

A, **B**, **D**, **E** - Simultaneous determination of water- and fat-soluble vitamins

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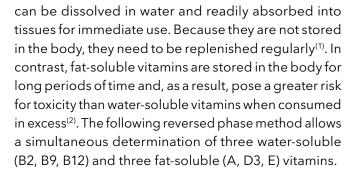


SUMMARY

Vitamins are divided into two different groups: water-soluble and fat-soluble. The fat-soluble vitamins are A, D, E and K whereas the water-soluble vitamins are the vitamin C and vitamin B complexes. Based on their different chemical properties, they are usually analyzed using different methods and modes. Reversed phase HPLC is very common for the water-soluble vitamins whereas normal phase mode is often applied for the fat-soluble vitamins. Nevertheless, it is possible that the analytes can occur in the same matrix. For this reason, a simultaneous determination of both water- and fat-soluble vitamins was developed in this work, resulting in a reduction in solvent consumption and total analysis time, is desirable.

INTRODUCTION

Vitamins are essential nutrients and are needed for a healthy metabolism in the human body. They are naturally found in many foods and are often present as additives in processed food products. They can be broadly classified into two groups: water-soluble and fatsoluble vitamins. But what is the difference? One major difference is that they are absorbed into the bloodstream and metabolized in a different manner and excreted from the body at different rates. Water-soluble vitamins





Additional Information

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SAMPLE PREPARATION

Individual stock solutions were prepared. Standards of vitamins B2, B9 and B12 were dissolved in deionized water. The solutions for vitamin E and vitamin D3 were ready to use ampules with a concentration of $\beta = 1 \text{ mg/ml}$ for each vitamin. Vitamin A palmitate stock solution was prepared by dissolving vitamin A palmitate in 100% methanol.

Analyte	Concentration (µg/ml)	Concentration (mg/100 g)
Vitamin B2 (riboflavin)	135	13.50
Vitamin B9 (folic acid)	115	11.50
Vitamin B12 (cyanocobalamin)	100	10.00
/itamin A palmitate	218	21.80
√itamin D3	1000	100.00
∕itamin E	1000	100.00

 Tab. 1 Concentrations of stock solutions

With these single standards a mixed standard was prepared.

Tab. 2 Concentrations of analytes in mixed standard

Analyte	Volume stock solution (ml)			Concentration Mix (mg/100 g)
Vitamin B2	1	4.30	31.40	3.14
Vitamin B9	1	4.30	26.74	2.67
Vitamin B12	1	4.30	23.26	2.33
Vitamin A palmitate	1	4.30	50.70	5.07
Vitamin D3	0.1	4.30	23.26	2.33
Vitamin E	0.2	4.30	46.51	4.65

RESULTS

The mixed standard was measured with the listed method in the Materials and Method section. A 5-point calibration was performed, and all analytes showed good linearity with values of R2 > 0.9996.

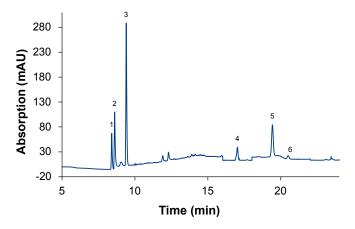


Fig. 1 Chromatogram of mixed standard; (1) vitamin B2; (2) vitamin B9; (3) vitamin B12; (4) vitamin A palmitate; (5) vitamin D3; (6) vitamin E, UV detection with wavelength switching

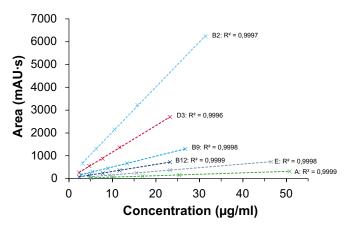


Fig. 2 Calibration curves

Based on the measurement of the lowest calibration level, the LOQ (S/N=10) and the LOD (S/N=3) were calculated.

