Top-Down Approach Making Anisotropic Cellulose Aerogels as Universal Substrates for Multifunctionalization

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ABSTRACT: Highly porous, strong aerogels with anisotropic structural properties are of great interest for multifunctional materials for applications including insulators in buildings, filters for oil cleanup, electrical storage devices, etc. Contemporary aerogels are mostly extracted from fossil resources and synthesized from bottom-up techniques, often requiring additional strategies to obtain high anisotropy. In this work, a universal approach to prepare porous, strong, anisotropic aerogels is presented through exploiting the natural hierarchical and anisotropic structure of wood. The preparation comprises nanoscale removal of lignin, followed by dissolution–regeneration of nanofibers, leading to enhanced cell wall porosity with nanofibrillated networks occupying the pore space in the cellular wood structure. The aerogels retain structural anisotropy of natural wood, exhibit specific surface areas up to 247 m²/g, and show high compression strength at 95% porosity. This is a record specific area value for wood aerogels/foams and even higher than most cellulose-based aerogels for its assigned strength. The aerogel can serve as a platform for multifunctional composites including scaffolds for catalysis, gas separation, or liquid purification due to its porous matrix or as binder-free electrodes in electronics. To demonstrate the multifunctionality, the aerogels are successfully decorated with metal nanoparticles (Ag) and metal oxide nanoparticles (TiO₂) by in situ synthesis, coated by the conductive polymer (PEDOT:PSS), and carbonized to yield conductive aerogels. This approach is found to be a universal way to prepare highly porous anisotropic aerogels.

KEYWORDS: aerogel, anisotropy, top-down, wood nanotechnology, biocomposite

Aerogels are porous solids often prepared from a gel form substrate where a solvent is exchanged for air or gas. The aerogels exhibit high surface area and low density, rendering them suitable in a broad range of materials applications, ranging from insulators in buildings to supercapacitors in electrical devices. A wide variety of aerogels are available, among which inorganic silica aerogels are most studied. However, the impaired strength properties limit their use. Synthetic polymers can overcome fragility problems, but the strength is limited and they are often derived from fossil resources. Aerogels based on natural polymers are of increasing interest due to potential eco-friendly characteristic. Reported biobased aerogels include bacterial cellulose, starch, alginate, lignocellulose, and chitosan-based aerogels. Cellulose nanofiber (CNF) aerogels are interesting due to high strength-to-density and large surface area-to-weight ratios. The low thermal conductivity renders them suitable for insulating purposes. Low density is an advantage, and the highly tailorable pores show promise for drug delivery, filters, and adsorbent applications. Through tailoring of the conductivity, the aerogels have also been used as components in electronic devices.

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the bottom-up approach, where nanocellulose is extracted from sources and prepared by reconstructing of the building blocks to form aerogels. This approach is time-consuming and energy intensive and usually results in aerogels with isotropic structural properties. An anisotropic structure is beneficial where directional conduction is needed, such as electrodes for batteries and substrates for solar steam evaporation.

However, assembly of anisotropic materials is challenging. For nanocellulose aerogels, a common method is directional freeze-casting, in which ice crystallites are formed by inducing a temperature gradient during freezing. Even though the method is useful, the addition of auxiliary processing steps results in time demanding, costly fabrication with substantial energy needs.8,20,21

An alternative approach is a top-down synthesis of aerogels. Instead of reassembly of nanosized building blocks, this method can take advantage of the hierarchical structure of wood. Wood consists of anisotropic and highly elongated cells (20–30 μm in diameter, 1–3 mm in length), organized in a parallel manner. This results in orthotropic mechanical properties, but the cells can also transport water and nutrients for biosynthesis.22 The wood cell wall is a nanostructured composite, comprised of layers of cellulose nanofibril reinforcement embedded in a hydrated matrix of molecularly mixed hemicelluloses and lignin. In the thickest S2 layer the cellulose fibrils are strongly oriented at a comparably small angle with respect to the axial direction of the wood cell. The structural organization of the cells and the directional alignment result in excellent mechanical properties. Functionalization of the structure at the molecular or nanoscale could result in multifunctional nanomaterials of potential for large-scale applications.23,24 One limitation is the low specific surface area of wood structure. Preparation of wood structured aerogels is one solution, where the structural anisotropy is retained. Thus far, one type of wood aerogel has been reported, where the cell wall was delignified and part of the hemicellulose was removed.9,25,26 However, none of these materials previously studied have a specific surface area comparable with nanocellulose-based aerogels.

