Mechanisms of Phosphorous Uptake Efficiency of Safflower and Sunflower Grown in Different Soils

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Abstract Plant species vary in their phosphorous (P) use efficiency under suboptimal P supplies using different strategies, but the mechanisms are not clearly documented for some alternative plant species. Safflower was considered as low input oil crop, but its P uptake efficiency mechanism was not fully investigated. Therefore P uptake efficiency of safflower was studied as compared to sunflower under semi-controlled conditions in sandy and loamy soils. Both species responded strongly to increasing P supplies in both soils and performed better in loamy soil. Both species had similar agronomic P efficiency in both soils, indicated by similar external P requirement under P-deficient conditions. Under P deficiency, safflower had less relative shoot and root production when they were grown in sandy soils and the opposite was found in terms of loamy soil. Safflower had the disadvantage of less root length and root shoot ratio in both soils under low and high P supplies but had the advantage of higher specific root density, less root radius, and slower shoot growth rate. Under P deficiency in both soil types, both species responded similarly in terms of P influx, depleting P from soil solution and P concentration in shoots. Under high P supply, P influx and P concentration in shoots was less in safflower in both soil types. Safflower was characterized by higher shoot demand on roots for P under low and high P supplies in both soil types. Therefore the cause of high root demand on P in safflower roots at low and high P supplies stems from the low root shoot ratio of safflower at both P supplies, low P concentration and low P influx of safflower at high P supplies not because of higher shoot growth rate of safflower at low and high P supplies. Therefore using different measures of utilization efficiency parameters to differentiate plant species and genotypes to superior and inferior could be in some cases misleading.

Keywords: P uptake efficiency, alternative crops, safflower, P influx, root shoot ratio, shoot demand


1. Introduction

Phosphorus (P) is an essential element for life, serving as an integral component of nucleic acids, lipids and a diverse range of other metabolites [1]. After nitrogen and potassium, P is quantitatively the most important inorganic nutrient for plant growth and crop productivity, unless supplied as fertilizer [2]. Therefore P is the most determinant nutrient for crop yield in many regions of the world [3]. Because P is a non-renewable resource, its global reserves is continuously depleting due to extraction by mining day and night [4], making P-fertilizer prices continue increasing [5,6]. The availability of P in soil is low [7] as a result of its fixation, being utilized by organisms forming organic P, and by sorption onto iron and aluminium [5,8]. The recovery of fertilizer P is very low [9], often below 15% in the first year of application and hardly reaches 50% after 30 years [10]. Although, in view of limited P resources [5] and serious environmental and economic consequences [11], a considerate use of P is mandatory to correct nutrient deficiencies [12,13]. P application has been shown to be particularly effective with respect to yield formation in safflower and sunflower [14,15,16], but in organic farming, where application of inorganic P fertilizers is not permitted [17,18], the P availability is not easily increased [19]. Plant species and even cultivars differ in their ability to grow or yield well at suboptimal P supply with remarkable ability to acquire sparingly available soil P, and to utilize internal P efficiently, that could be explored for future use in crops selection [5,12,13,15,16]. In developing countries, where the proportion of less fertile soils is particularly high, it may be difficult to fulfill the nutritional requirements of high-yielding crops [20]. It is thus desirable to aim for efficient use of P, both in view of resource limitations and environmental constrains for increasing the production potential on marginal land [21-27]. The use of alternative oil crops that differ in their response to P nutrition is a possibility to meet the increasing global demand for vegetable oil, and may be possible if phosphorus efficiency mechanisms are illustrated [12,13]. NUE involves various soil and plant mechanisms that contribute to the variability in uptake and utilization of nutrients by different plants in different soils [13,28,29]. Definitions of nutrient efficiency vary greatly [14,21], and in some cases may be even misleading in the quest for identification of mechanisms for enhanced nutrient acquisition and utilization [12,13,30,31,32].
Nutrient supply to plants results from interactions between plant roots and soil, depending on the nutrient quantity, nutrient availability and mobility in soil, and uptake kinetics of the root system [5,33,34,35]. As plant roots absorb a nutrient ion, soil solution concentration decreases at the root surface, the equilibrium in soil is thus disturbed, a gradient created, and the adjacent soil release nutrients from the solid soil phase into solution, and/or transport nutrient from the bulk of the soil to the root [36,37,38]. Phosphorous uptake by roots from the rhizosphere is affected by desorption of P from soil particle surface, transport of P in the soil solution towards the root surface and inflow of P into root [39,40,41]. These processes depend on soil parameters, and plant parameters as well as the nutrient characteristics [35,42], as a result of interaction between P availability (quantity and mobility) in soil and the ability of plant to acquire it [13,36,37,38]. A prerequisite of P uptake is the contact between plant roots and the nutrients in soil, which occur by root growth to the places where nutrients are located and in the same time the transport of nutrients through the soil to the root surface [36]. Therefore, plants develop large root systems to expose large areas of root surface to the soil [36,43,44,45].

The soil is a medium consists of three phases; solid, liquid, and gas, in which the liquid phase is the actual medium for ion transport [46]. Ion diffusion occur from surfaces of solid soil material, to the soil solution towards the root cell, a process that is important for the release of ions located in the soil particle to nourish the plant [36,47]. To make use of diffusion it is important to lower the initial ion concentration of the soil solution around roots in order to create a concentration gradient from soil toward the root to cause diffusive flux, and to disturb the equilibrium between nutrients on the solid phase with those in the liquid phase to cause their release from the matrix into solution [36,48,49]. The concentration of nutrient in soil solution, the volume of soil that is filled with water, and the geometry of the soil pore system are the major factors affecting diffusion ability of ions in soil [50]. Plant roots act as a sink for soil nutrients, and the amount of an ion that arrives at a plant root surface depend on the size of root system, length, or surface area, and root distribution in the soil profile [48]. It is the plant that initiates nutrient transport from soil to root [51]. In order to search for low input alternative oil crops and to understand factors affecting P uptake efficiency among plant species, this study aims to investigate the influence of different P supplies in two soil types (sandy and loamy) on the components of the uptake efficiency of safflower and sunflower.

2. Methodology

2.1. Experimental Design

A pot experiment was conducted to evaluate P uptake efficiency and P dynamics in the rhizosphere of safflower (Carthamus tinctorius L., variety ‘Sabina’) and sunflower (Helianthus annuus L., variety ‘Peredovick E’), grown in two low P status soil types (loamy and sandy), using three levels of P supply in a greenhouse having semi-controlled climatic conditions. Before beginning the experiment, field-moist soil samples were sieved to 2-mm particle size, from which, subsamples of soil were air dried and analyzed for extractable P, exchangeable K, Mg, and pH. Initially, the sandy soil (pH 5.6 by water extraction) contained 26 mg kg\(^{-1}\) CAL-extractable P, 22 mg kg\(^{-1}\) CAL-exchangeable K, and 28 mg kg\(^{-1}\) NH\(_4\)-acetate exchangeable Mg. The loamy soil (pH 7.0 by water extraction) contained 16.5 mg kg\(^{-1}\) CAL-extractable P, 28 mg kg\(^{-1}\) CAL-exchangeable K, and 141 mg kg\(^{-1}\) NH\(_4\)-acetate exchangeable Mg.

