

1 Are flow-vegetation interactions well represented by mimics?

2 A case study of mangrove pneumatophores

3 - Revised manuscript

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14 Keywords: vegetated flow; *Avicennia marina*; mangrove pneumatophores; vegetation

15 mimics; turbulence

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17 **Highlights (85 characters each)**

18 ~~— Real pneumatophores exhibit heterogeneity in shape, height and spatial distribution~~

19 ~~- Vegetation heterogeneity has minor effects on relative within-canopy velocities~~

20 ~~- Discrete canopy density transitions do intensify and confine the canopy shear layer~~

21 ~~- Uniform-height canopies ~~dowels generate stronger~~ enhance turbulence maxima and~~
22 ~~within-canopy turbulence than variable height canopies~~

23 ~~- Bed shear stress in pneumatophores two times smaller than in uniform-height dowels~~

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25 **Abstract**

26 Arrays of real mangrove pneumatophores and artificial dowel mimics were constructed in a
27 laboratory flume to compare differences in flow dynamics. Compared to the uniform-height
28 dowel canopy, the non-uniform height of the pneumatophores significantly reduced the
29 intensity of the canopy shear layer, and shifted the turbulence maxima from directly above
30 the dowels upwards by approximately the standard deviation of the pneumatophore heights.
31 Consequently, bed shear stresses were up to two times greater in the uniform-height dowel
32 canopy than in a pneumatophore canopy of similar density. At the same time, ratios of the
33 within-canopy velocity to the free-stream velocity above the canopies were not significantly
34 altered by heterogeneous heights, shapes and spatial distributions of the pneumatophores. Our
35 results emphasize that uniform dowels are poor proxies of real pneumatophore canopies and
36 may lead to underestimates of sediment-trapping efficiency.

37

38 **1 Introduction**

39 Aquatic vegetation imposes a significant drag force on the surrounding water, altering flow
40 velocity, turbulence and transport of suspended matter. Consequently, water motions through
41 vegetation canopies, such as mangrove root systems and saltmarsh grasses, have been widely
42 studied [reviewed by e.g. 1, 2-4]. While the vegetative drag tends to reduce horizontal current
43 velocities within the vegetation, continuity requires enhanced flow above (or around) the
44 vegetation, leading to increased velocity gradients over the canopy [2, 5, 6]. Resulting
45 velocity gradients at the interface of the canopy and the free-flowing water give rise to large-
46 scale turbulent motions, increasing the exchange of water and suspended matter between the
47 canopy and the water body [7-9].

48
49 Elevated turbulence levels over vegetation canopies are able to sustain enhanced
50 concentrations of suspended matter, however, these suspended materials may settle once
51 transported into the less dynamic zone within the vegetation. Consequently, enhanced
52 sediment deposition is often observed within (tidally) submerged canopies of mangrove roots,
53 saltmarsh grasses and seagrass [10-14]. The actual deposition is highly dependent on
54 vegetation properties, sediment supply and characteristics, as well as local hydrodynamic
55 conditions. For example, while Young & Harvey [15] observed a positive correlation
56 between accretion rates and root densities in mangroves, erosion has also been observed in
57 mangroves with relatively high root densities [16-18]. This contrast in accretionary trends in
58 mangroves is directly related to a balance between energy dissipation due to enhanced drag
59 forces and turbulence generation around these root systems [e.g. 19].

60
61 An abundance of data on drag and turbulence within aquatic vegetation exists, mostly from
62 experiments with uniform, rigid cylindrical dowels as vegetation mimics, positioned in

63 random, staggered or aligned arrangements [20-26]. In uniform emergent vegetation, flow
64 speeds within the vegetation monotonically decrease with the vegetation density (measured
65 by $a = Nd$, wherein a is the frontal vegetation cover per unit area [m^{-1}], N is the number of
66 stems per unit area [m^{-2}] and d the stem diameter [m]), the water depth-to-stem diameter ratio
67 (h/d) and the drag coefficient (C_D) induced by the surface roughness of the vegetation
68 elements [21, 27]. Both current velocities and turbulence intensity, expressed as the square-
69 root of the turbulent kinetic energy (k) divided by the flow velocity (u), are found to be
70 approximately constant over the water depth within the emergent canopy, with exception of
71 the near-bed region where peaks have been observed in both velocity and turbulence intensity
72 [6, 21, 27]. While turbulence intensity increases with the presence of sparse emergent
73 vegetation, there exists a critical vegetation density, beyond which the turbulence decreases
74 owing to the reduction in flow speeds occurring at higher densities [27].

75

76 Similar to the emergent scenario, in submerged canopies of uniform vegetation elements with
77 height h_c , within-canopy velocities tend to be constant with depth, provided the canopy
78 density is sufficiently large to create a drag force that exceeds the bed stress, which occurs at
79 $ah_c \gtrsim 0.23$ [2, 28]. Above such a dense submerged canopy, a characteristic logarithmic
80 velocity profile develops for submergence ratios $h/h_c > 1.25$ [21, 29], with an inflection point
81 of the velocity profile near the top of the canopy. Kelvin-Helmholtz instability in this canopy
82 shear layer can give rise to the production of coherent vortices, forming a dynamic mixing
83 layer and causing a local peak in the turbulence profile [20, 21, 30, 31]. The elevation of the
84 logarithmic velocity profile with respect to the top of the submerged canopy, and
85 consequently the penetration depth of the turbulence into the vegetation, depends on
86 vegetation density, the submergence ratio and the vegetation surface roughness [21, 22, 29].

87

88 In addition to the above studies that considered idealized artificial canopies, some studies
89 have also used real canopies with heterogeneous properties, both in the field and in the flume
90 [32-34]. However, studies in real canopies often suffer from an inability to comprehensively
91 resolve the hydrodynamics within the irregular (and dynamic) vegetation. This difficulty in
92 obtaining high-resolution data means that the effects of non-homogeneous vegetation
93 properties on the hydrodynamics remain largely unquantified. Liu et al. [35] showed that
94 turbulence production was different in a heterogeneous canopy by studying flow patterns
95 within a double-layer of idealized rigid vegetation. Throughout their experiments, the canopy
96 consisted of various densities of regularly-spaced short and tall acrylic dowels. For
97 sufficiently dense dowel configurations and sufficiently large relative height differences
98 between the short and tall dowels, the velocity and turbulence profiles in the composite
99 canopies resembled a superposition of the profiles for single-layer submerged canopies.
100 However, the shear across the top of each of the vegetation layers was reduced compared to
101 the shear at the top of a uniform-height dowel canopy of the same density, lowering the
102 maximum turbulence production.

103
104 These previous studies have highlighted the key principles of hydrodynamics in simplified
105 uniform-height and composite double-layer vegetation canopy mimics. Mangrove
106 pneumatophores are woody aerial roots growing upward from shallow lateral root systems of
107 e.g. *Avicennia* and *Sonneratia* species, reaching heights ranging from a few centimetres to
108 decimetres [36]. These pneumatophores show a considerable resemblance to the rigid
109 cylindrical dowels frequently used as vegetation mimics. However, pneumatophore heights
110 and diameters can vary substantially within the same field site [e.g. 37, 38]. The impacts of
111 such heterogeneity in vegetation characteristics has not been addressed in previous studies.

112

113 In a pilot study, we demonstrated that real pneumatophores may have a different effect on the
114 flow dynamics when compared to published studies using dowel mimics [39]. However, the
115 range of experimental conditions used in the published studies made it difficult to isolate and
116 quantify differences. Here, we use dowel canopies that have been designed to mimic the real
117 pneumatophores, allowing for a direct comparison of the flow dynamics in three resembling
118 vegetation canopies: real *Avicennia* pneumatophores (with three different densities), wooden
119 dowels with a variable canopy height equal to the pneumatophores, and wooden dowels of a
120 uniform height. Through these experiments, we explore: (i) the differences in flow dynamics
121 between submerged canopies of real, heterogeneous pneumatophores and idealized dowel
122 canopies of both uniform and variable height; and (ii) the hydrodynamic impacts of density
123 variations in submerged canopies of real pneumatophores, compared to previous results for
124 mimic canopies consisting of uniform dowels. These results provide an evaluation of
125 common practices in physical and numerical modelling that mimic natural vegetation by
126 uniform canopies and the observed differences will be used to identify potential avenues to
127 incorporate effects of the non-uniformity of natural vegetation in such models.

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128

129 2 Methods

130 2.1 Experimental set-up

131 Experiments were conducted in a 6.6 m long and 0.5 m wide recirculating acrylic flume with
132 a horizontal bottom [see 40]. The potential gradient across the flume was associated with a
133 water surface gradient only and a steady flow was generated by a variable-frequency impeller
134 that was mounted in the recirculating duct of the flume (Figure 1a). An insert of 0.3 m long
135 stacked 20 mm diameter PVC tubes was used to damp turbulence and rectify unidirectional
136 flow at the entrance of the flume.

Commented [EH2]: As requested by reviewer 2. Should I also include that hence 'the potential gradient was associated with water surface gradient only'? We don't do anything with these gradients later on, so I reckon we could skip this bit.

