SUPPLEMENTARY INFORMATION

An Environmentally Benign Approach to Achieving Vectorial Alignment and High Microporosity in Bacterial Cellulose/Chitosan Scaffolds

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BC and BC-Chitosan composites were subjected to a slow freezing process using an ultra-cold freezer (Sanyo, Japan); the samples were placed in a freezer before which ended up producing large random ice crystals; after which, all the samples were subjected to freeze drying shown in Figure S1.

**Figure S1.** SEM microphotograph of BC produced by *Komagataeibacter xylinus* that were freeze dried by a slow freezing process. Note the presence of large pores and the absence of ostensible alignment.
Figure S2. SEM microphotograph of BC-Chitosan composites that were freeze dried by a slow freezing process. Note the presence of large pores and the absence of ostensible alignment.
Figure S3. Determination of eutectic point using DSC

DSC (model Q200, TA Instruments, New Castle, Del.) was applied under nitrogen atmosphere at a flow rate of 20 mL min⁻¹. Samples (~13 mg) were equilibrated at -30 °C for 2 min. and heated to -20 °C at a heating rate of 10 °C min⁻¹.
**Table S1.** Selected physical properties of all samples.

<table>
<thead>
<tr>
<th>Sample Name</th>
<th>Mean pore diameter (µm)</th>
<th>Porosity (%)</th>
<th>Absorption properties (%)</th>
<th>Surface Area (m²/g)</th>
<th>Crystallinity Index (%)</th>
<th>Dry Compressive Modulus (MPa)</th>
<th>Wet Compressive Modulus (MPa)</th>
</tr>
</thead>
<tbody>
<tr>
<td>BC</td>
<td>2 ± 1</td>
<td>75 ± 2</td>
<td>4671</td>
<td>201</td>
<td>89.45</td>
<td>9.51±0.23</td>
<td>37.32±1.07</td>
</tr>
<tr>
<td>BC-Ch-1%</td>
<td>4 ± 2</td>
<td>87 ± 1</td>
<td>1625</td>
<td>29.5</td>
<td>83.57</td>
<td>8.04±0.47</td>
<td>0.58±0.17</td>
</tr>
<tr>
<td>BC-Ch-1.5%</td>
<td>2.5 ± 1</td>
<td>80 ± 1</td>
<td>1532</td>
<td>22.06</td>
<td>83.94</td>
<td>9.13±0.40</td>
<td>0.82±0.15</td>
</tr>
<tr>
<td>BC-Ch-2%</td>
<td>1.4 ± 1</td>
<td>74 ± 2</td>
<td>1246</td>
<td>19.85</td>
<td>79.46</td>
<td>4.74±1.00</td>
<td>9.64±0.46</td>
</tr>
</tbody>
</table>
MTT test and Schwann cell culture

Per 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) assay (Fig. S4), the membrane extract demonstrated good biocompatibility as indicated by good cell viabilities compared to that of the control group during the different time.  

![MTT test graph](image)

**Fig. S4.** The MTT test in different time (black represents blank control, red for membrane).

TBO staining was utilized to observe the Schwann cell behavior on the samples (Fig. S5), which showed the cells had spindle shape with projections from the first day to the third day of culture, and the cells were homogeneously distributed on the hydrogels surface, these results indicated that the scaffold is suitable for Schwann cells attachment and growth.  

![TBO staining images](image)

**Fig. S5.** Attachment and proliferation of Schwann cells on scaffolds (A 1d, B 3d).
References


