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## **Supplementary Material**

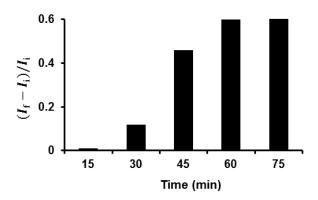
A rapid fluorescence detecting platform: applicable to sense carnitine and chloramphenicol in food samples

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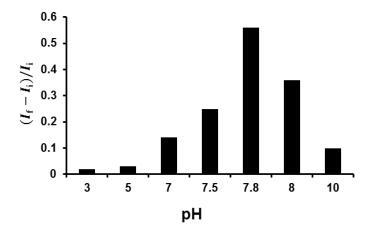
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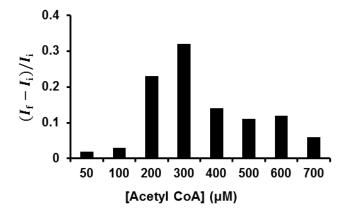
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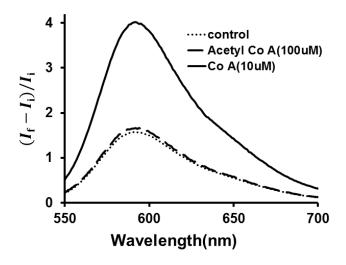
**Fig. S1** Optimization of incubation time:  $(I_f - I_i/I_i)$  vs. time (s). The experiments were carried out in 10% [v/v] DMSO in phosphate buffer (pH 7.8) containing 5  $\mu$ M BCC and 5  $\mu$ M CoA.



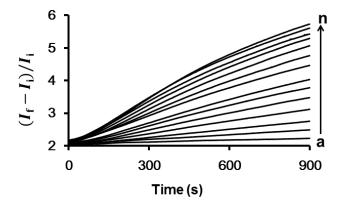
**Fig. S2.** pH Optimization plot:  $(I_f - I_i / I_i)$  versus pH. The experiments were carried out in 10% [v/v] DMSO in phosphate buffer containing 5  $\mu$ M BCC and 5  $\mu$ M CoA.



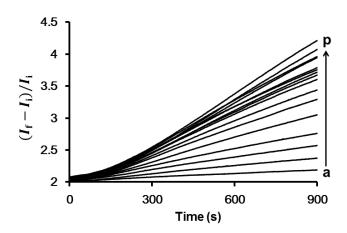
**Fig. S3.** Concentration of acetyl CoA optimization plot:  $(I_f - I_i/I_i)$  vs. [acetyl CoA] (μM). The experiments were carried out in 10% [v/v] DMSO, phosphate buffer solution (pH 7.8) containing 5 μM BCC, enzyme (0.1 unit of CrAT or 1 unit of CAT) and 5 μM of substrate (carnitine or chloramphenicol).



**Fig. S4.** Fluorescence spectra changes ( $\lambda_{ex} = 500 \text{ nm}$ ,  $\lambda_{em} = 595 \text{ nm}$ ) of BCC (5  $\mu$ M) with CoA (10  $\mu$ M), acetyl Co A and control sample. The experiments were carried out in 10% DMSO PBS (pH 7.8) (v/v).



**Fig. S5.** Kinetic plots for carnitine:  $(I_f - I_i/I_i)$  vs. time  $(0 - 600 \mu\text{M}; a=0, b=10, c-20, d=40, e= 60, f=80, g= 100, h=150, i= 200, j= 250, k= 300, l= 400, m= 500, n= 600 μM). The experiments were carried out in 10% DMSO PBS (pH 7.8) (v/v) containing 300 μM of acetyl CoA and 1 unit of CAT.$ 



**Fig. S6.** Kinetic plots for chloramphenicol:  $(I_f - I_i/I_i)$  vs. time  $(0 - 100 \,\mu\text{M}; a= 0, b= 5, c= 10, d= 20, e= 30, f= 35, g= 40, h= 45, i= 50, j= 55, k= 60, l= 65, m= 70, n= 80, o= 90, p= 100 <math>\mu$ M). The experiments were carried out in 10% DMSO PBS (pH 7.8) (v/v) containing 300  $\mu$ M of acetyl CoA and 0.1 unit of CrAT.