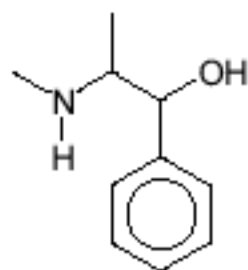
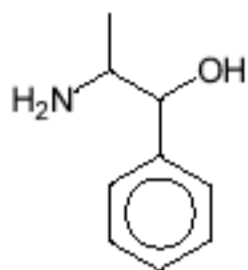


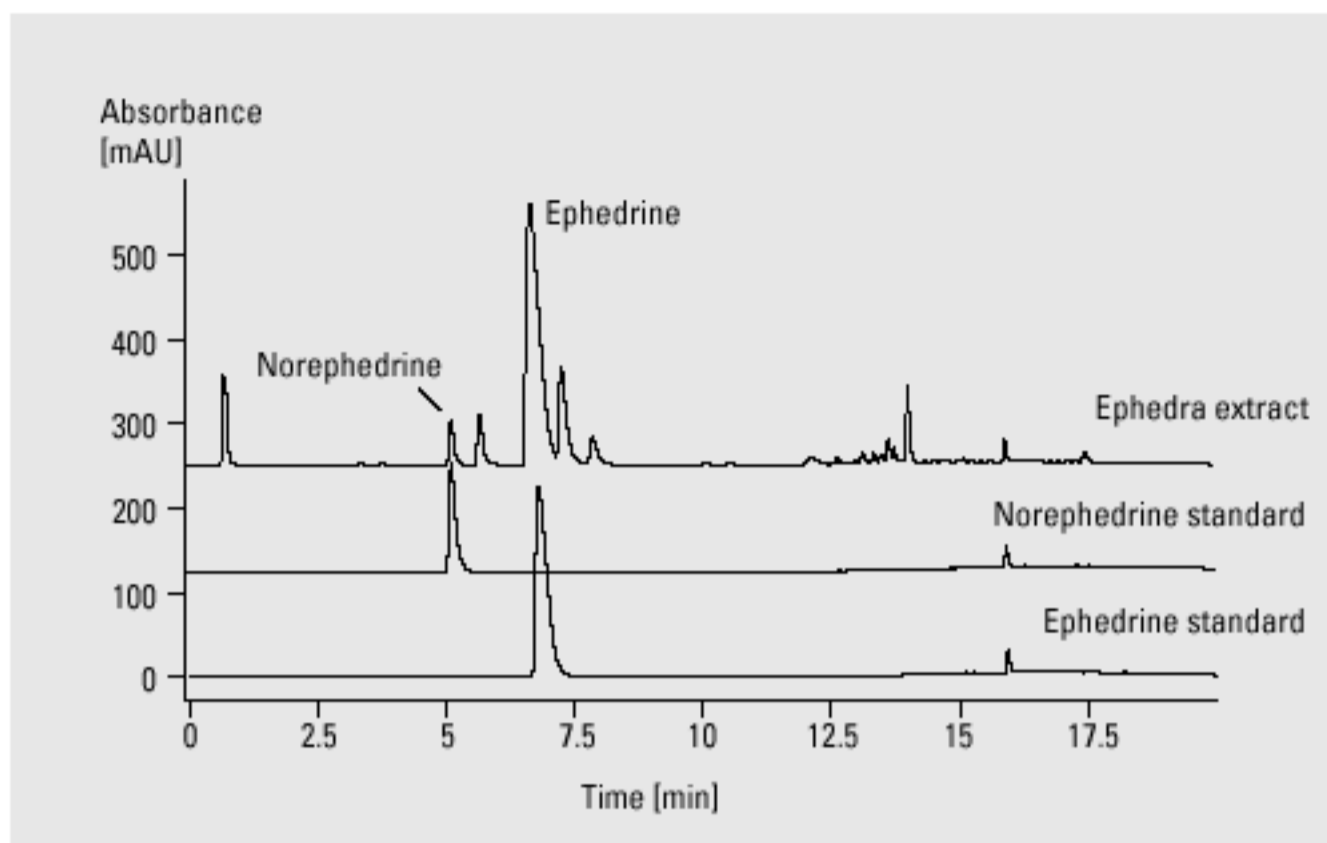
# Ephedra Sinica Stapf Extract



Ephedrine



Norephedrine



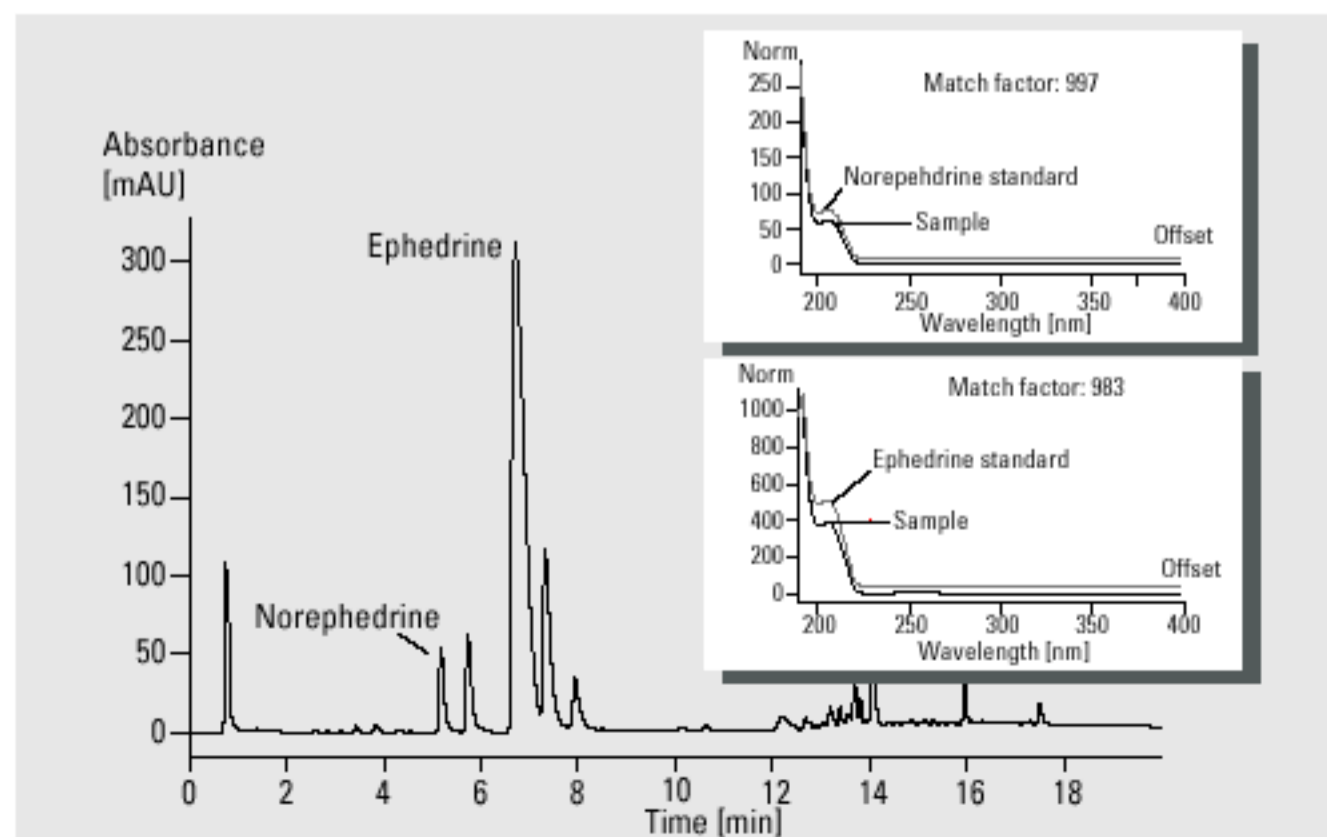
## Analysis of *Ephedra Sinica Stapf* extract

### Extraction

50 ml of 0.5 M H<sub>2</sub>SO<sub>4</sub> was added to 1 g dried plant and the mixture was stirred overnight. After filtration the pH was adjusted to 11-13 by adding 6 M NaOH. 8 g NaCl were added and the mixture extracted with ether (4 x 25 ml). The organic layer was dried over MgSO<sub>4</sub>, filtered and the solvent was removed *i. vac.* 5 ml H<sub>2</sub>O and 3 drops of 4 M HCl were added to the residue.

<b>Column</b>	4.6 x 75 mm Zorbax SB-C18, 3.5 µm
<b>Mobile phase</b>	A = 0.025 M KH <sub>2</sub> PO <sub>4</sub> in water (pH = 3), B = acetonitrile
<b>Flow rate</b>	1.0 ml/min
<b>Gradient</b>	at 0 min 2 % B at 10 min 10 % B
<b>Column wash</b>	at 15 min 80 % B at 18 min 80 % B at 20 min 2 % B
<b>UV detector</b>	variable wavelength detector 204 nm, standard cell
<b>Column compartment temperature</b>	25 °C
<b>Stop time</b>	20 min
<b>Post time</b>	5 min
<b>Injection volume</b>	5 µl

Instrumentation:  
see configuration example 2 on page 77



Comparison of sample and standard spectra of norephedrine and ephedrine