In this paper, a highly porous anisotropic wood aerogel is prepared by a top-down synthesis method based on nanostructural wood cell wall manipulation. The preparation comprises delignification followed by partial dissolution/regeneration of the cell wall using dimethylacetamide/lithium chloride (DMAc/LiCl). The cosolvent partially dissolves the cell wall and regenerates nanofibrils. As a result, a porous cell wall is obtained with a nanofibrous network in the central lumen pore space of wood cells. Figure 1 illustrates the wood aerogel preparation procedure. This approach preserves the structural anisotropy of wood and introduces exceptional specific surface areas. The combination of formed nanofibrils in the cell lumen and retention of highly aligned cellulose of the cell walls leads to a rare combination of large surface area with good mechanical strength, not commonly seen in aerogels. The aerogel is further modified by in situ synthesis of Ag nanoparticles and TiO2 nanoparticles that effectively deposited on both the cell walls and the formed nanofibrils of the cell lumen. The conductive polymer (PEDOT:PSS) (poly(3,4-ethylenedioxythiophene) polystyrenesulfonate) is seamlessly coated onto the aerogel, and carbonization of the aerogel is also performed to yield conductive carbon aerogels with an extensive surface area. This highly adaptable aerogel shows promise as an eco-friendly alternative for multifunctional scaffolds in several application areas.

**RESULTS AND DISCUSSION**

Balsa wood derived aerogels with retained hierarchical structure were prepared first by delignification to increase porosity and chemical accessibility. After delignification, the wood was treated with an 8 wt % solution of DMAc/LiCl for 24 h, which partially dissolved the wood cell wall. The samples were subjected to solvent exchange to precipitate, followed by freeze-drying or critical point drying (CPD) to form aerogels. This is a facile preparation of biobased anisotropic aerogels; however, from a perspective of large-scale production, the drying methods could potentially be an obstacle. The preparation is exemplified in Figure 2, depicting the change in chemical composition, material properties, and Brunauer–Emmet–Teller (BET) specific surface area (SSA). Images of typical dimensional changes of the specimens are shown in Figure S1a–c, and data of dimensional change, weight change, and density are shown in Table S1. The average material yield after delignification and freeze-drying was 76 wt %, whereas the wood aerogels obtained a yield of 60 wt %. This means removal of lignin represented the larger material loss, and the dissolution/regeneration process the lesser.

The delignification procedure was performed with NaClO2 in an acetate buffer solution of pH 4.6. Delignification takes place by radical action of ClO2, where aromatic rings are oxidized, resulting in ring-opening reactions that effectively decomposes the lignin. Nearly 90% of lignin was removed during the procedure (Figure 2). After the procedure, the light-absorbing chromophoric groups, mainly from lignin, were removed, resulting in a white texture (Figure 3a and b). As lignin was removed, the cell wall layers became partially delaminated and voids were generated in the previously lignin-rich middle lamella (Figure 3d and e). In addition, nanoscale pores are generated in the cell wall after delignification, as reported by Fu et al.27 The extensive loss of lignin and increased cell wall porosity resulted in a more accessible cell wall for chemical agents. Data recorded by the N2 sorption method and the calculated BET surface area support the observation of increased cell wall porosity (Figure 4a). The
BET surface area of the wood after delignification and freeze-drying (FD) was 9 m$^2$/g compared with native wood, exhibiting 2 m$^2$/g. Note that these values are still very low compared with typical nanocellulose aerogels. Following delignification, DMAc/LiCl treatment was introduced to partially dissolve the wood cell wall. After regeneration of the dissolved cell wall in acetone and drying, a wood aerogel was obtained. The final aerogel shows similar macroscopic appearance as delignified wood (Figure 3c) yet with vastly different nanostructure. From the SEM images of the wood aerogel cross-section, the lumen space of the fibers is filled by a nanofibril network (Figure 3f). DMAc/LiCl is reported with the ability to extract cell wall components, mainly by dissolving cellulose. When an antisolvent (acetone in this study) is added, intermolecular hydrogen bonds are re-formed and the dissolved cellulose chains coagulate to form nanofibril structures in situ in the lumen. In the regeneration bath, the following reasons limit the diffusion of formed nanofibril network out of the wood: (1) the small pores of the cell wall, (2) the interaction of the nanofibrils and the cell wall, and (3) the shrinkage of the wood structure during the solvent exchange. During the dissolution process, roughly 41% of the crystalline cellulose was dissolved. Wherein the regenerated cellulose inside the wood accounted for approximately 34% of the dissolved cellulose, the remaining portion was lost to the regeneration bath. Retention of nanofibril networks of the lumen space after drying is attributed to the high concentration of fibrils and the rapid freezing rate. The dense presence of fibrils has been reported to hinder growth of ice crystals.
preserving separation of fibrils, observed by Jin et al. A faster freezing rate was also observed to retain separation of the fibrils in nanocellulose aerogels, again through prevention of ice crystal growth.

