



Carbon Uptake and Air Pollution Mitigation of Different Evergreen Shrub Species

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Abstract. Three independent experiments assessed CO₂ assimilation and metals leaf deposition of seven evergreen shrub species (*Arbutus unedo* L., *Elaeagnus × ebbingei* L., *Laurus nobilis* L., *Ligustrum japonicum* Thunb., *Photinia × fraseri* Dress., *Viburnum tinus* subsp. *lucidum* L., and *Viburnum tinus* subsp. *tinus* L.). CO₂ assimilation and carbon allocation were determined in 2011 (Exp. 1) under optimal water availability and in 2012 (Exp. 2) under drought on potted plants. A third experiment (Exp. 3) measured seasonal leaf depositions of Cd, Cu, Ni, Pb, and Zn in 2011 on plants transplanted in proximity of a four-lane road.

E. × ebbingei showed the highest CO₂ assimilation under optimal water availability but one of the lowest under drought (Exp. 1, 2). Conversely, *P. × fraseri* had intermediate CO₂ assimilation but it declined less during drought compared to the other species. In Experiment 3, *E. × ebbingei* showed the highest metal deposition, mainly due to its greater leaf area. Greater rainfall and RH% decreased metal depositions, whilst greater wind velocity and air temperature increased leaf depositions. Species which drastically reduce CO₂ assimilation under drought (*V. tinus* subsp. *lucidum*, *L. japonicum*, *E. × ebbingei*) are not recommended in drought-prone environments, where drought-tolerant “mesic” species (*P. × fraseri*), should be preferred. *E. × ebbingei* could be used to optimize deposition of metals. The three experiments provide useful insights especially about CO₂ assimilation (Exp. 1, 2) and air pollution mitigation (Exp. 3) of widely used shrubs for application in urban areas and planning of roadside greening in southern Europe.

Key Words. *Arbutus unedo* L.; CO₂ Assimilation; Drought; *Elaeagnus × ebbingei* L.; Italy; *Laurus nobilis* L.; Leaf Deposition; *Ligustrum japonicum* Thunb.; Meteorological Parameters; *Photinia × fraseri* Dress.; Relative Growth Rate; Seasonal Trend; Shrub; Trace Metals; Traffic Pollution; *Viburnum tinus* subsp. *lucidum* L.; *Viburnum tinus* subsp. *tinus* L.

Human activities, such as vehicle traffic, industry, and the heating and conditioning of buildings, contribute to changing the atmospheric environment in terms of CO₂ and air pollution concentrations (Grubler 1994; O’Meara 1999; Suzuki 2006). Since the early 1900s, the average surface temperature of Earth has increased by 0.74°C (IPCC 2013). Global warming is mainly driven by CO₂, which is largely emitted from the combustion of fossil fuels (PBL Netherlands Environmental Assessment Agency 2013). Urban greening has been reported to contribute effectively to the uptake and storage of atmospheric CO₂ through its conversion into organic carbon mean photosynthesis. CO₂ is then stored as woody biomass for medium to long periods, depending on the average life span of the species involved and the use of the woody material after removal from the planting site (Pugh et al. 2012; Nowak et al. 2013). The potential of trees

and shrubs to remove CO₂ and pollutants is enhanced when they are planted within or near urban environments, where the concentration of CO₂ and pollutants is higher (Niinemets and Peñuelas 2007). In addition, they offer others benefits *in situ*, such as improving microclimate conditions and reducing energy consumption of buildings for air conditioning (McCarthy et al. 2010). Plant species differ in their relative capacity to assimilate and store CO₂ (Somogyi et al. 2008; Semwal et al. 2013). Previous research investigated CO₂ storage by several trees species and found that this is positively related to life span and size at maturity of the tree (Nowak and Crane 2002). However, CO₂ storage potential of shrubs has been poorly investigated, even though they may account for about 10%–11% of total carbon storage and for 21%–25% of total annual carbon uptake by urban vegetation (Jo and McPherson 1995).

In Mediterranean areas, the use of drought-tolerant shrubs can considerably increase carbon sequestration by local vegetation (Bombelli and Gratani, 2003). Unfortunately, the degree of drought tolerance and the mechanisms adopted to cope with drought have not been studied for most shrub species commonly planted in the urban environment. Particular interest should be given to determining which species are capable of tolerating (rather than avoiding) drought, thus maintaining high CO₂ assimilation during dry periods (Bussotti et al. 2013; Fini et al. 2014). Shrub species are suggested as valid complements, especially where there is not enough space for tree growth or where local legislation imposes specific distances for tree plantation along streets or buildings.

In parallel with CO₂ assimilation and carbon storage, plants may play a role in decreasing local air pollution. Particulate matter (PM) is the principal component of anthropogenic air pollutants. PM includes a mixture of elements, heavy metals, black carbon, polycyclic aromatic hydrocarbons, and other substances suspended in the atmosphere (Bell et al. 2011). Plants can adsorb and absorb PM, as well as other noxious gaseous pollutants such as SO_x, NO_x, and O₃ (Nowak et al. 2000; Nowak et al. 2006; Bealey et al. 2007), and pollution-capturing efficiency is species-specific (Dzierzanowski et al. 2011; Sæbø et al. 2012; Mori et al. 2015a). Generally, it has been found that coniferous species intercept higher amounts of particulates due to their complex leaf structures compared to broad-leaved species (Beckett et al. 2000; Freer-Smith et al. 2005). In addition, evergreen plants are able to intercept air pollutants during winter as well (Freer-Smith et al. 2005), when the concentration of air pollutants is usually higher (Pikridas et al. 2013). Species with large leaf areas and dense canopies can create higher air turbulence and consequently have a higher impact of suspended particles compared to species with low-dense canopies (Mori et al. 2015b). Trees can also act negatively, increasing air pollution concentrations at the pedestrian level by reducing air circulation (Buccolieri et al. 2009; Salmond et al. 2013).

Leaf anatomical and ultrastructural traits, such as texture, surface roughness, presence of

pubescence, and the quantity and size of stomata are positively related to the retention capacity of leaves toward PM (Huixia et al. 2013). Leaf surfaces are temporary retention sites for pollutants, the presence of which is influenced by meteorological factors (Nowak et al. 2006). Thus, an interesting aspect of the interaction between plants and air pollutants is how deposition of pollutants on leaves changes over time. For example, Anicic et al. (2011) found a seasonal accumulation, from May to September, of Cr, Fe, Ni, Zn, and Pb on leaves of *Aesculus hippocastanum* and *Tilia* spp., but similar information about shrub species in dry and warm climates is still lacking. Indeed, most of the research in the last 15 years has been carried out on temperate tree species (Beckett et al. 2000; Nowak et al. 2002; Nowak and Crane 2002; Pugh et al. 2012). Elevated concentrations of heavy metals are common in urban areas as a result of both a wide range of human activities and natural processes (Lu et al. 2010).

As heavy metals are strongly associated with PM (Duzgoren-Aydin et al. 2006), especially with the coarse fraction having diameters up to 10 µm (PM₁₀) (Araujo and Nel 2009), several studies have measured some heavy metals as descriptors of air pollutants (Fowler et al. 2003; Mori et al. 2015a). To improve knowledge of the benefits of shrubs for urban landscape, three experiments were carried out to characterize the following seven shrub species regarding their capacity for CO₂ uptake and storage and air pollution sequestration: *Viburnum tinus* subsp. *lucidum* L. cited as *Viburnum lucidum*, *Viburnum tinus* subsp. *tinus* L. cited as *Viburnum tinus*, *Arbutus unedo* L., *Photinia* × *fraseri* Dress., *Laurus nobilis* L., *Elaeagnus* × *ebbingei* L., and *Ligustrum japonicum* Thunb. CO₂ uptake and storage were evaluated under conditions of non-limiting moisture availability (Exp. 1) and drought stress (Exp. 2), while a third experiment (Exp. 3) aimed to identify the species with highest leaf deposition of trace metals (Cd, Cu, Ni, Pb, and Zn) (*Viburnum tinus* was not included in this experiment), to study the relationships between trace metal depositions trends and meteorological parameters, and to identify the possible sources responsible for the presence of metals on leaves.

