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1 Aerodynamic characterisation of green wall vegetation based on plant 2 morphology: an experimental and computational fluid dynamics 3 approach

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12 Abstract

13 The installation of urban green infrastructure, particularly green walls, has proven to be an effective strategy for
14 the mitigation of particulate matter (PM) pollution and the urban heat island effect. For the interaction between
15 vegetation, PM and the local microclimate, wind flow is the main driving force. In order to investigate these
16 interactions in detail, it is important to know how air flows through vegetation. This study proposes a method
17 based on the Darcy-Forchheimer equation, where vegetation is considered as a porous medium and several plant
18 species and the effects of plant morphological characteristics are examined both experimentally and using
19 computer simulations. Results showed that the Darcy-Forchheimer model is a simple and robust way to describe
20 air flow through vegetation regardless of its morphology. This research provides a new vision on studying
21 aerodynamic properties of vegetation in relation to their morphology and provides opportunities for modelling
22 the interaction between vegetation and its environment.

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26 Keywords: Darcy-Forchheimer; CFD; wind tunnel; air flow; vegetation morphology

27 Nomenclature table

A	Area	m ²
FLS	Functional leaf size	(-)
K	Permeability	m ²
K _i	Inertial permeability	m
LAD	Leaf area density	m ² m ⁻³
LDI	Leaf dissection index	(-)
LS	Leaf size	m ²
LSS	Leaf surface structure	(-)
LR	Leaf ratio	(-)
M	Mass	kg
P	Perimeter	m
PAD	Plant area density	m ² m ⁻³
Q	Flow rate	m ³ s ⁻¹
R ²	Coefficient of determination	(-)
SLA	Specific leaf area	m ² kg ⁻¹
q	Flux	m s ⁻¹
v	Fluid velocity	m s ⁻¹
V	Volume	m ³
ΔP	Pressure loss	Pa
ΔP _{norm}	Normalised pressure loss	(-)
Δx	Distance	m
ρ	Density	kg m ⁻³
φ	Porosity	(-)
τ	Kendall rank correlation coefficient	(-)
μ	Viscosity	Pa s

28 **1 Introduction**

29 Over the last decade green walls and façades have become an increasingly popular type of urban green
30 infrastructure (UGI). They provide an efficient way to introduce green environments in the limited space
31 typically available in the urban environment provided they are thoughtfully planned (Cameron &
32 Blanuša, 2016). UGI is able to fulfil many different ecosystem services (ES), including air quality
33 improvement (by particulate matter [PM] deposition) (Abhijith et al., 2017; Gallagher et al., 2015),
34 environmental cooling, carbon sequestration, increasing biodiversity and also serving social and cultural
35 purposes (Madre, Clergeau, Machon, & Vergnes, 2015; Tallis et al., 2015; White & Gatersleben, 2011).
36 Improvements in air quality and environmental cooling are significantly driven by physical interactions
37 between the plants and air flowing over and through them. Vegetation affects local air flow and it
38 therefore alters wind-driven processes such as PM deposition (K. P. Beckett, Freer-Smith, & Taylor, 1998;
39 Janhäll 2015; Litschike & Kuttler, 2008; Nowak, Crane, & Stevens, 2006; Ottelé, van Bohemen, & Fraaij,
40 2010; Pugh, Mackenzie, Whyatt, & Hewitt, 2012; Tallis et al., 2015), evaporation and heat transfer
41 (Demuzere et al., 2014; Hunter et al., 2014; Perini, Ottelé, Fraaij, Haas, & Raiteri, 2011; Tallis et al., 2015).

42 Several investigations have demonstrated the influence of vegetation on local air flow, based either on
43 field measurements (Hong, Lin, Wang, & Li, 2012) or wind tunnel studies using plant imitators
44 (Buccolieri, Gromke, Di Sabatino, & Ruck, 2009; Endalew et al., 2009; Gromke 2011 and references
45 therein; Gromke et al., 2016) or real plants (Molina-Aiz, Valera, Álvarez, & Madueño, 2006; Lin &
46 Khlystov, 2012a; Sase, Kacira, Boulard, & Okushima, 2012). Some studies were based on numerical
47 airflow models (Connell, Endalew, & Verboven, 2011; De Maerschalck, Maiheu, Janssen, & Vankerkom,
48 2010; Endalew et al., 2009; Hong et al., 2012) or empirical models (Raupach, Woods, Dorr, Leys, &
49 Cleugh, 2001; Tiwary, Morvan, & Colls, 2005; M. Lin, Katul, & Khlystov, 2012b) or a combination of
50 multiple approaches. Most model-based studies considered vegetation as rigid structures, omitting
51 realistic descriptions of micro-scale parameters, such as porosity and leaf morphology *within* vegetation
52 stands. This simplification of the vegetation structure increases the uncertainty in study outputs.
53 Moreover, few studies have been performed on the dynamics of air flowing *through* vegetation. This
54 knowledge gap can be explained by the complexity of a vegetation stand and the physics involved, with
55 randomly distributed air spaces between inhomogeneous foliage elements, causing flows with high
56 turbulence (Tiwary et al., 2005). Also, drag forces may affect the airflow. The aerodynamic effects of
57 plant stands are often expressed in terms of resistance to the flow or a drag coefficient (Molina-Aiz et
58 al., 2006; Sase, Kacira, Boulard, & Okushima, 2012). Endalew et al. (2009) modelled the interaction
59 between airflow and vegetation based on leaf drag/area and volume weighted drag (based on porosity),
60 but for both methods detailed 3D geometrical information is needed (Endalew et al., 2009 and
61 references therein). However, given the complexity of the systems under study, some simplification is
62 recommended when modelling the dynamics of air flowing through vegetation.

63 This paper proposes a method based on the Darcy-Forchheimer equation (Nield, 2000). This equation
64 originates from earth sciences where it is used to describe the flow of water through porous soils, but it
65 has been proven to be applicable in many more research fields involving fluid flow (including air) through
66 porous media (Bartzanas, Kittas, & Boulard, 2002; Bartzanas, Katsoulas, & Kittas, 2012; Mattis, Dawson,
67 Kees, & Farthing, 2012a; Miguel, van de Braak, & Bot, 1997; Verbruggen et al., 2016). The Darcy-
68 Forchheimer equation is an extension of the simple Darcy equation that relates the pressure drop
69 observed when a fluid flows through a porous medium to the flow rate of the fluid. The Forchheimer-
70 drag includes inertial effects that occur at high flow rates. Molina-Aiz et al., (2006) found that, when air

71 velocity and pressure drop are plotted, the Darcy-Forchheimer equation seemed to have the best fit
 72 with their wind tunnel data compared to drag equations.

73 The objectives of this study are (1) to assess whether the Darcy-Forchheimer equation is applicable to
 74 describe air flow through vegetation; (2) to investigate whether there are significant differences in the
 75 corresponding aerodynamic parameters between plant species and if so, (3) to check whether
 76 aerodynamic parameters can be related to plant species morphology.

77 2 Materials and methods

78 2.1 Selection of species and morphology

79 In order to cover the high variability in morphology typical for green wall vegetation, a broad range of
 80 plant species were selected according to their occurrence in green walls. Climbers/creepers as well as
 81 herbaceous species were chosen to cover both soil-bound and substrate-bound vertical garden systems,
 82 and a broad range in leaf morphological characteristics, like e.g. leaf size and shape (Table 1).

83 Table 1: Selected species and their morphological characteristics. Scientific name, leaf shape and length
 84 and width of an individual leaf are given.

Scientific name	Leaf morphology	Length x width	Scientific name	Leaf morphology	Length x width	Scientific name	Leaf morphology	Length x width
<i>Hedera helix</i>	Heart shaped	70 x 100 mm	<i>Campanula isophylla</i>	Heart shaped, serrated	40 x 45 mm	<i>Lonicera nitida</i>	Oval	15 x 5 mm
								
<i>Festuca glauca</i>	Very thin grass	200 x 2 mm	<i>Fragaria sp. 'Ostara'</i>	Ternate leaf, leaflets serrated	120 x 120 mm	<i>Berberia cordifolia</i>	Oval, large, thick	100 x 200 mm
								
<i>Geranium traversii x oxonianum</i>	Hand shaped, lobed	50 x 60 mm						
								

85

86 A series of directly measurable morphological plant parameters was examined. Morphological
 87 parameter selection was based on their potential to affect the air flow through vegetation. Each of these
 88 parameters is able to explain some aspects of leaf morphology into a single metric, which makes inter-

89 species comparison possible. An overview is given in Table 2. Methods were original unless stated
90 otherwise with a reference.

91 *Plant area density* (PAD) is the amount of one sided plant surface (including twigs; A_{plant}) per total bulk
92 volume (the volume of the cylinder holding the plants) (V_{total}) and a measure of how vegetation can fill
93 up a space. It is closely related to porosity which is the void volume fraction of a porous material (see
94 further). Plant surface was measured with a Li-3100c leaf area meter (Li-cor Inc., Lincoln, NE, USA). PAD
95 is similar to *leaf area density* (LAD), which is more commonly used in plant ecology and considers only
96 the one sided leaf surface. However, because in this study species with a highly diverse morphology
97 were used and branches also affect local air flow, PAD is a better choice, as it is the total portion of plant
98 material which contributes to pressure losses. Realistic PAD values were pursued without omitting
99 extremely sparse or dense vegetation.

