

NECTAR AND POLLEN PRODUCTION AND INSECT VISITATION ON ORNAMENTALS FROM THE GENUS *HOSTA* Tratt. (ASPARAGACEAE)

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Abstract

Properly arranged ornamental gardens in both urban and agricultural landscapes can be of a benefit to bees. In this study, we observed the flowering phenology, nectar, and pollen production of the ornamental *Hosta* species and varieties (*H. sieboldiana* Engler, *H. capitata* Nakai, *H. crispula* Maekawa, *H. fluctuans* Maekawa, syn. *H. sieboldiana* var. *fluctuans* hort., *H. undulata* var. *univittata* Miquel (Hylander), syn. *H. univittata*). Our experiment was conducted in the 2012 - 2014 time period, at the UMCS Botanical Garden in Lublin, Poland (51° 14' N, 21° 34' E). The total sugar yield varied almost 5-fold among *Hosta* ornamentals; the lowest amount was calculated for *H. fluctuans* (2.31 g per 10 m²) and the highest for *H. capitata* (11.80 g per 10 m²). The average pollen yield was from 0.24 g per 10 m² (*H. undulata* var. *univittata*) to 9.53 g per 10 m² (*H. capitata*). Pollen grains were bilaterally symmetrical, and large-sized. In polar view, they were prolatum (shape index 1.33 - 1.61), while in equatorial view, oblatum (shape index 0.5 - 0.7). *Hosta* species can complete the summer pasture mainly for bumblebees. In four of the five *Hosta* representatives, bumblebees accounted for 60 - 70% of the total visits. The honeybee predominated only on the flowers of *H. capitata* (54 - 71%). Solitary bees were rarely observed (3 - 9% of visits). The phenotypic traits of *Hosta* flowers (the accumulation of nectar in the deep and narrow perianth tube) reduced the access to nectar reward and restricted an array of insect visitors.

Keywords: bee pasture, bumblebees, honeybees, *Hosta*, nectar, pollen.

INTRODUCTION

The forage base for pollinators is subjected to continuous changes and nutritional stress is considered an important factor that impact on the decline of pollinators (Alaux et al., 2010; Gonzalez-Varo et al., 2013). The promotion of flowering plants that attract pollinators by nectar and/or pollen has been well known to help to overcome the problem (Garbuzov et al., 2015). Improvement of bee pastures can support pollinators and may counteract the alarming reduction of their diversity and abundance. One of the ways to help pollinators is to cultivate ornamental garden plants (Kołtowski, 2006; Masierowska, 2006; Garbuzov et al., 2015). Properly arranged ornamentals can benefit bees in urban areas (Matteson et al., 2012) as well as enhance pollinators in the surrounding of intensively managed farmlands (Samnegård et al.,

2011). The favourable effect of the presence of ornamental gardens on pollination in adjacent crops has been recently documented (Pereira-Peixoto et al., 2014). However, the usefulness of ornamentals for pollinator-friendly gardens differs considerably between species or even between cultivars (Denisow and Strzałkowska-Abramek, 2013). To maximize the efforts for the design of pollinator-friendly gardens we primarily require the knowledge of floral reward accessibility and the value of nectar and pollen provided by flowers. In addition, the flowering biology and ecology observations are necessary to help the gardener choose insect-friendly flowers. Non-native plants particularly need great attention to evaluate if they are appropriate for the pollinator groups present outside of their natural environment (Wróblewska and Stawiarz, 2012; Denisow and Strzałkowska-Abramek, 2013).

In this study, we determined the floral reward value of the *Hosta* species and varieties. These are popular ornamentals which are propagated in the modern garden design due to their versatility and usefulness for different arrangements (Grabowska and Kubala, 2012). In particular, we examined (i) blooming biology, (ii) nectar secretion, (iii) pollen production, (iv) pollen grain size and shape (useful for identification of pollen grains in microscopic analysis of bee products), and (v) the spectrum of insect visitors.

MATERIAL AND METHODS

Study site and study species

Our experiment was conducted in the 2012 - 2014 time period, in the Botanical Garden of Maria Curie-Skłodowska University in Lublin, Poland (51° 11' N, 22° 28' E). The plants singled out for the study were *H. sieboldiana* Engler, *H. capitata* Nakai, *H. crispula* Maekawa, *H. fluctuans* Maekawa, syn. *H. sieboldiana* var. *fluctuans* hort., *H. undulata* var. *univittata* Miquel (Hylander), syn. *H. univittata*. The experimental individuals were grown in the collection of ornamental plants by the pond, under the shade of surrounding trees.

