellín, *Helicotylenchus*, *Hoplolaimus*, and *Xiphinema* were frequent and abundant. *Ditylenchus* and *Pratylenchus* were also frequent but had low abundance. Other nematode genera were not recovered from the sampled sites. In San Remigio, all nematode genera were frequent except

432 *Filenchus* and *Meloidogyne*. In the sampled site, *Helicotylenchus* and *Ditylenchus* frequently
 433 occurred and were found to be abundant. *Helicotylenchus* was considered as potential
 434 pathogen as it was prevalent in all locations except in Bogo. The rest of the nematode genera
 435 were frequent but had very low abundance index from the different locations, which identified
 436 them as omnipresent nematodes.

437 **Table 3. Occurrence and abundance of plant-parasitic nematodes (PPNs) associated**
 438 **with sugarcane in different sampling locations in Northern Cebu, Philippines**
 439

Genus	Bogo	Medellin	San Remigio
<i>Criconema</i>	+		+
<i>Ditylenchus</i>		+	+++
<i>Filenchus</i>	+		
<i>Helicotylenchus</i>	+	+++	+++
<i>Hoplolaimus</i>	+	+++	+
<i>Meloidogyne</i>	+		
<i>Pratylenchus</i>		+	+
<i>Rotylenchulus</i>			+
<i>Rotylenchus</i>			+
<i>Tylenchus</i>	+		+
<i>Xiphinema</i>	+	+++	+

440 Notes: + = frequent (present in more than 30% of the fields surveyed); ++ = abundant (AI \geq
 441 1.3/5g roots; AI \geq 2.3/L soil); and +++ = frequent and abundant (F \geq 30% and AI \geq 2.3/L
 442 soil).
 443

444 The study of Martinha et al. (2022) stated that the difference between the frequency of
 445 occurrences may be associated with the environmental factors, temperature, water content of
 446 the soil, sugarcane cultivar, and management adopted by producers that interfere with the
 447 density and frequency of genera associated with the sugarcane rhizosphere (Bond et al.,
 448 2000; Matinha 2022).
 449

450 In terms of soil type, the result of this study showing high presence of different nematode
 451 genera in San Remigio (Table 3) may be associated with its sandy loam type of soil (Table 1)
 452 which nematodes prefer more than any other type of soil (Warner, 2009).
 453

454 In correlation with the physicochemical parameters of soil (water holding capacity, moisture
 455 content, pH, electrical conductivity, and temperature), Gade et al. (2018) surveyed the
 456 nematode diversity in sugarcane fields of Aurangabad District in India. It appeared that
 457 temperature and electrical conductivity provided positive correlation to nematode diversity.
 458 Whereas to pH, it had a negative correlation. It was noted in this study that the pH of the three
 459 municipalities varied (Table 1), leading to the idea that as extremophiles (Sapir, 2021), plant-
 460 parasitic nematodes can survive in soils over wide ranges of pH (Warner, 2009).

461 **4. CONCLUSION**

462
 463 Different plant-parasitic nematodes which differ in densities and prevalence are in association
 464 with sugarcane. Identified plant-parasitic nematodes include *Criconema*, *Ditylenchus*,
 465 *Filenchus*, *Helicotylenchus*, *Hoplolaimus*, *Meloidogyne*, *Pratylenchus*, *Rotylenchulus*,

466 *Rotylenchus*, *Tylenchus*, and *Xiphinema*. No nematode genera recovered are considered
467 prevalent in sugarcane roots due to low abundance index and frequency of occurrence.
468 *Helicotylenchus* are the most prevalent plant-parasitic nematodes in soil rhizosphere, which
469 could cause significant yield loss in sugarcane production if not properly managed. These
470 results can further farmers' knowledge on plant-parasitic nematodes associated with
471 sugarcane in their locality.

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