Figures 3g–i show the plane perpendicular to the fiber direction for native wood (Figure 3g), delignified wood (Figure 3h), and the regenerated wood aerogel (Figure 3i). The regenerated nanofibril network is present in the lumen space throughout the aerogel (Figure 3i), but is absent in the other materials. The regenerated nanofibers showed approximate diameters ranging from 10 to 20 nm (Figure S2). The nanofiber-filled lumen structure suggests high specific surface area. N2 sorption data showed that the BET surface area reached up to 159 m2/g for wood aerogels after FD, seen in Figure 4a. In addition, a higher value of 247 m2/g was obtained after CPD. From the results above, a hierarchically structured anisotropic aerogel with very high BET surface area is achieved. The BET surface area of wood after both FD and CPD had higher values than any other documented top-down prepared wood aerogel.

To investigate the partial dissolution mechanism, powder X-ray diffraction (XRD) was performed, displayed in Figure 4b. Native wood and delignified wood show cellulose I structures with crystallinity indices of 0.73 and 0.83, respectively. According to the literature, cellulose regeneration from DMAc/LiCl dissolution is often in a nonordered manner, consisting of both amorphous and paracrystalline cellulose. By analyzing the intensity changes of crystalline native cellulose I (14.7°, 16.6°, 22.8°, and 34.9°) and amorphous cellulose (21.5°), an indication of the dissolution/regeneration mechanism can be acquired. The regenerated wood aerogel displays lower intensities for cellulose I peaks and a broader amorphous peak, based on a peak deconvolution method. The partially retained peaks for cellulose I confirm its presence in the cell wall. In Figure 4b, the [200] peak for the wood aerogel has a slight shift toward lower angles. The occurrence is not fully known, but it could result from the increase of amorphous cellulose, in close proximity to the crystalline cellulose. Kentaru Abe and Hiroyuko Yamamoto postulated that the swelling matrix of wood interacts mechanically with microfibrils of the cell wall, giving rise to a peak shift upon drying.

Part of the cell wall is apparently not affected by the solvent during the DMAc/LiCl treatment. With the peak deconvolution method, the crystallinity index (CI) was calculated to be 73%, 83%, and 63% for native wood, delignified wood, and the wood aerogel, respectively (Table 1). The decreased CI of the wood aerogel is a consequence of dissolution and regeneration of cellulose. The broader amorphous peak indicates the formation of regenerated cellulose.

