

1 **Supplementary Information for**

2
3 **Foliar Graphene Sensor for Continuous, Non-Destructive Monitoring of Plant Tissue**
4 **Hydration, Drought Stress and Circadian Rhythm**

5 Ning Liu¹, Pascal Mahukpe Ayelo^{2,*}, Utkarsh Misra^{1,*}, Maria Lynn Ulatowski³, Caixing Xiong²,
6 Neelotpala Kumar¹, Michael Kolomiets², Henry Fadamiro², Jean Anne Incorvia¹, Ashley M.
7 Matheny³, Dmitry Kireev^{4,&}, and Deji Akinwande^{1,5,&}

8 ¹ Department of Electrical and Computer Engineering, University of Texas at Austin, Austin,
9 Texas 78712, USA

10 ² Department of Entomology, Texas A&M University, College Station, TX 77843

11 ³ Department of Earth and Planetary Sciences, The University of Texas at Austin, Austin, Texas
12 78712, USA

13 ⁴ Department of Biomedical Engineering, University of Massachusetts Amherst, Amherst, MA
14 01002, USA

15 ⁵ Department of Biomedical Engineering, University of Texas at Austin, Austin, Texas 78712,
16 USA

17
18 * These authors contributed equally

19 & Corresponding authors: deji@ece.utexas.edu and dkireev@umass.edu

20 **Supplementary Note 1. Comparison of Plant Water Content Monitoring Techniques**

21 Accurate measurement of plant water status is essential for understanding plant physiology
22 and improving irrigation management. Traditionally, leaf water content (LWC) has been quantified
23 using destructive gravimetric methods, in which leaves are excised and dried to determine water
24 loss. Although highly accurate, such approaches cannot provide continuous measurements and are
25 unsuitable for in situ monitoring in natural environments.

26 To address this limitation, recent studies have developed field-deployable sensors that
27 estimate plant hydration indirectly by measuring environmental parameters near the leaf surface,
28 such as relative humidity or vapor pressure deficit, using resistance-, capacitance-, or
29 voltage-based sensors¹⁻³. Even though these sensors provide a good estimate of tangential
30 hydration features such as relative humidity, they typically infer leaf hydration indirectly from
31 surrounding microclimate conditions rather than measuring the internal water status of the leaf
32 itself. In addition to the derived measurements, many of these systems utilize graphene oxide
33 (GO)^{1,4-6} or laser-induced graphene (LIG) films as the sensing material (Table S1)^{3,7}. Because
34 these sensors are intended for field deployment, minimizing interference with natural leaf
35 physiology is essential. However, many reported designs, particularly those based on GO or LIG
36 films, partially block light transmission and restrict gas exchange when attached to the leaf surface,
37 which may perturb plant physiology and limit their suitability for long-term monitoring. In
38 addition, most reported systems are configured to measure surface humidity rather than internal
39 leaf hydration, which can introduce delays or indirect responses relative to physiological water
40 dynamics.

41 Alternative non-contact approaches, including terahertz (THz) wave propagation and
42 quantum cascade laser techniques, have demonstrated the ability to probe internal leaf hydration
43 and pressure–volume relationships^{8,9}. Although these optical methods provide valuable
44 mechanistic insights, they typically rely on bulky and costly instrumentation and are therefore less
45 practical for continuous in-field monitoring.

46 The foliar graphene sensor (FGS) developed in this work addresses these challenges by
47 directly probing leaf hydration through four-probe bioimpedance measurements. In this
48 configuration, the excitation current passes through the leaf tissue itself, enabling interrogation of
49 internal hydration dynamics rather than relying on surface or environmental proxies. The
50 four-probe configuration minimizes the influence of contact resistance and parasitic impedance,
51 allowing accurate extraction of tissue impedance values^{10,11}. Because the measurement current
52 flows through the leaf tissue, the resulting impedance signal reflects the electrical properties of the
53 plant tissue and therefore provides a direct indicator of leaf hydration state. In addition, the
54 ultrathin and lightweight graphene-based sensing interface enables continuous operation for more
55 than two weeks without restricting leaf growth, outperforming many GO- and LIG-based
56 alternatives in both physiological compatibility and operational duration. In addition, the FGS
57 platform can be directly deployed under natural environmental conditions. Although outdoor
58 measurements exhibit slightly higher signal variability than controlled laboratory experiments, the
59 raw (unfiltered) data still clearly resolve variations in hydration states and diurnal physiological
60 dynamics (Figure 5e). Furthermore, optical techniques such as THz spectroscopy are typically

61 performed as discrete spot measurements rather than continuous monitoring. Compared with these
62 approaches, the system presented here provides a portable, low-cost, and scalable sensing platform
63 capable of real-time, long-term monitoring while maintaining minimal interference with plant
64 physiology. By combining direct tissue-level hydration sensing with a mechanically compliant and
65 wearable device architecture, the FGS platform enables practical and continuous monitoring of
66 plant water status in both controlled and field environments.

67 **Supplementary Note 2. Electrode Architecture and Mechanical Integration of the Foliar**

68 **Graphene Sensor (FGS)**

69 Establishing a mechanically compliant and non-perturbative interface between electronic
70 devices and living leaves presents a significant challenge due to the softness, continuous growth,
71 and dynamic motion of plant tissues. The electrode architecture of the FGS was therefore designed
72 to minimize mechanical stress, preserve natural leaf movement, and ensure stable electrical
73 contact.

74 The FGS employs bilayer graphene as the sensing electrode. Although graphene possesses
75 exceptional intrinsic mechanical strength, its atomic-scale thickness makes it mechanically fragile
76 and difficult to handle. To provide mechanical integrity and ease of handling, an ultrathin
77 polymethyl methacrylate (PMMA) support layer was incorporated (Figure S5). The resulting
78 graphene–PMMA laminate forms a conformal electrode that directly interfaces with the leaf
79 surface.

80 Electrical connection between the sensing electrode and the measurement instrumentation
81 was achieved using ultrathin gold tape fabricated by depositing a conductive gold layer onto a
82 flexible polymer substrate. Owing to its minimal thickness and high flexibility, the gold tape
83 exhibits mechanical compliance comparable to the graphene–PMMA laminate, thereby
84 minimizing stress concentration at the junction and preventing mechanical failure during leaf
85 movement.

86 Although gold provides excellent electrical conductivity, the extremely small thickness of
87 the gold tape leads to increases resistance over long connection lengths. To ensure reliable signal
88 transmission while preserving flexibility, silver epoxy was used to bond the terminal end of the
89 gold tape to an enameled copper wire, which then interfaced with the measurement instrument.

90 To minimize mechanical loading on the leaf, a 36 AWG enameled copper wire was
91 selected due to its ultralight weight (0.116 g/m) and small diameter (0.14 mm). Conventional
92 electrical leads and alligator clips were found to impose excessive weight, causing leaf bending
93 and restricting natural diurnal movements, including repositioning and orientation changes. The
94 lightweight wiring configuration enabled stable measurements while preserving natural leaf
95 motion and physiological function.

96 To quantify the mechanical load imposed by the sensing system, the total mass of the
97 complete setup was measured to be 0.2916 g, including four graphene sensors, four gold tape
98 interconnects, and four connecting wires. The combined mass of the four graphene sensors alone
99 was approximately 0.00035 g (Sartorius Entris II BCE124-1S analytical balance). For comparison,
100 a fully hydrated fifth leaf weighed 6.218 g, indicating that the sensing system accounts for
101 approximately 4.7% of the leaf mass. This minimal mass loading reduces gravitational bending
102 and mechanical perturbation, ensuring unobtrusive monitoring while preserving natural leaf
103 posture, movement, and physiological function.

