Smart water channelling through dual wettability by leaves of the bamboo *Phyllostachys aurea*

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**Highlights**

- Dual hydrophobic-hydrophilic wetting of young *Phyllostachys aurea* bamboo leaf surfaces leads to water channelling and self-cleaning.
- Nanoscale roughness of epicuticular waxes combined with very-long-chain alkyl compounds underpin localised leaf wetting characteristics.
- Bioinspired replication of such dual wetting may offer potential for fog collection and dew harvesting in water-scarce regions of the world.

**Abstract**

The young leaves of the bamboo plant, *Phyllostachys aurea*, exhibit a distinct dual wetting behaviour on their adaxial surface. Contact angle analysis, variable pressure (environmental) scanning electron microscopy, gas chromatography, time-of-flight secondary ion mass spectrometry, and X-ray photoelectron spectroscopy have shown that the epicuticular wax morphology/topography and the surface distribution of chemical species underpin this water-channelling behaviour. Envisaged bioinspired applications include fog and dew harvesting in water-scarce regions of the world.

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**1. Introduction**

The study of plant surfaces [1–3], has led to the discovery of some fascinating phenomena including the superhydrophobicity of *Nelumbo nucifera* (Indian lotus) leaf [4], water adhesion on *Rosa*...
mozesii (red rose) [5], dry adhesion by Galium aparine [6], fog harvesting by Cotula fallica [7], and water channelling by Thuja plicata [8]. The understanding and replication of such surfaces is paving the way to many everyday technological applications; for example, self-cleaning designs based upon the lotus leaf have been introduced into the paint, glass, automotive, and textile industries [9]. Plant species displaying a range of wettabilities have been widely studied [10,11]. These have included different wettabilities on opposite sides of the same leaf [12], as well at different stages of leaf growth [13,14]. Quite often, these properties can be related to the native environment of the plants, as for instance in the case of the Namib Desert grass, Stipagrostis sabulicola, whose leaves collect water from mist in lieu of uptake from soil [15]. Despite the number of plant surfaces studied, the range of wetting phenomena reported has been fairly limited (such as hydrophobicity, hydrophilicity, and fog capture) [16,17].

The present study focuses on the unusual dual wetting behaviour of the young leaves of Phyllostachys aurea, a bamboo that originates in East Asia. In order to further understand this mechanism, the physical and chemical properties of these surfaces have been compared between young and old leaves using contact angle analysis, (environmental) scanning electron microscopy, gas chromatography, time-of-flight secondary ion mass spectrometry, and X-ray photoelectron spectroscopy.

2. Experimental

2.1. Preparation of Phyllostachys aurea samples

Phyllostachys aurea plant cuttings were collected from outdoor beds, situated in Northern England and Western Canada climates (both temperate oceanic climates), and their stems kept in water to minimize dehydration during transportation to nearby analytical facilities.

2.2. Contact angle measurements

Contact angles were measured at room temperature using a video capture apparatus (VCA 2500 XE, AST Products Inc.) and 1 μL high-purity water droplets (ISO 3696 grade 1). Static water contact angle values less than 90° signify a hydrophilic surface, whilst greater than 90° corresponds to a hydrophobic surface [18]. Advancing and receding contact angle values were determined by respectively increasing or decreasing the liquid drop volume [19].

2.3. Variable pressure (environmental) scanning electron microscopy (ESEM)

An environmental scanning electron microscope (XL30 ESEM-FEG, FEI Company) operating in wet mode was used to monitor water vapor condensation onto individual leaf samples in real time. The leaves were mounted onto carbon disks and attached to a Pelletier cooling stage. Vapor condensation was controlled by varying the chamber pressure and sample temperature. Electron micrographs were taken using a gaseous secondary electron detector (GSED), in conjunction with a 15 kV accelerating voltage, and a working distance between 9 and 11 mm.

2.4. Scanning electron microscopy (SEM)

Individual leaf specimens were prepared for scanning electron microscopy analysis by using a glycerol substitution process [20]. Each sample was then mounted onto a carbon disk. No gold coating was required due to the conductive nature of the glycerol [21]. Plant surface structure images were taken on a scanning electron microscope (Stereoscan 240, Cambridge Instruments Company) operating in a secondary electron detection mode, in conjunction with an 8 kV accelerating voltage, and a working distance between 9 and 35 mm.

