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Growing Phenotype-controlled Phononic Materials from Plant Cells Scaffolds

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Biological composites offer self-healing properties, biocompatibility, high responsivity to external stimuli, and multifunctionality, due, in part, to their complex, hierarchical microstructure. Such materials can be inexpensively grown, and self-assembled from the bottom up, enabling democratized, sustainable manufacturing routes for micro- and nano-devices. While biological composites have been shown to incorporate rich photonic structures, their phononic properties have hitherto remained unexplored. In this study, we demonstrate that biological composites in the form of micron-thick decellularized onion cell scaffolds behave as an organic phononic material, with the presence of band gaps forbidding the propagation of elastic waves in select frequency ranges. We show that the onion cells’ phononic properties can be phenotypically tuned, and anticipate these findings will yield new biologically-derived, “green,” and genetically tailorable phononic materials.

From a sustainability perspective, it is critical that existing technologies incorporating inorganic compounds, such as rare earth metals, be replaced with biodegradable and recyclable organic materials.1 Because of this, the development of biological composites has become a societal necessity.2,3 As an added value, biological composites also offer self-healing properties,4 biocompatibility,5 high responsivity to external stimuli for applications such as sensing and soft robotics,6,7 and multifunctionality (for instance, as is exemplified by butterfly wings having both structural coloration8 and superhydrophobicity9). In contrast to inorganic systems, which are typically fabricated using non-scalable top-down approaches (where the fabrication time scales cubically with the system to microstructure size ratio), biological composites exhibiting complex structural hierarchy can be grown, and scalably self-assembled from the bottom up.10,11 Such naturally-occurring structural hierarchy makes biological composites a class of uniquely transformative materials for bridging functional nanocomposites to macroscale integrated devices in a green manner.12

Within the context of biological composites, a wide array of systems with rich functionalities have been implemented, such as cellulose-based flexible electronics,13 plasmonic wood,14 wool-based nano-patterns,15 lasers,16 and DNA-based mechanical metamaterials.17 Yet harnessing the power of biological composites to control elastic waves has remained an unmet challenge, partly because, in contrast with visible photons, there has been no observation of phonon-based biological function at the supramolecular scale. In the past decade, the use
of inorganic, man-made composites, often referred to as phononic materials, have enabled manipulation of elastic waves across frequencies ranging from tens of Hz to a few THz, with applications ranging from seismology to thermal transport, respectively. As a result of their structural features, inorganic phononic materials have been shown to exhibit unique properties such as negative Young’s modulus, mass density, and refraction. Integration of these materials in ultrasonic biosensors has provided sensitivity enhancements and enabled diagnostic applications. Miniaturized lab-on-a-chip devices for fluid shaping, nebulization, tweezing, and streaming and sorting have all shown benefit from the introduction of phononic microstructures. The use of phononic materials in ultrasonic signal processing, such as the SAW filters which are ubiquitous in modern communication devices, has been shown to reduce device size and enable functionalities such as wave guiding and logic.

However, these inorganic phononic materials operating at high frequencies (MHz and above) are subject to several major limitations, namely manufacturing scalability, fragility, sustainability, and biocompatibility. Leveraging the ability of biological composites to be grown rapidly, inexpensively, en masse from renewable resources would allow building biodegradable and recyclable phononic materials, and facilitate the democratization of state-of-the-art technologies in the areas of sensing, microfluidic control, acoustic signal processing, and potentially even thermal control materials. Moreover, in contrast to their inorganic counterparts, biological materials are soft, stretchable, and inherently biocompatible, making them uniquely suited to translate the above-mentioned applications into wearable sensors, health-care devices, and human augmentation devices. We also envision these biocompatible phononic materials could be used in ultrasonic biomedical imaging and wave-assisted regenerative medicine applications as integrated signal focusing and localization devices or acoustically-encoded reporters. In this paper, we demonstrate that biological composites in the form of decellularized plant cell scaffolds can behave as phononic materials, including forbidding the propagation of elastic waves in select frequency ranges (i.e. band gaps). Our discovery of phononic behavior in biological composites, in particular, plants, shows how biology can be directly used to transfer the existing physics of phononic materials into scalably manufacturable, biocompatible, sustainable, and democratizable forms.