Despite the chemical treatment, the aerogel still shows preserved macroscopic wood structure. Figure 3f and i show the main wood cell wall structure preserved, leading to anisotropic properties of the resulting aerogel. The orientation of crystalline cellulose I can provide further information about the orientation of microfibrils of the cell wall. Hence, information about the anisotropy of the material can be obtained. The orientation was extracted through azimuthal integration of the diffraction images obtained from 2D wide-angle X-ray scattering (WAXS), seen in Figure 4c. The degree of orientation and CI are summarized in Table 1. The degree of orientation and CI peak height

Table 1. Degree of Orientation and Crystallinity Index of Native Wood, Delignified Wood, and Wood Aerogel

| Material        | Degree of Orientation | CI Peak Height  | CI Deconvolution |
|-----------------|-----------------------|-----------------|------------------|
| native wood     | 0.893 ± 0.004         | 81 ± 1.2        | 73 ± 0.9         |
| delignified wood| 0.862 ± 0.025         | 85 ± 0.4        | 83 ± 2.0         |
| wood aerogel    | 0.836 ± 0.006         | 68 ± 2.0        | 63 ± 11.4        |

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of orientation of cellulose microfibrils in native wood, delignified wood, and the wood aerogel was 0.89, 0.86, and 0.84, respectively. For the aerogel, the degree of orientation is significantly higher than for CNF or CNC-based aerogels or foams prepared by freeze-casting.\textsuperscript{10,20} Moreover, according to Kriechbaum et al.\textsuperscript{10} a specific energy consumption of \textgreek{e}3 kWh/kg was necessary to reach a degree of orientation of 0.82. In contrast, no mechanical shear is necessary in this work.

The mechanical properties of the native wood, delignified wood, and wood aerogel were characterized by compressive testing in the longitudinal direction. The mechanical behavior is summarized in Table 2 and Figure 5a. Notably, for native wood, a prominent linear region is shown between 0 and 4% strain. A clear distinction is apparent for delignified wood, where an initial region probably represents buckling of the edges of the specimen at low strain, followed by the linear region at 2.5–5% strain. The wood aerogel displays a protracted linear region encompassing 0–6% of strain before reaching the plateau region, where the cellular structure is collapsing, possibly by cell wall buckling or folding. The Young’s modulus of each specimen was 295, 82, and 25 MPa for native wood, delignified wood, and the aerogel, respectively. The Young’s modulus (compression) of native wood is similar to data reported earlier.\textsuperscript{27} The value is low compared with data for higher density wood, since the volume fraction of wood tissue is only 9%. The structural collapse of native wood occurred at 12.9 MPa, in agreement with previous observations.\textsuperscript{34} The highly oriented microfibrils and anisotropy provided by the parallel honeycomb-like cells of native wood and the preserved lignin cell wall adhesive result in a strength significantly higher than for treated wood. Delignified wood attains a yield strength of 3.9 MPa, whereas the wood aerogel achieves a yield strength of 1.2 MPa. Compared to native wood, the lower strength of delignified wood is due to lignin removal. Furthermore, the delignification results in partial delamination of the cell wall (Figure 3e). For the wood aerogel, the DMAc/LiCl treatment leads to higher cell wall porosity and lower degree of crystallinity, so that yield strength

| density \[kg/m^3\] | Young’s modulus \[MPa\] | yield strength \[MPa\] |
|-----------------|-----------------|-----------------|
| native wood | 129 ± 2 | 295 ± 24 | 12.9 ± 0.9 |
| delignified wood | 90 ± 12 | 82 ± 32 | 3.9 ± 1.5 |
| wood aerogel | 79 ± 8 | 25 ± 13 | 1.2 ± 0.3 |

Figure 5. (a) Compressive stress–strain curves of native wood, delignified wood, and wood aerogel, at 50% relative humidity and 23 °C. (b) Yield strength against the specific surface area for documented cellulose-based aerogels and foams. (c) Yield strength versus relative density for documented cellulose-based aerogels and foams, compared to this work.\textsuperscript{3,25,36–38,44–52} Black symbols represent aerogels and foams prepared to obtain anisotropy, whereas red symbols are aerogels and foams prepared with isotropic synthesis methods. $\rho^*$ represents the density of the individual porous material \[kg/m^3\], and $\rho_s$ represents the density of the solid cell wall material. Numbers in parentheses represent the wt % of the component. Relative densities used were either retrieved from the article or calculated, where the solid density 1500 kg/m$^3$ was assumed for cellulose.\textsuperscript{34}
is lowered compared with delignified wood. Mechanical properties from compression perpendicular to the longitudinal direction (radial direction) can be seen in Figure S4.