Mitscherlich pots (6 L) were filled with 3 kg sand (0 mg kg\(^{-1}\) CAL-extractable P, 3 mg kg\(^{-1}\) CAL-exchangeable K, and 1.8 mg kg\(^{-1}\) NH\(_4\)-acetate exchangeable Mg, pH in water was 7.3) and 3 Kg either sandy or loamy soil. Three P levels (0, 0.2, and 1.0g P pot\(^{-1}\)) were added as Ca(H\(_2\)PO\(_4\))\(_2\)*H\(_2\)O, resulting in solution P (mg P L\(^{-1}\) soil solution) content of 0.2, 0.6, 8.2 for sandy soil and 0.0, 0.2, 45. 7 for loamy soil in consecutive added P levels. The extractable P content (mg P Kg soil\(^{-1}\)) of the soil after adding external P were 25.0, 50.8, 229.5 for sandy soil and 35.4, 54.3, 263.0 for loamy soil in respective P supplies (0, 0.2, and 1.0g P pot\(^{-1}\)). Other nutrients added per pot were 2g N (as NH\(_4\)NO\(_3\)), 3g K (as K\(_2\)SO\(_4\)), 0.8g Mg (as MgSO\(_4\)), micronutrients were added in adequate amount for both species in both soil types (mg pot\(^{-1}\): 17.5 B, 2.5 Mo, 8 Cu, 50 Mn, and 40 Zn). Three safflower or two sunflower plants were planted in each pot (because sunflower is larger than safflower). The treatments were replicated four times. Four additional pots per each P level for each soil type were left unplanted as control for the measurement of extractable and soil solution P concentrations during the experiment without be affected by plant species. The planted and the unplanted pots were watered daily to nearly volumetric soil water content of 35%. The experiment was conducted as a completely randomized design.

2.2. Harvesting and Analytical Procedures

The plants were harvested in two harvest times. The plants in one pot of each treatment (plant species and soil type) was harvested in the first harvest after 42 days from sowing for both crops in both soil types, and the rest three pots in each treatment were harvested in the second harvest after 56 days from sowing for both species in both soil types. At each harvest, the soil in each shoot harvested was weighed (moist soil with roots), and then soil was cut to two similar parts (also accurately weighed). One part of the soil in each pot was sieved to remove the roots and then was sub-sampled for the following measurements: a soil sample to measure the moisture content of the soil (around 100g), a soil sample for measuring soil solution P (around 350g), and finally a soil sample for measuring extractable P (around 100g). The second half of the soil of each pot was put in sealed plastic bags and kept at 6°C for collecting the roots within 48 hours.

Harvested plants were separated into stems, leaves and roots (half roots per pot were collected). Stems and leaves were measured for fresh and dry weights, then were analyzed for their P contents. The roots in half soil of the pot (precisely weighed) were separated from the soil by washing over a 0.2 mm sieve, then were preserved in...
plastic bottle at 6°C to be measured for their fresh weight and length within 24 hours.

**Shoot measurements and P analysis**

At harvest, the dry weight of plant parts were determined after drying at 70°C till constant weight. Dried plant materials were ground to pass a 1.5 mm sieve, of which, after thorough mixing, a sub-sample of 5 g was ball-milled to a fine powder. The plant samples were prepared for P analysis using wet microwave digestion using concentrated tri acid mixture (HNO₃, HClO₄, and H₂SO₄ with a volumetric ratio of 8:2:1). Total P of the plant material digest was measured using colorimetric method (Ammonium-Vanadate-Molybdate) [52].

**Measurement of soil solution and extractable P concentration, pH and water content**

The column displacement method was used [53] to collect the soil solution in order to determine initial soil solution P concentration. The method permits accurate determination of the unaltered composition of soil solution, in which a sample of moist soil equivalent to 350 g was packed into a plastic column with a pore in its bottom. Filter paper was placed in the bottom of each soil column to avoid soil particles losses during the collection. The samples were allowed to equilibrate for 24 h; then, deionized water was pumped to each column at a rate of 4 ml h⁻¹ until the soils reached field capacity water content. The displaced solution was collected till 25 ml to insure not to collect diluted solution, and then filtered through a 0.20μm filter. The solutions were analyzed for P by colorimetric method [52]. Soil solution concentration was measured for planted and unplanted pots immediately at the time of each harvest.

To determine solid phase (extractable) P, a 10-g subsample of soil from each pot was air dried then extracted with calcium acetate lactate (CAL) method [54]. Phosphorous concentration in the extracts was determined also using the colorimetric method [52].

The pH was measured using 0.01 M CaCl₂ (1: 2.5 soil: solution ratio). The gravimetric water content was determined in soil samples when soil samples were dried at 105°C to constant weight and the water content was calculated as the difference before and after drying.

**Root length, root radius, and specific root density measurement**

The roots were carefully collected by washing off the soil in a sieve with a 0.2mm wide mesh. Roots were cleaned of any foreign materials and then spread on paper towels. The surface moisture on the roots was removed manually by applying uniform pressure using paper towels. The root length in the fresh weight subsample was measured using a hand tally counter. The root length in the aliquot of the sample in cm, GD = Grid dimension (1.25 cm grid squares), N = Number of intercepts.

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**Root density, root length, and specific root density measurement**

The roots were carefully collected by washing off the soil in a sieve with a 0.2mm wide mesh. Roots were cleaned of any foreign materials and then spread on paper towels. The surface moisture on the roots was removed manually by applying uniform pressure using paper towels and finally the root fresh weight (RFW) was recorded. Afterwards, a representative fresh root material of different parts of the root system (upper, middle and apical) was cut into small pieces (0.5-1 cm). After fine cutting these root portions (1-3 mm), two sub-samples were taken accurately for the root length measurement, using the line intersection method [55]. The fresh weight of the rest of the roots was recorded and then oven-dried and grinded as described above for the shoots. Each fine-cut root sub-sample was dispersed in a known volume of water and an accurately measured volume of aliquot of the root soap was taken and poured in a plastic dish with a grid bottom with lines 1.25 cm apart. The total number of root intercepts with the vertical and horizontal grid lines was counted by means of hand tally counter. The root length in the aliquot of the sub-sample was calculated using the following equation:

\[ \text{RL} = \frac{(11/14)\times \text{GD} \times N}{\pi} \]

Where, \( \text{RL} \) = Root length of the sample in the plastic dish in cm, GD = Grid dimension (1.25 cm grid squares), N = Number of intercepts.

The root length in the fresh weight subsample was calculated from a volumetric relation between the aliquot and the subsample. The total root length of the plants was obtained from the weight relation between the subsample and the total weight. Assuming that the specific weight of roots is 1g cm⁻³, the mean root radius (r₀) was calculated as:

\[ r_0 = \sqrt{\frac{\text{Root fresh weight (RFW)} / \pi \times \text{Root length (RL)}}{}} \]

The specific root density or root length density (RLₚ) was calculated by dividing root length (RL) by the soil volume of the pot and interpreted as cm root cm⁻³ soil⁻¹.