104 **Supplementary Note 3. Biocompatibility and Physiological Compatibility of the Foliar**

105 **Graphene Sensor (FGS)**

106 Maintaining physiological compatibility is essential when interfacing electronic devices
107 with living leaves, as surface coverage may disrupt gas exchange, light transmission, or local
108 tissue function. The FGS was therefore designed to minimize interference with photosynthesis,
109 transpiration, and cellular integrity during prolonged attachment.

110 More than 20 maize plants were evaluated using the FGS. Across all experiments, no
111 visible chlorosis, lesions, or necrosis were observed in leaf regions covered with the FGS (Figure
112 S1), indicating that prolonged contact does not induce observable tissue stress or damage.

113 In contrast, mild chlorosis was observed in control regions covered with a non-breathable
114 polymer film. This response is consistent with restricted gas exchange and reduced light
115 transmission, highlighting the importance of breathable and optically transparent interfaces for
116 maintaining leaf physiological function.

117 The graphene sensing region is ultrathin, optically transparent, and permeable to gas
118 exchange. However, electrical interconnection requires ultrathin gold tape, which is not permeable.
119 To minimize potential physiological interference, the gold tape coverage area was carefully
120 limited during device assembly. The contact area was maintained below $4 \text{ mm} \times 12 \text{ mm}$ (< 0.5
121 cm^2), thereby minimizing obstruction of gas exchange and light exposure (Figure S1–S4).

122 After completion of measurements, the FGS was removed by gently wiping the surface
123 with a laboratory tissue (Kimwipes™, Kimtech) moistened with deionized (DI) water, followed by
124 careful peeling of the gold tape. Post-removal inspection revealed no visible differences between
125 previously covered and uncovered regions. The limited gold contact area produced no observable
126 physiological or morphological damage.

127 Together, these results demonstrate that the FGS provides a physiologically compatible
128 interface with living leaves, enabling prolonged monitoring without compromising tissue integrity
129 or function.

130 **Supplementary Note 4. Quantification of Leaf Color Difference (Figure 4a)**

131 To quantitatively assess potential color changes between FGS-covered and uncovered leaf
132 regions, digital image analysis was performed to provide an objective measure of visual
133 differences.

134 Regions of interest (ROIs) were selected using Photoshop (Adobe), excluding major veins
135 to avoid structural artifacts. Mean RGB intensity values were extracted for both covered and
136 uncovered regions.

137 Color difference was calculated as the Euclidean distance between the mean RGB vectors
138 in three-dimensional color space and normalized to the maximum possible distance within the
139 RGB color cube:

$$\Delta C = \frac{\sqrt{(R_1 - R_2)^2 + (G_1 - G_2)^2 + (B_1 - B_2)^2}}{\sqrt{3 \times 255^2}} \times 100\%$$

140 R,G,B represent the red, green, and blue intensity values of the selected regions.

141 This normalized metric provides a quantitative measure of color difference between the
142 two regions, enabling objective evaluation of potential chlorosis or visual tissue changes.

143 **Supplementary Note 5. Extraction of Impedance Values at Target Frequencies**

144 The raw bioimpedance data consist of frequency-resolved spectra spanning 10 Hz to 100
145 kHz, sampled at 201 logarithmically spaced frequencies. Each measurement file contains
146 impedance values corresponding to each excitation frequency.

147 For quantitative analysis and visualization, impedance values at specific target frequencies
148 were extracted from the full spectra. Single-frequency data were used to enable time-series
149 analysis, statistical comparison, and tracking of hydration dynamics.

150 Target-frequency values (e.g., 1 kHz) were obtained by filtering rows corresponding to the
151 desired excitation frequency and exporting the filtered data for subsequent analysis and plotting.

152 Data processing was performed using a custom Python script to ensure reproducibility and
153 consistency across datasets.

154 **Supplementary Note 6. Correlation Analysis Between Impedance and Environmental**

155 **Variables**

156 To evaluate the relationship between leaf impedance dynamics and environmental
157 conditions, correlation analysis was performed between normalized impedance changes ($\Delta Z/Z_0$ at
158 1 kHz) and measured environmental parameters(Figure S16).

159 The analyzed variables included light intensity, soil humidity, ambient temperature, soil
160 electrical conductivity (EC), and relative humidity. Raw datasets were converted to numerical
161 format and cleaned by removing missing values prior to analysis.

162 Pairwise correlations were quantified using the Pearson correlation coefficient to quantify
163 linear relationships between impedance and environmental variables. The resulting correlation
164 matrix was visualized as a heatmap to facilitate interpretation of multivariate relationships.

165 Correlation coefficients greater than 0.5 were considered indicative of strong positive
166 correlation, whereas values below -0.5 were considered strongly negative. Intermediate values
167 indicate moderate to weak associations.

168 All analyses and visualizations were performed using Python (pandas, NumPy, and
169 Seaborn) to ensure reproducibility.

Supplementary Note 7. Outdoor Environmental Validation

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171 Demonstrating sensor performance under natural environmental conditions is essential for
172 evaluating robustness beyond controlled laboratory settings. Because plants in field environments
173 experience wind-induced motion, fluctuating radiation, and heterogeneous microclimates, outdoor
174 measurements provide a stringent test of sensor stability and physiological relevance.

175 Outdoor impedance monitoring was conducted on a red oak (*Quercus buckleyi*) (Figure
176 S3), a native woody species, at an experimental agricultural field on the University of Texas at
177 Austin campus (Austin, TX, USA). Plants were exposed to ambient environmental conditions, and
178 impedance and environmental variables were recorded following the protocol described in the
179 main Methods.

180 Compared with controlled-environment measurements, increased signal variability was
181 observed, reflecting wind-induced motion, environmental fluctuations, and heterogeneous
182 radiation exposure typical of field conditions. Despite this variability, impedance dynamics
183 remained physiologically consistent with environmental drivers, demonstrating the feasibility of
184 outdoor monitoring and supporting cross-species deployment of the foliar graphene sensing
185 platform.

186 **Supplementary Note 8. Hydration Recovery Dynamics and Coupling with Soil Moisture**

187 Understanding how leaf hydration responds to soil water availability is essential for
188 interpreting plant water status and recovery dynamics. Because water transport occurs through an
189 integrated soil–plant hydraulic continuum, linking leaf impedance measurements to soil moisture
190 dynamics provides validation of the physiological relevance of the sensing platform.

191 Long-term in situ monitoring of maize leaves enabled quantitative characterization of plant
192 water uptake and recovery dynamics under varying hydration states. Following irrigation, leaf
193 impedance decreased rapidly, indicating immediate water uptake and hydraulic transport from
194 roots to leaf tissues. This initial response typically occurred within approximately 20 minutes,
195 reflecting rapid root–leaf hydraulic connectivity. Impedance then gradually stabilized as tissue
196 hydration approached equilibrium (Figure S11).

197 Hydration recovery times depended strongly on drought severity. In drought-stressed
198 plants exhibiting visible leaf curling, impedance remained elevated prior to irrigation and declined
199 slowly during recovery, requiring nearly 24 hours to return to baseline levels. This gradual decline
200 coincided with visible physiological recovery, including leaf unfolding and restoration of normal
201 diurnal movement patterns. Such delayed recovery is consistent with previous studies showing
202 that severe or prolonged drought slows restoration of leaf water status and gas exchange^{12–15}.

203 In contrast, well-watered plants re-equilibrated much more rapidly after irrigation.
204 Although minor impedance fluctuations occurred due to transient soil moisture variation and
205 internal redistribution of water, impedance stabilized at a fully hydrated state within
206 approximately 5 hours¹⁶ (Figure S11). Under adequate hydration, plants typically require only tens
207 of minutes to a few hours to re-establish water balance, consistent with established plant hydraulic
208 behavior. Recovery time, defined as the interval between irrigation and impedance return to
209 baseline, differed significantly between hydrated and drought-stressed plants, confirming that
210 drought severity strongly influences rehydration kinetics (Figure S12).