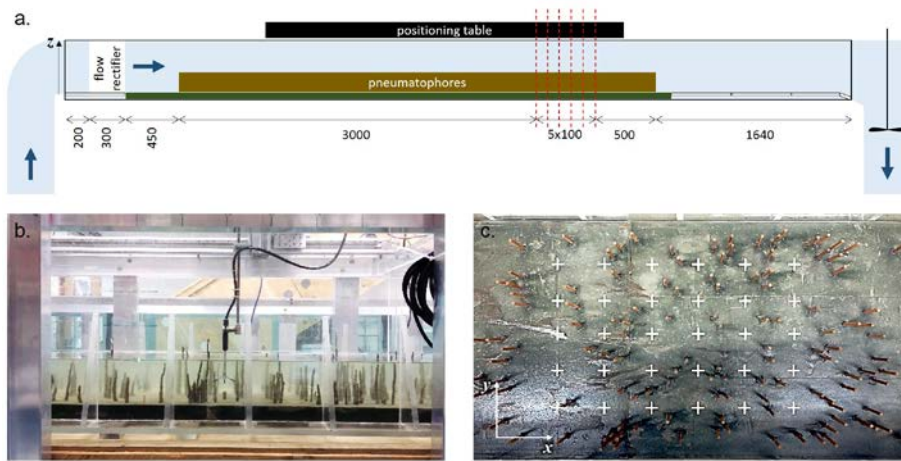
137

138 Vegetation canopies were mounted in large sheets of floral foam that were incorporated in the
139 false floor of the flume, providing a smooth bed over the full flume length. The canopies
140 were 4.0 m long, starting from 0.45 m downstream of the flow rectifier and extended across
141 the full width of the flume (Figure 1a). All data were collected along a series of six cross-
142 sections between 3.0 m and 3.5 m downstream of the leading edge of the canopies (Figure 1a)
143 and flow velocities prior to the implementation of the canopies were obtained from ten
144 spatially averaged observations of the free-stream velocity at either end of the data
145 acquisition area (see section 2.4 for data processing procedures).

146

147 Experiments were run with freshwater at three different water depths: 0.12 m, 0.21 m and
148 0.30 m, mimicking various tidal inundation stages of the canopies. At each water depth
149 experiments were run for two flow rates equating to free-stream flow velocities of 0.10 m/s
150 and 0.15 m/s prior to the implementation of the canopies. Impeller frequencies of the pump
151 were not modified after the introduction of the canopies, in order to maintain a constant flow
152 forcing throughout experiments with the same water depth and flow rate. These experimental
153 conditions are representative of the tidal dynamics in sheltered intertidal mangrove fringes
154 [e.g. 41, 42].

155



156
 157 *Figure 1 – Experimental setup: (a) Side view of the flume (dimensions in mm) including the vegetation canopy, (b) side view*
 158 *of the data collection area in the flume for the high-density pneumatophore experiment ($h = 0.12$ m), and (c) top-view of the*
 159 *data collection area for the experiment with uniform-height dowels. Locations for collecting vertical velocity profiles are*
 160 *marked with dashed lines in (a) and white '+'s in (c).*

161

162 **2.2 Vegetation canopies**

163 Experiments were undertaken for a plane bed and for five different canopies: three different
 164 densities of real pneumatophores and two high-density dowel canopies with variable and
 165 uniform heights (Table 1). Pneumatophores were harvested from the Waikareao Estuary in
 166 Tauranga Harbour, New Zealand. In order to obtain a natural spread of the pneumatophores
 167 in the flume, scaled photographs of 1×4 m² patches of sparse, average and dense
 168 pneumatophore covers were collected at the field site prior to harvesting (Figure 2a). These
 169 surveys revealed a patchiness of the pneumatophores that was reconstructed in the flume by
 170 applying a 0.10×0.10 m² raster overlay on both the photo reconstruction and the flume bed
 171 (Figure 2b,c).

172

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173 The reconstructed pneumatophore canopies had low (Pneum-LD), average (Pneum-AD) and
174 high (Pneum-HD) densities N of 71, 154 and 268 m^{-2} , respectively. The pneumatophores
175 generally featured a tapered shape with an average basal diameter of 8.1 mm, decreasing to
176 6.6 mm at mid-height and 4.0 mm at the tip of the pneumatophore, with a depth-mean
177 diameter d of 6.2 mm. The mean pneumatophore diameter remained constant in all three
178 reconstructions (Table 1). The resulting mean cross-sectional pneumatophore density $\varphi =$
179 $N\pi d^2/4$ ranged from 0.0023 to 0.0086 and the frontal canopy cover per unit area (a) ranged
180 from 0.44 m^{-1} to 1.67 m^{-1} for the low and high pneumatophore densities, respectively (Table
181 1). These densities were similar to values observed in lower latitude mangrove systems [16,
182 37, 38].

183
184 The average pneumatophore heights h_c differed slightly between densities and across the
185 different experiments the ratio of the water depth to the mean canopy height (h/h_c) ranged
186 from 1.9 to 4.7 (Table 2). Owing to the variability in heights of individual elements (standard
187 deviations of pneumatophore height were up to just over half of the mean canopy height;
188 Table 1), not every pneumatophore was fully submerged for the lowest water depth.

189 Throughout the experiments the pneumatophores behaved as rigid structures that did not
190 move or bend due to the hydraulic forcing, which is similar to their static behaviour in the
191 field when exposed to tidal currents only (personal observation).

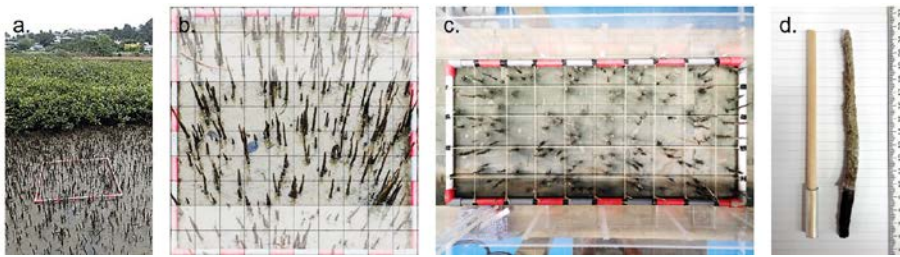
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192
193 The high density experiments were reproduced with rigid dowel mimics. Firstly, each
194 pneumatophore was replaced with a dowel of the same height (Figure 2d), resulting in a
195 variable-height dowel canopy (DowVar) with the same horizontal distribution as the
196 pneumatophores (Table 1). Secondly, all dowels were replaced with dowels of a uniform
197 height (DowUni) equal to the mean height of the high-density pneumatophore and variable-

Commented [EH5]: Rigid dowels

198 height dowel canopies. The cross-sectional density ϕ for both dowel canopies was slightly
 199 smaller than for the high-density pneumatophore canopy owing to the marginally smaller
 200 mean diameter of the dowels relative to the pneumatophores (Table 1).

201



202

203 Figure 2 – Canopy reconstruction in the flume: (a) high-density pneumatophore cover in the Waikareao Estuary in New
 204 Zealand, (b) a section of the rectified photo resulting from the field survey, and (c) reconstructed canopy in the flume and (d)
 205 pneumatophore with a dowel mimic of equal height (the black part of the pneumatophore and the tubed part of the dowel
 206 were fixed in the bed).

207

208 Table 1 – Vegetation properties of the experiments: heights h_c (mean \pm SD of all pneumatophores); diameters d are
 209 averages of the top, middle and bottom diameter of each pneumatophore (mean \pm SD of >100 pneumatophores);
 210 frontal canopy cover per unit area a ; and cross-sectional pneumatophore densities ϕ (as per definition computed with the
 211 root-mean-squared pneumatophore diameter).

	Pneumatophores			Dowels	
	low density Pneum-LD	average density Pneum-AD	high density Pneum-HD	variable height DowVar	uniform height DowUni
N [m ⁻²]	71	154	268	268	268
h_c [mm]	64 \pm 25	65 \pm 34	71 \pm 38	71 \pm 38	71 \pm 0
d [mm]	6.2 \pm 1.4	6.2 \pm 1.4	6.2 \pm 1.4	6.0 \pm 0	6.0 \pm 0
a [m ⁻¹]	0.44	0.96	1.67	1.61	1.61
ϕ [-]	0.0023	0.0049	0.0086	0.0076	0.0076

212

213 2.2.2.3 Data collection

214 For each of the canopies, experiments were run at the three different water depths and two
 215 flow rates (experimental conditions summarized in Table 2). A fully automated 3D

216 positioning table (Animatics SmartMotors) allowed for precise positioning and accurate
217 repetitions of hydrodynamic measurements (Figure 1a). Flow and turbulence data were
218 collected with a 10 MHz Nortek Vectrino Profiler (VP) that was mounted on the positioning
219 table (Figure 1b). The VP collected velocity profiles comprising 35 vertical cells of 1 mm
220 each, starting at 40 mm below the probe. Data were collected at a sampling rate of 50 Hz for
221 60 s at any position (tests conducted with a range of time windows showed that 60 s was long
222 enough to obtain consistent flow characteristics). The VP was moved along the vertical (z -
223 direction) to obtain semi-continuous vertical velocity profiles over the full water depths. To
224 account for the inhomogeneity of the velocity field both inside and above the canopy, these
225 vertical velocity profiles were collected at 30 positions within the data collection area (Figure
226 1a), covering 6 transects with a 100 mm interval in the streamwise x -direction and 5 positions
227 with a 75 mm spacing in the lateral y -direction along every transect (Figure 1c). The
228 observations were located no less than 0.10 m away from the flume walls to prevent shear
229 from the side walls impacting the measurements. For the plane bed experiments, the
230 streamwise spacing was increased to 500 mm and 10 vertical velocity profiles were collected
231 within the data collection area in order to account for minor streamwise and lateral velocity
232 variations due to imperfections in the flume.