Our biological composite is composed of a micron-thick onion cell epidermis with slender
cell walls extruding up from it that resemble blind bore beehive structures, and is adhered to a glass substrate. We measure the dispersion of sub-GHz surface acoustic waves (SAWs) in the onion composite, and reveal their interaction with compressional and flexural resonances of the wall structure, which open deep and wide band gaps. We also demonstrate that the SAW dispersion can be phenotypically controlled by selecting plants at different developmental stages. We thus foresee that, in the long-term, synthetic biology, whereby artificial devices are built from biological parts (cells, DNA strands...), and the ability to tailor the genome by controlled mutations or gene editing, could provide a plausible, scalable manufacturing route for future phononic materials design. This is a clear advantage compared over previously studied biological systems that have limited potential for genetic tailoring and mass production.

Fresh onion scales were selected from different depths of an onion bulb, as is shown in Fig. 1a. The outer half of selected layers was peeled off and dehydrated over 1-2 days at 4°C prior to testing (Supplementary note 1). Figure 1b shows optical microscope images of the peeled organic layers, which show different phenotypes corresponding to different developmental stages (indexed as a function of consecutive layer number, starting from the outermost epidermis). The scales were placed onto a photoacoustic transducer composed of a 150 nm layer of gold coated onto a thick 1 mm glass substrate, as is illustrated in Fig. 1c. The remaining cell walls, constituting the bottom of the onion scale and the vertically protruding walls, resemble a borehole beehive structure. We show a 3D profilometry image of a representative sample surface in Fig. 1d, as well as typical line profiles recorded along the minor and major axes of the cavities in Fig. 1e (which corresponds to the sample shown in Fig. 1d). The surface topography of the decellularized epidermal layers reveals that the vertical portions of the cell walls decrease in size (height and full width at half maximum) as a function of the age of the layer, ranging from about 16 μm × 30 μm for the oldest layer to 3 μm ×10 μm for the youngest layer, as is shown in Fig. 1f. To analyze the sub-GHz acoustic properties of this organic structure, we illuminate the bottom of the metal film with a pulsed laser (400 ps pulse duration, 532 nm wavelength, and 1.0 mW average power) focused to a ~10 μm diameter spot (Fig. 1c and Supplementary note 2). The subsequent rapid thermoelastic expansion in the metal generates propagating SAWs with a broad frequency spectrum extending up to 400 MHz. We use the time-dependent deflection of a continuous probe beam (577 nm wavelength, 12 mW average...
FIG. 1: (a) Sample preparation. Epidermal layers were obtained from different depths of an onion bulb, going from the outermost (oldest) to the innermost scale (youngest), shown in the inset. (b) Optical microscope images of the peeled organic layers showing different morphogenetic profiles as a function of their developmental stage. Scale bar is 100 μm. The diagrams show profiles of the pillar-like structure of the wall as a function of the developmental stage (index denotes layer number). (c) Opto-acoustic setup used to generate and detect sub-GHz SAWs in the sample. (d) Profilometry image of the onion epidermal surface with a superimposed illustration of the propagating SAWs. (e) Representative profiles along the minor and major axis of the cells, respectively. (f) Variation of the height and width of the wall’s vertical portions as a function of the developmental stage of the plant. (g) Snapshots at different times showing the propagation of SAWs on a bare substrate. Scale bar is 100 μm.

power) by the acoustic-induced surface ripples (sketched in Fig. 1d) to monitor the propagation of the SAWs (Supplementary note 2). By scanning the probe beam across the bottom of the metal film over a 300 × 300 μm² area, we obtain movies of the propagating SAWs. These animated maps reveal largely circular wavefronts, as is shown in Fig. 1g, which depicts SAWs propagating along a substrate without a cell layer on top (hereafter referred to as “bare”).
FIG. 2: (a) Snapshots at $t = 40$ ns, filtered to display wave propagation in limited frequency ranges. Scale bar is 100 $\mu$m. (b) Acoustic power calculated at each frequency for the bare (black) and for the onion epidermis (red). Dispersion curve measured on (c) the bare substrate and (d) the organic layer in the $-136^\circ$ direction. The black lines in (c) correspond to the dispersion of surface and longitudinal wave speeds calculated using a layered-half-space model. The black dashed line corresponds to the transverse wave speed in the bare substrate. (e) Dispersion curve obtained from the numerical analysis using the wall dimensions of the onion layer with index number 8. The dispersion curve was computed by evaluating the spatio-temporal diagram at the gold-glass interface. The horizontal red and green lines are the compressional and flexural resonant modes of the wall respectively. The mode shapes of the resonant modes are illustrated on top of dispersion curve. The figure on the right side of the dispersion curve shows the frequency response calculated by summing the $k_r$-components of the dispersion maps. The red region shows the phononic band gap evaluated at 70% of the maximum amplitude.

