



explore

LUNA®



phenomenex®
...breaking with tradition™





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Becoming the world's #1 HPLC column meant no shortcuts.

The Luna® brand of columns and media is more than just a product line from Phenomenex. It is a pledge to provide you with the highest level of satisfaction for your chromatographic goals. Every aspect of Luna products has been engineered to meet the exacting demands placed on today's chromatographers.

Luna products continue to uphold the quality our customers depend on. If you have never tried Luna columns or media, this brochure will guide you through the various solutions to fit your needs.

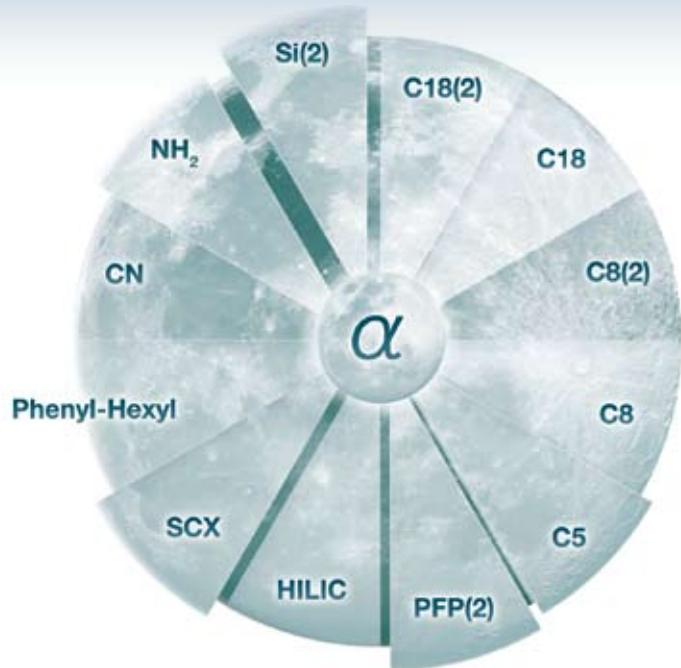
For those who use Luna products daily, thank you for making Luna columns #1 in the world.

Explore Successful Separations

Your success begins with our commitment to provide the essential solutions to HPLC separations in the Luna brand. Some of the highest quality and performance standards are incorporated into Luna products, making them an indispensable platform for all areas of HPLC.

Explore Resolution with Luna Selectivities

Phase selectivity has the strongest impact on overall chromatographic resolution. Choosing the optimal selectivity can drive your separation to success. Luna phases span through 12 different chemistries, each offering its own unique selectivity.



Bonded Phases

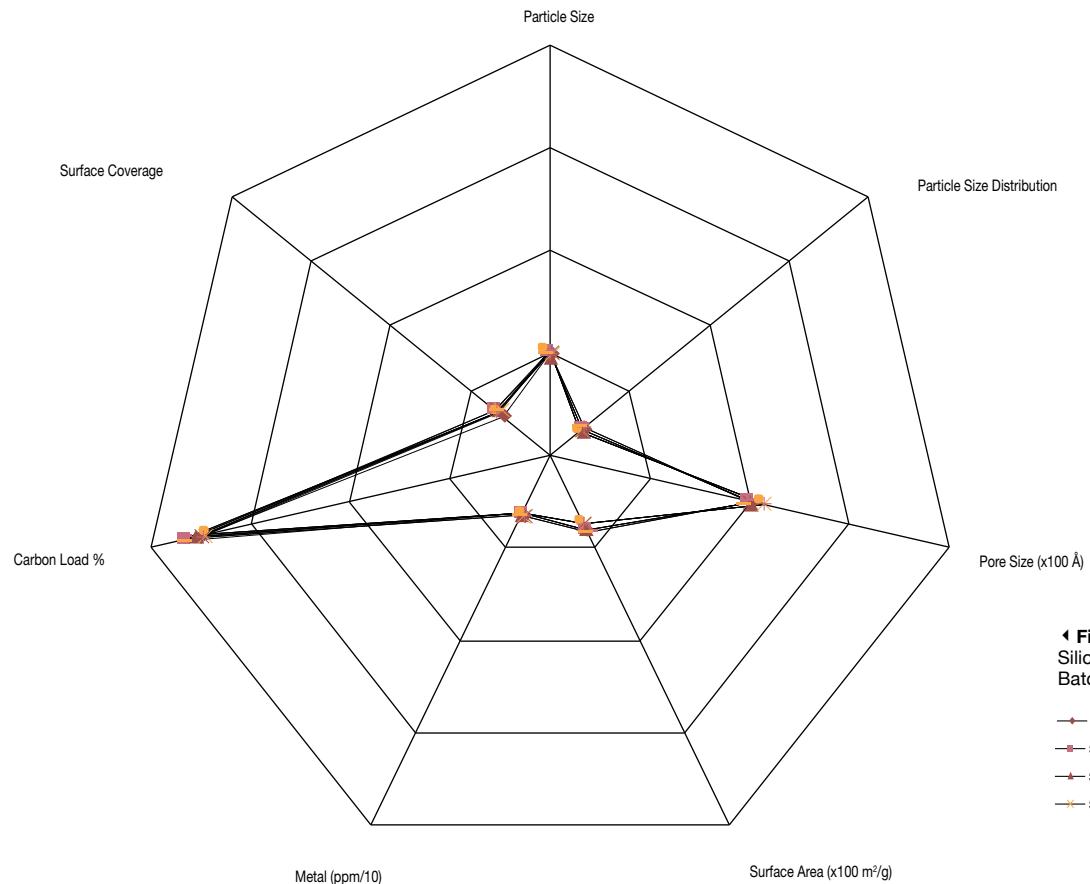
LUNA PHASES	Description	Particle Size (μm)	Pore Size (\AA)	Surface Area (m^2/g)	Carbon Load (%)	Bonded Phase Coverage ($\mu\text{mole}/\text{m}^2$)	pH Stability	Application	Reversed Phase	Normal Phase	HILIC	IEX
Silica(2)	Unbonded silica	3, 5, 10, 10-PREP, 15	100	400	—	—	—	Non-polar compounds		<input checked="" type="radio"/>		
C5	5 Carbon ligand	5, 10	100	440	12.5	7.85	1.5 - 10	Good alternative to C8 when less retention is desired	<input checked="" type="radio"/>			
C8	Highly retentive, original C8 phase	5, 10	100	440	14.75	5.50	1.5 - 10	For use when a more retentive C8 is desired	<input checked="" type="radio"/>			
C8(2)	C8 ligand optimized for improved peak shape	3, 5, 10, 10-PREP, 15	100	400	13.5	5.50	1.5 - 10	Great starting phase for method development	<input checked="" type="radio"/>			
C18	Highly retentive, original C18 phase	5, 10	100	440	19.0	3.00	1.5 - 10	Use when maximum retention of non polars is desired	<input checked="" type="radio"/>			
C18(2)	C18 ligand optimized for improved peak shape	2.5, 3, 5, 10, 10-PREP, 15	100	400	17.5	3.00	1.5 - 10	From capillary LC/MS to process scale OUR MOST POPULAR PHASE	<input checked="" type="radio"/>			
CN	Versatile CN phase	3, 5, 10	100	400	7.0	3.80	1.5 - 7.0	For improving the retention of polar compounds	<input checked="" type="radio"/>	<input checked="" type="radio"/>		
NH₂	Rugged and reproducible NH ₂	3, 5, 10	100	400	9.5	5.80	1.5 - 11	Sugar alcohols, anionic or hydrogen bonding compounds	<input checked="" type="radio"/>	<input checked="" type="radio"/>		<input checked="" type="radio"/>
Phenyl-Hexyl	Phenyl phase attached to C6 (hexyl) ligand	3, 5, 10, 10-PREP, 15	100	400	17.5	4.00	1.5 - 10	Unique selectivity for very polar and aromatic compounds	<input checked="" type="radio"/>			
SCX	Benzene sulfonic acid	5, 10	100	400	Binding Capacity: 0.15 meq/g		2.0 - 7.0	Amine and polyamine containing compounds				<input checked="" type="radio"/>
HILIC	Reproducible, cross-linked diol	3, 5	200	200	5.7	4.30	1.5 - 8.0	Increased retention and MS sensitivity of polar compounds			<input checked="" type="radio"/>	
PFP(2)	Pentafluorophenyl with a C3 (propyl) linkage	3, 5	100	400	11.5	2.2	1.5 - 8.0	Highly polar compounds, halogenated compounds and isomers	<input checked="" type="radio"/>			

Explore Robust Methods

Successful methods depend on results that can tolerate minor variations in chromatographic parameters. The base silica of Luna is 99.999 % pure and meticulous care is given to quality control over all aspects of silica structure and chemistry. This ensures that Luna columns will always perform consistently, resulting in method reproducibility you can trust.

Reliable Performance

Almost no variation is observed among the batches of Luna. **Figure 1** shows quality control test data designed to monitor the slightest differences that may affect reproducibility - particle shape and smoothness, porosimetry, bonding consistency and pH stability.

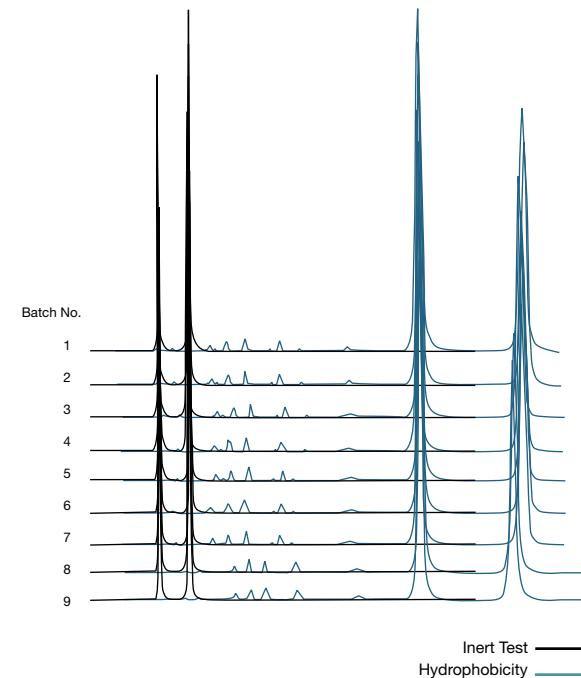


◀ **Figure 1**
Silica Reproducibility.
Batch No:

- ◆— 5291-1 —★— 5291-5
- 5291-2 —○— 5291-6
- ▲— 5291-3 —△— 5291-7
- ×— 5291-4 —— 5291-8
- 5291-9

Column to Column Reproducibility

The chromatograms in **Figure 2** show consistency of inertness (black) and hydrophobicity (blue) for Luna 5 μm C18(2) columns from 9 different batches. Almost no variation is observed.