The genus *Hosta* includes about 45 species native to northeast Asia (China, Korea, Japan), they are mostly grown in wetlands and moist meadows. Several species have been introduced to Europe and widely cultivated as shade-tolerant foliage plants (Grabowska and Kubala, 2012). The genus is currently placed in the family Asparagaceae (APG III). *Hosta* species are herbaceous perennials, very popular, and frequently used as border and ground cover plants, suitable for planting near water bodies (Marcinkowski, 2007; Grabowska and Kubala, 2012). The advantage is that they grow well in a polluted city environment. In Poland, hostas are commonly cultivated. *Hosta sieboldiana* develops huge round, heavy textured greenish-blue leaves and light blue flowers. *Hosta capitata* has ovate leaves and purple flowers. The leaves of *H. crispula* are dark green, broadly margined with creamy-white, and the flowers are pale lavender. The leaves of *H. fluctuans* possess particularly protruding veins, and the flowers are light lavender. *Hosta undulata* var. *univittata* develop ovate, strongly twisted, deep green leaves, striped creamy-white in the centre; the flowers are light purple (Marcinkowski, 2007). In the natural environment, hostas are known to attract insect visitors (Takahashi et al., 1994).

Flowering

The protocols described by Denisow (2009; 2011) were applied during flowering observations. The phenology of flowering was established by means of the onset and length of the blooming period. To determine the flower lifespan, the buds ($n = 6 - 10$ per year) were randomly chosen and marked on different individual plants ($n = 3 - 4$). The flower lifespan was defined as the period from bud opening until perianth closing (observed only in 2012 - 2013). The number of flowers per raceme ($n = 24 - 30$) and the number of inflorescences per individual ($n = 5 - 7$) was established. The data were converted to the number of flowers per 10 m² of the surface and used to estimate the total nectar and pollen yield.

Nectar secretion

Nectar production was only measured in 2012 and 2013. Nectar secreted in single flowers was collected using the pipette method (Jabłoński, 2002). Prior to nectar collecting, flowers were covered by tulle isolators to exclude insect visitors. Nectar collection was conducted in 3 - 5 replications during the blooming period. In each replication, 4 - 6 samples were collected. A single sample contained nectar from 4 - 7 flowers collected from 3 - 4 different individual plants. The total sugar concentration was measured with an Abbe refractometer. Nectar volume and sugar concentration were used to calculate the total sugar mass in each sample. Relevant calculations allowed us to determine the amount of sugars produced per 10 flowers (in mg) and per 10 m² (in g). We excluded data for *H. undulata* var. *univittata* (during the study course some isolators were destroyed, and the remaining samples seemed to be unrepresentative).

Pollen production

For the full-blooming phase, pollen production was determined using the ether-ethanol method described in detail by Denisow (2011). The anther dry mass and the mass of produced pollen was assessed. Anthers were dissected and placed in tarred vessels prepared for this purpose. Samples were collected every year in four replications ($n = 60$ anthers in each replication). The samples were inserted in the ELCON CL 65 dryer, at ca. 33°C. The pollen from anthers was rinsed once with pure ether (1 - 2 mL) and then 2 - 3 times with 70% ethanol (2 - 8 mL). The mass of produced pollen was calculated for 10 flowers (in mg), and per 10 m² (in g).

Pollen grain dimensions were determined in glycerol-gelatine slides (Erdtman, 1954) in 2013 and 2014. The lengths of the polar axis (P) and two equatorial axes with distinct lengths (longitudinal = E_1 and transversal = E_2) were determined ($n = 4 \times 50$ per year). The shape coefficients (P/E_1 and E_1/E_2) were defined. These observations were conducted using a Nikon Eclipse 200 light microscope.

Insect activity

Insects visits were recorded from 7:00 a.m. until 6:00 p.m. (GMT + 2.00 h) in two (three) hours intervals on random plots ($n = 3$; 1 m² per species). These observations were conducted for three consecutive days, at the full-bloom phase of the species. During each census of observation (3 - 6 min), the total number of visiting insects was recorded and the type of forage (nectar vs. pollen) noted.