100 Volumetric *porosity* (ϕ) is the void volume fraction ($1 - [V_{\text{plant}} V_{\text{total}}^{-1}]$) of a porous material. In this context,
101 it is the volume fraction occupied by the air filled spaces in between the leaves and branches of the
102 vegetation, and is typically higher than 93% (Gromke, Buccolieri, Di Sabatino, & Ruck, 2008). It should
103 not to be confused with the 'optical porosity', which is a two-dimensional parameter that ignores some
104 spatial information. Plant volume was determined through displacement by putting the vegetation in a
105 graduated cylinder filled with water. Despite being the best empirical method to estimate the volume
106 of an irregular shaped object, this method is quite crude. Therefore, as a quality control for each species,
107 the volume was correlated with dry mass.

108 The *leaf ratio* (LR) is the amount of leaf material with respect to the total amount of plant material,
109 calculated both in mass or volume units. The mass based leaf ratio was assessed by weighing leaves and
110 stems that were dried in an oven (Memmert UF 260, Memmert, Eeklo, Belgium) separately on an
111 analytical balance (S-234, Denver Instruments, NY, USA, accuracy= 0.1 mg). These masses were
112 averaged for every species, with between 8 and 15 replicates per species, and the correlation between
113 weight and volume based leaf ratios was identified.

114 *Specific leaf area* (SLA) (Cornelissen et al., 2003) is expressed as the one-sided leaf surface (A_{leaf}) with
115 respect to leaf dry mass (M_{leaf}). Dry matter was measured on the analytical balance (accuracy = 0.1 mg).
116 This parameter can be used as a measure for leaf structure/thickness (Vile et al., 2005). One average
117 SLA value was calculated for every species out of 8 to 15 replicates. Including mass in a dataset can be
118 beneficial, as most other parameters only cover a two dimensional representation of vegetation, while
119 mass is also indicative for the occupied volume.

120 *Leaf dissection index* (LDI) can be described as the ratio of a leaf's perimeter (P_{leaf}) to the square root of
121 its surface area ($\sqrt{A_{\text{leaf}}}$) (McLellan & Endler, 1998). This is a dimensionless parameter which is used to
122 describe leaf shape. Plant species that have strongly incised or elongated leaves have a higher LDI than
123 plants which have leaves with a more circular shape. The minimum value for LDI is 3.545, this is the value
124 for a perfect circle. The square root of leaf area is taken in order to normalize for leaf size, as the test
125 plants had leaves with similar shapes but different sizes. LDI was determined for every species by
126 measuring six undamaged leaves over their leaf size range, of which the mean value was taken as the
127 species' LDI. It should be noted that for *Fragaria* sp. only one leaflet was considered, as one full leaf
128 consist of three fully separated petioles.

129 *Functional leaf size* (FLS) is the surface of the largest full circle that can be drawn into a leaf (A_{circle}),
130 normalized by leaf surface (A_{leaf}). This parameter is the major determinant of the leaf boundary layer

131 and thus strongly influences PM deposition and transpiration. Per species one value for FLS was
 132 retrieved by taking the mean of six leaves of varying sizes as used for LDI. For species of which the leaves
 133 were consisting of multiple leaflets (e.g. *Fragaria* sp.), again only the single leaflet was considered.

134 *Leaf size* (LS) is the one-sided surface area of a leaf (A_{leaf}). Per species one value for leaf size was taken
 135 as an average from six individual undamaged leaves over their size range.

136 *Leaf surface structure* (LSS) was assessed by categorization. Hairy leaves and leaves with rough
 137 (irregular) surfaces, at microscopic level, are known to sustain a thicker boundary layer than non-hairy
 138 and smooth leaves. This could have an influence on local air flow (turbulence), PM deposition and gas
 139 exchange processes. Of each species, SEM (scanning electron microscope) images were taken with a
 140 Quanta FEG 250 at high vacuum. For full explanation of the methodology we refer to Castanheiro,
 141 Samson, & De Wael (2016). Surface characteristics were divided into 3 categories and categorized as
 142 very hairy and/or irregular (considered as '2'), semi-hairy and/or semi-irregular ('1'), and smooth ('0').

143 For LR, SLA, LDI, FLS and LS results are presented as mean values with a standard deviation.

144 Table 2: Overview of determined plant morphological parameters. V: volume (m^3), A: leaf surface (m^2),
 145 P: leaf perimeter (m), M: dry leaf mass (kg)

Parameter	Formula	Unit
Plant Area Density (PAD)	$PAD = \frac{A_{plant}}{V_{total}}$	$m^2 m^{-3}$
Porosity (ϕ)	$\phi = 1 - \left(\frac{V_{plant}}{V_{total}}\right)$	dimensionless
Leaf Ratio (LR)	$LR = 1 - stem\ ratio$	percentage
Specific Leaf Area (SLA)	$SLA = \frac{A_{leaf}}{M_{leaf}}$	$m^2 kg^{-1}$
Leaf Dissection Index (LDI)	$LDI = \frac{P_{leaf}}{\sqrt{A_{leaf}}}$	dimensionless
Functional Leaf Size (FLS)	$FLS = \frac{A_{circle}}{A_{leaf}}$	dimensionless
Leaf size (LS)	A_{leaf}	m^2
Leaf Surface Structure (LSS)		

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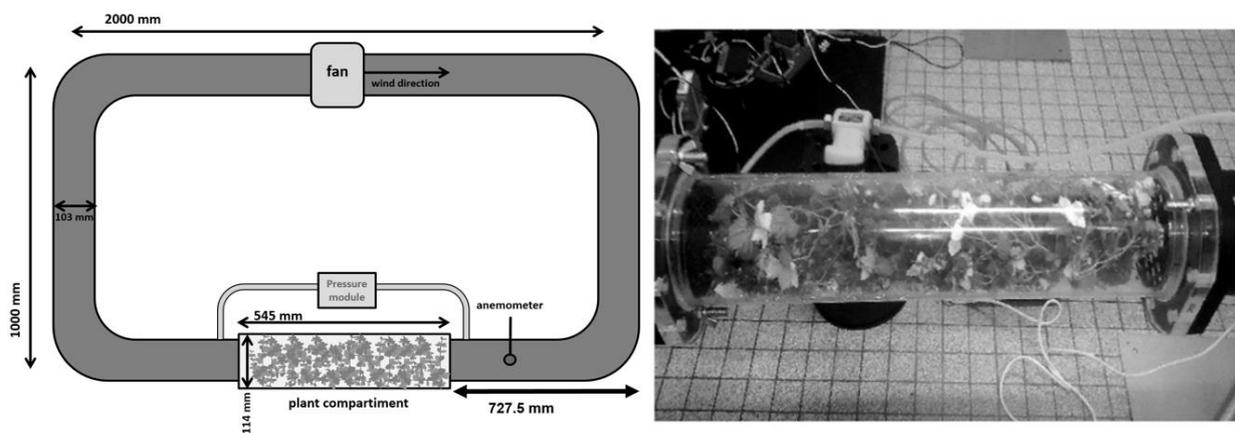
148 2.2 Experimental procedure

149 2.2.1 Wind tunnel experiments

150 Pressure losses over vegetation were measured in a continuous flow, closed circuit wind tunnel (Fig. 1
 151 left). The circuit consists of circular air ducts (inner diameter 103 mm) with a total length of 6 m. A plant
 152 compartment consisting of a transparent pipe (length 545 mm, inner diameter 114 mm) was fitted in
 153 the closed circuit. Air movement was created by a continuously controllable inline duct fan which could
 154 induce wind speeds varying from 1.14 to 4.40 $m s^{-1}$ (for an empty compartment), which is a realistic but
 155 low range of wind speeds. Air velocity induced by the duct fan was measured by a hotwire air velocity
 156 sensor (CTV 110, KIMO Instruments, Chevry-Cossigny, France). Instead of continuously varying the
 157 airflow rate during an experiment, the power supply to the duct fan was controlled and fixed at five
 158 power supply settings: the minimum and maximum power supply (corresponding to the minimum (but

159 not zero) and maximum wind speed), and approximately the 1st, 2nd (median) and 3th quartile. The
160 advantage of selecting five specific power supplies was that for each setting, a corresponding fan curve
161 (i.e. the relation between the pressure loss and the flow rate for the fan used) could be determined from
162 all pressure loss and velocity measurements (for all species under study) corresponding to the specific
163 power supply. The fan curves could then be used as boundary conditions in the models (see further).
164 Obviously, the air velocity v is not uniform over the cross section of the flow. Mostly, average velocities
165 are used in the analysis of duct flow. Air velocity profiles were derived from measurements with the hot
166 wire velocity sensor located at different distances from the duct wall and the average air velocity was
167 calculated by integrating the profile over the flow cross section.

