Reveleris® X2 Flash System

Reveleris® X2 System Purifies Ferulic Acid in Rhizoma Chuanxiong Extract with Greater Recovery

Introduction

Medicinal plants are an important resource for drug development. They are commonly analyzed in biochemical, pharmaceutical, and clinical research for their complex constituents and low quantities of bioactive ingredients.

Ferulic acid is known to be one of the main bioactive components found in Rhizoma chuanxiong. The plant has been extracted here and used for the separation and identification of its scarce ingredients using the Reveleris® X2 flash chromatography system.

Extraction Conditions:

Extract Type: Pulverized

Weight: 2g

Extraction Solvent: Ether Solvent Volume: 40mL Ultra-sonication: 20 min

Flash Chromatography Conditions:

Cartridge: Reveleris® silica 12g

Solvent A: Hexane
Solvent B: Ethyl acetate
Flow Rate: 25mL/min
UV1 Wavelength: 254nm
UV2 Wavelength: 280nm
Cartridge Equilibration: 3.0 min

Injection Type: Liquid

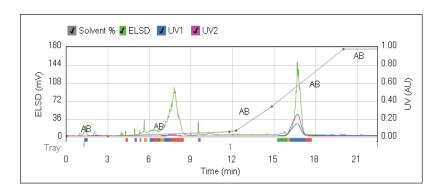


Figure 1: Separation of Rhizoma chuanxiong extract shows the detection of peaks using the UV and the evaporative light scattering detectors.

Gradient Table		
Step	Time (min.)	%В
1	0.0	0
2	3.0	0
3	8.8	5
4	0.5	6
5	2.6	33
6	5.2	98
7	2.6	98



HPLC Conditions:

Cartridge: Reveleris® silica 12g

Column: Alltima[™] silica 5μm, 250 x 4.6mm **Mobile Phase A:** 1% Formic acid in Hexane

Mobile Phase B: 1% Formic acid in

Ethyl acetate

Flow Rate: 1.0mL/min UV Detector: 280nm

Gradient: Time: 0 | 10 | 20

%B: 0 | 50 | 100

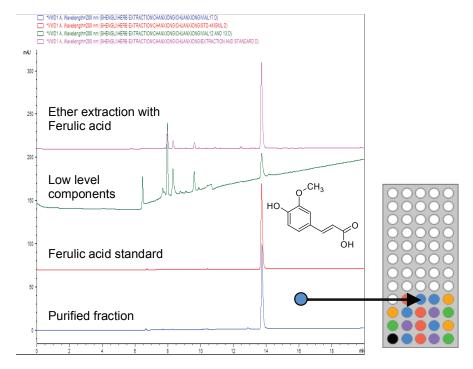


Figure 2: Ferulic acid recovery = 97% (calculated by peak area)

Conclusion

Using the sensitive Reveleris® X2 flash chromatography system, the critical components of the plant extract at low levels have been isolated and collected. The RevealX™ technology also allows non-chromophoric peaks to be detected with the chromophoric ones in the same run for fraction collection and further analysis. This allows purification of compounds present in the natural product extract, compounds that are not always easily detectable using traditional flash chromatography system and are present at low amounts and useful for medicinal purposes.

References

1. Chen, X.; Kong, L.; Su, X.; Fu, H.; Ni, J.; Zhao, R.; Zou, H.; Separation and identification of compounds in Rhizoma chuanxiong by comprehensive two-dimensional liquid chromatography coupled to mass spectrometry; J. Chromatograpy A, 1040, (2004), pp. 169-178.

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