2.5. Gas chromatography (GC)

2.5.1. Extraction of adaxial leaf wax

To selectively extract cuticular wax only from the adaxial side of the leaf, one end of a glass cylinder (Schott, Standard expansion adapter NSE 19/14, open at both ends) was gently pressed onto the leaf surface with sufficient force to create a seal between cylinder and leaf whilst not damaging the plant material, and filled three times with 1.5 mL of fresh chloroform (Aldrich, ≥99%, 0.75% ethanol as stabilizer) at room temperature, with intermittent agitation of the chloroform using a Pasteur pipette. The chloroform extracts from five randomly chosen leaf surface areas were combined, concentrated at 50 °C under a stream of N2 (Praxair, ≥99.998%), and stored prior to analysis. The total surface area per sample was calculated from the diameter of the cylinder to be 3.6 cm² [13].

2.5.2. Analysis of wax extracts

Before GC–MS/FID analysis, wax extracts were spiked with 10 μL of a 1.02 mg/mL solution of n-tetracosane (Alfa Aesar, ≥99%) in chloroform as an internal standard and subjected to derivatization with 10 μL of N,O-bis(trimethylsilyl)trifluoroacetamide (BSTFA, Aldrich, GC grade) in 10 μL pyridine (Aldrich, ≥99.8%, anhydrous) for 30 min at 70 °C. Then, they were concentrated to dryness at 50 °C under a stream of N2 and re-dissolved in 20 μL chloroform to a total volume of 20 μL. Under these conditions, amides did not undergo silylation, resulting in poor chromatographic peak shape for these wax constituents. Therefore, after re-evaporation to dryness, samples were derivatized for 1 h at 70 °C in 20 μL of benzyl bromide (Aldrich, ≥98%) in the presence of 0.1 mg NaH (Aldrich, anhydrous, ≥95%). Then, excess NaH was quenched with distilled water, the basic aqueous layer discarded and the organic layer containing benzylated derivatives taken to dryness. Finally, the resulting mixture was again subjected to silylation with BSTFA/pyridine as described above, dried, and brought to a final volume of 20 μL with fresh chloroform.

Two different GC instruments were used for separation and detection of wax constituents, both equipped with the same type of capillary GC column (6890N, Agilent, Avondale PA, USA; 30 m long; type HP-1: 100% PDMS; 0.32 mm i.d.; df = 0.1 μm) and following the same temperature program (on-column injection at 50 °C, constant for 2 min, ramp 40 °C min⁻¹ to 200 °C, constant for 2 min, ramp 3 °C min⁻¹ to 320 °C, constant for 30 min). The first instrument employed He gas (Praxair, ≥99%) as the mobile phase, at a flow rate of 1.4 mL min⁻¹, and was equipped with an MS detector (5973N, Agilent, EI-70 ev), primarily serving the purpose of qualitative identification of the separated wax compounds. The second instrument used H2 carrier gas (Praxair, ≥99.95%) at 2.0 mL min⁻¹ and an FID detector for the quantification of individual wax homologs/isomers, based on normalization of their peak areas against that of the internal standard. It had been shown previously that most wax compounds have relative response factors of 1.00 to this standard under very similar GC-FID conditions [22]. Quantitative compositions of aliphatic ester metamers were determined from GC–MS data as described by Lai et al. [23].

2.6. Time-of-Flight secondary ion mass spectrometry (ToF-SIMS)

The leaf samples for ToF-SIMS analysis were mounted onto a piece of double-sided tape (3M ‘Scotch Tape’, grade 665) attached to a piece of clean PET film (Melinex, ‘O’ grade). A second piece of PET film, containing a square hole in the centre, was mounted over the leaf in an attempt to minimize the tendency for the leaf
margins to curl over. This assembly was attached to the base plate of a sample holder using stainless steel clips.