211 Despite differences in recovery timescales, both drought-stressed and well-watered plants
212 exhibited a rapid initial impedance decrease across frequencies following irrigation. In
213 drought-stressed plants, comparable responses at low and high frequencies suggest simultaneous
214 replenishment of extracellular and intracellular water compartments (Figure S10, S15). In
215 well-watered plants, changes were more pronounced at higher frequencies, consistent with rapid
216 intracellular hydration. Over subsequent hours, transient impedance fluctuations likely reflect
217 redistribution of water between extracellular and intracellular spaces, ultimately yielding a
218 uniform hydration profile.

219 Impedance dynamics also exhibited systematic relationships with soil humidity (Figure
220 S13). During the active recovery phase immediately following irrigation, transient deviations from
221 steady-state behavior were observed; therefore, recovery-phase data were excluded from curve
222 fitting to avoid bias. The remaining data were well described by a logistic relationship ($R^2 =$
223 0.835), indicating nonlinear coupling between soil moisture availability and leaf hydration state.

224 This behavior reflects the integrated hydraulic pathway linking soil water availability, root uptake,
225 and tissue hydration.

226 Together, these observations demonstrate that impedance monitoring captures hydration
227 recovery dynamics and their coupling with soil moisture in real time. The ability to detect early
228 dehydration and track recovery suggests a framework for irrigation management in which
229 watering is triggered before visible stress symptoms emerge, enabling faster recovery and
230 improved water-use efficiency (Figure S14).

Supplementary Note 9. Hydraulic Coupling Between Sap Transport and Leaf

Hydrodynamics

Linking leaf hydration dynamics to whole-plant water transport provides critical insight into how water is redistributed within plants. Because water movement occurs along a continuous soil–plant hydraulic pathway, simultaneous monitoring of sap transport and leaf impedance provides insight into the temporal coordination of water redistribution within the plant.

To quantify stem-level water transport, mini heat field deformation sensors (HFD) based on the design from previous research¹⁵ were used to measure relative sap flux (RSF) in small plants. These mini HFD sensors enable bidirectional sap flow measurements with high temporal resolution through continuous heating and thermocouple-based temperature gradients. The design and calculation of RSF are based on commercially available sensors^{17–20}. HFD sensors were designed in kiCad and printed on flexible PCBs (PCB gogo, Shenzhen, Guangdong, China). HFD thermocouples were connected to a datalogger (CR1000x, Campbell Scientific, Logan UT, USA), which recorded temperatures every 30 seconds and averaged them to 30-minute intervals. HFD sensors were placed on the outside of stems below the lowest branching junction. They contain a top, bottom and side thermocouple adjacent to a heating element (SMD 100 ohm resistor) that all make direct contact with the stem when attached. To convert RSF to absolute sap flow (g/h), excised stems were connected to a mass-balance hydraulic conductivity system¹⁵ while a mini HFD sensor was simultaneously applied to the stem to record RSF.

Multi-day averaged diurnal profiles revealed that sap flow is primarily driven by sunlight intensity, exhibiting strong alignment with daily radiation patterns (Figure S17). This behavior reflects transpiration-driven water transport regulated by stomatal conductance and atmospheric demand.

Following irrigation events, sap flow dynamics exhibited a delayed response. Unlike soil moisture, which increases immediately after watering, sap flow peaks typically increased on the following day (Figure S18). This temporal offset indicates that sap transport responds after internal water redistribution and adjustment of transpiration demand rather than directly to soil water availability.

The direct correlation between sap flow and leaf impedance yielded a Pearson correlation coefficient of 0.289 ($p = 3.83 \times 10^{-50}$). Although the correlation strength is modest ($r \approx 0.3$), this is expected because sap flux is governed by multiple interacting environmental and physiological factors, including radiation, vapor pressure deficit, temperature, wind, internal hydration status, and hormonal regulation. Given this complexity, it is unlikely that any single variable fully explains sap flow variance. Thus, the observed correlation strength is reasonable, and the high statistical significance (p-value) indicates a statistically significant relationship despite apparent variability.

Lagged correlation analysis further revealed a temporal offset between sap transport and leaf hydration dynamics. The correlation peaked at approximately 4.5 hours when sap flow was shifted forward relative to impedance, indicating that increases in sap flow precede decreases in

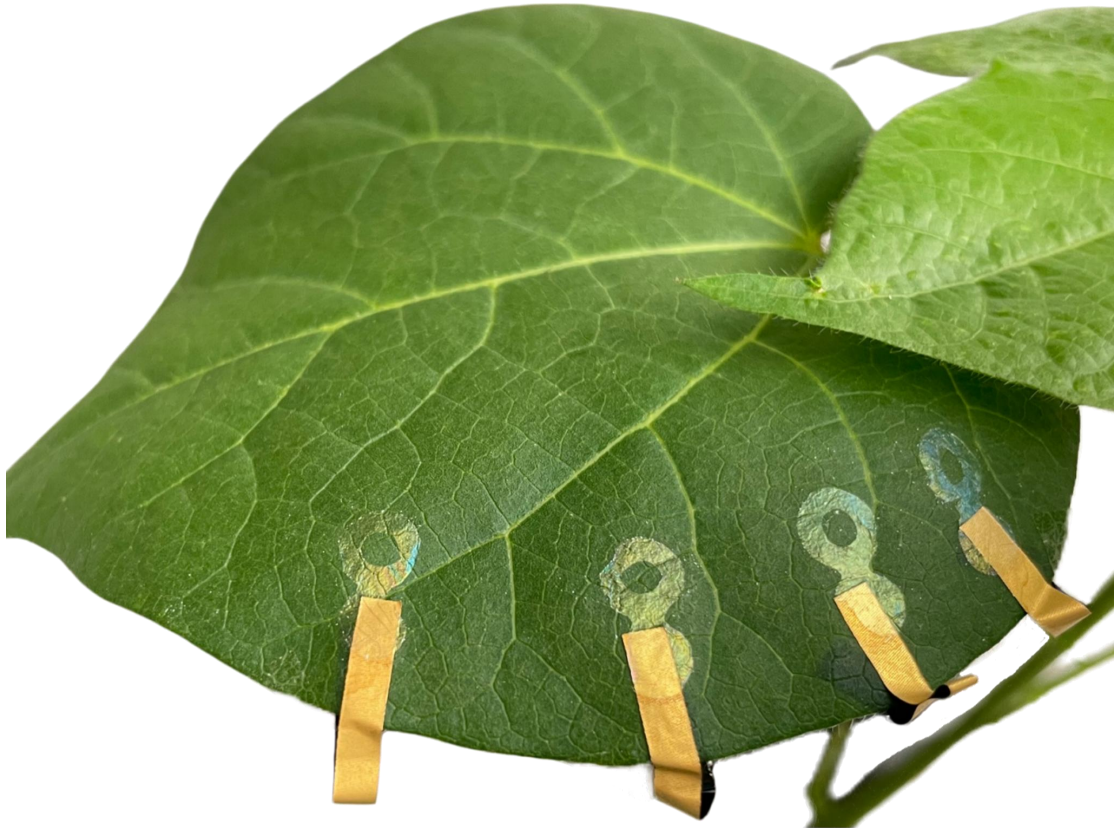
270 leaf impedance (Figure S19). This delay is consistent with the time required for water transported
271 through the xylem to redistribute into leaf tissues and restore cellular hydration. Multiple lag
272 processes coexist in plant water transport. Sap flux is known to lag environmental drivers such as
273 radiation and vapor pressure deficit^{21,22}, while leaf hydration reflects a later stage of the hydraulic
274 pathway. The observed delay between sap transport and impedance therefore supports the
275 interpretation that impedance measurements capture tissue-level hydration dynamics rather than
276 immediate hydraulic flow.

