

Application Note

Simultaneous Analysis of Vitamin A and D₃ in Vitamin Premixes and Concentrates by Convergence Chromatography/PDA Detection

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Abstract

In this application note, we report a method for analysis of vitamin A and D₃ in vitamin premixes and concentrates in one analytical run without purification or dilution. The metrological properties of the UPC² and the advantages of the method compared to the HPLC are also discussed.

Benefits

- Simultaneous determination of vitamin A and D₃ in premixes and concentrates.
- Simplified sample preparation – no purification or dilution in sample preparation.
- Direct injection of sample extract – no reconstitution.
- Much lower solvent waste compared to LC.

Introduction

Simultaneous analysis of fat-soluble vitamins in foods is challenging due to their different properties and concentrations. A typical method involves extraction and saponification of fat, followed by high performance liquid chromatography (HPLC) with UV/Vis detection. After saponification the extracts can be analyzed for vitamin A directly, but they have to be diluted due to high abundance of the vitamin A in foods and its high molar extinction. Unfortunately, the dilution makes it impossible to detect vitamin D₃ in the same solution. To measure vitamin D₃, the extracts have to be cleaned on a semi-preparative chromatograph and concentrated. For these reasons vitamins A and D₃ had to be analyzed separately. The HPLC of these compounds suffers from a long runtime, slow equilibration, and poor reproducibility.

As the extraction and saponification of separate vitamin A and D₃ methods are identical, we investigated whether it would be possible to apply Waters UltraPerformance Convergence Chromatography (UPC²) to analyze extract for vitamin A and D₃ in a single chromatographic run.

UPC² is a separation technique that uses compressed carbon dioxide as the primary mobile phase. It takes advantage of sub-2 μm particle chromatography columns, the low viscosity of CO₂, and an advanced chromatography system. This differs from traditional HPLC and improves the sensitivity of this assay. UPC² also generates much less solvent waste compared to conventional liquid chromatography.¹ In this application note, we report a method for analysis of vitamin A and D₃ in vitamin premixes and concentrates in one analytical run without purification or dilution. The metrological properties of the UPC² and the advantages of the method compared to the HPLC are also discussed.

Experimental

Sample description

Retinyl acetate (vitamin A acetate) was purchased from Sigma-Aldrich, cholecalciferol (vitamin D₃) and ergocalciferol (vitamin D₂) were purchased from the US pharmacopeia. The vitamin premix was from DSM Nutritional Products, Switzerland. Figure 1 shows the structures of relevant compounds used in this study.

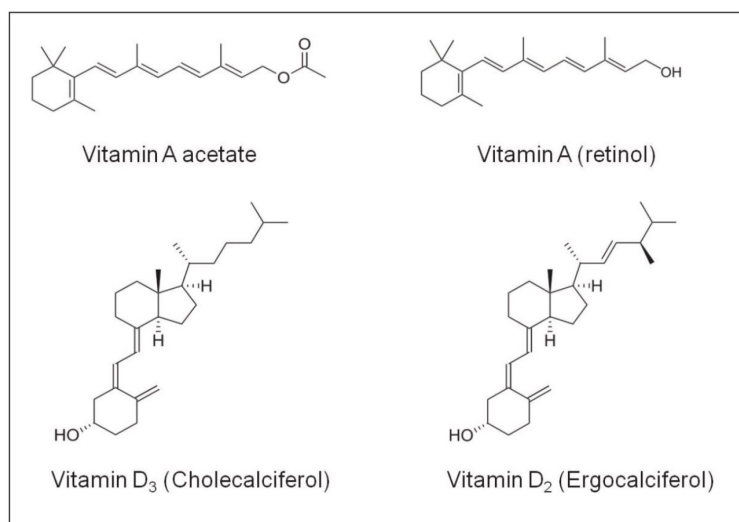


Figure 1. Structures of retinyl acetate, retinol, cholecalciferol, and ergocalciferol.

Sample preparation

1.5 g vitamin raw sample was weighed and spiked with 1 mL of vitamin D₂ solution (internal standard) into a 250-mL Erlenmeyer flask. 50 mL ($\pm 10\%$) ethanol was added, along with 10 mL of 50% KOH, and 2 mL of 33% sodium ascorbate. The samples were saponified for about 1 hour at 80 °C to 85 °C in a water bath. After saponification, the samples were cooled down to room temperature and extracted with a n-hexane diethyl ether mixture. The extract was washed with de-ionized water until neutral, and the extracts were made up to 50-mL with n-hexane. The solution was filtered through a 0.2 μm syringe filter into 2-mL autosampler vials and analyzed by UPC²/PDA.

UPC² conditions

System:	ACQUITY UPC ²
Detector:	PDA
Software:	Empower 3
Column:	ACQUITY UPC ² BEH 3.0 mm x 100 mm, 1.7 µm
Mobile phase A:	Compressed CO ₂
Mobile phase B:	isopropanol
Wash solvent:	Methanol
Flow rate:	1.7 mL/min
APBR:	2,000 psi
Column temp.:	55 °C
Sample temp.:	10 °C
Injection volume:	7.0 µL
Detection:	UV 260 nm
Gradient:	0.5% to 20% B in 9.9 min Hold at 20% for 2 min, re-equilibrate for 3 min

Results and Discussion

Due to the fact that the molar absorbance of vitamin D₃ is low and its concentration in samples is ten times lower than vitamin A, both vitamins A and D₃ were quantified at the wavelength of the maximum absorbance of D₃. The typical chromatograms obtained at 260 nm UV wavelengths for vitamin standards in solvent are shown in Figure 2. Chromatograms of the sample extract are shown in Figure 3.