Static SIMS analysis was carried out using an instrument of single-stage reflectron design (TOF-SIMS IV – 200, ION-TOF GmbH) [24]. Positive and negative ion mass spectra and images of the leaf samples were obtained using a 20 keV Bi\textsuperscript{32+} focused liquid metal ion beam, incident at 45° to the surface normal, and operated in ‘bunched’ mode for high mass resolution (20 ns wide ion pulses at a 6.7 kHz repetition rate). Charge compensation was provided by a low-energy (ca. 20 eV) electron flood gun. The total ion dose density was kept to less than 1 \times 10^{16} ions m\textsuperscript{−2} in all cases. Sample surface topography and the ion gun mode of operation limited the mass resolution to ca. \textit{m}/\Delta \textit{m} = 1000. Positive and negative ion static SIMS spectra of the samples were recorded at room temperature with a 128 \times 128 pixel raster and a field of view of 200 \mu m \times 200 \mu m, and the sample imaged with a field of view of 500 \mu m \times 500 \mu m. In addition, images and spectra were recorded from larger areas in the range 7 mm \times 7 mm up to 12.5 mm \times 12.5 mm by moving the sample under the ion beam (the so-called ‘stage raster’ mode of operation).

2.7. X-Ray photoelectron spectroscopy (XPS)

X-ray Photoelectron Spectroscopy (XPS) surface characterization was carried out using an electron spectrometer (AXIS Nova, Kratos Analytical Ltd), equipped with a monochromated Al Kα X-ray source (1486.6 eV) and a concentric hemispherical analyser combined with a spherical mirror analyser. Photoelectron spectra were collected at a take-off angle of 0° from the substrate normal (mean escape depth = 0.5–1.5 nm) [25], with electron detection in the constant analyser energy mode (CAE mode, pass energy = 160 eV for wide scan & 20 eV for high resolution spectra, step size = 1.0 eV & 0.1 eV, respectively). Rectangular samples, spanning the width of the leaf, were cut from fresh *Phyllostachys aurea* leaves and mounted, adaxial side facing upwards, onto an aluminum sample plate. Beryllium-copper clips were used to secure the samples. The leaves were preloaded into the system at a pressure of less than 90°), Table 1. These contrasting wettabilities are most noticeable in the movement of water across the leaf surface. Once large enough, droplets of water on the margin of the leaf roll towards the middle, assisted by the low contact angle hysteresis.

![Fig. 1](image)

Fig. 1. Images of the adaxial surface of *Phyllostachys aurea* leaves following rainfall: (a) a young leaf exhibiting dual wettability – the combination of hydrophobic margins and a hydrophilic middle (the image in the top corner shows a magnified view of the area outlined by a dashed box); (b) a young leaf showing the results of coalescing droplets in the middle of the leaf moving downwards to the leaf tip; (c) a dead young leaf still showing the same dual wetting property; (d) an older leaf not exhibiting the dual wettability (the image in the top corner shows a magnified view of the area of the surface outlined by a dashed box); and (e) schematic illustrating dual wetting behaviour on young leaf.
Table 1

Water contact angle values for the adaxial surface of Phyllostachys aurea leaves.

<table>
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<tr>
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<th>Water Contact Angle/°</th>
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<tr>
<td></td>
<td>Static</td>
<td>Advancing</td>
<td>Receding</td>
<td>Hysteresis</td>
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<tr>
<td>Phyllostachys aurea</td>
<td></td>
<td></td>
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<td></td>
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<tr>
<td>Adaxial Surface</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Young Leaf (Margin)</td>
<td>141 ± 3</td>
<td>142 ± 2</td>
<td>136 ± 4</td>
<td>5 ± 4</td>
</tr>
<tr>
<td>Young Leaf (Middle)</td>
<td>77 ± 13</td>
<td>84 ± 12</td>
<td>62 ± 10</td>
<td>23 ± 8</td>
</tr>
<tr>
<td>Older Leaf (Margin)</td>
<td>69 ± 12</td>
<td>83 ± 12</td>
<td>51 ± 11</td>
<td>32 ± 6</td>
</tr>
<tr>
<td>Older Leaf (Middle)</td>
<td>69 ± 14</td>
<td>87 ± 12</td>
<td>60 ± 10</td>
<td>27 ± 7</td>
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</table>

inherent to the margin, Table 1, and the slight gradient downwards due to the concave shape of the leaf, Fig. 1(a). Pools of water collecting in the middle eventually coalesce over time, forming a downward stream-like film of water, Fig. 1(b–c). As the stream increases in volume, the water flow begins to move towards the leaf tip, assisted by the hanging gradient present between the petiole and the apex of the leaf. Eventually, the droplet formed at the apex falls to the ground. In contrast, the abaxial (underside) surface of young Phyllostachys aurea leaves consists of a different physical structure visible to the naked eye and displays an absence of dual wetting (superhydrophobicity everywhere with water static contact angle exceeding 160°).