**Shoot growth rate**

This ratio relates the difference in shoot growth between the two harvests divided by the number of days between the two harvests: Shoot growth rate (GRₚ) = \( \ln{(SW_2 - SW_1) / (t_2 - t_1)} \)

Where, SW₁ and SW₂ are shoot dry weight at the first and the second harvests respectively, and t₁ and t₂ are number of days of the plants at the first and the second harvests respectively.

**Shoot demand (SD): shoot growth rate in relation to average root length**

This ratio relates the K acquisition load imposed by shoot growth to each root segment. It was calculated by dividing the shoot growth rate (GRₚ) by the average root length (aRL) assuming exponential root growth: Shoot growth rate/Root length (GRₚ/RL) = \( (SW_2 - SW_1) / (RL_2 - RL_1) \) X \( \ln{(RL_2 / RL_1)} \)

Where RL is the root length [cm] and SW is the shoot dry weight [g] at two harvest dates (t₂-t₁).

**Net P influx**

The influx is the net amount of a nutrient that is taken up per unit root length (or root surface area) per unit time. Since direct measurement of the influx is not possible, only an average influx can be calculated for a given time period. At least two harvests are needed in which the nutrient content and root length of the plants are known. Assuming that the roots of young plants show exponential growth, the average influx was calculated [56]: \( \ln{U} = [(U_2 - U_1) / \ln{(RL_2 / RL_1)}] / (RL_2 - RL_1) \)

Where U is the influx, U₀ is the shoot P content [mol] at two harvest dates (t₂-t₁) related to the root length between the two harvests (RL₂-RL₁).

### 2.3. Statistical Analysis

All statistical analyses were carried out using SAS (SA Institute Inc., Cary, USA, Release 8.02, 2001). Comparisons of means between different treatments were carried out using the GLM procedure considering a fully randomized design. With multiple t-test, the Bonferoni procedure was employed in order to maintain an experiment-wise α of 5%.
3. Results

3.1. Growth and Morphology

Both species responded strongly to increasing P supply in terms of fresh weight of all growth parameters (Table 1). Both species produced significantly higher fresh weight of all growth parameters when grown in loamy soil as compared to sandy soil. At zero added P supply, both species showed better relative growth in terms of fresh weight of all growth parameters when grown in sandy soils in comparison with that grown in loamy soils. Under very low P supply, safflower leaves were reduced more than stems in both soil types, the same response was observed for sunflower grown in sandy soil but the opposite was recorded in loamy soil. Comparing both species under low P supply (0 and 0.2g P pot⁻¹), relative leaves, stems, shoot, root, and total fresh weight of safflower was higher than that of sunflower, the only exception was recorded for root fresh weight in loamy soil at 0.2g P pot⁻¹.

The dry weight of the plant parts of both species responded strongly to increasing P supply (Table 2). Under low P supplies, safflower leaves dry weights were affected more than stems in both soil types, and for sunflower plants when were grown in sandy soils only. Both species performed better in terms of dry weight production of shoot components when were grown in loamy soil as compared to sandy soils, while the opposite was found in sunflower plants when were grown with no added P supply. The relative shoot dry weight production was found not significantly different in both species under zero added P supply in sandy soil, while under 0.2 g P pot⁻¹, safflower was found superior as compared to sunflower grown in sandy soil. The same figure in loamy soil proved the superiority of safflower as compared to sunflower in terms of relative dry weight production at both 0 and 0.2g P pot⁻¹.

Table 1. Effect of P supply on fresh matter (g pot⁻¹) of safflower and sunflower as absolute value (without brackets) and relative values (between brackets). For a given species and a given soil type, means within each column followed by the same letter are not significantly different, * indicates significant difference for a given plant species and a given P level within soil types. P< 0.05, n=3

<table>
<thead>
<tr>
<th>P supply (g pot⁻¹)</th>
<th>Leaves</th>
<th>Stem</th>
<th>Shoot</th>
<th>Fine roots</th>
<th>TFW</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Sandy</td>
<td>Loamy</td>
<td>Sandy</td>
<td>Loamy</td>
<td></td>
</tr>
<tr>
<td>Safflower</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>0</td>
<td>3.9 B *</td>
<td>4.8 C</td>
<td>2.3 B *</td>
<td>2.7 C</td>
<td>6.2 B *</td>
</tr>
<tr>
<td></td>
<td>(32.5)</td>
<td>(9.8)</td>
<td>(46.9)</td>
<td>(13.1)</td>
<td>(36.5)</td>
</tr>
<tr>
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<td>8.0 AB *</td>
<td>24.0 B</td>
<td>4.1 A *</td>
<td>10.5 B</td>
<td>12.2 BA *</td>
</tr>
<tr>
<td></td>
<td>(40.0)</td>
<td>(49.0)</td>
<td>(83.7)</td>
<td>(51)</td>
<td>(71.8)</td>
</tr>
<tr>
<td>1.0</td>
<td>12.0 A *</td>
<td>49.0 A</td>
<td>4.9 A *</td>
<td>20.6 A</td>
<td>17.0 A *</td>
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<tr>
<td></td>
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<tr>
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<td>105.6 B</td>
<td>58.4 B *</td>
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<td></td>
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<td>(37.4)</td>
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<td>(34.6%)</td>
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<td>214.5 A</td>
<td>82.2 A *</td>
<td>306.2 A</td>
<td>168.8 A *</td>
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</table>

Table 2. Effect of P supply on dry matter (g pot⁻¹) of safflower and sunflower as absolute value (without brackets) and relative values (between brackets). For a given species and a given soil type, means within each column followed by the same letter are not significantly different, * indicates significant difference for a given plant species and a given P level within soil types. P< 0.05, n=3

<table>
<thead>
<tr>
<th>P supply (g pot⁻¹)</th>
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<th>Stem</th>
<th>Shoot Dry Weight</th>
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</thead>
<tbody>
<tr>
<td></td>
<td>Sandy</td>
<td>Loam</td>
<td>Sandy</td>
</tr>
<tr>
<td>Safflower</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>0</td>
<td>0.67 C *</td>
<td>0.86 C</td>
<td>0.56 A *</td>
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<tr>
<td></td>
<td>(33.0)</td>
<td>(11.7)</td>
<td>(56.6)</td>
</tr>
<tr>
<td>0.2</td>
<td>1.22 B *</td>
<td>4.12 B</td>
<td>0.83 A *</td>
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<tr>
<td></td>
<td>(60.0)</td>
<td>(55.9)</td>
<td>(83.8)</td>
</tr>
<tr>
<td>1.0</td>
<td>2.03 A *</td>
<td>7.37 A</td>
<td>0.99 A *</td>
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<td></td>
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<tr>
<td>Sunflower</td>
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<td>9.55 A *</td>
<td>22.73 A</td>
<td>4.84 A *</td>
</tr>
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</table>


**Growth rate of roots and shoot**

Root growth rate (cm root/day) for both species grown in sandy soil were statistically similar in different P supplies (Table 3). In loamy soils, root growth rate (GRr) of safflower increased significantly at low P supplies, while in sunflower the significantly lowest value was found in the lowest P supply. Comparing the same species at different soil types, safflower showed similar values in both low and high P supplies, but in intermediate P levels, GRr was significantly higher when safflower was grown in sandy soil as compared to loamy soil. Sunflower GRr was similar in both soil types at both intermediate and high P supplies, while at low P levels, plants grown in sandy soil had significantly higher values than that grown in loamy soil.