Weather conditions

In 2012 and 2013, air temperatures in Lublin in the period of blooming of *Hosta* species were 2.9 and 0.8°C higher than the long-term average. Total precipitation was approx. 26% lower than normal in 2012, while it was on the average level in 2013.

Statistical analysis

Data are presented as means with SD. The Kruskal-Wallis H-test was used to determine the species effect for the flower life span (non-normally distributed data). Parametric statistical ANOVA was applied for the number of flowers and nectar and pollen traits. A post hoc comparison of the means was tested by the HSD Tukey's test. The level of statistical significance for all analyses was at $P = 0.05$. All analyses were performed using Statistica ver. 6.0 (StatSoft Poland, Kraków).

RESULTS

The blooming of the *Hosta* species and varieties took place from the last decade of June to the first week of August. *Hosta sieboldiana* was the first to bloom, and was followed by *H. fluctuans*, *H. capitata*, *H. crispula*, and by *H. undulata* var. *univittata* (Tab. 1). The blooming duration for particular species differed 7 - 14 days between the years of the study. Irrespective of the year, the shortest blooming was characteristic for *Hosta crispula* (only 16 - 20 days), the longest for *H. capitata* (25 - 36 days). In 2014, the flowering onset of every species/variety was accelerated (10 - 14 days) as compared to the flowering times recorded in the previous years.

The flowers of *Hosta* representatives are arranged in raceme inflorescences. Within an inflorescence, flowering progresses acropetally from the bottom to the top positioned ones, with little flowering overlap among flowers. The individual flowers are perfect, actinomorphic. Perianth form a narrow tube, 10 - 30 mm in length. *Hosta* flowers opened before 8:00 a.m. The life span of the single flower in the raceme did not differ significantly in individual taxa (Kruskal-Wallis test: $H = 12.11$, $P = 0.114$). The longevity of individual flowers ranged between 20-38 hours.

The number of flowers per inflorescence and the number of stems per unit area was species-related ($F_{4,104} = 4.21$, $P = 0.043$ and $F_{4,36} = 2.21$, $P = 0.013$, respectively). The lowest number of flowers in inflorescences was noted for *H. undulata* var. *univittata* and *H. crispula*, and the highest for *H. sieboldiana* (Tab. 1). The number of flowers per inflorescence was not affected by the year of observation in any of the studied taxa. Among taxons, disparities in the number of flowers produced per 10 m² were 2 - 7 fold.

The *Hosta* flowers attracted insect visitors by primary attractants (nectar and pollen) and secondary attractants (shape - funnel-bell; colour - white, lavender, light violet; soft scent). The fused perianth form a long tube (15 - 20 mm), which is narrow at the base. Nectar is accumulated and hidden in the narrow tube. The intermediate flower part is formed like a broad tube. The nectar-secreting tissue is located in three septal nectaries in the flowers of *Hosta* species and varieties. The nectar is exudated outside the pistil via outlets and is accumulated in the spaces between the ovary and the staminal filaments. The level of secreted nectar usually reaches to 1/4 - 1/2 of the narrow tube. Nectar was secreted already in the bud. In the first hours of the life-span, the pollen is released from the anthers and is presented to the insect visitors, pollen exposition under sunny and dry weather lasts for about 5 - 7 hours per flower. The sugar concentration in nectar differed between studied *Hosta* representatives ($F_{3,54} = 3.11$, $P = 0.021$). The sugar concentration in the nectar of *H. fluctuans* was almost 2-fold higher compared with that of *H. crispula* (Tab. 2). No species effect was found for the sugar-mass, produced in flowers ($F_{3,54} = 21.5$, $P = 0.096$). The sugar mass differed considerably between years in the flowers of *H. sieboldiana* ($F_{1,8} = 2.11$, $P = 0.047$) and *H. capitata* ($F_{1,8} = 3.16$, $P = 0.031$). The average sugar yield per 10 m² varied almost 2-5-fold among *Hosta* representatives (2.31 - 11.80 g).

Table 1.

