

# Supporting Information

## Magnetic Graphene Quantum Facilitate Dots Closed-Tube One-Step Detection of SARS-CoV-2 with Ultra-Low Field Nuclear Magnetic Resonance

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## Supplementary Figures

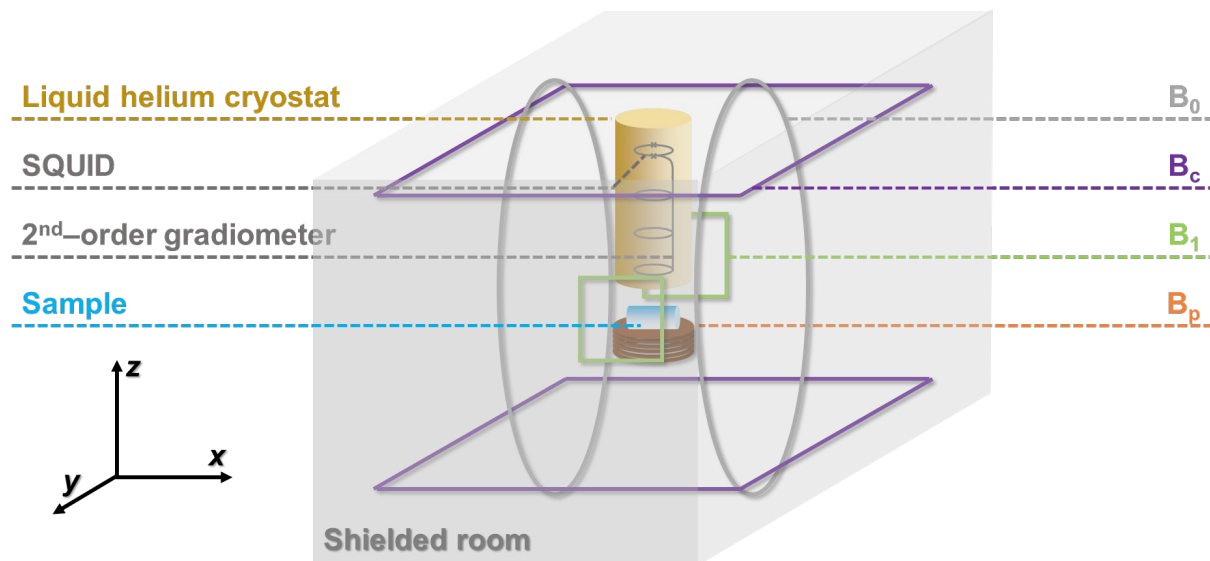


Fig. S1. Depiction of the homemade ULF NMR system.

