

Characterization and adsorption potential of Lignosulfonate-based activated carbon fibers

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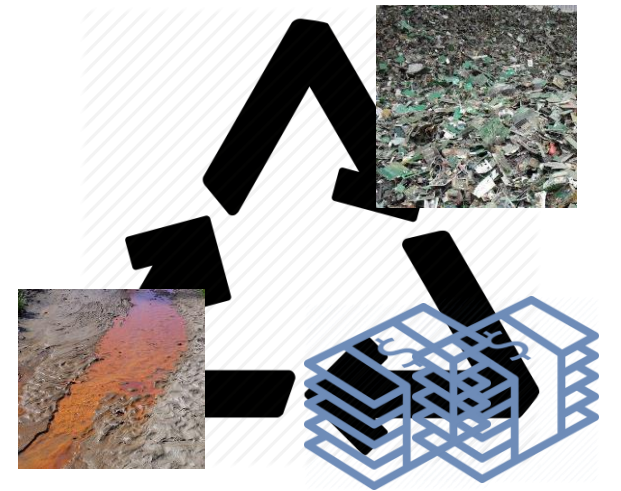
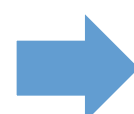
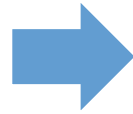
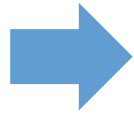


Outline

- Introduction
- Materials & Methods
- Results & Discussions
- Future Works

Introduction

- Why lignosulfonate-based activated carbon fiber & adsorption?



- Goal—Characterize properties → investigate its adsorption potential

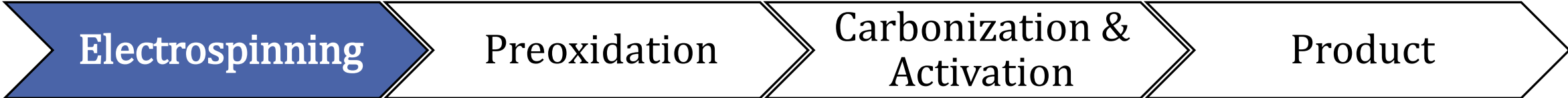
Preparation of Activated Carbon Fiber

Materials

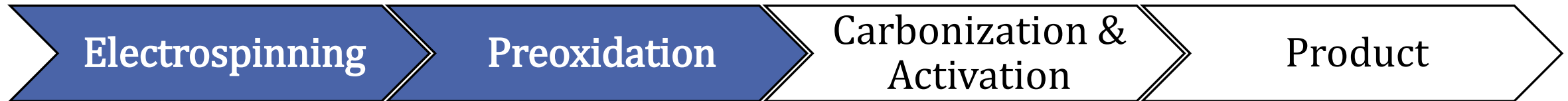


- Hardwood lignosulfonate (HLS)
Polyethylene oxide (PEO) (95:5)
+dH₂O
→30 wt% mixture
- @Room Temp, stir until dissolve

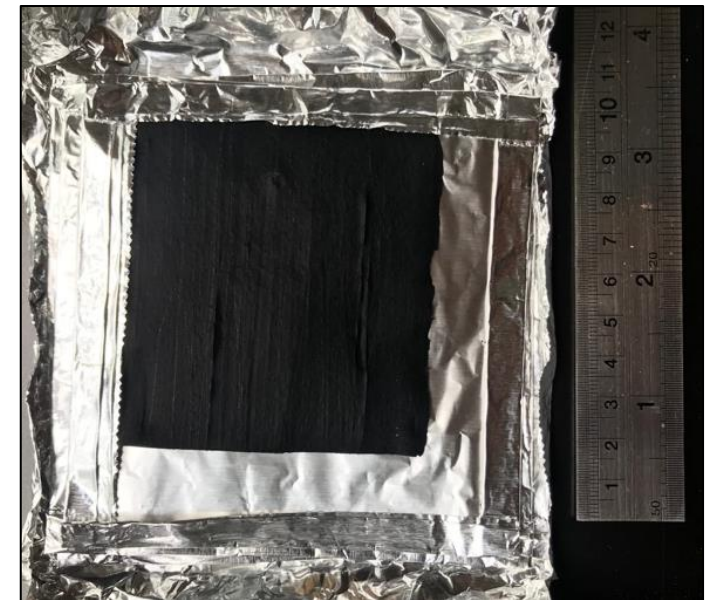
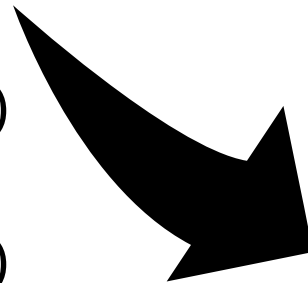
Methods—Production of Activated Carbon Fiber