Phenology of flowering and abundance of blooming for *Hosta* representatives studied in the 2012 - 2014 time period. Mean values \pm SD are given

Species / variety	Year	Flowering		Number of flowers per inflorescence		Number of inflorescences per 1 m ²		Number of flowers per 10 m ² thous.	
		Date	Length (days)	Mean	\pm SD	Mean	\pm SD	Mean	\pm SD
<i>H. sieboldiana</i>	2012	3.07 - 25.07	23	30.9 _a	4.8	7.1 _a	4.0	2.19	0.9
	2013	7.07 - 26.07	20	37.4 _a	11.8	7.8 _a	2.8	2.92	1.1
	2014	19.06 - 16.07	28	41.2 _a	9.4	9.7 _b	2.1	4.00	1.5
	Mean		23.7	36.5 _c		8.2 _A		3.04	
<i>H. fluctuans</i>	2012	4.07 - 27.07	24	24.8 _a	3.9	6.2 _a	3.1	1.54	1.8
	2013	9.07 - 28.07	20	30.0 _a	6.8	5.8 _a	2.4	1.74	1.2
	2014	23.06 - 21.07	29	28.3 _a	5.3	6.8 _a	1.3	1.92	2.0
	Mean		24.3	27.7 _B		6.3 _A		1.73	
<i>H. capitata</i>	2012	7.07 - 3.08	28	26.4 _a	3.2	30.3 _a	1.5	8.00	4.2
	2013	8.07 - 01.07	25	30.6 _a	4.2	29.8 _a	3.2	9.12	3.1
	2014	20.06 - 25.07	36	29.5 _a	4.9	32.8 _a	5.4	9.68	5.5
	Mean		27.7	28.8 _B		31.0 _C		8.93	
<i>H. crispula</i>	2012	8.07 - 26.07	19	22.1 _a	8.2	11.2 _a	2.4	2.48	1.1
	2013	12.07 - 27.07	16	27.2 _a	10.6	14.3 _{ab}	1.3	3.89	0.8
	2014	28.06 - 17.07	20	25.7 _a	8.0	15.8 _b	3.7	4.06	2.1
	Mean		18.3	25.0 _{AB}		13.8 _B		3.48	
<i>H. undulata</i> var. <i>univittata</i>	2012	10.07 - 29.07	20	16.6 _a	3.2	13.2 _a	4.5	2.19	2.3
	2013	14.07 - 30.07	17	19.0 _a	2.7	10.9 _a	3.2	2.07	1.7
	2014	26.06 - 21.07	26	18.2 _a	2.5	11.2 _a	2.6	2.04	2.0
	Mean		21.0	17.9 _A		11.8 _B		2.10	

Means within columns with the same small letter do not differ significantly between years within a species, and the means with the same capital letter do not differ significantly between species at $P < 0.05$, based on HSD Tukey's test.

Table 2.

Concentration of nectar and amount of sugars produced by *Hosta* representatives during the two study years. The mean values \pm SD are given

Species	Year	Number of examined flowers	Concentration of sugars in nectar (%)		Total sugars yield			
			Mean	\pm SD	per 10 flowers (mg)		g /m ²	
<i>H. sieboldiana</i>	2012	125	18.5 _a	5.5	9.4 _a	3.2	2.06	0.7
	2013	103	23.6 _a	2.4	17.1 _b	2.5	4.99	1.3
	Mean		21.05 _{AB}		13.2 _A		3.53	
<i>H. fluctuans</i>	2012	110	32.4 _a	6.3	13.8 _a	4.3	2.12	1.4
	2013	120	37.2 _a	4.8	14.3 _a	2.6	2.49	1.6
	Mean		34.80 _B		14.0 _A		2.31	
<i>H. capitata</i>	2012	121	18.5 _a	8.5	9.2 _a	3.8	7.36	2.8
	2013	108	26.6 _b	6.9	17.8 _b	5.2	16.23	6.2
	Mean		19.85 _A		13.5 _A		11.80	
<i>H. crispula</i>	2012	115	15.3 _a	7.1	16.6 _a	5.4	4.11	0.5
	2013	135	20.5 _b	5.8	15.3 _a	3.9	5.95	2.7
	Mean		17.90 _A		15.9 _A		5.03	

ANOVAs procedures were performed separately for each analysed feature. Means with the same small letter do not differ significantly between years within a species, and the means with the same capital letter do not differ significantly between species, at $P < 0.05$, based on HSD Tukey's test.