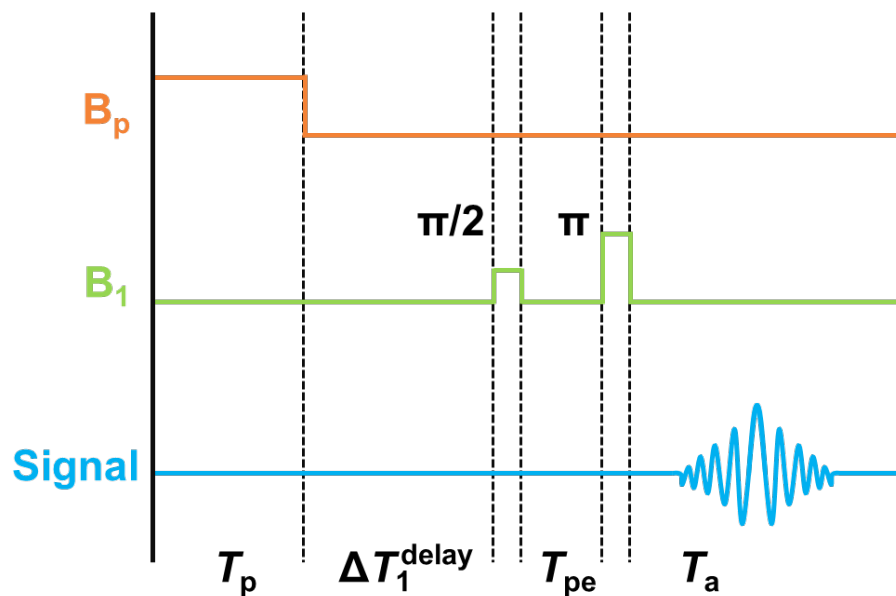
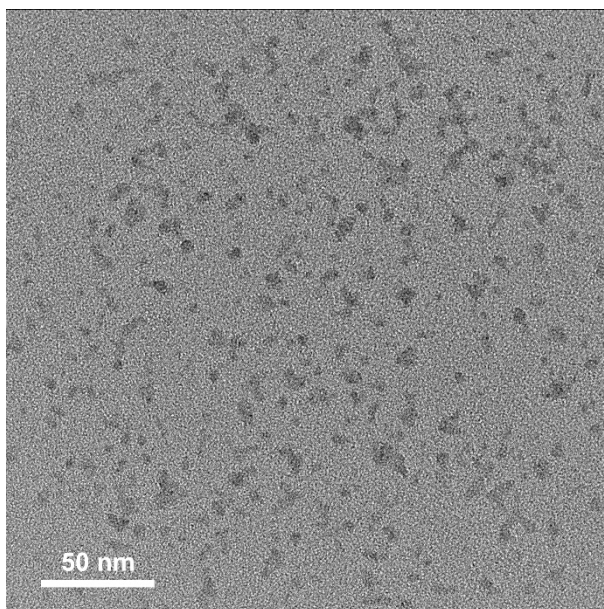
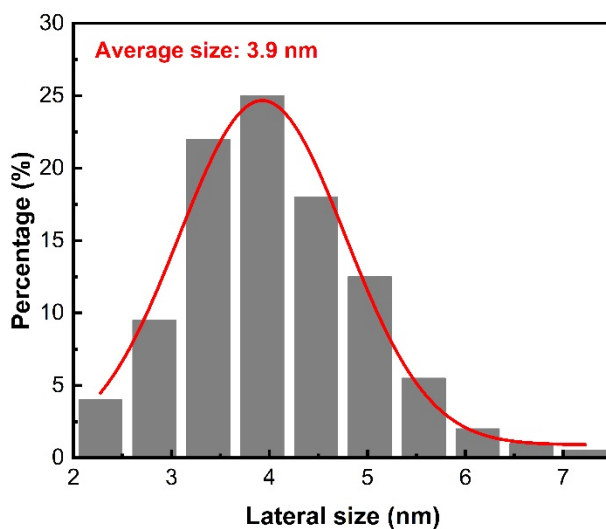


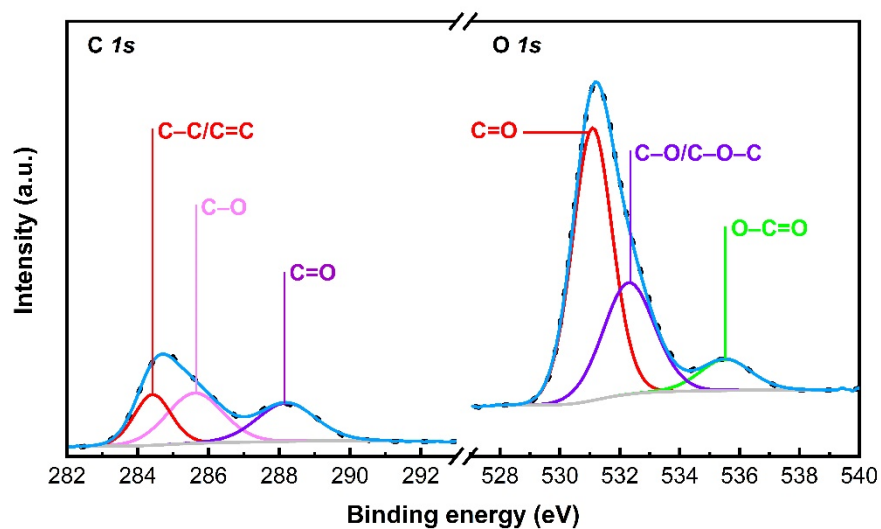
Fig. S2. Pulse sequence for  $T_1$  measurement in ULF NMR system.



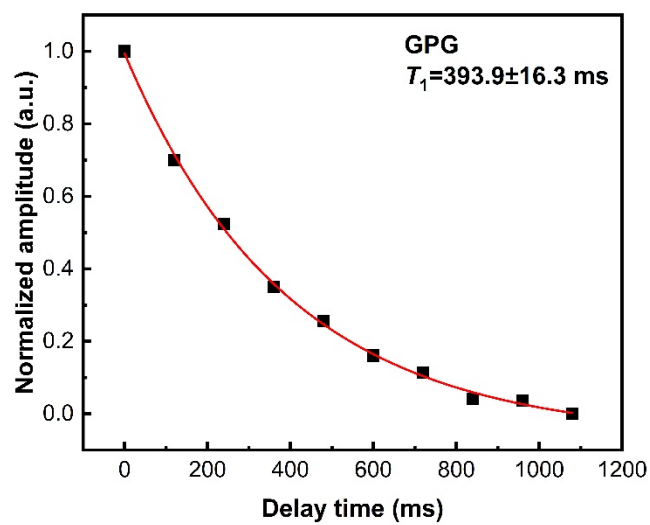
**Fig. S3.** Transmission electron microscopy (TEM) image of GQDs. The image shows that GQDs have a great dispersity, which is proven by the zeta potential measurement ( $-46.2$  mV).