MATERIALS AND METHODS

Experiment 1: Carbon Uptake and Storage Under Optimal Water Availability

In autumn 2010, 84 uniform, two-year-old shrubs (12 per species) of *Arbutus unedo*, *Elaeagnus × ebbingei*, *Laurus nobilis*, *Ligustrum japonicum*, *Photinia × fraseri*, *Viburnum lucidum*, and *Viburnum tinus*, were potted in 3L containers. The substrate was a peat:pumice mixture 4:1, (v/v), added with 3 kg m⁻³ of a controlled release fertilizer (Osmocote Exact, 18-2,5-7, 8–9 months, Everris, Intl. B.B, Geldermalsen, The Netherlands). Shrubs were arranged in a randomized block design, with six blocks of two shrubs per species in each block, and grown outside in Sesto Fiorentino (43.816652 N, 11.199673 E), Florence, Italy, under full sunlight. Average maximum temperature of the area from 01 June to 01 September, calculated over the last 30 years, is 30.2°C with peaks of 42°C. The average rainfall (June to September for the same period) is 149.8 mm (Wikipedia 2016).

Shrubs were irrigated daily to container capacity to avoid water stress for the entire experimental period.

Daily trends of net photosynthesis and transpiration were measured for 12 consecutive days during summer 2011, from 9:00 am to 6:30 pm, using an infrared gas analyzer (Ciras 2, PP-system, Amesbury, Massachusetts, U.S.). Measurements were conducted at 380 ppm of CO₂ and ambient irradiance on 18 current-season, fully expanded leaves per species. Water use efficiency (WUE, μmol CO₂/mmol H₂O) was calculated as the ratio between CO₂ assimilation and transpiration for all 12 days of measurement (Gebhardt et al. 2014). Mean daily assimilated CO₂ per unit leaf area was obtained applying integral analysis of the daily CO₂ assimilation curve as reported by Fini et al. (2012). In detail, total assimilated CO₂ was estimated by plotting the mean daily net CO₂ assimilation versus time and, thereafter, applying an integral analysis to calculate the area underlying the curve. To obtain daily CO₂ assimilation per plant, the mean CO₂ assimilation per unit leaf area of each species was multiplied per the total leaf area of shrubs, assuming that self-shading was not relevant because of the small size of the shrubs.

Biomass measurements were carried out at the beginning and at the end of the experiment

on four shrubs per species. In November 2011, shrubs were cut off at the root flare and roots were cleaned from the medium using an air flush. All leaves were separated from stems and immediately scanned with an A3 scanner. An image analysis software (Image Tool 1.3, UTHSCSA) was then used to measure leaf area. To determine dry weight, leaves, stems, and roots were oven dried at 70°C until constant weight was reached (≈72 hours). Root-to-shoot ratio was calculated as the ratio between root dry weight and leaves + stem dry weight (Fini et al. 2011). Biomass allocation (%) was calculated as the change in dry mass of a specific plant organ (stems, roots, and leaves) over the change in dry mass of the whole plant (stems + leaves + roots) with the aim of identifying the species that allocated and stored more carbon into more durable organs compared to leaves.

Experiment 2: Carbon Uptake Under Drought Stress

In October 2011, 144 three-year-old shrubs of the same species used in Experiment 1 (24 shrubs per species; *Laurus nobilis* was not used), were potted in 5L containers. Experimental site, substrate, and fertilization were the same as in Experiment 1, but a novofol foil (see Fini et al. 2012) was used to protect shrubs from undesired rainfall. Shrubs were arranged in a randomized block design of three blocks with eight shrubs per species in each block. Container capacity, wilting point, and water-holding capacity of the substrate were determined using a gravimetric method (Sammons and Struve 2008) as described in other works (Fini et al. 2011; Fini et al. 2013). All shrubs were watered daily to container capacity (CC) until August, and then exposed to water stress according to the following treatments: 1) WW, 72 shrubs (four shrubs per species per block) were watered daily to container capacity throughout the experiment; and 2) WS, (72 shrubs), water withheld from four shrubs per species for two periods of five days each (drought phase 1 and 2), separated by 14 days of partial relief. During partial relief the substrate moisture was maintained at 30% of CC, corresponding to mild to moderate drought (Ruiz-Sanchez et al. 2000; Fini et al. 2011) (Figure 4A). Physiological parameters were measured six times during the 24 days of the water stress experiment according to the following plan:

1) leaf gas exchange [CO_2 assimilation (A); transpiration (E); water use efficiency ($\text{WUE} = A/E$)] were measured at T-0 (before the onset of drought), T-1 (one day after imposition of drought), T-2 (three days after imposition of drought), T-3 (five days after imposition of drought), T-4 (end of partial relief), T-5 (end of experiment), using the same methods and instruments as described in Experiment 1;

2) pre-dawn water potential was measured at T-3 using a Scholander type pressure bomb (PMS Instruments, Albany, Oregon, U.S.) between 3:00 and 5:00 am on four leaves per species, treatment, and block;

3) maximum quantum yield of PSII photochemistry (Fv/Fm) and chlorophyll a fluorescence induction (OJIP) curve were performed with HANDY PEA (Hansatech Instruments Ltd, Norfolk, United Kingdom) on four leaves per species, treatment, and block (44 leaves in total);

4) drought-induced changes in growth rate were measured calculating the relative growth rate (RGR) of each species from the beginning to the end of the experiment, as described in a previous work (Fini et al. 2010).

Experiment 3: Leaf Surface Accumulation of Trace Metals

In spring 2010, 50 three-year-old shrubs of *A. unedo*, *E. × ebbingei*, *L. nobilis*, *L. japonicum*, *P. × fraseri*, and *V. lucidum* (300 shrubs in total) were planted in a field next to a high-traffic road. Before planting, soil was plowed to 40 cm. Shrubs were planted into hand-dug planting holes (as deep as the rootball and 1.5 times the width of the rootball) and irrigated as required to avoid any drought stress during the first year. Shrubs were planted to form two vegetation belts, each 30 m long and 5 m wide. These belts were parallel to a four-lane road (average daily number of vehicles was 12,000) in Pescia, Italy (N 43°52'57.7992", E 10°40'58.0692"), and were at a distance of 1.5 m from the road verge. Distance between belts was 30 m. Twenty-five shrubs per species formed a 5 m

× 5 m section in each belt. The experimental area is located 35 km from the Ligurian Sea and the land is dedicated mainly to floricultural and residential uses (Figure 1). Climate parameters for the area during the experiment are reported in Table 1. The dominant wind in the experimental plot is from the sea (west to east), thus the barriers are on the leeward side of the road (Figure 1). At planting, shrubs were pruned to uniform size of 150 cm, with the exception of *L. japonicum* and *E. × ebbingei* which were 130 cm tall. Initial total leaf area was measured on four shrubs per species. To measure leaf area, shrubs were fully defoliated and leaf fresh weight was immediately determined; 30% (in weight) of the leaves of each shrub were subsampled and used for the leaf area measurements, which were carried out using a RS2XA illumination and camera system (Kaiser, Germany) and A3 Lightbox (G.C.L., Durham, UK) with WinDIAS software (Dynamax Inc., Houston, Texas, U.S.). Each subsample (30% in weight) was then dried separately and the specific leaf area (SLA) was calculated on those leaves as the ratio between their area and dry weight. SLA values were then multiplied per the total leaf dry weight of samples to obtain the total leaf area of each plant (Table 2).



Figure 1. Aerial photo of the area surrounding the experimental site. The orientation of the two vegetation belts (indicated with dark rectangle) and the main land use areas (agricultural and residential) are shown. Black arrow indicates north.

Table 1. Meteorological data for the experimental area of the three periods prior to the 2011 samplings.

Sampling periods	19 May – 21 June	21 June – 03 August	03 August – 04 October
Precipitation (mm)	10.80	11.20	80.00
Wind direction	SW	SW	WSW
Wind speed (m s^{-1})	0.80	0.80	0.62
Temperature ($^{\circ}\text{C}$)	20.78	23.33	23.24
Relative humidity (RH%)	71.45	63.94	61.33

Table 2. Leaf area (m²) of the six shrub species used for trace metals analysis at planting date (autumn 2010).

	Mean	SD
<i>Viburnum lucidum</i>	0.41	0.04
<i>Arbutus unedo</i>	0.30	0.01
<i>Photinia</i> × <i>fraseri</i>	0.35	0.05
<i>Laurus nobilis</i>	0.44	0.09
<i>Elaeagnus</i> × <i>ebbingei</i>	0.50	0.06
<i>Ligustrum japonicum</i>	0.20	0.05

In 2011, leaf trace metal deposition was measured three times (June, August, and October) on leaves formed in the current growing season in order to avoid any confusing age-related effects. Samplings were conducted after at least 10 rainless days. Two replicates of 10 fully expanded leaves per species per belt (40 leaves in total) were collected, including leaves at different heights. The leaf area of every sample was measured. Healthy leaves of each species were collected at a distance of at least 1.5 m from the adjacent species in order to reduce the influence of position. During the experiment, 72 samples were collected and immediately taken to the laboratory for subsequent analysis. In the laboratory, leaves were washed in 50 ml of a solution composed of distilled water and HCl 0.01 M (pH 2.0), per 30 minutes, placing samples on an orbital shaker. The concentrations of Zn, Cd, Pb, Ni, and Cu in the solutions of the different samples were directly analyzed by Inductively Coupled Plasma – Optical Emission Spectrometer – Dual Vision (Optima 2000, Perkin Elmer, Waltham, Massachusetts, U.S.). The results were expressed as weight of metals per unit of leaf area.