Methods—Production of Activated Carbon Fiber



- (1) @ 1 °C/min,
100°C (30 min)
- (2) @ 1 °C/min,
300°C (30 min)



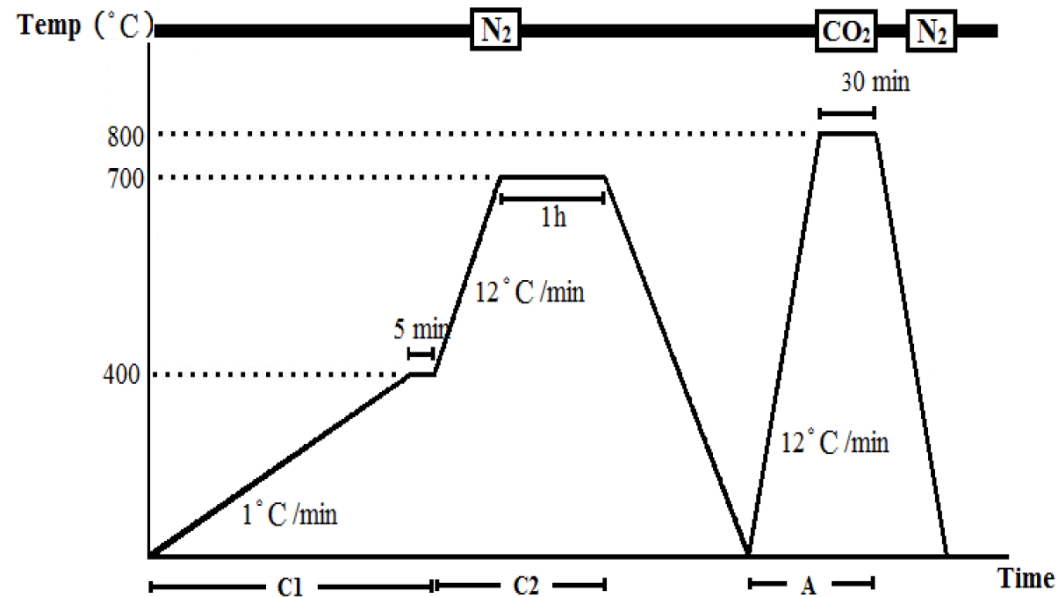
Methods—Production of Activated Carbon Fiber

Electrospinning

Preoxidation

Carbonization & Activation

Product



Analytical Methods

Factors influence adsorption ability

- Morphology → affect interactions between adsorbates and adsorbents
- Specific surface area → larger, greater potential
- Pore size distribution → adsorb substances with similar diameter
- Surface properties → electrons on the surface are helpful

