# CHROMABOND® HR-X

#### Technical data

Hydrophobic polystyrene	-divinylbenzene	copolymer	(PS/DV	B)
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SPE mode:	Reversed phase
Interactions:	Hydrophobic and $\pi$ – $\pi$
Particle shape:	Spherical
pH stability:	1–14
Particle size:	85 µm and 45 µm
Pore size:	55–60 Å
Specific surface:	1000 m²/g
RP capacity:	390 mg/g (caffeine in water)



#### Recommended application

- Pharmaceuticals / active ingredients from tablets, creams and water
- Drugs and pharmaceuticals from urine, blood, serum and plasma
- Trace analysis of pesticides, herbicides, phenols, PAH and PCBs from water

## Standard protocol for CHROMABOND<sup>®</sup> HR-X

#### MN Appl. No. 304310

Column type: CHROMABOND<sup>®</sup> HR-X/3 mL/200 mg, REF 730931

#### Sample pretreatment:

Individual sample preparation in reference to the compounds and matrix (adjust pH value if necessary).		
Conditioning:	5 mL methanol, then 5 mL water (do not let run the column dry!)	
Sample aspiration:	The prepared sample is passed through the column by vacuum or pressure (max. 1000 mL sample volume)	
Washing:	5 mL water / methanol (95:5, v/v)	
Drying:	With nitrogen or air	
Elution:	3 x 2 mL methanol	
Further analysis:		
Evaporation and reconstitution (if necessary); HPLC or GC		
<b>T</b> I IV.		

These conditions are a starting point for SPE method development. Further optimization may be required to improve results.

#### Good to know

- A possible replacement for:
- Nexus
- ENVI-Chrom P
- Bakerbond H<sub>2</sub>O-phobic DVB

Strata<sup>™</sup>-X



### Applications

Dete	rmination of	pyrrolizidine alkaloids	
MN A	ppl. No. 30662	20	
Chron	natographic co	nditions	
	Columns:	CHROMABOND <sup>®</sup> HR-X/85 µm/3 mL/200 mg	140
	MN REF:	730921	120
V	Pretreatment:	The following analysis were performed with standard solutions	
	Conditioning:	5 mL methanol, 5 mL water	
	Application:	10 mL neutralized standard solution with a flow rate of 3 mL/min	
	Washing:	2 x 5 mL of water with a flow rate of 3 mL/min	
	Drying:	5–10 min with vacuum	40 40 40 40 40 40 40 40 40 40 40 40 40 4
	Elution:	5 mL methanol	20
	Eluent exchange: Add 1.0 mL water as keeper. Evaporate eluate to a volume of 0.5 mL at 40 °C under a stream of nitrogen and fill up to 1.0 mL with water / methanol (95:5, v/v).		○ 缶 荒 山 岙 교 중 壬 롯 ¥ 톤 우 령 그 号 기 중 왕 중 8 중 8 중 8 중 8 중 8 중 8 중 8 중 8 중 8
	Further analys	sis:	ECHROMABOND® C <sub>18</sub> ec, 500 mg, 6 mL ECHROMABOND® HR-X, 85 µm, 200 mg, 3 mL
	HPLC determ NUCLEOSHE to MN Appl. N	ination of recovery rates with EC 150/2 ELL <sup>®</sup> RP 18plus, 2.7 μm (REF 763236.20) in reference No. 127480	Superior to silica based RP phase CHROMABOND <sup>®</sup> HR-X shows higher recovery rates for most tested pyrrolizidine alkaloids than CHROMABOND <sup>®</sup> C18 ec under the given conditions.

## Enrichment of opiates

### MN Appl. No. 306710

#### Chromatographic conditions

Columns:	CHROMABOND <sup>®</sup> HR-X/45 µm/3 mL/60 mg
MN REF:	730936P45
Pretreatment:	$400~\mu L$ methanolic standard solution were diluted with 50 mmol/L phosphate buffer pH 7.0 to 20 mL 2.5 mL of this solution are equal to 5 ng of each analyte
Conditioning:	3 x 1 mL methanol, 3 x 1 mL water, then 3 x 1 mL 50 mmol/L phosphate buffer pH 7.0
Aspiration:	2.5 mL of pretreated sample solution is passed through the column at a flow of 1–2 mL/min
Washing:	3 x 1 mL 50 mmol/L phosphate buffer pH 7.0, 3 x 1 mL water
Drying:	5 mL air by pushing with a syringe
Elution:	3 x 1 mL 0.1 % formic acid in methanol

Solvent change: Eluate is evaporated to dryness at 30 °C under a stream of nitrogen and then redissolved in organic solvent suited for the subsequent analysis.

#### Further analysis:

HPLC determination of recovery rates with EC 100/2 NUCLEOSHELL® Biphenyl, 2.7  $\mu m$  (REF 763634.20) in reference to MN Appl. No. 128880

Compound	Recovery rate [%]	Standard deviation [%]
Ecgonine methyl ester	94	0
Morphine	77	3
Dihydrocodeine	101	1
Codeine	97	1
6-Acetylmorphine	89	1
Benzoylecgonine	102	0
6-Acetylcodeine	100	0
Cocaine	109	1
Noscapine	95	1
Papaverine	98	2