Shoot growth rate (g TDW/day) of safflower plants was found the highest in plants grown in intermediate P supplies in both soils. In loamy soil, shoot growth rate (GRs) of safflower was similar in both low and high P supplies, while in sandy soil, this figure was found lower in plants grown in high P supply as compared to low P supply. Sunflower GRs didn’t change significantly with increasing P supply when plants were grown in sandy soil, while the same figure in loamy soil was significantly similar in intermediate and high P supplies, while at low P level the value was significantly lower than the other P levels (Table 3). GRr and GRs for the same plant species responded similarly in different soil types; both traits were found inferior in safflower as compared to sunflower in sandy soil in both low and high P levels but not in the intermediate level (both species are similar). In loamy soil both traits were similar in both species at low P supply but safflower was interior as compared to sunflower at intermediate and high supplies.

### 3.2. P Uptake Efficiency Parameters

Uptake efficiency parameters discussed in this investigation where those related to plant (root parameters) and those related to soil. Root parameters influencing the P uptake are: root length, root diameter, specific root density, P influx in roots, shoot demand for P on roots, and root shoot ratio. Soil parameters include soil solution P, extractable P, and pH.

**Root length (RL), specific root density (RLv), and radius (r₀)**

Safflower root length (cm pot⁻¹) increased with increasing external P supply in both soil types (Figure 1), and it was longer in loamy soil than that in sandy soil (the difference at 0 added P was not significant). Roots of sunflower plants grown in loamy soil increased in length significantly with progressive P supplies, while that grown in sandy soils were longer at the highest P supply (1.0 g P pot⁻¹), and shortest at the intermediate P supply (0.2 g P pot⁻¹), but under 0 added P, root length were intermediate. At 0 added P level, sunflower grown in sandy soils enlarged the root length more than that in loamy soil, while the opposite was recorded at 0.2 and 1.0 g P pot⁻¹.

### Table 3. Effect of P supply on relative root growth rate (cm root/day), relative shoot growth rate (g TDW/day) for safflower and sunflower.

For a given species and a given soil type, means within each column followed by the same capital letter are not significantly different, means in the same soil type and the same P level and different plant species followed by the same small letter are not significantly different, * indicates significant difference for a given plant species and a given P level within soil types. P<0.05, n=3

<table>
<thead>
<tr>
<th>P supply (g pot⁻¹)</th>
<th>Root growth rate (cm root/day)</th>
<th>Shoot growth rate (g TDW/day)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Sandy</td>
<td>Loam</td>
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<tr>
<td>Safflower</td>
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</tr>
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<td>0.104 BA, b</td>
<td>0.118 A, a</td>
</tr>
<tr>
<td>0.2</td>
<td>0.193 A, a *</td>
<td>0.098 A, b</td>
</tr>
<tr>
<td>1.0</td>
<td>0.011 B, b</td>
<td>0.019 B, b</td>
</tr>
<tr>
<td>Sunflower</td>
<td></td>
<td></td>
</tr>
<tr>
<td>0</td>
<td>0.183 A, a *</td>
<td>0.081 B, a</td>
</tr>
<tr>
<td>0.2</td>
<td>0.187 A, a</td>
<td>0.158 A, a</td>
</tr>
<tr>
<td>1.0</td>
<td>0.215 A, a</td>
<td>0.179 A, a</td>
</tr>
</tbody>
</table>

*Figure 1. Effect of P supply on root length (cm pot⁻¹) of safflower (A) and sunflower (B) in sandy soil and loamy soil. For a given species and a given soil type, means within each column followed by the same capital letter are not significantly different. * indicates significant difference for a given plant species and a given P level within soil types. P<0.05, n=3*
Specific root density (cm root/ g root) decreased significantly with improving P supply in both soil types in safflower, and in sunflower when grown in loamy soil, while the opposite was found in sunflower grown in sandy soil (Table 4). Specific root density for each plant species were similar in different soil type. This trait was higher in safflower as compared to sunflower in sandy soil, while the opposite was recorded in loamy soil. Root radius was significantly reduced at low P supplies (0 and 0.2 g P pot⁻¹) as compared to high P supply (1.0 g P pot⁻¹) in safflower grown in both soil types and in sunflower grown in loamy soil, while the values for sunflower grown in sandy soil were reduced but this reduction was statistically not significant (Table 4). Root radius of safflower was significantly less than that of sunflower at all respective P supplies in both soil types, but this trait remained similar in each plant species and different soil type.

**Root-shoot ratio (RSR)**

Both species had higher root shoot ratios at low P supplies in both soil types (Figure 2). RSR was found significantly lower in safflower as compared to sunflower in both soil types. This trait didn’t change significantly for safflower in respective P levels among soil types. In sunflower, this parameter was similar at low P supply in both soil types, lower in sandy soil at intermediate P supply and the opposite was recorded at high P level.

### Table 4. Effect of P supply specific root density (cm/g root) and root radius (cm x 1000) of safflower and sunflower. For a given species and a given soil type, means within each column followed by the same capital letter are not significantly different, means in the same soil type and the same P level and different plant species followed by the same small letter are not significantly different, * indicates significant difference for a given plant species and a given P level within soil types. P< 0.05, n=3

<table>
<thead>
<tr>
<th>K supply (g pot⁻¹)</th>
<th>Specific root density (cm/g root)</th>
<th>Root radius (cm)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Sandy</td>
<td>Loam</td>
</tr>
<tr>
<td>Safflower</td>
<td></td>
<td></td>
</tr>
<tr>
<td>0</td>
<td>2500.1 A, a</td>
<td>2899.4 A, a</td>
</tr>
<tr>
<td>0.2</td>
<td>2166.1 B, a</td>
<td>2143.2 B, a</td>
</tr>
<tr>
<td>1.0</td>
<td>1968.8 B, a</td>
<td>1800.6 C, a</td>
</tr>
<tr>
<td>Sunflower</td>
<td></td>
<td></td>
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<tr>
<td>0</td>
<td>1498.7 B, b</td>
<td>1619.3 A, b</td>
</tr>
<tr>
<td>0.2</td>
<td>1357.3 B, b</td>
<td>1547.3 A, b</td>
</tr>
<tr>
<td>1.0</td>
<td>1605.2 A, b</td>
<td>1373.5 B, b</td>
</tr>
</tbody>
</table>

**Figure 2.** Effect of P supply on root-shoot ratio (cm root/ g shoot) pot⁻¹ of safflower and sunflower in sandy soil and loamy soil. For a given species and a given soil type, means within each column followed by the same capital letter are not significantly different. * indicates significant difference between the two columns at each P supply (A and B) or a given plant species (C and D) at the same P level. P< 0.05, n=3
**Phosphorous influx**

Phosphorous influx (pmol P/cm root/second) increased significantly at high P supply in both species in both soil types as compared to low P supplies (Figure 3). This figure for safflower grown in sandy soil was found the highest in the intermediate P supply and was similar in both high and low P supplies. Both species showed similar P influx when grown in sandy soils at low P supply, while safflower was more efficient than sunflower in loamy soils. At intermediate P levels, safflower roots were more efficient in P uptake as compared to that of sunflower in both types of soil. At high P supplies safflower roots were inferior in P uptake as compared to that of sunflower roots in both soil types. Comparing each crop in different soil types, P influx was similar at each respective P level, except for safflower at high P supply, where the value was significantly less in sandy soil as compared to that grown in loamy soil.