Analyte	Level 1 (µg/ml)	Level 2 (µg/ml)	Level 3 (µg/ml)	Level 4 (µg/ml)	Level 5 (µg/ml)
Vitamin B2	3.14	6.28	10.47	15.70	31.40
Vitamin B9	2.67	5.35	8.92	13.37	26.74
Vitamin B12	2.33	4.65	7.75	11.63	23.26
Vitamin A palmitate	5.07	10.14	16.90	25.35	50.70
Vitamin D3	2.33	4.65	7.75	11.63	23.26
Vitamin E	4.65	9.30	15.50	23.26	46.51



Tab.4 Calculated LOQ and LOD

Analyte	LOQ (µg/ml)	LOQ (μg/100 g)	LOD (µg/ml)	LOD (µg/100 g)
Vitamin B2	0.010	1.00	0.003	0.30
Vitamin B9	0.029	3.00	0.009	0.90
Vitamin B12	0.038	4.00	0.011	1.10
Vitamin A palmitate	0.668	67.00	0.200	20.00
Vitamin D3	0.040	4.00	0.012	1.20
Vitamin E	0.465	47.00	0.140	14.00

CONCLUSION

The described method allows the simultaneous determination of water-soluble and fat-soluble vitamins. The challenge with this application is not the determination itself but finding an appropriate sample preparation procedure. The simultaneous method still requires the extraction of samples using two different methods that are suitable for each class of vitamins (dependent on their solubility). Nevertheless, no additional method for analysis is needed, meaning a reduction of solvent consumption and total analysis time is achievable. Furthermore, the analysis can be performed on a single reversed phased HPLC-System. A second normal phase HPLC-system is not necessary. In consequence, not only the cost for solvent consumption and analysis time are reduced, but also the cost for laboratory devices.

Method parameters

Parameter	Description				
Eluent A	Water + 0.01 % TFA (pH=4)				
Eluent B	Methanol				
Flow rate	0.5 ml/min				
Temperature	30° C				
Gradient	Time (min)	% A	% B		
	0	95	5		
	3	95	5		
	12	0	100		
	22	0	100		
	22.02	95	5		
	28	95	5		
Injection volume	5 μl				
Detection	Wavelength program				
	Time (min)	Wavelengt	h (nm)		
	0	360			
	8.50	270			
	16	318			
	18	270			
	20	292			
	22-28	360			
Data rate	20 Hz				
Time constant	0.05 s				

MATERIALS AND METHODS

System configuration

Instrument	Description	Article No.
Pump	AZURA P 6.1L, 10 ml, LPG	APH34EA
Autosample	r AZURA AS 6.1L with cooling/heating	AAA51AA
Detector	AZURA DAD 6.1L	ADC11
Flow cell	High Sensitivity KNAUER LightGuide UV Flow Cell Cartridge, 50 mm, 6 μL, 50 bar	AMD59XA
Thermostat	AZURA CT 2.1	ATC00
Column	Eurospher II 100-3 C18P with precolumn, 150 x 3 mm ID	15XE182E2G
Software	ClarityChrom 8.2.3 - Workstation, autosampler control included	<u>A1670</u>
Software	ClarityChrom 8.2.3 - PDA extension	A1676



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REFERENCES

[1] https://www.verywellhealth.com/fat-vs-watersoluble-998218, 2021-09-27

[2] Fat-Soluble Vitamins: A, D, E, and K; Fact Sheet No. 9.315, Food and Nutrition Series/Health, <u>https://</u> <u>extension.colostate.edu/topic-areas/nutrition-</u> <u>food-safety-health/fat-soluble-vitamins-a-d-e-</u> <u>and-k-9-315/</u>, 2021-09-27

RELATED KNAUER APPLICATIONS

VFD0172 - A D E K - Easy separation of fat-soluble vitamins using GPC/SEC

VFD0162 - Separation of ascorbic acid and vitamin B complexes - essentially required nutrients