Interestingly, this dual wetting behaviour persists in leaves which have died early in their development but are still attached to the stem, Fig. 1(c). Furthermore, this characteristic water collection mechanism associated with the younger leaves is found to be absent for older leaves of the same species, Fig. 1(d). Their adaxial surfaces appear to be hydrophilic throughout, Table 1, and there-

![Fig. 2. SEM of the adaxial surface of (A) young and (B) old Phyllostachys aurea leaves. Micrographs at increasing magnifications of the: (a) margin and (b) middle of the leaf; the “double-humped” papillae which make up the troughs on the: (c) margin and (d) middle; a tilted view of the “double-humped” papillae on the: (e) margin and (f) middle; and the nanoscale epicuticular wax crystals which cover the surface on the: (g) margin (inset shows the platelet-shaped wax crystals) and (h,i) middle. The horizontal axis in (a)–(d) is parallel to the leaf axis.](image-url)
fore water spreads to wet both the middle and margins of these leaves.

3.2. Electron microscopy

Comparison of scanning electron microscopy (SEM) micrographs for the margin and the middle of young Phyllostachys aurea leaves revealed distinct differences in surface micro- and nano-scale structure. In both regions, there were perpendicularly orientated “double-humped” papillae running parallel to the macroscopic longitudinal leaf grooves between the petiole and the leaf tip. At the margin, these papillae displayed deep troughs in between their peaks as well as between the individual pairs of papillae. In contrast, the troughs in the middle appeared shallow and considerably less defined (lower roughness). These differences were further evident for the nanoscale epicuticular wax crystals covering some of the micro-relief in both areas. The surface at the margin of the leaves is covered in a dense “carpet” of upright platelet-shaped wax crystals, approximately 0.5–1 μm in height, in contrast, the wax crystals in the middle are granular and approximately 0.1–0.5 μm high.

Scanning Electron Microscopy (SEM) showed a smooth surface in both the margin and centre of old Phyllostachys aurea leaves, as well as a layered surface with a smooth crust on the middle of young leaves. The features seen for the margin and middle of young leaves appear to have been retained. Noticeable differences when compared to the young leaves is the softening/suppression (lower roughness) in young leaves. In contrast, the troughs in the middle appeared shallow and considerably less defined (lower roughness).

3.3. Gas chromatography (GC)

To establish the identities and abundances of all lipids coating the adaxial side of Phyllostachys aurea leaves, total wax mixtures were extracted from young and old plant leaves. The waxes were identified and quantified using gas chromatography/mass spectrometry (GC–MS) and gas chromatography–flame ionization detection (GC-FID), respectively.

3.3.1. Classes of cuticular wax constituents

The total adaxial wax coverage was very similar on young and old Phyllostachys aurea leaves: 3.6 ± 0.4 μg cm⁻² and 3.3 ± 0.1 μg cm⁻², respectively. In both cases, more than 90% of the GC-detectable compounds could be identified (Fig. 4A), and cyclic terpenoids constituted almost half of the wax mixtures (1.5 ± 0.2 μg cm⁻² and 1.6 ± 0.2 μg cm⁻², respectively). They were accompanied by very-long-chain (VLC) fatty acids, primary alcohols, alkyl esters, aldehydes, alkanes, and amides. Most of these compounds were present in similar amounts on young and old P. aurea leaves. However, free alcohols were found in higher abundance on the surface of young plants as compared to old ones (0.30 ± 0.04 μg cm⁻² vs. 0.18 ± 0.01 μg cm⁻²). Overall, terpenoids together with alkyl esters and fatty acids made up three quarters of the wax mixture, in a ratio of ca. 4:1:1, irrespective of plant age.

3.3.2. Homolog/Isomer distributions within cuticular wax classes

All VLC aliphatic classes comprised a series of homologous compounds. Alkyl esters, fatty acids, alcohols, and amides were dominated by homologs with even carbon numbers, and alkanes by odd-numbered homologs. Generally, the chain length distributions of linear aliphatics were similar between adaxial surface waxes of young and old Phyllostachys aurea leaves. Free alcohols exhibited bimodal distributions, peaking at the C38 and C40 homologs irrespective of age, with slightly higher amounts of C16, C18, and C20 acids on old P. aurea leaves. The alcohol and alkane series were fairly evenly distributed, peaking at C28 and C29, respectively, in young and old leaves alike.

