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Highlights

Vitamin A is an essential micronutrient. An uncorrected vitamin A deficiency can lead to eye, skin, and reproductive disorders. Vitamin A is also being studied as a potential cancerfighting agent. Current chemical methods of measuring total vitamin A, or retinol, are complex and lengthy.

A sensitive and specific method of measuring retinol is normal phase high performance liquid chromatography with fluorescence detection. Saponification and solvent extraction of the sample converts the vitamin A esters acetate and palmitate to retinol. Then after a filtration step, the sample can be directly injected into the HPLC system as the extraction solvents are compatible with the HPLC mobile phase.

Operating Conditions

Sample preparation: Saponification, extraction, and filtration Column: Waters™ Nova-Pok™ Silica, 3.9 x 150 mm Mobile phase: 0.5% Isopropyl aicohoi in iso-octane Flow rate: 1.0 ml/min.

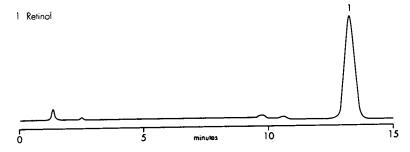
Detection: Waters 470 Scanning Fluorescence Excitation: 365 nm,

Note:

Emission: 510 nm

If interested in separating vitam:n A esters, use similar conditions without saponification step.

Vitamin A in Feed Sample



Reference

- Thorpe, V.A., J. Assoc. Off. Anal. Chem., 1990, 73(3), 463
- 2 Ball, G.F.M. Fat Soluble Vitamin Assays in Food Analysis, Elsevier Applied Science. Chapter 8 (1988)

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Ion Chromatography



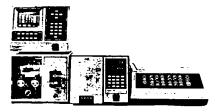
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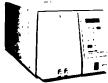
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UV/Vis: photodiode array, fixed, variable and programmable wavelength. Refractive index. Conductivity. Electrochemical. Fluorescence: fixed and programmable/scanning wavelength. Waters offers the food technologist the best choice of detectors to solve separations problems now and in the future.

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