Methods—Characterization of Activated Carbon Fiber

Physical Characterization

Scanning Electron Microscopy
(SEM)
Fiber Morphology

BET Analysis
(N₂@77K)
Pore structure & Surface Area

Quenched solid density
functional theory (QSDFT)
Pore Size Distribution

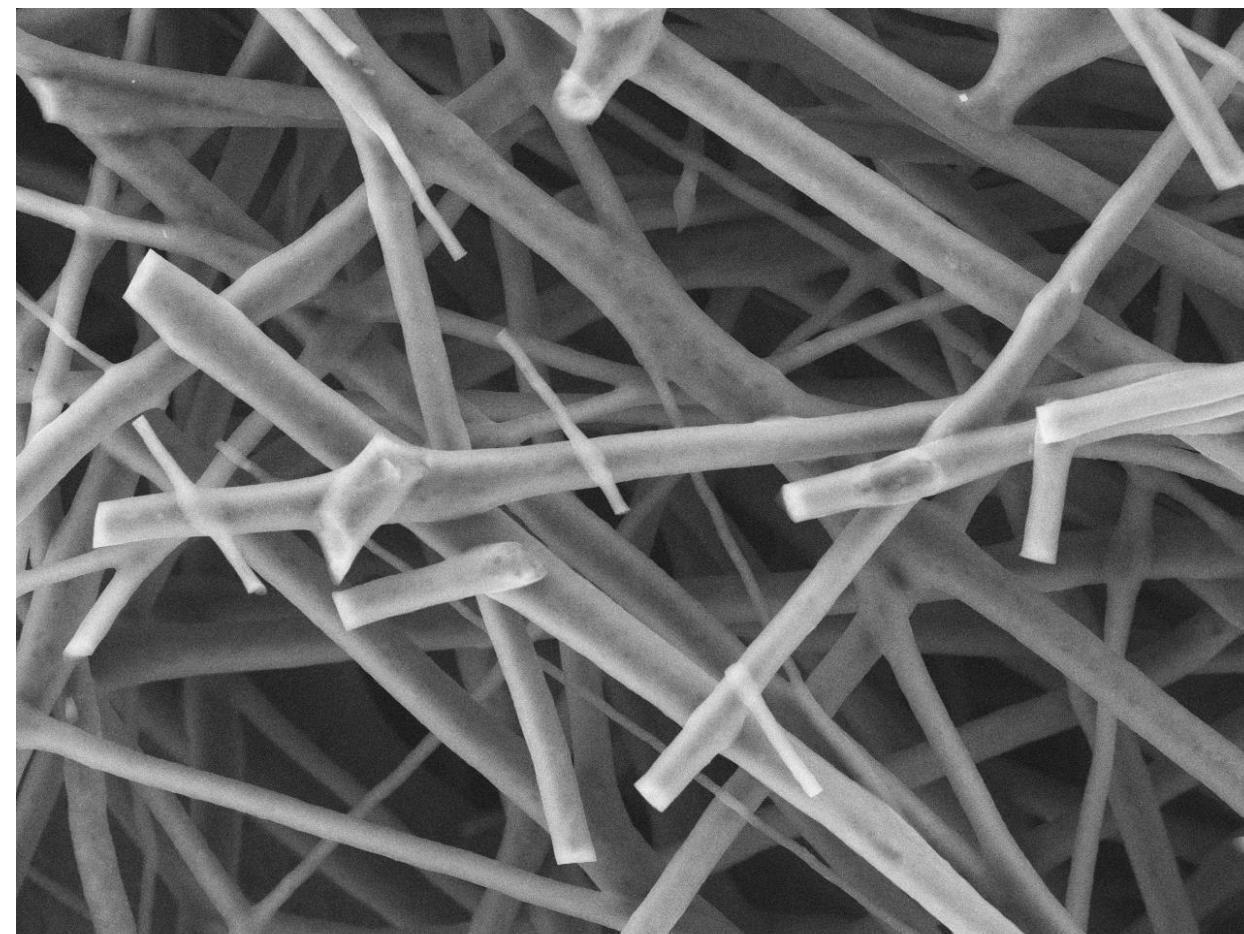
Chemical Characterization

Raman Spectroscopy
sp² electron

Results and Discussions

Physical Characterization – Fiber Morphology (SEM)