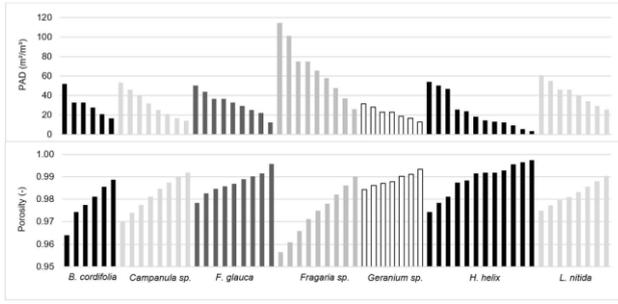
168 Freshly cut plant specimens were brought into the plant compartment of the wind tunnel and stacked
169 as homogeneously as possible with the least possible clustering (Fig. 1 right), approximating the
170 composition of the plants in a real green wall. After selecting one of the five power supply settings
171 (always from low to high), the inline duct was switched on and the system was allowed to equilibrate
172 until the pressure loss was stationary. Pressure loss (ΔP) over the plant compartment was assessed by a
173 Pressure Module (750PD2, FLUKE Corporation, Gent, Belgium) with a range of ± 7 kPa and a Pressure
174 Calibrator (717 30G, FLUKE Corporation). Test experiments showed similar results for the same
175 vegetation specimen if brought multiple times into the wind tunnel in different configurations, providing
176 that stacking was done uniformly. Only when heavily clustered, resulting in an inhomogeneous
177 distribution within the plant compartment, did the results differ from each other. Therefore, each
178 specimen was only tested once for a particular PAD under different airflow rates. Nevertheless, each
179 plant species was tested at different PADs (Fig. 2 top) and porosities (Fig. 2 bottom) by adding or
180 removing a certain amount of plant material. Vegetation was held in place by two grates, whose
181 pressure loss was measured in a separate run in order to subtract the values from the vegetation
182 measurements. Runs through an empty tunnel were used as a reference.



184 Fig. 1: Left panel: setup of the wind tunnel with the plant compartment filled with vegetation, right panel: plant
185 compartment filled with fresh *Campanula* sp.

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Fig. 2: PAD values for each run (top) and porosity values for each run (bottom). Species are sorted alphabetically.

190

191 2.2.2 Aerodynamic characterisation

192 In recent literature, an explicit approach is advised to simulate wind flow in urban areas where
 193 vegetation is present (Buccolieri et al., 2018). In most computation fluid dynamics (CFD) studies
 194 vegetation is considered as a porous medium, and additional terms are added to the momentum and
 195 turbulence equations to account for the loss of energy (refs from Buccolieri et al., 2018). While most
 196 published work focused on aerodynamic effects in the vicinity of trees, a characterisation of the air flow
 197 through vegetation was performed, and parameterised in a CFD model by adding pressure loss terms
 198 described in terms of permeability (for viscous losses) and ‘inertial permeability’ (for inertial losses).
 199 Since the intention was to correlate plant morphological parameters to aerodynamic parameters, a
 200 limited set of aerodynamic parameters was needed.

201 From pressure loss measurements over the plant compartment stacked with vegetation, it is possible to
 202 calculate the permeability of the vegetation to fluid flow. The permeability of a porous material is its
 203 ability to allow a fluid to pass through it and is closely related to its structure and porosity. For
 204 vegetation, which always has high values for porosity (often over 95%), leaf size and morphology were
 205 expected to be the determining parameters. At low fluid velocities (Reynolds numbers up to 10), the
 206 permeability of a porous material (K) can be simply derived from Darcy’s law as:

$$207 \quad K = \frac{v A \mu \Delta x}{\Delta P} \quad (\text{Eq. 1})$$

208 where v is the bulk fluid speed (m s^{-1}) (the mean speed within the pores of the porous media), μ the
 209 viscosity of the fluid (Pa s), Δx the length of the porous medium (m), A the cross section area of the flow
 210 (m^2) and ΔP the pressure loss (Pa). The simple Darcy equation assumes a linear relation of fluid flow rate
 211 (or velocity) and pressure loss, and a unique value for the permeability. However, for very high velocities
 212 (Reynolds numbers being multitudes of 10) in porous media, inertial effects become important and an
 213 inertial quadratic term, known as Forchheimer-drag, needs to be added to the Darcy equation. This term
 214 accounts for the kinetic energy and inertia of the fluid and the typical non-linear relation of fluid flow
 215 rate and the pressure drop (S. A. Mattis, Dawson, Kees, & Farthing, 2012b):

$$216 \quad \frac{\Delta P}{\Delta x} = - \left(\frac{\mu}{K} \right) q - \left(\frac{\rho}{K_1} \right) q^2 \quad (\text{Eq. 2})$$

217 where K_1 is the inertial permeability (m), ρ the fluid density (kg m^{-3}) and q the fluid flux (m s^{-1}), calculated
 218 as:

$$219 \quad q = v \varphi \quad (\text{Eq. 3})$$

220 where φ represents the porosity (-). The term $\frac{\rho}{K_1}$ is called the Forchheimer-drag coefficient. Notice that
221 in all calculations, the average air velocity (see section 3.2) was used.

222 For each of the species, the pressure loss was measured as a function of average air velocity. The
223 permeability (K) and Forchheimer-drag ($\frac{\rho}{K_1}$) were obtained by regression of Eq. 2. Obviously, the amount
224 of plant material brought in the compartment affects the PAD, the porosity and the aerodynamic
225 parameters. Experiments were repeated with 8 to 15 different PAD's per species, depending on how
226 easily the vegetation could be stacked and how long the branches stayed fresh (see Fig. 2). A wind tunnel
227 experiment for one species took approximately one day. This was no problem with species such as
228 Hedera, but more sensitive species, for example Fragaria became weak towards the end of the
229 experiment so fewer wind tunnel tests could be performed.

230 In some cases, regression led to negative values for permeability and/or inertial permeability. This was
231 solved by modifying the polynomial (Eq. 2) to a linear equation as follows:

$$232 \quad \frac{\Delta P}{\Delta x \ q} = - \left(\frac{\mu}{K} \right) - \left(\frac{\rho}{K_1} \right) q \quad (\text{Eq. 4})$$

233 K and $\frac{\rho}{K_1}$ could then be retrieved by simple linear regression. Samples for which parameters remained
234 negative were left out, as these physical impossible outcomes were attributed to flaws in those
235 particular runs. In order to compare the aerodynamic characteristics of different species under similar
236 conditions of wind velocity and PAD, pressure losses were normalised for wind velocity and PAD, so that
237 a dimensionless, normalized pressure loss (ΔP_{norm}) can be defined as:

$$238 \quad \Delta P_{\text{norm}} = \frac{\Delta P}{\rho \ v^2 \ \text{PAD} \ \Delta x} \quad (\text{Eq. 5})$$

239 where $\rho \ v^2$ is a measure of kinetic energy per unit volume, also called dynamic pressure (Pa).

240

241 **2.2.3 Calculations and statistical analysis**

242 All calculations and analysis were performed in R version 3.3.1 (<https://cran.r-project.org/>) and an Excel
243 spreadsheet. Within species morphological parameters, significant differences are identified with a
244 Tukey-HSD test and these differences are indicated with the letters A, B and C. Due to the nature of the
245 permeability and Forchheimer-drag data we performed a log (ln) transformation for normalization.
246 Herewith an ANOVA was executed for looking at the variance between permeability and Forchheimer-
247 drag between species. A Tukey-HSD test was performed to see where the differences are between
248 species. A Kendall rank correlation test was performed for testing differences between permeability and
249 Forchheimer-drag against porosity and PAD. A Kendall rank correlation test is comparable to a
250 Spearman's test but stronger and more robust considering outliers as our data is highly skewed. Larger
251 tau-values (always between 0 and 1) mean stronger correlations. Finally, to explain variance in
252 permeability and Forchheimer-drag, linear models of the mean and median of permeability and
253 Forchheimer-drag against SLA, LDI, FLS, LR, LS and LSS were made.

254

255 2.3 Numerical modelling

256 To test whether the aerodynamic parameters derived from the wind tunnel experiments can be used in
257 a modelling approach, a numerical model of the wind tunnel was developed. The advantage of a
258 numerical modelling approach such as CFD is that geometrical effects (realistic green walls with complex
259 geometries) as well as (dynamic) climate boundary conditions can be tested. CFD simulations were
260 conducted using the commercially available software package COMSOL Multiphysics® version 5.2a
261 (COMSOL Inc., MA, USA). The geometry consisted of the entire wind tunnel (Fig. 3) including the plant
262 compartment (which is slightly wider than the other ducts). Considering the symmetry of the geometry,
263 only half of the geometry was meshed as shown in the figure. The computational grid consisted of
264 approximately 170,000 tetrahedral cells with a boundary layer at the duct walls. Grid size independency
265 was ensured by gradually refining the mesh until further refinement did not affect the results. The
266 average mesh quality of the geometry was 0.72. Mesh quality is a compound quantity representing some
267 quality attributes of the mesh elements (skewness, volume versus circumradius, growth rate, ...). A
268 quality of 1 indicates the best possible fit and an optimal element. At the other end of the interval, 0
269 represents a degenerated element. Under the flow rate conditions, a turbulent flow regime is expected
270 and a k- ω turbulent, incompressible flow model at standard conditions (101325 Pa, 293.15 K) was
271 considered in the ducts. In contrast to the laminar flow model, the k- ω model solves for extra variables:
272 the turbulence kinetic energy and the rate of dissipation of turbulence kinetic energy ω . A steady-state
273 solution was generated with a direct stationary solver (relative tolerance 0.001), by solving the
274 governing equations of momentum and mass continuity in the wind tunnel ducts (Comsol 2017):