233
234 *Table 2 – Experimental conditions: Canopy types as in Table 1; water depth h ; free-stream velocity U for the plane bed*
235 *scenario at the same water depth and flow rate (low or high) prior to inserting vegetation; canopy submergence h/h_c ; free-*
236 *stream velocities U_w at 0.05 m below the water surface (for experiments with 0.12 m water depth these observations*
237 *coincided with the top of the canopies, hence no free-stream velocity could be resolved); depth-averaged within-canopy*
238 *velocities U_c for $0 \leq z \leq h_c$ and the equivalent depth-averaged velocities U_0 for the plane bed experiments (presented*
239 *values for U_0 for $h_c = 71$ mm); ratios of within-canopy velocity to free-stream velocity U_c/U_w ; spatially averaged near-bed*

Commented [EH6]: Leaving out U_{inf} for the 12 cm water depths as requested by reviewer 2.

240 $(z=0.2h_c)$ Reynolds stresses $\langle u'w' \rangle$; the flow Reynolds number R ; the stem Reynolds number of the vegetation Re_s ; and the
 241 estimated drag coefficients C_D of the vegetation elements.

Canopy type	h [m]	U [m/s]	h/h_c [-]	U_c [m/s]	$U_c(U_0)$ [m/s]	U_c/U_c [m/s]	$\langle u'w' \rangle_{z=0.2h_c}$ [$\times 10^{-5} \text{ m}^2/\text{s}^2$]	Re [$\times 10^4$]	Re_s [-]	C_D [-]
PlaneBed	0.12	0.10	-	0.10	0.08	-	-1.70	1.09	-	-
PlaneBed	0.12	0.15	-	0.15	0.12	-	-3.54	1.65	-	-
PlaneBed	0.21	0.10	-	0.10	0.07	-	-1.33	1.88	-	-
PlaneBed	0.21	0.15	-	0.15	0.11	-	-3.15	2.85	-	-
PlaneBed	0.30	0.10	-	0.10	0.07	-	-1.23	2.65	-	-
PlaneBed	0.30	0.15	-	0.15	0.10	-	-2.52	3.97	-	-
Pneum-LD	0.12	0.10	1.9	0.09 n.a.	0.07	0.78 n.a.	-1.05	0.98	425	0.7
Pneum-LD	0.12	0.15	1.9	n.a.0.13	0.10	n.a.0.77	-2.23	1.47	632	0.7
Pneum-LD	0.21	0.10	3.3	0.11	0.06	0.52	-0.92	1.86	354	1.3
Pneum-LD	0.21	0.15	3.3	0.16	0.09	0.52	-1.93	2.76	527	1.2
Pneum-LD	0.30	0.10	4.7	0.11	0.05	0.46	-0.91	2.61	307	1.7
Pneum-LD	0.30	0.15	4.7	0.16	0.08	0.47	-1.88	3.93	468	1.6
Pneum-AD	0.12	0.10	1.8	n.a.0.08	0.07	n.a.0.78	-0.98	0.93	405	0.7
Pneum-AD	0.12	0.15	1.8	n.a.0.13	0.10	n.a.0.79	-2.10	1.40	612	0.7
Pneum-AD	0.21	0.10	3.2	0.11	0.05	0.46	-0.81	1.78	312	1.3
Pneum-AD	0.21	0.15	3.2	0.16	0.07	0.46	-1.85	2.64	465	1.2
Pneum-AD	0.30	0.10	4.6	0.11	0.04	0.37	-0.71	2.57	260	1.7
Pneum-AD	0.30	0.15	4.6	0.17	0.06	0.37	-1.53	3.86	391	1.6
Pneum-HD	0.12	0.10	1.7	n.a.0.07	0.06	n.a.0.79	-0.53	0.80	354	0.8
Pneum-HD	0.12	0.15	1.7	n.a.0.10	0.08	n.a.0.80	-1.02	1.17	521	0.9
Pneum-HD	0.21	0.10	3.0	0.11	0.04	0.37	-0.42	1.70	260	1.3
Pneum-HD	0.21	0.15	3.0	0.17	0.06	0.38	-0.97	2.51	385	1.3
Pneum-HD	0.30	0.10	4.2	0.12	0.03	0.27	-0.33	2.53	203	1.7
Pneum-HD	0.30	0.15	4.2	0.18	0.05	0.27	-0.79	3.81	312	1.6
DowVar	0.12	0.10	1.7	n.a.0.07	0.06	n.a.0.80	-0.21	0.81	350	0.5
DowVar	0.12	0.15	1.7	n.a.0.11	0.09	n.a.0.80	-0.71	1.25	541	0.4
DowVar	0.21	0.10	3.0	0.11	0.04	0.41	-0.25	1.65	263	0.8
DowVar	0.21	0.15	3.0	0.16	0.07	0.42	-0.55	2.48	400	0.7
DowVar	0.30	0.10	4.2	0.12	0.04	0.31	-0.23	2.49	216	1.1
DowVar	0.30	0.15	4.2	0.18	0.06	0.31	-0.55	3.77	330	0.9
DowUni	0.12	0.10	1.7	n.a.0.08	0.06	n.a.0.74	-0.86	0.87	342	1.5
DowUni	0.12	0.15	1.7	n.a.0.12	0.09	n.a.0.74	-2.06	1.35	532	1.4
DowUni	0.21	0.10	3.0	0.11	0.04	0.39	-1.09	1.73	264	3.0
DowUni	0.21	0.15	3.0	0.17	0.07	0.40	-2.54	2.64	404	2.9
DowUni	0.30	0.10	4.2	0.11	0.04	0.33	-0.92	2.53	227	3.6
DowUni	0.30	0.15	4.2	0.17	0.06	0.34	-1.99	3.84	352	3.5

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Commented [EH7]: Removed these values, as requested by reviewer 2, because no observations above canopies at 12 cm water depth.

242

243 [2.32.4](#) Data processing

244 Data were filtered for correlations $\geq 70\%$ and signal-to-noise ratios ≥ 15 dB [cf. 43] and the
 245 procedure of Hurther & Lemmin [44] was applied to remove Doppler noise from the

246 turbulent velocities by compensating for the covariance between the two vertical velocity
247 signals that were obtained by the VP. Profiles of Doppler noise exhibited a parabolic shape,
248 with a minimum around the ‘sweet spot’ of the instrument and larger values towards the
249 extremities of the profiles [45]. Doppler noise only affects the turbulence intensities and
250 hence estimates of turbulent kinetic energy. The Reynolds stress, computed with the
251 horizontal and vertical velocities obtained through the same beam pair of the VP, is
252 unaffected by this noise term [44].

253
254 Time-averaged ($\bar{u}, \bar{v}, \bar{w}$) and turbulent (u', v', w') velocities were extracted from each pre-
255 processed data record, with u , v and w denoting the streamwise, spanwise and vertical
256 velocity components, respectively. Turbulent kinetic energy $k = \frac{1}{2}(u'^2 + v'^2 + w'^2)$ and
257 Reynolds stress $u'w'$ (i.e. Reynolds shear stress normalized by the density ρ) were computed
258 for each location at every time. Parameters were then time-averaged over the length of the
259 data records (denoted with overbars). Spatial averages (denoted with angle brackets) were
260 obtained by averaging over all 30 separated vertical profiles (Figure 1c), or 10 profiles for the
261 plane bed experiments [cf. 46].

262
263 After application of the above filtering procedures, temporally- and spatially-averaged
264 velocity profiles showed a smooth pattern over the water depth. The turbulence profiles,
265 however, still showed substantial variability along the vertical after correction (to a lesser
266 extent also shown in the Reynolds stress profiles; see e.g. Figure 3b,c), even though Brand et
267 al. [45] found that Hurther & Lemmin’s noise removal procedure effectively improved VP
268 data accuracy. Consequently, this paper focusses on the point observations of turbulence
269 properties deemed to be most reliable, located in the ‘sweet spot’ of the VP at 50 mm below

270 the probe [47], thus reducing data profiles to a discrete series of point observations with a
271 vertical spacing of 45 mm.

