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Effect of irrigation on some growth parameters of cup plant and dehydrogenase activity in soil

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Abstract: *Effect of irrigation on some growth parameters of cup plant and dehydrogenase activity in soil.* A field experiment carried out in two vegetation seasons in the years 2012 and 2013 on very light soil at Kruszyn Krajeński, in the vicinity of Bydgoszcz (Poland). Investigation of the influence of drip irrigation on some growth parameters of three- and four-year-old cup plant (*Silphium perfoliatum* L.) and on the activity of dehydrogenase participating at cycling in soil was examined. The cup plants were cultivated from the micropropagation seedlings. Experiments were performed as one factorial in four replications. The factor of the experiment was the following variants: O – without irrigation (control plots), D – with drip irrigation. Drip irrigation was scheduled according to tensiometers indications. Irrigation significantly increase the height of the plants, the length of internode, the thickness of the stalk, the fresh weight of the shoots, and the number of the leaves and flowers. Irrigation also increased the transpiration leaf area and the dry matter content. In the soil sampled under cup plant in 2012 there was found a greater activity of dehydrogenases in soil derived from non-irrigated objects.

Key words: drip irrigation, light soil, micropropagation, *Silphium perfoliatum*, soil enzymes

INTRODUCTION

Representing family *Asteraceae*, cup plant (*Silphium perfoliatum* L.) is a perennial reaching up to 2.5 m height. In the first

year of growth the plant creates a rosette, generative shoots develop in successive years. Due to the production of big over-ground mass, the plant demonstrates high water and irrigation requirements. Cup plant is a plant of temperate climate with little soil requirements. In the natural environment it occurs in the central and eastern part of the United States as well as in the south of Canada. The optimal growth and development of cup plant require sunny sites and the temperature of air about 25°C. It reacts to long-term periods of drought with dying of lower leaves and browning of buds. In an extreme case it stops growing, creates small flowers and a little amount of seeds (Stanford 1990, Kowalski and Wierciński 2004). Under the climatic conditions of Poland irrigation is the method of supplementing water deficits. Due to decreasing resources of disposable water for irrigation, more and more attention is paid to water-saving irrigation systems, the so-called microirrigation, including drip irrigation which is characterized with the high influence of the agricultural productivity effects (Pierzgalski and Jeznach 1993, Jeznach 2007 and 2009, Rolbiecki

and Rolbiecki 2008). The preliminary results of the present research by Figas et al. (2011) point to the fact that cup plant is a plant predestined for growing under supplemental irrigation.

The plant can be grown for feed, ornamental, honey-producing and energy-generating purposes (Majtkowski 2007, Piłat et al. 2007, Decourtye et al. 2010, Țiței et al. 2013, Wróbel et al. 2013, Jasinskas et al. 2014) and as it has low soil requirements, it can be recommended as a pioneer plant for the recultivation of degraded areas (Klimont 2007). The herbage of cup plant is also a potentially precious material for food and pharmaceutical industries. The extracts from that plant show pain-killing, anti-inflammatory properties, promote sweating, as well as restorative, antibacterial, antifungal, expectorant properties as well as they can lower the level of cholesterol (El-Sayed et al. 2002, Kowalski and Wolski 2003a,b, Kowalski and Wierciński 2004, Kowalski 2005, Jemiołkowska and Kowalski 2012).

Irrigation, by changing water conditions, affects the population of microorganisms in soil. Dehydrogenases (EC 1.1.1) catalyze the transferring of hydrogen from oxidized substrates to acceptors and the dehydrogenases are active in all soil microorganisms; aerobic as well as anaerobic. Assays for dehydrogenase activity in soil have often been used to achieve the value of the index of total soil microbial activity (Brzezińska 2004).

Numerous functional properties of the cup plant and climatic conditions similar to those it occurs naturally can enhance the cultivation of that species in the country. However, the source of the sowing material, however, which occurs in

Poland do not satisfy a growing interest of the breeders. The technology which facilitates obtaining vegetative propagation material is the micropropagation of plants in cultures *in vitro*.

The aim of the present research was to investigate the effect of drip irrigation on selected parameters of growth of three- and four-year-old cup plants (*Silphium perfoliatum* L.), derived from micropropagation grown on light soil as well as defining the activity of dehydrogenases in soil.