◀ **Figure 2**
Column-to-column
reproducibility for 9 batches of Luna

Explore Options for Every Development Route

Luna media is available in a wide variety of particle sizes and formats, each with different attributes that can be optimized for nearly any stage of development.



Fast LC-MS Methods

Luna media is available in MercuryMS™ Cartridges and online columns for quick, cost-effective screening methods.



High-Speed-Technology

Luna 2.5 µm C18(2)-HST columns deliver highly efficient separations without the need for expensive high-pressure instruments.



Develop Robust Analytical Methods

Analytical HPLC columns are the most widely used format and are available in a wide variety of dimensions and particle sizes.



Lab-Scale Purification Redefined

Axia™-packed Luna® preparative columns provide industry-leading lifetimes and efficiencies.



Beyond our largest preparative column dimensions, Luna phases are available in bulk quantities for HPLC purification at the process, pilot, and commercial scale. The highly reproducible manufacturing process makes scaling to large scale purification extremely straight-forward.

The wide range of Luna phases provides you with the selectivity choices to optimize parameters such as retention time and resolution. Additionally, the high surface area ($400 \text{ m}^2/\text{g}$) of Luna materials gives you greater loadability than most other medias. For those challenging purifications where chromatography is the best option, the Luna family offers an excellent platform for all purification challenges.



Contact your Phenomenex technical consultant or local distributor for bulk media sales.

Luna C18(2), C8(2), C5

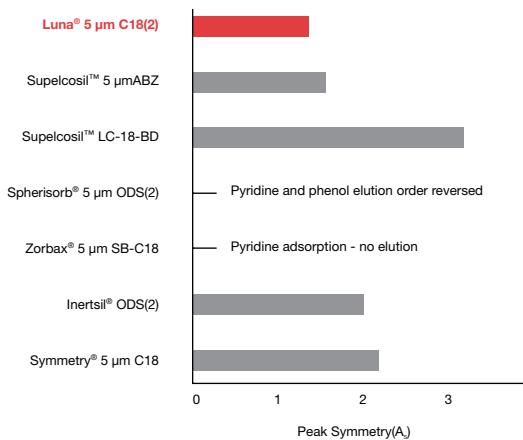
Your Starting-Point for All Reversed Phase Methods

Luna has found a place as one of the world's top reversed phase columns because it can help optimize two important chromatographic properties: resolution and peak shape. The high efficiencies and bonded phase surface coverage provide for sharp peaks. Whether you need a column for USP methods or just general method development, Luna C18(2) and C8(2) should be your first choice every time.

The result:

- » Free exposed silanols virtually eliminated by complete bonding and endcapping
- » Sharp peak shape for good method sensitivity
- » pH stable from 1.5 to 10.0 for over 10,000 hours

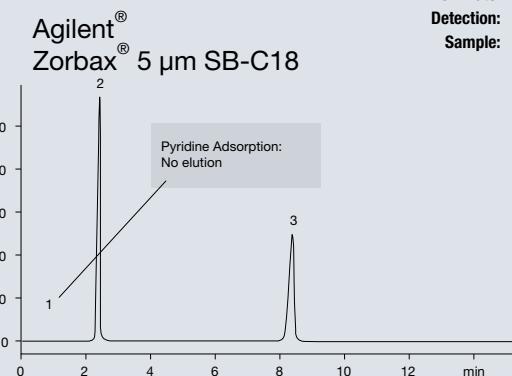
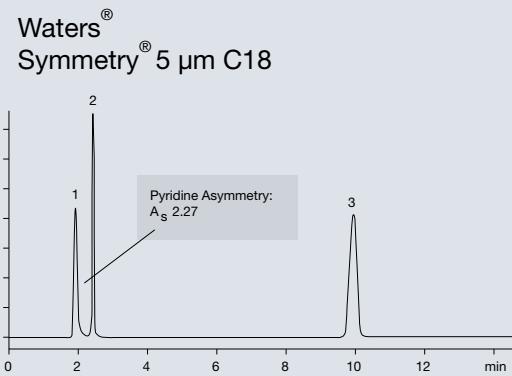
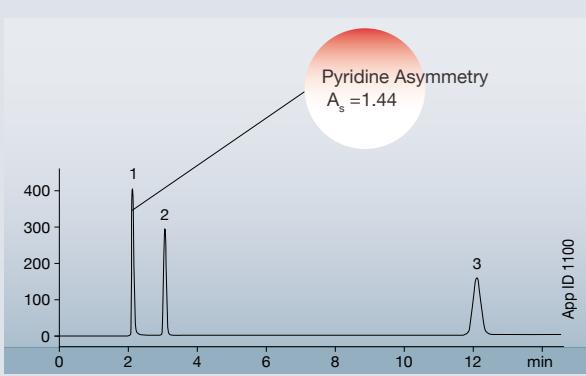
Pyridine Peak Asymmetry Comparison**



Comparison of 7 different 5 μm reversed phase columns. This survey measures the degree of silanol activity on the surface of each silica. In this survey, Luna 5 μm C18(2) material demonstrates the lowest silanol activity.

PEAK ASYMMETRY COMPARISON OF COMPETING COLUMNS**

Phenomenex® Luna® 5 μm C18(2)



Conditions for all columns

Dimension: 150 x 4.6 mm
Mobile Phase: Acetonitrile/Water (50:50)
Flow Rate: 1.0 mL/min
Detection: UV @ 254 nm
Sample: 1. Pyridine
2. Phenol
3. Toluene

** The comparative data presented here may not be representative for all applications.

Polar, Acidic Drugs

■ Phenomenex® Luna® 3 µm C18(2)
■ Waters® Symmetry® 3.5 µm C18

Conditions same for both columns

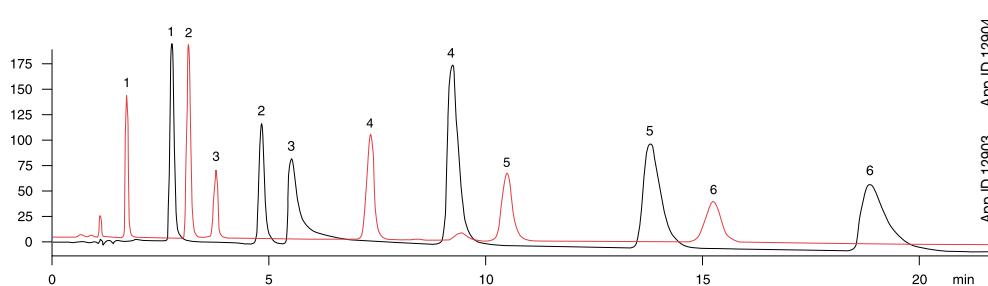
Dimension: 75 x 4.6 mm
Mobile Phase: 20 mM KH₂PO₄/Acetonitrile(70:30)

Flow Rate: 0.75 mL/min

Detection: UV @ 202 nm

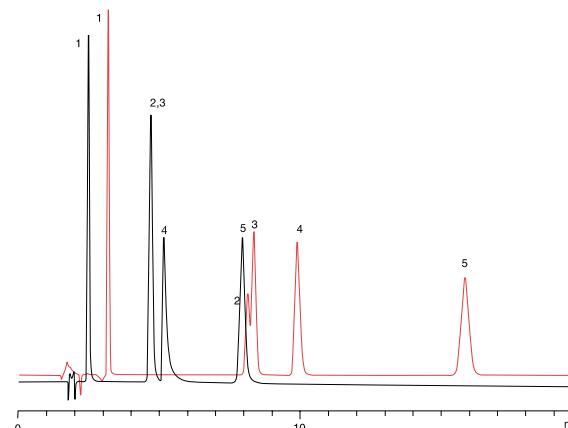
Sample:

1. Tolmetin
2. Naproxen
3. Diflunisal
4. Fenoprofen
5. Indomethacin
6. Ibuprofen



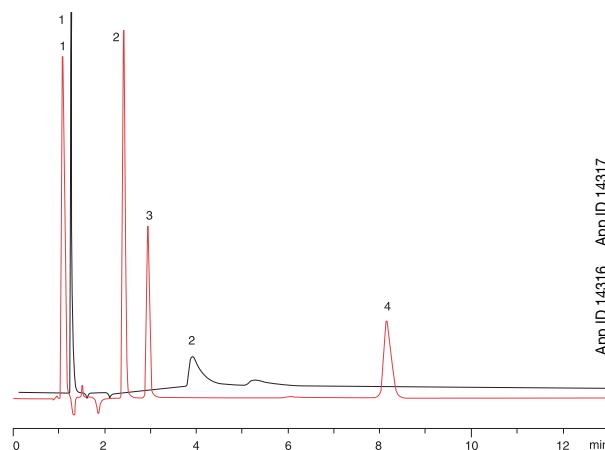
Hydrophobic, Acidic Compounds

■ Phenomenex® Luna® 5 µm C18(2)
■ Thermo Hypersil-Keystone® HyPURITY™ Elite 5 µm C18



Basic Compounds

■ Phenomenex® Luna® 5 µm C18(2)
■ Macherey Nagel® Nucleosil® 5 µm C18



Conditions same for both columns

Dimension: 150 x 4.6 mm
Mobile Phase: 20 mM Potassium phosphate, pH 2.5 / Acetonitrile (75:25)