272

273 The free-stream velocity over the canopy U_∞ was defined as the (horizontally-averaged)
274 velocity observed nearest to the water surface, i.e. at 0.05 m below the water surface
275 accounting for the instrument size and blanking distance. To calculate the depth-averaged
276 velocity within the canopy U_c , the velocity profiles were linearly interpolated across the data
277 gaps in the vertical and averaged over $0 \leq z \leq h_c$, (Table 1). These interpolations were based
278 on the 10 data points around the sweet spots (to avoid outliers impacting the interpolations),
279 while data from the lowest observation towards the bed were all maintained as there was no
280 interpolation possible. The same procedure was applied for the plane bed experiments to
281 obtain the equivalent average velocity U_o over each of the heights h_c . Additionally, standard
282 logarithmic velocity profiles were fitted to the spatially-averaged velocity data from the plane
283 bed experiments which resulted in a roughness height z_o of 1.5×10^{-4} m for the foam bed in the
284 flume [see e.g. 48].

285

286 Wake structures behind individual canopy elements depend on the stem Reynolds number
287 $Re_d = U_c d / \nu$, where ν is the kinematic viscosity of water. Nepf [27] defines the transition
288 from a laminar to turbulent flow regime in vegetated flows as $Re_d \approx 200$. In the present study
289 Re_d varied between 200-650 (Table 2). Hence, all experiments with vegetation were in the
290 turbulent regime. These stem Reynolds numbers were similar to the vegetation Reynolds
291 numbers observed in tidal marshes [27, 49].

292

293 The wakes of individual canopy elements will interact at increasing vegetation densities,
294 enhancing local turbulence and reducing the drag coefficient of individual canopy elements

295 [27, 50]. Hence, the drag coefficient (C_D) of canopy elements is a function of local flow
296 conditions within the vegetation canopy and the structural vegetation characteristics: density
297 N and diameter d . Following Tanino & Nepf (2008), the depth-averaged drag coefficient can
298 be determined by balancing the viscous dissipation of turbulent kinetic energy and the work
299 done by form drag within a vegetation canopy, which yields:

$$\frac{\sqrt{\langle \overline{k} \rangle}}{\langle \overline{u} \rangle} = 1.1 \cdot \left[C_D \frac{2\varphi}{(1-\varphi)\pi} \right]^{1/3}, \quad (1)$$

300 noting that in the present study, the average spacing between the pneumatophores or dowels
301 is >2 times their mean diameter across all densities. Drag coefficients for the vegetation
302 canopies could not be resolved directly through the momentum equation as the water level
303 gradient along the flume was too small to be measured accurately in these experiments.
304

305 **3 Experimental results**

306 3.1 Flow dynamics in canopies of real pneumatophores and dowel mimics

307 To examine the changes in flow dynamics arising from the shape variations and roughness of
308 natural pneumatophores, relative to the commonly-used smooth dowel mimics with a
309 constant diameter (Figure 2d), the results from the experiments with the high-density real
310 pneumatophore canopy are compared to the artificial canopies with the variable-height and
311 uniform-height dowel mimics (Table 1).
312

313 Spatially-averaged depth-integrated velocities within the canopy U_c decreased by up to 51%
314 in the pneumatophore canopy compared to the plane bed velocities U_0 which are depth-
315 averaged over the same vertical extent. Compared to the real pneumatophore canopy, U_c
316 increased with 10 % and 17 % for the variable-height and uniform-height dowel canopies,
317 respectively, at $h = 0.30$ m (Figure 3a). Within-canopy flow velocities were faster for a

318 reduction of the water depth and hence the submergence h/h_c of the pneumatophore canopy
319 (Figure 3d); a trend that was observed for all canopies and flow rates (Table 2). The ratio
320 U_c/U_0 was inversely correlated with water depth and hence with canopy submergence for all
321 canopies (Figure 4a), as a greater fraction of the flow got forced through the canopy in the
322 shallower flows.

323

324 The increase of the within-canopy velocities in both artificial dowel canopies compared to the
325 pneumatophore canopy could be explained by the slightly smaller density of the dowels
326 compared to the pneumatophores, reducing the total canopy resistance (Figure 4a). Although
327 the dowels had an average diameter and density very close to the real pneumatophores (Table
328 1), the tapered shape of the pneumatophores led to a change in the gradient of the vertical
329 velocity profile. In particular close to the bed, where the real pneumatophores were widest,
330 faster flow speeds were observed in the experiments with the variable-height dowels (Figure
331 3a). Within the uniform-height dowel canopy, the velocity increase was more pronounced in
332 the region around the top of the dowel canopy (at $z = h_c$) when compared to the high-density
333 pneumatophore canopy. With both dowel canopies having the same density, these differences
334 of within-canopy flow patterns are owing to the varying height structure of the canopies.

335

336 Flow velocities above each of the three canopies were higher than those over a plane bed for
337 heights exceeding two times the canopy height ($z \gtrsim 2h_c$; Figure 3a). In this region, velocities
338 above the variable-height dowel canopy were smaller than the flow velocities over the
339 pneumatophores, due to the fixed flow forcing and the higher within-canopy velocity in the
340 dowels. For the uniform-height dowel canopy, only the free-stream velocities above $z/h_c \approx$
341 2.5 were reduced compared to velocities over the pneumatophore canopy.

342

343 Distortion of the vertical flow profile due to the canopies was expressed by the ratio of the
344 within-canopy velocity U_c to the free-stream velocity U_∞ [cf. 22, 51]. For the pneumatophore
345 cover, this ratio was 0.27 for $h = 0.30$ m, whereas for the plane bed the equivalent ratio U_0/U_∞
346 was 0.67. Compared to the experiments with real pneumatophores, the ratio U_c/U_∞ was 10-
347 15% greater for the variable-height dowels and up to 25% greater for the uniform-height
348 dowels (Table 2). The flow forcing, as imposed by the fixed impeller frequencies for the low
349 and high flow regimes at each water depth, had no effect on the ratio of the flow velocity
350 within the vegetation and the free-stream velocity above the vegetation canopy and U_c/U_∞
351 was constant across each set of experiments with the same canopy and water depth (Table 2).
352 When normalized by the spatially averaged velocity at the top of the canopy $\langle \bar{u} \rangle_{z=h_c}$ [cf. 23],
353 the velocity profiles for each canopy collapsed for the three water depths and two flow rates
354 (Figure 3g).

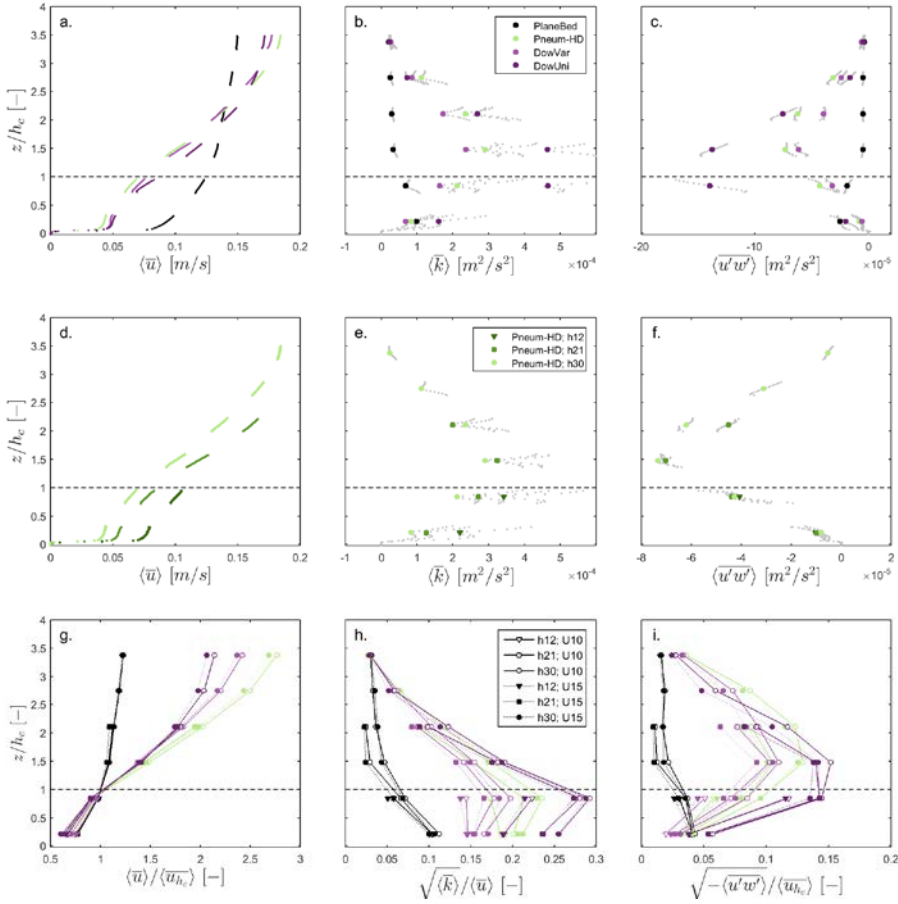
355
356 The vertical shear at the top of the canopies is associated with the generation of Kelvin-
357 Helmholtz vortices [29]. Consequently, the turbulence and Reynolds stresses peaked around
358 the inflection points of the velocity profiles. Turbulence maxima occurred at $z \approx 1.5h_c$ for the
359 pneumatophores and variable-height dowels (Figure 3b,c), a level that equated to the mean
360 plus one standard deviation height of these canopies (Table 1). However, for the uniform-
361 height dowel canopy, turbulence maxima were observed closer to the canopy at $z \approx h_c$, similar
362 to results from past studies with uniform-height dowels [e.g. 20, 22, 29, 52].