277 Together, these observations demonstrate that leaf impedance dynamics are hydraulically
278 coupled to stem sap transport and reflect downstream redistribution of water within plant tissues.



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280 **Figure S1. FGS conformally attached to a corn leaf.** Photograph showing four FGS
281 conformally attached to the adaxial side (top) of a single corn leaf, forming the sensing interface.
282 Gold tape serves as electrical interconnects, and external wires (routed on the backside and not
283 visible) complete the measurement setup.



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Figure S2. FGS attached on a cotton leaf. The sensing configuration shown in Figure S1 is applied to a cotton leaf. Four FGS are attached to the leaf surface and connected via gold adhesive tape to external wiring on the reverse side.



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289 **Figure S3. FGS attached to a red oak (*Quercus buckleyi*) leaf under natural outdoor**

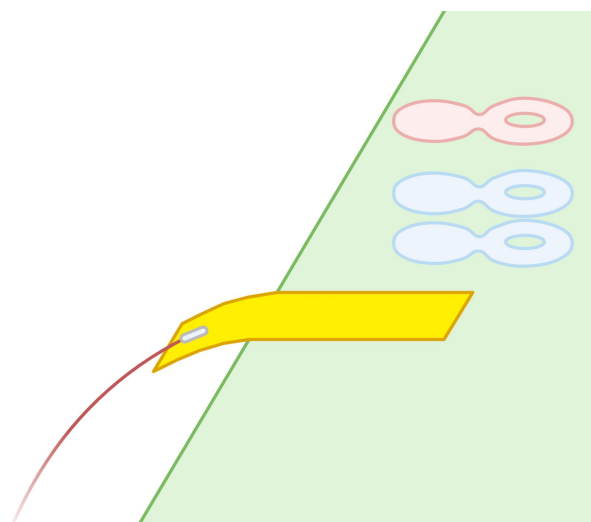
290 **conditions.** Four FGS are conformally attached to the leaf surface and connected to external

291 wiring using gold tape, following the sensing configuration shown in Figure S1.



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293 **Figure S4. A single FGS electrode and its electrical interconnect.** Photograph showing a single
294 FGS unit and its gold tape interconnect. The left region of the graphene film contacts the leaf
295 surface, while the right region overlaps the gold tape to form the electrical interconnect.



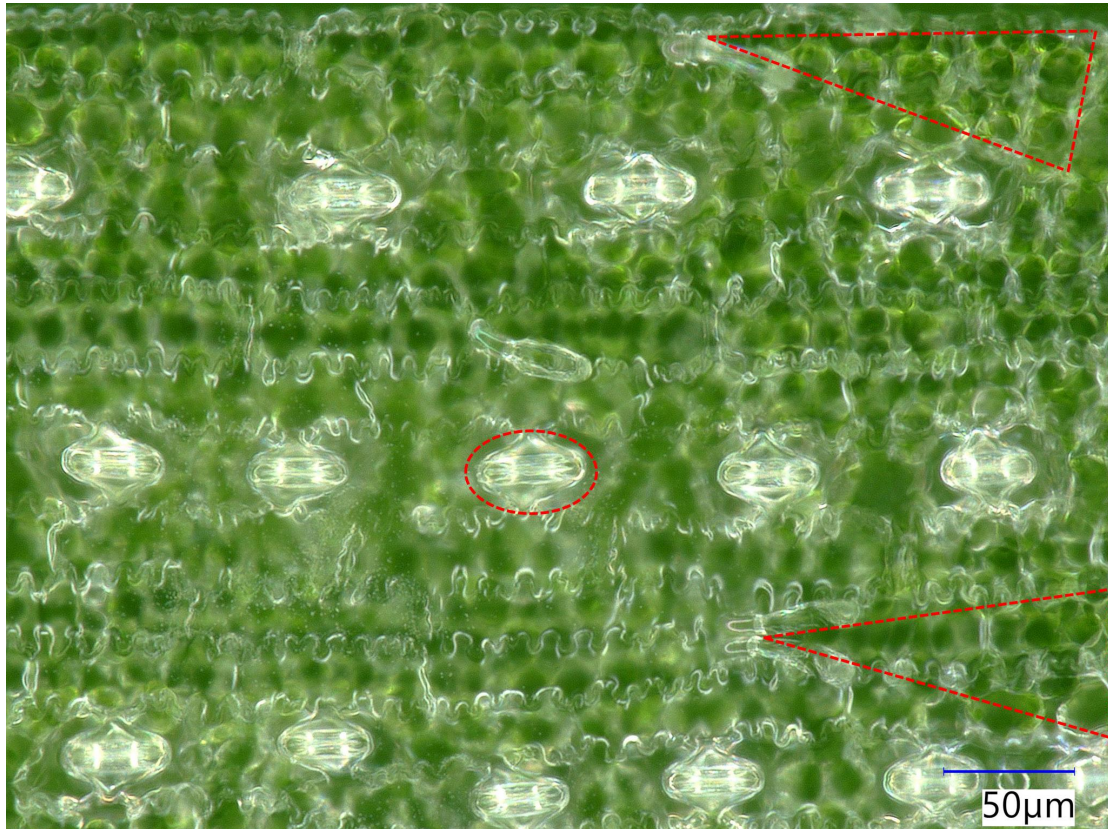
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297 **Figure S5. FGS architecture and measurement configuration.** Schematic illustration of the

298 sensing structure and measurement setup. Pink represents the PMMA supporting layer; blue

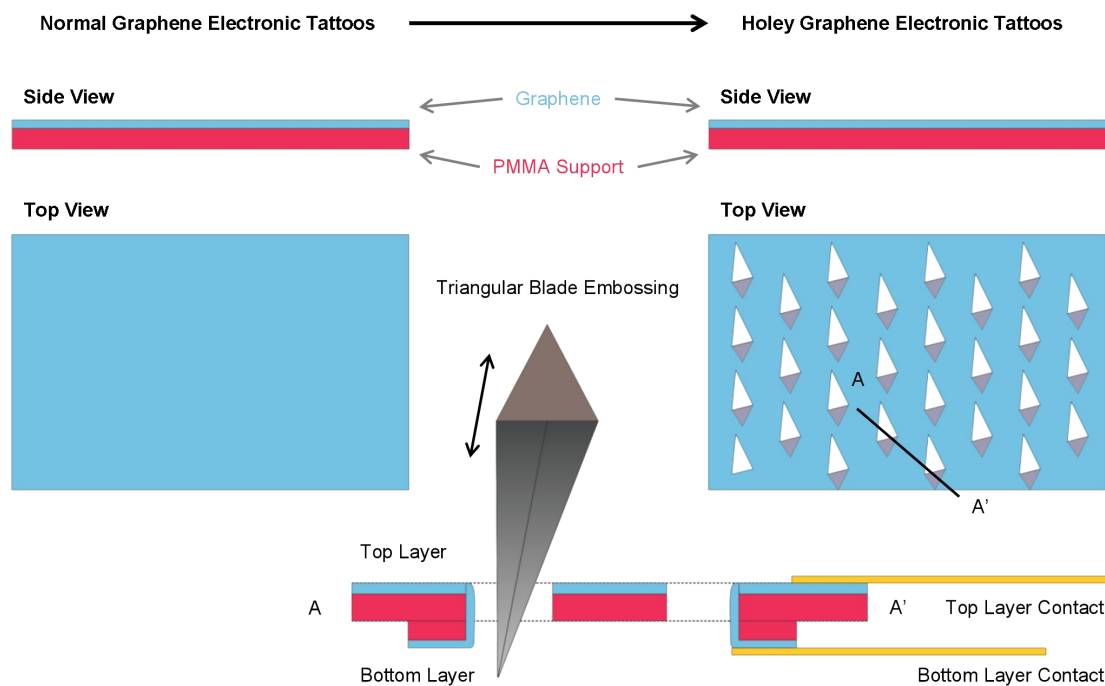
299 indicates monolayer graphene; gold corresponds to gold tape; silver denotes silver epoxy; red lines

300 represent enameled copper wires; and green indicates the leaf surface.



