Full Length Article

Phytotoxicity of three Plantago species on germination and seedling growth of hairy beggarticks (Bidens pilosa L.)

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ABSTRACT

Plantago genus has a wide geographical distribution all over the world. It has been widely used in folk medicine for various purposes, where it is used as anti-inflammatory, antimicrobial and antitumor agent. In this research, total phenols, tannins, saponins, flavonoids and alkaloids were determined in Plantago lagopus, Plantago major and Plantago squarrosa. Furthermore, concentrations of 2.5, 5, 7.5 and 10 mg ml$^{-1}$ of both alcoholic and aqueous extracts were prepared to study their phytotoxic effect on the germination and seedling growth of the noxious weed Bidens pilosa. P. major expressed the highest values of total phenolics, tannins and flavonoids. However, P. lagopus has the highest content of alkaloids. The germination of B. pilosa was completely inhibited under treatment of P. lagopus and P. major methanolic extracts at 7.5 mg ml$^{-1}$ and 10 mg ml$^{-1}$, respectively. However, the allelopathic effect of P. lagopus aqueous extract showed a complete inhibition of B. pilosa germination followed by P. major. The germination inhibition of B. pilosa increased with the increase in the extracts concentration. In addition, both radicle and plumule were strongly inhibited under the same treatment. Our results showed a potent allelopathic effect on B. pilosa that could be valorized in managing this noxious weed as an ecofriendly bio-control method. However, other studies are needed for the identification, characterization of the responsible allelochemicals and for the demonstration of their modes of action.

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1. Introduction

Due to the high food requirement of the ever-growing population of the world and the oriented productivity of agriculture, the introduction of synthetic herbicides to control noxious weeds was a favorable way below the threshold limit to reduce the yield loss. Besides improving the crop production, a predatory impact on the environment quality and on human health was generated and increased the number of herbicide resistant
2. Materials and methods

2.1. Preparation of plant materials

P. lagopus, P. major and P. squarrosa aerial parts were harvested at a vegetative stage. The plant tissues were clipped 1 cm above the soil, washed with distilled water and left to dry in room temperature (25 °C) in a shaded place for several days until completely dried. The dried samples were grounded to pass a 1 mm screen, packed in a polyethylene bag and stored in a refrigerator until use.

2.2. Phytochemical analysis

Total phenolics, tannins, alkaloids, flavonoids and saponins of P. lagopus, P. major and P. squarrosa samples were determined spectrophotometrically [14-17].

2.3. Allelopathic bioassay

2.3.1. Weed seed source

The seeds of B. pilosa were collected from the orchards habitat in El-Sharkia Governorate in Egypt, sterilized by 0.3% sodium hypochlorite, rinsed by distilled water, dried on the filter paper in the laboratory at room temperature for 7 days and packed in a paper bag until use [18,19].

2.3.2. Preparation of plant extract

For bioassay tests, aqueous and methanolic stock extracts (10% w/v) were diluted with distilled water to obtain concentrations of 7.5%, 5% and 2.5% (v/v) test extracts. All osmotic concentrations of bioassay solutions were less than 0.1 Mpa and hence not considered a factor affecting germination. The solutions were filtered through double layers of muslin cloth followed by a Whatman No.1 filter paper, the pH values were adjusted to 7 and these were kept in the refrigerator at 4 °C until further use [20].

2.3.3. Germination bioassay

Two layers of Whatman No.1 filter paper were placed in 90 mm diameter glass Petri dishes. In each dish, 25 seeds were placed and 10 ml of each plant extract were added in concentrations of 10, 7.5, 5 and 2.5% (v/v). In case of methanolic extract, seeds were placed after alcohol evaporation and then 10 ml of distilled water were added. A check treatment was assigned with distilled water and left at room temperature. Starting from the first day of the experiment, germinated seeds were counted and removed daily. A seed with 2 mm of radicle was considered germinated. Experiment designed was randomized complete block with three replicates and the experiment was repeated twice. The inhibition percentage was calculated.

2.3.4. Seedling growth bioassay

The seeds of B. pilosa were germinated on filter paper in the dark at room temperature for 2 days. Twenty five germinated seeds were transferred to Petri dishes lined with two layers of Whatman No.1 filter paper and 10 ml of different extracts were added in concentrations of 10, 7.5, 5 and 2.5% (v/v). In addition a check treatment was assigned with distilled water and left at room temperature. Experiment designed was randomized complete block with three replicates and the experiment was repeated twice. The shoot and root lengths of seedlings were measured on the tenth day and growth inhibition for radicle and plumule lengths was calculated.