363
364 Turbulence was less intense for the experiments with variable-height dowels than with real
365 pneumatophores, but the much greater gradient of the velocity profile at the interface of the
366 uniform-height dowel canopy resulted in a stronger shear-layer with significantly greater
367 vortex-generating capacity (Figure 3b,c). The maximum reduction of turbulence and

368 Reynolds stress in the variable-height dowel canopy compared to the pneumatophores was
369 observed at $z \approx 2.1h_c$ and ranged around 25% and 34%, respectively, for the experiments with
370 0.30 m water depth. Conversely, near the top of the uniform dowel canopy, at $z \approx 0.8h_c$,
371 turbulence increased by about 114% and Reynolds stresses had increased by 228% compared
372 to the pneumatophore canopy. Differences in turbulence and Reynolds stresses right at the
373 top of the dowel canopy may have been even more pronounced, but these could not be
374 resolved from the data (see section 2.3). The enhanced turbulence in the uniform-height
375 dowels was observed throughout the canopy, causing near-bed Reynolds' s stresses to be of
376 similar magnitude as over the plane bed, whereas both variable height canopies (natural and
377 mimics) caused a much greater attenuation of these stresses (Figure 3c).

378
379 A reduction of the canopy submergence increased the turbulent kinetic energy (Figure 3e)
380 within and just above the canopies [cf.29], owing to the enhanced within-canopy velocities
381 occurring at shallower water depths. Vertical profiles of the turbulence intensity $\sqrt{\langle \bar{k} \rangle} / \langle \bar{u} \rangle$
382 and the normalized Reynolds' stresses $\sqrt{-\langle u'w' \rangle} / U_c$ were similar between experiments with
383 different flow rates (Figure 3h,i), but profiles did not collapse for the different water depths
384 (cf. the normalized velocity profiles in Figure 3g). However, the reduction of the near-bed
385 turbulence and Reynolds stress in the variable-height dowels and their increase in the
386 uniform-height dowels, compared to the high-density pneumatophores, was consistent for all
387 water depths and flow regimes (Table 2).

388



389

390

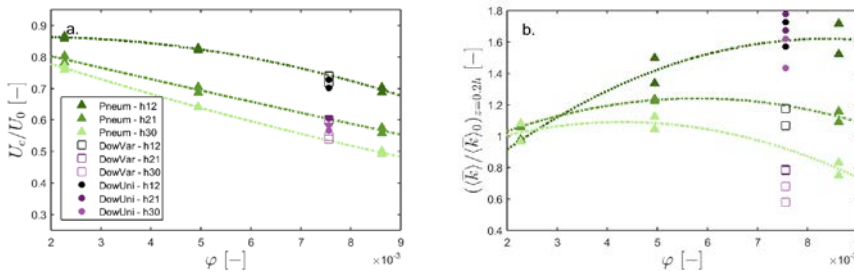
391

392 *Figure 3 – Combined results of the flume experiments for high-density real and artificial canopies. Spatially-averaged*
 393 *vertical profiles of (a) streamwise velocity, (b) turbulent kinetic energy, and (c) Reynolds stress, for a plane bed, the high-*
 394 *density canopy of real pneumatophores, dowels with variable heights that are similar to the pneumatophore heights, and*
 395 *dowels with a uniform height equal to the average pneumatophore height (see Table 1), all at 0.30 m water depth and for the*
 396 *high flow rate ($U = 0.15$ m/s). (d-f) Show the same profiles for the pneumatophore canopy as (a-c) but at multiple water*
 397 *depths (and the high flow rate $U = 0.15$ m/s). Spatially-averaged vertical profiles of (d-g) streamwise velocity scaled by the*
 398 *velocity at the top of the canopy, (e-h) turbulence intensity, and (f-i) friction velocity scaled by the velocity at the top of the*
 399 *canopy, for the different canopies (different-colours as in a-c) at all water depths (symbols as in d-f) and at both flow rates.*
 400 *Dashed horizontal lines indicate the average canopy height. Panels (b,c,e,f) present the reliable turbulence observations*
 401 *from the sweet spots (solid symbols) and the noise-affected observations for the remainder of the observed profiles (grey*

402 *dots; see text for details). Legend key: canopy types as in Table 1; $h_{12}/h_{21}/h_{30} = 0.12/0.21/0.30$ m water depth; $U_{10}/U_{15} =$
 403 $low/high$ flow rate comparing to a free-stream velocity of 0.10/0.15 m/s over the plane bed.*

404
 405 Drag coefficients for the dowels in the variable-height dowel canopy were 37-51% lower
 406 than those computed for the high-density pneumatophores, other conditions kept equal (Table
 407 2), likely owing to the smooth dowel surface compared to the relatively rough surface of the
 408 pneumatophores (Figure 2d). Conversely, the computed element drag coefficients for the
 409 uniform-height dowels were 3-4 times higher than those obtained for the variable-height
 410 dowels, even though the material was the same, and around 2 times higher than those for the
 411 real pneumatophores (Table 2). These canopy drag coefficients computed from (1) were
 412 mostly unaffected by the flow forcing (Table 2). However, the increasing within-canopy
 413 turbulence at shallower water depths resulted in a decrease of the vegetation drag coefficient
 414 with a reduction of the canopy submergence.

415



416
 417 Figure 4 – Comparison of within-canopy flow conditions relative to the no-vegetation case:(a) ratios of the within-canopy
 418 velocity to the plane bed velocity ($0 < z < h_c$) and (b) ratios of the near-bed turbulent kinetic energy ($z = 0.2h_c$) in vegetated
 419 and plane bed (subscript 0) conditions. Dotted lines are fits (cubic spline) illustrating the different trends in the flow
 420 dynamics for the real pneumatophores at the various water depths.

421

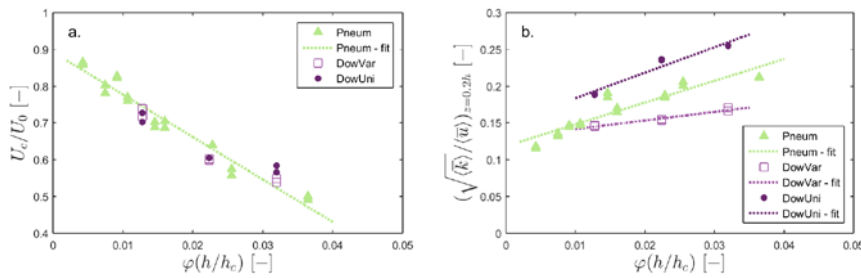
422 3.2 Effects of canopy density in real pneumatophores

423 Pneumatophore canopies with different densities were used for further comparison of the
424 hydrodynamic behaviour of canopies composed of real pneumatophores versus dowel mimics
425 (results provided in supplementary material). Spatially-averaged depth-integrated velocities
426 within the different pneumatophore canopies U_c decreased by up to about 24%, 36% and 50%
427 compared to the plane bed velocities U_0 , for the low, average and high pneumatophore
428 densities, respectively (Figure 4a). Near-bed turbulence was largest for the average canopy
429 density, with lower values observed in both the sparse and dense pneumatophore canopies
430 (Figure 4b). A similar dependency of turbulence on canopy density was found in flume
431 experiments with submerged bamboo vegetation mimics [12] and in a natural stream with
432 submerged and emergent macrophytes [53]. These results of the pneumatophore experiments
433 show evidence of the competing effects of an increasing vegetation density, causing the flow
434 to slow down while at the same time facilitating greater wake production [27]. The computed
435 drag coefficients for the pneumatophores were largely independent of the canopy density and
436 the imposed flow forcing and reduced from 1.7 to 1.3 and to 0.8 for water depths of 0.30,
437 0.21 and 0.12 m, respectively.

438
439 Depth-averaged within-canopy velocities decreased monotonically with increasing vegetation
440 densities across all experiments (Figure 4a). To enable the comparison of the results over the
441 full range of experiments, we defined an adjusted vegetation density that is the product of the
442 vegetation density ϕ and the submergence ratio h/h_c . The within-canopy velocities relative to
443 the plane bed velocities (U_c/U_0), demonstrated a significant and negative correlation ($r = -$
444 $0.98, p \ll 0.01$) with this submergence-adjusted density parameter $\phi(h/h_c)$ across all
445 experiments with pneumatophores (Figure 5a). This relationship is a result of the fact that
446 both larger vegetation densities and larger submergence ratios led to a decrease in within-

447 canopy flow speeds (Figure 4a). The same submergence-adjusted density parameter also
 448 showed a significant but positive correlation ($r = 0.92$, $p < 0.01$) with the near-bed turbulence
 449 intensities in the pneumatophore canopies (Figure 5b).

