

Supporting Information

Fast immobilization of human carbonic anhydrase II on Ni-based metal-organic framework nanorods with high catalytic performance

Mengzhao Jiao¹, Jie He¹, Shanshan Sun¹, Frank Vriesekoop², Qipeng Yuan¹, Yanhui

Liu^{1} and Hao Liang^{1*}*

¹ State Key laboratory of Chemical Resource Engineering, Beijing University of

Chemical Technology, 100029, Beijing, P.R. China

² Department of Food Technology and Innovation, Harper Adams University,

Newport, TF10 8NB, United Kingdom.

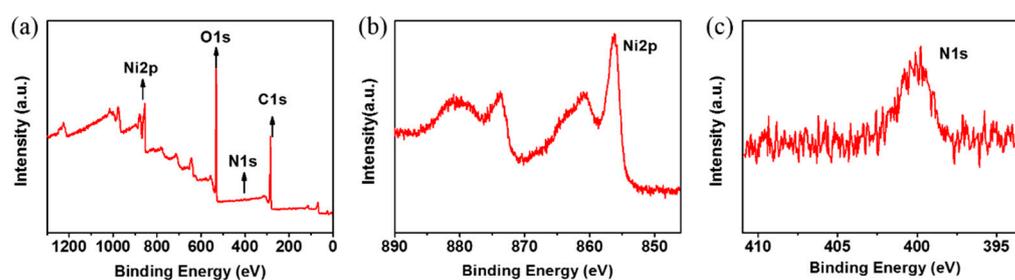
*Corresponding Author: lianghao@mail.buct.edu.cn; liuyhmail.buct.edu.cn

Table S1 The amount of immobilized His-hCA II.

Free Enzyme		Immobilized Enzyme				
Protein (mg/ml)	Specific Activity (U/mg)	Protein in washing	Bound Protein	Total Activity(U)	Specific Activity (U/mg)	Protein Yield (%)
0.4	1.96	0.273	0.127	1054.1	8296.3	31.76

Table S2 Michaelis–Menten kinetics parameters of immobilized and free enzymes

	K_m (mmol/L)	V_{max} (mmol/min)
Free enzyme	1.82	0.037
Immobilized enzyme	1.96	0.035

**Figure S1** XPS spectra of Ni-BTC nanorods: (a) full scan, (b) Ni 2p, (c) N 1s.

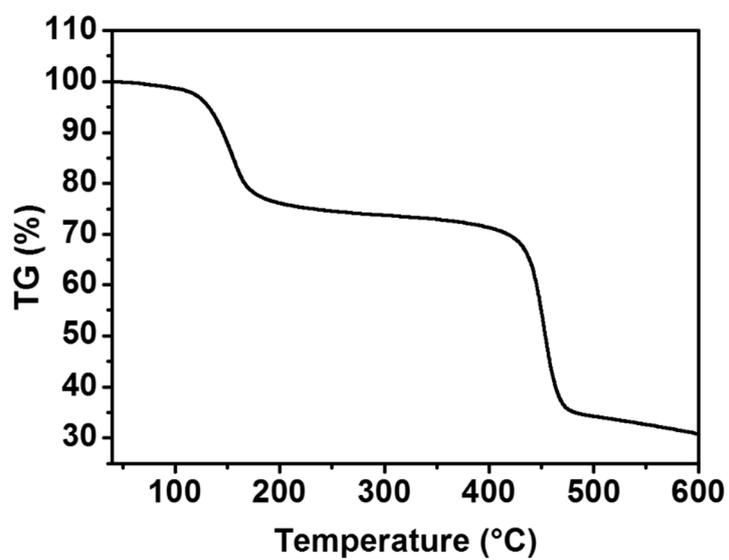


Fig. S2. TG curves of Ni-BTC.

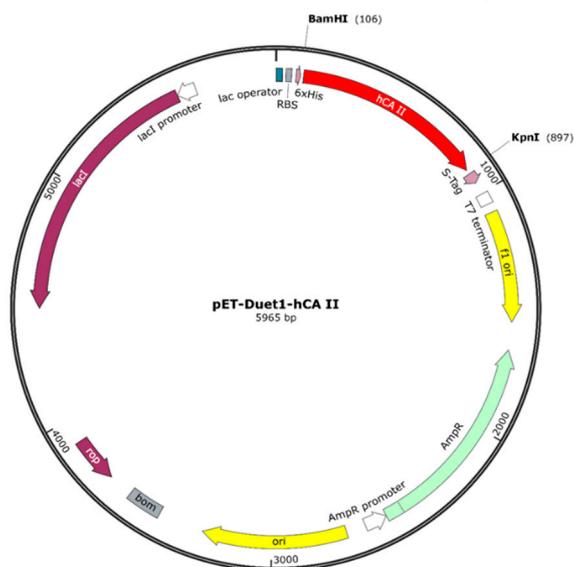


Fig. S3. The plasmid map of pETDuet-1-His-hCA II.