Table 3.

Pollen production and pollen viability of *Hosta* representatives in the 2012 - 2014 time period.
The mean values \pm SD are given

Species / variety	Year	Pollen production				
		per 10 flowers (mg)			g/m ²	
		Min - max	Mean	\pm SD	Mean	\pm SD
<i>H. sieboldiana</i>	2012	1.1 - 5.7	3.6 _a	3.2	0.79	0.6
	2013	1.5 - 4.1	2.9 _a	2.6	0.85	0.5
	2014	1.2 - 5.0	3.6 _a	1.4	1.44	0.7
	Mean		3.4 _B		1.02	
<i>H. fluctuans</i>	2012	0.6 - 5.9	3.9 _a	2.6	0.60	0.2
	2013	1.7 - 5.0	3.6 _a	1.0	0.63	0.3
	2014	1.2 - 6.0	4.4 _a	1.8	0.85	0.5
	Mean		4.0 _B		0.69	
<i>H. capitata</i>	2012	4.2 - 18.1	9.7 _a	3.5	7.76	1.9
	2013	3.5 - 14.7	11.7 _a	5.7	10.67	4.3
	2014	4.3 - 17.0	10.5 _a	4.5	10.16	4.9
	Mean		10.6 _D		9.53	
<i>H. crispula</i>	2012	1.9 - 8.5	6.2 _a	4.8	1.53	3.6
	2013	1.3 - 6.9	7.5 _b	3.8	2.92	4.6
	2014	2.2 - 9.4	5.9 _a	2.3	2.40	5.3
	Mean		6.5 _C		2.28	
<i>H. undulata</i> var. <i>univittata</i>	2012	0.3 - 1.6	0.8 _a	0.5	0.18	0.2
	2013	0.8 - 2.0	1.6 _b	1.0	0.33	0.1
	2014	0.2 - 1.6	1.1 _b	0.5	0.22	0.1
	Mean		1.2 _A		0.24	

ANOVAs procedures were performed separately for each analysed feature. The means with the same small letter do not differ significantly between years within a species and the means with the same capital letter do not differ significantly among species, at $P < 0.05$, based on HSD Tukey's test.

Table 4

Morphological characters of pollen grains of five *Hosta* species (the mean for 2013 - 2014)

Species	Length of axis (μ m)						Shape index	
	Polar (P)		(E ¹)		Equatorial transverse		P/E ¹	E ¹ /E ²
	(E ²)	Mean (\pm SD)	Min - max	Mean (\pm SD)	Min - max	Mean (\pm SD)		
<i>H. sieboldiana</i>	43.4 - 59.9	52.4 (6.7)	78.4 - 107.6	95.4 (8.6)	44.2 - 71.6	59.2 (10.0)	0.5	1.6
<i>H. fluctuans</i>	32.5 - 58.8	44.4 (10.6)	66.2 - 88.1	80.0 (7.6)	43.5 - 68.2	53.7 (8.7)	0.6	1.5
<i>H. capitata</i>	30.7 - 51.6	41.2 (6.6)	65.9 - 93.7	81.9 (8.0)	38.3 - 81.7	51.4 (10.9)	0.5	1.6
<i>H. crispula</i>	44.6 - 67.1	53.9 (8.4)	64.4 - 90.7	82.1 (8.3)	47.4 - 69.3	61.7 (6.5)	0.7	1.3
<i>H. undulata</i> var. <i>univittata</i>	44.9 - 50.6	48.7 (2.5)	66.5 - 103.4	85.8 (9.0)	43.4 - 66.6	56.1 (6.5)	0.6	1.5

E - equatorial longitudinal axis



Fig. 1. Light micrographs of the pollen grains of *Hosta capitata* in a. equatorial view b. distal polar view and c. proximal polar view; insect visitors d. *Apis mellifera*, e. *Halictus* sp.

The flowers of the *Hosta* species contained 6 stamens that extended over six lobes of the perianth and curved inward. The anther size differed between species ($F_{4,58} = 7.6$, $P = 0.022$) and consequently, the mass of the pollen produced in flowers ($F_{4,58} = 4.8$, $P = 0.012$). The amount of pollen produced was the highest in the flowers of *H. capitata*, and the lowest in *H. undulata* var. *univittata* (Tab. 3). The average pollen yield per 10 m² was from 0.24 g (*H. undulata* var. *univittata*) to 9.53 g (*H. capitata*).