450



451

452 *Figure 5 – Correlations of the relative within-canopy flow conditions with the submergence-adjusted vegetation densities:*
 453 *(a) the ratios of the within-canopy velocities to the near-bed velocities over a plane bed (U_c/U_0) correlate significantly with*
 454 *the density parameter $\varphi(h/h_c)$ for the pneumatophore data ($r = -0.98$, $p < 0.01$) and (b) the near-bed turbulence intensity, at z*
 455 *$= 0.2h$, correlates positively with the submergence-adjusted vegetation density in each of the different canopies with $r =$*
 456 *0.92 , 0.97 and 0.97 ($p < 0.01$) for the experiments with pneumatophores (Pneum), variable-height dowels (DowVar) and*
 457 *uniform-height dowels (DowUni), respectively.*

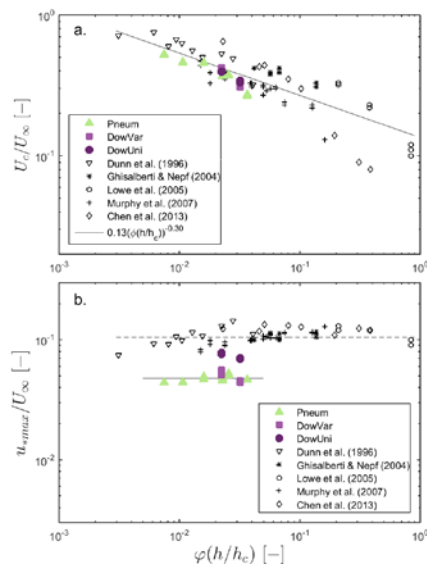
458

459 4 Discussion

460 The variable shape and height of the pneumatophores was found to have no substantial
 461 impact on the attenuation of the within-canopy velocity in comparison to the canopies of
 462 dowel mimics, as the results for the dowel experiments collapse well onto the obtained
 463 relationship between the submergence-adjusted vegetation density and the velocity
 464 attenuation U_c/U_0 (Figure 5a). This result is confirmed by a comprehensive comparison with
 465 data from prior studies using uniform-height dowels [20, 22-24, 51]. These data cover 62
 466 experiments and encompass a wide range of canopy densities and submergence ratios and
 467 demonstrate a significant and negative correlation between the calculated (or estimated)

468 ratios of the within-canopy velocity to the free-stream velocity above the vegetation (U_c/U_∞)
 469 and the submergence-adjusted vegetation density (Figure 6a). In general, the ratios U_c/U_∞
 470 from the present experiments are in good agreement with the data obtained from the past
 471 experiments with uniform-height dowels. In both the present and previous studies, an
 472 increasing density, an increasing submergence ratio of the canopy, or a combination thereof,
 473 resulted in a non-linear decrease of the ratio U_c/U_∞ , expressed by the fitted power law
 474 $U_c/U_\infty = 0.13(\varphi(h/h_c))^{-0.30}$ ($R^2 = 0.72$). Given the similarity of the ratios of the within-canopy
 475 velocity to the free-stream velocity in the pneumatophore canopies with those observed in
 476 uniform-height dowel canopies in the present and past studies, it can be concluded that the
 477 variable shape and height of the pneumatophores in this study does not result in a
 478 significantly different vertical distribution of the streamwise velocities than the idealized
 479 uniform-height dowels.

480
 481



483
 484 Figure 6 – Data from the present work in comparison to data compiled from previous studies in artificial canopies of
 485 uniform-height dowels (experiments with 0.12 m water depth were omitted from this analysis as U_∞ was unresolved). (a)

Commented [EH8]: Removed 12 cm water depth data

486 Ratios of the within-canopy velocity to the free-stream velocity were found to correlate significantly ($r=-0.64$, $p<<0.01$) with
487 the submergence-adjusted canopy density $\phi(h/h_c)$. The solid line presents a power law fit for the results of both present and
488 previous studies ($R^2 = 0.76$). (b) The maximum friction velocity scaled by the free-stream velocity varied around a constant
489 value of 0.048 ($SD = 0.004$) for the experiments with pneumatophores and variable-height dowels (solid line), which is
490 significantly lower (one-sided t-test, 95% confidence level, $p<<0.01$) than the average of 0.105 ($SD = 0.016$) throughout the
491 present and previous results for uniform-height dowels (dashed line).

492

493 The observed unique relationship between the velocity reduction and the submergence-
494 adjusted vegetation density emphasizes the governing role of the canopy submergence on the
495 dynamics of vegetated flows. All data presented consider shallow submergence ratios ($h/h_c <$
496 5) for which the flow within the canopy is determined by the balance between pressure
497 gradients, canopy drag and turbulent stresses [54]. In emergent canopies ($h/h_c \leq 1$), water
498 flow through the canopy is primarily the result of the balance between the pressure gradient
499 and the vegetative drag only, as the contribution by (stem-scale) turbulence often is
500 negligible. For submerged vegetation ($h/h_c > 1$), the shear layer at the top of the vegetation
501 contributes to the turbulent stresses within the vegetation. In a canopy consisting of uniform
502 dowels, the characteristic scale of the shear layer turbulence and, hence, the penetration of
503 turbulent stresses into the vegetation, is limited for submergence ratios $1 < h/h_c < 2$ [29].

504

505 The similarity between the experiments with dowels and with pneumatophores did not extend
506 to the turbulence in the canopy shear layer (Figure 5b) and our results showed that the height-
507 variation of the vegetation density had a substantial effect on the turbulence production. The
508 maximum value of the spatially-averaged non-dimensional friction velocity, i.e.

509 $u_{*max} = \sqrt{-\langle u'w' \rangle_{max}}$, scaled by the free-stream velocity, was found to be approximately

510 constant over the full range of conditions in both the collection of past experiments and in the
511 present experiments with uniform-height dowels, with a mean value of 0.105 (Figure 6b).

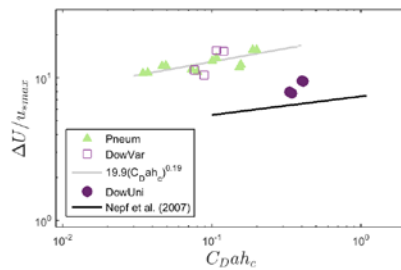
512 This finding was consistent with observations from Lowe et al. [22], who concluded that the
513 non-dimensional canopy-induced friction velocity was about 0.1 throughout their
514 experiments. However, the present experiments with variable-height canopies of
515 pneumatophores and dowels produced significantly lower values (one-sided t-test, 95%
516 confidence level, $p \ll 0.01$) for this non-dimensional friction velocity, with a mean value of
517 0.048 (Figure 6b).

518
519 The reduction of the maximum friction velocity over the variable-height canopies, even
520 though the vertical distribution of the streamwise velocity profile for these canopies was
521 similar to the uniform-height dowel canopies, can be attributed to changes in the canopy
522 shear layer owing to the gradual decline of the vegetation density along the vertical. The
523 turbulent exchange of momentum at the top of the canopy is controlled by shear-scale
524 vortices, and so the friction velocity at the top of the canopy $u_{*hc} = \sqrt{-\langle u'w' \rangle_{z=hc}}$ scales with
525 the velocity gradient across the water column, or the vortex velocity scale, ΔU [28]. Based on
526 a subset of the data collated in Figure 6, Nepf et al. (2007) obtained a positive relationship
527 between the ratio $\Delta U/u_{*hc}$ and the canopy drag parameter C_{Dahc} (Figure 7). This relationship
528 indicates that the exchange of momentum becomes less efficient as the vortex scale decreases
529 for denser canopies, diminishing the Reynolds stress for a given velocity gradient. We
530 computed the same parameters for our experiments (Figure 7), substituting the friction
531 velocity at the top of the canopy with the maximum friction velocity u_{*max} (where u_{*max} equals
532 u_{*hc} for uniform-height canopies), and using $\Delta U = U_\infty - U_c$. The uniform-height dowel
533 experiments of the present study compared well with the relationship derived by Nepf et al.
534 (2007). However, the variable-height canopies produced much higher values for $\Delta U/u_{*max}$
535 (Figure 7). This difference coincides with the consistently lower values of the normalized

536 maximum friction velocity in Figure 6b for these canopies, due to its similarity with the
 537 inverse of the ratio $\Delta U/u_{\max}$. The reduced efficiency of the vertical momentum exchange
 538 across the pneumatophores and variable-height dowel canopies is associated with the
 539 heterogeneity of the canopy height, preventing the formation of a confined canopy shear layer
 540 as it is observed in the case of a discrete density transition at the top of uniform-height dowel
 541 canopies (Figure 3c; cf. Ghisalberti & Nepf, 2004).

542
 543 The canopy shear layer is associated with an inflection point in the velocity profile, which
 544 was observed in all velocity profiles for the two highest water depths used here (Figure 3g,
 545 Figure 9d), similar to the observations of King et al. [5]. The canopy drag parameter for the
 546 present experiments was as small as $C_{Dah_c} = 0.03$, whereas previous studies only identified a
 547 well-defined shear layer and inflection point in the velocity profile for $C_{Dah_c} > 0.1$ [23, 28,
 548 52]. Moreover, previous studies identified velocity inflection points and turbulence maxima
 549 at the top of the uniform-height vegetation at $z = h_c$ [e.g. 5, 28], but for the variable-height
 550 canopies in the present study inflection points and turbulence maxima were observed at
 551 height $z \approx 1.5h_c$, equating to the mean pneumatophore (or dowel) height plus one standard
 552 deviation of the height of the canopy elements (Table 1).