MATERIAL AND METHODS

Micropropagation *in vitro*

The initial research material involved the seeds of cup plant (*Silphium perfoliatum* L.), which came from the Botanical Garden of the National Center of Plant Gene Resources of the Institute of Plant Breeding and Acclimatization (IHAR) in Bydgoszcz (Poland). To induce germination, sterilized embryos were placed onto the Murashige and Skoog medium – MS (Murashige and Skoog 1962), diluted at the ratio of 1 : 1 and solidified with 0.7% agar. From six-week sterile seedlings there were isolated apical parts of shoots. The explants were put into the test-tubes on the MS regeneration medium contained growth regulators: 1 mg·dm⁻³ NAA (1-naphthaleneacetic acid) and 5 mg·dm⁻³ BAP (6-enzylaminopurine). After about 6–8 weeks the shoots were isolated and transferred onto MS rooting medium without growth regulators. *In vitro* culture of plants was conducted in a phytotron under controlled environmental conditions: a 16-hour photoperiod at a light intensity of 40 μmol·m⁻²·s⁻¹ and a constant temperature of 25 ±2°C.

The rooted plants were transferred into the mixture of sterile soil and perlite (1 : 1) and acclimatized under greenhouse conditions.

Field trial

The field experiment with drip irrigation of the cup plants was carried out in the years 2012–2013 at the very light soil (type – Mollisoil; texture – Fine sand) in Kruszyn Krajeński in the vicinity of Bydgoszcz (53°04'53" N, 17°51'52" E). The soil characteristic is shown in Table 1. The water reserve to soil depth of 1 m at field capacity was 87 mm and the available water quantity was 67 mm. The soil was characterized by the low organic matter content (1.5%). The cup plants were cultivated from the micro-propagation seedlings planted in 2010. The experiment was performed as a one-factorial in four replications. The factor of the experiment was the following variants: O – without irrigation (control plots), D – with drip irrigation. Drip irrigation was scheduled according to tensiometers indications, installed at 25 cm depth. The irrigation was started when the soil water pressure was up to -0.04 MPa. The irrigation water rates were strictly connected with the rain distribution and amounted to 116 and 108 mm for 2012 and 2013 respectively. The single rates of water ranged from 6 to 12 mm. Irrigation

was done with the drip line “T-Tape” with the distance among the drippers 30 cm. The agrotechnical practises and fertilization adopted were the standard used across the country. The mineral fertilization was applied at the rates of 500 kg N : P : K · ha⁻¹ at the ratio of 2 : 2 : 3. Doses of potassium (potash salt) and phosphorus (superphosphate) fertilization were dependent on the abundance of these nutrients in the soil. The nitrogen fertilization (ammonium nitrate) was supplied at three single rates. The area of single experimental plot was equal to 11 m². The seedlings were planted with a row spacing of 1 m. The plant spacing was 0.7 m.

The height and plant parameters of cup plant and dehydrogenases activity analysis

Measurements of the growth parameters were performed in the first decade of October, in each of the years of research. In the experiment rated the height of the plants (cm), the length of internode (cm), the fresh matter of the shoots (kg·plant⁻¹), dry matter of the leaves (%), the number of the leaves, the thickness of the stalk (mm), number of flowers on shoot and leaf area (cm²). Measurements of the stem thickness measured by caliper. The height of the plants was measured in cm from the soil surface to the top of the

TABLE 1. Physical properties of the soil

| Genetic horizon | Depth (cm) | Texture | Bulk density | | | Porosity | Moisture |
|-----------------|------------|---------------------|--------------------|-------|--------|----------|----------|
| | | | Specific density | temp. | actual | | |
| | | | Mg·m ⁻³ | | | % vol. | |
| Ap | 0–33 | slightly loamy sand | 2.290 | 1.426 | 1.324 | 42.2 | 10.2 |
| AC | 33–60 | loose sand | 2.680 | 1.620 | 1.591 | 40.6 | 2.9 |
| C | 60–150 | loose sand | 2.740 | 1.691 | 1.653 | 39.7 | 3.8 |

shoot apex. For leaf area measurement the digital planimeter was used. Samples of the leaves were oven-dried at 105°C and their dry weights were determined.

In the adequately prepared plant material the following were assayed: total organic carbon (TOC) and total nitrogen (TN) were determined with the analyser TOC Primacs provided by Scalar.

Soil samples were taken from the top of 0–30 cm using an auger during cup plant vegetation in August 2012 and 2013. Soils were, respectively. Then the air-dried soils were sieved using sieve with holes of below 2 mm in diameter. Meanwhile, the sampled soil water content was determined using an oven-drying method, so as to calculate soil enzymatic activities per g of dry soil. Dehydrogenase activity was measured using triphenyltetrazolium chloride as a substrate; samples were incubated for 24 h at 37°C with the Thalmann (1968) method.