**Shoot demand (SD)**

Shoot demand (SD) on the root is the P acquisition load imposed by shoot growth on each cm of the root and is calculated by dividing the shoot growth rate by the average root length, assuming that the roots of plants grow exponentially (Figure 4). Shoot demand on roots increased significantly with decreasing P supply in both species and both soil types. It was higher in safflower than that of sunflower at each respective P supply for each soil type separately. It was higher in sandy soil as compared to loam soil at respective P supplies for each species separately.

**P concentration in dry matter**

Phosphorous concentration (g P 100g⁻¹ DM) increased significantly in leaves, stems and shoots of both species with increasing P supplies in both soils (Table 5). At 0 and 0.2g added P supplies, safflower grown in both soil types contained significantly similar P concentration in their leaves, while at high P supplies safflower grown in loamy soil concentrated more P in their leaves as compared to those grown in sandy soil. Sunflower leaves concentrated more P in their leaves when they were grown in sandy soils as compared to that in loamy soils at very low P supply (0g P pot⁻¹) and high P supply (1 g P pot⁻¹) while the opposite was found at the intermediate P level (0.2g P pot⁻¹). When grown in sandy soil both species concentrated similar values of leaves P% at respective low P levels (0 and 0.2g P pot⁻¹), while the same figure was significantly lower in safflower leaves as compared to sunflower at high P supply. In loamy soil, sunflower leaves contained significantly higher P% than safflower at respective 0.2 and 1.0 g P pot⁻¹ while at 0 added P, the opposite was found.

---

**Figure 3.** Effect of P supply on P influx (pmol P/cm root/second) for safflower and sunflower in sandy (A) and loamy (B) soil. For a given species and a given soil type, means within each column followed by the same capital letter are not significantly different. * indicates significant difference between columns at the same P level. P< 0.05, n=3

**Figure 4.** Effect of P supply on shoot demand; shoot growth rate (GR)/average root length (aRL) ratio for safflower and sunflower in sandy (A) and loamy (B) soil. For a given species and a given soil type, means within each column followed by the same capital letter are not significantly different. * indicates significant difference between the two columns at the same P level. P< 0.05, n=3
P concentrations in stems of both species were found lower than that of leaves at all respective P levels in both soil types. Both species contained significantly similar stem P% in the same soil type and respective low P supplies (0 and 0.2 g P pot⁻¹), while safflower concentrated less P at high P supply (1 g P pot⁻¹) in respective soil type. When grown in sandy soil, safflower stems contained higher, similar, and lower P% as compared to that grown in loamy soil when grown at 0, 0.2, and 1.0 g P pot⁻¹ respectively. The same figure concerning sunflower showed that sunflower stems contained similar P concentrations at respective P supplies within the two soil types.

When grown in sandy soil, safflower stems contained higher, similar, and lower P% as compared to that grown in loamy soil when grown at 0.0, 0.2, and 1.0 g P pot⁻¹ respective. The same figure concerning sunflower showed that sunflower stems contained similar P concentrations at respective P supplies within the two soil types.

Shoots of both crops contained similar P% at respective 0 and 0.2 g P pot⁻¹ in both soils, while safflower shoot P% was significantly lower than that of sunflower in both soil types at high P supply. Comparing P% in shoots of each plant species within soil type reveals that both species concentrated significant higher values when grown in sandy soils as compared with loamy soil at 0 added P supplies. At 0.2 g P supply, P% was similar in safflower among soil types, while that of sunflower was higher in loamy soil. At high P supply, shoots of safflower concentrated more P in loamy soil while the opposite was recorded for sunflower.

Table 5. Effect of P supply on P concentration (g 100g⁻¹ DM) of safflower and sunflower. For a given species and a given soil type, means within each column followed by the same capital letter are not significantly different, means in the same soil type and the same P level and different plant species followed by the same small letter are not significantly different, * indicates significant difference for a given plant species and a given P level within soil types. P< 0.05, n=3

<table>
<thead>
<tr>
<th>K supply (g pot⁻¹)</th>
<th>Leaves</th>
<th>Stem</th>
<th>Shoot Dry Weight</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Sandy</td>
<td>Loam</td>
<td>Sandy</td>
</tr>
<tr>
<td>Safflower</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>0</td>
<td>0.289 B, a</td>
<td>0.207 C, a</td>
<td>0.107 B, a *</td>
</tr>
<tr>
<td>0.2</td>
<td>0.361 BA, a</td>
<td>0.395 B, b</td>
<td>0.172 B, a</td>
</tr>
<tr>
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<td>0.693 A, b</td>
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<tr>
<td>Sunflower</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>0</td>
<td>0.307 B, a *</td>
<td>0.153 C, a</td>
<td>0.122 B, a</td>
</tr>
<tr>
<td>0.2</td>
<td>0.337 B, a *</td>
<td>0.432 B, a</td>
<td>0.143 B, a</td>
</tr>
<tr>
<td>1.0</td>
<td>0.917 A, a *</td>
<td>0.791 A, a</td>
<td>0.693 A, a</td>
</tr>
</tbody>
</table>

Table 6. Effect of P supply on P soil solution (mg K L⁻¹), extractable (CAL) P (mg 100g⁻¹), and pH. For a given species and a given soil type, means within each column followed by the same capital letter are not significantly different, means in the same soil type and the same P level and different plant species followed by the same small letter are not significantly different, * indicates significant difference for a given plant species and a given P level within soil types. P< 0.05, n=3

<table>
<thead>
<tr>
<th>K supply (g pot⁻¹)</th>
<th>Soil solution P (mg P L⁻¹)</th>
<th>CAL P (mg 100g⁻¹)</th>
<th>Soil pH (CaCl₂ extract)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Sandy</td>
<td>Loam</td>
<td>Sandy</td>
</tr>
<tr>
<td>Safflower</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>0</td>
<td>0.14 C, a *</td>
<td>0.07 C, a</td>
<td>2.42 C, a *</td>
</tr>
<tr>
<td>0.2</td>
<td>0.24 B, b *</td>
<td>0.50 B, a</td>
<td>3.29 B, a *</td>
</tr>
<tr>
<td>1.0</td>
<td>0.79 A, a *</td>
<td>2.38 A, a</td>
<td>10.73 A, a *</td>
</tr>
<tr>
<td>Sunflower</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>0</td>
<td>0.18 C, a *</td>
<td>0.05 C, a</td>
<td>2.43 C, a *</td>
</tr>
<tr>
<td>0.2</td>
<td>0.33 B, a *</td>
<td>0.24 B, b</td>
<td>3.00 B, a *</td>
</tr>
<tr>
<td>1.0</td>
<td>0.86 A, a *</td>
<td>2.05 A, a</td>
<td>7.83 A, b *</td>
</tr>
</tbody>
</table>