Flow Rate: 1.5 mL/min

Temperature: 30 °C

Detection: UV @ 210 nm

Sample:

1. Maleic acid
2. Tripolidine*
3. Chlorpheniramine*
4. Diphenhydramine*

*Peaks 2-4 adsorb on Nucleosil C18

α- and β-acids in Hop Extract

Column: Luna 5 µm C18(2)

Dimension: 250 x 4.6 mm

Part No.: 00G-4252-E0

Mobile Phase: Methanol with 0.1 % H₃PO₄ / Water with 0.1 % H₃PO₄ (90:10)

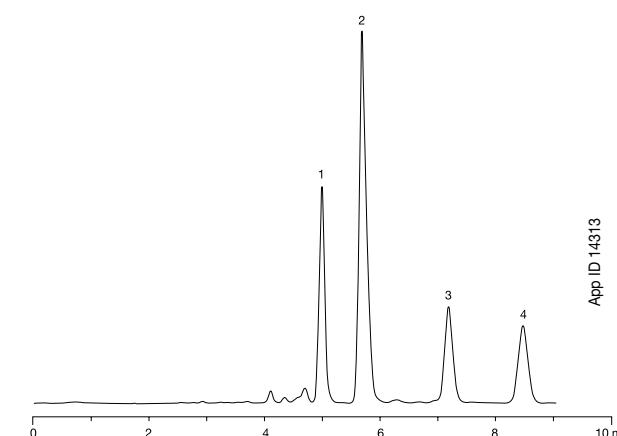
Flow Rate: 1.5 mL/min

Temperature: 30 °C

Detection: UV @ 314 nm

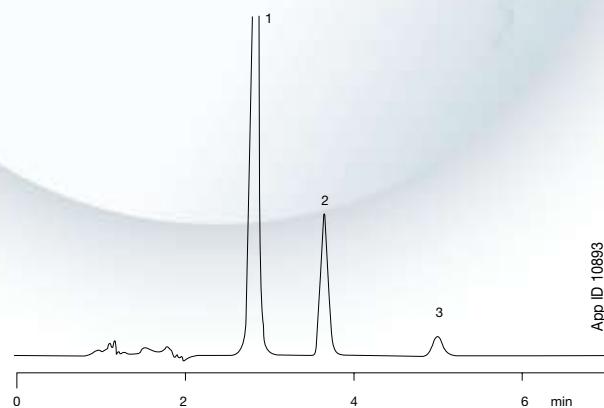
Sample:

1. Cohumulone
2. Ad-+humulone
3. Colupulone
4. Ad-+lupulone



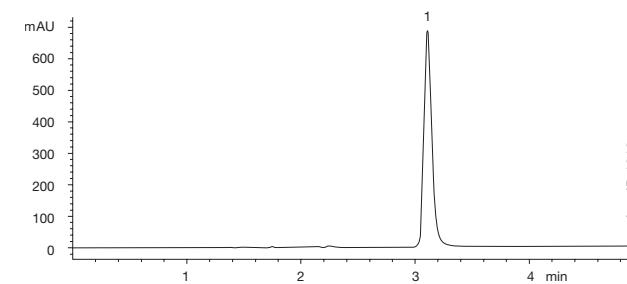
Luna C18(2), C8(2), C5 (cont.)

USP METHOD: ESTRADIOL



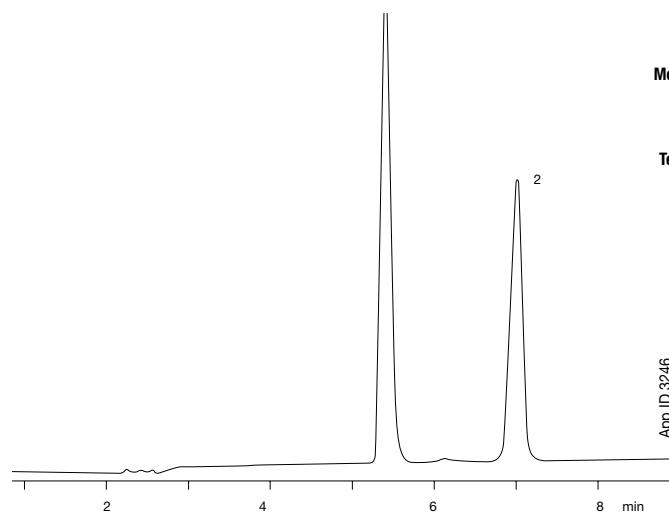
Column: Luna 5 μ m C18(2)
Dimension: 150 x 4.6 mm
Part No.: 00F-4252-E0
Mobile Phase: Acetonitrile/Water (55:45)
Flow Rate: 1 mL/min
Temperature: 30 °C
Detection: UV @ 254 nm
Sample: 1. Ethylparaben
2. Estrone
3. Estradiol

COCAINE-NARCOTIC DRUG



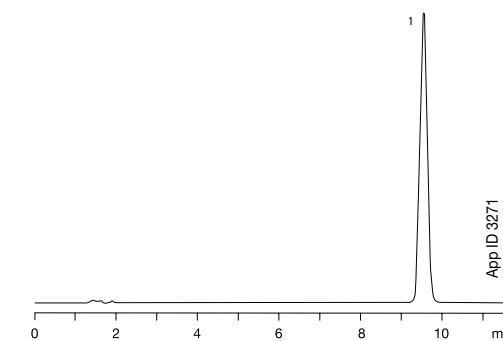
Column: Luna 5 μ m C18(2)
Dimension: 150 x 4.6 mm
Part No.: 00F-4252-E0
Mobile Phase: Phosphate Buffer, pH 2.5/
Acetonitrile (75:25)
Flow Rate: 1 mL/min
Temperature: 30 °C
Detection: UV @ 233 nm
Sample: 1. Cocaine hydrochloride

USP METHOD: PHENYLEPHRINE HYDROCHLORIDE INJECTION



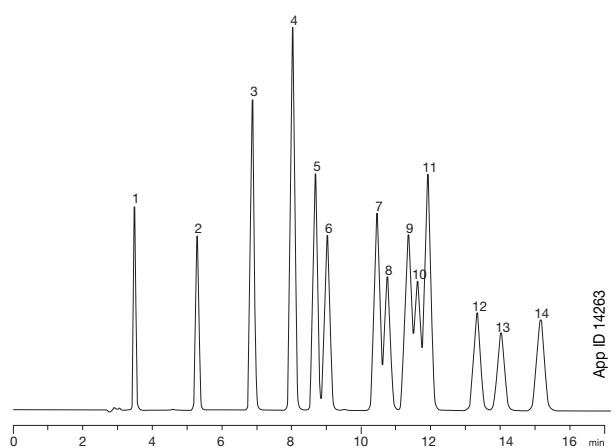
Column: Luna 5 μ m C18(2)
Dimension: 250 x 4.6 mm
Part No.: 00G-4252-E0
Mobile Phase: Water/Methanol with 1.1 %
1-Octanesulfonic
acid pH 3.0 (50:50)
Flow Rate: 1 mL/min
Temperature: 22 °C
Detection: UV @ 280 nm
Sample: 1. Phenylephrine
hydrochloride
2. Epinephrine bitartrate

USP METHOD: HYDROCORTISONE CREAM



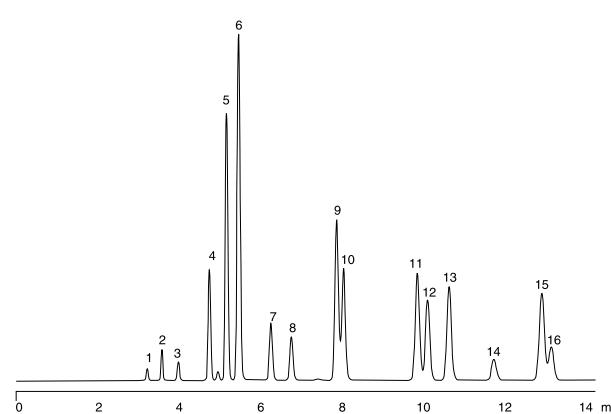
Column: Luna 5 μ m C18(2)
Dimension: 250 x 4.6 mm
Part No.: 00G-4252-E0
Mobile Phase: Water/Acetonitrile (75:25)
Flow Rate: 2 mL/min
Temperature: 30 °C
Detection: UV @ 254 nm
Sample: 1. Hydrocortisone

EPA 8330 - EXPLOSIVES



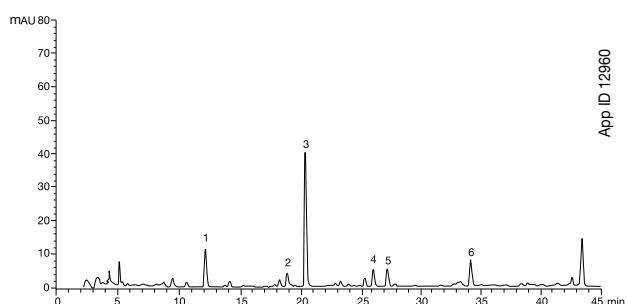
Column: Luna 5 μm C18(2)
Dimension: 250 x 4.6 mm
Part No.: 00G-4252-E0
Mobile Phase: Methanol/Water (55:45)
Flow Rate: 1 mL/min
Temperature: 35 °C
Detection: UV @ 254 nm
Sample: 1. HMX
 2. RDX
 3. 1,3,5-Trinitrobenzene
 4. 1,3-Dinitrobenzene
 5. Tetryl
 6. Nitrobenzene
 7. 2,4,6-Trinitrotoluene
 8. 4-Amino-2,6-Dinitrotoluene
 9. 2-Amino-4,6-Dinitrotoluene
 10. 2,6-Dinitrotoluene
 11. 2,4-Dinitrotoluene
 12. 2-Nitrotoluene
 13. 4-Nitrotoluene
 14. 3-Nitrotoluene