To measure the quantity of metals washed off from the canopy by precipitations, two containers (50 cm × 100 cm) were placed at the base of four replicate shrubs of each species. Control containers were placed at the same distance from the street in an area without vegetation. Rainwater in the containers was collected, measured, and stored in the dark at 0°C from June to October 2011, after each significant rain event (>5 mm). In November 2011, each rainwater sample was mixed until complete re-suspension of the sediments was reached and greater homogeneity within the sample was obtained; 20 ml of each sample was collected and fully evaporated at 50°C to collect the dry matter. The dry matter was then digested with 5 ml of a mixture of HNO₃ (65%) :

HClO₄ (68%), (5:2) v/v at 180°C per 60 minutes using a heating digester DK 6 (VELP® Scientifica, Usmate, Italy) to obtain the metal concentration, which was, for each sample, multiplied by the total volume of collected rainwater in order to obtain the total quantity of each metal. The primary aim was not to compare the effect of species but to compare the effect of presence or absence of shrubs on the metal content in runoff water.

Plant height and crown diameter were measured on six shrubs (three per belt) per species in May, July, and October 2011. On the same sampling dates, leaf area index (LAI) was measured by a ceptometer (AccuPAR LP-80, Decagon Devices, Inc., Pullman, Washington, U.S.) at 1 m above the ground. In October, the total leaf area of four shrubs per species was measured by destructive harvest. The depositions of trace metals on the whole plant were then calculated by multiplying the depositions per unit leaf area and the whole plant leaf area.

Morphology of the leaf surface and presence of particulate matter on the abaxial and adaxial surfaces were observed on ten leaves per species from one sampling event in July using an ESEM (electron scanning environmental microscope) Quanta 200, (Fei Corporation, Eindhoven, The Netherlands), operating in low-vacuum mode (chamber pressure was kept at 1 Torr) and 25 kV. Leaves to be observed by ESEM were harvested using separate disposable gloves for each species; leaves were harvested from the petioles, avoiding contact with the leaf blade. After harvesting, leaves were carefully placed in disposable paper bags and immediately processed.

Meteorological parameters (rainfall, wind speed, wind direction, relative humidity, and air-temperature) were measured continuously from June to October 2011 at 1.5 m height, using an iMetos® SD meteorological station (Pessl Instruments, Weiz, Austria) positioned between the vegetation belts (15 m distant from each belt).

Statistical Analysis

Growth parameters and water relationships were analyzed using one-way (Exp. 1) or two-way ANOVA (Exp. 2). Leaf gas exchange data were analyzed using repeated measures ANOVA. Mean separation was carried out using Duncan's MRT (SPSS statistics 19, IBM Company, New York, U.S.). Differences in trace metal deposi-

tions (Exp. 3) were tested with repeated measures ANOVA and separation of the means was carried out using Duncan's MRT (CoStat 6.303, CoHort Software, Monterey, California, U.S.). The relationships between metal depositions on leaves and climate parameters were tested using Partial Least Square Regression (PLSR) for each metal separately. PLSR is useful when a high level of multi-collinearity among the explanatory variables is assumed (Rosipal and Trejo 2001). Correlations between the deposited metals were tested using Pearson product-moment correlation coefficients (r) and only the r values above 0.5 were considered. Multivariate methods were used to identify the possible sources of different metals. Cluster Analysis (CA) was performed using Ward's method and Euclidean distance as metric (Oliva and Espinosa 2007). Factor Analysis (FA) was accomplished with Varimax rotation (Lorenzini et al. 2006). The outliers were identified and removed from the data set.

RESULTS

Experiment 1: Carbon Uptake Under Optimal Water Availability

Elaeagnus × ebbingei, *A. unedo*, and *L. nobilis* had the highest mean daily CO₂ assimilation per unit leaf area compared to the other species when shrubs were grown under optimal water availability (Figure 2A). Conversely, *V. lucidum*, *L. japonicum*, and *V. tinus* showed the lowest values. All species showed the highest values of CO₂ assimilation during the morning hours and then underwent a significant midday depression. *Elaeagnus × ebbingei* and *L. japonicum*, in contrast with the other species, showed a significant recovery of CO₂ assimilation, reaching values similar to those in the morning, after 3:00 pm.

CO₂ assimilation of the whole plant (a parameter that considers both the assimilation per unit leaf area and the total leaf area of the plant) was higher in *E. × ebbingei* and *L. nobilis* than in *V. lucidum*, *V. tinus*, and *P. × fraseri*; *A. unedo* and *L. japonicum* showed the lowest values (Figure 2B).

Species had different WUE (*V. lucidum* = 3.87 ± 2.35; *A. unedo* = 3.93 ± 1.86; *P. × fraseri* = 4.71 ± 2.51; *L. nobilis* = 4.86 ± 2.14; *E. × ebbingei* = 4.76 ± 2.34; *L. japonicum* = 4.09 ± 2.06; *V. tinus* = 4.36 ± 2.60 expressed as μmol CO₂ mmol⁻¹ H₂O).

Laurus nobilis and *E. × ebbingei* gave the highest woody biomass production (data not shown) and the highest relative growth rate calculated over the experimental period (Figure 2C). A positive strong correlation ($P < 0.01$; $R^2 = 0.95$) was found between whole-plant CO₂ assimilation and relative growth rate (Figure 2D).

Allocation of photosynthates among plant organs differed across species. *V. lucidum* allocated more carbon to leaves compared to all other species. On the contrary, *V. tinus*, *E. × ebbingei*, and *L. japonicum* displayed lower percentages of C allocation to leaves compared to the other species considered, even if differences were significant only when compared to *V. lucidum* (Figure 3). At the end of the experiment, *E. × ebbingei*, *L. nobilis*, *V. lucidum*, and *V. tinus* had higher total leaf area (0.29 ± 0.05; 0.32 ± 0.04; 0.27 ± 0.04; 0.30 ± 0.05 m², respectively) than *A. unedo*, *P. × fraseri*, and *L. japonicum* (0.18 ± 0.03; 0.19 ± 0.05; 0.16 ± 0.05 m², respectively).

Experiment 2: Carbon Uptake and Storage Under Drought Stress

On average, substrate moisture of WS shrubs declined to 15% of its holding capacity by the end of "drought phase 1" (T-3), where it was maintained around 30% by irrigation until T-4 and finally declined to 12% at the end of "drought phase 2" (T-5) (Figure 4A). CO₂ assimilation of WW shrubs largely confirmed the values observed in Experiment 1 (data not shown) and did not statistically differ from data collected from WS shrubs at T-0. As drought progressed (T-1, T-2, T-3), CO₂ assimilation declined in all species, but the magnitude of the decline was species-specific, with species such as *E. × ebbingei* and *V. lucidum* experiencing greater drought-induced reductions in CO₂ assimilation than *V. tinus* and *A. unedo*. During partial relief (T-4), CO₂ assimilation of some species (i.e., *P. × fraseri*, *V. tinus*, *L. japonicum*) exhibited some recovery, while in other species (i.e., *E. × ebbingei*, *A. unedo*, *V. lucidum*) CO₂ assimilation showed little or no recovery. At the end of drought phase 2 (T-5), *A. unedo*, *P. × fraseri*, *V. tinus*, and *L. japonicum* had higher CO₂ assimilation than *E. × ebbingei* and *V. lucidum*. However, as drought progressed, CO₂ assimilation of *E. × ebbingei* was strongly reduced (Figure 4B). A similar trend was found in *V. lucidum*, which