553



554
 555 *Figure 7 – Comparison of the ratio of the vortex velocity scale ΔU and the maximum friction velocity u_{\max} for different*
 556 *vegetation types (for uniform-height canopies this maximum occurs at $z \approx h_c$, for variable-height canopies at $z \approx 1.5h_c$). The*

557 *black line represents the relationship obtained by Nepf et al. (2007) for uniform-height dowels. The ratios for the*
558 *pneumatophores and the variable-height dowels also correlate positively with the canopy drag parameter C_{Dah_c} as*
559 *indicated by the grey line ($r=0.69$, $p<0.01$), but are substantially higher than those for the uniform-height dowels.*
560 *(Experiments with 0.12 m water depth were omitted from this analysis as U_w was unresolved.)*

561

562 The spatial distribution of the vegetation elements within canopies was found to have no
563 significant impact on the flow dynamics. All canopies in the present study were based on the
564 natural patchy spatial distribution of the pneumatophores. In Figure 6, the results of the
565 uniform-height dowel experiments from the present study were in good agreement with
566 results from previous studies using random [24, 51] as well as staggered dowel placements
567 [20, 22, 23]. This similarity implies that the non-uniform spatial distribution of the vegetation
568 elements within the canopy is of lesser importance relative to the impact of the non-uniform
569 height of the dowels or pneumatophores.

570

571 The results provide evidence that simulations of a natural non-uniform canopy with uniform-
572 height dowel mimics could significantly over-estimate the maximum Reynolds stress in the
573 canopy shear layer and hence its contribution to transport of sediment over and into the
574 vegetation canopy. The canopy shear layer that develops at the top of the submerged canopies
575 enhances turbulent mixing and plays a crucial role in the exchange of both momentum and
576 mass between the canopy and the flow above [8]. The maximum Reynolds stress for the
577 dense pneumatophore canopy in this study was observed to be up to 2.9 times greater than the
578 maximum stress over a plane bed under the same flow conditions (Figure 3c). This result is
579 consistent with observations by Lacy and Wyllie-Echeverria [33] of Reynolds stresses at the
580 top of macro-tidal eelgrass canopies being 2-4 times greater than the maximum stress over
581 the plane sea bed. However, when the pneumatophores were replaced by uniform-height

582 dowels, the maximum stress in the canopy shear layer was up to 5.6 times greater than over
583 the plane bed (Figure 3c).

584

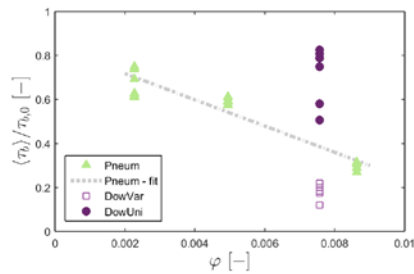
585 Penetration of the turbulence from the canopy shear layer into the vegetation canopy is a key
586 parameter in assessing the influence on sediment entrainment and has been shown to be
587 reduced for high density canopies [55]. We estimated the spatially-averaged bed shear stress
588 from the near-bed Reynolds stress: $\langle \tau_b \rangle = -\rho \langle \overline{u'w'} \rangle_{z=0.2h_c}$ at 0.015 m above the bed ($z = 0.2h_c$).

589 The ratio of the bed shear stresses in the pneumatophore canopies and the bed shear stress
590 over the plane bed $\langle \tau_b \rangle / \tau_{b,0}$ showed a significant negative correlation with the pneumatophore
591 density (Figure 8), irrespective of the level of submergence of the canopy, indicating that
592 erosion potential was reduced for higher pneumatophore densities. Similarly, bed shear
593 stresses in eelgrass beds were observed to be about 5.7 times smaller than at a nearby bare
594 site [56]. The reduction of the bed shear stress to sub-critical values and hence the reduction
595 of sediment transport capacity within (dense) vegetation canopies has been observed to
596 enhance sediment deposition in both rigid and flexible submerged canopies [12, 56, 57].

597

598 Reduced vertical variability of the canopy density within a canopy of variable-height dowels
599 compared to the (tapered) pneumatophores, combined with the smoother surface of the
600 dowels, was found to reduce the near-bed turbulence intensity (Figure 5b), giving rise to a
601 substantial decrease in the erosion potential within the canopies (Figure 8). Compared to the
602 high-density canopy of real pneumatophores, we observed markedly lower near-bed
603 turbulence in the variable-height dowel canopies (Figure 4b) and bed shear stresses were
604 about 2 times lower than the expected value for a real pneumatophore canopy of similar
605 density (Figure 8), even though the scaled maximum friction velocity was similar for those

606 two canopy types (Figure 6b). The lower near-bed turbulence in the variable-height dowels is
 607 a combined effect of the impact of the tapered shape of the pneumatophores, enhancing the
 608 vertical gradient of the vegetation density, on the within-canopy velocity distribution (Figure
 609 3a) and the reduced turbulence production around the smoother surface of the mimics (Figure
 610 2d; Figure 3b,c).
 611



612
 613 *Figure 8 – The ratio of the spatially-averaged bed shear stresses in the pneumatophore canopies $\langle \tau_b \rangle$ to the bed shear*
 614 *stress on the plane bed $\tau_{b,0}$ declines significantly at higher pneumatophore densities, as shown by the linear fit ($r=-0.95$,*
 615 *$p \ll 0.01$). Bed shear stresses in the artificial dowel canopies do not comply with the relative bed shear stress predicted for a*
 616 *pneumatophore cover of the same density, as shown by the fitted line.*

617
 618 Variability in the height of the canopy reduced the near-bed turbulence intensity (Figure 5b)
 619 and scaled friction velocity (Figure 6b), causing a substantial decrease of the erosion potential
 620 in such canopies compared to a uniform-height dowel canopy. Computed bed shear stresses
 621 in the uniform-height dowels were 3-5 times higher than those for the variable-height dowels,
 622 and they spanned a much wider range than any of the other canopies did for a single density
 623 (Figure 8). The much greater exposure of the bed in uniform-height canopies compared to
 624 variable-height canopies follows naturally from the observations that (i) turbulence intensity
 625 and Reynolds stresses in the canopy shear layer are much greater, and (ii) maximum
 626 Reynolds stresses were observed closer to the bed, at $z \approx h_c$ instead of $z \approx 1.5h_c$ (Figure 3b,c).

627 Consequently, bed shear stresses in the uniform-height dowel canopy were up to two times
628 greater than would have been expected for a real pneumatophore canopy of similar density
629 (Figure 8), particularly for the deeper submerged canopies. Based on these observations we
630 can extend the findings by Liu et al. [35], who observed a reduction of the maximum
631 turbulence intensity in a double-layer canopy compared to a uniform-height canopy, to
632 natural heterogeneous vegetation canopies.

633
634 Previous experimental studies with idealized uniform-height canopies [e.g. 20, 22, 28, 55, 57]
635 may not provide appropriate results to directly infer sediment dynamics in natural aquatic
636 canopies that typically have a non-uniform height distribution. Likewise, turbulence-
637 resolving numerical model simulations often assume uniform vegetation properties, including
638 vegetation height [e.g. 12, 24, 52, 58, 59], thereby not accounting for the identified impacts of
639 heterogeneity in vegetation canopies on flow dynamics. Our findings suggest that a constant
640 scaling parameter of ~ 0.45 could be applied to the maximum friction velocity (scaled by the
641 free-stream velocity) for a uniform-height canopy to obtain a value representative of a natural
642 variable-height canopy, regardless of the canopy density and submergence (Figure 6b),
643 keeping in mind that at the same time this maximum translates from the top of the canopy (z
644 $\approx h_c$) to a height similar to the average plus the standard deviation of the canopy height ($z \approx$
645 $\mu(h_c) + \sigma(h_c)$). Following these modifications to the turbulent properties, a similar scaling
646 could be developed for near-bed turbulence intensities (c.f. Figure 5b) and bed shear stresses
647 (c.f. Figure 8) in natural canopies, but these relations would require further investigation to be
648 confirmed across a broader range of uniform-height dowel canopy densities.

649

650 **5 Conclusions**

651 Natural canopies of mangrove pneumatophores often exhibit a heterogeneous distribution
652 with substantial variability in both height and diameter of vegetation elements [e.g. 37, 38].

653 Such heterogeneous natural canopies deviate significantly from the uniform dowel canopies
654 that are frequently used in experiments to study flows in aquatic vegetation.