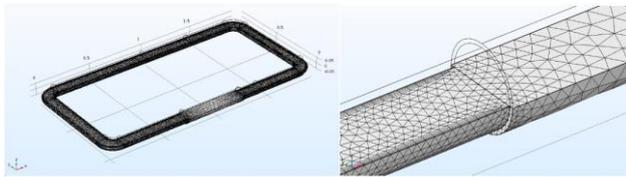
$$275 \quad \rho(\mathbf{u} \cdot \nabla)\mathbf{u} = \nabla \cdot \left(-P\mathbf{I} + (\mu + \mu_T)(\nabla\mathbf{u} + (\nabla\mathbf{u})^T) \right)$$

$$276 \quad \rho\nabla(\mathbf{u}) = \mathbf{0}$$

277 With ρ the air density (1.2044 kg m⁻³), \mathbf{u} the velocity vector (m s⁻¹), \mathbf{I} an identity matrix, P the pressure
278 (Pa), μ the dynamic viscosity (Pa s) and μ_T the eddy viscosity (Pa s), which is calculated by the k- ω model
279 for a turbulent flow and equals zero in the case of a laminar flow. Second order discretisation was set
280 by default in all equations. In the plant compartment the air velocity and pressure fields were modelled
281 using the Darcy–Forchheimer equation (Eq. 2), which considers single-phase flow in a porous medium.
282 To couple the physics, the pressures at the boundaries between the air ducts (turbulent flow) and the
283 plant compartment (Darcy-Forchheimer) were equalised. This is a convenient and computationally
284 profitable approximation for single phase fluid flow in a porous medium (Bejan, 2013). For the duct walls
285 a no-slip condition is used which means that the fluid velocity adjacent to the wall is zero. Wall roughness
286 was chosen to be generic with standard roughness height and roughness parameter. The inline duct fan
287 was modelled as an ‘interior fan’. This boundary condition considers the static pressure curve data (the
288 fan curve) of the actual duct fan. For the fan used in this study, five pressure/flow rate curves were
289 determined corresponding to the five different power supply settings. Due to the different densities in
290 plant material during the experiment, a trend line could be fitted through the data points on the pressure
291 loss versus flow rate curve. This means that the more plant material that is present, the higher the
292 pressure loss between the front and rear of the vegetation sample and the lower the air flowrate. Curves
293 are at various positions for each chosen wind speed; at higher wind speeds there is a shift towards higher
294 values. However, this relationship retains the same slope. The advantage of using the fan boundary
295 condition is, that the model automatically finds the working point of the fan, corresponding to the

296 pressure losses imposed by the vegetation and all duct walls. Consequently, the velocity measured by
 297 the hot wire anemometer in the experiment can be used to validate the accuracy of the model, by
 298 plotting experimental and modelled values against each other. The same applies to the pressure loss
 299 measured by the pressure module and the modelled pressure loss. Permeability, porosity and
 300 Forchheimer-drag datasets were provided by the wind tunnel experiments. Apart from Darcy-
 301 Forchheimer, a simple Darcy model was tested as well. Model results for all experiments were obtained
 302 using a parametric sweep, which is the easiest way to find the solution for a sequence of input
 303 parameters. The mesh of the plant compartment was built at a larger element size, as a finer mesh
 304 would take too much computational time and computer capacity. The pressure losses, obtained by the
 305 model, were then compared with the experimentally measured values to test the method. As a
 306 validation, the modelled velocity was compared to the value measured by the anemometer, at the
 307 corresponding measuring point.

308



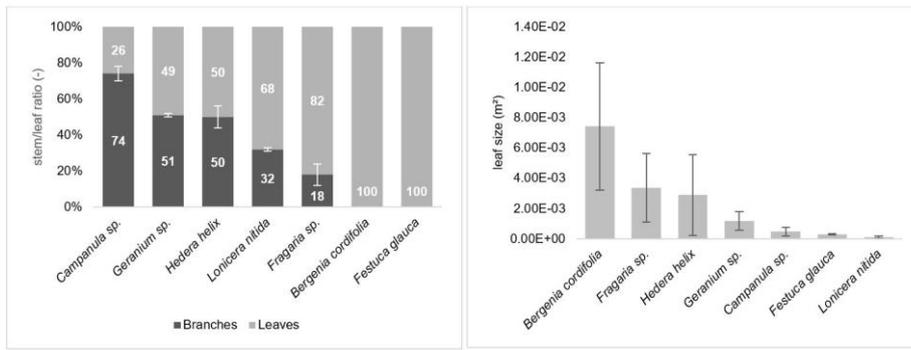
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310 Fig. 3: Geometry of the wind tunnel (left) and the computational mesh used in the model (right)

311 **3 Results**

312 **3.1 Species morphology**

313 A series of species morphological parameters was determined in order to explain potential variance in
 314 pressure and permeability data. Fig. 4 on the left shows the mean LR per species. For *Campanula* ($26\% \pm 0.04$)
 315 the stem volume overrules the leaf volume. For *Geranium* ($49\% \pm 0.01$) and *Hedera* ($50\% \pm 0.06$),
 316 the ratio is 50/50. *Lonicera* ($68\% \pm 0.01$) and *Fragaria* ($82\% \pm 0.06$) possess more leaf than stem material
 317 in terms of volume. *Bergenia* (rozet plant) and *Festuca* (grass) both consist solely of leaves. LS (Fig. 4
 318 right panel) shows *Bergenia* ($741.2 \text{ mm}^2 \pm 421.3$) to have the largest leaves, followed by *Fragaria* (335.5
 319 $\text{mm}^2 \pm 236.6$) and *Hedera* ($288.4 \text{ mm}^2 \pm 265.4$) which are close together. *Geranium* has small leaves
 320 ($118.2 \text{ mm}^2 \pm 62.9$) followed by *Campanula* ($47.6 \text{ mm}^2 \pm 27.8$). *Festuca* is a very fine grass ($29.6 \text{ mm}^2 \pm$
 321 3.0) and *Lonicera* has the smallest leaves ($11.1 \text{ mm}^2 \pm 6.6$). SLA varied from $20.60 \text{ m}^2 \text{ kg}^{-1} \pm 1.99$ for
 322 *Fragaria*, down to $5.60 \text{ m}^2 \text{ kg}^{-1} \pm 1.31$ for *Hedera* (Fig. 5). Furthermore, FLS varied from $8.87 \times 10^{-1} \pm 0.08$
 323 for *Fragaria* down to $3.22 \times 10^{-3} \pm 0.001$ for *Festuca* (Fig. 5). LDI of the species varied from 4.16 ± 0.3 for
 324 *Lonicera*, which has the roundest leaves to 33.8 ± 2.5 for *Festuca*, which is a grass. LSS varied widely
 325 between species, which made comparison complicated. Hence, surface structure was categorised into
 326 robust classes (Table 3). Based on an ANOVA-analysis no statistically significant correlations could be
 327 found between the species' morphological parameters.

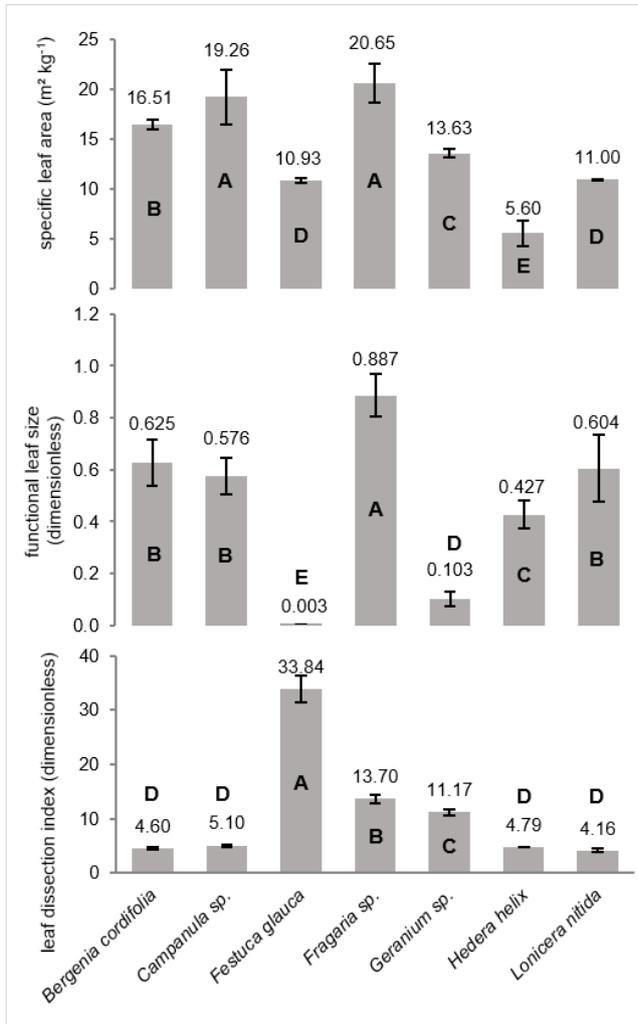


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Fig. 4: Volumetric leaf ratio (LR) of the considered plant species, classified from lowest to highest

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Fig. 5 (from top to bottom): Mean Specific Leaf Area (SLA), Functional Leaf Size (FLS) and Leaf Dissection Index (LDI) with standard deviations for all species, classified alphabetically according to species name. Different letters indicate significant differences.