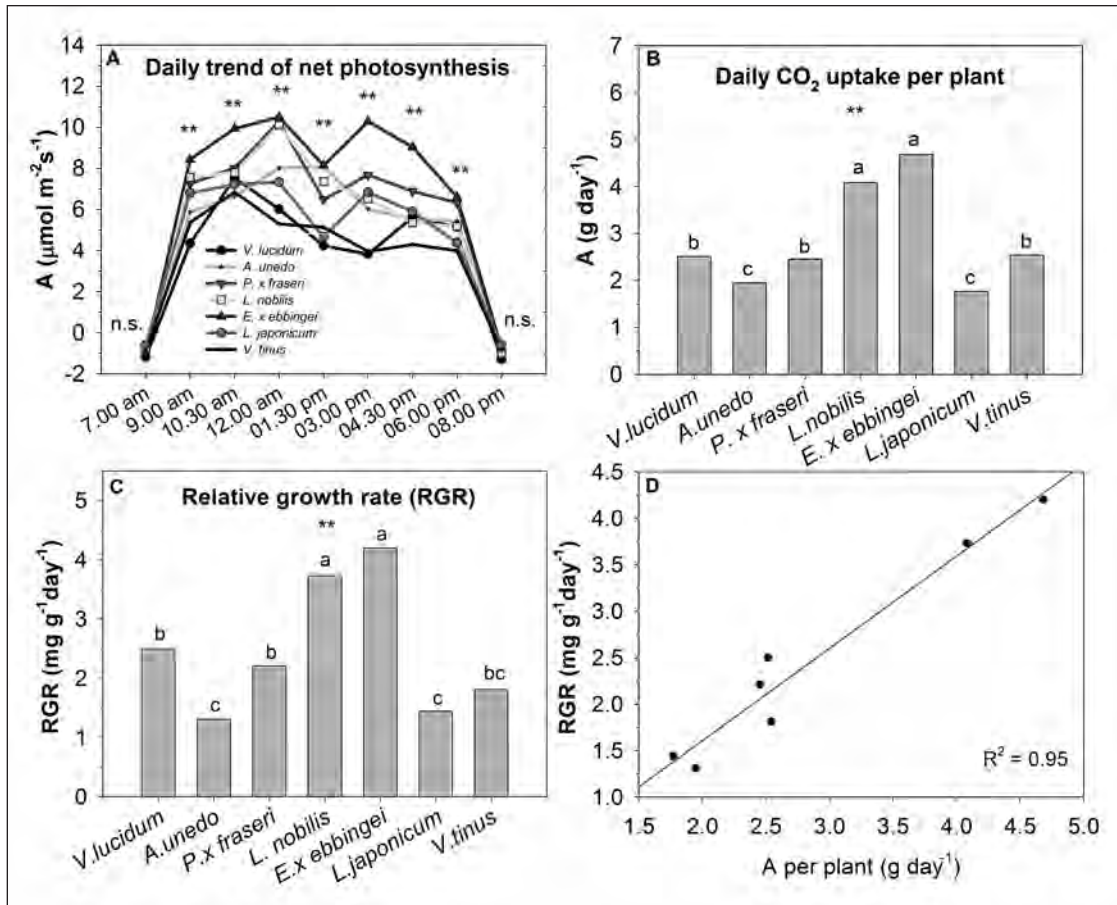


Figure 2. A) Daily trend of net photosynthesis expressed as $\mu\text{mol CO}_2 \text{ m}^{-2} \text{ (leaf area) s}^{-1}$; B) daily CO₂ uptake of the whole plant expressed as $\text{g CO}_2 \text{ assimilated day}^{-1}$; C) relative growth rate (RGR); and D) relation between RGR and whole-plant daily CO₂ assimilation of seven different shrubs in 2011, under optimal water availability. Data of CO₂ uptake are the average of 12 daily measurements of net photosynthesis. Double asterisk (**) indicates significant differences among species at $P < 0.01$, different letters indicate different homogeneous groups using Duncan's MRT.

had a very low CO₂ assimilation during drought. *Viburnum tinus* and *A. unedo* had the smallest reduction in CO₂ assimilation under drought stress.

Drought differentially reduced the maximum quantum yield of PSII photochemistry (Fv/Fm) during drought phase 1 (Figure 5). *Ligustrum japonicum*, *E. x ebbingei*, and *V. lucidum* showed the highest depletion in Fv/Fm, whereas *P. x fraseri* displayed the lowest deviation. In particular, Fv/Fm decreased below 0.70 in *L. japonicum* and *V. lucidum*, indicating a condition of stress.

Experiment 3: Leaf Deposition of Trace Metals

Species differed for Pb deposition on unit leaf surfaces while differences were not statistically significant for other metals (Table 3). *Ligustrum*

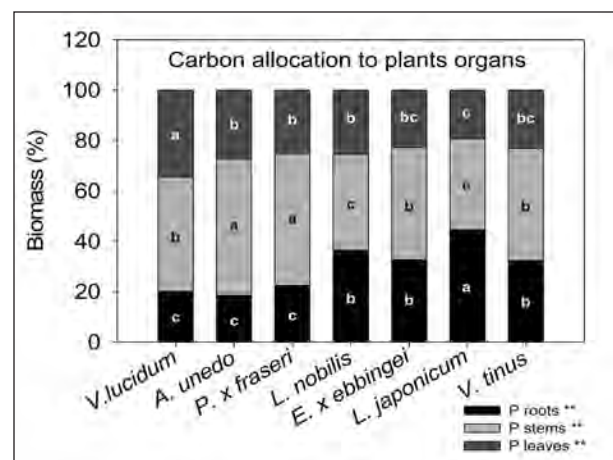


Figure 3. Carbon allocation to different plant organs in seven shrub species in 2011 under optimal water availability. Different letters within an organ (i.e., leaves, stem, and roots) indicate significant differences among species at $P < 0.01$ using Duncan's MRT.

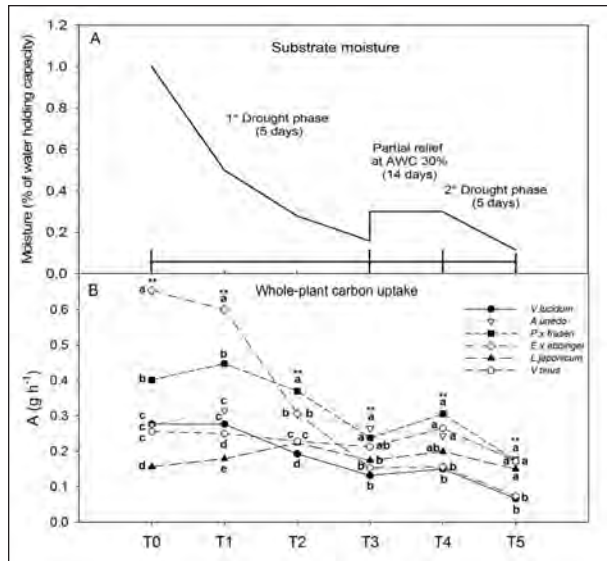


Figure 4. A) Irrigation scheduling and substrate moisture (expressed as percentage of the water holding capacity of the container), for plants under drought stress (WS); and B) Whole plant carbon uptake in six shrubs grown under water stress in 2012 expressed as g (CO₂ assimilated) per h⁻¹. Data of substrate moisture are the average 12 plants per species for six species. T indicates when physiological measurements were carried out. Different letters within the same sampling date indicate significant differences among species at $P > 0.01$ using Duncan's MRT.

japonicum, *E. × ebbingei*, and *V. lucidum* had higher Pb depositions per unit leaf area than *L. nobilis*. The quantity of Cu, Ni, Pb, and Zn deposited on the whole canopy was significantly higher in *E. × ebbingei* than in the other species.

Generally, deposition of metals increased from early to mid-summer (June–August) and thereafter decreased to October levels comparable to those in June (Figure 6). All metals, with the exception of Pb, presented a significantly different seasonal trend ($P < 0.001$). Partial least square regression analysis (Table 4), with four out of five significant results, suggests that the leaf metal depositions are negatively related to precipitation and RH% and positively related to air temperature and wind velocity.

The photos taken by ESEM (Figure 7) reveal the clear presence of particulate matter on the upper and lower leaf surfaces of all species with a greater quantity of trapped PM in the hairy leaves of *E. × ebbingei* when compared to the other (smooth-leaved) species.

Metal Washout from the Canopy

The rainwater collected at the base of all species, except *V. lucidum*, showed greater quantities of Zn, Cu, and Pb compared to control samples (Figure 8).