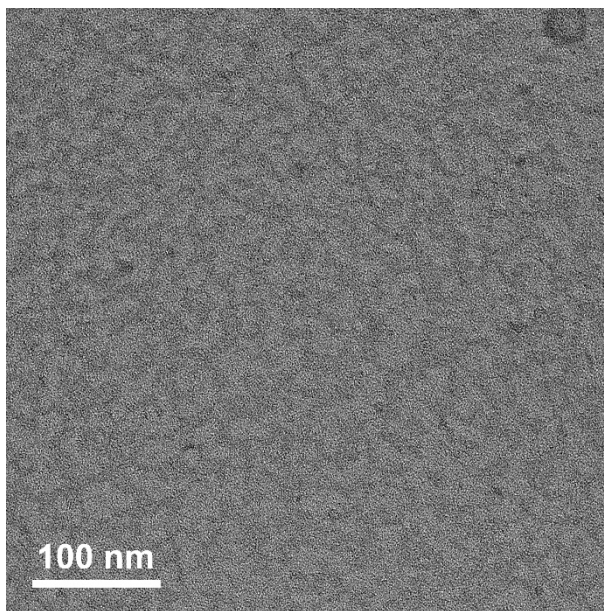
**Fig. S4.** Size distribution histogram of GQDs derived from **Fig. S3**. A Gaussian distribution fit was used to analyse the average size of GQDs. The average size of GQDs is found to be 3.9 nm.



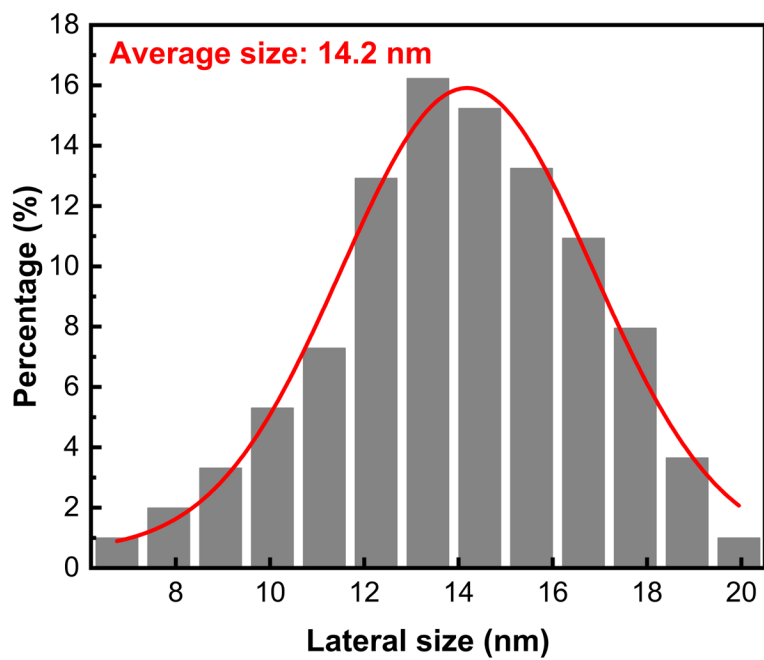
**Fig. S5.** High-resolution X-ray photoelectron spectroscopy (XPS) C *1s* and O *1s* spectra of GQDs. In the C *1s* spectrum of GQDs, peaks located at 284.4, 285.6, and 288.2 eV are recognized as C–C/C=C, C–O, and C=O bonds, respectively, while the peaks located at 531.1, 532.3, and 535.5 eV in the O *1s* spectrum can be attributed to C=O, C–O/C–O–C, and O–C=O bonds, respectively.