2.4. Statistical analysis

All values of phytochemistry and allelopathy experiments are the mean of three replicates ± standard error. Data were subjected to ANOVA and the mean values were separated based on the least significant difference (LSD) test.
3. Results

3.1. Phytochemical constituents of the studied Plantago species

The phytochemical constituents of the aerial parts of *P. lagopus*, *P. major* and *P. squarrosa* are presented in Table 1. *P. major* attained the highest significant values of phenolics, tannins and flavonoids compared to *P. lagopus* and *P. squarrosa*. However, *P. lagopus* exhibited the highest values of flavonoids and saponins. Tannins, alkaloids and saponins did not show significant variation between *P. lagopus* and *P. major* (*P* ≤ 0.05).

3.2. Allelopathic effect of various Plantago extracts on *B. pilosa* germination

The allelopathic effect of the aqueous and methanolic extracts of *P. lagopus*, *P. major* and *P. squarrosa* on the germination of *B. pilosa* four days after treatment is illustrated in Fig. 1. The obtained data revealed that there was a slight significant variation between the three studied *Plantago* species (*P* ≤ 0.05); however, the degree of inhibition was significantly increased in a concentration-dependent manner.

The aqueous extracts of *P. major*, *P. lagopus* and *P. squarrosa* at 10 mg ml⁻¹ inhibited the germination of *B. pilosa* by about 72.41%, 62.96% and 62.96%, respectively. On the other hand, the lowest concentration (2.5 mg ml⁻¹) of *P. lagopus*, *P. major* and *P. squarrosa* extracts showed the lowest inhibition percentage of germination (10.34%, 7.41% and 3.70%, respectively). The IC₅₀ values (the concentration of a substance that is required for 50% inhibition of a specific biological or biochemical function) of the germination of *B. pilosa* were 7.50 mg ml⁻¹, 7.88 mg ml⁻¹ and 7.91 mg ml⁻¹, respectively for *P. squarrosa*, *P. major* and *P. lagopus* extracts (Fig. 1).

On the other hand, *P. lagopus* methanolic extract showed a complete inhibition of germination at 7.5 mg ml⁻¹, while *P. major* extract showed a complete inhibition of germination at 10 mg ml⁻¹. *P. squarrosa* extract showed the highest inhibition percentage (85.19%) at 10 mg ml⁻¹. At the lowest concentration (2.5 mg ml⁻¹), *P. lagopus*, *P. major* and *P. squarrosa* extracts showed the following inhibition percentages: 72.41%, 62.96% and 62.96%, respectively.

Table 1 – The composition of the active secondary chemical constituents (mg/g dry weight) of the three studied *Plantago* species.

| Plant species     | Phenolics  | Tannins   | Alkaloids | Flavonoids | Saponins |
|-------------------|------------|-----------|-----------|------------|----------|
| *Plantago lagopus*| 50.3 ± 1.32| 20.3 ± 0.53| 12.5 ± 0.26| 10.0 ± 0.28| 10.8 ± 0.33|
| *Plantago major*  | 123.9 ± 3.26| 21.3 ± 0.56| 12.0 ± 0.42| 16.0 ± 0.27| 10.4 ± 0.32|
| *Plantago squarrosa*| 40.8 ± 1.07| 16.7 ± 0.44| 10.0 ± 0.36| 13.0 ± 0.18| 6.8 ± 0.26|
| LSD₀.₀⁵          | 6.54       | 1.58      | 0.94      | 1.07        | 0.77     |

Different letters in each column indicate values with significant variation (*P* ≤ 0.05).

Fig. 1 – The inhibitory effect and IC₅₀ of both aqueous and methanolic *Plantago* extracts on the germination percentage (mean value) with the error bars of *Bidens pilosa* four days after treatment. Different letters indicate values with significant variation (*P* ≤ 0.05).
showed the lowest inhibition percentage of germination expressed 17.24%, 14.81% and 7.41%, respectively. The IC₅₀ values of the germination of *B. pilosa* were 4.32 mg ml⁻¹, 4.52 mg ml⁻¹ and 5.89 mg ml⁻¹, respectively for *P. lagopus*, *P. major* and *P. squarrosa* (Fig. 1).

### 3.3 Allelopathic effect of various Plantago extracts on *B. pilosa* radicle growth

The allelopathic effect of both aqueous and methanolic extracts on *B. pilosa* radicle growth after ten days of treatment revealed that there was not any significant variation between the three studied *Plantago* species (*P* ≤ 0.05). However, the degree of inhibition significantly increased in a dose-dependent manner (Fig. 2).