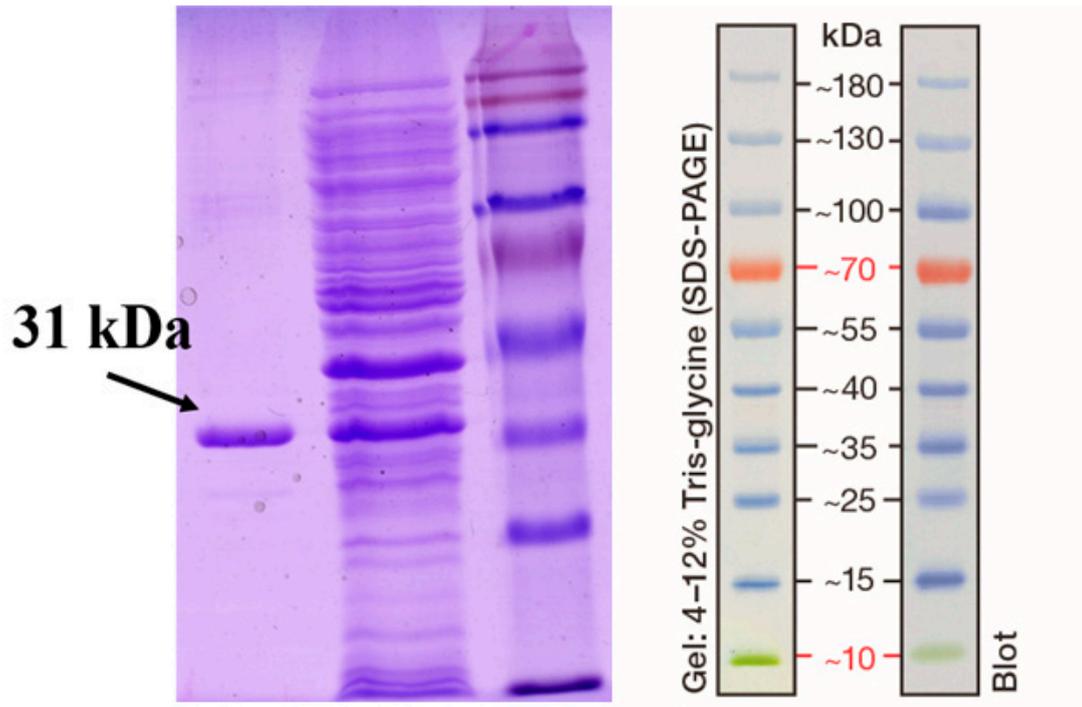


Fig. S4. SDS-PAGE of purified the recombinant hCA II from the culture supernatant. (Lane 1: His6-tagged hCA II purified using Ni-NTA column, Lane 2: supernatant of hCA II cell lysate, Lane 3: molecular weight marker)

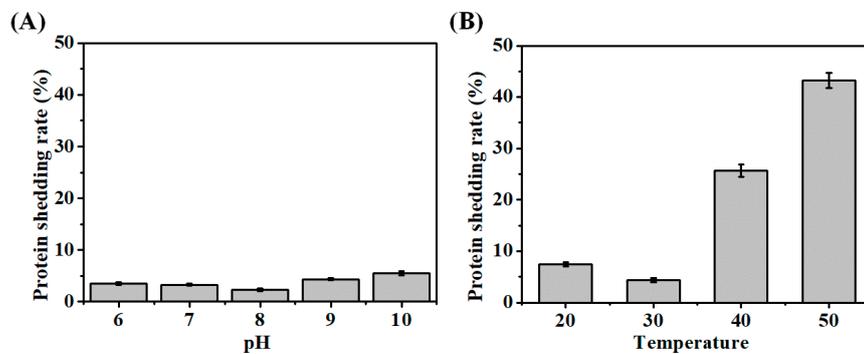


Fig. S5. (A) The protein shedding rate of immobilized enzyme at different pH and (B) the protein shedding rate of immobilized enzyme at different temperatures.

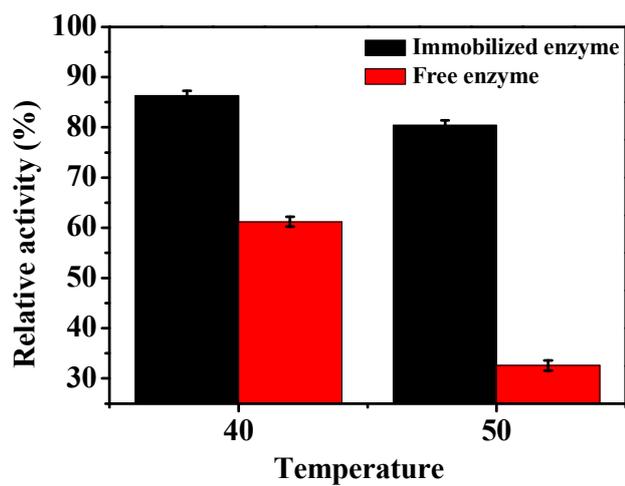


Fig. S6. The catalytic activity of His-hCA II@Ni-BTC and free enzyme at the same protein concentration was studied at 40 °C and 50 °C.

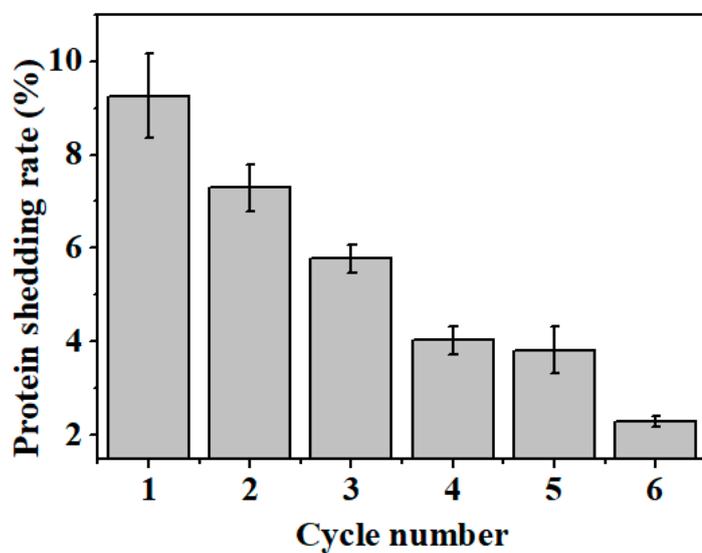


Fig. S7. The protein shedding rate of cycle process