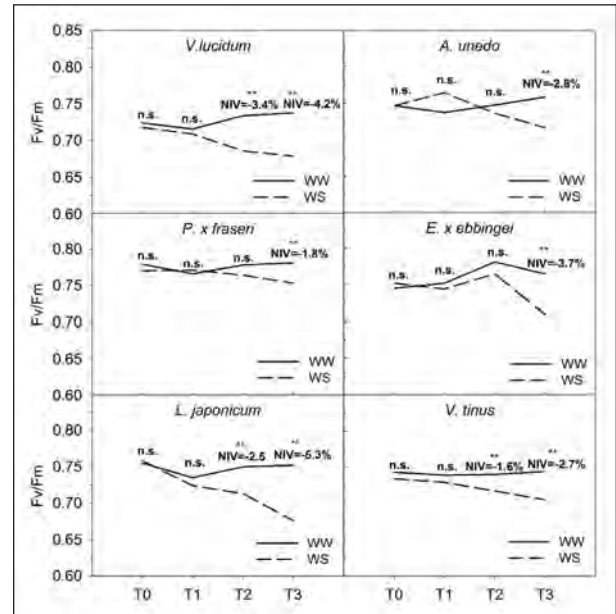


Figure 5. Maximum quantum yield of PSII photochemistry during the “drought phase 1” in six shrubs grown under either well watered (WW) or water stress (WS) conditions. Double asterisk (**) indicates significant differences between WW and WS plants of a species for $P > 0.01$ using Duncan's MRT. The normalized index of variation (NIV) was calculated as: $[(Fv/Fm)_{ws} - (Fv/Fm)_{ww}] / [(Fv/Fm)_{ws} + (Fv/Fm)_{ww}]$ (Tattini et al. 2006).

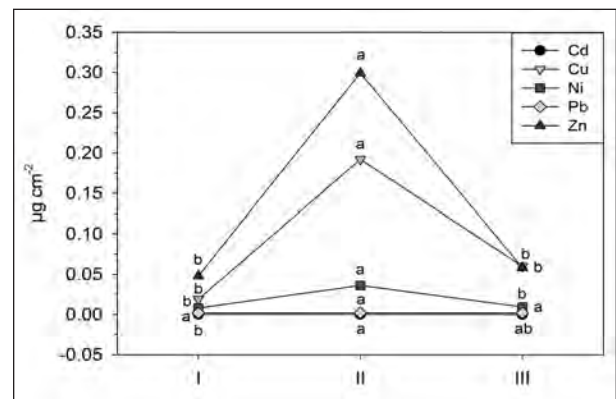


Figure 6. Metal depositions on leaf surface, expressed as µg (metal) cm⁻² (leaf area), in the different periods during growth season of 2011: 21 June 2011 (I); 03 August 2011 (II); 04 October 2011 (III); 30 August values are means of all species (µg cm⁻²). Different letters within the same sampling date indicate significant differences among species at $P > 0.01$ using Duncan's MRT.

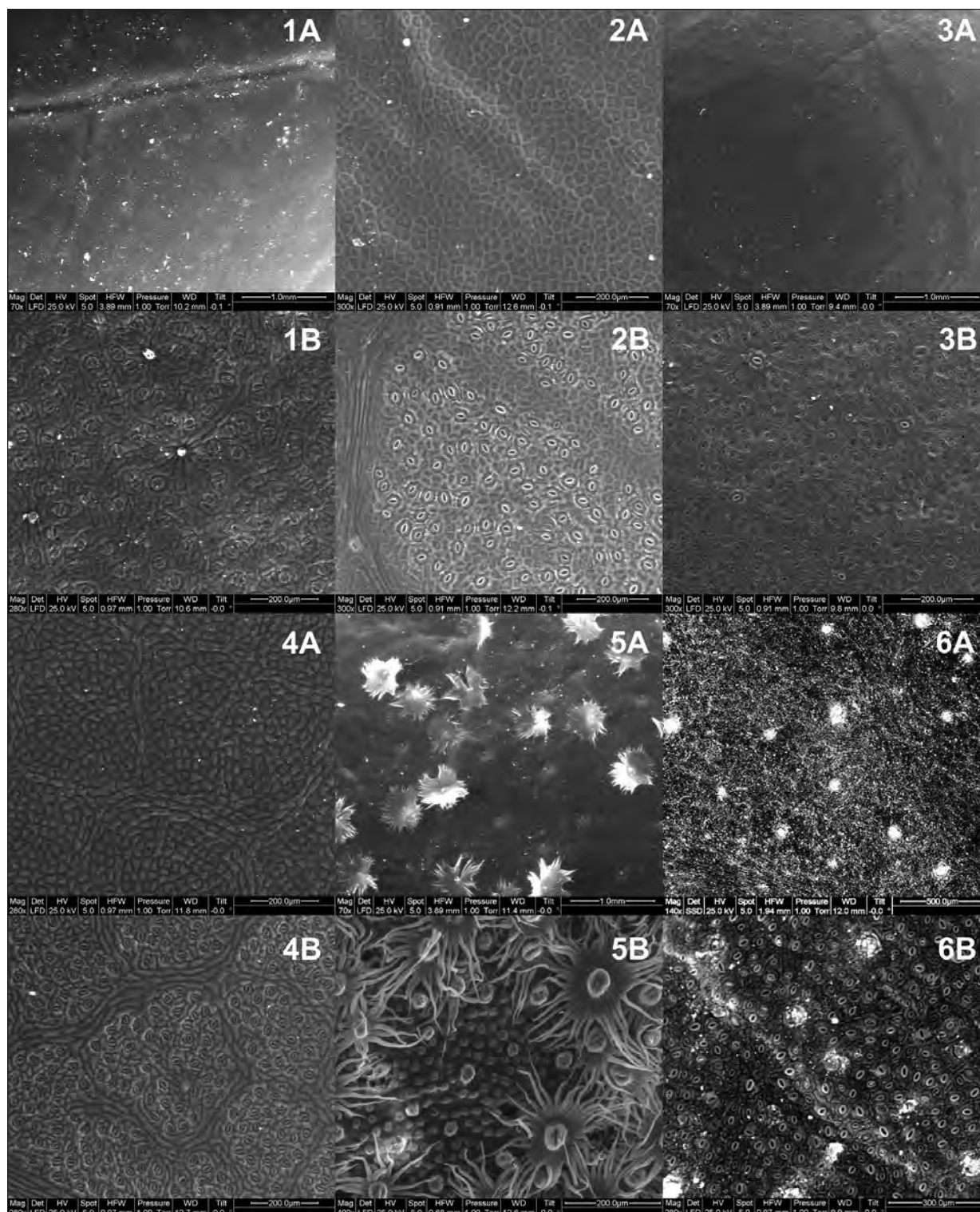


Figure 7. Environmental scanning electron microscope (ESEM) micrographies: A) adaxial and B) abaxial leaf surface of (1) *Viburnum lucidum*, (2) *Arbutus unedo*, (3) *Photinia x fraseri*, (4) *Laurus nobilis*, (5) *Elaeagnus x ebbingei*, and (6) *Ligustrum japonicum*. Images were taken at different magnifications as reported on the images.

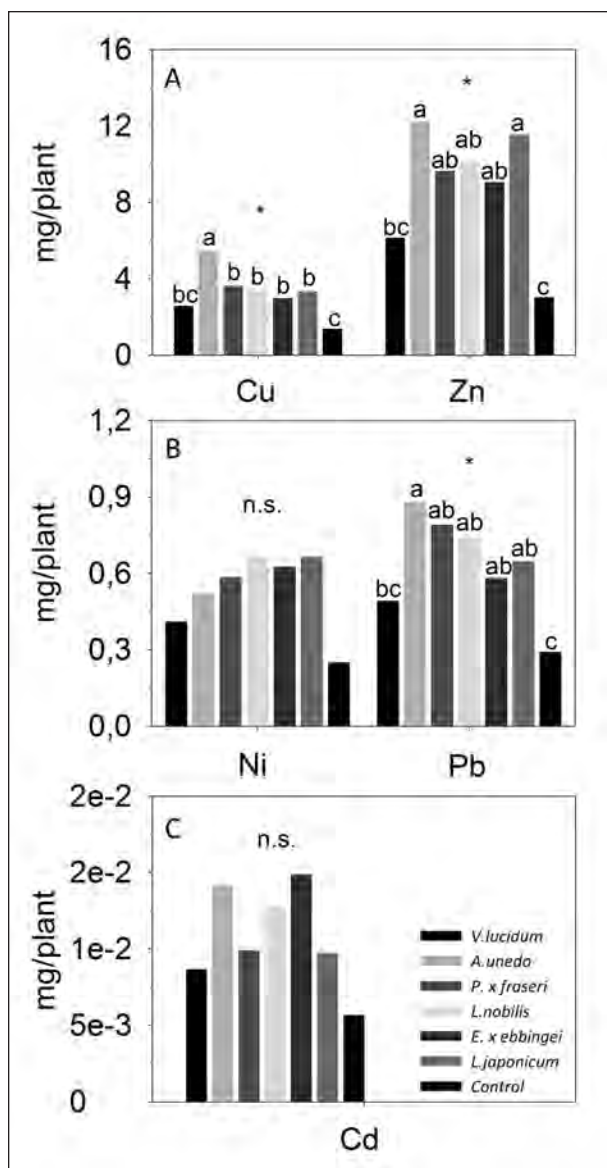


Figure 8. A) Accumulation of Cu and Zn; B) Ni and Pb; C) and Cd expressed as mg (metal) / plant, from June to October 2011 in the water collected at the base of *Viburnum lucidum*, *Arbutus unedo*, *Photinia x fraseri*, *Laurus nobilis*, *Elaeagnus x ebbingei*, *Ligustrum japonicum*, and control. Values are means (mg/plant). Different letters within the same sampling date indicate significant differences among species at $P < 0.05$ (*) using Duncan's MRT.