Three compound classes had homolog profiles changing with age. In the aldehyde fraction, the predominant homolog (C20) was accompanied by slightly more of the shorter homologs in young leaves and longer homologs in old leaves. An identical age-related chain length effect was observed for amides, also peaking at C30 and with shorter homologs more abundant for young and longer ones for old plant leaves. Further evidence for amide structure elucidation and related biology will be presented elsewhere.

Finally, the ester homolog profiles peaked at C48 on leaf surfaces of both ages, accompanied by near-identical amounts of the C60 and C44, and C22 homologs, and higher proportions of C60 and C38 homologs on young leaves versus higher abundance of the C50 homolog on old leaves.

Pentacyclic triterpenoids were found in similar amounts within the wax mixtures of young and old Phyllostachys aurea leaves, Fig. 4B. Free triterpene alcohols dominated, with β-amyrin and isomultiflorol together accounting for more than half of the terpenoid load. Other free triterpenoids (lupeol, α-amyrin, glutinol and epifriedelanol) were less abundant. Additionally, triterpenoid esters amounted to ca. 25–30% of all terpenoids, corresponding to approximately an eighth of the total adaxial wax coverage for both old and young leaves, and thus making triterpenoid and VLC alkyl esters equally abundant (compare Figs. 4A and B). Palmitates of β-amyrin, isomultiflorol, glutinol and α-amyrin were found in a 3:3:3:1 ratio on young and old leaves alike. Additionally, trace amounts of α- and γ-tocopherols were detected, Fig. 4B.
Fig. 4. (A) Compound class distribution in the wax mixtures on the adaxial side of Phyllostachys aurea leaves. Total coverages (μg cm$^{-2}$) of compound classes are given for the wax mixtures of young and old P. aurea leaves. (B) Relative compositions within wax compound classes within the wax mixtures on the adaxial side of Phyllostachys aurea leaves. Relative abundances (%) of individual homologs or isomers (in the case of terpenoids) from each compound class are given for young and old P. aurea leaves. Numbers on the x-axis indicate homolog chain length, for alkyl esters, the sum of alkyl and acyl carbon numbers. Minor odd-numbered acid and alcohol homologs as well as even-numbered alkane homologs are omitted for clarity. Bars represent mean ± standard deviation (n = 4).

To provide complete reference data for ToF-SIMS peak assignment, the distributions of acyl and alkyl moieties within the wax esters were determined. Each of the aliphatic ester homologs represented a complex mixture of isomers derived from numerous pairwise combinations of acid and alcohol chain lengths. Since such metamers could not be chromatographically resolved and thus quantified by GC-FID, their distributions were determined instead based on abundances of characteristic fragments in the mixed mass spectrum associated with the GC peak of each ester homolog [23,31–35].

Three different patterns of isomer distributions could be discerned depending on overall ester homolog chain length, Fig. 5. In young Phyllostachys aurea leaves, the shorter ester homologs (C$_{36}$–C$_{42}$) were formed primarily by C$_{20}$ alcohol and C$_{16}$–C$_{22}$ acids, the mid-range esters (C$_{42}$–C$_{44}$) included a large portion of isomers formed by C$_{26}$–C$_{28}$ alcohols and C$_{16}$ acid, while the longer esters (C$_{44}$–C$_{52}$) contained mainly C$_{22}$–C$_{30}$ alcohols and C$_{22}$ acid. In the leaf wax mixtures of old leaves, similar patterns were found, with the C$_{36}$–C$_{38}$ esters formed by C$_{20}$ alcohol and C$_{16}$–C$_{18}$ acids, the C$_{40}$–C$_{44}$ esters by C$_{24}$–C$_{28}$ alcohols and C$_{16}$ acid, and the C$_{46}$–C$_{52}$ esters by C$_{24}$–C$_{30}$ alcohols and C$_{22}$ acid. Overall, C$_{22}$ acid was the most abundant acid in the linear VLC esters, independent of leaf age (compare Figs. 4B and 5A/C). The second-most abundant acid in alkyl esters was palmitic acid (C$_{16}$), most notably for old bamboo leaves (Figs. 4B and 5C).

