

Electronic Supplementary Information (ESI)

1. Calculation of Quantum Yield

Fluorescence quantum yields (Φ) were estimated by integrating the area under the fluorescence curves using the equation¹,

$$\phi_{\text{sample}} = \phi_{\text{ref}} \times \frac{\text{OD}_{\text{ref}} \times A_{\text{sample}} \times \epsilon_{\text{sample}}}{\text{OD}_{\text{sample}} \times A_{\text{ref}} \times \epsilon_{\text{ref}}}$$

where A was the area under the fluorescence spectral curve and OD was optical density of the compound at the excitation wavelength. Complex tris(2,2'-bipyridyl)ruthenium(II) ($\Phi = 0.042$ in water)² has been used as quantum yield standard for measuring the quantum yields of **APC** and its arsenate assembly .

2. Calculation of detection limit

To determine the detection limit, fluorescence titration of **APC** with arsenate is carried out by adding aliquots of micro-molar concentration of arsenate. From the concentration at which there was a sharp change in the fluorescence intensity multiplied with the concentration of **APC** gave the detection limit.³

3. Equations used for calculating detection limit (DL)

$$\text{DL} = C_L \times C_T$$

C_L = Conc. of **APC**; C_T = Conc. of arsenate at which fluorescence enhanced.

Thus;

$$\text{DL} = 1 \mu\text{M} \times 0.001 \mu\text{M} = 0.001 \mu\text{M} = 1 \times 10^{-9} \text{ M}$$

