



💫 🔶 🔶 Analytical Sciences		
The Japan Society for Analytical Chemistry		
Available Issues Japanese	>> <u>P</u>	ublisher Site
Author: <u>ADVANCED</u> Volume Page		
Keyword: Search		Go
Add to Favorite / Citation Add to Articles Alerts Add to Publications Alerts	ister ts	? My J-STAGE HELP

<u>TOP</u> > <u>Available Issues</u> > <u>Table of Contents</u> > Abstract

ONLINE ISSN : 1348-2246 PRINT ISSN : 0910-6340

JST Link Cen

Analytical Sciences

Vol. 26 (2010), No. 7 p.797

[PDF (500K)] [References]

Spectrophotometric Determination of Trace Phosphate Ions by Amplitude-Modulated Flow Analysis Coupled with Malachite Green Method

<u>Takeshi UEMURA</u>¹⁾, <u>Takeshi OGUSU</u>²⁾, <u>Masaki TAKEUCHI</u>³⁾ and <u>Hideji</u> TANAKA³⁾

1) Graduate School of Pharmaceutical Sciences, Tokushima University

2) Faculty of Pharmaceutical Sciences, Tokushima University

3) Institute of Health Biosciences, Tokushima University

(Received April 16, 2010) (Accepted May 6, 2010)

An amplitude-modulated flow analysis coupled with lock-in detection is proposed for the determination of trace phosphate. The flow rate F_S of sample solution is varied in response to a periodic signal V_c of alternating waveform. A coloring reagent solution (ammonium molybdate + Malachite Green) is delivered at a constant flow rate F_R . Under the constant total flow rate F_T , both solutions are merged with a diluent (water). Phosphate ion in the sample reacts with molybdate and then with Malachite Green to form green ion pairs in acidic media. Downstream, the absorbance of the mixed solution is measured at 625 nm. The output voltage V_d from the detector is sent to a lock-in amplifier, where the wave component of V_d that has the same frequency as that of V_c is distinguished from background signals. Phosphate ion can be determined from the amplitude of the component thus extracted. The calibration curve is linear ($r^2 > 0.998$) and the limit of detection (3.3 σ) is 0.17 µmol dm⁻³. Compared with a conventional flow-based method with no modulation, the present method is less susceptible to the baseline drift due mainly to the adsorption of the ion-pair on the optical window, because the quantification is based not on the

absorbance itself but on the amplitude of the absorbance. Good recoveries around 100% are obtained for the phosphate ions spiked into real water samples.

[PDF (500K)] [References]



Download Meta of Article[<u>Help</u>] <u>RIS</u> <u>BibTeX</u>

To cite this article:

Takeshi UEMURA, Takeshi OGUSU, Masaki TAKEUCHI and Hideji TANAKA, Anal. Sci., Vol. 26, p.797, (2010).

doi:10.2116/analsci.26.797 JOI JST.JSTAGE/analsci/26.797

Copyright (c) 2010 by The Japan Society for Analytical Chemistry