During its propagation, the SAW sets the onion epidermis atop in motion, and this coupled movement alters the SAW dispersion. To show this, we plot snapshots of the traveling SAWs in Fig. 2a, filtered to display wave propagation in three limited frequency ranges (Supplementary note 3). We observe very low acoustic amplitude $u_f(x, y)$ in the 160-180 MHz range, compared to the other two frequency ranges. Such frequency
dependence is not present for SAWs propagating across the bare substrate. In Fig. 2b, we plot the acoustic power \( P_f = \sum_{x,y} |u_f(x,y)|^2 \) calculated at each frequency \( f_i \), for the bare substrate (black) and for the onion epidermis characterized in Fig. 2a (dashed red line) at a time \( t \sim 50 \text{ ns} \). This plot indicates the presence of a phonon stop band, or resonant attenuation zone, in the 160-180 MHz range with 94% extinction compared to the bare substrate caused by the presence of the organic layer.

To better understand this observation, we calculate the acoustic dispersion curves \( |\tilde{u}(k_r, \theta, f)| \) using a 2D Fast Fourier Transform (FFT) in the radial wavevector-frequency domain \( k_r-f \) for each angular direction \( \theta \) (see Fig. 1g and Supplementary note 3). Figure 2c shows a dispersion curve measured on the bare substrate for one wavevector direction. We observe two distinct modes, corresponding to surface skimming longitudinal and Rayleigh SAW modes. We calculate analytical dispersion curves using a slow-on-fast layered-half-space model for the bare substrate (dotted red lines in Fig. 2c),\(^{42}\) with typical acoustic properties for silica (\( v^s_L = 5700 \text{ m/s}, v^s_T = 3400 \text{ m/s}, \text{ and } \rho^s = 2500 \text{ kg/m}^3 \))\(^{43}\) and gold (\( v^g_L = 3200 \text{ m/s}, v^g_T = 1200 \text{ m/s}, \text{ and } \rho^g = 19320 \text{ kg/m}^3 \)),\(^{44}\) and find good agreement with the experimentally identified branches.

The dispersion curve (Fig. 2d) corresponding to a location containing an epidermal layer shows a starkly different behavior. We observe avoided crossing behavior due to the coupling of Rayleigh waves with local resonances in the organic layer, that opens a wide gap (likely aided by resonant attenuation effects) around 170 MHz. The lower branch starts as a Rayleigh wave at low-\( k_r \) values (red dotted line) and approaches a horizontal asymptote near the resonance frequency. The upper branch tends to the Rayleigh wave speed at high wavenumbers, but deviates from the Rayleigh line near the gap frequency and vanishes beyond the \( v^T_T \) threshold (white dotted line), below which it becomes evanescent.\(^{45}\) Such dispersion curves are characteristic of locally resonant metamaterials, and have previously been observed at the geophysics scale,\(^{46}\) wherein trees in a forest were used as local resonators, and down to the microscale via the interaction of SAWs with the contact resonances of microsphere monolayers.\(^{47}\)

By analogy with observations made on inorganic, man-made phononic metamaterials,\(^{48,49}\) we suspect the slender vertical portions of the wall may serve as the local resonators that couple to the SAWs. To verify this assumption, we conduct 2D finite element (FE) analysis using COMSOL Multiphysics. We model SAWs propagating on a gold-coated glass substrate by
applying a point-like load with a step-like excitation in the time domain (Supplementary note 4). To mimic the response of the organic layer, we add a uniform onion layer of 3 \( \mu m \) thick on top of the substrate with rectangular ridges extruding up from it, and equally spaced by the measured average walls’ periodicity (Supplementary note 1) as is illustrated in the schematic of Fig. 2e. The ridges have a height of 6 \( \mu m \) and a width of 16 \( \mu m \) (thus mimicking the onion layer with index number 8). The ridges and the onion layer underneath are given the effective elasticity of dry cell walls (Supplementary note 4). From our simulated spatiotemporal diagrams, we plot the dispersion curve of the SAWs using a 2D FFT, as is shown in Fig. 2e. We plot the frequency response averaged over all \( k \)-values on the right side of the dispersion curve. The red region denotes the width of the phononic band gap evaluated at 70% of the maximum amplitude. Our computational analysis reveals a wide gap in a good agreement with our observations.