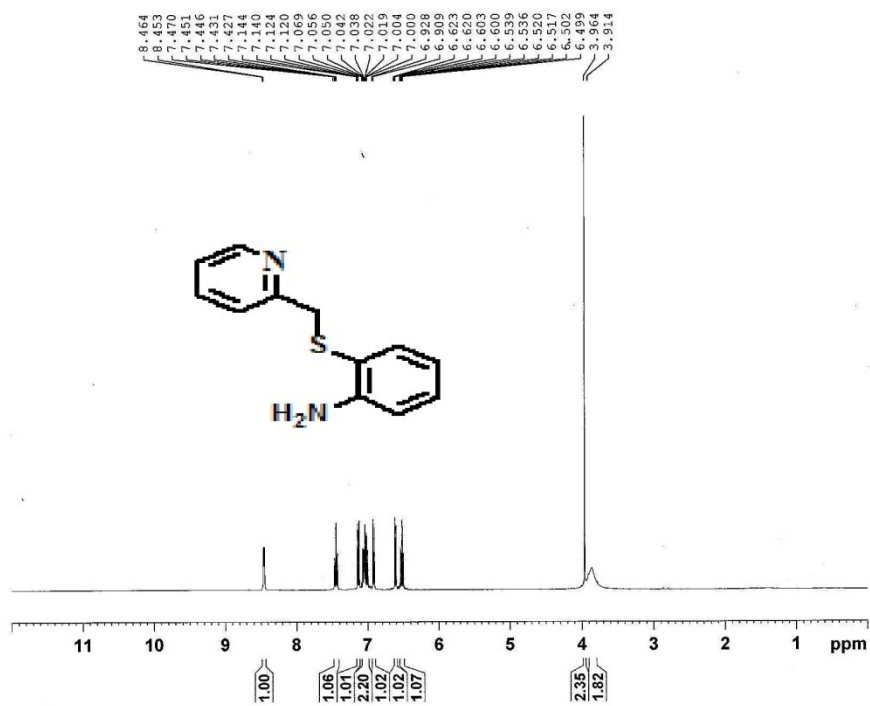


Figure S1. ¹H NMR spectrum of AP in CDCl₃

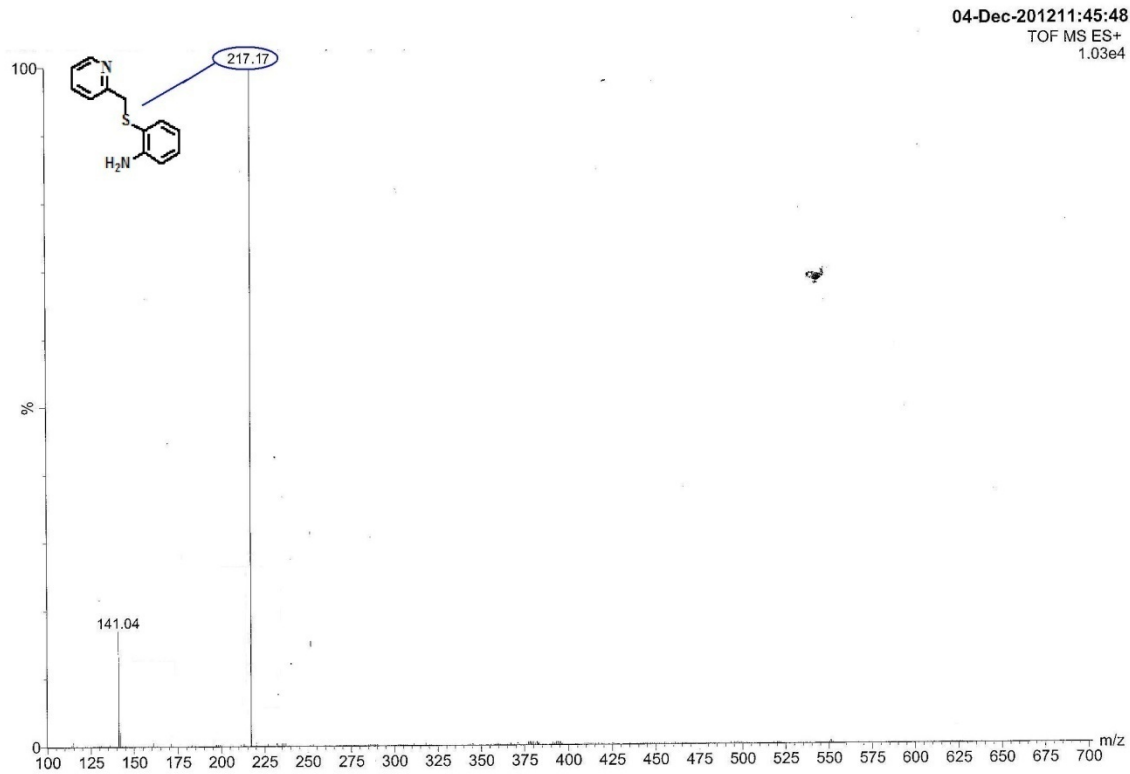


Figure S2. QTOF –MS ES⁺ spectrum of **AP**

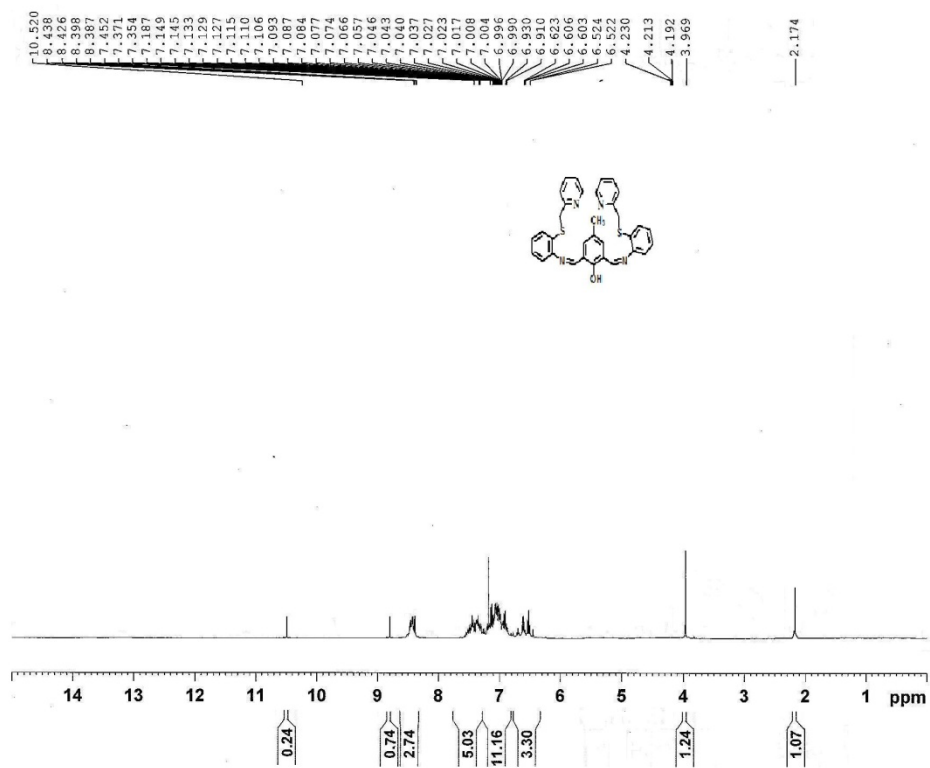


Figure S3. ¹H NMR spectrum of APC in CDCl₃

