

Fabrication of Porous Fe₂O₃/PTFE Nanofiber Membranes and Their Application as Catalyst for Dye Degradation

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TG analysis

Thermal stability PTFE/PVA/BA nanofiber membrane of was investigated by TG/DTA 6300 thermogravimetric analyzer with 2 mg sample at the heat rate of 10 °C/min in the range of 25-500 °C.

XRD analysis

The changes of the crystal phase of PTFE nanofiber membrane were characterized with D/MAX-2500 X-ray diffractometer (XRD, Japan Rigaku).

BET analysis

The Brunauer-Emmet-Teller specific surface area of the as-spun PTFE/PVA/BA nanofiber membrane and the Fe₂O₃/PTFE nanofiber membranes were determined using a high-speed automated area and pore size analyzer (3H-2000PS1, Beijing).



Fig. S1 The photo of the electrospinning device

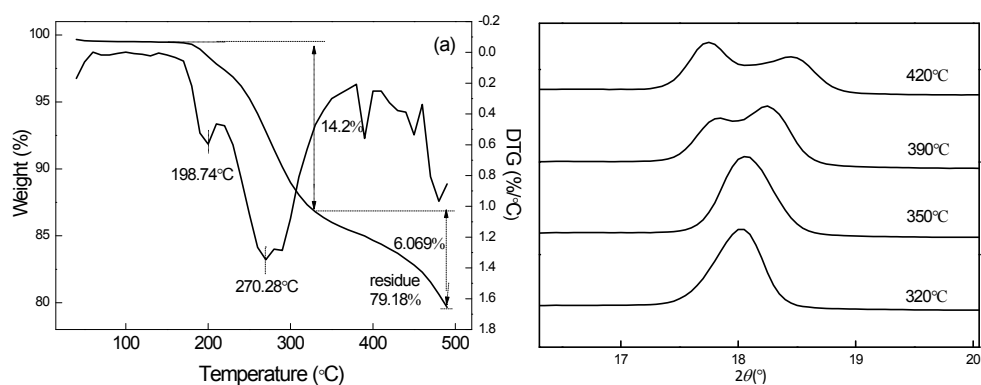


Fig. S2 The TG curve of and XRD patterns of PTFE/PVA/BA nanofiber membrane

Discussion of Fig. S2 The peak at 198.74 °C was attributed to the decomposition of BA and the peak at 270.28 °C was due to the decomposition of PVA. The PTFE nanofiber membrane was obtained via calcining the as-spun PTFE/PVA/BA nanofiber membrane. In the XRD patterns, the crystal structure changed due to the molecular chain motion when the calcination temperature higher than 390 °C. Then the calcination temperature was set as 390 °C.

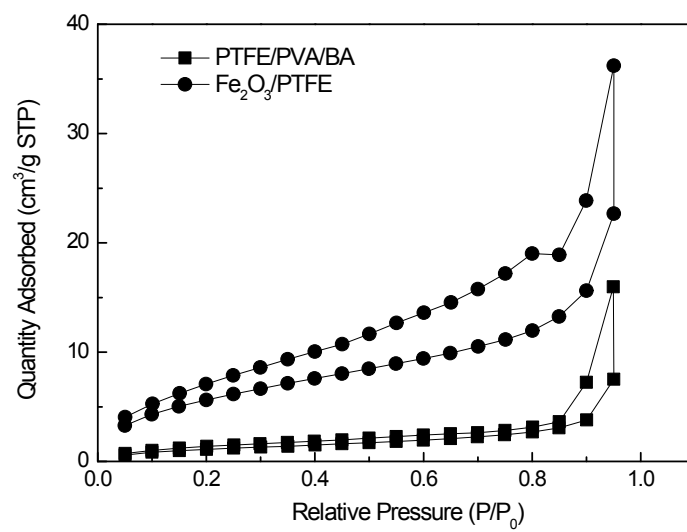


Fig. S3 The N₂ adsorption isotherms at 77 K of samples