Supporting information

Grafting redox-active molecule on graphene oxide through a diamine linker:

length optimization for electron transfer

Rizwan Khan^{*a,b*} and Yuta Nishina^{*a,b**}

a Graduate school of natural science and technology, Okayama University, 3-1-1 Tsushimanaka, Kita-ku, Okayama 700-8530, Japan b Research Core for Interdisciplinary Sciences, Okayama University, 3-1-1 Tsushimanaka, Kita-ku, Okayama 700-8530, Japan

E-mail: nisina-y@cc.okayama-u.ac.jp.

Fable S1. Detail quar	ntitative analysis of	wide scan XPS spectra	of GO 1a, and	d GO 2a.
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Sample	at.%			
	С	0	N	
GO 1a	70.0	25.8	3.6	
GO 2a	80.0	17.1	2.9	



Fig. S1. High-resolution N 1s XPS spectra of (a) GO 1a, and (b) GO 2a



Fig. S2. XPS survey spectra of (i) GO 1b, (ii) GO 1c, (iii) GO 1d, and (iv) GO 1e.

Table S2. Detail quantitative analysis of wide scan XPS spectra of GO 1b, GO 1c, GO 1d, andGO 1e.

Sample	Element at.%		
	С	0	N
GO 1b	71.7	25.5	2.7
GO 1c	71.7	25.2	2.9
GO 1d	71.3	25.1	3.5
GO 1e	71.9	24.6	3.3



Fig. S3. High-resolution N 1s XPS spectra of (a) GO 1b, (b) GO 1c, (c) GO 1d, and (d) GO 1e.



Fig. S4. XPS survey spectra of (i) GO 2b, (ii) GO 2c, (iii) GO 2d, and (iv) GO 2e.

Table S3. Detail quantitative analysis of wide scan XPS spectra of GO 2b, GO 2c, GO 2d, and GO 2e.

Sample	Element at.%		
	С	0	Ν
GO 2b	81.3	16.1	2.5
GO 2c	79.7	17.1	2.8
GO 2d	79.1	18.1	2.8
GO 2e	78.1	19.3	2.5



Fig. S5. High-resolution N 1s XPS spectra of (a) GO 2b, (b) GO 2c, (c) GO 2d, and (d) GO 2e.





Fig. S6. SEM images of (a) GO, (b) GO 1a, and GO 2a

Fig. S7. (a) Cycling stability test of **GO 2a-e** at a current density of 10 A g⁻¹; (b) rate capability test of **GO 2a-e**.



Fig. S8. Galvanostatic charge discharge curve of GO 2a at different current density.



Fig. S9. (a) CV study of **GO 2a** before and after hydrolysis of amide bond at a scan rate of 50 mV s⁻¹, (b) FTIR analysis of **GO 2a** before and after hydrolysis of amide bond.