In order to reveal the origin of the observed band gaps, we computed the first five eigen-modes of the ridges and plotted their mode shapes in Fig. 2e. We denoted the flexural and compressional modes by green and red lines, respectively. We observe that both the first compressional mode and the second flexural mode fall within the range of the gap. Although this observation suggests that the width of the gap is due to the interaction of the SAW with both compressional and flexural resonances, we note that SAWs have previously been observed to hybridize predominantly with compressional resonances\(^{46}\), which may suggest here that the second flexural resonance plays a marginal role in the formation of the gap. This can also explain the absence of hybridization with the first flexural mode at a lower frequency. We note that the periodicity of the wall structures (on the order of \( a = 50 \mu m \)) in the samples can open Bragg gaps at \( k \sim \pi/a \sim 0.06 \mu m^{-1} \), that is \( f \sim 30 \text{ MHz} \), however this is significantly below our acoustic measured frequency.

The structure of plants, which is one element of their phenotype, can be controlled by environmental or genetic cues. Such modifications, which have enabled crop optimization, are also at the core of plant synthetic biology.\(^{51}\) To illustrate the ability to control the structure of the organic resonators without any physical or chemical intervention, we select epidermises at different developmental stages. As the plant grows, the phenotype changes and the ridges become thicker and higher and further spaced, as observed in the profilometry images (Fig. 1f). In Fig. 3a, we plot the spatial average of the displacement amplitude \( \tilde{u}_{av}(f) = \sum_{k,r,a} \tilde{u}(k,r, \theta, f) \) calculated at each frequency, normalized by the response of the
FIG. 3: (a) Spatial average of the frequency response normalized by the response of the bare substrate for seven layers of increasing age. Red dots indicate the center of the gap, also reported in Fig. 3b (b) Variation of the center frequency of the gap as a function of the cell growth stage. The dashed line shows the variation of the first compressional mode (fixed-free organ-pipe mode) of a continuous onion pillar as a function of the normalized height. The red circles indicate the simulated frequency of the gap. (c) Variation of the resonator spacing with the developmental stage of the cells. (d) Simulated (red) and measured (blue) band gap width as a function of the developmental stage. As a guide to the eye, the simulated results are fitted to a linear curve (dashed line) (e) Snapshot at $t = 60$ ns taken from an acoustic movie filtered between 120-140 MHz, that shows directionality in the SAW propagation. As a guide to the eye, we indicate with a dashed line the major axis of the cells (at $\sim 40^\circ$). $f - \theta$ maps corresponding to (f) the bare substrate and (g) the onion epidermis.
bare substrate for seven layers of increasing age from the same onion bulb (Supplementary note 3). We observe a gap whose center frequency decreases with the age of the epidermis by up to a factor of two, from \(\sim 230 \text{ MHz} \) to \(\sim 120 \text{ MHz} \) (also plotted in Fig. 3b with blue squares).

Among the different parameters that can influence the gap tunability, let us first discuss the influence of the geometry of the cell wall (resonators) and consider a simple analytical description. Rayleigh waves, having an elliptical polarization, are known to hybridize strongly with compressional resonances.\(^{47}\) We thus estimate the frequency \(f = \frac{v_L^L}{4h} \) of the first out-of-plane compressional mode (organ-pipe mode) of a continuous onion pillar, where \(v_L^L = 3400 \text{ m/s} \) is the average longitudinal sound speed measured in onion cells at high frequencies\(^{52}\) and \(h \) is the height of the ridges measured by profilometry. We plot in Fig. 3b the frequency of this mode as a function of normalized ridge height and developmental stage (dotted red line). We observe a qualitative agreement in the frequency range and trend of the measured gaps as a function of ridge height, illustrating that the increase in the height of the vertical portions of the wall, as a result of tissue growth, correlates to a decrease in the gap frequency. Similarly, a decrease in the average stiffness of the epidermis could also result in a decrease in the gap frequency. The plant cell wall is mainly composed of highly oriented cellulose microfibrils cross-linked by hemicelluloses, embedded in a hydrated gel-like matrix of pectins.\(^{53-55}\) To regulate cell growth, new layers of cellulose are deposited with highly oriented microfibrils, which result in older cell walls being stiffer. We suggest that this change in elasticity may partially counteract the ridge-height-induced decrease in band gap frequency with increasing cell age.