655

656 Heterogeneity of the canopy height was found to significantly reduce the strength of the
657 canopy shear layer in comparison to a canopy with uniform height. Maximum friction
658 velocities, scaled by the free-stream velocity, were two times greater over uniform-height
659 dowel canopies in the present and previous studies than over a natural pneumatophore canopy
660 or dowel mimics with a similar, natural height distribution. Apart from the reduced
661 turbulence over the variable-height pneumatophores and dowels, the turbulence maxima were
662 also observed to be shifted upward, at $z \approx 1.5 h_c$ which equated to the mean plus the standard
663 deviation of the height of the canopy elements, as opposed to at h_c for uniform-height dowel
664 canopies. These two effects combined caused variable-height canopies to be less susceptible
665 to enhanced near-bed turbulence levels. Turbulent stresses in the pneumatophore canopies
666 were substantially lower and bed shear stresses were up to two times smaller in
667 pneumatophore canopies compared to uniform-height dowel canopies with similar densities.

668

669 Conversely, the variability in height and shape in pneumatophore canopies were found to
670 have a very limited impact on the vertical velocity distributions in vegetated flows, as did the
671 spatial distribution of the vegetation elements. The ratio of the within-canopy velocity to the
672 free-stream velocity above the canopies showed a monotonous decrease with an adjusted
673 density parameter $\phi(h/h_c)$ that incorporates the submergence ratio of the canopy. This trend

674 was confirmed across present and past experiments with real pneumatophores and vegetation
675 mimics that covered a wide range of flow conditions and canopy arrangements.

676

677 The present findings demonstrate substantial differences in the hydrodynamics between
678 canopies of real pneumatophores and dowel mimics, which may have significant implications
679 for predictions of sediment transport and deposition within vegetated regions. Based on the
680 observed reductions of maximum friction velocities and bed shear stresses in variable-height
681 canopies, real pneumatophores may be much more efficient sediment traps than can be
682 predicted based on uniform-height dowel experiments and model simulations.

683

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689

690 **Notation**

a	Frontal canopy density per unit area [m^{-1}]
C_D	Canopy element drag coefficient [-]
d	Canopy element diameter [m]
h	Water depth [m]
h_c	Canopy height [m]
k	Turbulent kinetic energy [m^2/s^2]
N	Canopy density [m^{-2}]
Re	Flow Reynolds number [-]
Re_d	Stem Reynolds number of the canopy [-]
u, v, w	Streamwise (x), spanwise (y) and vertical (z) velocities [m/s]
u', v', w'	Turbulent streamwise (x), spanwise (y) and vertical (z) velocities [m/s]
$u'w'$	Reynolds stress [m^2/s^2]
u_*	Friction velocity [m/s]
u_{*h_c}	Friction velocity at the top of the canopy $z = h_c$ [m/s]

Commented [EH9]: As suggested by reviewer 1.

u_{\max}	Maximum friction velocity over the full water depth h [m/s]
U	Free-stream velocity prior to canopy introduction [m/s]
U_c	Depth-averaged within-canopy velocity for $0 \leq z \leq h_c$ [m/s]
U_0	Depth-averaged plane bed velocity for $0 \leq z \leq h_c$ [m/s]
U_∞	Free-stream velocity (at $z = h - 0.05$ m) [m/s]
z	Height above bed [m]
ΔU	Canopy vortex velocity scale $U_\infty - U_c$ [m/s]
ρ	Mass-density of water [kg/m^3]
τ_b	Bed shear stress [N/m^2]
$\tau_{b,0}$	Bed shear stress for plane bed without canopy [N/m^2]
φ	Cross-sectional canopy density [-]
$\varphi(h/h_c)$	Submergence-adjusted canopy density parameter [-]
$\bar{\quad}$	Temporally averaged variable
$\langle \dots \rangle$	Spatially averaged variable

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860

861

862 6 Supplementary material

863 Density effects in real pneumatophore canopies

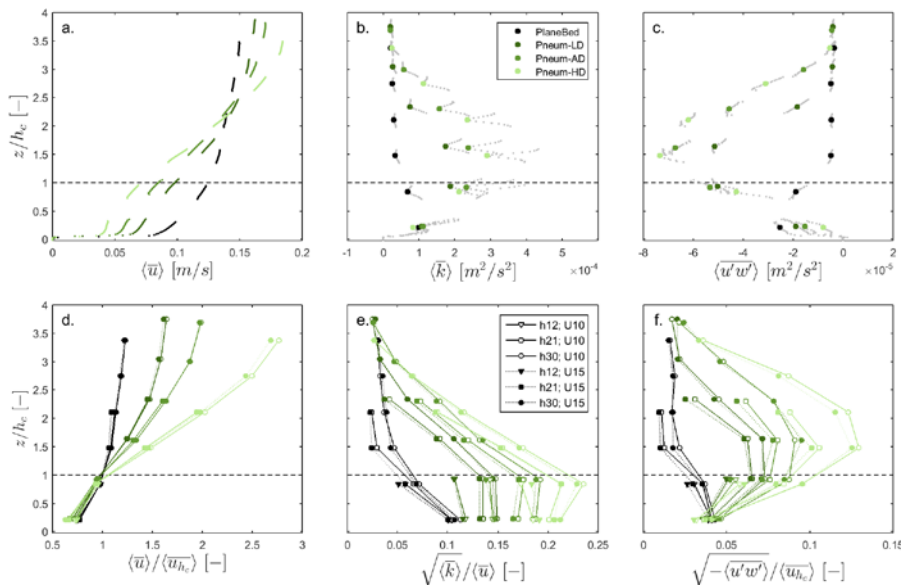
864 This supplement presents the results of the experiments with the different pneumatophore
865 densities (Pneum-LD, Pneum-AD, Pneum-HD; see Table 2). Significant reductions of the
866 within-canopy flow velocities were observed for all three densities of the real pneumatophore
867 canopies, compared to the plane bed scenario, accompanied by enhanced over-canopy flow
868 (Figure 9a). Velocity reductions occurred for $z > 2h_c$, above which faster flow velocities were
869 observed compared to the plane bed scenario. For the experiments with 0.30 m water depth,
870 the ratios of the within-canopy velocity to the free-stream velocity decreased from $U_c/U_\infty =$
871 0.46 for the lowest pneumatophore density, to 0.37 for the average density pneumatophore
872 cover and 0.27 for the highest pneumatophore density (Table 2). For comparison, for the
873 plane bed scenario with $h = 0.30$ m, the ratio of the depth-averaged velocity over the
874 equivalent canopy height compared to the free-stream velocity U_0/U_∞ was 0.67.

875
876 Turbulence generation in the canopy shear layer increased with the velocity gradient across
877 the top of the canopy and hence with the pneumatophore density, in agreement with previous
878 observations by e.g. Dunn et al. (1996). Conversely, the near-bed Reynolds stresses at $z =$
879 0.015 m ($z = 0.2h_c$) decreased with vegetation density, by up to 22% and 63% (compared to
880 the low pneumatophore density) for the average and high densities, respectively, owing to the
881 lower within-canopy velocities (Figure 9c).

882
883 The velocity profiles for each canopy densities, scaled by the velocity at the top of the
884 canopy, collapsed for the three water depths and two flow rates (Figure 9d), similar to the
885 results presented in Figure 3g. The profiles of the turbulence intensity and the normalized

886 Reynolds' stresses were similar for experiments with different flow rates, but did not collapse
 887 for the different water depths (Figure 9h,i).
 888
 889 The amplification of within-canopy turbulence with a reduction of the canopy submergence
 890 (Figure 3e) became more pronounced with an increase in canopy density. Near-bed ($z/h_c =$
 891 0.2) turbulent kinetic energy at the shallowest water depth was up to 25%, 72% and 166%
 892 larger than for the deepest submergence, for the lowest, average and highest pneumatophore
 893 densities, respectively. Similarly, the near-bed Reynolds stresses increased up to 18%, 38%
 894 and 61% for the low, average and high pneumatophore densities, respectively, between the
 895 deepest and the shallowest canopy submergence.
 896

897



898

899 *Figure 9 – Combined results of the flume experiments for all pneumatophore canopies. Spatially-averaged vertical profiles*
 900 *of (a) streamwise velocity, (b) turbulent kinetic energy, and (c) Reynolds stress, for all vegetation densities at 0.30 m water*
 901 *depth and the high flow rate (0.15 m/s). (d-f) Show the same as (a-c) for the greatest pneumatophore density and high flow*
 902 *rate ($U = 0.15$ m/s) at multiple water depths. Spatially-averaged vertical profiles of (gd) streamwise velocity scaled by the*
 903 *velocity at the top of the canopy, (he) turbulence intensity, and (if) friction velocity scaled by the velocity at the top of the*

904 canopy, for all canopy densities (different colours as in a-c), at all water depths and at both flow rates. Dashed *horizontal*
905 black lines indicate the average canopy height. Panels ~~(b,c,e and f)~~ present the reliable turbulence observations from the
906 sweet spots (solid symbols) and the noise-affected observations throughout the remainder of the observed profiles (grey
907 dots; see text for details). Legend key: canopy types as in Table 1; $h_{12}/h_{21}/h_{30} = 0.12/0.21/0.30$ m water depth; $U_{10}/U_{15} =$
908 low/high flow rate comparing to a free-stream velocity of 0.10/0.15 m/s over the plane bed.

909
910