Statistical analysis

The analysis of variance was performed using all the results applying FR-ANALWAR software based on Microsoft Excel (Rudnicki 2011). The significance of differences was evaluated by the Tukey's test and LSDs were calculated at a significance level of $\alpha = 0.05$.

RESULTS AND DISCUSSION

The mean height of cup plants on control plots (without irrigation) in the research period of 2012–2013 was equal to 98.63 cm (Table 2). Drip irrigation had a significant effect on an increase in the mean plant height by 35.08 cm (35.6%). The increase in the cup plant height was due to an increase in the length of inter-

nodes. Their mean length was 14.23 cm, both on the objects irrigated and non-irrigated. Drip irrigation increased the length of internodes by 45.3%.

Drip irrigation significantly affected the fresh weight of shoots in 3- and 4-year-old cup plants (Table 2). The experimental factor applied resulted in, on average over the research period of 2012–2013, its significant increase by 2.56 kg. The mean fresh weight of shoots from the plants collected from the irrigated objects was equal to 4.41 kg·plant⁻¹. Similar tendencies of the growth of fresh weight of leaves due to the application of drip irrigation was found in the case of one-year-old cup plants (Figas et al. 2011).

In the experiment the content of dry weight in the leaves of cup plant accounted for, on average in the research years and on average for the variants of the experiment of 19.17% (Table 2). The application of irrigation significantly increased the content of dry weight in leaves by 35.4%. Figas et al. (2011) performing research with the irrigation of one-year-old cup plants recorded a significant increase in the value of that parameter by 17.6%. In this experiment it ranged from 14.33 to 22.95%, respectively on the control objects and the drip-irrigated ones. Similar values of the content of dry weight in the leaves of cup plant are reported by Stanford (1990), accounting for 21.5%. The tendencies to increase the content of dry weight in the irrigate plants coincide with the reports on other plant species (Pronk et al. 2005, Wichrowska et al. 2007).

The mean number of leaves per stem was equal to 16.45 (Table 2). The number of leaves on control objects was 14.34 and on the objects treated with drip

TABLE 2. Influence of drip irrigation on the height and plant parameters of cup plant (*Silphium perfoliatum* L.) in the years 2012 and 2013

| Parameter | Year | | | | | | Mean |
|--|------------------------------|--------|--------|--------|-----------------------------|--------|--------|
| | 2012 | | 2013 | | Mean in the years 2012–2013 | | |
| | D | 0 | D | 0 | D | 0 | |
| Height of plants (cm) | 101.50 | 68.50 | 170.25 | 121.75 | 133.71 | 98.63 | 116.17 |
| | LSD _{0.05} – 12.93 | | | | | | |
| Length of internode (cm) | 14.91 | 10.01 | 18.81 | 13.19 | 16.86 | 11.60 | 14.23 |
| | LSD _{0.05} – 1.900 | | | | | | |
| Number of the leaves (pcs) | 19.67 | 15.17 | 17.50 | 13.50 | 18.56 | 14.34 | 16.45 |
| | LSD _{0.05} – 3.234 | | | | | | |
| Fresh matter of the shoots (kg plant ⁻¹) | 3.90 | 1.75 | 4.92 | 1.95 | 4.41 | 1.85 | 3.13 |
| | LSD _{0.05} – 0.331 | | | | | | |
| Dry matter of the leaves (%) | 22.95 | 18.23 | 21.15 | 14.33 | 22.05 | 16.28 | 19.17 |
| | LSD _{0.05} – 3.354 | | | | | | |
| Number of flowers on the shoot (pcs) | 15.67 | 4.83 | 93.75 | 31.25 | 54.71 | 18.08 | 36.40 |
| | LSD _{0.05} – 4.85 | | | | | | |
| Leaf area (cm ²) | 409.00 | 298.00 | 425.25 | 320.25 | 417.13 | 309.13 | 363.13 |
| | LSD _{0.05} – 70.918 | | | | | | |
| Thickness of the stalk (mm) | 8.50 | 6.17 | 12.25 | 8.25 | 10.38 | 7.21 | 8.80 |
| | LSD _{0.05} – n.s. | | | | | | |
| Nitrogen content in the leaves (%) | 1.61 | 1.60 | 2.21 | 2.35 | 1.97 | 1.91 | 1.94 |
| | LSD _{0.05} – n.s. | | | | | | |
| Carbon content in the leaves (%) | 41.71 | 41.24 | 25.77 | 27.60 | 33.74 | 34.42 | 34.08 |
| | LSD _{0.05} – n.s. | | | | | | |

