

A light-up probe for detection of adenosine in urine samples by a combination of an AIE molecule and an aptamer

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Table S1 The recipe of different concentrations of the TPE-2N+ solutions for the examination of the emission behavior of TPE fluorogen in tris buffer

Sample TPE-2N+ (μ M)	100 μ M TPE-2N+ (μ L)	Tris (μ L)
0	0	200
5	10	190
10	20	180
20	40	160
50	100	100
100	200	0

Table S2 The recipe for the investigation of the viability of (TPE-2N+) + ABA probe

curve	Tris- HCl (μ L)	100 μ M TPE (μ L)	100 μ M ABA (μ L)	1 μ M Adenosine (μ L)
a	200	0	0	0
b	180	20	0	0
c	180	20	0.2	0
d	160	20	0	20
e	160	20	0.2	20

Table S3 The recipe of different concentrations of nucleotide samples the specificity experiment

Sample nucleotides (μM)	100 μM TPE (μL)	100 μM ABA (μL)	0.001 μM nucleotides (μL)	10 mM Tris (μL)
0	20	0.2	0	180
0.00001	20	0.2	2	178
0.00005	20	0.2	10	170
0.0001	20	0.2	20	160
0.0005	20	0.2	100	80
1 μM nucleotide (μL)				
0.001	20	0.2	0.2	180
0.005	20	0.2	1	179
0.01	20	0.2	2	178
0.05	20	0.2	10	170
0.1	20	0.2	20	160

Table S4 The intra-day precision of the detection by the probe for urine samples

Adenosine Added (μM)	sample1 Found (μM)	sample2 Found (μM)	sample3 Found (μM)	RSD (% , n=3)
0.01	0.0107	0.0024	0.0147	2.0
0.05	0.0472	0.0371	0.0448	1.5
0.1	0.0762	0.0885	0.0933	2.3

Table S5 The inter-day precision of the detection by the probe for urine samples

Adenosine Added (μM)	sample1 Found (μM)	sample2 Found (μM)	sample3 Found (μM)	RSD (% , n=3)
0.01	0.0069	0.0160	0.0036	2.1
0.05	0.0434	0.0481	0.0399	1.2
0.1	0.0986	0.0870	0.0916	1.5