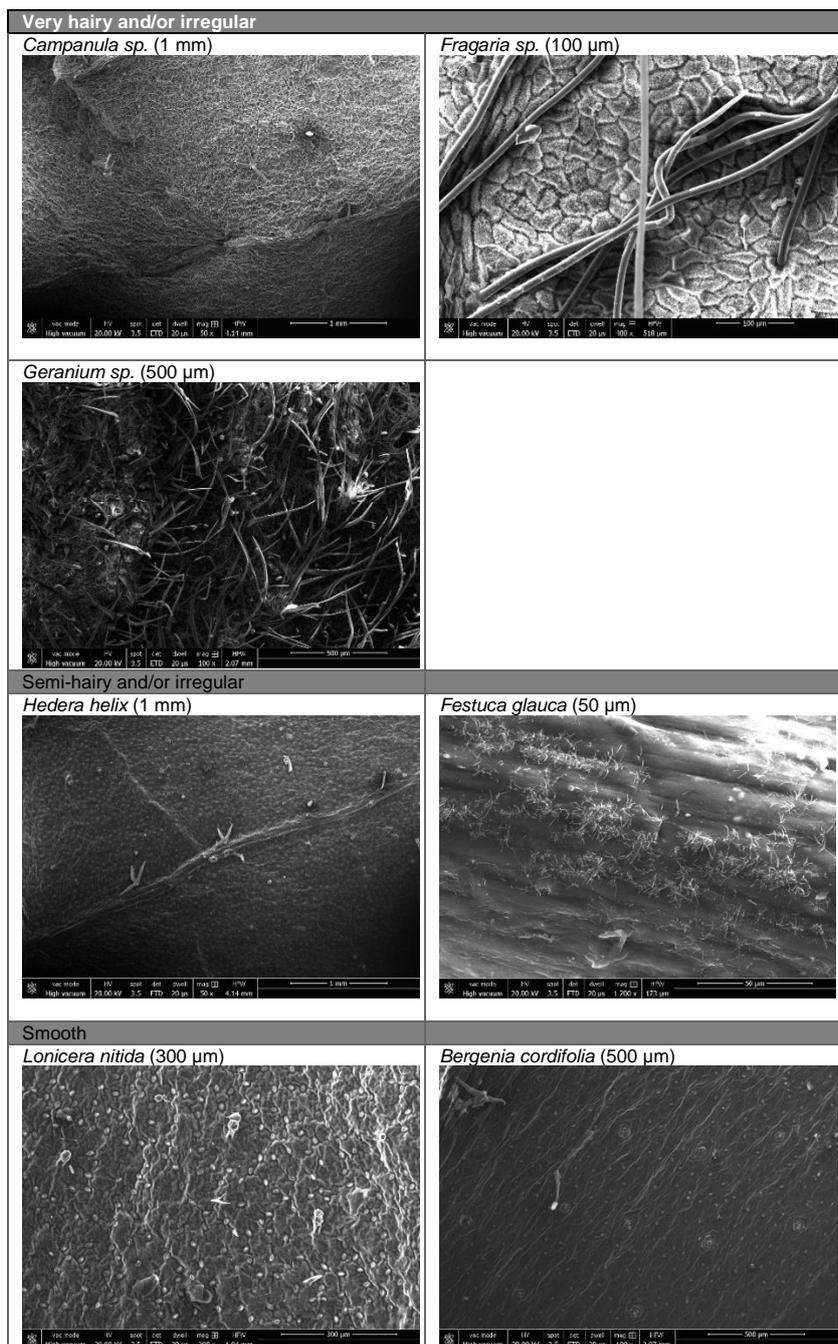
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337 Table 3: Leaf surface structure (LSS) divided into 3 categories: very hairy and/or irregular, semi-hairy and/or irregular, and smooth. Images were taken at 10 μm to 1 mm, depending on the size of the
 338 and/or irregular, and smooth. Images were taken at 10 μm to 1 mm, depending on the size of the
 339 structures showed.



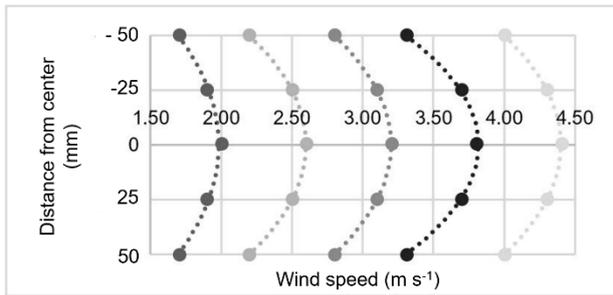
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342 3.2 Aerodynamic characterization using wind tunnel experiments

343 Figure 6 shows the shape of the air velocity profiles, derived from measurements at the five power
 344 supply settings and three different positions in the tunnel duct (centre, 0.025 and 0.05 m from the
 345 centre). During the pressure loss experiments, air velocity was only measured in the centre of the duct
 346 and the average velocity was calculated by integration over the cross section surface of the duct,
 347 according to the profiles in Fig. 6. For further determination of the aerodynamic parameters, the average

348 velocity was used in the equations. Reynolds numbers were calculated for each run using the kinematic
349 viscosity of air at 20° C which is $1.51 \times 10^{-5} \text{ m}^2 \text{ s}^{-1}$. Reynolds numbers varied from 2.33×10^4 to $1.50 \times$
350 10^5 . These values indicate turbulent air flow, which justifies the use of Darcy-Forchheimer for further
351 calculations.

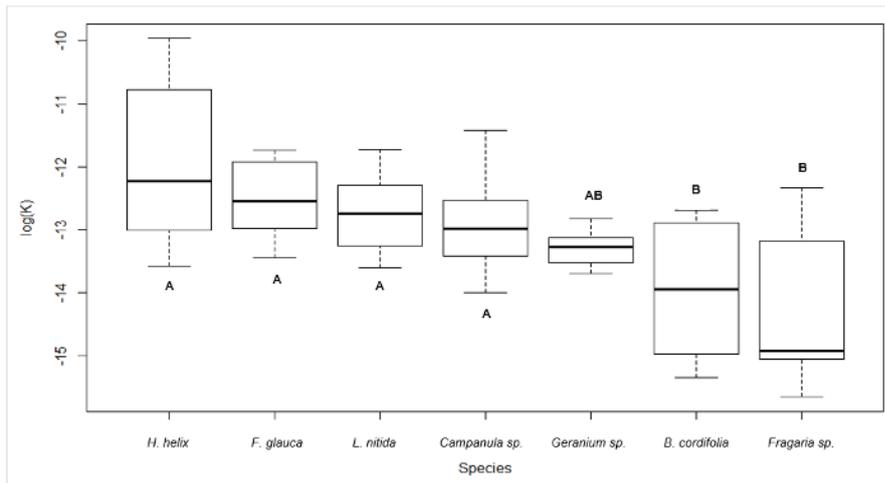


352
353 Fig. 6: Air velocity profiles at the five used power supply settings and three different positions in the tunnel duct
354 (centre, 0.025 and 0.05 m from the centre). The dotted lines indicate polynomials plotted through the data points.

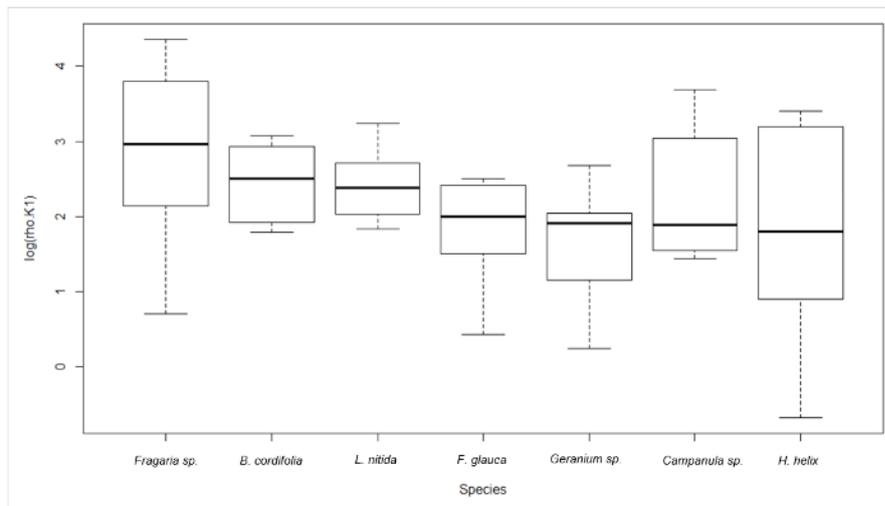
355
356 From the pressure loss measurements at a range of average air velocities, the permeability (K) and
357 Forchheimer-drag (ρ/K_1) were obtained by regression using Eq. (2). Values of permeability and
358 Forchheimer-drag varied from $1.59 \times 10^{-7} \text{ m}^2$ to $4.71 \times 10^{-5} \text{ m}^2$ and $5.09 \times 10^{-1} \text{ kg m}^{-4}$ to $7.80 \times 10^{-1} \text{ kg m}^{-$
359 4 , respectively. Data distribution for both permeability and Forchheimer-drag is heavily skewed.
360 Therefore, to investigate the difference between species, a natural log transformation was performed.
361 Normality could be confirmed by a Shapiro-Wilk normality test. Figure 7 shows differences in
362 permeability (top) and Forchheimer-drag (bottom) between species. An ANOVA proved that log (K) is
363 different between species ($p = 1.63 \times 10^{-5}$). The TukeyHSD test indicated where the differences are
364 between species (A and B). On the other hand, log (ρ/K_1) did not differ between the species ($p = 0.09$).

