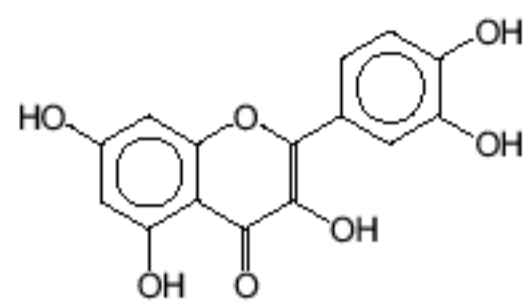
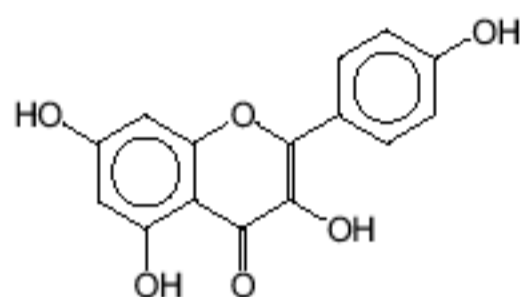


Ginko Biloba Extract and Tablets



Quercetin



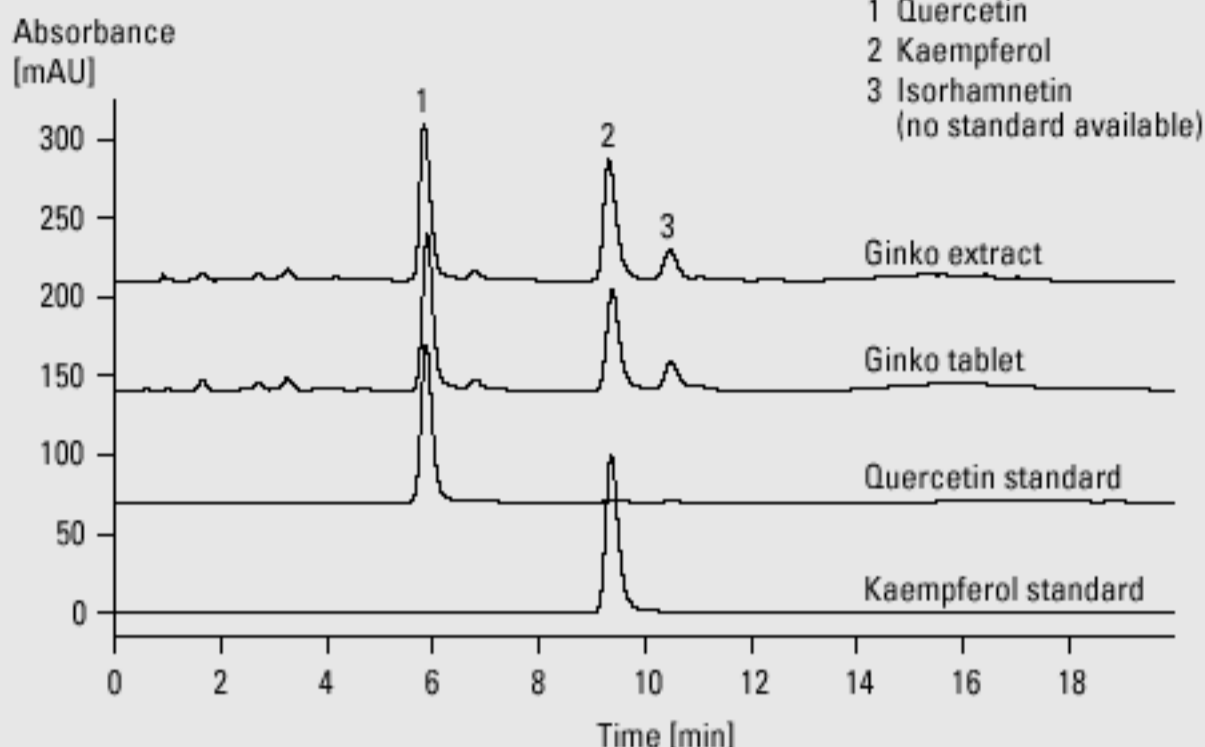
Kaempferol

Extraction

4 g of *Ginko Biloba* extract (from Caesar & Loretz GmbH, Germany) were refluxed for 30 min in 70 ml methanol and 10 ml 25 % HCl. After cooling to room temperature the mixture was filtered and the filter washed with approximately 100 ml methanol. The solvent was partly removed *i. vac.* and diluted with methanol to 100 ml in a volumetric flask. 5 ml of this solution were filtered through a C18 disposable cartridge. The cartridge was washed with 4 ml methanol and the filtrate diluted to 10 ml in a volumetric flask. The same procedure was used to extract 10 ginko tablets ('Promod Ginko biloba L.', 40 mg ginko biloba extract per tablet, Sine Laboratories', Shanghai, China)

