**Supporting Information** 



Fig. S1 XRD (A) and FT-IR (B) spectra of the PCN and SC-PCN.

Table S1 BET surface areas and corresponding pore sizes of different synthetic processes
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Samples	Processing step	BET Surface Area (m <sup>2</sup> g <sup>-1</sup> )	Pore Size (nm)
1	Primary PCN	171.7	7.3
2	PCN treated with $HNO_3$	74.2	8.5
3	Primary SC-PCN	222.3	9.6
4	SC-PCN treated with $HNO_3$	177.8	8.8
5	SC-PCN with HNO <sub>3</sub> pretreatment	178.0	8.9

Table S2 Organic elemental analysis of  $g-C_3N_4$  with different treating processes

Sample	C(wt%)	N(wt%)	O(wt%)	H(wt%)
Primary PCN	34.51	59.42	4.39	1.68
Primary SC-PCN	33.92	58.60	5.67	1.81
SC-PCN treated with HNO <sub>3</sub>	30.67	55.08	11.94	2.31



**Fig. S2** (A) Fluorescence spectra of excitation and emission PCN. (B) UV-vis absorption spectra of  $HNO_3$  and TNP in different pH values. (C) Fluorescence intensities of PCN in different pH values.



**Fig. S3** (A) Fluorescence determination of TNP by PCN in pH 3 with linearity range from 0.01 to 54  $\mu$ M at excitation wavelength of 300, 330 and 350 nm. (B) Fluorescence determination of TNP by PCN in different pH solutions (pH=3, 4 and 7) with linearity range from 0.01 to 54  $\mu$ M with excitation wavelength of 350 nm.



**Fig. S4** (A) The fluorescence spectra of SC-PCN in the presence of different concentrations of TNP in pH 7 (0.0001, 0.001, 0.01, 0.05, 0.1, 0.5, 1, 2, 4, 6, 8, 10, 20, 30, and 54  $\mu$ M ). (B) Fluorescence determination of TNP by SC-PCN in pH 7, with the magnification of determination for 0.1 nM-4  $\mu$ M as the inset. (C) Fluorescence determination of TNP by SC-PCN with linearity ranges from 4 to 54  $\mu$ M by normal SVE. (D) Fluorescence determination of TNP by SC-PCN with linearity ranges from 0.1 nM to 4  $\mu$ M by double logarithmic fitting of *Ig[(F<sub>0</sub>/F)-1]* and *IgC<sub>TNP</sub>*.



**Fig. S5** Fluorescence response of the SC-PCN in the presence of 10  $\mu$ M TNP and other co-existing metal ions. *F*<sub>0</sub> and *F* represent the fluorescence intensity of SC-PCN in the absence and presence of TNP and other metal ions.

		Pseudo-second-order kinetics					
Adsorbent	Temperature	Equation	Qe	k <sub>2</sub>	R <sup>2</sup>		
		- 4	(mg g⁻¹)	(g mg <sup>-1</sup> min <sup>-1</sup> )			
	room	v-0.0152v+0.00150	65.49	0.15	0.997		
3C-PCN	temperature	y=0.0133X+0.00139					
DCN	room	v-0.0170v+0.0414	EE 00	0.0077	0.009		
PCN	temperature	y-0.01/9X+0.0414	55.99	0.0077	0.998		

Table S3 Parameters of adsorption	h kinetics fits for TNP	adsorption
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## Table S4 Parameters of Henry, Freundlich fits for TNP adsorption

					Henry- Freundlich					
Sample	He	Henry		Freundlich		Henry (0-0.87 μM)		Freundlich (0.87-44 μM)		
	K <sub>h</sub> (L g <sup>-1</sup> )	R <sup>2</sup>	K <sub>F</sub> (L g <sup>-1</sup> )	N	R <sup>2</sup>	K <sub>h</sub> (L g <sup>-1</sup> )	R <sup>2</sup>	K <sub>F</sub> (L g <sup>-1</sup> )	N	R <sup>2</sup>
SC-PCN	4.3661	0.964	8.68	1.19	0.9776	18.611	0.9999	10.77	1.58	0.9957
PCN	3.1399	0.9961	3.88	1.06	0.9987					



Fig. S6 Zeta potential measurements in different pH solutions by PCN and SC-PCN.

Water samples	Concentration added (µM)	Concentration found (μM)	Recovery (%) <sup>a</sup>	RSD (%)		
Comple 1	0.02	0.0236	118.0	10.03		
Sample 1	1	0.946	94.6	5.62		
	10	10.080	100.8	2.39		
Comple 2	0.02	0.0231	115.5	11.18		
Sample 2	1	1.051	105.1	5.75		
	10	10.020	100.2	2.48		
<sup>a</sup> Recovery (%)= 100 × (concentration found/concentration added).						

Table S5 Determination of TNP spiked in water samples (n=6)

Table S6 Performance comparison with other reported methods for TNP sensing

Methods	Linear detection	Detection	Water samples	Reference	
Wethous	range	range limit		nererenee	
Silicon nanoparticles(SiNPs)	0.087 523 μM	29 nM	River and tap water	7	
Chemically oxidized and liquid	0 0.5 μM	9.2 mM	Lake water and sea	17	
exfoliated g-C <sub>3</sub> N <sub>4</sub> nanosheets	0.5 10 μM	0.2 1111	water		
Terbium-doped blue	500 mM 100 mM	200 - 14	Running water and	10	
carbon dots	500 hivi 100 μivi	200 110	lake water	10	
Fluorescence quantum dots					
ZnS:Mn <sup>2+</sup> @allyl mercaptan	0.22 35 μM	3.4nM	Lake water	S1	
(QDs@AM)					
Cd-Metal–Organic	0 40 μM	1.3µM		9	
Fluorescent method with	0.218 305 μM	0.141 μM		S2	

8-hydroxyquinoline Aluminum nanospheres				
Fluorescent method with MoS <sub>2</sub> quantum dots	0.099 36.5 μM	95 nM	Lake water	6
Carbon dots	0 30 μM	75.6 nM		S3
Fluorescent method with phosphonated pyrene derivatives	0 35 μM	61nM		S4
Luminescent graphene quantum dots (GQDs)	1 60 μM	0.3 μM	Lake water	S5
Zn(II)/Cd(II) mixed ligand coordination polymers (CPs)	0 0.2 mM	0.06 µM		S6
Co(II)/Cd(II) Metal Organic Frameworks		0.15µM		S7
Boron nitride quantum dots (BNQDs).	0.25 200 μM	0.14 μM	River water	S8
Rotational paper-based microfluidic chips (RPADs)	17.2 87.3 μM	0.31µM	Lake and sea water	S9
Hydrazine-substituted BODIPY probe	0 40 μM,	0.44 μM		S10
This work	0.1 nM 4 μM 4 μM 54 μM	0.04 nM	Ground water and lake water	

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