The wood aerogel is special in its combination of mechanical strength and high BET surface area, compared with cellulose-based aerogels and foams reported in the literature.3,25,35−52 Figure 5b,c show the yield strength and BET surface area for documented cellulose-based aerogels and foams. In Figure 5c, the yield strength increases with higher relative density. The wood aerogel is positioned slightly higher than the anisotropic aerogel trend would suggest; see black symbols in Figure 5c.

A substantial advantage for the wood aerogel compared with anisotropic aerogels is the BET surface area of up to 247 m²/g, compared with 20−30 m²/g25,41 for anisotropic nanocellulose aerogels, Figure 5b. The penalty in reduced strength for the wood aerogel is small. For isotropic aerogels, red symbols, a much lower yield strength at given relative density is observed. On the other hand, such aerogels can have BET surface areas of up to 340 m²/g.49 Here, the preserved wood honeycomb structure and high microfibril orientation contribute to the strength, so that high strength and high specific surface area are combined.

The combination of high specific surface area and good mechanical properties is difficult to achieve; resolving that challenge for wood, a hierarchically structured template for multifunctional materials can be realized. The wood aerogel obtained in this work can be used to create nanostructured materials for high-technology applications. To demonstrate the possibility, four systems were used to functionalize the wood aerogel. They include in situ synthesis of metal nanoparticles (Ag) and metal oxide nanoparticles (TiO₂) and internal coating by conductive PEDOT:PSS polymer in the wood aerogel structure, followed by carbonization. Reduction of Ag from aqueous metal salts resulted in nanoparticle formation (Figure 6a) throughout the aerogels. The nanoparticles are present in both the fiber cell walls and the nanofiber networks in the lumen pore space; see SEM and energy-dispersive X-ray spectroscopy (EDX) images in Figure 6a. The TiO₂ nanoparticles were also successfully precipitated inside the wood aerogel as confirmed by EDX data; see Figure 6b.

Ag aerogels have been investigated as functional materials for solar steam generation53 and antibacterial applications,54 and TiO₂ aerogels have been studied as photocatalytic templates.55 In addition to nanoparticle functionalization, the wood aerogel could also be functionalized by layers of conductive polymer coatings inside the structure. The electrically highly conductive polymer PEDOT:PSS is infiltrated into the cellulosic matrix and forms a layer on the fiber cell walls (Figure 6c). A cross-section along the fiber direction after PEDOT:PSS infiltration is presented in Figure S5. The image confirms that the polymer is distributed throughout the fibers of the aerogel.

As a final demonstration, the unmodified, highly porous anisotropic aerogel was carbonized to form porous carbon aerogels; see Figure 6d. The carbonized wood aerogel shows a BET surface area of 555 ± 3 m²/g, more than double that of carbonized native wood (235 ± 4 m²/g). BET surface area isotherms are presented in Figure S6. The carbonized aerogel retains the anisotropic morphology of the wood aerogel, but the fiber lumen pore channels show a significantly smaller diameter. The overall shrinkage of the carbon aerogel is illustrated in Figure S1d. Native balsa wood has a fiber lumen diameter of approximately 18 μm,22 but after carbonization the lumen diameter is decreased and the nanofibril structures in the cell lumen are still present. The carbon aerogel shows promise as a self-supporting, nonadditive, binder-free electrode.
with potential in supercapacitor applications. Capacitance measurements from cyclic voltammetry can be seen in Figure S8, where the carbon aerogel exhibited 66 F/g compared to 5 F/g for native carbonized wood at a scan rate of 10 mV/s. Supporting SEM images of the carbon aerogel can be seen in Figure S7. Both the PEDOT:PSS-infiltrated wood aerogel and the carbon aerogel show good electrical conductivity with values of 213.7 and 371 ± 5 S/m, respectively. In all cases, the partial nanofibrillation of the cell wall is maintained.