Finally, the overall distribution of acid and alcohol moieties across all esters could be calculated, including alkyl esters and alicyclic (triterpenyl) esters. Thus, the chain length distribution of esterified acids was found to be bimodal, with maxima at C$_{16}$ and C$_{22}$ irrespective of plant age, Fig. 6A. The esterified acids had gradually decreasing abundances from C$_{24}$ to C$_{28}$, in sharp contrast to the chain length profile of free acids gradually increasing towards C$_{28}$ (compare Figs. 4A and 6A). The absolute amounts of individual esterified acid homologs were independent of leaf age, Fig. 6A.

In contrast to the profile of esterified acids, the chain length distribution of esterified alcohols differed between young and old
Fig. 5. Relative compositions of esterified acids and alcohols within the wax mixtures on the adaxial side of Phyllostachys aurea leaves. Relative abundances (%) of individual isomers are given by chain length of their acid moiety (A/C) and alcohol moiety (B/D) for each ester homolog, in young (A/B) and old (C/D) P. aurea leaves. Bars represent mean ± standard deviation (n = 4). Each group of bars adds up to 100%.

Table 2

XPS analysis of Phyllostachys aurea leaf surfaces. The elemental composition is averaged using data from three different positions at the margin and middle of two young and two old leaves – 24 sampled regions in total. In the case of silicon, which was not detected for all six analysis runs of each leaf region, the digit in brackets corresponds to the number of leaf samples which did show the presence of silicon. The \(-\text{C}_x\text{H}_y\) hydrocarbon component at 285.0 eV is reported as a percentage of the overall C(1s) peak (set at 100%).

<table>
<thead>
<tr>
<th>Phyllostachys aurea Adaxial Surface</th>
<th>Relative% Elemental Composition</th>
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<tbody>
<tr>
<td></td>
<td>C(1s)</td>
</tr>
<tr>
<td></td>
<td>(C_{\text{total}})</td>
</tr>
<tr>
<td>Young Margin</td>
<td>96.0 ± 0.7</td>
</tr>
<tr>
<td>Middle</td>
<td>93.7 ± 0.6</td>
</tr>
<tr>
<td>Old Margin</td>
<td>84.0 ± 3.6</td>
</tr>
<tr>
<td>Middle</td>
<td>84.4 ± 1.8</td>
</tr>
</tbody>
</table>
leaves. Higher levels of C18–C22 alcohols were found in esterified wax of young Phyllostachys aurea leaves, and substantially more C26 alcohol in the wax esters of old leaves (Fig. 6B). The overall profile of esterified alcohols thus differed from that of corresponding free alcohol distributions, with the predominant chain length of C26 in the ester alcohols and C28 in the free alcohols (compare Figs. 4B and 6B). Similarly, the profiles of esterified and free triterpenols also differed (independent of age), most notably for glutinol (compare Figs. 4B and 6B). The C(1s) higher binding energy tail region comprised –C–CX at 285.7 eV (where X represents: =O, (=)NHR, –OOR, or –OOH) [36], –C–N at 286.0 eV, –OH at 286.6 eV, O=C=O[CONH] at 287.9 eV [37], and –C(=O)OR (where R is C or H), at 289.1 eV [36]. These values were consistent with those previously reported in the literature for leaf waxes of other species [38,39].

The O(1s) spectra were consistent with the C(1s) spectra, comprising complementary peak components: –CONHR at 531.6 eV, –C=O at 532.4 eV, –OH at 533.1 eV, and –(=O)OR (where R is C or H) at 533.9 eV, (Supporting Information Fig. S 3) [36]. The N(1s) envelope consisted of two peaks at 400.2 eV and 401.7 eV, characteristic of amide (–CONR) and protonated-amide (–NHR2) functionalities, (Supporting Information Fig. S 3) [36,39,40]. The latter may also be associated with hydrogen-bonding between the amide nitrogen atom and nearby hydrogen-containing functional groups, such as alcohols [41,42]. The possibilities of N(1s) amine (–NH2) or nitrate functionalities were excluded because they typically appear below 400 eV [42,43] and around 407–408 eV, respectively. Nitrogen has been detected in XPS studies of other leaf surface waxes [39].