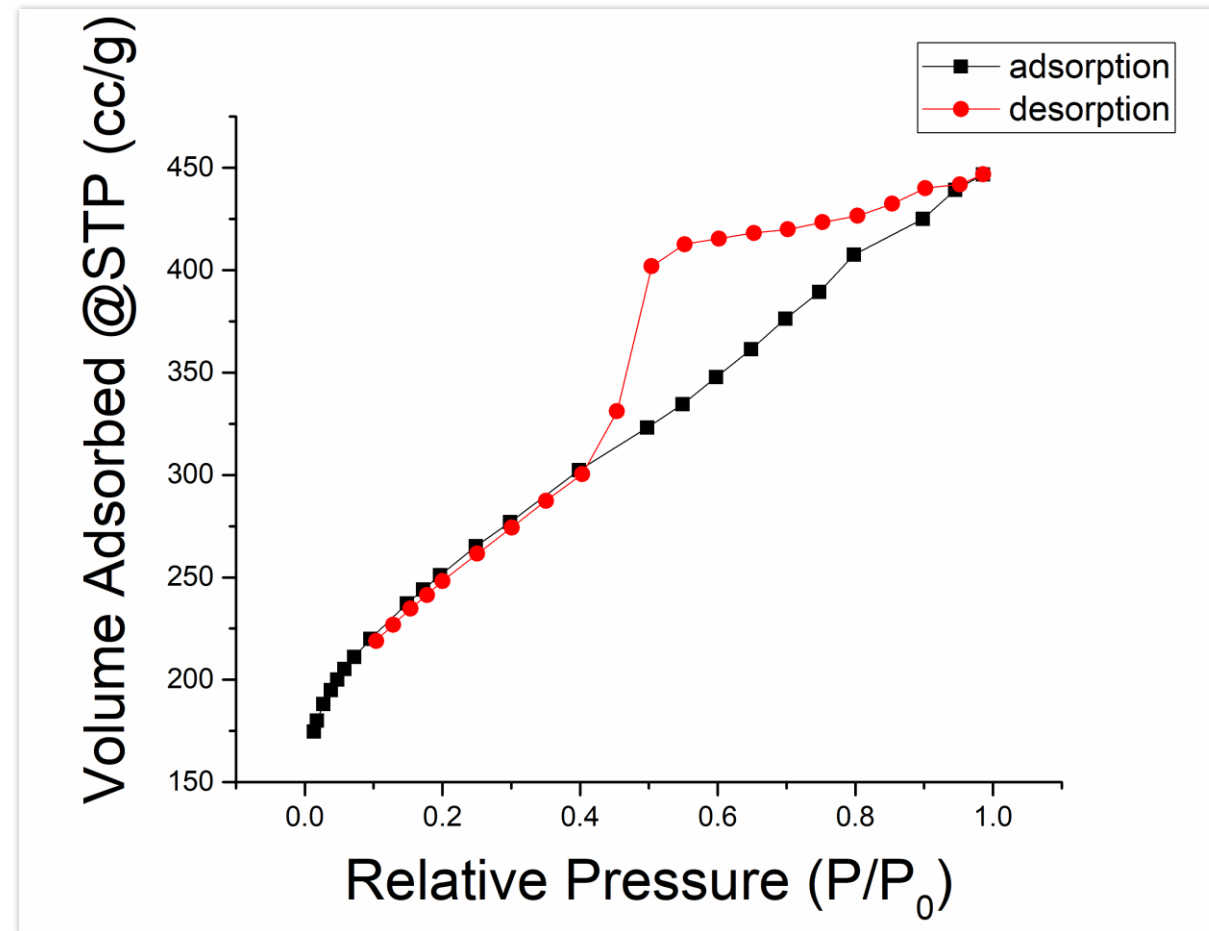
- Morphology
- 1) Fiber Fusion → diameter varies
→ Various pore size
- 2) Defective nest-like structure →
Dispersive forces



NMU x1.0k 100 μm

Physical Characterization – Structure (BET)