The aqueous extract of *P. lagopus* expressed a complete inhibition of radicle growth of *B. pilosa* at 10 mg ml⁻¹. However, *P. major* and *P. squarrosa* extracts showed 90.48% and 87.39%, respectively. At 2.5 mg ml⁻¹, *P. lagopus*, *P. major* and *P. squarrosa* extracts showed the lowest inhibition percentage of radicle growth 42.22%, 32.14% and 34.23%, respectively. The IC₅₀ values for *P. lagopus*, *P. major* and *P. squarrosa* extracts on the radicle development of *B. pilosa* were 4.18 mg ml⁻¹, 4.85 mg ml⁻¹ and 5.14 mg ml⁻¹, respectively (Fig. 2).

On the other side, the methanolic extracts from *P. major*, *P. lagopus*, and *P. squarrosa* at 10 mg ml⁻¹ inhibited the radicle growth of *B. pilosa* by 98.10%, 97.93%, and 93.15%, respectively. At the lowest concentration (2.5 mg ml⁻¹), *P. lagopus*, *P. major* and *P. squarrosa* extracts showed the lowest inhibition percentage of radicle growth (93.79%, 92.38% and 75.34%, respectively). The IC₅₀ values of *P. lagopus*, *P. major* and *P. squarrosa* extracts on *B. pilosa* were 1.82 mg ml⁻¹, 1.97 mg ml⁻¹ and 2.89 mg ml⁻¹, respectively (Fig. 2).

### 3.4 Allelopathic effect of various Plantago extracts on *B. pilosa* plumule growth

The phytotoxic effect of both methanolic and aqueous extracts from the studied *Plantago* species on *B. pilosa* plumule growth revealed slight significant variation between the three studied *Plantago* species. Nevertheless, there was a highly significant variation (*P* ≤ 0.05) between the different concentrations (Fig. 3).

The aqueous extracts from *P. lagopus*, *P. major* and *P. squarrosa* showed the highest inhibition percentage of *B. pilosa* plumule growth (88.44%, 69.62% and 49.37%, respectively) at 10 mg ml⁻¹. However, at 2.5 mg ml⁻¹, *P. lagopus*, *P. major* and *P. squarrosa* extracts inhibited the plumule growth by 11.56%, 18.35% and 18.36%, respectively. The IC₅₀ values of the plumule development of *B. pilosa* were 5.18 mg ml⁻¹, 7.71 mg ml⁻¹ and 10.43 mg ml⁻¹, respectively for *P. lagopus*, *P. major* and *P. squarrosa* extracts (Fig. 3).

However, the methanolic extracts of *P. lagopus* and *P. major* completely inhibited *B. pilosa* plumule growth at 7.5 mg ml⁻¹ and 10 mg ml⁻¹, respectively. However, *P. squarrosa* extract expressed 93.75% at 10 mg ml⁻¹. On the other hand, the lowest concentration (2.5 mg ml⁻¹) of *P. major*, *P. squarrosa* and *P. lagopus* extracts inhibited the plumule growth by 60.95%, 33.09% and 22.92%, respectively. The IC₅₀ values for *P. major*, *P. lagopus* and *P. squarrosa* extracts were 2.11 mg ml⁻¹, 4.81 mg ml⁻¹ and 4.81 mg ml⁻¹, respectively (Fig. 3).

### 4 Discussion

Medicinal plants are used as old as human civilization and continuous efforts of scientists around the world are being made...
to isolate and characterize novel bioactive compounds from these plants. The present investigation revealed that, *P. major* is rich in phenolics, tannins and flavonoids compared to that reported by Kobeasy et al. [21]. However, the phenolics of *P. lagopus* was lower than reported in Turkish ecotype [12]; this could be corroborated to the variation in the habitat, climate and/or genetic pool [22,23]. On the other hand, it attained the highest content of flavonoids and saponins compared to *P. major* and *P. squarrosa*.

The allelopathic assay of the present study showed that the germination of *B. pilosa* was completely inhibited under treatment of *P. lagopus* and *P. major* methanolic extracts at 7.5 mg ml\(^{-1}\) and 10 mg ml\(^{-1}\), respectively. In addition, the aqueous extracts of *P. major*, *P. lagopus* and *P. squarrosa* at 10 mg ml\(^{-1}\) inhibited the germination of *B. pilosa* by about 72.41%, 62.96% and 62.96%, respectively. Moreover, the germination inhibition of *B. pilosa* increased with the increase in the extracts concentration [5,24–26]. Many plant species showed inhibitory effects on *B. pilosa* germination such as Cajanus cajan, maize roots and rice husks [27], Lantana camara [28] and Ipomoea cairica [29]. In addition, many identified allelochemicals had inhibitory effects on *B. pilosa* germination such as eugenol [30] and parthenin [1].

