Supporting Information

One-pot construction of Cu and O co-doped porous g-C₃N₄ with enhanced

photocatalytic performance towards the degradation of levofloxacin

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Experiment

The FT-IR spectrum of CN and Cu/O-PCN-3 were identified on a Fourier transform infrared spectrophotometer (Avater 370, Nicolet Co., USA). The sample about of 2.0 mg was evenly mixed with dried KBr powder and pressed into pellets for FT-IR determination over the wavelength range of 4000 - 400 cm⁻¹.



Fig. S1 FT-IR spectra of CN and Cu/O-PCN-3 nanomaterials.

The FT-IR spectra of CN and Cu/O-PCN-3 are given in Fig. S1. For CN and Cu/O-PCN-3 samples, a series of peaks ranging from 1200 to 1650 cm⁻¹ are attributed to the typical stretching vibration of CN heterocycles, while the sharp peak located at 812 cm⁻¹ is assigned to the vibrational mode of repeating triazine units, indicating that heptazine units are retained after Cu and O co-doping.¹ Compared with CN, an extra peak at 2144 cm⁻¹ corresponds to the nitrile groups appeared in Cu/O-PCN-3,² which maybe result from the incomplete polymerization when adding Cu(NO₃)₂·3H₂O. Obviously, the peaks appears at about 576 and 630 cm⁻¹ corresponds to the stretching vibration of Cu (II)-O and Cu (I)-O bonds are not detected in Cu/O-PCN-3. The result of FT-IR further confirms that Cu was ion-doped into the g-C₃N₄.³

Reference

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