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Reverse flow injection spectrophotometric determination of iron(III) using norfloxacin

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Abstract

A reversed flow injection colorimetric procedure for determining iron(III) at the μg level was proposed. It is based on the reaction between iron(III) with norfloxacin (NRF) in 0.07 mol l^{-1} ammonium sulfate solution, resulting in an intense yellow complex with a suitable absorption at 435 nm. Optimum conditions for determining iron(III) were investigated by univariate method. The method involved injection of a $150 \mu\text{l}$ of 0.04% w/v colorimetric reagent solution into a merged streams of sample and/or standard solution containing iron(III) and 0.07 mol l^{-1} ammonium sulfate in sulfuric acid (pH 3.5) solution which was then passed through a single bead string reactor. Subsequently the absorbance as peak height was monitored at 435 nm. Beer's law obeyed over the range of $0.2\text{--}1.4 \mu\text{g ml}^{-1}$ iron(III). The method has been applied to the determination of total iron in water samples digested with $\text{HNO}_3\text{--H}_2\text{O}_2$ (1:9 v/v). Detection limit (3σ) was $0.01 \mu\text{g ml}^{-1}$ the sample through of 86 h^{-1} and the coefficient of variation of 1.77% ($n = 12$) for $1 \mu\text{g ml}^{-1}$ Fe(III) were achieved with the recovery of the spiked Fe(III) of 92.6–99.8%.

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