Pollen grains of the *Hosta* species were bilaterally symmetrical (Fig. 1a - c). The length of the polar axis (P) ranged from 30.7 µm to 67.1 µm (Tab. 4). The longitudinal equatorial axis (E₁) was the longest for *H. sieboldiana* (mean = 95.4 µm), the shortest for *H. fluctuans* (mean = 80.0 µm). In polar view, the E₁/E₂ ratio ranged between 1.33 - 1.61. In equatorial view, the ratio of the polar to equatorial longitudinal axis (P/E₁) ranged between 0.5 and 0.7.

On average, the most frequent insect visitors recorded on the flowers of *H. sieboldiana*, *H. fluctuans*, *H. crispula*, and *H. undulata* var. *univittata* were bumblebees, responsible for 60 - 70% of total insect visits (Fig. 2). We observed only *B. terrestris* (L.), *B. lapidarius* (L.), *B. pascuorum* (Scopoli), and *Bombus hortorum* (L.). Regardless of the study year, the honeybee predominated on the flowers of *H. capitata*, comprising 54 - 71% of

the visits (Fig. 1d). Solitary bees were rare visitors (3 - 9%) and were almost exclusively interested in pollen (Fig. 1e). The activity of bumblebees on *Hosta* flowers started in the early morning hours about 7:00 a.m. and was relatively constant throughout the entire day until 6:00 p.m. Most honey bees and solitary bees were observed on the flowers between 10:00 a.m. and 3:00 p.m. Honeybees collected both nectar and pollen, with nectar being the more attractive goal.

DISCUSSION

The value of the nectar and pollen of the *Hosta* species for insect visitors was not empirically evaluated, therefore our results have no equivalent in the literature data. If arranged together in ornamental gardens, the species will provide nectar and pollen from the second decade of June (*H. sieboldiana*) until the first week of August (*H. capitata*). The blooming period of the *Hosta* species established in our study, is similar to that stated for Poland by Marcinkowski (2007). It is worth noting, that the flowering phenology of particular ornamentals studied did not differ considerably among the growing seasons, indicating that the studied hostas are not sensitive to abiotic factors, e.g. cyclic changes in weather as regards