EPA 8310 - POLYNUCLEAR AROMATIC HYDROCARBONS (PAH S)



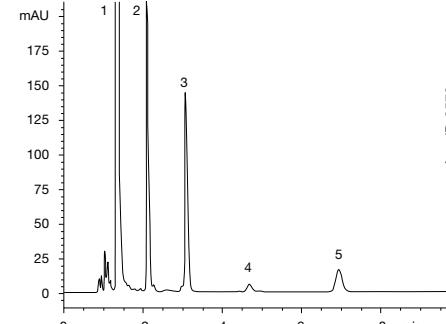
Column: Luna 5 μm C18(2)
Dimension: 250 x 4.6 mm
Part No.: 00G-4252-E0
Mobile Phase: A: Water B: Acetonitrile
Gradient: A/B (25:75) to 100 % B in 25 min
Flow Rate: 2 mL/min
Temperature: 22 °C
Detection: UV @ 254 nm
Sample: 1. Naphthalene
 2. Acenaphthalene
 3. Fluorene
 4. Phenanthrene
 5. Anthracene
 6. Fluoranthracene
 7. Pyrene
 8. Benzo(a)anthracene
 9. Chrysene
 10. Benzo(e)pyrene
 11. Benzo(b)fluoranthene
 12. Benzo(k)fluoranthene
 13. Benzo(a)pyrene
 14. Dibeno(a,h)anthracene
 15. Benzo(g,h,i)perylene
 16. Indeno(1,2,3-c,d)pyrene

GINGER PUNGENTS



Column: Luna 5 μm C18(2)
Dimension: 250 x 4.6 mm
Part No.: 00G-4252-E0
Mobile Phase: A: Water B: Acetonitrile
Gradient: A/B (55:45) to A/B (50:50)
 in 8 min, A/B (35:65) in 15 min,
 A/B (10:90) in 40 min
Flow Rate: 1 mL/min
Temperature: 50 °C
Detection: UV @ 282 nm
Sample: 1. 6-Gingerol
 2. 8-Gingerol + isomer
 3. 6-Shogaol
 4. 10-Gingerol
 5. 8-Shogaol
 6. 10-Shogaol

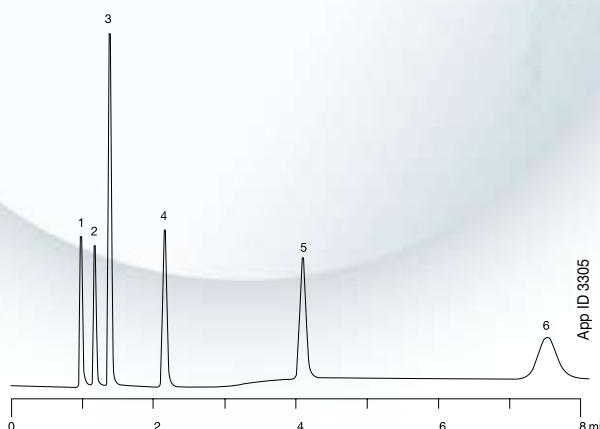
COLD MEDICINE



Column: Luna 5 μm C8(2)
Dimension: 150 x 4.6 mm
Part No.: 00F-4249-E0
Mobile Phase: Methanol/Acetonitrile
 with 0.1 % H_3PO_4 /Water
 with 0.1 % H_3PO_4 and
 0.1% Heptane
Flow Rate: 1.5 mL/min
Temperature: 22 °C
Detection: UV @ 214 nm
Sample: 1. Acetaminophen
 2. Pseudoephedrine
 3. Benzoic acid
 4. Chlorpheniramine
 5. Dextromethorphan

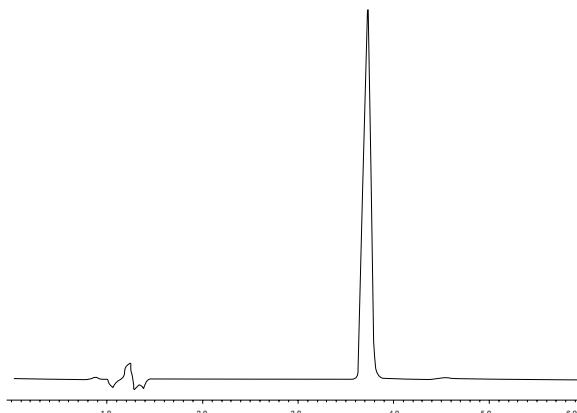
Luna C18(2), C8(2), C5 (cont.)

WATER SOLUBLE VITAMINS



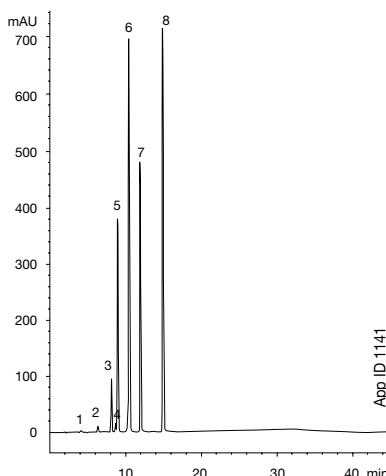
Column: Luna 5 μm C18(2)
Dimension: 150 x 4.6 mm
Part No.: 00F-4252-E0
Mobile Phase: 20 mM Potassium Phosphate, pH 3.0/Acetonitrile (95:5)
Flow Rate: 1.5 mL/min
Temperature: 22 °C
Detection: UV @ 214 nm
Sample:
1. Thiamine
2. Cyanocobalamin (Vitamin B12)
3. Ascorbic acid
4. Pantothenic acid
5. Niacinamide
6. p-Aminobenzoic acid

USP METHOD: LORAZEPAM TABLETS



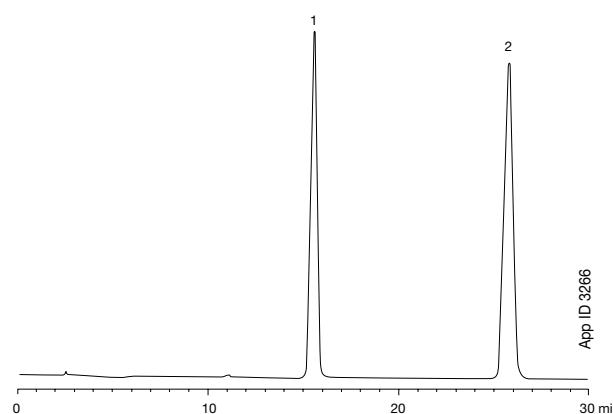
Column: Luna 5 μm C18(2)
Dimension: 250 x 4.6 mm
Part No.: 00G-4252-E0
Mobile Phase: Water/Methanol/Acetic acid (54:44:2)
Flow Rate: 2 mL/min
Temperature: 22 °C
Detection: UV @ 254 nm
Sample: 1. Lorazepam

PHARMACEUTICAL PRESERVATIVES



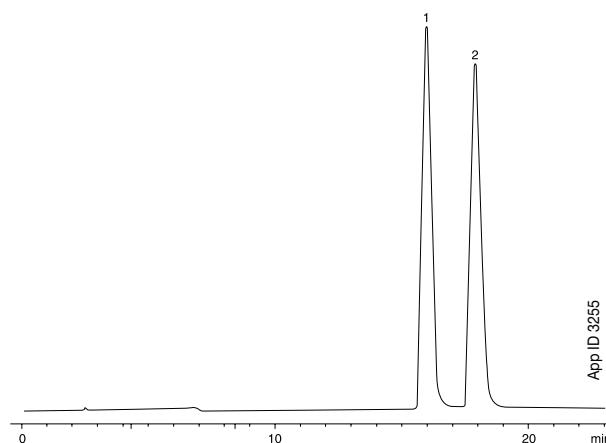
Column: Luna 5 μm C5
Dimension: 150 x 4.6 mm
Part No.: 00F-4043-E0
Mobile Phase: A: 0.5 % Acetic acid in water /Acetonitrile (80:20)
B: 0.5 % Acetic acid in water /Acetonitrile (20:80)
Gradient: A/B (100:0) to A/B (0:100) in 30 min
Flow Rate: 1 mL/min
Temperature: 25 °C
Detection: UV @ 254 nm
Sample:
1. Propylparaben impurity
2. Benzyl alcohol
3. Phenol
4. Benzoic acid
5. Methylparaben
6. Benzaldehyde
7. Ethylparaben
8. Propylparaben

USP METHOD: MINOXIDIL



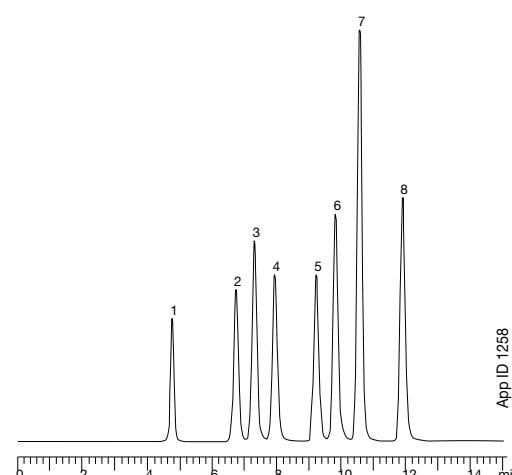
Column: Luna 5 μm C18(2)
Dimension: 250 x 4.6 mm
Part No.: 00G-4252-E0
Mobile Phase: Methanol/Water/Acetic acid with 7 mM Docusate sodium pH 3.0 (69.3:29.7:1)
Flow Rate: 1 mL/min
Temperature: 22 °C
Detection: UV @ 254 nm
Sample: 1. Medroxyprogesterone acetate
2. Minoxidil