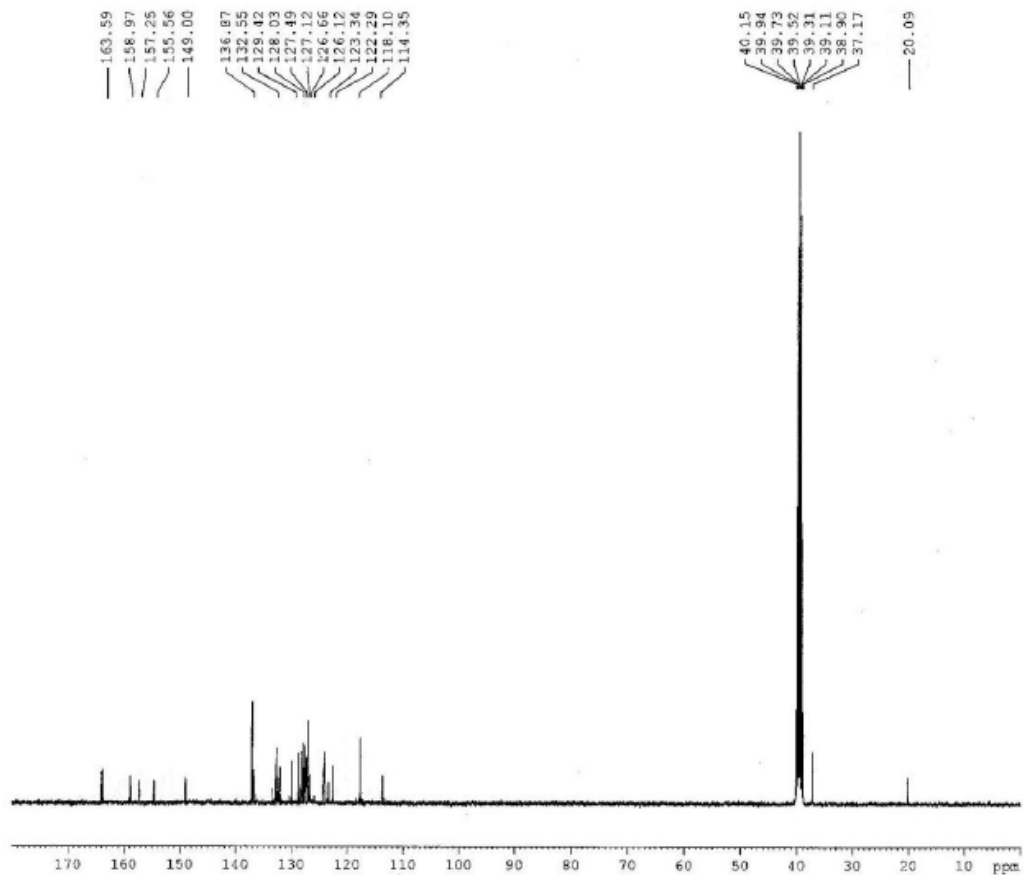


Figure S4. ^{13}C NMR spectrum of APC in DMSO-d_6

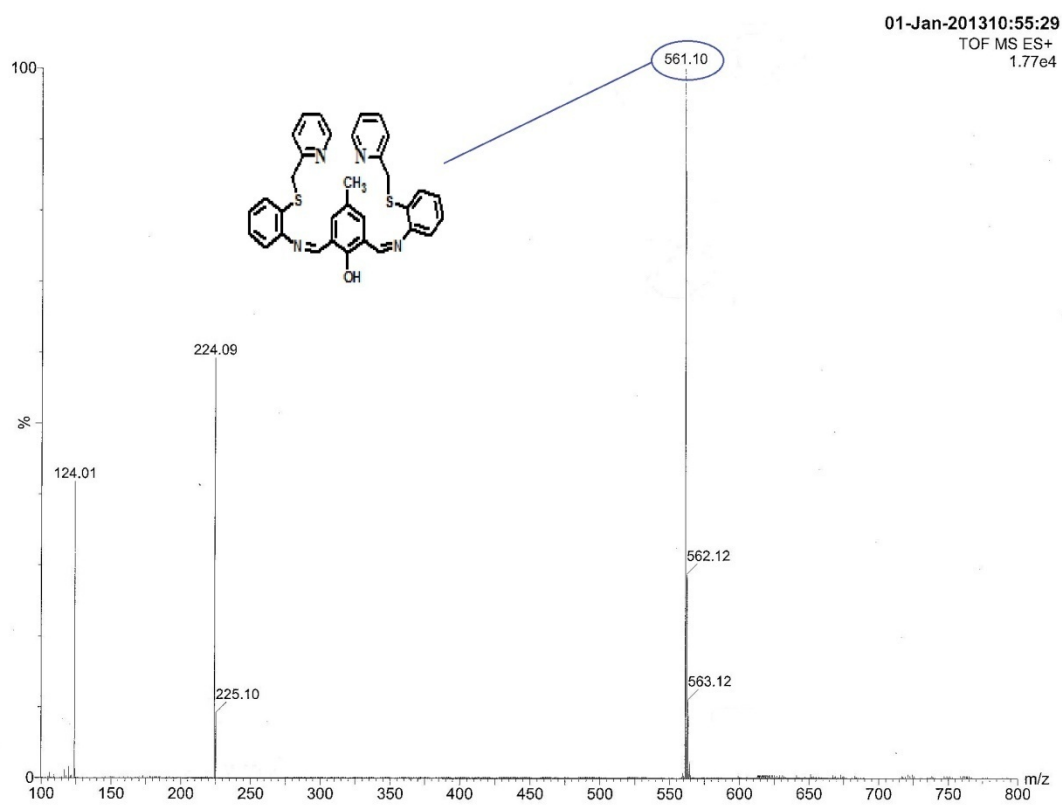


Figure S5. QTOF –MS ES⁺ spectrum of APC

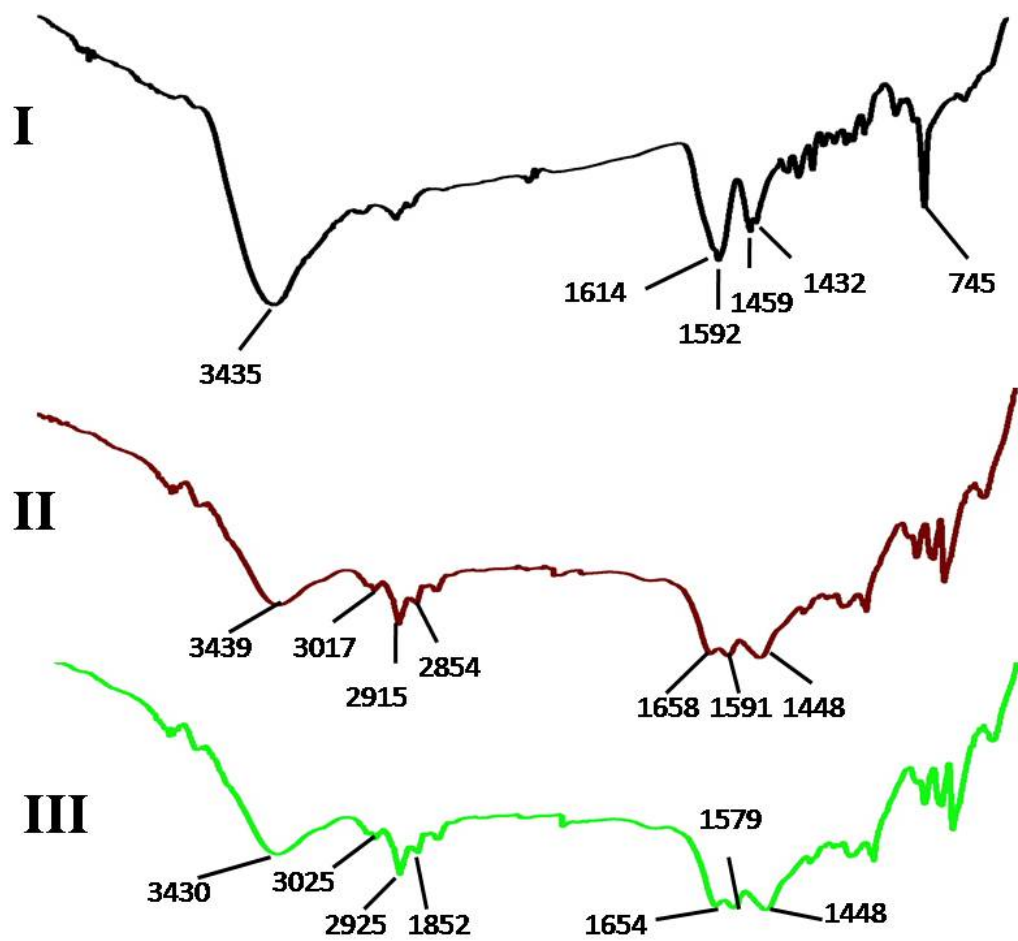


