A novel carbazole-based mitochondria-targeted ratiometric fluorescent probe for bisulfite in living cells

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Fig.S1 The absorption spectra of **DCI** before and after reaction with bisulfite (10 equiv.) in PBS buffer (pH 7.4, 10 mM). [**DCI**] =10 μ M. λ ex = 350 nm. Slit: 5nm/5 nm.



Fig.S2 Fluorescence spectra of **DCI** before and after reaction with bisulfite (30 equiv.) in PBS buffer (pH 7.4, 10 mM). [**DCI**] =10 μ M. λ ex = 350 nm. Slit: 5nm/5 nm.



Fig. S3 The response of **CZ-Id** and **DCI** to bisulfute in PBS buffer (pH 7.4, 10 mM). For **DCI**: [**DCI**] =10 μ M, [HSO₃⁻] = 100 μ M, λ ex = 350 nm, the ratio (I₅₀₇/I₆₂₈) was achieved 10 min after addition of bisulfite; For **CZ-Id**: [**CZ-Id**] = 10 μ M, [HSO₃⁻] = 100 μ M, λ ex = 350 nm, the ratio (I₄₉₀/I₅₉₀) was achieved 10 min after addition of bisulfite.



Fig. S4 The line relationship between the fluorescent intensity ratios of **DCI** and the concentration of the bisulfite in PBS buffer (pH 7.4, 10 mM). [**DCI**] =10 μ M. λ ex = 350 nm. Slit: 5nm/5 nm.



Fig.S5 The kinetic response of the bisulfite (30 equiv.) to **DCI** in PBS buffer (pH 7.4, 10 mM). $[\mathbf{DCI}] = 10 \ \mu\text{M}. \ \lambda\text{ex} = 350 \ \text{nm}.$ Slit: 5nm/5 nm.



Fig.S6 ¹H NMR of DCI in the abstrance and presence of bisulfite.



Fig S7. HRMS spectral change of **DCI** in the presence of 15 equiv of NaHSO₃ in $CH_3OH / H_2O = 9:1$.



Fig. S8 HPLC result of **DCI** in the presence of bisulfite at different time. Waters e2695 Separations Module using Waters 2998 PDA detector equipped with a Symmetry C18 column(4.6×150 mm, 5µm). Water 20% and methanol 80% were used as eluents with a flow rate of 1ml/min. 295nm was used as wavelength.



Figure S9Effects of **DCI** on the viability of A549 Cells. The results are the mean standard deviation of three separate measurements.





