

A study on the use of Bathophenanthroline as a fluorescence sensor for iron(II)

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Although Fe²⁺ is present as a trace element, it plays a vital role in living systems. Therefore measurement of trace amounts of Fe²⁺ is very important for many biological and ecological studies. Bathophenanthroline (BPhen) forms a stable 1:3 red color complex with Fe²⁺ and appears as a promising colorimetric probe for Fe²⁺. Using colorimetry, it has been possible to detect 10 parts of Fe²⁺ in 10⁹ parts of water (540 nM).^{1,2} Although fluorescence is much more sensitive than absorbance, no reported attempts were made to detect Fe²⁺ with BPhen, using a fluorescence method. Therefore this study was conducted to study the capability of BPhen as a fluorescence sensor for Fe²⁺.

In this study, the fluorescence emission by BPhen was studied in 50% ethanol. At the excitation wavelength of 272 nm, BPhen gave a very high emission intensity with an emission maximum at 385 nm. A maximum fluorescence intensity was observed at a concentration of 0.15 μM, with 5 nm slit width for both excitation and emission. A 0.15 μM solution of BPhen was titrated with a 0.51 μM solution of Fe²⁺. A linear calibration curve could be obtained for this quenching process with a limit of detection of 9.0 nM of Fe²⁺. The linear range for the curve was 25 nM to 530 nM. The stoichiometry for the static quenching process was 1:3 as predicted according to literature.¹ The binding constant for the complex was calculated using the experimental data as $8.1 \times 10^{20} \text{ mol}^{-3} \text{ dm}^9$. An interference study was conducted with common fluorescence quenchers Ni²⁺, Co²⁺ and Cu²⁺ as well as Fe³⁺. The tolerance limits were 2 for Ni²⁺ and Co²⁺ while the values were 3 and 10 for Cu²⁺ and Fe³⁺, respectively, in terms of molar ratio of Fe²⁺. Further investigations should be carried on pH effect, solvent effect and temperature effect for this fluorimetric method although its sensitivity was impressive relative to the colorimetric method.

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References

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