0 – control (without irrigation), D – drip irrigation, LSD – the lowest significant difference (Tukey's confidence half-interval) for $p = 95\%$ ($\alpha = 0.05$), n.s. – non-significant differences.

irrigation – 18.56. Stanford (1990) claims that the plant can have 8 to 14 pairs of stem leaves. The increase in the number of leaves as a result of the irrigation used was significant and it was equal to 4.22. The regularity recorded coincides with earlier research results on the drip irrigation of the one-year-old cup plants (Figas et al. 2011) as well as other nursery plants (Rolbiecki et al. 2005, Klimek et al. 2009).

The drip irrigation applied increase the average leaf area three- and 4-year-old cup plants as compared with the control objects on average by 108 cm² (Table 2). The leaf area measured from control objects was equal to 309.13 cm², on the irrigated objects was higher and equal to 417.13 cm². Similar tendencies in affecting the leaf area were reported by Rolbiecki et al. (2005), Klimek et al. (2009) and Figas et al. (2011).

Over 2012–2013 years the mean number of flowers per plant both on the irrigated and non-irrigated objects was equal to 36.40 (Table 2). In this experiment drip irrigation significantly increased the number of flowers per plant by as much as 202.6% (36.63). The result is confirmed by Koszański et al. (2008)

on the irrigation of northern highbush blueberry (*Vaccinium corymbosum* L.) as a result of which the authors recorded a two-fold increase in the mean number of flowers on irrigated objects (Table 3).

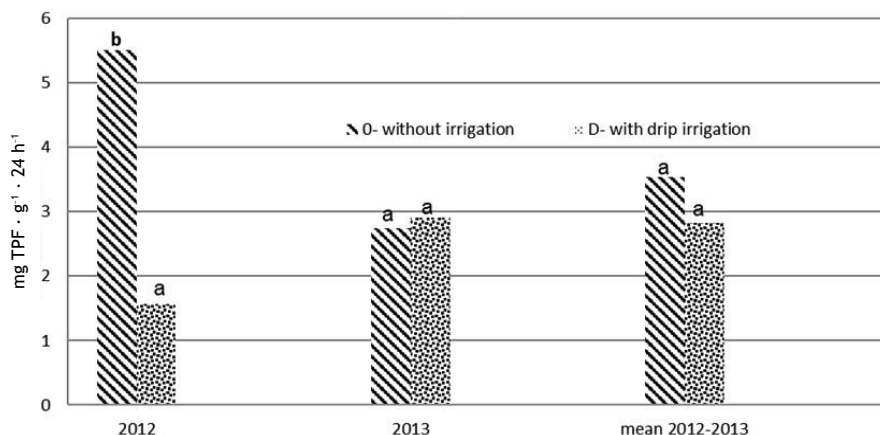
In the analyzed two years of this research there was found no effect of drip irrigation on the content of organogenic elements such as carbon and nitrogen in cup plants (Table 2). As reported by Trawczyński (2012), the irrigation treatments can affect the yield size more considerably than the content of nutrients in plants.

The water-air conditions in soil regulate the activity of microorganisms, affecting their type of metabolism as well as a number of adaptations they must develop as a response to the existing environmental conditions (Gleeson et al. 2008). Soil dehydrogenase activity increases under anaerobic conditions (Brzezińska 2004).

In the soil sampled under cup plant in 2012 there was found a greater activity of dehydrogenases in the soil from the non-irrigated objects (Fig. 1). The activity was 71% higher as compared with the objects exposed to drip irrigation. Such a high difference of the index of micro-

TABLE 3. Air temperature and rainfall data during the two vegetation period of cup plant by monthly measurements in the period of 1981–2010 and the years 2012 and 2013

| Specification | Air temperature (°C) | | | | | |
|-----------------------|----------------------|-------|-------|------|------|-------|
| | V | VI | VII | VIII | IX | V–IX |
| 2012 | 14.5 | 15.2 | 18.8 | 17.6 | 13.3 | 15.9 |
| 2013 | 14.2 | 14.4 | 18.9 | 18.1 | 10.7 | 15.3 |
| Long period 1981–2010 | 13.1 | 16.0 | 18.5 | 17.9 | 13.2 | 15.7 |
| × | Rainfall (mm) | | | | | |
| 2012 | 25.4 | 133.8 | 115.6 | 51.8 | 25.1 | 351.7 |
| 2013 | 91.7 | 49.3 | 79.0 | 56.6 | 64.1 | 340.7 |
| Long period 1981–2010 | 49.3 | 52.8 | 69.8 | 62.6 | 46.0 | 280.5 |



a, b – values followed by the same letter are not significantly different at $p < 0.05$