- Pore structure
 - 1) IUPAC: Mesopores (Major)
 - 2) Bottleneck-like pore connection
- Specific Surface Area
 $\approx 350 - 900(\text{m}^2/\text{g})$



Physical Characterization – Pore Size (QSDFT model)

○ Pore Size Distribution

1) Total Pore Volume

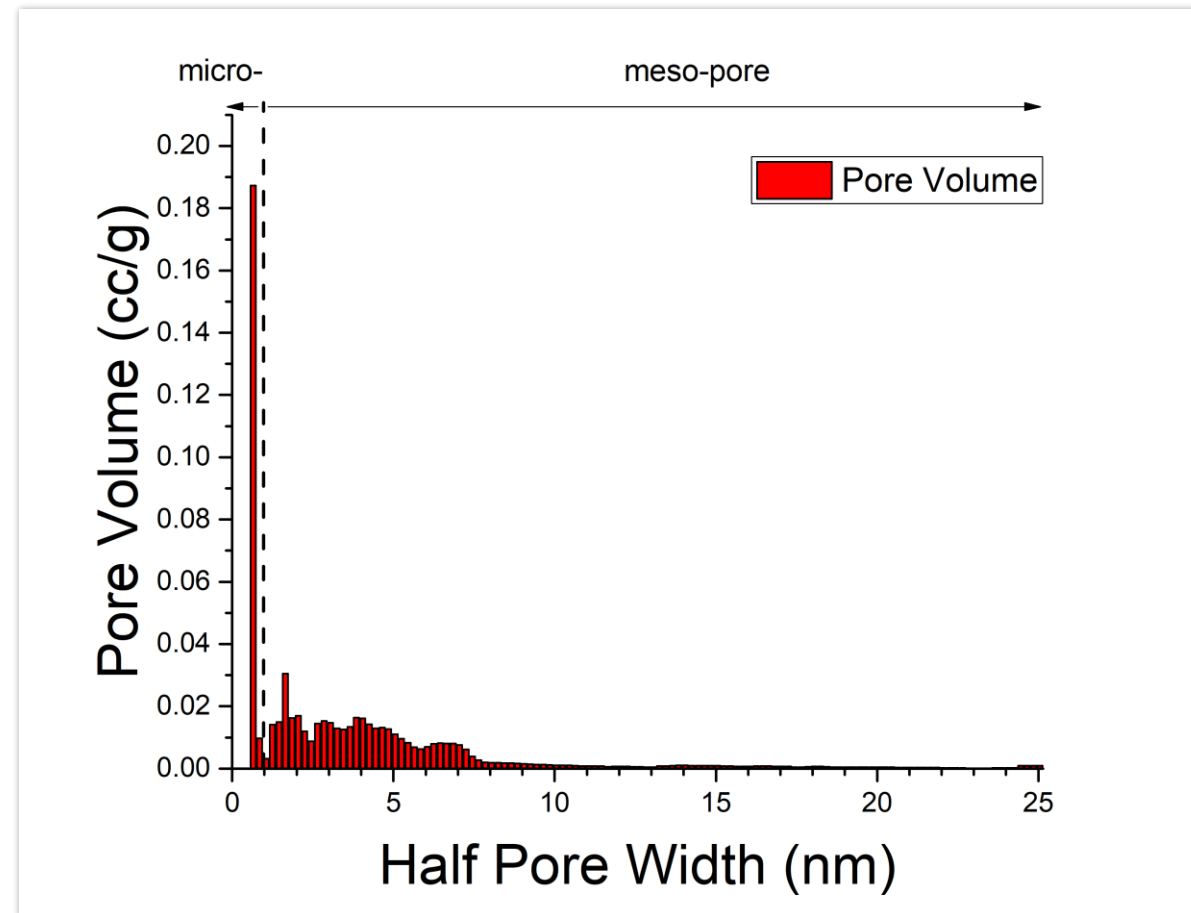
≈ 0.21 – 0.65 cm³/g

2) Micro/Mesopore (%)