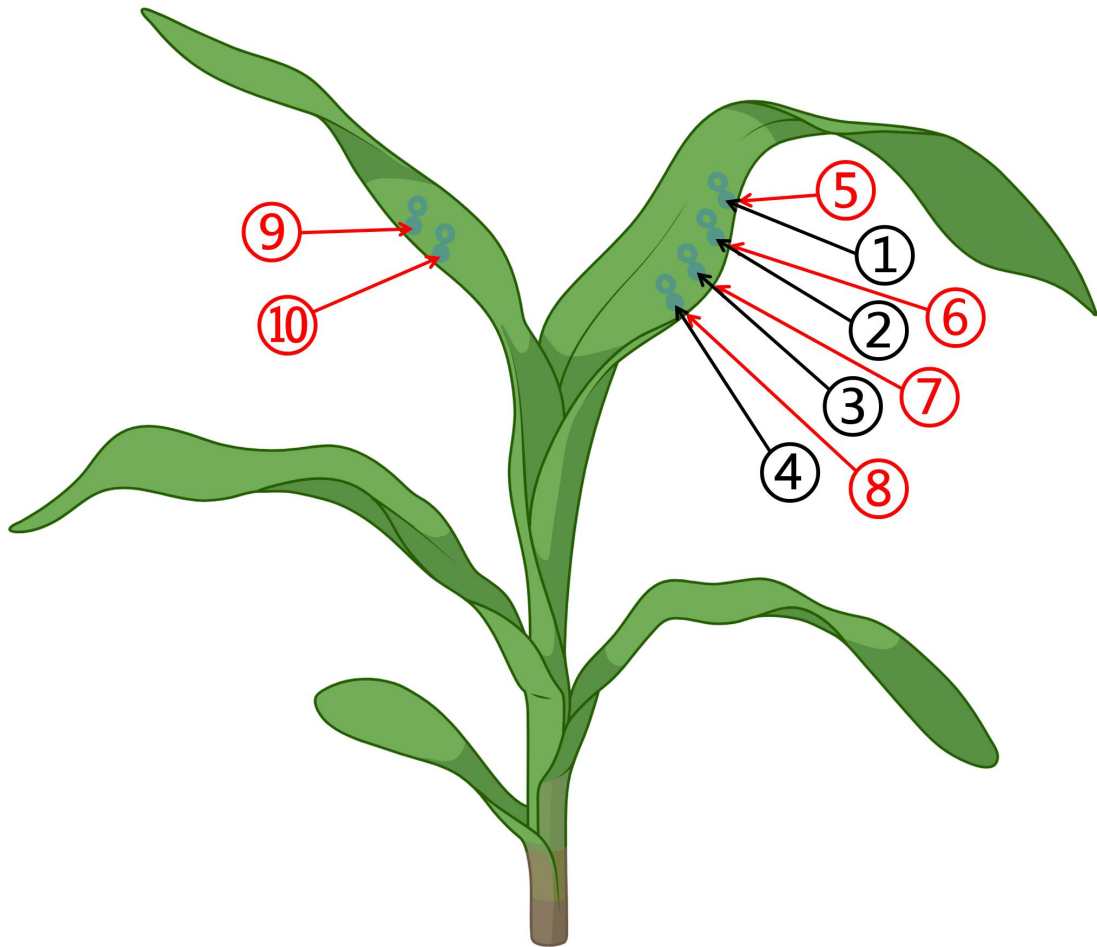
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Figure S6. Microporous structure of the FGS. Optical micrograph (VHX-X1, Keyence) of an FGS attached to the leaf surface. Triangular micropores in the graphene are outlined with red dashed lines, and a native stomatal pore of the leaf is marked with a red dashed ellipse. Although the graphene micropores do not always align perfectly with the stomata (as shown here), their high density ensures sufficient proximity to allow gas molecules to diffuse through the interfacial gap between the graphene and the leaf surface and enter the micropores, thereby maintaining gas exchange.



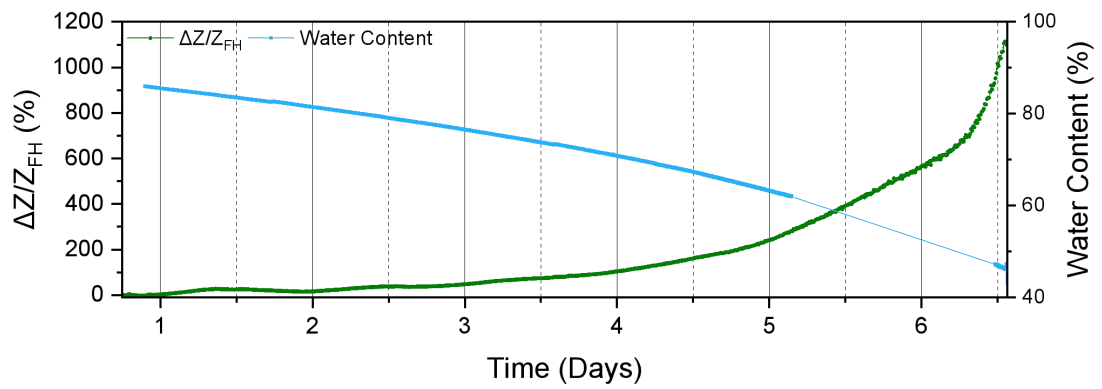
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310 **Figure S7. Formation of triangular micropores on graphene via embossing.** The
 311 graphene/PMMA is embossed using a Silhouette Cameo cutter. Because the process involves
 312 embossing rather than cutting, the pressed regions adhere to the back side of the film, partially
 313 delaminating the graphene and exposing a larger graphene area on the PMMA side. This creates
 314 triangular micropores on the graphene surface and enables electrical access to the FGS not only
 315 from the top (graphene) surface but also from the bottom (PMMA) side. The detailed fabrication
 316 process and electrical characterization of this method have been reported previously²³.



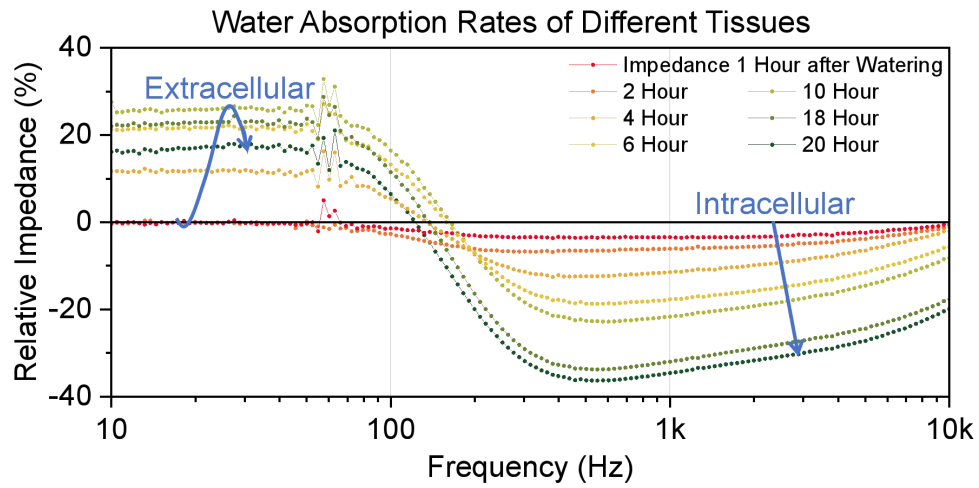
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318 **Figure S8. Placement of FGS on corn leaves.** Black arrows and numbers indicate sensors
319 positioned on the adaxial leaf surface, while red arrows and numbers indicate sensors on the
320 abaxial surface. Sensors 1–4 are placed on the adaxial surface (top), and sensors 5–8 are placed on
321 the abaxial surface (bottom). Sensors 2, 3, 6, and 7 placed on opposite surfaces of the same leaf
322 (across), whereas sensors 6, 7, 9, and 10 are placed on different leaves to measure (between).