FIGURE. Dehydrogenase activity in the instigated soil under cup plant in the years 2012 and 2013

biological soil activity can be due to the occurrence of anaerobic conditions in the soil sampled from control objects with cup plant. In 2012 most rainfall coincided in the summer season (June – August), while in spring the rainfall did not practically occur at all (Table 3). Water stress caused by periodical drought and then rapid soil environment irrigation get quickly reflected in the nature of the community of microorganisms (Young and Ritz 2000). According to Gleeson et al. (2008), under soil irrigation conditions, after a long period of drought, there occurs decay of the cells of microorganisms connected with a release of intracellular enzymes from them. After the spring period of low rainfall, in June there was reported an intensive soil irrigation (rainfall of 133.8 mm) and intensified microbiological activity of soil. In the irrigate soil the stabilizing availability of water did not affect so intensively on the activity of dehydrogenases in the soil. Xiang et al. (2008) found a five-fold higher activity of dehydrogenases in the flooded soil, as compared with the con-

ditions of dry soils. Such a direction of transformations points to an increased share of anaerobic bacteria, the habitat of which are moist soil environments in the formation of the activity of dehydrogenases of soils (Pascual et al. 2007). In the successive year of the vegetation period of cup plant, there was found no significant effect of drip irrigation on the activity of dehydrogenases in soil (Fig.).

The activity of dehydrogenases in the soil collected from the objects exposed to drip irrigation, was 6% lower, as compared with the control. Most probably high rainfall (91.7 mm) and temperature (14.2°C) in May at Kruszyn Krajeński eliminated the stabilizing effect of irrigation on the development of soil microflora, which did not cause such a high variation in the activity of dehydrogenases.

CONCLUSIONS

1. Under Bydgoszcz climatic conditions the plant is capable for the cultivation under field conditions on light soil and it shows a significant increase in the

parameters of growth of aboveground parts as a result of drip irrigation.

2. It was shown that the effect of drip irrigation and rainfall during wet plant vegetation cup plant in 2012 increased on the activity of dehydrogenases which are a sensitive indicator of the changes which occur in the environment of soil microorganisms. The air-water status in soil plays an important role in the regulation of the composition and metabolic activity of soil microorganisms.

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Streszczenie: Wpływ nawadniania na wybrane parametry wzrostu roznika przerośniętego i zawartość dehydrogenazy w glebie. Doświadczenie polowe przeprowadzono w dwóch sezonach wegetacyjnych w latach 2012 i 2013 na glebie bardzo lekkiej w Kruszynie Krajeńskim w pobliżu Bydgoszczy. Badano wpływ nawadniania na niektóre parametry wzrostu 3- i 4-letnich roślin roznika przerośniętego (*Silphium perfoliatum* L.)

oraz na aktywność dehydrogenaz uczestniczących w przemianach zachodzących w środowisku glebowym. Doświadczenie założono jako jedno-czynnikowe w czterech powtórzeniach. Czynnikiem stanowiącym źródło zmienności było nawadnianie kropłowe. Zastosowano dwa warianty doświadczenia: O – bez nawadniania (kontrola), K – nawadnianie kropłowe, przeprowadzane na podstawie wskazań tensjometrów ($-0,04$ MPa). Materiałem do badań były rośliny roznika przerośniętego (*Silphium perfoliatum* L.), które używano w procesie mikrorozmnażania. Nawadnianie istotnie zwiększyło wysokość roślin, długość międzywęźli, grubość łodygi, świeżą masę pędów oraz liczbę liści i kwiatów. Nawadnianie zwiększyło również powierzchnię transpiracyjną liścia oraz zawartość suchej masy. Większą aktywność dehydrogenaz spod uprawy roznika stwierdzano w glebie pobranej w 2012 roku z obiektów nie-nawadnianych.

Słowa kluczowe: nawadniania kropłowe, gleba lekka, mikrorozmnażanie, *Silphium perfoliatum*, enzymy glebowe

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