

Tables:

Table 1: Effect of the interfering agents concentration on the selectivity of the NRD photo probe for TYR

Interfering agents			
Materials	ΔF_1 (380 nm) Quenching	ΔF_2 (610 nm) Enhancement	Recovery \pm RSD %
Arginine	4	5	99.2 ± 2.66
Cysteine	4	3	99.5 ± 2.87
Glycine	2	4	98.3 ± 3.10
Histidine	6	4	98.6 ± 2.12
Lysine	5	2	99.8 ± 2.11
Phenylalanine	4	2	99.34 ± 2.98
Proline	3	6	99.45 ± 1.98
Serine	3	5	99.78 ± 2.70
Threonine	2	3	97.4 ± 4.20
Tryptophan	15	14	99.7 ± 2.12
Valine	2	3	99.4 ± 2.00
Glucose	3	4	99.8 ± 0.78
protein	2	2	99.8 ± 1.78

ΔF_1 is the difference between the fluorescence intensity of the $\text{NRD}^+\text{-TRY}^-$ ion pair in acetonitrile in absence and in the presence of interfering species at $\lambda_{\text{em}} = 380$ nm.

ΔF_2 is the difference between the fluorescence intensity of the $\text{NRD}^+\text{-TRY}^-$ ion pair in acetonitrile in absence and in the presence of interfering species at $\lambda_{\text{em}} = 610$ nm.

Table 2. Sensitivity and regression parameters for photo probe.

Parameter	values
λ_{em} , nm	610
Linear range, mol L ⁻¹	$1 \times 10^{-5} - 5 \times 10^{-9}$
Limit of detection (LOD), mol L ⁻¹	2.7×10^{-9}
Limit of quantification (LOQ), mol L ⁻¹	8.9×10^{-9}
Regression equation, Y*	
Intercept (a)	15.05
Slope (b)	6.4×10^9
Standard deviation	0.053
Variance (Sa ²)	2.8×10^{-3}
Regression coefficient (r)	0.993

*Y=a+bX, Where Y is luminescence intensity, X is concentration in n mol L⁻¹, a is intercept, b is slope.

Table 3: Comparison of NRD photo probe method with different methods of the determination of (TYR)

Sample (TYR)	Method	Linear range (mol L ⁻¹)	Detection limit (mol L ⁻¹)	References
Compound amino acid injection	Microcolumn electrophoresis	0.60–90 mol L ⁻¹	0.20 mol L ⁻¹	[21]
Serum	UV spectrophotometrically	0–64 mol L ⁻¹	5 mol L ⁻¹	[22]
Urine	HPLC–APCI–MS/MS	0.03–10 mol L ⁻¹	0.025 mol L ⁻¹	[23]
Pharmaceutical	Cyclic voltammetry	1×10 ⁻⁷ to 5×10 ⁻⁵ mol L ⁻¹	8×10 ⁻⁸ mol L ⁻¹	[24]
Serum and urine samples	Spectrofluorimetry	1.1×10 ⁻⁷ to 1.1×10 ⁻⁵ mol L ⁻¹	6.8×10 ⁻⁸ mol L ⁻¹	[25]
Spirulina and food samples	Spectrofluorimetry	0.3–20.0 gm L ⁻¹	0.094 gm L ⁻¹	[26]
Pharmaceutical and serum samples	NRD Photo Probe	1 x 10 ⁻⁵ – 5 x 10 ⁻⁹ mol L ⁻¹	2.7 X 10 ⁻⁹ mol L ⁻¹	Present work

Table 4. Evaluation of intra-day and inter-day accuracy and precision.

Method	TYR taken*	Intra-day accuracy and precision (n=3)			Inter-day accuracy and precision (n=3)		
		TYR Average Found* ±CL	%RE	%RSD	TYR average found* ±CL	%RE	%RSD
Tyrosine, USA	3.0	3.07 ± 0.07	2.33	0.007	3.11± 0.27	2.75	0.036
	6.0	5.97 ± 0.05	0.50	0.006	6.08± 0.19	1.33	0.013
	9.0	8.92 ± 0.09	0.88	0.002	9.09± 0.22	1.00	0.010
Serum sample	4.0	3.99 ± 0.20	0.25	0.02	4.08± 0.20	2.00	0.020
	6.0	5.98 ± 0.15	0.33	0.01	6.09± 0.22	1.50	0.015
	9.0	8.97 ± 0.22	0.33	0.01	9.12± 0.29	1.33	0.013

.The values are multiplied by 10^{-7} mol L⁻¹ for method*

RE. Percent relative error, %RSD. relative standard deviation and CL. Confidence limits were calculated % from: $CL = \pm tS/\sqrt{n}$. (The tabulated value of t is 4.303, at the 95% confidence level; S = standard deviation and .(n = number of measurements

Table 5: Determination of (TYR) in pharmaceutical and serum preparations using NRD^+ -TYR $^-$ photo probe.

Drug	Taken ($\times 10^{-7}$ M)	Found ($\times 10^{-7}$ M)	Average e^*	Average recovery \pm R.S.D. (%)	B.P. (LC)
Tyrosine, USA	3.0	3.08, 3.05, 3.10	3.07	102.3 \pm	99.7 \pm 0.055
	6.0	6.02, 5.94, 5.96	5.97	0.007	
	9.0	8.91, 8.92, 8.94	8.92	99.5 \pm 0.006 99.1 \pm 0.002	
Serum sample	4.0	3.94, 3.97, 4.08	3.99	99.75 \pm 0.02	99.5 \pm 0.050
	6.0	6.01, 6.05, 5.90	5.98	99.66 \pm 0.01	
	9.0	8.91, 8.91, 9.10	8.97	99.66 \pm 0.01	

***Average of nine measurements.**