USP METHOD: IMPIRAMINE



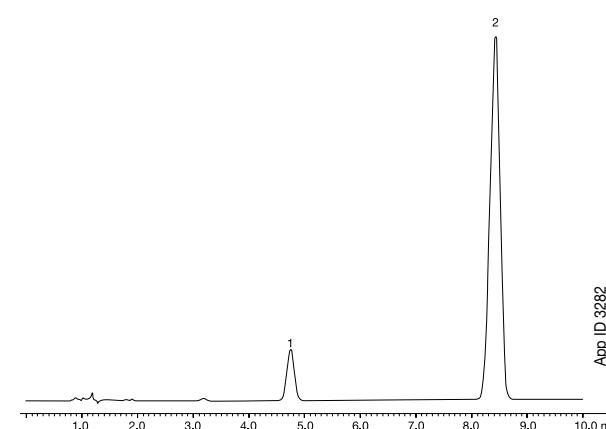
Column: Luna 5 μ m C18(2)
Dimension: 250 x 4.6 mm
Part No.: 00G-4252-E0
Mobile Phase: 0.06 M Sodium perchlorate, pH 2.0/Acetonitrile/Triethylamine (62.5:37.5:0.1)
Flow Rate: 1.5 mL/min
Temperature: 22 °C
Detection: UV @ 269 nm
Sample: 1. Imipramine
2. Desipramine

FATTY ACIDS



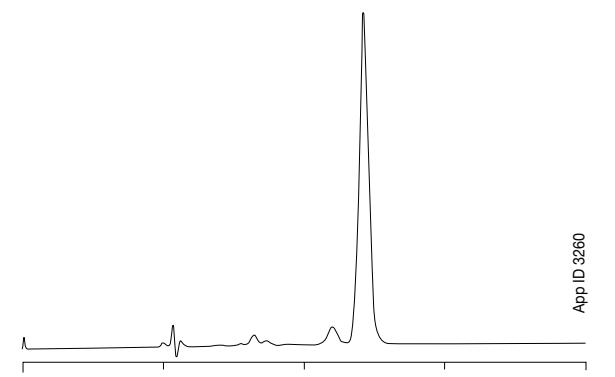
Column: Luna 5 μ m C8(2)
Dimension: 150 x 4.6 mm
Part No.: 00F-4249-E0
Mobile Phase: A: Acetonitrile
B: Water (18 Mohms DI)
Gradient: A/B (70:30) to A/B (90:10) in 10 min, A/B (90:10) to A/B (70:30) in 2 min, hold for 4 min
Flow Rate: 0.3 mL/min
Detection: Evaporative Light Scattering (ELSD)
Temperature: 22 °C
Sample: 1. Lauric acid
2. Myristic acid
3. Palmitoleic acid
4. Linoleic acid
5. Palmitic acid
6. Oleic acid
7. Heptadecanoic acid
8. Stearic acid

USP METHOD: NAPROXEN TABLETS



Column: Luna 5 μ m C18(2)
Dimension: 150 x 4.6 mm
Part No.: 00F-4252-E0
Mobile Phase: Acetonitrile/Water/Glacial acid, pH 3.0 (50:49:1)
Flow Rate: 1.2 mL/min
Temperature: 22 °C
Detection: UV @ 254 nm
Sample: 1. Naproxen
2. Butyrophenone

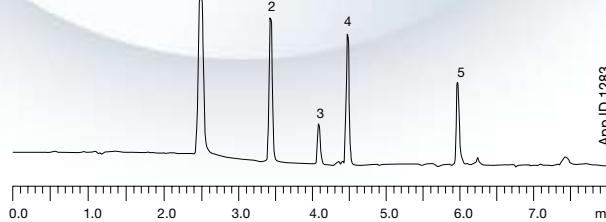
USP METHOD: ALBUTEROL TABLETS



Column: Luna 5 μ m C18(2)
Dimension: 150 x 4.6 mm
Part No.: 00F-4252-E0
Mobile Phase: Methanol/Water with 5 mM Hexane sulfonic acid and 1 % Glacial acetic acid (40:60)
Flow Rate: 1.5 mL/min
Temperature: 22 °C
Detection: UV @ 276 nm
Sample: 1. Albuterol

Luna C18(2), C8(2), C5 (cont.)

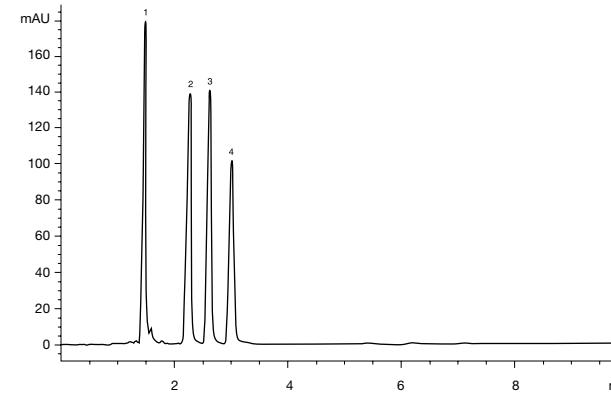
ANTIOXIDANTS



Column: Luna 5 µm C18(2)
Dimension: 100 x 4.6 mm
Part No.: 00D-4252-E0
Mobile Phase: A: Acetonitrile
 B: Phosphate Buffer
Gradient: A/B (30:70) to A/B (70:30) in 5 min
Flow Rate: 1 mL/min
Temperature: 22 °C
Detection: UV @ 254 nm
Sample:

- 1. PG
- 2. TBHQ
- 3. DMT
- 4. BHA
- 5. BHT

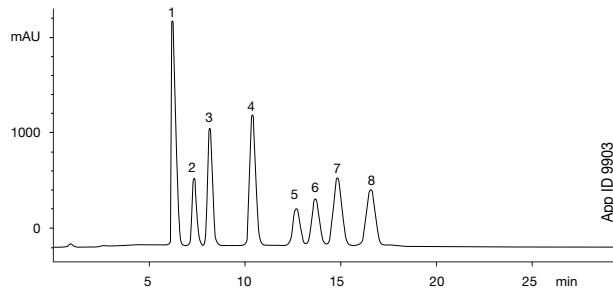
CEPHALOSPORIN ANTIBIOTICS



Column: Luna 5 µm C18
Dimension: 150 x 4.6 mm
Part No.: 00F-4041-E0
Mobile Phase: Water with 0.5% Acetic acid/
 Aetonitrile (85:15)
Flow Rate: 1.0 mL/min
Temperature: 22 °C
Detection: UV @ 254 nm
Sample:

- 1. Cefadroxil
- 2. Cephalexin
- 3. Cephradine
- 4. Cefachlor

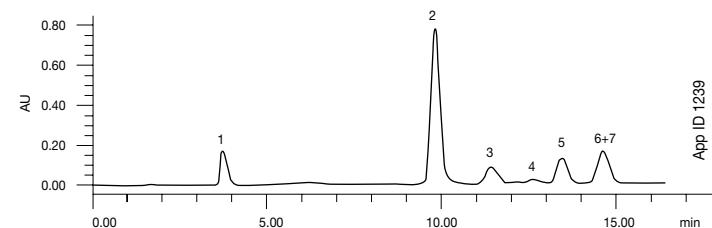
HORMONES



Column: Luna 5 µm C18
Dimension: 100 x 4.6 mm
Part No.: 00D-4041-E0
Mobile Phase: Water/Tetrahydrofuran/Acetonitrile
 (60:22.5:17.5)
Flow Rate: 1.2 mL/min
Temperature: 40 °C
Detection: UV @ 244 nm
Sample:

- 1. Norgestrel
- 2. syn-Norgestrel oxime
- 3. anti-Norgestrel oxime
- 4. Norgestrel acetate
- 5. syn-Norgestimate
- 6. syn-Norgestrel oxime diacetate
- 7. anti-Norgestimate
- 8. anti-Norgestrel oxime diacetate

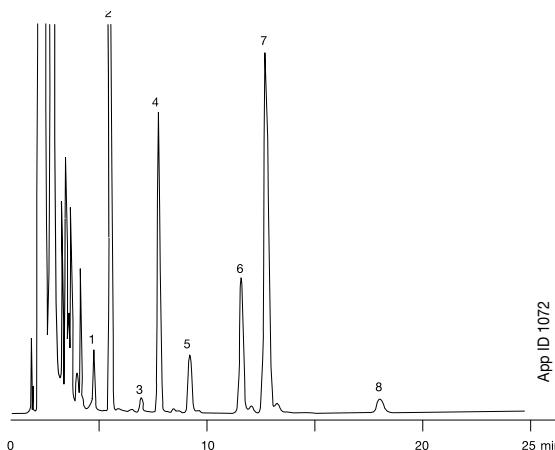
SUN SCREENS



Column: Luna 5 µm C18
Dimension: 250 x 4.6mm
Part No.: 00G-4041-E0
Mobile Phase: Methanol/Water (80:20)
Flow Rate: 1.5 mL/min
Temperature: 22 °C
Detection: UV @ 240 nm
Sample:

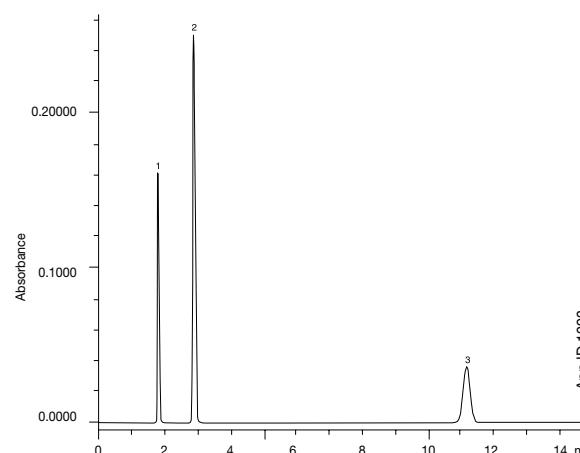
- 1. Benzophenone 3
- 2. DCHP
- 3. Parsol MCX
- 4. Homosalate 2
- 5. Octyl salicylate
- 6. Homosalate 1
- 7. Homosalate