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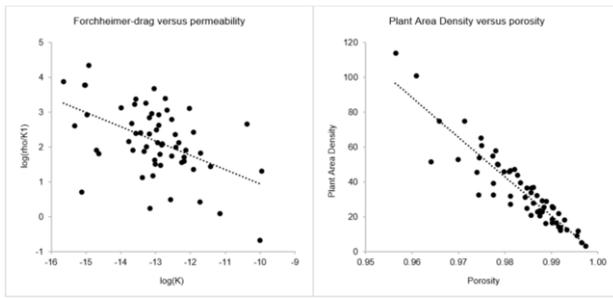


373 Fig. 7: Dispersion of permeability $\log(K)$ in m^2 (top) and Forchheimer-drag $\log(p/K_1)$ in $kg\ m^{-4}$ (bottom) for all
374 species, classified from high to low. For $\log(K)$ differences between species are indicated with the letters A and B.

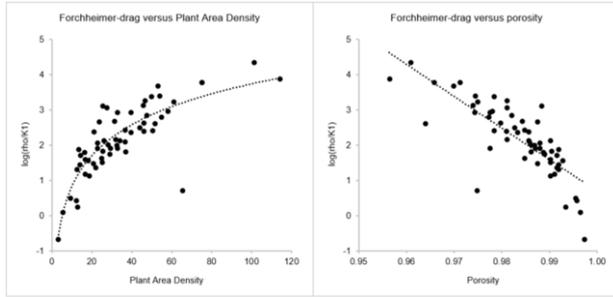
375

376 Furthermore, Kendall rank correlation tests were performed to test the correlations between K and ρ/K_1
377 on one hand and PAD and porosity on the other hand and between PAD and porosity reciprocally (Fig.
378 8). Tau and p-values are given in Table 4. Permeability is positively correlated to porosity and negatively
379 correlated to PAD, while for the Forchheimer-drag, the opposite is observed (as seen on the middle and
380 bottom row, respectively). Furthermore, porosity can be seen as the best indicator (highest values of
381 tau) for both permeability and Forchheimer-drag. In general, porosity was close to 1 for all experiments
382 (0.956 to 0.997). Linear regression for PAD as a function of porosity for individual species showed R^2
383 values between 0.99 (*Campanula*) and 0.96 (*Fragaria*). For aggregation of all samples, also a strong
384 correlation was found ($p < 2.2 \times 10^{-16}$, $R^2 = 0.85$, $\tau = -0.77$), with more scatter at higher PAD and lower
385 porosity values. Consequentially, one linear relationship could be derived: $\varphi = 0.997 - 3.78 \times 10^{-4} \times PAD$.
386 It appears that the Forchheimer-drag becomes more important with decreasing permeability, as
387 evidenced by the top left figure (although the data is scattered).

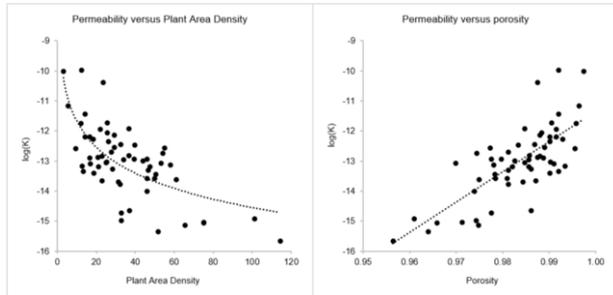
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391 Fig. 8: Kendall rank correlations between permeability (m^2), Forchheimer-drag ($kg\ m^{-4}$), plant area density ($m^2\ m^{-3}$)
 392 and porosity (dimensionless). Each points represents one wind tunnel run for a specific species under different
 393 wind speeds. Trendlines are also given (dotted line). The equations of the trend lines, from left to right and top to
 394 bottom, are $y = -0.4107x - 3.167$, $y = -2206 \ln(x) - 1.6602$, $y = -1.248 \ln(x) - 8.803$, $y = 101.3x - 112.63$, $y = 1.2676$
 395 $\ln(x) - 2.0972$ and $y = -88.04 \ln(x) + 0.7038$, respectively.

396

397 Table 4: Kendall rank correlation coefficients (τ) and p-values or correlations between permeability (m^2),
 398 Forchheimer-drag ($kg.m^{-4}$), plant area density ($m^2.m^{-3}$) and porosity (dimensionless).

	Permeability (m^2)	Forchheimer-drag ($kg\ m^{-4}$)	PAD ($m^2\ m^{-3}$)
Forchheimer-drag ($kg\ m^{-4}$)	$\tau = -0.42$ p-value = 1.58E-06		
PAD ($m^2.m^{-3}$)	$\tau = -0.43$ p-value = 1.58E-06	$\tau = 0.65$ p-value = 3.88E-13	
Porosity	$\tau = 0.52$ p-value = 6.79E-09	$\tau = -0.69$ p-value = 1.58E-14	$\tau = -0.77$ p-value < 2.2E-16

399

400

401 3.3 Correlation between aerodynamic and morphologic parameters

402 For each distinct species, the morphological parameters SLA, LDI, FLS, LR, LS and LSS were analysed as a
 403 function of permeability and Forchheimer-drag. Because morphological parameters have only one value
 404 per species and permeability and Forchheimer-drag have a series of values per species, we need also
 405 one value for permeability and Forchheimer-drag to compare them with the morphological parameters.
 406 P-values obtained for the linear models are shown in Table 5. Since data was skewed, both mean and

407 median values of permeability and Forchheimer-drag were used. Note that LSS should be considered as
 408 a factor.

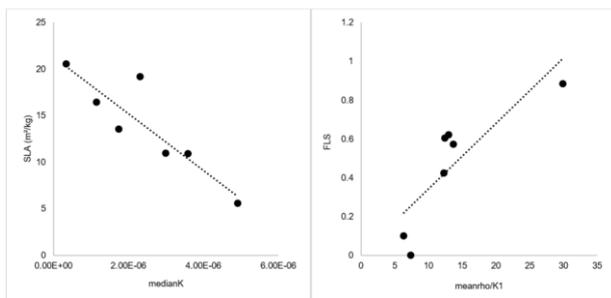
409 Table 5: Morphological parameters Specific Leaf Area (SLA), Lead Dissection Index (LDI), Functional Leaf
 410 Size (FLS), Leaf Ratio (LR), Leaf Size (LS) and Leaf Surface Structure (LSS) as a function of mean and
 411 median permeability (K) and Forchheimer-drag (ρ/K_1) for all species. The table shows p-values and R²-
 412 values of the correlations as result of an ANOVA. Stars indicate significant p-values ($p \leq 0.05$).

	Mean K	Mean ρ/K_1	Median K	Median ρ/K_1
SLA	$p = 0.04^*$ $R^2 = 0.53$	$p = 0.17$ $R^2 = 0.21$	$p = 0.007^*$ $R^2 = 0.79$	$p = 0.15$ $R^2 = 0.23$
LDI	$p = 0.75$ $R^2 = -0.17$	$p = 0.73$ $R^2 = -0.16$	$p = 0.82$ $R^2 = -0.19$	$p = 0.93$ $R^2 = -0.20$
FLS	$p = 0.71$ $R^2 = -0.16$	$p = 0.018^*$ $R^2 = 0.71$	$p = 0.27$ $R^2 = 0.083$	$p = 0.05^*$ $R^2 = 0.47$
LR	$p = 0.50$ $R^2 = -0.088$	$p = 0.78$ $R^2 = -0.18$	$p = 0.62$ $R^2 = -0.14$	$p = 0.28$ $R^2 = 0.07$
LS	$p = 0.85$ $R^2 = -0.19$	$p = 0.52$ $R^2 = -0.09$	$p = 0.36$ $R^2 = -0.0002$	$p = 0.34$ $R^2 = 0.017$
LSS	$p = 0.19$ $R^2 = 0.34$	$p = 0.71$ $R^2 = -0.26$	$p = 0.10$ $R^2 = 0.51$	$p = 0.61$ $R^2 = -0.18$

413

414

415 The variance of mean and especially median values of K could be explained by SLA ($R^2 = 0.79$) and the
 416 variance of both mean and median values of ρ/K_1 could be predicted by FLS ($R^2 = 0.71$). These
 417 correlations are plotted in Fig. 9. No other statistically significant correlations could be found.