Figure S6. FTIR spectra of (I) APC, (II) APC- Merrifield polymer and (III) APC- Merrifield polymer + H₂AsO₄⁻.

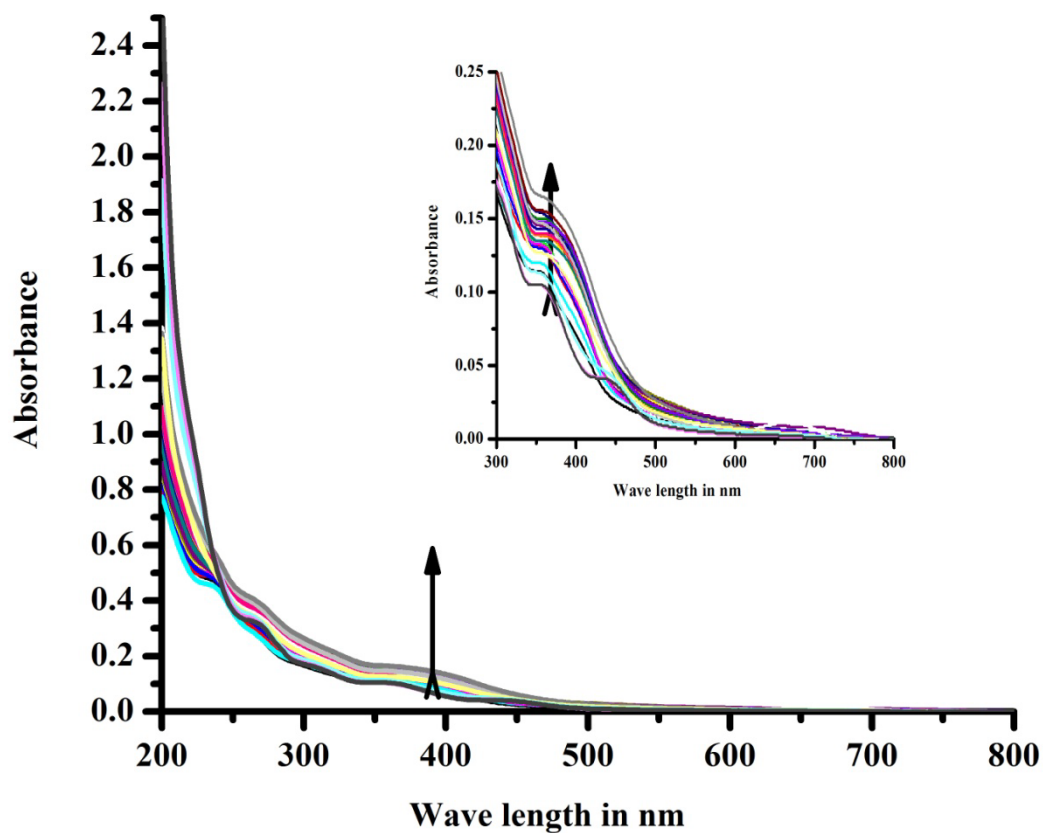


Figure S7. Changes in the absorbance of APC (10 μM) in HEPES buffered (0.1 M; EtOH– H_2O , 1:99 v/v; pH 7.4) solution upon gradual addition of H_2AsO_4^- (0.010, 0.025, 0.050, 0.075, 0.100, 0.250, 0.500, 0.750 1.000, 5.000, 10.000, 20.000, 30.000, 40.000, 50.000, 60.000, 70.000, 80.000, 90.000, 100.000, 300.000, 500.000 μM).