CONCLUSIONS

A highly porous, anisotropic aerogel of exceptionally high specific surface area combined with good mechanical properties is reported. It has a distinctive nanostructure, where a cellulose nanofibril network occupies the porous lumen space of wood cells. It is prepared from native wood using a top-down synthesis approach. Lignin is removed from the native wood cell wall to form a porous cellulose template for facile permeation of a cellulose solvent. The cosolvent partially dissolves the cell walls, with cellulose precipitation in the form of nanofibril networks inside the lumen. The aerogel exhibits a crystallinity index of 63%, and the degree of crystalline cellulose I orientation is 0.84.

The increased porosity and partial cell wall fibrillation results in specific surface areas as high as 247 m²/g for critical point drying and 159 m²/g for freeze-drying. The oriented honeycomb structure and high fibril orientation result in a yield strength of 1.2 MPa, at a porosity of 95%. This is higher than for comparable isotropically structured nanocellulose aerogels.

The multifunctionality of the wood aerogel is demonstrated by three concepts: (a) through reduction of Ag and TiO₂ for nanoparticle formation in the wood cell wall, (b) infiltration of the PEDOT:PSS conducting polymer, and (c) wood aerogel carbonization. Reduced nanoparticles form inside the oriented nanofibril cell wall as well as in the fibril networks inside the 20 μm lumen pore space of the aerogel. The electrically conductive PEDOT:PSS was readily impregnating the aerogel, and the carbonized aerogel also showed anisotropic character; both materials showed enhanced electrical conductivity. The present synthesis approach is a facile method to prepare anisotropic aerogels from wood of potential for multifunctional materials for applications such as sensors, chemical reactors, and electrical devices.

EXPERIMENTAL SECTION

Materials and Chemicals. Balsa wood (Ochroma pyramidale) with a density of 129 ± 2 kg/m³ was bought from Material AB, Sweden, and subsequently cut to the dimensions of 15 × 15 × 10 mm³ (tangential × radial × axial). The delignification chemical, sodium chlorite (NaClO₂), was purchased from Sigma-Aldrich, and the delignification buffer was prepared from sodium acetate and acetic acid, bought from Sigma-Aldrich, Sweden. The solvent exchange chemicals ethanol and acetone were purchased from VWR, Sweden. Lithium chloride (LiCl) and dimethylacetamide (DMAc) for wood treatment was bought from Sigma-Aldrich, Sweden. Titanium(IV) butoxide (97%), AgNO₃ (>99%), polyvinylpyrrolidone (molecular weight 10k), benzyl alcohol, and PEDOT:PSS (1.3 wt %, conductive grade) were also purchased from Sigma-Aldrich, Sweden.

Delignification of Balsa Wood. This was performed by immersing the native wood into a solution containing 1 wt % sodium chlorite in an acetate buffer of pH 4.6. The delignification occurred for a duration of 18 h, in which the buffer solution was renewed every 6 h. After delignification, the balsa templates were thoroughly rinsed with Milli-Q water.

Preparation of DMAc/LiCl-Treated Aerogels. Washed and delignified templates were introduced to an ethanol solvent exchange, followed by a DMAc solvent exchange. Solvent exchange was repeated four times, respectively. When the templates were completely infiltrated with DMAc, the samples were submerged in the DMAc/LiCl solvent containing 8 wt % LiCl. The templates were treated for 24 h and subsequently introduced to a regeneration bath of acetone. The resultant templates were rinsed in Milli-Q water and rapidly frozen in liquid nitrogen (−196 °C), followed by freeze-drying (−83 °C) for at least 48 h to yield aerogels. Aerogels were also obtained from critical point drying, in which the templates were first solvent exchanged to ethanol followed by CPD in an Autosamdr-815.