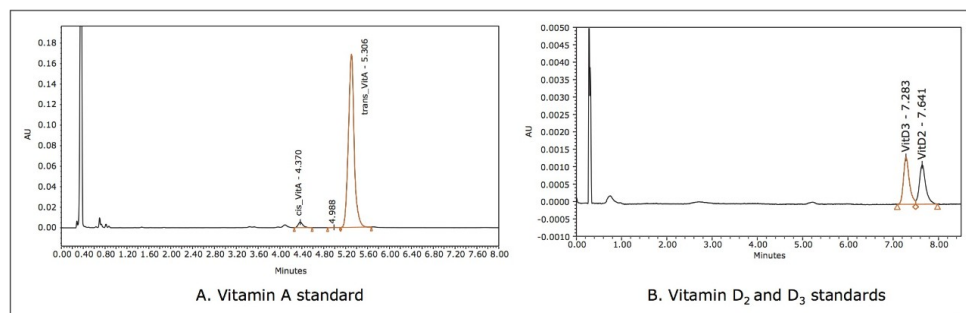


Figure 2. Chromatograms for the vitamin standard solutions at 260 nm.

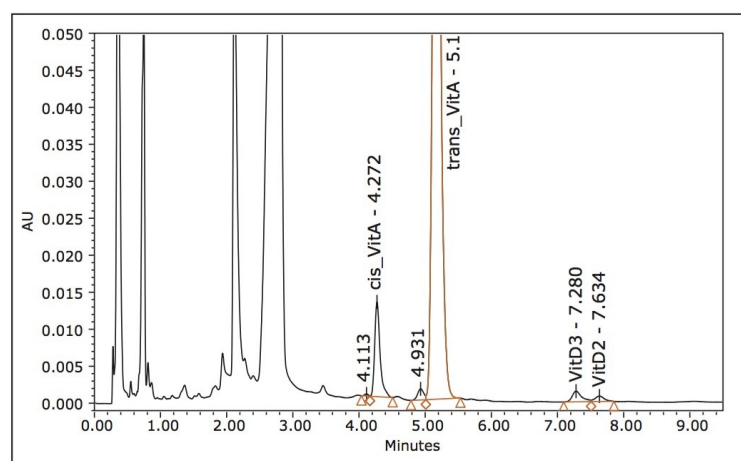


Figure 3. Chromatogram for the sample extract at 260 nm.

The linearity of the assay was investigated using standard solutions. The average peak areas of triplicate injections of seven concentrations were used. Vitamin D₂ was used as the internal standard for the quantification of vitamin D₃; there was no internal standard used for vitamin A. The coefficient of the determinations R^2 for vitamins A and D₃ were 0.9998 and 1.0000, respectively. The intermediate precision and reproducibility of the assay over a three-month period are shown in Table 1. The intermediate precision of the method for vitamin A was 5.1% and 5.7% for vitamin D₃, both falling well within the requirement of $\leq 8\%$.

Analyte	Intermediate precision, %	Requirement %, ^{2,3}	Product range, IU/g
Vitamin A	5.1	≤8	38,000 to 55,800
Vitamin D3	5.7	≤8	2,280 to 4,230

Table 1. Metrological properties of the method evaluated over three-month period.

To evaluate ruggedness, changes were made to the flow rate, system pressure, and column temperature to observe their effects on the results. There was no statistically significant difference to the $\pm 2\%$ changes on the results, and the chromatographic resolution of the critical D₂/D₃ pair was maintained above 1.6 (requirement for R ≥ 1.2).

The utilization of supercritical fluid chromatography using CO₂ combined with separation on sub 2- μm particle size columns provided fast and sufficient separation of the vitamins A and D within the same chromatographic run in a broad range of concentrations. The excellent selectivity of UPC² provided baseline separation of analytes from interfering impurity peaks and allowed quantification of the cis- and trans- retinols and the vitamin Ds at 260 nm UV in the same analytical run. Vitamins D₃ and D₂ (internal standard) were separated from each other, and eluted before the matrix peak, in contrast to our traditional method. The extracts in n-hexane can be injected directly into the system, which eliminates the solvent exchange step previously required in our laboratory.

Conclusion

A simple and cost-effective assay to analyze vitamin A and D₃ in different vitamin raw materials, premixes, and concentrates using UPC² has been developed and validated. The UPC²/PDA method demonstrates excellent linearity, resolution, and repeatability. The intermediate precision of the method calculated over a three-month period was less than 6%. The introduction of the ACQUITY UPC² System significantly reduced the consumption of HPLC solvents. Beside CO₂, the only other solvent required was isopropanol. The consumption of isopropanol was 0.12 mL per test. The laboratory has therefore significantly reduced the consumption of HPLC solvents as well as the disposal of waste solvents. By simplifying the procedures and eliminating the repartitioning steps, the reduction of solvent usage has streamlined our workflow and decreased the potential exposure of both laboratory staff and the environment to harmful solvents.

References

1. Aubin A, Analysis of Fat Soluble Vitamin Capsules using UltraPerformance Convergence Chromatography UPC.² Waters Application Note No. 720004394EN (2012).
2. Standard Method Performance Requirements for Vitamin A in Pre-Blends, Pre-Mixes, and Pure Materials – AOAC SMPR 2012.003.
3. Standard Method Performance Requirements for Vitamin D₃ in Pre-Blends, Pre-Mixes, and Pure Materials – AOAC SMPR 2012.004.

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720005220, November 2014

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