Metal Source Identification

Cluster analysis, which is generally used to group variables (metals) based on their similarity/dissimilarity, grouped the metals in two principal clusters. The first cluster included Cu and Ni, while Cd, Zn, and Pb were located in the second (Table 5). Factor Analysis (Table 5), which has the aim of describing variability among variables by a lower

number of artificial variables (factors), indicated that approximately 85% of the variance may be explained by the first two factors. Factor 1, which accounted for 58% of total variance, showed positive loadings with Cu, Ni, and Zn. In factor 2 (27% of total explained variance), Cd and Pb were dominant metals. Correlation analysis between leaf metal depositions showed that Cd is correlated with Pb ($r = 0.51$), Cu with Ni and Zn (respectively $r = 0.92$ and 0.83), and Ni with Zn ($r = 0.77$).

DISCUSSION

Experiments 1 and 2: CO₂ Uptake Under Contrasting Water Availability

Experiments 1 and 2 were aimed at investigating CO₂ assimilation and storage of seven widely used evergreen shrub species under either optimal moisture availability (Exp. 1) or water stress (Exp. 2). The results of these experiments corroborate the idea that the “carbon benefit” of shrubs is largely species-specific and dependent on a species-planting site interaction, as previously observed with tree species (Nowak and Crane 2002; McPherson and Simpson 2003). In particular, the ecological behavior of the species and the mechanisms evolved to cope with environmental stress are key determinants of carbon assimilation and carbon storage (Penuelas et al. 2001; West et al. 2012). The data highlight different growth strategies by the seven species investigated, which may be critical for proper use of these shrubs in the landscape. Some shrub species (i.e., *E x ebbingei*, *V. lucidum*, and *L. japonicum*) displayed a profligate/opportunistic water use (in the sense intended by Moreno-Gutiérrez et al. 2012), characterized by high water use and CO₂ assimilation per unit leaf area when water is not limiting, and drastic reduction of stomatal conductance, transpiration, and CO₂ assimilation after the onset of stress. Other species (i.e., *A. unedo* and *V. tinus*), instead, have evolved a conservative water-use strategy, consisting in relatively low leaf gas exchange during the favorable periods, which was only slightly depressed during stress (Moreno-Gutiérrez et al. 2012).

These findings have important applications for sustainable and benefit-based planning of urban green areas. In moist and favorable sites, or where irrigation water is readily available, CO₂ assimilation and carbon storage are highly correlated

Table 3. Foliar deposition, per unit leaf area and per the whole canopy, of six shrub species of five trace metals (Cd, Cu, Ni, Pb, and Zn). Data are the mean of three samplings performed in June, August, and October. Data are means ± SD. ANOVA tests (A) of differences between species are also included. Values as 0.000 are less than 0.001.

	<i>V. lucidum</i>			<i>A. unedo</i>			<i>P. × fraseri</i>			<i>L. nobilis</i>			<i>E. × ebbingei</i>			<i>L. japonicum</i>			F	P
	Mean	SD	A	Mean	SD	A	Mean	SD	A	Mean	SD	A	Mean	SD	A	Mean	SD	A		
	Depositions per unit leaf area (µg cm ⁻² *1000)																			
Cd	0.05	0.06	a	0.04	0.05	a	0.03	0.04	a	0.04	0.06	a	0.03	0.05	a	0.03	0.05	a	0.31	0.90
Cu	81.4	83.7	a	96.4	109	a	70.1	65.6	a	116	158	a	108	123	a	69.6	56.2	a	0.83	0.55
Ni	14.8	17.7	a	17.1	17.5	a	11.8	13.1	a	25.1	31.7	a	24.4	26.6	a	18.0	14.9	a	1.02	0.44
Pb	2.37	1.35	ab	1.62	1.25	bc	1.89	0.96	abc	1.25	1.03	c	2.09	0.72	ab	2.55	0.75	a	3.71	0.02
Zn	92.3	82.4	a	140	131	a	114	127	a	182	201	a	164	155	a	146	133	a	2.33	0.09
	Depositions on the whole plant (mg)																			
Cd	0.00	0.00	a	0.00	0.00	a	0.00	0.00	a	0.00	0.00	a	0.01	0.01	a	0.00	0.00	a	0.44	0.82
Cu	4.08	4.20	b	3.84	4.35	b	3.91	3.66	b	3.06	4.48	b	10.1	11.6	a	2.45	2.22	b	2.46	0.04
Ni	0.74	0.89	b	0.68	0.70	b	0.66	0.73	b	0.73	0.92	b	2.30	2.51	a	0.57	0.59	b	3.16	0.01
Pb	0.11	0.07	ab	0.06	0.05	bc	0.11	0.05	ab	0.03	0.03	c	0.16	0.10	a	0.10	0.03	b	5.53	0.00
Zn	4.62	4.13	b	5.60	5.23	b	6.37	7.06	b	5.30	5.85	b	15.5	14.6	a	5.13	5.17	b	3.08	0.02

Table 4. Partial least square regression analysis (PLSR) coefficients showing how deposition of different metals depends on wind velocity (Wind), air temperature (Air T°), relative humidity (RH%), and precipitation (Prec.). Values as 0.000 are not equal to 0 but less than 0.001.

	F	P	r ²	Const.	Wind	Air T°	RH%	Prec.
Cd	10.74	0.002	15	-0.000	0.000	0.000	-0.000	-0.000
Cu	34.41	0.000	53	-0.732	0.486	0.048	-0.009	-0.001
Ni	19.98	0.000	40	-0.121	0.086	0.007	-0.001	-0.000
Pb	3.24	0.077	5	-0.001	0.002	0.000	-0.000	-0.000
Zn	128.18	0.000	67	-1.159	0.801	0.071	-0.012	-0.002

(Herms and Mattson 1992). Under these conditions, the planting of profligate/opportunistic species with a high assimilation rate is highly advisable because they allow maximization of both CO₂ assimilation and growth. Among the species studied here, *E. × ebbingei* and *L. nobilis* outperformed the others under non-limiting moisture in terms of carbon uptake and growth. In detail, *E. × ebbingei* showed the highest CO₂ assimilation rates per unit leaf area throughout the day, and particularly during the afternoon, when photosynthesis of most of the other species was greatly constrained by midday depression. Furthermore, this species, when grown with adequate moisture, can sustain these rates over a large leaf area, resulting in higher daily carbon uptake per individual plant compared to most of the other species investigated. On the contrary, other species, such as *P. × fraseri* and *L. japonicum*, had high CO₂ assimilation rates but low total leaf area per plant, resulting in lower daily CO₂ assimilation and growth rates compared to *E. × ebbingei* and *L. nobilis*.

It must also be considered that, for a given increase in plant biomass, species that invest more in leaves than in woody biomass store carbon for a shorter period, particularly in the urban environment where fallen leaves are artificially removed

Table 5. Clusters derived from cluster analysis and factor loadings of two factors identified from factor analysis for metals on leaves during the entire sampling period. Varimax rotation was applied to factor analysis.

	Cluster	Factor 1	Factor 2
Cd	2	0.25	0.83
Cu	1	0.97	0.09
Ni	1	0.96	0.01
Pb	2	-0.02	0.89
Zn	2	0.88	0.28

(Nowak et al. 2002). Therefore, species such as *V. lucidum*, with a high carbon allocation to leaves, may not be advisable for maximizing carbon storage.

Drought stress reduces plant carbon gain by acting on both stomatal and mesophyll conductance, and affecting the activity of Rubisco and the rate of ribulose regeneration (Chaves et al. 2002; Medrano et al. 2002). Furthermore, drought can alter photosynthate partitioning to primary (growth) and secondary (defense) metabolism, thereby decreasing the amount of assimilated carbon stored as woody biomass. For example, in a recent experiment, when shrubs of the isoprene emitter species *Platanus × acerifolia* were moved from a moist to a dry environment, the amount of photosynthate allocated to isoprene (a defense compound) was unchanged,

despite a five-fold reduction of photosynthesis (Tattini et al. 2015). Clearly, if photosynthesis is impaired, secondary metabolism is fed by reserves that then become unavailable for growth (Loreto and Schnitzler 2010); the more the stress is perceived as severe by the plant, the more primary metabolism is likely to be affected. Finally, when a plant is killed by drought, the carbon stored in its woody organs will return to the atmosphere, unless the woody material is properly managed (i.e., soil amendment) (Nowak and Crane 2002). Therefore, when planting harsh sites with the aim of improving CO₂ sequestration, it is necessary to plant species that are “horticulturally tolerant”; that is, drought-tolerant species capable of sustaining carbon gain and growth under stressful conditions (see Fini et al. 2013), or their benefits will be greatly reduced or neglected.