SAW PALMETTO BERRY, P-BROMOPHENACYL ESTERS



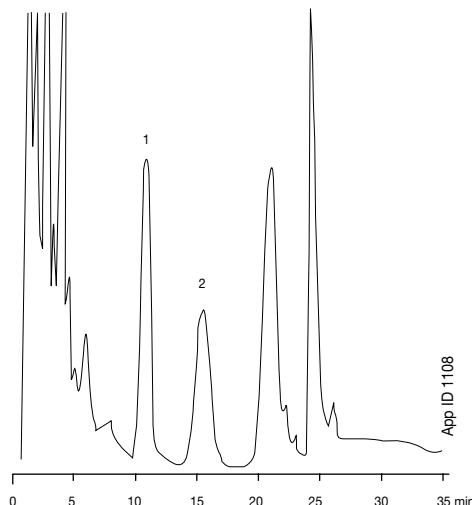
Column: Luna 3 μm C8(2)
Dimension: 150 x 4.6 mm
Part No.: 00F-4248-E0
Mobile Phase: Acetonitrile/Water (87:13)
Flow Rate: 1.5 mL/min
Temperature: 25 °C
Detection: UV @ 254 nm
Sample: 1. Capric acid
 2. Lauric acid
 3. Linolenic acid
 4. Myristic acid
 5. Linoleic acid
 6. Palmitic acid
 7. Oleic acid
 8. Stearic acid

ACETAMINOPHEN, USP METHOD



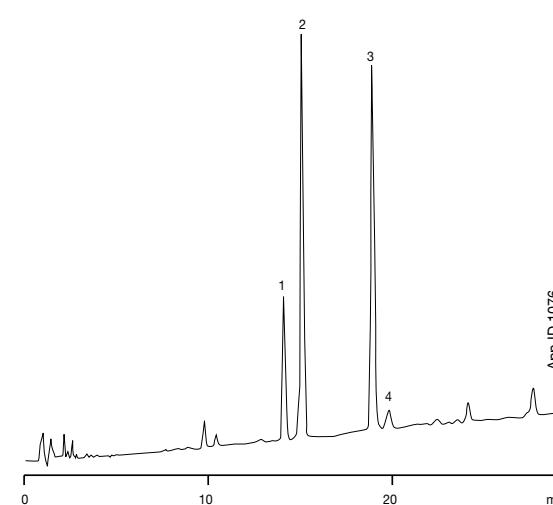
Column: Luna 5 μm C18(2)
Dimension: 150 x 4.6 mm
Part No.: 00F-4252-E0
Mobile Phase: Water/Methanol/Acetic Acid (69:28:3)
Flow Rate: 1.5 mL/min
Temperature: 45 °C
Detection: UV @ 275 nm
Sample: 1. Acetaminophen
 2. Caffeine
 3. Benzoic Acid

CYCLOSPORIN - IMMUNOSUPPRESSANTS



Column: Luna 5 μm C18(2)
Dimension: 150 x 4.6 mm
Part No.: 00F-4252-E0
Mobile Phase: Acetonitrile/Water, pH 3.1 w/1 mM H₃PO₄ (70:30)
Flow Rate: 1.3 mL/min
Temperature: 75 °C
Detection: UV @ 210 nm
Sample: 1. Cyclosporin A
 2. Cyclosporin D

CAPSAICIN



Column: Luna 5 μm C18(2)
Dimension: 150 x 4.6 mm
Part No.: 00F-4252-E0
Mobile Phase: A: Acetonitrile/Water (35:65)
 B: Acetonitrile/Water (60:40)
Gradient: 100 % A in 1 min to 100 % B in 29 min
Flow Rate: 1.5 mL/min
Temperature: 75 °C
Detection: UV @ 227 nm
Sample: 1. Nordihydrocapsaicin
 2. Capsaicin
 3. Dihydrocapsaicin
 4. Homocapsaicin

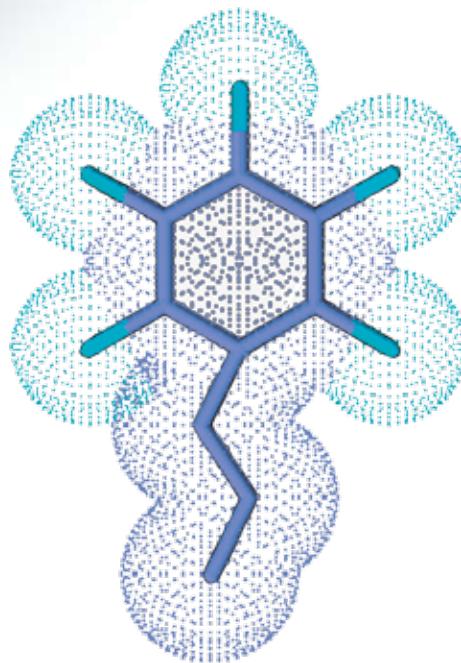
Luna PFP(2)

Powerful Selectivity for Reversed Phase Methods

Luna PFP(2) columns provide unique selectivity for highly polar compounds, complex natural products, isomers and other closely related compounds. This is achieved by using a propyl-linked pentafluorophenyl, which provides multiple retention mechanisms unique to typical reversed phase medias.

Luna PFP(2) selectivity is achieved through 4 mechanisms of interaction

- 1 Hydrogen Bonding
- 2 Dipole-Dipole Interactions
- 3 Aromatic and π - π Interactions
- 4 Hydrophobic

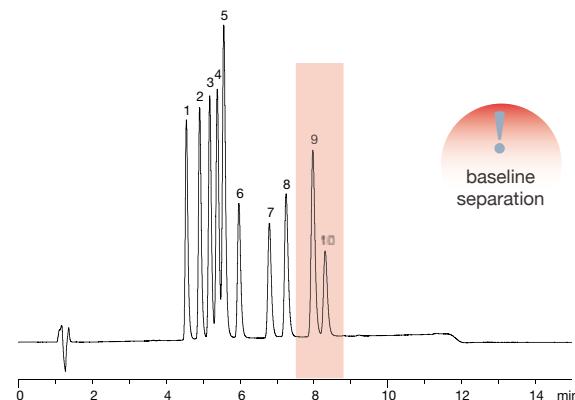


- » Achieve unique selectivity using four mechanisms of solute/stationary phase interactions
- » Extremely discerning for halogenated, aromatic, and conjugated compounds
- » Provides orthogonal selectivity, even using traditional reversed phase solutions

Halogens can radically increase the polarity of compounds, thus decreasing typical retention characteristics. Luna PFP(2) columns retain, discriminate, and separate halogens easily.

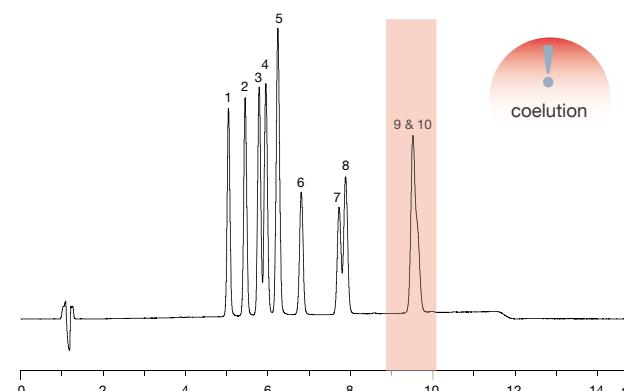
POSITIONAL ISOMERS OF HALOGENATED PHENOLS

Luna 3 μ m PFP(2)



Column: Luna 3 μ m PFP(2)
Dimension: 150 x 4.6 mm
Part No.: 00F-4447-E0
Mobile Phase: A: 0.1 % Formic acid in Water
B: 0.1 % Formic acid in Acetonitrile
Gradient: A/B (60:40) to (50:50) in 10 min
Flow Rate: 1 mL/min
Temperature: 22 °C
Detection: UV @ 254 nm
Sample: 1. 2,3-Dimethylphenol
2. 2,5-Dimethylphenol
3. 2,6-Dimethylphenol
4. 3,4-Dimethylphenol
5. 3,5-Dimethylphenol
6. 2,5-Dichlorophenol
7. 2,6-Dichlorophenol
8. 3,4-Dichlorophenol
9. 3,5-Dichlorophenol
10. 2,4-Dibromophenol

Luna 3 μ m C18(2)



Column: Luna 3 μ m C18(2)
Dimension: 150 x 4.6 mm
Part No.: 00F-4251-E0
Mobile Phase: A: 0.1 % Formic acid in Water
B: 0.1 % Formic acid in Acetonitrile
Gradient: A/B (60:40) to (50:50) in 10 min
Flow Rate: 1 mL/min
Temperature: 22 °C
Detection: UV @ 254 nm
Sample: 1. 2,3-Dimethylphenol
2. 2,5-Dimethylphenol
3. 2,6-Dimethylphenol
4. 3,4-Dimethylphenol
5. 3,5-Dimethylphenol
6. 2,5-Dichlorophenol
7. 2,6-Dichlorophenol
8. 3,4-Dichlorophenol
9. 3,5-Dichlorophenol
10. 2,4-Dibromophenol

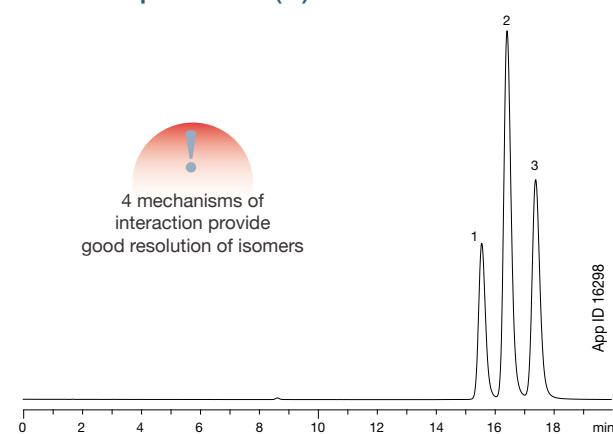
Isomeric Compounds

Positional changes on an analyte of interest may effect the compound's dipole moment. This change can be readily noticed by the highly electronegative fluorine (F) atom and other retention mechanisms.