In Situ Synthesis of Nanoparticles (NPs) in the Aerogel. This was performed with a temperature-controlled microwave oven (Biotage Initiator). Time and temperature was adjusted according to the specimen and was subsequently cooled with compressed air to 50 °C after completion. Silver NPs were synthesized by mixing a solution of 2 mL of 50 mM AgNO₃, 10% polyvinylpyrrolidone and 2 mL of 25 mM AgNO₃. The aerogels were immersed into the mixture under vacuum for at least 4 h. The Ag reduction was executed at a temperature of 120 °C for 10 min in the microwave. For the titania synthesis, the aerogel was solvent exchanged from water to benzyl alcohol for 12 h, to ensure complete infiltration. A 14 μL (0.04 mmol) amount of titanium butoxide was mixed into 4 mL of benzyl alcohol, and the aerogel was then submerged into the mixture. The heating was then performed in two consecutive steps. First, the mixture was heated to 50 °C for 5 min followed by 120 °C for 10 min. The samples were subsequently cleaned in acetone for 20 min under vacuum; this was repeated four times. Lastly, the samples were solvent exchanged for water and freeze-dried for 24 h. Ag and TiO₂ reduced on the cellulose template following a similar approach detailed by Roig et al.⁵⁶ The proposed reaction mechanism can be seen elsewhere.⁵⁵–⁵⁸

Wood Carbonization. Native wood and freeze-dried DMAc/LiCl-treated delignified wood was pyrolyzed in a tubular furnace (model: Nabertherm, tube furnace R 50/250/13). The pyrolysis transpired in a N₂ atmosphere (100 mL/min flow) with a heating rate of 5 °C/min from RT to 1000 °C, in which an isothermal treatment occurred for 3 h in a nitrogen atmosphere. Thereafter, the samples were cooled at a rate of 10 °C/min.

PEDOT:PSS Aerogel Preparation. Aerogels were infiltrated with the conductive polymer PEDOT:PSS by filtering a 1.3 wt % PEDOT:PSS solution dropwise onto the aerogel, using a Büchner funnel. The PEDOT:PSS retentate during filtration was reintroduced to the aerogel three consecutive times to ensure complete infiltration of the polymer. The infiltrated aerogels were thereafter washed in water and subsequently freeze-dried to obtain the conductive aerogels.

Characterization. Morphology of native wood, delignified wood, and the wood aerogel was analyzed using a field emission scanning electron microscope (Hitachi S-4800, Japan). The samples were sputtered with a platinum/palladium conductive layer using a Cressington 208HR, UK. For 20 s, resulting in a layer thickness of approximately 3 nm. Positioned on the FE-SEM an EDX instrument (Oxford Instruments, X-MAX N 80, UK) was equipped. EDX was used to evaluate the presence of elements and was carried out with an accelerating voltage of 15 kV during TiO₂ sample examination and 30 kV for the Ag specimen. The working distance was 15 mm, and the composition was gathered with a point ID analysis, taken from six different regions of the EDX images. The density of the specimens was obtained by drying the samples in a 105 °C oven overnight, followed 30 min under vacuum before weighing. Thereafter, the porosity was calculated by utilizing eq 1, where the solid density of balsa wood was assigned as 1500 kg/m³ and solid density of cellulose as 1500 kg/m³.⁴₄
Inside the aerogel; compressive stress—strain curves in the radial direction of specimens; digital image of the PEDOT:PSS-infiltrated aerogel; adsorption and desorption isotherms of carbonized wood and carbonized wood aerogel; SEM images showing the maintained nanofibrillated networks inside the carbonized aerogel; specific capacitance from cyclic voltammetry of the carbon aerogel and native carbonized wood (PDF)

ASSOCIATED CONTENT

Supporting Information

The Supporting Information is available free of charge at https://pubs.acs.org/doi/10.1021/acsnano.0c01888.

Digital images representing dimensional changes of the material throughout the preparation procedure; data of dimensional changes, weight, and density during the preparation procedure; diameter distribution of regenerated fibrils; SEM images of nanofibrillated networks inside the aerogel; compressive stress—strain curves in the radial direction of specimens; digital image of the PEDOT:PSS-infiltrated aerogel; adsorption and desorption isotherms of carbonized wood and carbonized wood aerogel; SEM images showing the maintained nanofibrillated networks inside the carbonized aerogel; specific capacitance from cyclic voltammetry of the carbon aerogel and native carbonized wood (PDF)

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Notes

The authors declare no competing financial interest.

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