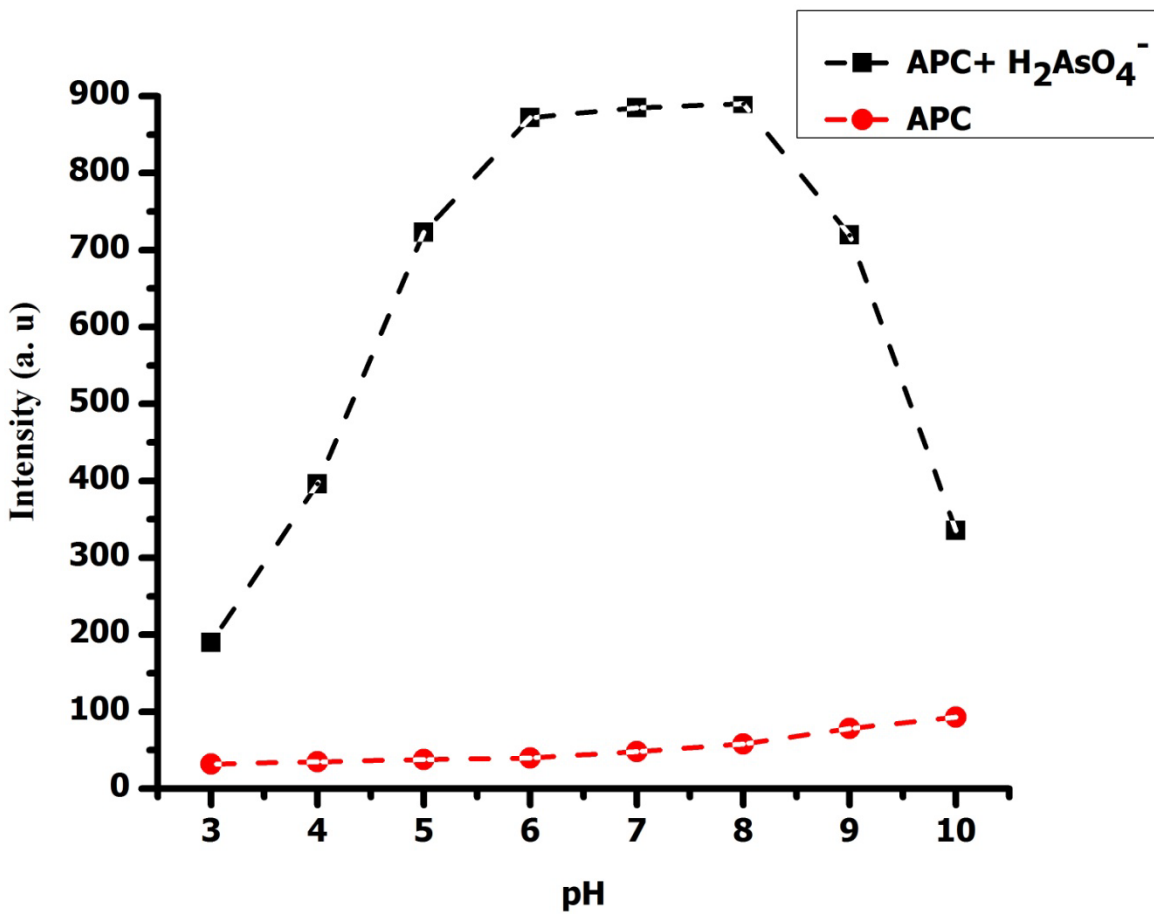


Figure S8. Influence of pH on the emission intensities of free APC (1 μ M) and its H₂AsO₄⁻ (5 μ M) adduct