Excellent choice for positional isomers due to the multiple retention characteristics of Luna PFP(2) columns

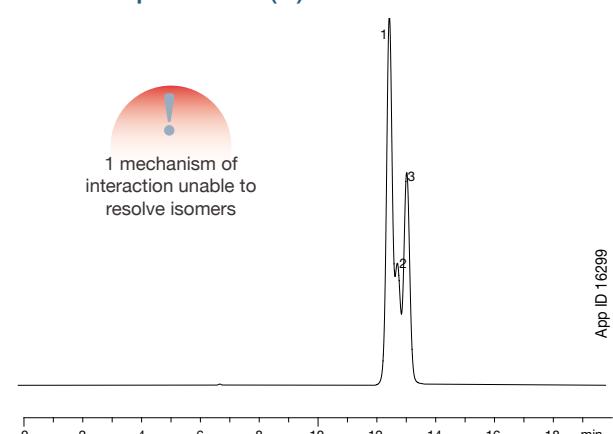
POSITIONAL ISOMERS OF METHYLACETOPHENON

Luna 3 μ m PFP(2)



Column: Luna 3 μ m PFP(2)
Dimension: 150 x 4.6 mm
Part No.: 00F-4447-E0
Mobile Phase: Water/ Methanol (50:50)
Flow Rate: 1 mL/min
Temperature: 22 °C
Detection: UV @ 254 nm
Sample: 1. o-Methylacetophenone
2. m-Methylacetophenone
3. p-Methylacetophenone

Luna 3 μ m C18(2)



Column: Luna 3 μ m C18(2)
Dimension: 150 x 4.6 mm
Part No.: 00F-4251-E0
Mobile Phase: Water/ Methanol (50:50)
Flow Rate: 1 mL/min
Temperature: 22 °C
Detection: UV @ 254 nm
Sample: 1. o-Methylacetophenone
2. m-Methylacetophenone
3. p-Methylacetophenone

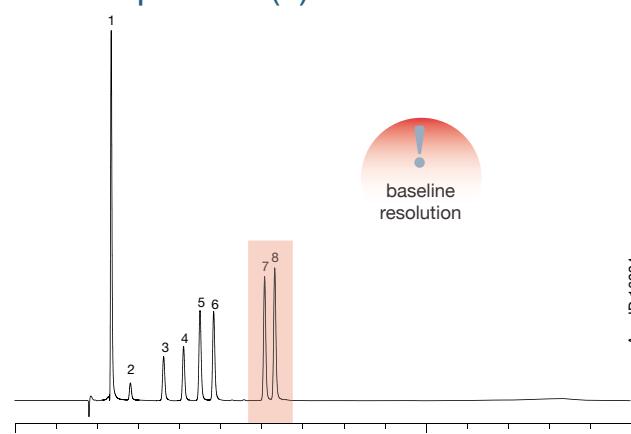
Aromatic Compounds

Aromatic compounds show unique retention characteristics on Luna PFP(2) compared to traditional reversed phase columns. The presence of the aromatic benzene ring in Luna PFP(2) increases the relative attraction between the stationary phase and aromatic analytes, leading to increased retention for these types of compounds.

Closely related polyphenolic compounds are readily separated with Luna PFP(2) columns

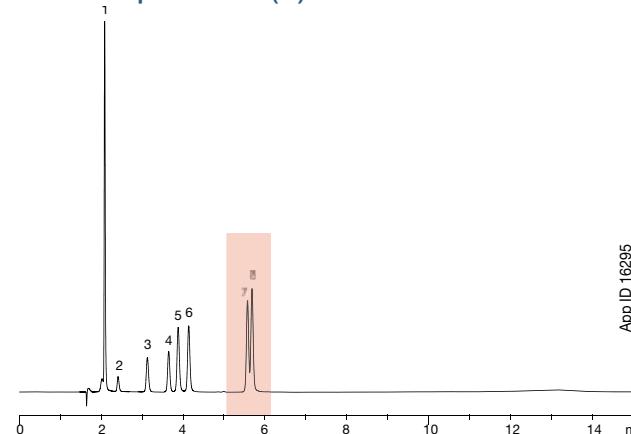
CATECHINS

Luna 3 μ m PFP(2)



Column: Luna 3 μ m PFP(2)
Dimension: 150 x 4.6 mm
Part No.: 00F-4447-E0
Mobile Phase: A: 0.1 % Formic acid in Water
B: 0.1 % Formic acid in Acetonitrile
Gradient: A/B (80:20) to (55:45) in 10 min
Flow Rate: 1 mL/min
Temperature: 22 °C
Detection: UV @ 280 nm
Sample: 1. Gallic acid
2. Epigallo catechin
3. Catechin
4. Epicatechin
5. Epigallocatechin gallate
6. Gallocatechin gallate
7. Epicatechin gallate
8. Catechin gallate

Luna 3 μ m C18(2)



Column: Luna 3 μ m C18(2)
Dimension: 150 x 4.6 mm
Part No.: 00F-4251-E0
Mobile Phase: A: 0.1 % Formic acid in Water
B: 0.1 % Formic acid in Acetonitrile
Gradient: A/B (80:20) to (55:45) in 10 min
Flow Rate: 1 mL/min
Temperature: 22 °C
Detection: UV @ 280 nm
Sample: 1. Gallic acid
2. Epigallo catechin
3. Catechin
4. Epicatechin
5. Epigallocatechin gallate
6. Gallocatechin gallate
7. Epicatechin gallate
8. Catechin gallate

Luna Phenyl-Hexyl

Engineered for Stability

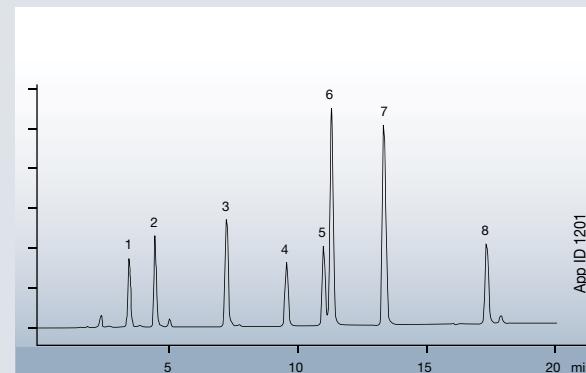
The patented Luna Phenyl-Hexyl is a reproducible, extremely stable phenyl phase. Most other phenyl phases use a short propyl (3 carbon) linker, which limits the phase stability. This patented phase uses a hexyl (6 carbon) linker to attach the phenyl group to the silica surface.

The result:

- » Highly reproducible phenyl phase
- » Dual selectivity of both phenyl phase and a short alkyl phase (such as a C8)
- » Excellent retention of amine and polar aromatic compounds
- » 1.5 to 10 pH stability for 10,000 hours

ANTIBACTERIALS: COMPARISON OF PHENYL COLUMNS**

Phenomenex® Luna® 5 µm Phenyl-Hexyl



Conditions same for all columns

Dimension: 150 x 4.6 mm

Mobile Phase: A: 20 mM KH₂PO₄, pH 2.5
B: Acetonitrile

Gradient: A/B (80:20) to A/B (75:25) in 5 min, then to A/B (55:45) in 15 min

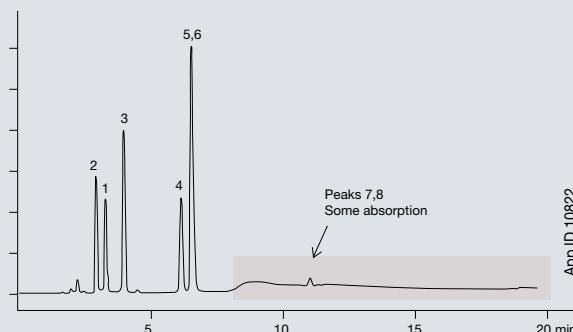
Flow Rate: 1.0 mL/min

Detection: UV @ 254 nm

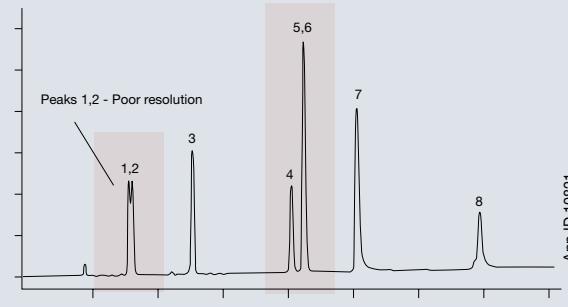
Temperature: 22 °C

Sample:
1. Carbadox
2. Thiamphenicol
3. Furazolidone
4. Oxolinic acid
5. Sulfadimethoxine
6. Sulfaquinoxaline
7. Nalidixic acid
8. Piromidic acid

