

Supporting Information

Precursor-dependent distinctive polymerization process controls optoelectronic properties in graphitic carbon nitride photocatalyst

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Table S1: XRD peak positions and calculated d-spacing values obtained for GCN-D and GCN-C after the fit.

Sample	(002)		(100)	
	2 θ (degree)	d-spacing (nm)	2 θ (degree)	d-spacing (nm)
GCN-D	27.45	0.324	13.13	0.673
GCN-C	27.46	0.324	13.23	0.668

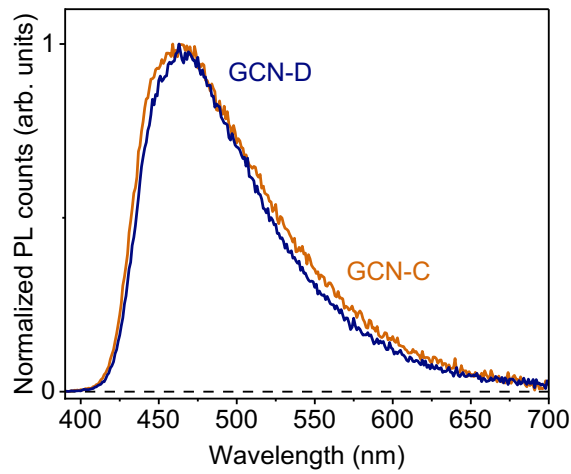


Figure S1: Steady state photoluminescence emission spectra of GCN derived from DCDA and CYN.

Table S2: Lifetime of the photoexcited charge carriers obtained from tr-PL spectra shown in Figure 1d.

Sample	τ_1 (ns)	A_1	τ_2 (ns)	A_2
GCN-D	1.176 ± 0.021	5.287 ± 0.133	4.927 ± 0.155	0.426 ± 0.024
GCN-C	1.222 ± 0.021	4.925 ± 0.112	5.001 ± 0.161	0.407 ± 0.024

The two-exponential function,

$$y_0 + A_1 \times \exp\left(\frac{-x}{\tau_1}\right) + A_2 \times \exp\left(\frac{-x}{\tau_2}\right) \quad (1)$$

was used to fit the tr-PL decay profiles.

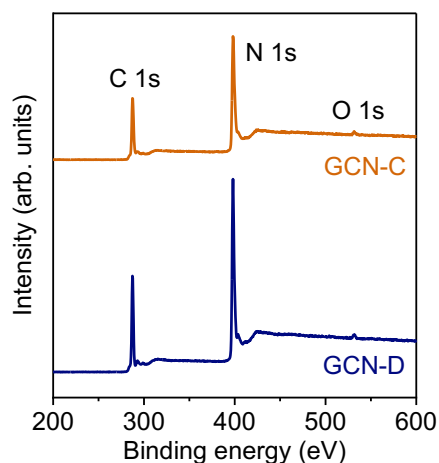


Figure S2: Survey spectra of synthesized samples from XPS analysis.

Table S3: Peak parameters obtained after C and N 1s spectral deconvolution of GCN derived from CYN and DCDA.

a. C 1s spectrum of GCN-D and GCN-C.

	Adv. C		C-NH _x (C ₁)		N-C=N (C ₂)	
	Position (eV)	Peak area %	Position (eV)	Peak area %	Position (eV)	Peak area %
GCN-D	284.8	3	286.8	1	288.3	96
GCN-C	284.8	3	286.5	2	288.3	95

b. N 1s spectrum of GCN-D and GCN-C.

	C=N-C (N ₁)		N-(C) ₃ (N ₂)		-NH _x (N ₃)	
	Position (eV)	Peak area %	Position (eV)	Peak area %	Position (eV)	Peak area %
GCN-D	398.8	76	400.2	14	401.3	10
GCN-C	398.9	64	400.1	21	401.3	15

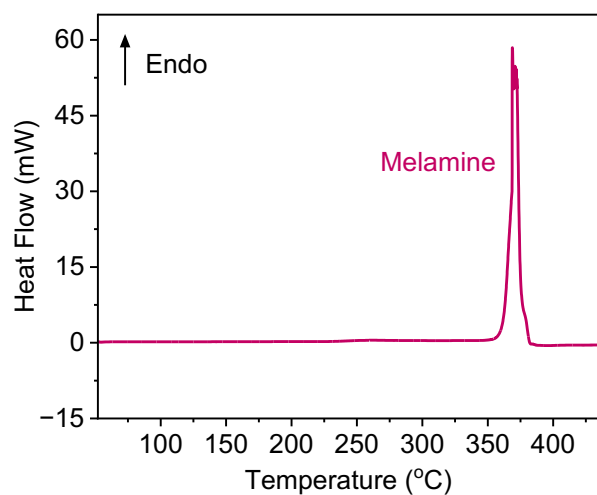


Figure S3: DSC thermogram of melamine.

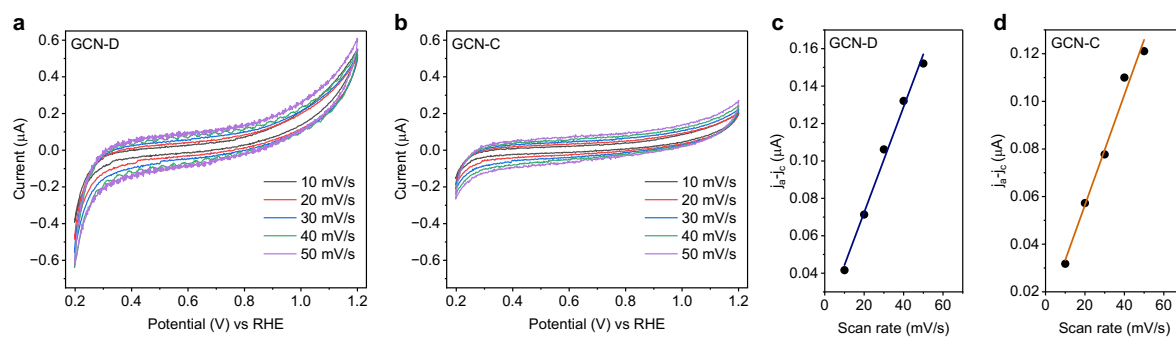


Figure S4: a,b) CV of GCN-D and GCN-C at different scan rates to measure ECSA. c,d) Plots of current and scan rate to calculate the C_{dl} value.