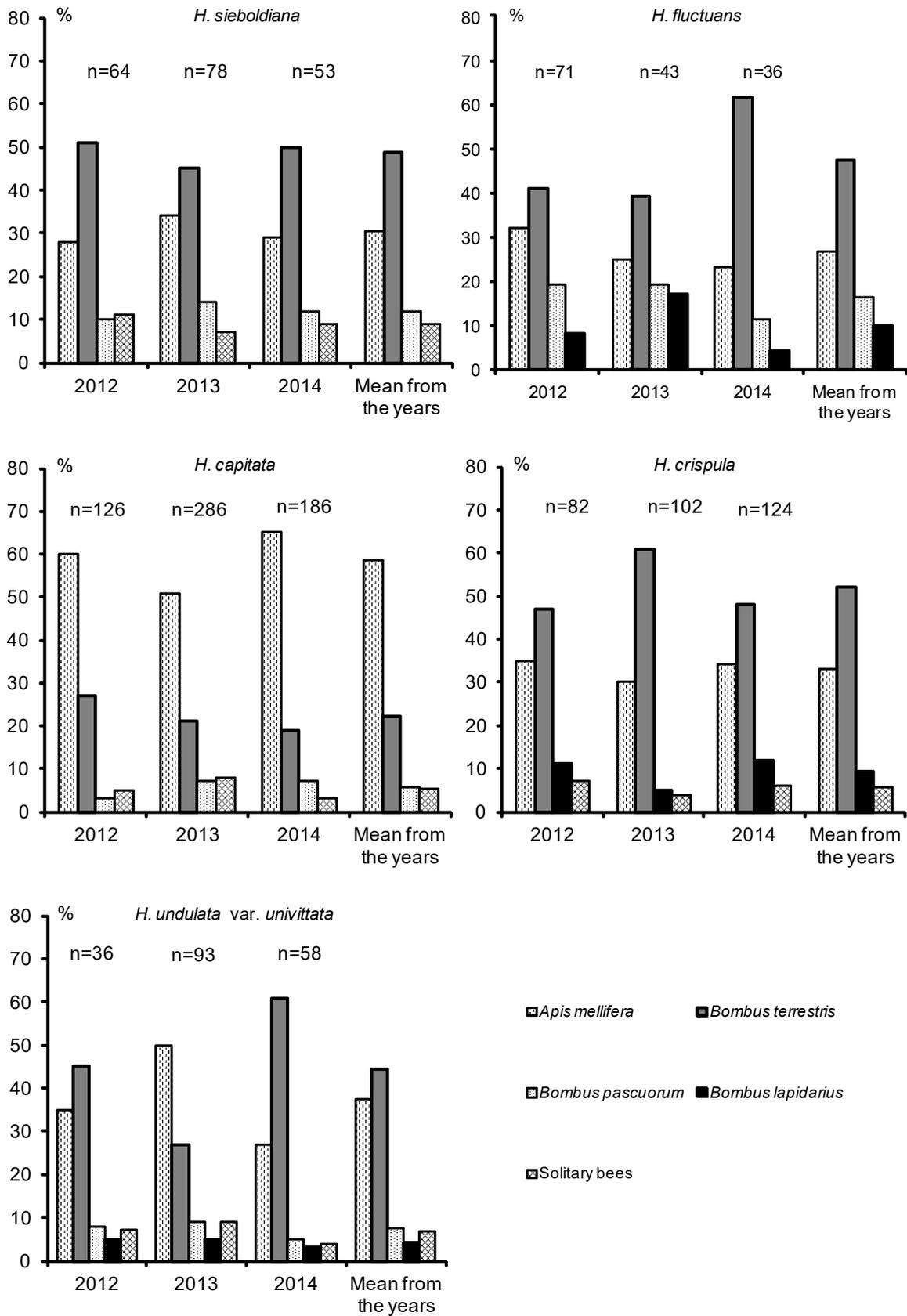


Fig. 2. The percentage participation of insect visitors on flowers of *Hosta* representatives. Mean values for the years of study are given; n = the total number of insect visitors recorded.

the conditions noted over the years of the study. Consequently, there is a high degree of probability of year-to-year repeatable food resources for insects, during a period when the assortment of blooming ornamentals for pollinator-friendly gardens is markedly declining in Poland (Kołtowski, 2006).

Our observations concerning the inter-species differences in the number of flowers per inflorescence are in accordance with descriptions of Marcinkowski (2007), and Grabowska and Kubala (2012), and indicate that the trait is highly genetically determined. Interestingly, the number of flowers per inflorescence did not differ substantially among growing seasons for any of the species studied. These results may suggest a high resistance of bud formation to exogenous conditions (e.g. air temperature, water balance). This is a unique feature. In a great number of species, the flower-bud set has been described as sensitive to weather factors (Chmiel, 2010). In perennial species, the onset of flowering is highly dependent on the resources accumulated in the stored organs as well as of resource allocation and/or current foliar photosynthetic efficiency during bud formation (Cao et al., 2007).

Open perianth of the studied *Hosta* species lasted only 20 - 36 hours, which is consistent with the findings of Cao et al. (2007) for *H. rectifolia* (only 1 day of flower life span) and with the observation of Chwil (2009) for *H. fortunei* (approx. 2 days of flower life span). The flower life span is species-specific (van Doorn, 1997; Masierowska, 2006; Denisow et al., 2014b), but is also under the control of various exogenous factors, e.g. the air temperature, air humidity, and whether or not the flower is pollinated (Ashman and Shoen, 1994; Denisow, 2009). Generally, flower life span has an ecological, genetic, and physiological significance (Waser et al., 1996). It impacts on the number of open flowers per inflorescence (= display size) and influences the rate of availability of floral reward for flower visitors (van Doorn, 1997). On a given day, only one or two *Hosta* flowers per inflorescence are in anthesis and expose nectar and pollen available for flower visitors. Accordingly, the limited quantity of the reward offered at a given time may partly explain the low frequency of insect visitors to hostas flowers.