Figure 5. Effect of P supply (g/ pot) on soil solution P (mg P/ L soil solution) (A) and CAL-P (mg P/ 100g soil) (B) in unplanted pots loamy and sandy soil.
3.3. Soil Parameters

Available (soil solution) P, extractable (Calcium Acetate Lactate (CAL)) P, and pH

Soil solution around roots of both plants increased significantly with increasing P supply in both plant species and both soil types (Table 6). Soil solution was found similar in both species in the same soil type at very low and very high P supplies, while it was lower in safflower at intermediate P supply in sandy soil and the opposite was found in loamy soil. In safflower, soil solution was significantly higher in sandy soil as compared to loamy soil at all respective P supplies, and similar response was recorded for sunflower except for high P supply where the opposite was observed. Calcium acetate lactate (CAL) extractable P increased significantly with increasing external P supply in both soils for both crops (Table 6). CAL-P was significantly higher in sandy soil in all respective P supplies for both plant species. The values were significantly lower for sunflower in loamy soil at all respective P supplies and in sandy soil at high P fertilization level, while values were similar for both species at 0 and 0.2g pot⁻¹ in sandy soil. Soil pH in sandy soils for both species decreased in the highest P supply only (Table 6). pH in loamy soil for safflower was significantly reduced in the intermediate P supply and was reduced significantly in both 0.2 and 1.0g pot⁻¹ for sunflower.

In the unplanted pots of both types of soil, the soil solution contained increasing amounts of P as a function of increasing external P supply. The solution obtained from loamy soils had significantly higher than that obtained from sandy soil at high added P levels (Figure 5, material and methods section). Both soils had significantly similar extractable P at all respective P supplies (Figure 5, Figure 6, material and methods section). Buffer power is normally calculated as the ratio of soil exchangeable P (mol cm⁻³ soil) and the soil solution P concentration (mol cm⁻³ soil solution). Figure 6 demonstrates the relation between both sources of P and the relation for both soil types fit the Langmuir isotherm showing the relationship between the adsorbed and equilibrium phosphorous concentration (quantity/intensity).

4. Discussion

4.1. Growth and Morphology

Biomass is an important plant trait in growth analysis and the key parameter in many allometric relationships [57,58]. Repeated measurements of biomass are the basis for the calculations of net primary production and growth rates [59], and thus a basis for quantifying physiological responses of plants to environmental conditions and their developmental processes. Thus the production of shoot...
Phosphorus is needed most by young, fast-growing tissues, and performs a number of functions related to growth, development, photosynthesis, and use of carbohydrates; hence, P-deficient soils produces poor plant growth and yield. P deficiency reduces leaf expansion, auxiliary bud growth and shoot canopy, therefore, reduces the plant's photosynthetic area and carbohydrate utilization [1]. Since cell and leaf expansion are more retarded than chloroplast and chlorophyll formation [62], low P supply increases the soluble protein and chlorophyll content per unit leaf area, resulting in small and darker green leaves [63]. It is stated that the decrease in leaf number and size is one of the earliest and most reliable responses of P-deficient plants [64]. Leaf expansion occurs due to cell multiplication and elongation of the newly formed cells in plants and turgor pressure is a crucial factor for cell expansion [65]. There were instances, where P-deficiency decreased the hydraulic conductivity of water in the roots [66, 67], and reduced the water potential of the plant, possibly by lowering the activity of the water channel proteins, aquaporins [67]. It is also possible that P-deficiency induced the closure of stomata [66], improving the water potential of these organs temporarily. The decrease in hydraulic conductivity of the root and stomatal conductance of the leaf, result in a severe reduction of leaf expansion under P-deficiency [67]. In addition to the above mentioned effects on vegetative growth, low-P supply also limits the formation of reproductive organs, results in premature leaf senescence, delayed flower initiation [15], decreased number of flowers [14] and restricted seed formation [15,16], and all contribute to yield reductions under P-limited conditions.

In agreement with our results concerning dry weight (Table 1) and fresh weight (Table 2), P nutrition was reported to have a positive influence on dry matter production in sunflower [12,68] and safflower [13,15]. The reduction of leaf biomass of both species in both soils under study (Table 1, Table 2) was particularly strong (more than stems) and was more pronounced in loam soil (relative DM production). Indeed, in plants suffering from P deficiency, reduction in leaf area [69] and leaf number [64] is the most striking effect. Although the contents of chlorophyll per unit leaf area are often increased under P deficiency [63], the photosynthetic rate per unit area is typically reduced [68], suggesting that both effects (reduction of leaf area and reduction of net photosynthesis per unit of leaf area) may contribute to the final reduction of biomass production. Unlike other reports, sunflower was more sensitive to P deficiency than safflower in terms of relative dry matter accumulation (Table 2) in loam soil under the very low and intermediate P supply and in sandy soil at intermediate P supply [15,70] and the same response was reflected in the relative fresh and dry weights of leaves, stems, and roots (fresh weight). The contribution of the stem in reducing dry matter as affected by sub-optimal external P was less than that of leaves and may be caused by the reduction of stem diameter and the height of the plants [16]. The effect of P supply on increasing the number of branches per plant in safflower was reported [15].

### 4.2. P Uptake Efficiency

Phosphorus acquisition efficiency is defined partly in terms of total uptake per plant and is related to root size, root morphology and P mobility in soil making P acquisition by the plant very dependent on soil exploration in time and space [33, 34, 81]. Nutrient uptake by plants starts with contact between plant roots and the nutrients in soil due to large root system [43]. Safflower produced less root size and root length than sunflower interpreted as absolute root fresh weight (Table 1, Figure 1) because safflower is small plant as compared to sunflower. In the other hand, the relative root size and also relative root length (value at a particular P supply related to the value at the highest P supply) of safflower were much higher than that of sunflower when both species were grown at low P supply in either sandy or loamy soil. This indicate that safflower can increase the relative root size (weight and length) under low P supply which enable the plant to overcome the low P availability by increasing its root size to explore more soil volume. This agree with a previous findings stated that the total root-length production of Beta vulgaris in field plots at harvest was 120 km m−2 in high-P plots, and 200 km m−2 in low-P plots [72]. In this line with our findings for safflower in both soil types concerning relative root size (Table 1), relative root length, specific root density, root radius (Figure 1) and also root-shoot ratio (Figure 2), researchers found plants grown under low P supply can modify their root system (length,
Different nutrient uptake rates per unit root and time [91]. Phosphorus uptake kinetics parameters include maximum net influx per centimeter of root ($I_{\text{max}}$), Michaelis-constant ($K_m$) and minimum soil solution concentration ($C_{\text{min}}$) [12]. Under conditions in which the rate-determining step in P uptake is related to the root, P uptake will increase if root length per unit plant weight and $I_{\text{max}}$ increase, and $K_m$ and $C_{\text{min}}$ decrease [92]. These parameters vary with P concentration in the soil solution. In contrary with our findings (Figure 3), other researchers reported an increase in $I_{\text{max}}$ values under P deficient supply [93,94]. In our research, the P influx in the roots of both species was reduced under low P supply which indicate that both species don’t use this mechanism to enhance P uptake under P deficiency. The P influx of safflower was higher than sunflower at low P supply but was the opposite at high P supply. Other researchers documented that $K_m$ and $C_{\text{min}}$ values were not affected by P supply [94].