418

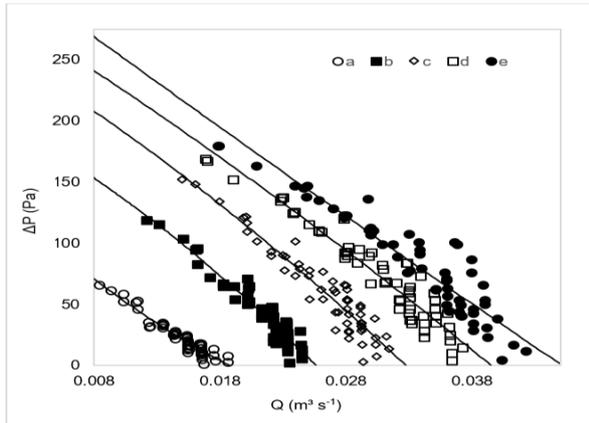
419 Fig. 9: Left: The correlation between SLA ($m^2 kg^{-1}$) and the median values for permeability K (m^2). Equation of trend
 420 line: $SLA (m^2 kg^{-1}) = -(3 \times 10^{-6}) median + 21.323$. Right: The correlation between FLS (dimensionless) and mean values
 421 for Forchheimer-drag ($kg m^{-4}$). Equation of trend line: $FLS = 0.0338mean + 0.004$. Each point represents a species.
 422 Trend lines are shown as dotted lines.

423

424 3.4 Numerical modelling

425 Before running the CFD model, the fan curves of the duct fan, corresponding to the five power supply
 426 settings, needed to be derived. These were the boundary conditions considered in the model. Fan curves
 427 corresponding to the five power settings were derived by plotting ΔP values against the flow rate (Q)
 428 for the total number of wind tunnel runs, including all species and densities (Fig. 10). Flow rates varied from
 429 the lowest to the highest wind speed the fan could provide.

430



431

Fig. 10: Fan curve derived from the experiments with plant material in the wind tunnel, for lowest (a) to highest wind speed (e) setting. ΔP = dynamic pressure loss between before and behind the plant compartment, Q = flow rate. The full lines are the linear approximations of the fan curves, obtained

432

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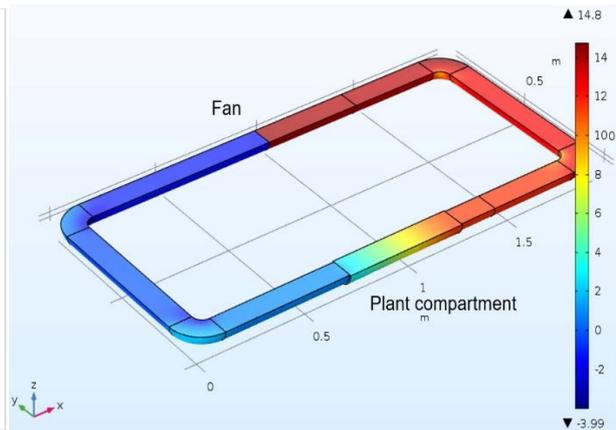
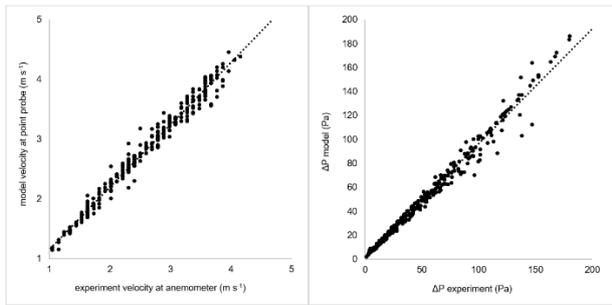


Figure 11: Typical pressure distribution obtained by the CFD model for the specific case of *Geranium* sp. ($PAD = 12.64 \text{ m}^2 \text{ m}^{-3}$, $\phi = 0.9933$, $K = 1.9039 \times 10^{-6} \text{ m}^2$, $\rho/K_1 = 1.2835 \text{ kg m}^{-4}$, $v = 1.36 \text{ m s}^{-1}$)

434 A typical result obtained by the CFD model is shown in Fig. 11. The figure shows the distribution of
435 pressure in the system for the k- ω coupled turbulence model, coupled with the Darcy-Forchheimer
436 equation for air flow through the porous vegetation in the plant compartment. The figure clearly shows
437 the sudden pressure drop over the fan, corresponding to the fan curve, and the gradual pressure drop
438 over the plant compartment. Apart from the Darcy-Forchheimer model, a simple Darcy model was
439 applied with values of K retrieved from standard Darcy's Law (Eq. 1). However, this model showed a
440 consistent underestimation of the pressure drop compared to the experiments, because no inertial term
441 was accounted for and porosities were high. The k- ω coupled turbulence model, which included the
442 Forchheimer-drag for air flow through the porous vegetation section, performed best when pressure
443 drop outcomes were correlated to experimental pressure drop results (R^2 between 0.96 and 0.98).

444 Model validation was performed by comparing the velocity measured with the anemometer and the
445 velocity calculated by the model, whereby the model velocity was evaluated at the exact anemometer
446 location (Fig. 12 left, $R^2 = 0.98$, $NMSE = 0.016$). Since the fan curve was used as a boundary condition, no
447 velocity boundary was considered, i.e. the experimentally measured velocity was not used as a model
448 input. Further validation of the model is evidenced by the comparison of measured and modelled ΔP
449 values (Fig. 12 right, $R^2 = 98$, $NMSE = 0.014$). Data points tend to be more scattered at higher pressure
450 differences (and thus wind velocities). A simple t-test showed no significant differences between model
451 and experimental pressure drops for every separate species.



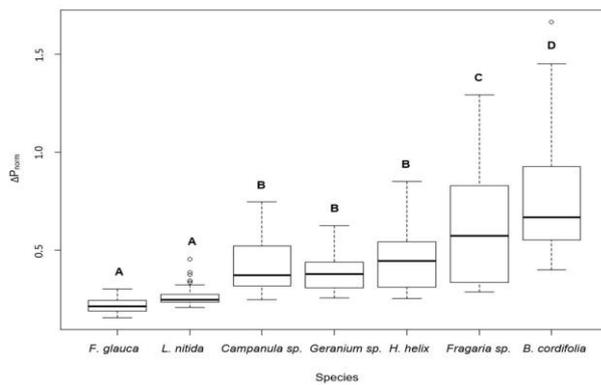
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453 Fig. 12: Left panel: correlation between experimental and modelled wind velocities at anemometer position. Each
 454 dot represents an experimental velocity respective to its model output. Right panel: modelled versus experimental
 455 pressure drop for all species considered. Each dot represents an experimental wind tunnel run under a certain PAD
 456 of a certain species with a certain velocity, respective to its modelled counterpart. The dotted lines represent the
 457 trend line of the correlations between parameters. Respective trend line equations: $y = 1.0439x + 0.1177$ and $y =$
 458 $0.9611x + 0.6972$.

459

460 For all species, differences between modelled and experimental normalised pressure loss were not
 461 significant. Consequentially, only modelled values are given in Fig. 13, which shows the ranking of all 7
 462 species from low to high normalised pressure loss according to Eq. 5. A clear ranking of species can be
 463 seen, even though a normalisation for air and plant density, wind velocity and distance (length of the
 464 plant compartment) was performed. *Festuca* shows the lowest value, while *Bergenia* is on the other side
 465 of the figure, showing the largest normalized pressure loss. Moreover, an ANOVA showed a significant
 466 difference between species ($p < 2.2 \times 10^{-16}$). For every species a mean normalized pressure loss (ΔP_{norm})
 467 was calculated for both experimental and modelled values for every species in order to test against
 468 morphological parameters. An ANOVA showed that the variance in normalised pressure loss could not
 469 be explained by either one of the considered morphological parameters.

470



471

472 Fig. 13: Boxplots of simulated normalised pressure drop for all considered species

473

474

475

476 4 Discussion

477 4.1 Wind tunnel experiments

478 Wind tunnel experiments are a powerful way to study the aerodynamics of air flow through vegetation
479 (e.g. K. Paul Beckett, Freer-Smith, & Taylor, 2000; Endalew et al., 2009, Gromke 2011). Relatively small
480 and thus, cheap wind tunnels are sufficient and experiments are not too time and labour intensive. For
481 these experiments plants were brought into a removable compartment of a wind tunnel under a range
482 of packing densities. Compared to Lin & Khlystov, (2012a), who tested packing densities (plant volume
483 to wind tunnel compartment volume) ranging from 0.017 to 0.040 for Pine and 0.037 to 0.055 for Juniper
484 (both coniferous tree species), our packing density range was much wider, from 0.0018 to 0.043 (both
485 for *Festuca*) with a median value of 0.0126 (*Lonicera*).

486 Regarding porosity, our values (0.9564 to 0.9973) are in agreement with the study of Buccolieri et al.
487 (2009) who used synthetic wadding material in wire cages ($\varphi = 0.96 - 0.975$), which is assumed typical
488 for tree crown porosities, and in agreement with Gromke et al. (2016), who used foam materials with
489 pore volume fractions of 96.1% and 94.5%. In contrast with these studies, our research included a whole
490 range of different porosities between and within species instead of using only one or two species with
491 fixed porosities.

492 In general, most wind tunnel studies considering vegetation aerodynamics are conducted with
493 vegetation simulators (model plants) (Buccolieri et al., 2009; Endalew et al., 2009; Gromke 2011 and
494 references therein; Gromke et al., 2016; Wuyts et al., 2008) instead of real vegetation. The main
495 disadvantage of using simulators is that they are artificial and rigid and thus, their structure is not
496 affected by wind velocities, whereas the flexible structure of plants enables them to bend and even
497 reduce drag at high wind speeds by changing their shapes. Wind tunnel experiments concerning real
498 vegetation have mostly been used in particulate pollution capture studies (Beckett et al., 2000; Huang,
499 Lin, Khlystov, & Katul, 2013; M.-Y. Lin & Khlystov, 2012a) but rarely in aerodynamic research, making
500 this study innovative in terms aerodynamic research of air flow through vegetation.