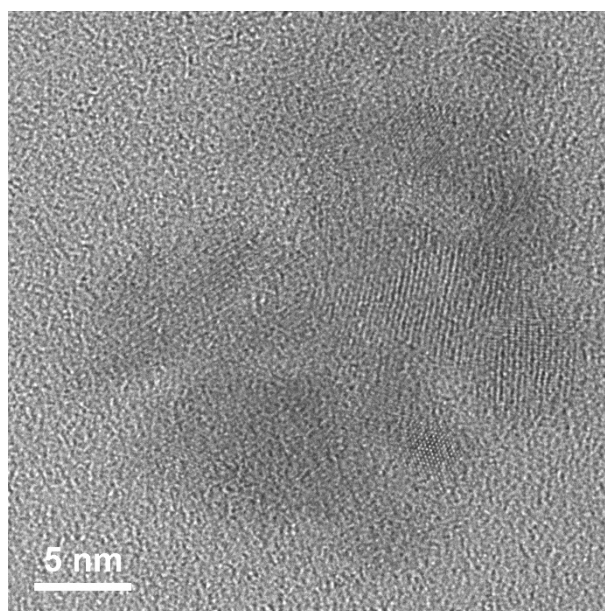
**Fig. S6.**  $T_1$  fitting of GPG in PBS buffer.



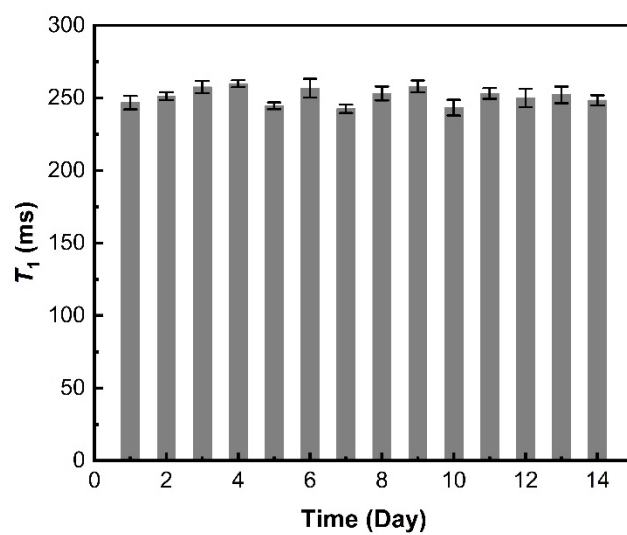
**Fig. S7.** TEM image of GPG-Abs. The scale bar is 100 nm in the image.



**Fig. S8.** Size distribution histogram of GPG-Abs. The average lateral size of GPG-Abs is 14.2 nm, which is larger than that of GPG.



**Fig. S9.** HR-TEM image of GPG-Abs. The scale bar is 5 nm in the image. GPG can be recognized as the assembly after the conjugation with Abs.



**Fig. S10.** Stability of GPG-Abs.  $T_1$ s of GPG-Ab measured at different times after the preparation.

## Supplementary Table

**Table S1.** Comparison of newly developed methods for SARS-CoV-2 detection. <sup>a)</sup>

Method	Targeting molecule	Pretreatment	Direct virus detection	Detection time	LOD	Ref.
MRSw	S protein	No	Yes	2 min	248 Particles mL <sup>-1</sup>	This work
Colorimetry	RNA	Yes	No	10 min	0.18 ng μL <sup>-1</sup>	[1]
FET	S protein	No	Yes	few min	242 copies mL <sup>-1</sup>	[2]
LSPR	RNA	Yes	No	10 min	0.22 pM	[3]
CRISPR	RNA	Yes	No	20 min	4.6 copies	[4]
LAMP	RNA	Yes	No	30–90 min	42 copies/reaction	[5]

<sup>a)</sup> Abbreviations in this table: LOD, limit of detection; MRSw: magnetic relaxation switches; FET, field-effect transistor; LSPR, localized surface plasmon resonance; CRISPR, clustered regularly interspaced short palindromic repeats; LAMP, loop-mediated isothermal amplification.

**Table S2.** The detailed costs of GPG preparation for a single test.

Category	Item	Usage	Price per unit	Cost
Material	Gd(NO <sub>3</sub> ) <sub>3</sub>	0.1 μmol	USD 2/mmol	USD 0.0002
	PEG <sub>6</sub>	0.01 mmol	USD 10/mmol	USD 0.1
	GQDs	1.5 mg	USD 300/g	USD 0.45
Power	Electricity	6 kW·h <sup>a)</sup>	USD 0.1/kW·h	USD 0.6
Storage	Vial	1	USD 0.1	USD 0.1
<b>Total cost</b>				<b>USD 1.2502</b>

<sup>a)</sup> In this work, about 300 kW·h of electricity is consumed for an entire process of hydrothermal reaction. Practically, 50 portions of the probe can be synthesized at one time, so the electricity used for synthesis is 6 kW·h per portion of the probe.



## Supplementary References

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