≈ 30/70

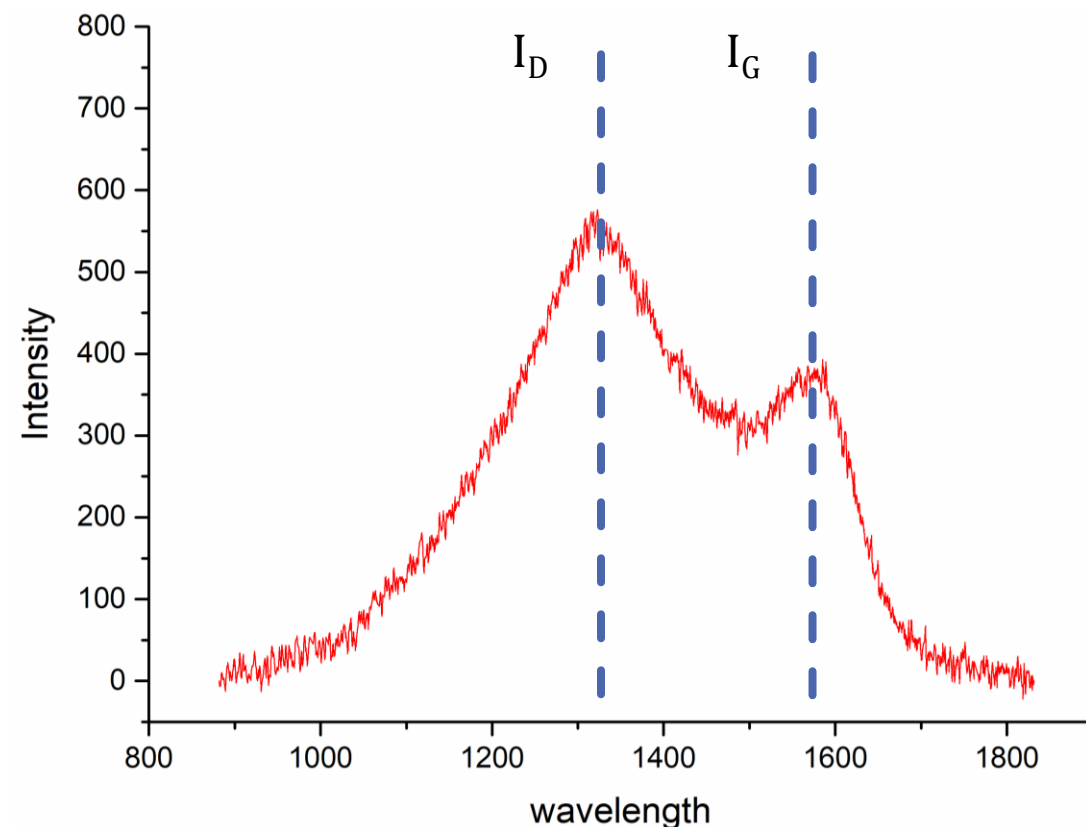
3) Mainly 1.6 – 20 nm

4) Hydrated metal ions' diameter → within micropore region



Chemical Characterization – π electron on surface

- Carbon Content ($\approx 50\%$)
- Raman Spectroscopy
 - (1) Further investigate C's structure
 - (2) I_D (Graphite, C-C, sp^3), around 1350 cm^{-1}
 - (3) I_G (Graphene, C=C, sp^2), around 1580 cm^{-1}
 - (4) $I_D/I_G < 1$, though I_G less \rightarrow electron rich regions exist \rightarrow beneficial to adsorption



Conclusions and Future

Conclusions and Future works

- Morphology → fiber structure creates forces to assist adsorption
- Specific surface area → large enough to perform adsorption
- Pore size distribution → similar diameter to adsorbates of interest
- Surface properties → electrons on surface assist adsorption
- LACF is a potential adsorbent!
- Future works: adsorption column test
 - study adsorption kinetics (metal ions)

Thank you