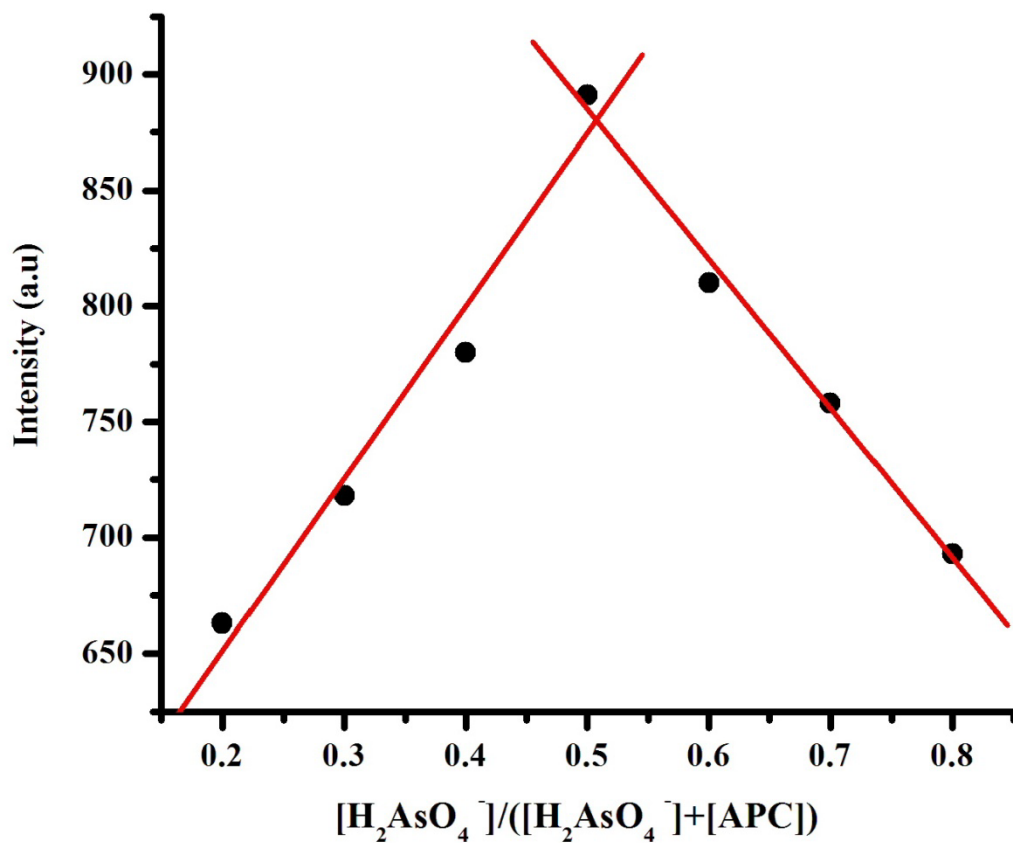


Figure S9. Job's plot for stoichiometry determination of APC- H_2AsO_4^- adduct, $\lambda_{\text{ex}} = 440\text{nm}$.