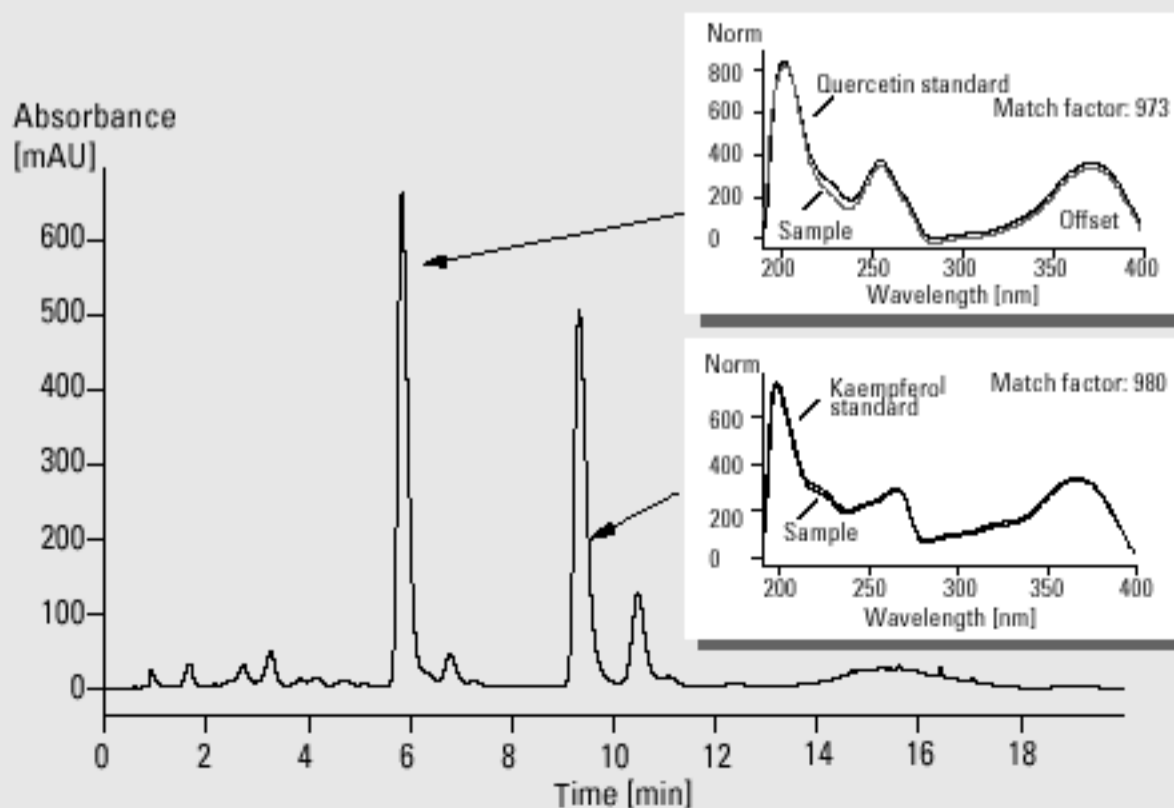
Method and extraction from:
A. Hasler, O. Stichler, *J. Chromatogr.* 508 (1990), 236-240

Instrumentation:
see configuration example 2 on page 77



Analysis of *Ginko Biloba* extract

Column	4 x 125 mm Hypersil ODS, 5 µm
Mobile phase	A = 0.5 % H ₃ PO ₄ in water, B = methanol
Flow rate	2.0 ml/min
Gradient	at 0 min 38 % B at 12 min 48 % B
Column wash	at 17 min 100 % B at 20 min 38 % B 100 % B to 38 % B in 3 min
UV detector	variable wavelength detector 370 nm, standard cell
Column compartment temperature	25 °C
Stop time	20 min
Post time	5 min
Injection volume	10 µl



Comparison of sample and standard spectra of quercetin and kaempferol