Species with conservative water use, such as the Mediterranean shrubs *A. unedo* and *V. tinus*, had lower stomatal conductance and carbon assimilation than profligate/opportunistic species under non-limiting water availability, but maintain relatively high leaf gas exchange during drought. The capacity of *A. unedo* and *V. tinus* to preserve the functionality of the photosynthetic apparatus during drought is confirmed by the lower decline in the maximum quantum yield of photosystem II (Fv/Fm) compared to profligate/opportunistic species, such as *E. × ebbingei*, *L. japonicum*, and *V. lucidum*. When CO₂ assimilation is largely constrained by water shortage, as occurs in profligate/opportunistic species, usage of irradiance in the photosynthetic process is also reduced, and leaves may experience severe excess light stress, leading to photo-inhibition and photo-oxidative damage (Chaves et al. 2008; Fini et al. 2012). The large drop in Fv/Fm observed in *E. × ebbingei*, *L. japonicum*, and *V. lucidum* indicates that photo-inhibition probably occurred in leaves of these species during drought (Percival 2005; Kalaji et al. 2012; Fini et al. 2013).

Although severe drought induced a decline of photosynthesis, *P. × fraseri* displayed the lowest decline in Fv/Fm. The high concentration of anthocyanins in this red-leafed species may explain its notable capacity to counter excess light stress and avoid severe photo-inhibition during water stress (Tattini et al. 2015).

In summary, the results of Experiments 1 and 2 indicate that, if maximizing CO₂ assimilation and

carbon storage is a project goal, species with a profligate/opportunistic water use and fast growth rate, such as *E. × ebbingei* and *L. nobilis*, should be planted in moist/irrigated planting sites, while species with a conservative water use or effective photoprotective mechanisms (*A. unedo*, *V. tinus*, *P. × fraseri*) are recommended for drought-prone environments.

A potential limitation of this study is that the life spans of the different species were not considered. Thus, while the results may be valuable for understanding CO₂ assimilation and carbon storage of different shrub species per unit time, they may not reflect the total carbon stored over the whole life of the plant. However, with the exception of *A. unedo*, which can live for up to 400 years, the life span of the other shrub species is similar, ranging from 20 to 50 years (Clemson Cooperative Extension 2015; AUB Landscape Plant Database 2016; European Trees 2016). In addition, all the three species (*A. unedo*, *V. tinus*, *P. × fraseri*) have similar size at maturity, ranging from 3 to 7 m (Brikell 2008).

These life spans fit the normal turnover of green areas in urban environments well, where plants often decline because of multiple stresses and are not likely to complete their life cycles (Kozłowski 1999).

Experiment 3: Leaf Deposition of Trace Metals

Leaf deposition per unit leaf area of Pb differed among species, while for the other metals adsorption was not affected by species. In addition, the species that showed the highest quantities of Pb (*L. japonicum*, *V. lucidum*, and *E. × ebbingei*) had no specific common characteristics (e.g., presence of leaf trichomes, high growth parameters) differentiating them from the other species (Table 6; Figure 7). *Elaeagnus × ebbingei* is the only species with a visible presence of trichomes that can be related to a higher capacity of pollutant retention (Beckett et al. 2000). *Ligustrum japonicum* and *V. lucidum*, unlike the other species, showed the presence of secretory glands, especially on the lower leaf surface (Figure 8). Secretory activity is evident by the light color of the glands, indicating of the presence of salts (Sakai 1974), which may be related to the higher values of Pb in the aforementioned species. Lavid et al. (2001) found that Pb may be excreted by leaves through epidermal glands, and this is consistent with the limited

seasonal variability observed across samplings in this study. In fact, the secretory activity may be considered independent from meteorological conditions, which on the other hand, affect the leaf depositions of metals from the PM in the air.

The results of whole-plant leaf deposition suggest that total leaf area had a greater effect on the interception capacity of shrubs than the morpho-anatomical traits of individual leaves (Table 6). Rapid-growing species with large total leaf area, and high LAI, crown diameters, and height, have been reported to adsorb more pollutants than slower-growing species (Bunzl et al. 1989; Beckett et al. 2000; Fowler et al. 2003; Freer-Smith et al. 2005).

The large variation in metal deposition during samplings confirmed that leaves are a temporary retention site for pollutants (Nowak et al. 2006). The results of partial least square regression, especially for Cu and Zn (r^2 above 50), make it possible to predict the deposition of metals with moderate accuracy by using climatic parameters (Table 4). The negative relationship between metal deposition and total rainfall indicates that metals on leaves decreased as rain increased, confirming the washout effect that rain quickly plays on pollutants adsorbed on leaf surfaces (Nowak et al. 2006). The inverse relationship between metal deposition and RH% may be considered more a consequence of the strong relationship between rain and RH% than a direct effect of RH% on metal deposition. Despite this, it is reported that RH% causes an increase in PM diameter and that when PM diameters are greater than 0.1 μm , an increase in diameter leads to an increase of deposition velocities (Fowler et al. 2003; Litschke and Kuttler 2008). The positive relationship between metals and wind speed could be due to the fact that increasing wind speed leads to increasing deposition velocities of PM (Beckett et al. 2000).

Temperature was also found to have a positive relationship with the deposition of pollutants, as previously reported (Cavanagh et al. 2009).

Washout by Rain of Metals from the Canopy

The protocol used for rainwater collection does not allow exclusion of possible contamination of samples because of bird droppings, insects, or other undesired animals. Therefore, researches focused mainly on differences between the presence and absence of plants, rather than on differences among species. The greater quantities of metals in containers under shrubs is further demonstration of the role of shrubs in intercepting air pollution (Nowak et al. 2000; Bealey et al. 2007). In addition, the effect of washout caused by rain, and the consequent deposition on the ground of metals, helps reduce the resuspension of pollutants in the air.

Metal Source Identification

The results of correlation, FA, and CA were compared in order to identify the possible source(s) of different metals. Cu, Ni, and Zn have high loadings in factor 1 of FA and are all strongly correlated (Table 5). Meanwhile CA clustered Cu and Ni in a different group compared to Zn; Zn was then excluded from the analysis. The relationship between Cu and Ni is then supported by three different statistical methods. Cd and Pb are dominant in factor 2 of FA and are significantly correlated. Moreover, in CA, they are clustered together in Group 2. Thus, the relationship between Cd and Pb is confirmed by three different methods. Considering these results, two different groups were created: Group 1 (Cu and Ni) and Group 2 (Pb and Cd). All four of these metals have previously been attributed to traffic emissions (Pant and Harrison, 2013; Valotto et al. 2015). This division

Table 6. Leaf area of the whole plant (LA), leaf area index (LAI), average lamina size (LS), number of leaves per plant (N°L), crown diameter (CD), and height of plants (HP), of six shrub species planted in a field near a heavily polluted road (Exp. 3). Values are means \pm SD. ANOVA tests (A) of differences between species are also included.

	<i>V. lucidum</i>			<i>A. unedo</i>			<i>P. × fraseri</i>			<i>L. nobilis</i>			<i>E. × ebbingei</i>			<i>L. japonicum</i>			F	P
	Mean	SD	A	Mean	SD	A	Mean	SD	A	Mean	SD	A	Mean	SD	A	Mean	SD	A		
LA (m ²)	5.01	0.67	bc	3.99	0.39	ab	5.58	1.58	c	2.9	0.24	a	9.43	0.68	d	3.87	0.56	ab	24.3	0.00
LAI	5.43	1.12	c	3.9	0.97	a	5.32	0.88	c	4.47	1.26	b	6.9	1.14	d	4.15	1.08	ab	17.6	0.00
LS (cm ²)	31.9	5.66	c	10.5	2.68	a	19.5	1.45	b	20.8	4.17	b	12.6	1.49	a	17.3	2.04	b	31.6	0.00
N°L	1608	265	ab	3967	898	d	2876	226	c	1443	271	a	7611	1045	e	2268	284	bc	88.1	0.00
CD	102	7.06	d	93.1	6.45	c	102	13.2	d	71.6	12.2	a	108	11.7	d	81.5	14.4	b	25.2	0.00
HP	146	16.8	b	136	11.5	a	194	28.7	e	166	9.99	cd	170	31.8	d	162	24.9	c	13.4	0.00

performed by statistical methods may indicate the presence of two different sources for the studied metals. Cu use is widespread in Pescia in the floriculture industry. Ni is a micronutrient for plants and its presence may be related to soil fertilization practices (Barker and Pilbeam 2006). On this basis, Group 1 may be classified as an “agriculture-related group.” Group 2 (Pb and Cd) can be attributed to traffic emissions (Pulles et al. 2012). Since one of the aims of the research was to identify the possible pollution source, it appears important to consider not only the singular metal itself, but also relationships between different metals and characteristics of the sampling site. By comparing different methods the results are more reliable.