The present results showed that the three studied Plantago species showed higher inhibition percentage on *B. pilosa* germination than that attained by maize roots, rice husks and *C. cajan* at 10 mg ml\(^{-1}\) [27]. Additionally, they are more effective against *B. pilosa* germination compared to that of *I. cairica* [29]; however the concentration used in the present study was lower than the *I. cairica* extract. The allelochemicals inhibited germination perhaps by affecting the cell division and elongation process that are very important at this stage or by interfering with oxidative enzymes [31] involved in the mobilization of nutrients necessary for germination [6,24] or by increasing ion leakage by altering membrane permeability [32].

*P. lagopus* aqueous extract showed a complete inhibition of *B. pilosa* radicle growth at 10 mg ml\(^{-1}\); moreover, *P. major* and *P. squarrosa* showed the maximum inhibitory effect of *B. pilosa* radicle growth at the same concentration. The same observation was reported for plumule growth, although *B. pilosa* radicle is more sensitive to the Plantago extracts than the plumule which could be attributed to the direct contact of radicle to the allelochemicals [24]. Additionally, the inhibition of *B. pilosa* seedling growth was concentration-dependent. This is similar to the effect of eugenol and parthenin on *B. pilosa* seedling growth [1,30].

The reduction in the seedling growth of *B. pilosa* in this study may be attributed to reduction in cell division of the seedlings, altering the ultrastructure of the cells [33]. The reduction of protein and nucleic acids, as well as the alteration of the ion uptake, water balance, phytohormone balance, photosynthesis, respiration and inactivate several enzymes in *B. pilosa* seedling growth [24,26,34].

The methanolic extract of *P. major*, *P. lagopus*, and *P. squarrosa* inhibited the radicle growth of *B. pilosa* by 98.10%, 97.93%, and 93.15%, respectively at 10 mg ml\(^{-1}\). On the other hand, *P. lagopus* and *P. major* extracts showed a complete inhibition of *B. pilosa* plumule growth at 7.5 mg ml\(^{-1}\) and 10 mg ml\(^{-1}\), respectively. The radicle growth of *B. pilosa* was more sensitive than plumule under treatment of the methanolic extracts of the three studied Plantago species. This observation is comparable to other studies [35]. Previous studies have indicated that weed species might vary in their response tolerance to phytotoxicity [4,5].

The allelopathic effect of the studied Plantago species could be attributed to several bioactive compounds that act in a synergistic manner or to compounds which regulate one another

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**Fig. 3** – The allelopathic effect and IC\(_{50}\) of both aqueous and methanolic Plantago extracts on the plumule growth inhibition percentage (mean value) with the error bars of Bidens pilosa ten days after treatment. Different letters indicate values with significant variation (*P* ≤ 0.05).
such as flavonoid, phenolic acids, saponin, alkaloids and tannins. *Plantago* species was reported to contain several bioactive secondary metabolites such as vanillic acid, iridoid glycoside (aucubin), caffeic acid derivatives, chlorogenic acid, ferulic acid, *p*-coumaric acid and triterpenes (oleanolic acid, ursolic acid) [36–38]. Many of these compounds were reported as allelochemicals [24,39].

According to the IC₅₀ values, the methanolic extracts of the three *Plantago* species were more phytotoxic on the germination of *B. pilosa* than the aqueous extract. Moreover, the allelopathic effects of the methanolic extracts on the growth of both radicle and plumule were higher than aqueous extracts. This could be attributed to the degree of solubility of the allelochemicals in the studied *Plantago* species [40]. Methanol has the ability to extract a wide variety of active components compared to water.

5. Conclusion

The germination of *B. pilosa* was completely inhibited under treatment of *P. lagopus* and *P. major* methanolic extracts at 7.5 mg ml⁻¹ and 10 mg ml⁻¹, respectively. Moreover, both radicle and plumule were strongly inhibited under the same treatment. This could be attributed to the high content of bioactive constituents. Therefore, these two species could be possible candidates to be used in managing this noxious weed in an ecofriendly bio-control method. Moreover, further studies are needed to identify and characterize the proper allelochemicals and demonstrate their modes of action.

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