We also quantify the width of the band gap (whose edges are defined as 70% of the average amplitude level outside of the gap) and plot it as a function of the developmental stage (indexed as the normalized height of the organic layers’s vertical portions), plotted as blue markers in Fig. 3d. We see that the gap becomes narrower as the organic layer ages. To investigate this, we consider the spatial density of the resonators. We estimate the averaged spacing \(\Lambda\) between the cells by counting the number of vertical wall portions along the minor axis of the cell on the profilometry images. In Fig. 3c, we plot \(1/\Lambda\) as a function of normalized ridge height. We see that \(1/\Lambda\) decreases with the age of the organic layers, i.e. the spatial density of the resonators decreases, which is consistent with the observed narrowing of the band gap with increased age. We also compared in Fig. 3d the observed
widths (blue squares) to the simulated ones (red circles) by accounting for the increased resonator spacing in our time-domain analysis. As a guide to the eye, we added a linear fit to the numerical results (dotted line). Our simulations show an increase in the gap width as a function of the normalized cell wall height, in agreement with our measurements. Our observations show that geometry, stiffness and number of resonators are key in adjusting the features—center frequency and width—of this phenotype-controlled band gap. These parameters of the biological composite could be manipulated by selecting the developmental stage (as we did), or by mutations or gene editing to tune phonon dispersion precisely, opening sustainable routes to engineer scalably manufacturable metamaterials from grown plants.

The sample has in-plane anisotropy due to the preferential orientation of the cells (which can be described by stripes aligned with the major axis of the cells), but also due to the anisotropy of the cell wall material itself. Both anisotropies are linked because the growth of oriented microfibrils contributes to cell elongation, and thus a transition from isotropic to orthotropic structure. In a perfect stripped system such as the one we modelled above, the gap should completely close along the stripes. In most of the samples we discussed, we did not observe any strong dependence of the center frequency of the gap, partly because the cell walls are interconnected by T-junctions. In some of our samples, we observed direction-dependence of the band gap. We plot in Fig. 3d a snapshot of propagating SAWs in the first onion layer (i.e. the oldest), filtered in the 120-140 MHz range. In the lower left quadrant, we see that acoustic energy propagates around -120°, while no acoustic energy remains after -150°. To explain this, for each angle θ, which denotes the direction of the radial wave vector, we project the $f-k$ dispersion maps on the $f$-axis to obtain the average displacement amplitude as a function of angle, as is shown in Figs. 3e, 3f (see Supplementary note 3). For the bare substrate (Fig. 3e), we observe two regions with low energy around 100° and -80° that are due to a lower sensitivity of the laser-deflection probe. This artifact was visible on all the epidermises we probed, as well as on the bare substrate (Fig. 3e). In contrast, in Fig. 3f, which corresponds to the oldest onion layer we probed, we observe a gap extending almost over all the $f-θ$ map that has a clear θ-dependence. We highlight in Fig. 3f with a white square a region corresponding to the lower left quadrant in Fig. 3d (between -180 and -90°). There, we observe that the gap opens and closes successively in the 120-140 MHz range, supporting the observation we made in the filtered image (Fig. 3d).
While natural anisotropy of the plant epidermis can lead to such directional band gaps, we note that microfibril reorientation can also be forced by repeated loading cycles (making the epidermis stiffer in the direction of the applied strain). Such conditioning may offer yet another means to control plant-based metamaterials. By leveraging the multiple advantages provided by biological composites, we anticipate our advances will lead to a wide range of new green and sustainable microdevices with tailored functionalities.

We have shown the repeatable occurrence of phononic band gaps in wild type onion epidermis, with extinction ratios comparable to those obtained in pillar-based locally resonant metamaterials. These results suggest a significant potential for acoustic manipulation of SAWs using organic surfaces. Most of micro- or nano-structured materials, such as cantilever-shaped MEMS or pillar-based resonators, are fragile and are thus designed in such a way that they do not support static load and remain sensitive. Conversely, organic layers, as well as many other plant samples, are stiff cellulose-based composites, wherein individual fibers have a rigidity comparable to steel that confer a high resistance to tensile stresses. For demanding applications, one can even envisage to preserve the structure of the plant samples, by drying the samples for long term storage, fixation or boiling to inactivate endogenous enzymes. At the same time their structure is dynamic, in that it can be modified when the plant is alive, for instance by stretching the epidermis to align the cellulose fibers, and it is conformable, meaning that they can be shaped to follow the corrugation of a substrate. This ability paves the way to soft metamaterial with an increased robustness compared to their inorganic counterparts. In addition, altering the cell structure in plants with high reproducibility (e.g. changing the shape of the cell walls) is quite standard using known genetic mutations, as is classically done in Arabidopsis, for instance. Based on more recent works, one could also use bio-hybrid approaches, such as chemical infiltration, to endow new or augmented properties in the plant-derived structures.

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