## **Supporting Information**

for Plasma Process. Polym.

## Plasma-Treated Organic Solutions for Enhanced Electrospun Nanofibers

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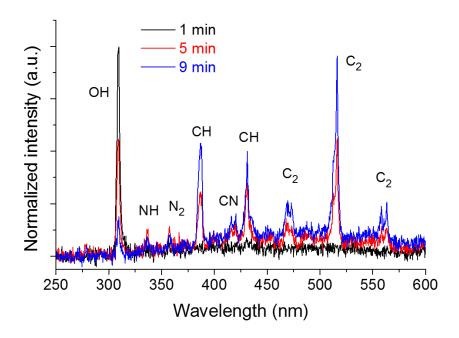


Figure S1. Effect of plasma treatment time on the optical emission spectrum of the plasma jet afterglow sustained in a 6% w/v PLA solution (PEPT parameters: 0.5 L/min, 2 kV).

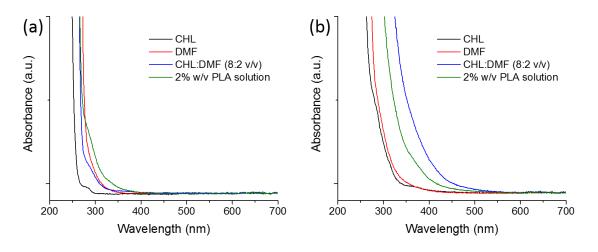


Figure S2. UV-vis absorbance spectra of (a) untreated solvents and a 2% w/v PLA solution, (b) plasma-treated solvents and a plasma treated 2% w/v PLA solution (PEPT parameters: 5 min, 0.5 L/min, 2 kV).

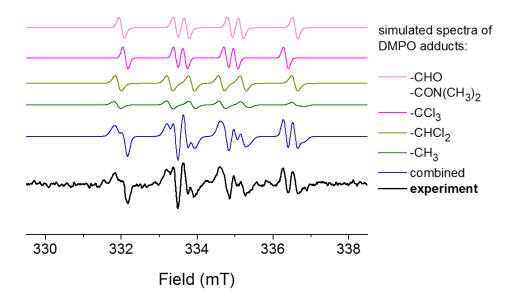


Figure S3. DMPO adducts in a 2% w/v plasma-treated PLA solution. Adducts: DMPO-CHO/CON(CH<sub>3</sub>)<sub>2</sub> ( $a_N = 1.41$  mT,  $a_H = 1.81$  mT); DMPO-CCl<sub>3</sub> ( $a_N = 1.33$  mT,  $a_H = 1.59$  mT); DMPO-CH<sub>2</sub> ( $a_N = 1.36$  mT,  $a_H = 1.93$  mT); DMPO-CH<sub>3</sub> ( $a_N = 1.52$  mT,  $a_H = 2.04$  mT).

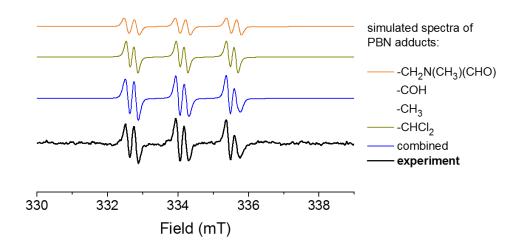


Figure S4. PBN adducts in a 2% w/v plasma-treated PLA solution. Adducts: PBN-CHCl<sub>2</sub> ( $a_N = 1.42 \text{ mT}$ ,  $a_H = 0.21 \text{ mT}$ ); PBN-CH<sub>2</sub>N(CH<sub>3</sub>)(CHO)/-COH/-CH<sub>3</sub> ( $a_N = 1.45 \text{ mT}$ ,  $a_H = 0.28 \text{ mT}$ ).

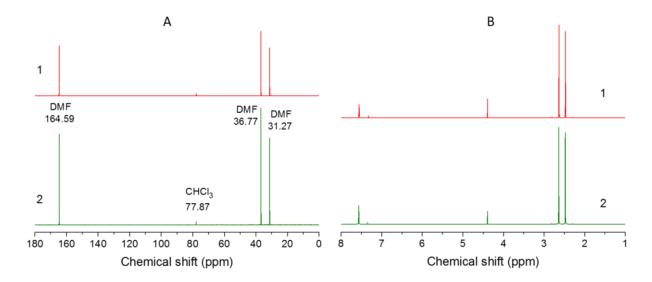
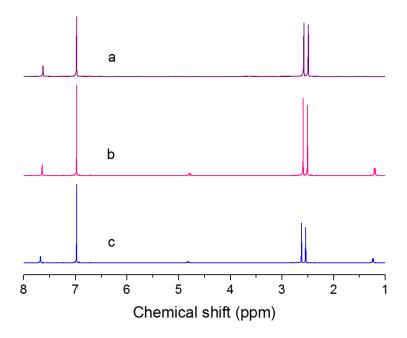


Figure S5. <sup>13</sup>C (A) and <sup>1</sup>H (B) NMR spectra of the aqueous extract from (1) an untreated and (2) a plasma-treated 2% w/v PLA solution.

In the <sup>13</sup>C NMR spectra shown in Figure S5, only the chemical shifts of carbon atoms of DMF and residual CHCl<sub>3</sub> were detected, with chemical shifts of 31.27, 36.77, 164.59 ppm (DMF) and 77.87 ppm (CHCl<sub>3</sub>). No differences in <sup>13</sup>C NMR spectra before and after plasma treatment can be revealed. <sup>1</sup>H-NMR analysis demonstrated signals with chemical shifts at 2.47 (NH<sub>2</sub>), 2.63 (OC=O), 4.39 (H<sub>2</sub>O), 7.32 (CHCl<sub>3</sub>), and 7.56 (DMF) ppm and the same signals were present in the D<sub>2</sub>O extract after PEPT (Figure S5B). In the extract of the PEPT solution, the H<sub>2</sub>O signals originated either from impurities in D<sub>2</sub>O or from generated HCl.



*Figure S6.* <sup>1</sup>H-NMR spectra of (a) a plasma-treated mixture solvent, (b) a plasma-treated 2% w/v PLA solution, and (c) an untreated 2% w/v PLA solution in CDCl<sub>3</sub>.

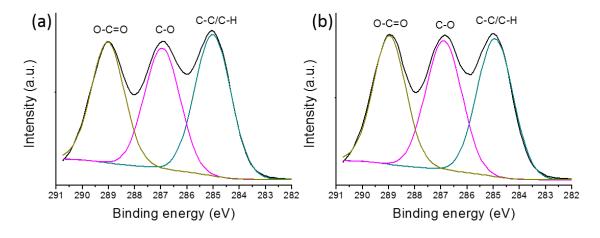


Figure S7. Fitted C1s peak of PLA nanofibers produced from (a) untreated and (b) plasma-treated (PEPT parameters: 5 min, 0.5 L/min, 2 kV) 6% w/v PLA polymer solutions.