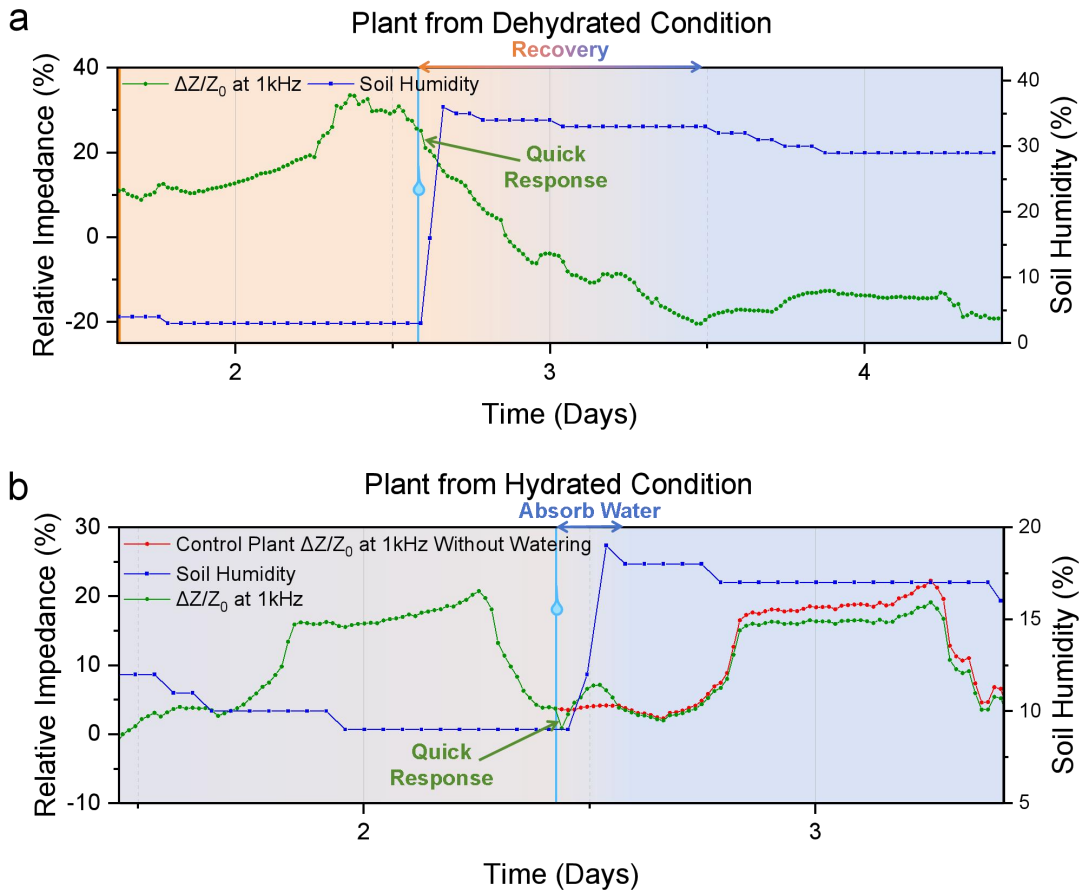


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324 **Figure S9. Correlation between LWC and impedance.** A detached corn leaf was continuously
 325 monitored for impedance and weight loss over time. LWC was determined gravimetrically from
 326 the fresh and dry weight. Solid vertical lines indicate midnight, dash vertical lines indicate noon.

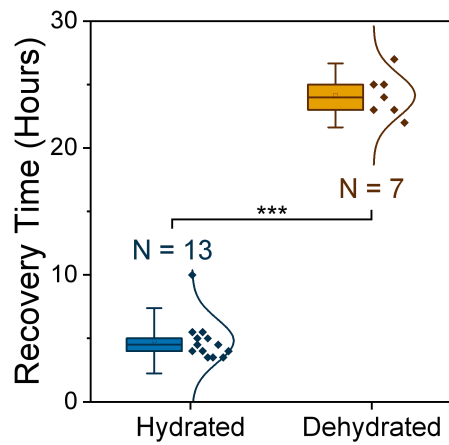


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 328 **Figure S10. Frequency-dependent impedance dynamics during water uptake.** Within 20
 329 hours following irrigation, relative impedance ($\Delta Z/Z_0$) changes vary across frequencies, indicating
 330 differential water uptake dynamics among leaf extracellular and intracellular compartments.



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Figure S11. Long-term dynamic monitoring of leaf impedance under different hydration states. **a**, Close-up view of Figure 2c showing the 24 h preceding and 48 h following an irrigation event. A rapid impedance decrease occurs immediately after irrigation, followed by approximately one day of recovery. The water drop icon indicates the irrigation event, and orange vertical lines denote visually observable water stress (leaf curling). Semi-transparent orange and blue shading represent periods of water stress and full hydration, respectively. **b**, Leaf impedance and soil humidity under hydrated conditions. Water absorption persists for approximately 5 hours following irrigation. The red curve represents a non-irrigated control plant. Because the two datasets originate from different plants, baseline impedance values differ. For visual comparison, a constant offset was applied to the control curve beginning at the irrigation time to align baselines; this adjustment does not affect the temporal response dynamics.



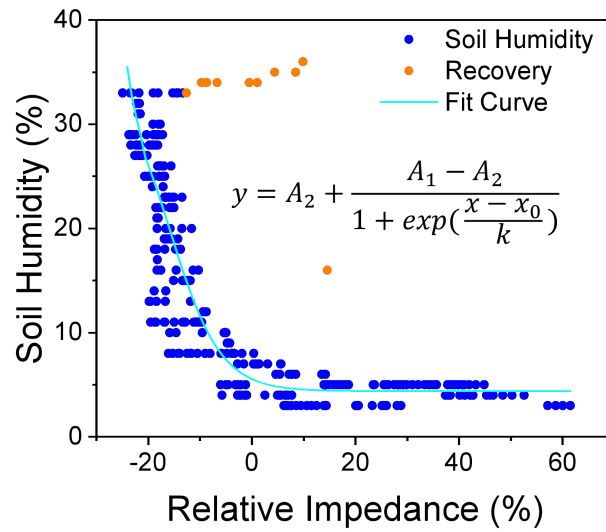
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344 **Figure S12. Comparison of recovery times under hydrated and drought-stressed conditions.**

345 Recovery time was defined as the interval between irrigation and impedance return to baseline.

346 Hydrated plants (N = 13) recovered significantly faster than drought-stressed plants (N = 7). The

347 difference in recovery time was statistically significant ($p = 5.05 \times 10^{-12}$).