In the genus *Hosta*, flowers tend to secrete a low to medium concentrated nectar. Chwil (2009) measured 23 - 30% sugar concentration in the nectar of *H. fortunei*. Sugar concentration could be related to nectary characteristics and the type

of sucrose transport across the nectary tissue (Nicolson and Thornburg, 2007). It is also related to changeable weather parameters (Denisow et al., 2014a). Variations in meteorological conditions (air humidity, air temperature) may explain the year-to-year differences in the nectar sugar concentration noted in two out of three *Hosta* species studied.

Bumblebees predominated on the flowers of five out of six *Hosta* representatives studied. Different *Bombus* species were described as the most frequent visitors to *H. sieboldiana* by Suzuki et al. (2003), who consider them the true pollinators. It is unclear why insects favour certain flowers. Our observations suggest that flower morphology rather than nectar characteristics restrict insect visitors to the flowers of *Hosta* species. It is accepted that nectar traits have an impact on the spectrum of insect visitors (Galletto and Bernardello, 2004). According to different authors, the optimal range of nectar concentration for bumblebee forage is 50 - 65%, and for honeybee forage is 25 - 40% (Nicolson and Thornburg, 2007). In our study, nectar 'suitable for honeybees' was avoided by honeybee foragers, whereas nectar of a lower concentration than 'optimal for bumblebees' was willingly collected by bumblebees. In general, a majority of studies have documented that the behaviour of foragers is complex. The broad space at the intermediate part of funnel-bell shaped *Hosta* flowers allows both honeybees and bumblebees to enter the perianth when nectar probing. On the other hand, the gradually narrowing perianth between the narrow- and broad-tube parts of the flower might enable insect species with a short proboscis to enter the narrow tube and to reach the nectar hidden deeply inside. Presumably, due to floral morphology, the foraging behaviour of bumblebees and honeybees differed. The vast majority of bumblebees, after one visit to the *Hosta* flower, continued nectar-feeding at subsequent flowers. But most honeybee individuals, after one floral visit, escaped from the flowers and did not continue to forage for nectar in other flowers. The only exception was *H. capitata*, whose flowers were predominantly foraged by honeybees. The interest of honeybees in the flowers of *H. capitata* was presumably attributable to the relatively high amount of sugars produced per unit area (2 - 5 fold more compared to other hostas). The energy intake rate is considered an important factor in honeybee diet selection (Fewell and Winston, 1996; Osborne et al., 2008). In general, the relationship between phenotypic floral traits and/or floral-reward characteristics and insect-visitor

frequency are common interdependences (Willmer and Stone, 2004). Year-to-year changes in visitor frequency observed in our study, is also a commonly reported phenomena and is usually directly related to fluctuations in the insect population size between growing seasons (Goulson, 1999).

There is more and more evidence that the limitation of floral resources is among the reasons for the pollinators' decline (Alaux et al., 2010). Therefore, the quantity of floral reward offered, is an important factor to consider during the selection of plants for bee-friendly gardens. Overall, the total sugar yield of the *Hosta* species studied was low, only 2.31 - 11.80 g per 10 m². Likewise, the pollen production amounted to 0.24 - 9.53 g per 10 m². These values place the hostas among poor sugar and pollen yielding plants (Kottowski, 2006).

Pollen grains of the *Hosta* species and varieties are large in size, as they are within the range of 50 - 100 µm (Erdtman, 1954). According to Liu et al. (2011), pollen grains of the *Hosta* species are long-ellipsoidal or ellipsoidal. In our study, the mean shape index (Erdtman, 1954) in polar view, classify the pollen grains as *prolatum* (shape index 1.33 - 1.61), while in equatorial view as *oblatum* (shape index 0.5 - 0.7). The pollen shape of *Hosta* representatives indicates a similarity with that of related genera, i.e. *Hemerocallis*, *Agave*, *Yucca* (Chung and Jones, 1989).

CONCLUSION

Under the environmental conditions of SE Poland, the ornamental *Hosta* species flower continuously from the second decade of June until the first week of August. The phenotypic traits of the *Hosta* flowers, including a deep, narrow perianth tube, reduce the access to nectar reward and restrict an array of insect visitors. Although the total sugar mass and pollen production are low in the *Hosta* species, the flowers can complete the summer pasture, particularly for bumblebees which are the key visitors.

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