The influx can be used to express plant’s nutrient demand on the roots. This sink property can be described as follows [33]: $I = X \frac{GRs}{RL} \frac{W}{RL}$. The sink intensity depends on the nutrient concentration in the plant ($X$), the relative growth rate (RGR), and the ratio of total plant weight ($W$) to root length ($RL$). All these parameters change with plant species [95,96], variety [82,97], and are more pronounced at developmental stage [96]. The ability of plants to adapt their morphological and physiological root characteristics to variable nutrient availability is genetically determined [98,99]. Previous reports found that, the influx of P per unit root length greatly enhanced by root hairs [89], a trait was not investigated in this research.

The assessment of the kinetic parameters ($I_{\text{max}}$, $K_m$ and $C_{\text{min}}$) by characterizing the uptake mechanism of a genotype is complicated by at least three factors [30]: (i) the plasticity of the system in response to the P status of the plant [89]; (ii) the differences in P uptake along roots [100]; and (iii) the dependence of P uptake on plant growth rate [101]. Thus there is general agreement that the efficiency of the uptake system is of minor importance for P acquisition from soils because transport of P to the root surface rather than the uptake is the limiting step [34]. Therefore it is less likely that selection for efficient P uptake kinetics will contribute to more efficient P acquisition from low-P soils, and accordingly, choosing this trait is not applicable in selecting safflower and sunflower for P uptake efficiency.

**Shoot demand for P on roots**

As mentioned above, roots have mainly to meet the nutrient demand exerted by shoot growth [77,101]. Hence, the shoot growth rate together with the required P concentration in the shoot is a measure of the demand, the shoot is putting on each root segment. Therefore, shoot demand (SD) on the root is interpreted as the P acquisition load imposed by shoot growth on each cm of root and is calculated by dividing the shoot growth rate by the average root length ($RL$), assuming that the roots of the plants grow exponentially: $GRs/RL = ((SW_{2}-SW_{1})/t_{2}-t_{1})X ((RL_{2}-RL_{1})/RL_{2}-RL_{1})$. Table 3 shows the shoot growth rate per unit of safflower and sunflower root length calculated during the first and second harvest when grown in sandy and loamy soils as affected by increasing P supplies. The higher SD ratio of safflower as compared
to sunflower at all respective P supplies and more pronouncedly at low P supply (Figure 4), was attributed to the lower values of root length/shoot dry matter ratio in safflower as compared to sunflower (Figure 2) not to a faster shoot growth rate of the former as compared to the later (Table 3). Since the shoot P concentration in safflower was less than that of sunflower (Table 5), and P root influx was found similar in both species (Figure 3), the higher P demand per unit root length of safflower as compared to sunflower could be attributed mostly to the lower values of root length/shoot dry matter ratio in safflower as compared to sunflower (Figure 2).

**P concentration and accumulation**

Phosphorus efficient crops, adapted to low P-supplying soils, are often characterized by low P requirements [83]. A number of crop species can grow normally with low tissue P concentrations due to efficient use of P among the major biochemical fractions (soluble-P, lipid-P, and residue-P) and was found to be more tolerant to low P conditions than that, which exhibited high P concentrations in the tissues [102]. The plant that maintains relatively low tissue concentration of P due to the efficient incorporation of the external P into residue-P [103], and because the vacuole acts as a P reservoir to maintain a constant cytoplasmic P concentration [104] are more tolerant to low P conditions.

Both species under investigation had the same internal P concentration in their plant parts and shoot at low and intermediate P supplies (Table 5) but this figure was less in safflower tissues as compared to sunflower when the crops were grown in high added P soil which could be explained by the dilution effect because of the large biomass production of sunflower plants [105,106]. Plant parts of both species had more P concentration when they were grown in sandy soils as those were cultivated in loamy soil because of the higher availability of P in sandy soil having low fixing ability of P as compared to loamy soil. As the nutritional status of the plant can be characterized by the P concentration in the dry matter, optimal plant growth requires P in the range of 0.3 to 0.5% of dry matter during the vegetative growth stage [12]. It is in agreement of our findings where the P deficient plants had P concentration in their plant parts and shoots lower these limits (Table 5). The productive efficiency of P for grain or seed is higher at early growth stages than at later stages because P is needed for tillering or branching. If sufficient P is absorbed at early growth stages, it will be redistributed to other growing organs [1,12,15,16,104].

Phosphorus supply significantly affected the P-concentration [P] (Table 5) and the accumulated P in tissues of the tested species was also reported in Brassica spp. [85], and for both species studied earlier [12,13]. More root growth [74] and more distribution of P to the roots as root are good sink for P under P-stress [107] are among reasons that reduce P concentration and accumulation in vegetative parts and make plants retain more P in their roots than shoots [85] in plants under P-starvation conditions.

**4.3. Soil Parameters**

Genotypic differences in P efficiency can be examined in field as well as in pot experiments with soil or with nutrient solution [108,109]. However, contradictory results may be obtained when a plant species or genotypes are evaluated using these three experimental systems due to different growth conditions. Results from pot trials with soil and especially from field trials can be not easily repeatable due to soil heterogeneity and complexity, and even using the same soil in pot experiments, the results are often not repeatable because the availability of nutrient can change during the soil storage. However, pot trials compared to field trials have the advantage that uniform growth conditions can be set regarding fertilization and soil homogeneity and also, that weather effects can be largely controlled. On the other hand, although nutrient solution experiments can be easily repeated, this can cover only part of the factors, which can be responsible for genotypic differences in nutrient efficiency by plants growing in soil. For instance, the root growth conditions and P uptake are substantially different between nutrient solution experiments and pot soil trials. Additionally, the relevance of different plant and non plant factors (soil) in P uptake would be different according to the experimental methodology used. To illustrate these differences between different growth conditions, it was reported that in early growth stages under field conditions, groundnut was not limited by low P soil (1.9 μM P), whereas maize only yielded 15-35% of its maximum yield [39]. In contrast with these findings, using the same plant material under flowing solution culture, the same researchers reported that maize was more P efficient since it was able to produce up to 90% of its maximum yield at only 1μM P concentration in the nutrient solution, whereas groundnut was inefficient, producing only 20% of its maximum yield at this P concentration [110].

Plant species and even varieties of same species differ in their ability to grow in soil low in nutrients [12,111]. Efficient species and cultivars are those that can utilize mobile, available, fixed nutrients in soil and can exploit more soil in order to maintain required rate of nutrient uptake by roots [111]. As discussed earlier, plant properties affecting uptake of nutrients from soil were kinetics of ion absorption by roots, the size of root system and morphological root properties [36]. Other properties are related to soil in which the supply of mineral nutrients to plants is the result of interactions between the nutrient availability in soil and the ability of plants to absorb this nutrient. Both soil and plant properties are therefore, important for the nutrition of plants.