The Si(2p) spectra could be fitted with spin-orbit splitting components Si(2p3/2)O2 and Si(2p1/2)O2 at 103.3 eV and 103.9 eV respectively, corresponding to silica-like species (Supporting Information Fig. S 3) [36,44]. No peak was present at ~101.5 eV, ruling out polysiloxanes as a potential source of silicon contamination [45]. Previous XPS studies have identified silicon in leaf surface waxes, citing silica as one of its possible forms [39].

3.4. Time-of-Flight secondary ion mass spectrometry (ToF-SIMS)

ToF-SIMS spectra of the leaf samples comprised peaks associated with whole and fragmented molecules (some of which were homologous series separated by 14/28 amu corresponding to differing numbers of CH2/CH2 units within alkyl chains), Fig. 7. Respective positive and negative ToF-SIMS spectra displayed different fragments at the margin of young leaves compared to the similar fragments detected for both the middle of young leaves and all across old leaves. The predominant species at the margin of young leaves were C22–acid esters containing mainly C24, C26, and C28 esterified alcohols (based on GC–MS ester metamer analysis) together with very-long-chain C28, C30, and C32 amides. In contrast, towards the middle of young leaves and across the whole old leaves, there was a large proportion of the shorter chain C16 carboxylic acid in addition to free alcohols. This correlates well with the higher proportion of C16–C18 free acids observed for old leaves through GC analysis, Fig. 4B.

Imaging ToF-SIMS (using the stage raster mode) showed that the very-long-chain amide species were concentrated towards the margin region of young leaves, whilst the shorter C16 carboxylic acid was localised along the middle (axial centre) of the leaf, Fig. 8. In contrast, no such spatial distribution of the same species was observed for the case of the older leaves.

3.5. X-Ray photoelectron spectroscopy (XPS)

XPS survey spectra (0–1200 eV binding energy range) of margin/middle of young/old Phyllostachys aurea leaves detected the presence of the following main peaks: C(1s), O(1s), N(1s), Si(2p) and Si(2s). The percentage carbon content was highest overall in the young leaves, whilst the amount of oxygen and nitrogen was greatest in the old leaves, Table 2.

The high-resolution C(1s) envelope contained a main –C-H hydrocarbon component at 285.0 eV (Supporting Information Fig. S 2). The C(1s) higher binding energy tail region comprised –C–CX at 285.7 eV (where X represents: =O, (=)NHR, –OOR, or –OHH) [36], –C–N at 286.0 eV, –OH at 286.6 eV, O=C=O[CONH] at 287.9 eV [37], and –C(=O)OR (where R is C or H), at 289.1 eV [36]. These values were consistent with those previously reported in the literature for leaf waxes of other species [38,39].

The O(1s) spectra were consistent with the C(1s) spectra, comprising complementary peak components: –CONHR at 531.6 eV, –C=O at 532.4 eV, –OH at 533.1 eV, and –(=O)OR (where R is C or H) at 533.9 eV, (Supporting Information Fig. S 3) [36]. The N(1s) envelope consisted of two peaks at 400.2 eV and 401.7 eV, characteristic of amide (–CONR) and protonated-amide (–NHR2) functionalities, (Supporting Information Fig. S 3) [36,39,40]. The latter may also be associated with hydrogen-bonding between the amide nitrogen atom and nearby hydrogen-containing functional groups, such as alcohols [41,42]. The possibilities of N(1s) amine (–NH2) or nitrate functionalities were excluded because they typically appear below 400 eV [42,43] and around 407–408 eV, respectively. Nitrogen has been detected in XPS studies of other leaf surface waxes [39].

The Si(2p) spectra could be fitted with spin-orbit splitting components Si(2p3/2)O2 and Si(2p1/2)O2 at 103.3 eV and 103.9 eV respectively, corresponding to silica-like species (Supporting Information Fig. S 3) [36,44]. No peak was present at ~101.5 eV, ruling out polysiloxanes as a potential source of silicon contamination [45]. Previous XPS studies have identified silicon in leaf surface waxes, citing silica as one of its possible forms [39].

4. Discussion

In contrast to earlier studies, where differences in wax structure and chemistry have been noted between different plant species [46,47], or leaves on the same plant [48], sides of the same leaf [49], or stages of development [13,50], the dual wettability observed in the present study for young Phyllostachys aurea leaves appears to be a hitherto unreported wetting behaviour. This initial dual wettability displayed during the early stage of P. aurea leaf growth may ensure self-cleaning properties that prevent microbial growth and damage.