CONCLUSION

The present investigation was aimed at evaluating the potential of some shrubs to sequester air pollutants and assimilate CO₂ from the atmosphere. This information may be relevant for planners and technicians involved in the management of urban green areas.

Elaeagnus × ebbingei had the highest carbon storage under optimal water availability but revealed its inability to thrive also in drought-prone environments. If water availability is a limiting factor in a site where a planting is designed, other species, such as *P. × fraseri*, *A. unedo*, and *V. tinus*, should be preferred.

A clear discrimination among species regarding their relative capacity for deposition of trace metals was not obtained. However, *E. × ebbingei* had the highest whole-plant leaf deposition of all the measured metals, primarily because of its extremely fast growth rate.

The study authors are aware that only a limited number of shrub species were tested in this study of CO₂ uptake and air pollution interception potential, and further research is needed to evaluate different species for different climatic areas. However, the results may offer initial insights regarding CO₂ assimilation and trace metal deposition potential of widely used shrubs for urban green areas in southern Europe.

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Résumé. Trois expériences indépendantes ont évalué l'assimilation du CO₂ et le dépôt de métaux sur les feuilles de sept espèces d'arbustes à feuilles persistantes (*Arbutus unedo* L., *Elaeagnus × ebbingei* L., *Laurus nobilis* L., *Ligustrum japonicum* Thunb., *Photinia × fraseri* Dress., *Viburnum tinus* subsp. *lucidum* L. et le *Viburnum tinus* subsp. *tinus* L.). L'assimilation du CO₂ et l'allocation du carbone ont été caractérisées sur des plants en pot sous des conditions optimales de disponibilité en eau en 2011 (Exp. 1) et sous des conditions de sécheresse en 2012 (Exp. 2). Une troisième expérience (Exp. 3) a mesuré le dépôt saisonnier sur les feuilles de Cd, Cu, Ni, Pb et Zn en 2011 sur des plantes transplantées à proximité d'une route à quatre voies.

Le *E. × ebbingei* a démontré la plus élevée assimilation du CO₂ sous les conditions optimales en eau, mais une des plus faibles lors des conditions de sécheresse (Exp. 1, 2). A l'inverse, *P. × fraseri* avait une assimilation moyenne du CO₂, mais celle-ci diminuait moins en période de sécheresse par rapport aux autres espèces. Dans l'expérience 3, *E. × ebbingei* a démontré le taux de dépôt de métaux le plus élevé, principalement en raison de sa plus grande surface foliaire. Les pluies fréquentes et le pourcentage de RH réduisaient le dépôt de métaux tandis qu'une plus grande vitesse des vents et la température de l'air augmentaient le dépôt sur les feuilles. Les espèces qui réduisent considérablement l'assimilation du CO₂ sous des conditions de sécheresse (*V. tinus* subsp. *lucidum*, *L. japonicum*, *E. × ebbingei*) ne sont pas recommandées dans les aménagements sujets à la sécheresse, alors que les espèces "mésiques" tolérantes à la sécheresse (*P. × fraseri*) devraient être préférées. *E. × ebbingei* pourrait être utilisé pour optimiser le dépôt de métaux. Les trois expériences fournissent des indications utiles concernant particulièrement l'assimilation du CO₂ (Exp. 1, 2) et la capacité de réduction de la pollution atmosphérique (Exp. 3) d'une gamme d'arbustes largement utilisés dans les zones urbaines et la planification du verdissement le long des routes en Europe méridionale.

Zusammenfassung. Drei unabhängige Experimente untersuchten die CO₂ Assimilation und Schwermetallbelastung auf den Blättern von 7 immergrünen Sträuchern (*Arbutus unedo* L., *Elaeagnus × ebbingei* L., *Laurus nobilis* L., *Ligustrum japonicum* Thunb., *Photinia × fraseri* Dress., *Viburnum tinus* subsp. *lucidum* L., und *Viburnum tinus* subsp. *tinus* L.). Die CO₂ Assimilation und die Kohlenstoffeinlagerung wurde in 2011 (Experiment 1) unter optimaler Wasserverfügbarkeit und in 2012 (Exp. 2) unter Trockenheitsbedingungen bei getopferten Pflanzen bestimmt. Ein drittes Experiment maß die saisonale Ablagerung von Cd, Cu, Ni, Pb, und Zn in 2011 auf Blättern von Pflanzen, die in die Nähe einer vierspurigen Straße gepflanzt wurden.

Elaeagnus × ebbingei zeigte die höchste CO₂ Assimilation unter optimalen Wasserbedingungen, aber eine der schlechtesten unter Trockenheitsbedingungen (Exp. 1, 2). Entgegengesetzt hatte *P. × fraseri* eine mittlere CO₂ Assimilation, aber sie sank während der Trockenheit im Vergleich zu anderen Arten. Im dritten Experiment zeigte *E. × ebbingei* die höchste Schwermetallablagerung, hauptsächlich wegen der großen Blattfläche. Großer Regenfall und RH% reduzierte die die Metallablagerungen, während größere Windbewegung und Lufttemperatur die Blattablagerungen vermehrten. Arten, die ihre CO₂ Assimilationen unter Trockenheitsbedingungen drastisch reduzieren (*V. tinus* subsp. *lucidum*, *L. japonicum*, *E. × ebbingei*), sind in einer Umgebung mit häufiger Trockenheit

nicht zu empfehlen, wo trockenheitstolerante „mesische“ Arten (*P. × fraseri*) bevorzugt werden sollten. *E. × ebbingei* könnte verwendet werden, um die Ablagerung von Metallen zu optimieren. Die drei Experimente liefern nützliche Einsichten über die CO₂ Assimilation (Exp. 1,2) und Minderung der Luftverschmutzung (Exp. 3) von flächendeckend verwendeten Sträuchern bei Pflanzungen in urbanen Räumen und Planung von Straßenbegleitgrün in Südeuropa.

Resumen. Tres experimentos independientes evaluaron la asimilación de CO₂ y la deposición de metales en la hoja de siete especies de arbustos de hoja perenne (*Arbutus unedo* L., *Elaeagnus × ebbingei* L., *Laurus nobilis* L., *Ligustrum japonicum* Thunb., *Photinia fraseri* × Dress., *Viburnum tinus* subsp. *lucidum* L. y *Viburnum tinus* subsp. *tinus* L.). La asimilación de CO₂ y la asignación de carbono se determinaron en 2011 (Exp.1) bajo óptima disponibilidad de agua y en el año 2012 (Exp.2) en condiciones de sequía en las plantas en maceta. Un tercer experimento (Exp.3) midió las deposiciones de hojas estacionales de Cd, Cu, Ni, Pb y Zn en 2011 en las plantas trasplantadas en la proximidad de una carretera de cuatro carriles.

E. × ebbingei mostró la mayor asimilación de CO₂ con buena disponibilidad de agua, pero uno de los más bajos en condiciones de sequía (Exp.1, 2). A la inversa, *P. × fraseri* tuvo asimilación de CO₂ intermedio pero disminuyó durante la sequía en comparación con las otras especies. En el Experimento 3, *E. × ebbingei* mostró la mayor deposición de metales, principalmente debido a su mayor área foliar. Las mayores precipitaciones y la humedad relativa disminuyeron las deposiciones de metal, mientras que una mayor velocidad del viento y la temperatura del aire aumentaron las deposiciones de las hojas. Las especies que redujeron drásticamente la asimilación de CO₂ en condiciones de sequía (*V. tinus* subsp. *lucidum*, *L. japonicum*, *E. × ebbingei*) no se recomiendan en entornos propensos a la sequía, donde las especies "mésicas" tolerantes a la sequía (*P. × fraseri*), deben ser preferidas. *E. × ebbingei* podría ser utilizado para optimizar la deposición de metales. Los tres experimentos proporcionan información útil sobre todo acerca de la asimilación de CO₂ (Exp. 1, 2) y la mitigación de la contaminación del aire (Exp. 3) de arbustos ampliamente utilizados para su aplicación en áreas urbanas y la planificación del enverdecimiento a borde de carretera en el sur de Europa.