Soils, characterized by having loamy texture may contains more than 20% iron or aluminium oxides in their clay particles, which “fix” or sorb rapidly large quantities of added phosphorus, transforming them into slowly soluble iron and aluminium phosphates that are not available to the plants [112]. Moreover, soils with a high P sorption capacity are able to absorb up to 5600 kg ha⁻¹ P until they are able to provide satisfactory crop growth [113]. On the other hand, diffusion of phosphorus “flux” through the soil to the plant’s roots, is - in many soils- the mechanism governing 90 to 98% of the P supply to the roots [114].

Mass flow and diffusion are the two mechanisms for nutrient movement from soil to root [115]. Diffusion is of fundamental importance for the availability of nutrients to plants growing in soil [82], because at low as well as
optimum soil nutrient concentrations, diffusion supplies much higher ion quantities from soil to roots than mass flow. Because a concentration gradient is required for diffusion to occur, the plant root takes up nutrients, lowers the nutrient concentration on its outside, and thus creates a gradient unless mass flow counteracts the process. Therefore the decrease of the nutrient concentration at the root surface is determined by the uptake properties of roots. Both species under study, in both soils had similar soil solution P at respective low and high P levels (Table 6) which indicate that both species have similar ability to deplete available P at respective P supplies. This was also reflected by the similar values of P influx among species in both soils at low P supplies (Figure 3). The extension of the depleted zone and the degree of depletion depends also on the nutrient mobility in the soil. Hence, ion diffusion from soil to root is basically the result of interactions between plant and soil. The plant efficient species is that can maintain higher nutrient influx by increasing diffusion towards roots due to its capability to use/release more soil nutrients than other species which resulted in the increase of concentration gradient [116].

The P supply represented as soil solution used in this investigation tested in pots before planting was 6.9, 19.6, and 256 μmol P L\(^{-1}\) in sandy soil and 0.2, 7.3, 419 μmol P L\(^{-1}\) in loamy soils after adding P levels (0, 0.2, and 1 g P pot\(^{-1}\)). The normal concentration of P in soil solution in the field, reported previously [117,118] was in the order of 0.32 – 19.37 μmol P L\(^{-1}\), and this concentration can be depleted rapidly by growing roots in soil. As solution P falls below its equilibrium concentration, it is replenished by labile P desorbed from loamy mineral surfaces adjacent to the roots [119]. Consequently, P moves from the adsorbed forms into solution and along a concentration gradient to the depletion zone of the root where the P concentration is low. However, in P-limited soils, the quantity of labile P may be insufficient to maintain P solution concentration against depletion by plant root. Thus this specific soil condition influences the movement of P toward the root surface because gradient is the driving force of diffusive P flux. On the other hand, P inflow depends on the concentration at the root surface, for that P depletion may imply severe restriction of P inflow into plants. It has been reported that the P concentration in soil solution (external P requirement) necessary to achieve maximum growth differs widely among crops. Using flowing solution cultures, a 25-fold difference in external P requirements among eight plant species and a 200-fold difference for other 18 species was reported [120]. In the field using adsorption isotherms, the external P requirements of a range of crops and vegetables were variable [119]. The external P requirements for several crop species varies in the range of 2 to 22 μmol L\(^{-1}\) [117]. Hence, at a low P concentration in soil solution, P efficient plants may be either those with a low external P requirement or those which are able to achieve their external requirement by developing of morphological and/or physiological root mechanisms.

Besides the relationship between P concentration and growth of plants, extractable P in the soil can be a measure of its availability. Ions not readily released from the soil matrix when the ambient solution concentration is low could be of minor importance to plants. But there is some evidence that phosphate fractions not detected by conventional soil test methods may play a role in supplying P to plants [79]. In utilizing these sources, plant roots function as more than sinks for diffusing ions. Both species depleted similar amounts of extractable P in sandy soil at both low and intermediate P supplies indicating a similar efficiency in solubilizing P from unavailable P pool in the soil, but sunflower was superior as compared to safflower at high P supply in sandy soil. The efficiency in accessing more P from unavailable P pool in loamy soil was more for sunflower as compared to safflower at all respective P supplies as the extracted P in soil hosting sunflower was lower than that for safflower at all respective P levels (Table 6).

The pH of the soil has a major influence on P solubility. Table 6 shows the pH of the soil as affected by plant species in increasing P supply in both soils. As an efficiency to solubilize scarce P under very low P supply, the pH of soil should be reduced, which was not applicable for both species and both soil types under investigation, where pH in both soils for both crops was significantly higher in deficient P supply under study. The pH was reduced by safflower at the highest P supply in sandy soil and at intermediate P supply in loamy soil. The pH value of soil hosting sunflower was reduced in sandy soil at the highest P supply and at both intermediate and high P levels in loamy soil. Sunflower reduced the pH of the sandy soil as compared to that of safflower but pH of the soil in loamy soil was not affected among plant species at low and intermediate P supplies. In contrary with our findings, it was reported that, the application of fertilizer did not produce any effect on the pH value [80]. It is well known that, the roots of many, if not all, plant species are able to exude substances which increase the solubility of soil P [121]. Exudates include protons that decrease the soil pH [122], reducing and chelating agents [123], organic acids [124] and phosphatase enzymes [125]. As the reduction of pH in the rhizosphere depends on the root activities to solubilize more P, the reduction of pH in soil at higher P supplies in both species and both soil was because the larger root systems of both species at higher P supplies (Table 1), therefore higher solubilizing efficiency, however they have less shoot demand of P on roots (Figure 4) under high P supplies in both soils.

5. Conclusion

New crops need to be developed, to acquire soil P more efficiently by focusing on species which represent uptake efficiency traits. Plant species vary in their P use efficiency at different P supplies and different soils by using different strategies related to uptake efficiency that could be used in selecting or breeding programs. P uptake efficiency depends on those factors related to plant parameters and those related to soil parameters. Efficient crop in terms of plant parameters increases root size, root length, specific root density, root-shoot ratio, nutrient influx and reduces root diameter, shoot growth rate and shoot demand on roots. The ability of the crop species to increase P solubility in the rhizosphere (P intensity and capacity) and depleting more soil solution and extractable P, along with acidifying the rhizosphere are considered as...
mechanisms of P uptake efficiency in terms of soil parameters. Under low P supplies (0, 0.2 external P), both species had similar agronomic P efficiency in both soils. Safflower had less absolute root length, higher relative root length, higher specific root density, thinner root diameter, less root shoot ratio in both soils. Safflower was superior as compared to sunflower in terms of P influx in both soil soils (except at 0 P supply in sandy soil where both species were similar). Safflower had higher shoot demand on root because of the less root length and higher shoot growth rate, although it had higher K influx. At very low P supply, both species depleted the same soil solution P in both soils and CAL-P in sandy soil, while sunflower depleted more CAL-P in loamy soil. Both species under study share both efficiency and inefficiency traits concerning both plant and soil parameters in different soils. Therefore using different measures of uptake efficiency parameters to differentiate plant species and genotypes to superior and inferior could be in some cases misleading.

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Competing Interests

None declared.

Abbreviations

P- Phosphorus, NUE- Nutrient use efficiency, CAL- Calcium acetate lactate, DM- Dry matter, RL- root length, RFW- root fresh weight, rgr root radius, RLV- specific root density, GRs- shoot growth rate, GRr- root growth rate, ln- influx, SW- shoot weight, t- time, SD- shoot demand, RSR- root shoot ratio.

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