The mechanisms underpinning wetting comprise a chemical and physical component, Table 3. Assuming a wax density of ≈1 g cm−3, the adaxial wax coverage measured by gas chromatography on young and old Phyllostachys aurea leaves (3.6 ± 0.4 μg cm−2 and 3.3 ± 0.1 μg cm−2, respectively), corresponds to at least 30 nm effective wax layer thickness. The GC results provide a complete inventory of wax constituents, some or all of which may be present at the outermost surface. An important finding is the identification of C28–C32 amides in both young and old leaves. Another is the detection of relatively high percentages of C22–acid esters in young leaves and C16–C18 free acids in wax of old leaves. These GC data have served as a reference for
compound assignments in the corresponding XPS and ToF-SIMS datasets which provided highly surface-sensitive information with sampling depths comparable to contact angle analysis (wetting). XPS showed high carbon content on the adaxial surface of the young leaves, in accordance with the greater proportion of very long-chain aliphatic molecules, detected by GC, relative to old-leaf samples. The presence of oxygen and nitrogen XPS signals also agrees well with the GC assignments made for alcohols, carboxylic acids, esters, and amides. All of the imaging XPS data analyses correlated well with the corresponding ToF-SIMS results and ESEM images displaying dual wetting. Furthermore, imaging ToF-SIMS showed that longer-chain alkyl species with relatively low oxygen content dominate the young leaf margin region, whereas shorter-chain carboxylic acids and alcohols are found in greater relative concentrations towards the middle as well as throughout old leaves.

The longer hydrocarbon chains can help explain the greater hydrophobicity at the young leaf margin [51], however this alone cannot account for the very high (+140°) water contact angles observed. There also exists a contributing physical factor, attributed to the dense array of fine nanoscale epicuticular wax platelets found only in the margin regions of young leaves. The longer-chain species present towards the margin of young leaves, as shown by imaging ToF-SIMS, may have a strong tendency to form wax crystals due to their relatively narrow chain length distribution. These crystals may trap small air pockets, and thus facilitate the greater
Given the inherent hydrophobicity of alkyl compounds found on the surface in the middle of young leaves, other factors must account for the observed hydrophilic nature in this region. This will include, the diminished surface roughness in the middle lowering hydrophobicity associated with the Cassie-Baxter wetting state \[4,52,53\].

Given that it has been reported that dual wetting hydrophobic – hydrophilic arrays can yield much higher water collection efficiencies compared to straightforward hydrophobic or hydrophilic surfaces \[59,60\], the replication of Phyllostachys aurea bamboo dual wetting hydrophobic – hydrophilic channels surfaces may offer potential scope for fog collection and dew harvesting in water-scarce regions of the world \[61\]. The observed surface wetting behaviour of leaves may also have relevance to crop spraying.

5. Conclusions

The dual wettability behaviour displayed by young leaves of the Phyllostachys aurea bamboo can be attributed to a combination of the chemical and physical differences observed between the leaf margin and middle. The epicuticular nanoscale roughness combined with the very-long-chain alkyl compounds present in the margin regions of the leaves, gives rise to much higher water contact angles (droplet movement) compared to the middle of the leaves. These differences underpin the self-cleaning properties of the bamboo. Bioinspired replication of such dual wetting surfaces may have potential for water harvesting.

### Fig. 8. Negative ion ToF-SIMS imaging of young (a–b) and old (c–d) Phyllostachys aurea leaves: (a)/(c) very-long-chain C30 amide \((m/z = -451, \text{red})\) versus long chain C22 free/esterified carboxylic acid \((m/z = -339, \text{green})\); and (b)/(d) C22 free/esterified carboxylic acid \((m/z = -339, \text{green})\) versus short chain C16 free/esterified carboxylic acid \((m/z = -255, \text{yellow})\) (for interpretation of the references to colour in this figure legend, the reader is referred to the web version of this article).
Data accessibility
Supporting data can be accessed at http://collections.durham.ac.uk.

Competing interests
We have no competing interests.

Author contributions

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Appendix A. Supplementary data
Supplementary data associated with this article can be found in the online version, at http://dx.doi.org/10.1016/j.colsurfa.2016.06.058.

References