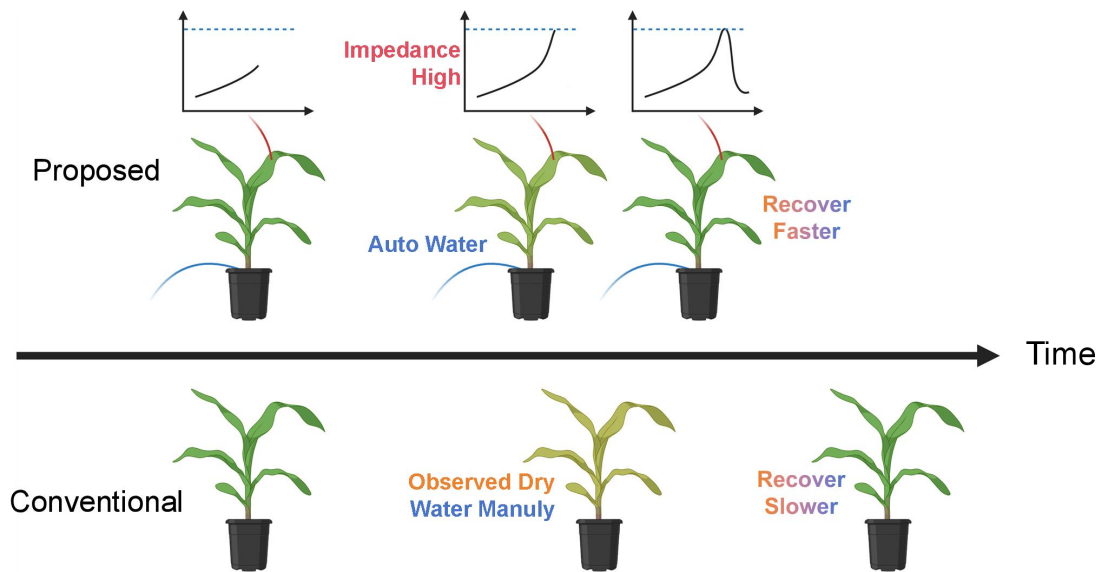
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349 **Figure S13. Relationship between leaf impedance and soil humidity.** Orange points indicate
 350 the active recovery phase during water uptake, while blue points represent steady-state conditions.

351 Recovery-phase data were excluded from curve fitting to avoid bias from transient dynamics. A

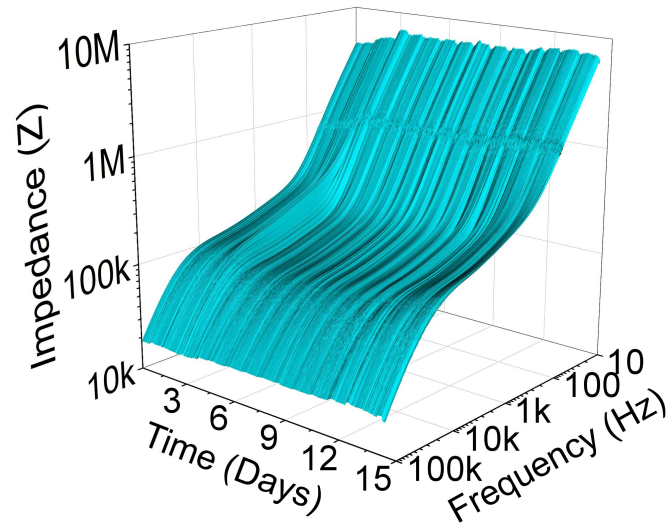
352 logistic fit captures the nonlinear relationship between soil moisture and leaf hydration state.

353 Logistic model parameters: $A_1 = 4.388$, $A_2 = 35.335$, $x_0 = -15.832$, $k = -4.879$.



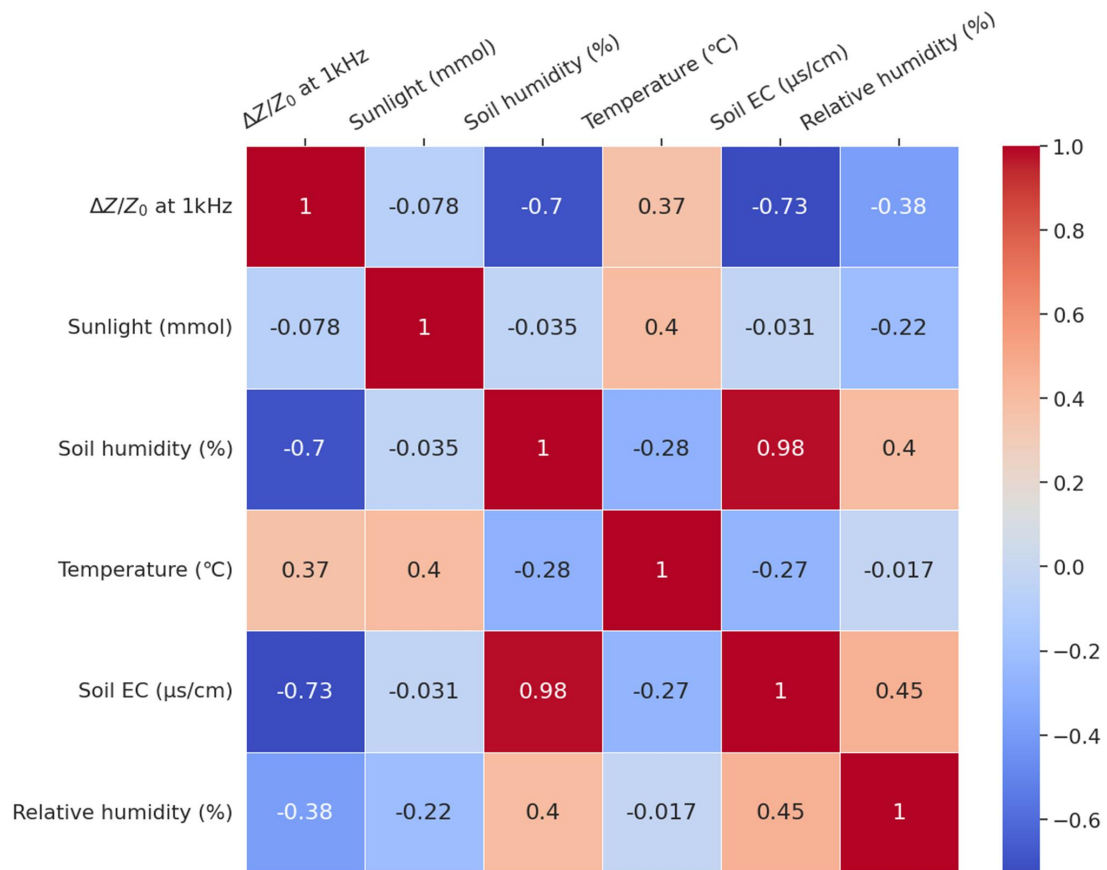
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355 **Figure S14. Conceptual illustration of irrigation management enabled by impedance**
 356 **monitoring.** In the upper diagram, crops are equipped with FGS linked to an automated irrigation
 357 system. When impedance rises beyond a threshold, indicating early signs of dehydration, water is
 358 delivered preemptively, allowing for faster recovery. In contrast, the lower diagram shows
 359 conventional practice in which irrigation is performed after visible leaf curling, indicating more
 360 severe dehydration and resulting in slower recovery.