501

502 4.2 Aerodynamic and morphological parameters and their correlation

503 Our results showed significant differences between species for permeability (Fig. 7 top) ($p = 1.63 \times 10^{-5}$)
504 ⁵), but not for the Forchheimer-drag (Fig. 7 bottom) ($p = 0.09$). This suggests that permeability is sensitive
505 to specific differences between species, while Forchheimer-drag apparently works in a similar way
506 regardless of species. Because permeability is dependent on porosity, pore size and detailed plant
507 geometry (Nield, 2000), variations in means of permeability are expected to be caused by morphological
508 and architectural differences between species. The spread of permeability values per species (see Fig.
509 8) can be explained by stacking convenience in the plant compartment of the wind tunnel. Species like
510 *Hedera* and *Fragaria* could be stacked in a wider series of PAD's (thus values for PAD lying further apart,
511 see Figure 2.3). While for species like *Geranium* and *Festuca* the variation in PAD was more limited as
512 these species were structurally lighter and more difficult to spread homogeneously. In accordance with
513 Sase et al. (2012), larger values of permeability were found with decreasing plant/leaf area density.
514 However, the plants used in the experiment of Sase et al. (2012) were much further from each other,
515 resulting in lower densities, so these values would be reported on the left side of the graph in Figure 8.

516 Furthermore, because of the strong correlation between PAD and porosity ($p < 2.2 \times 10^{-16}$) (Fig. 9),
517 porosity has a strong positive correlation with permeability ($p = 0.79 \times 10^{-9}$). PAD and porosity also seem

518 to have a strong influence on Forchheimer-drag with p-values of 3.88×10^{-13} and 1.58×10^{-14} ,
519 respectively. This is logical as larger amounts of plant material (higher PAD, lower porosity) cause more
520 drag and resistance to free air flow, which causes more turbulence and thus higher values for
521 Forchheimer-drag.

522 Our experiments suggest that SLA can be an indicator of permeability (Table 5 and Fig. 9 left). This means
523 that it could be possible to at least estimate the permeability of an herbaceous plant species (without a
524 woody stem) by only assessing SLA. This makes sense, because species with large, thin and broad leaves
525 (large SLA) cause a stronger decrease in air flow in contrast to compact, rigid leaves (small SLA), where
526 it's easier for air flow to pass around. Even though thin leaves would bend easier, this effect seems to
527 be subordinate. Moreover, Figure 9 (right) shows significant correlations between Forchheimer-drag
528 and FLS. This could be explained by the leaf boundary layer, which becomes larger with increasing FLS.
529 Thicker boundary layers signify thicker stagnant layers of air and thus an increased amount of air inertia.
530 Please note that the correlations between SLA and permeability, and FLS and Forchheimer-drag might
531 be subject to change. Although care was given to select representative species with a wide variety in
532 leaf morphology, it might be that adding more species, be they shrubs or trees, would reinforce the
533 given correlations. Therefore, still further research is needed to study these interactions.

534

535 **4.3 Model vs. experiment**

536 In recent years, CFD has become an attractive tool to predict the effects of green infrastructure on the
537 local microclimate (heat island effect, PM deposition) and the demand for CFD studies (and appropriate
538 parameters) for analysis of different urban planning scenarios is rising. However, CFD models can be
539 computationally demanding, which puts limitations on this method. As a result, most CFD models
540 represent vegetation in a rather rudimentary and theoretical way, which can simplify complex processes
541 too much; e.g. for the assessment of PM removal capacity of vegetation, where vegetation is modelled
542 as a simple body with theoretical leaf area densities (LAD), leading to an underestimation of PM removal
543 capacity (Hofman et al., 2016). CFD modelling is widespread and many aerodynamic studies exist on
544 street level (Bartzanas et al., 2002; Hong et al., 2012), but fewer work is done on vegetation level, i.e.
545 within vegetation stands. This does not come as a surprise because detailed assessment of wind flow
546 through vegetation is challenging. Compared to a solid body, around which the wind flow is predictable,
547 plants are porous and flexible. Also, individual plants of the same species can have different geometries
548 and most plants experience a shift in structure throughout the year, with as a pronounced example the
549 seasonal leaf dynamics of deciduous species, and throughout their lifetime. The complexity of air flow
550 through vegetation makes a simplification inevitable: in the best case, a minimum of parameters are
551 used (here: permeability, Forchheimer-drag and porosity) in the model approach and a compromise
552 between accuracy and computational demand needs to be found.

553 Figure 12b shows a strong correlation ($R^2 = 0.98$) of modelled and experimental pressure outcomes with
554 no significant differences between modelled and experimental results for all species considered. The
555 correlation becomes less obvious at higher pressure drop values (so also at higher wind velocities and
556 plant densities). This can be explained by movement or compression of the vegetation in the wind tunnel
557 at higher wind speeds, where in the model the vegetation is considered to be a static, non-moving body.
558 Higher wind speeds cause changes in permeability due to plants being pliable. More specifically, on one
559 hand, strong wind flow tends to bend foliage and make it more streamlined, thus increasing permeability
560 but on the other, permeability lowers at the point where vegetation in the vegetation compartment is

561 blown together due to an increase in local density or porosity. Tiwary et al. (2005) suggest that grass-
562 like and fine-leaved species appear to be more likely to be bent or compressed, while species possessing
563 more rigid leaves and twigs are only affected by strong winds. Thus, Fig. 12b suggests that the Darcy-
564 Forchheimer model is suitable to describe wind flow through herbaceous vegetation, with higher
565 reliability under lower pressure drops and wind velocities. This model is a solid base for further
566 experiments considering interactions between plants and wind-driven processes like PM deposition and
567 heat exchange. However, it is suggested that future validation of this model should be performed with
568 more plant functional types (e.g. shrubs and trees) with special features like very large leaves or needles,
569 as this will contribute to the model's robustness. Furthermore, higher wind speeds should be considered
570 to detect if the model performs also well under these conditions.

571 Pressure losses were normalised in order to assess differences between species. Even after
572 normalisation for wind velocity and PAD, a measure of plant packing density, a clear difference between
573 species was found (Fig. 13). The figure shows a ranking from small to large values for ΔP_{norm} . This is
574 expected to be caused by species-specific morphological characteristics. However, differences in ΔP_{norm}
575 between species could not be explained by the morphological parameters. Further research, with more
576 species and different species traits in terms of leaf shape and size, and perhaps rigidity, should be
577 conducted. Also no differences in mean pressure drops were found within species between modelled
578 and experimental outcomes. This means that the model is equally able to predict pressure drop over a
579 vegetation stand for the wide variety of species used in this research. This is promising, as living wall
580 systems often contain a combination of different species, which points towards a broad applicability.

581 To describe vegetation and its aerodynamic characteristics, different measures are used in literature,
582 which makes comparison difficult (Janhäll, 2015). Because of the setup used in our experiment it is
583 possible that the model provided values are overestimating pressure drop in a real-life situation. This
584 agrees with the comments of Janhäll (2015), who stated that pressure drop found in reality will be less
585 than that found in a wind tunnel. This is because in a wind tunnel air is restricted to flowing through the
586 vegetation whereas in reality air will partly pass around the structure due to resistance. Likewise, Tiwary
587 et al. (2005) state that high porosity barriers are more penetrable by wind while vegetation with lower
588 porosities forces air to pass around them.

589 **5 Conclusions**

590 This paper studied the interactions of vegetation and air flow and the influence of plant morphological
591 characteristics. For the first time, several herbaceous plant species were aerodynamically characterized.
592 A Darcy-Forchheimer model to describe air flow through the vegetation was proposed. The
593 permeability, Forchheimer-drag and porosity of the vegetation could be used as an input for CFD-
594 modelling. It was concluded that:

- 595 • The use of real plants instead of artificial materials is innovative in terms aerodynamic research
596 of air flow through vegetation.
- 597 • Strong correlations between modelled and experimental data show us that the Darcy-
598 Forchheimer model is a simple and robust way to describe air flow through vegetation, with
599 higher reliability under lower pressure drops.
- 600 • Our results illustrate that permeability is sensitive to variation between species, while
601 Forchheimer-drag is not.

- 602 • Because of the correlation between mean and median permeability values and SLA, it can be
603 assumed that the differences in permeability between species can be partly explained by their
604 SLA.
- 605 • Even though Forchheimer-drag showed no significant difference between species, mean and
606 median values for Forchheimer-drag seem to be correlated with FLS. Therefore, we suggest FLS
607 could be an indicator for Forchheimer-drag.
- 608 • A significant variance in normalized pressure drop exists between species, which could not be
609 explained by the investigated morphological parameters. This suggest that more research needs
610 to be conducted by making use of more species and other plant morphological characteristics.

611 It can be stated that the proposed method is useful for future modelling of vegetation and its interaction
612 with air flow, more specifically helping investigate the process of PM deposition and thermal behaviour
613 in green walls.

614

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