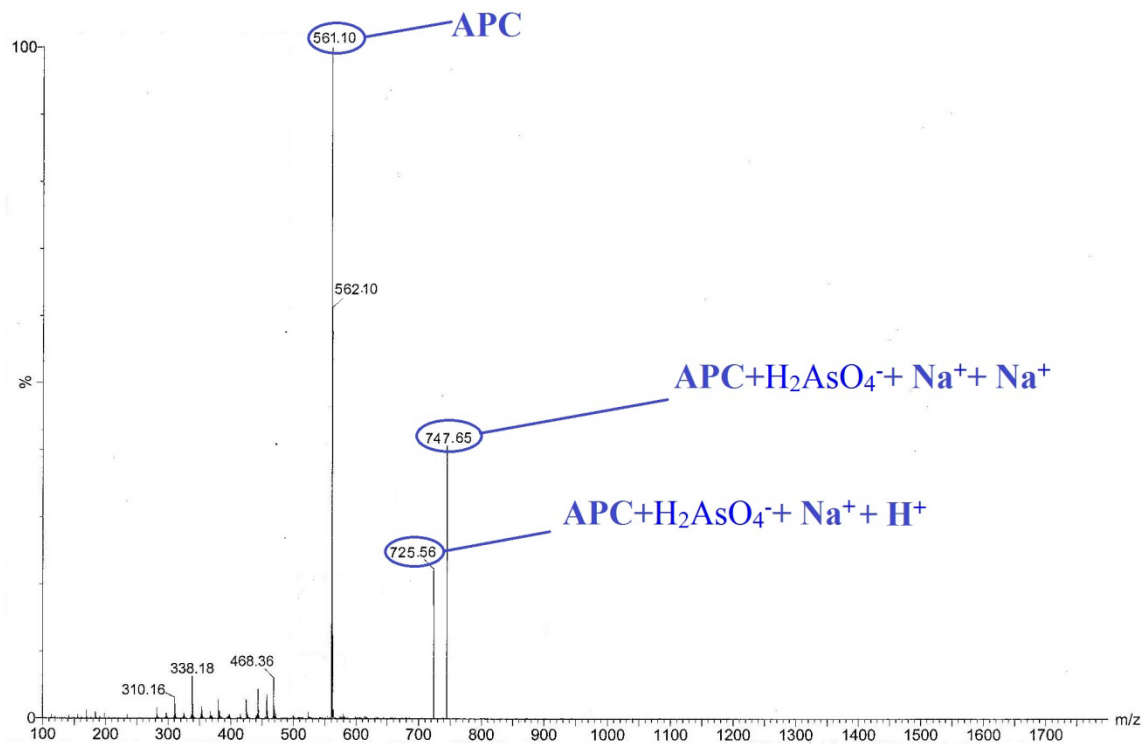


Figure S10. QTOF –MS ES⁺ spectrum of APC- H₂AsO₄⁻ adduct

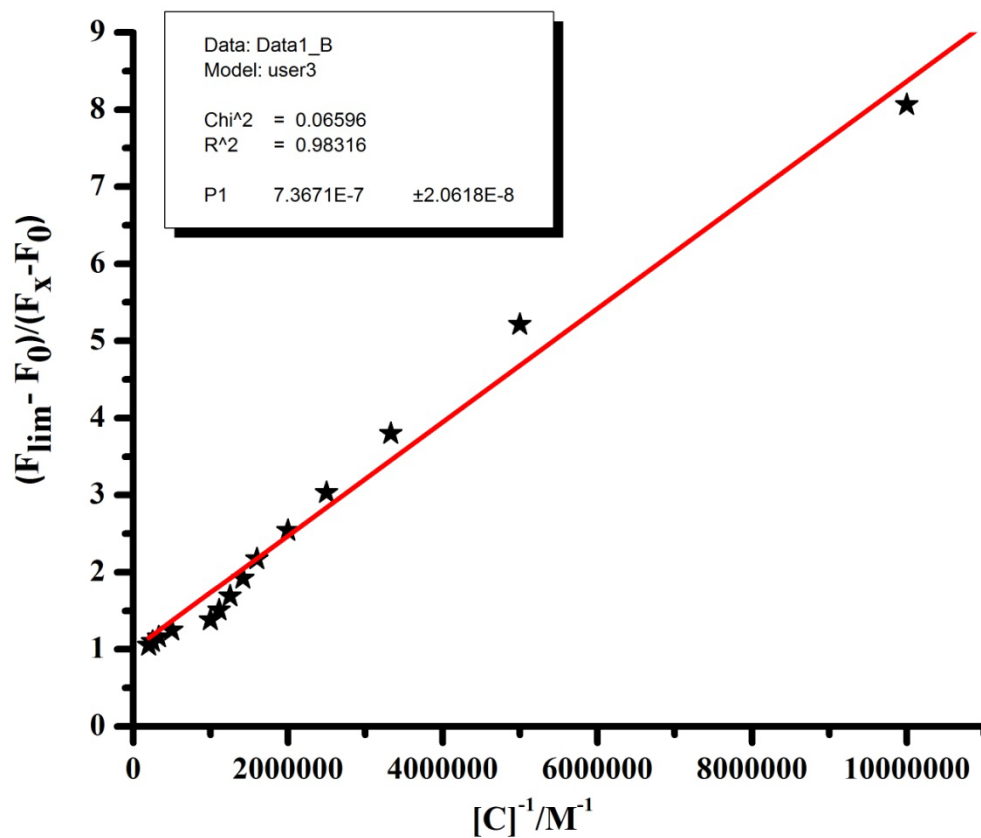


Figure S11. Estimation of binding constant of APC with $H_2AsO_4^-$ by fluorescence method

References

1. E. Austin, M. Gouterman, *Bioinorg. Chem.*, 1978, **9**, 281.
2. J. V. Houten, R. J. Watts, *J. Am. Chem. Soc.*, 1976, **98**, 4853.
3. G. L. Long, J. D. Winefordner, *Anal. Chem.*, 1983, **55**, 712A.