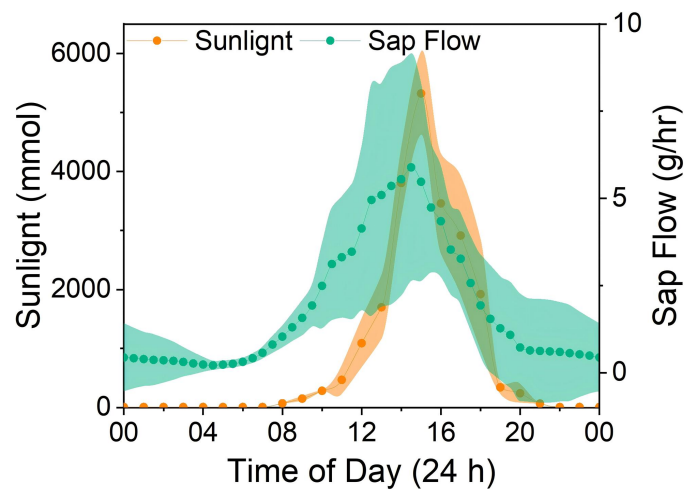
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362 **Figure S15. Continuous impedance monitoring across the large frequency range (10 Hz–100**
363 **kHz) over 14 days.**



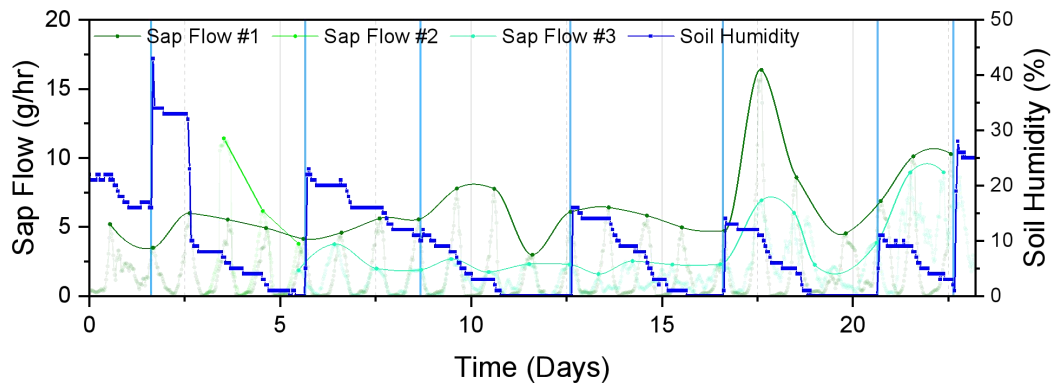
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365 **Figure S16. Correlation heatmap between leaf impedance and environmental parameters.**
 366 Pearson correlation coefficients quantify linear associations among impedance ($\Delta Z/Z_0$ at 1 kHz),
 367 sunlight intensity, soil humidity, temperature, soil electrical conductivity (EC), and relative
 368 humidity. Coefficients greater than 0.5 are indicative of strong positive relationships, whereas
 369 values below -0.5 are indicative of strong negative relationships.



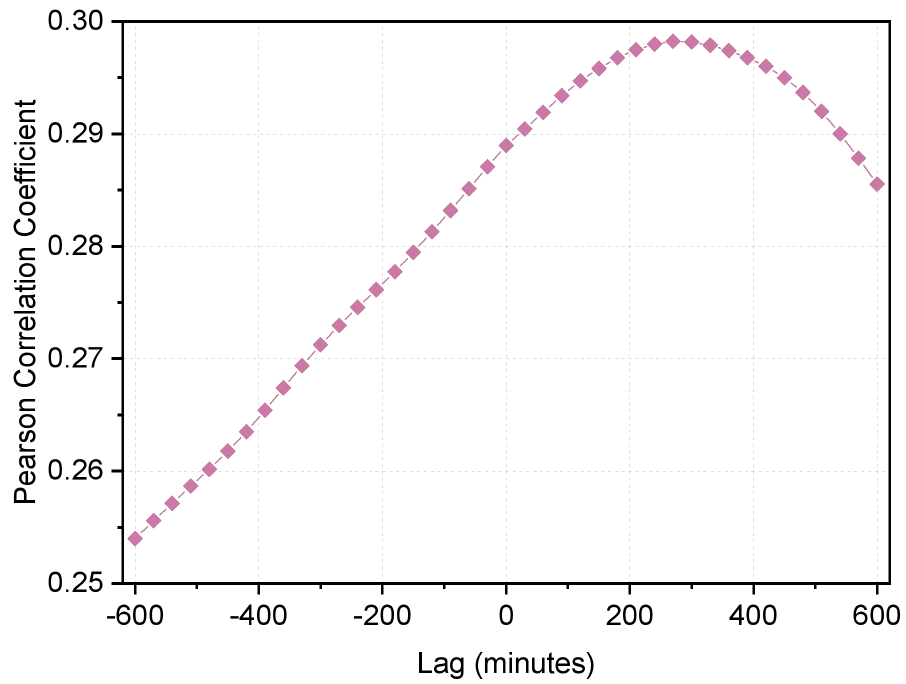
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371 **Figure S17. Diurnal patterns of sunlight intensity and sap flow.** Twenty-day averaged 24-hour
 372 profiles show strong alignment between sap flow and sunlight intensity. Solid lines represent
 373 means, and semitransparent regions represent \pm standard deviation (SD).



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375 **Figure S18. Sap flow response to irrigation events.** Semi-transparent green curves show
 376 real-time sap flow, while solid green lines indicate daily peak values. Light blue vertical lines
 377 denote irrigation timing. Sap flow increases typically occur on the following day, indicating
 378 delayed hydraulic adjustment.



379
380 **Figure S19. Lagged correlation between sap flow and leaf impedance.** Pearson correlation
381 coefficients (r) between sap flow and impedance are plotted for lag times from -600 to +600
382 minutes. Positive lag values indicate sap flow shifted forward in time. The peak correlation at ~ 4.5
383 hours indicates that increases in sap flow precede decreases in impedance.

384 **Table S1. Examples of emerging plant hydration monitoring sensors.**

<i>Study Reference</i>	<i>Measured Quantity</i>	<i>Measurement Method</i>	<i>Sensor Materials</i>	<i>Sensor Thickness</i>	<i>Sensor Weight</i>	<i>Invasiveness</i>	<i>Continuous Monitoring Duration</i>
<i>This Work</i>	Intra-/Extracellular LWC	4-Probe Impedance	2 Layer Graphene	< 300 nm	< 0.1 mg	Non-Invasive	16 Days
11	LWC	4-Probe Impedance	Aluminum Foil	--	--	Invasive	2 Days
7	Leaf Surface RH	DC Resistance	LIG	--	--	Invasive	16 Days
24	Leaf Surface RH	Capacitance	LIG and GO	--	--	Invasive	11 Days
25	Leaf Electrical Potential	Capacitance	Polymer + CNT	50 um	--	Non-Invasive	1 Month
5	Leaf Surface PWC and RH	2-Probe Impedance	GO	--	--	Invasive	4000 Seconds
6	Leaf Surface RH	DC Resistance	GO	--	--	Invasive	Yes
30	Leaf Hydration	Conductance	Graphene	< 0.3 um	9 mg	Non-Invasive	8 Days
26	Leaf Surface RH	Voltage	GO	10 um	--	Invasive	No
9	LWC	THz Waves	--	--	--	Non-Invasive	--
8	LWC	THz Waves	--	--	--	Non-Invasive	--
27	Sap Flow Rate	Temperature	Cu/Pt	--	240 mg	Non-Invasive	16 Days
28	Leaf Surface Water Content	2-Probe Impedance	Cu/SWCNT	30 um	17 mg	Non-Invasive	1 Day
29	Electrophysiology	2-Probe Impedance	EPC Thermogel	1 mm	--	Non-Invasive	--

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Supplementary Methods: Statistical Analysis

387 Statistical analyses were performed in Python using SciPy and statsmodels. Sample sizes
388 (N) refer to independent biological replicates and are reported in figure legends.

389 Statistical comparisons between two independent groups were performed using Welch's
390 two-tailed t-test, which does not assume equal variances. Statistical significance was defined at α
391 = 0.05. p-values ≥ 0.001 are reported exactly; values < 0.001 are reported as $p < 0.001$. Graphical
392 annotations use the following convention: * ($p < 0.05$), ** ($p < 0.01$), *** ($p < 0.001$), and n.s.
393 (not significant) for $p \geq 0.05$.

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