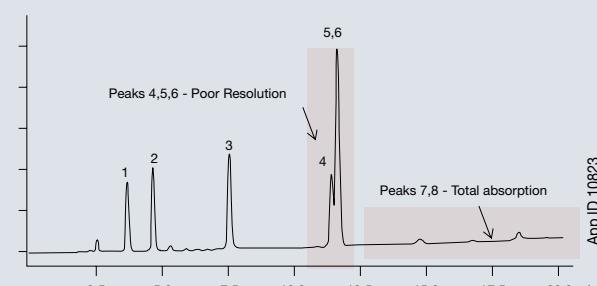
Waters® Spherisorb® 5 µm Phenyl



Agilent Technologies® Zorbax® 5 µm SB-Phenyl

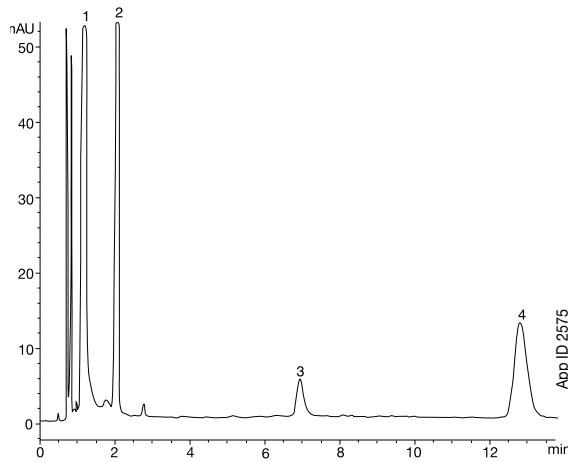


Agilent Technologies® Zorbax® 5 µm Phenyl



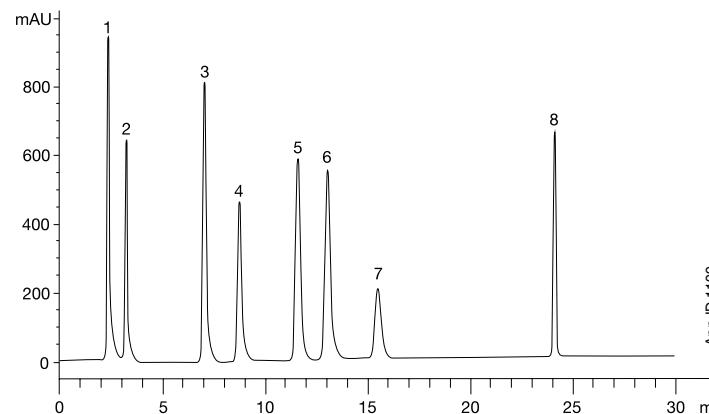
** The comparative data presented here may not be representative for all applications.

COUGH AND COLD-USP METHOD



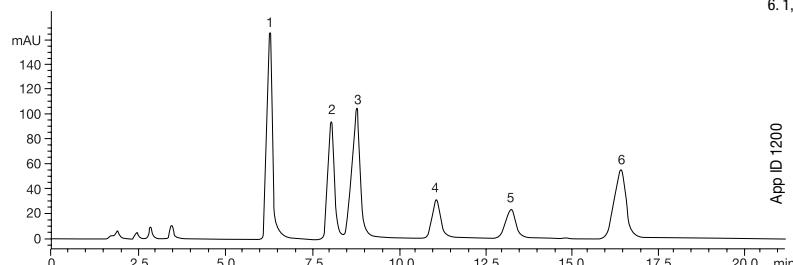
Column: Luna 5 μ m Phenyl-Hexyl
Dimension: 150 x 4.6 mm
Part No.: 00F-4257-E0
Mobile Phase: Methanol/Water with 0.1 % H_3PO_4 and 0.1 % Heptane Sulfonate/Acetonitrile with 0.1 % H_3PO_4 (35:55:10)
Flow Rate: 2.05 mL/min
Detection: UV @ 214 nm
Temperature: 22 °C
Sample:
 1. Acetaminophen
 2. Pseudoephedrine
 3. Chlorphenamine
 4. Dextromethorphan

FOOD ADDITIVES



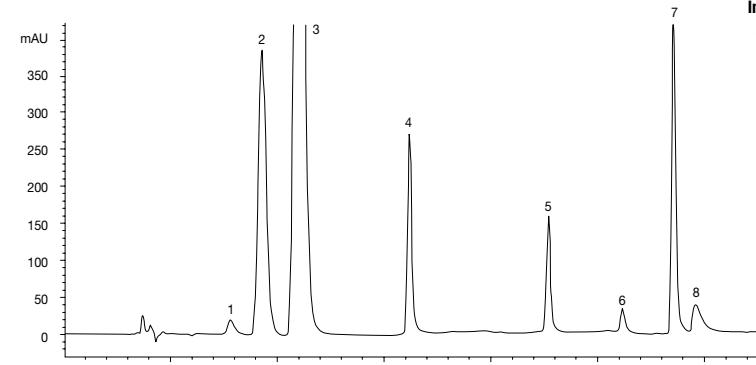
Column: Luna 5 μ m Phenyl-Hexyl
Dimension: 150 x 4.6 mm
Part No.: 00F-4257-E0
Mobile Phase: A: 50 mM KH_2PO_4 + 0.1% H_3PO_4 , B: Acetonitrile
Gradient: A/B (75:25) to A/B (25:75) in 18 min, hold at A/B (25:75) for 12 min
Flow Rate: 1.0 mL/min
Detection: UV @ 230 nm
Temperature: 22 °C
Injection: 20 μ L
Sample:
 1. Saccharin
 2. p-Hydroxybenzoic acid
 3. Sorbic acid
 4. p-Hydroxybenzoic acid methyl ester
 5. Dehydroacetic acid
 6. p-Tolue acid
 7. p-Hydroxybenzoic acid ethyl ester
 8. n-Propyl p-hydroxybenzoate

CHLOROBENZENES



Column: Luna 5 μ m Phenyl-Hexyl
Dimension: 150 x 4.6 mm
Part No.: 00F-4257-E0
Mobile Phase: A: Water
 B: Acetonitrile
Gradient: A/B (60:40) to A/B (45:55) in 10 min
Flow Rate: 1.0 mL/min
Detection: UV @ 254 nm
Temperature: 22 °C
Sample:
 1. Chlorobenzene
 2. 1,2-Dichlorobenzene
 3. 1,4-Dichlorobenzene
 4. 1,2,3-Trichlorobenzene
 5. 1,3,5-Trichlorobenzene
 6. 1,2,3,4-Tetrachlorobenzene

COUGH AND COLD MEDICINE



Column: Luna 3 μ m Phenyl-Hexyl
Dimension: 75 x 4.6 mm
Part No.: 00C-4256-E0
Mobile Phase: A: Acetonitrile
 B: 20 mM KH_2PO_4 / Methanol(80:20) pH 9.0
Gradient: A/B (0:100) to A/B (80:20) in 5 min
Flow Rate: 1.0 mL/min
Detection: UV @ 214 nm
Temperature: 22 °C
Injection: 20 μ L
Sample:
 1. p-Aminophenol
 2. Benzoic acid
 3. Acetaminophen
 4. Pseudoephedrine
 5. Butyl paraben
 6. Chlorphenamine
 7. Diphenhydramine
 8. Dextromethorphan

Luna (CN) Cyano

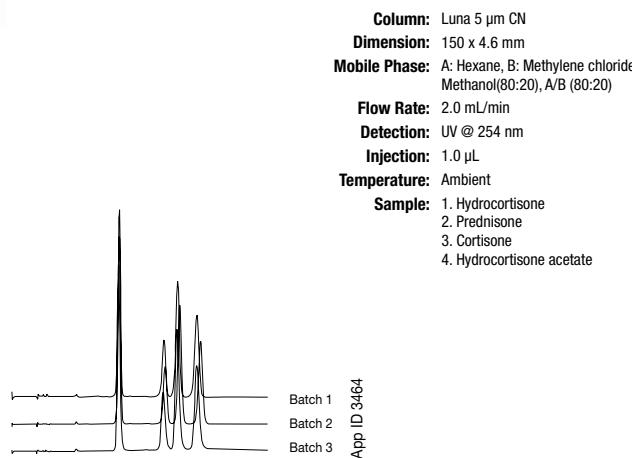
Proven Reproducibility

Luna CN columns were developed to provide reproducible chromatography from run-to-run, column-to-column and batch-to-batch. Luna® high-purity silica provides a ridged and dense column bed that allows for improved CN bonding techniques to make a stable CN phase.

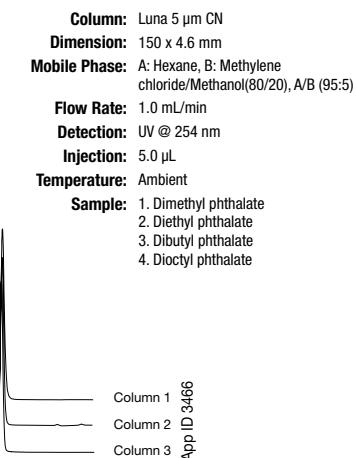
The result:

- » One of the most stable CN columns under both reversed phase or normal phase conditions.
- » Reproducible from run-to-run, column-to-column, batch-to-batch.
- » pH stable from 1.5 to 7.0

Batch-to-Batch Reproducibility

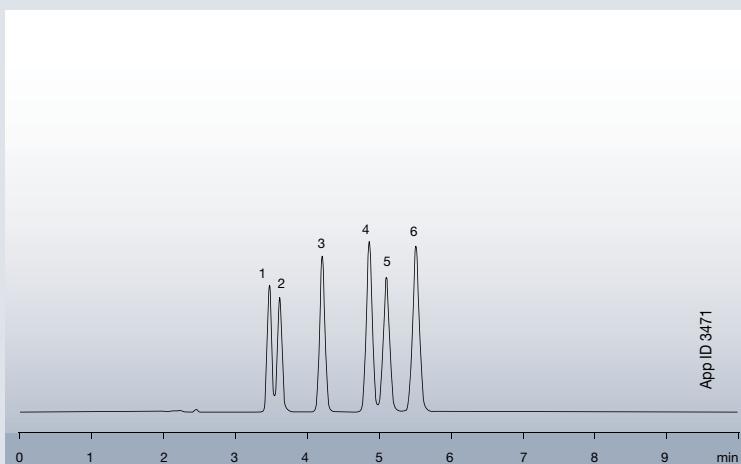


Column-to-Column Reproducibility

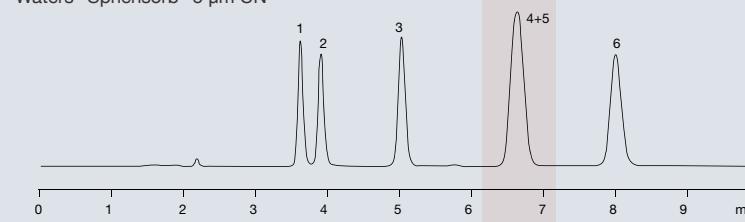


Phthalate Esters: A Comparison of CN Columns**

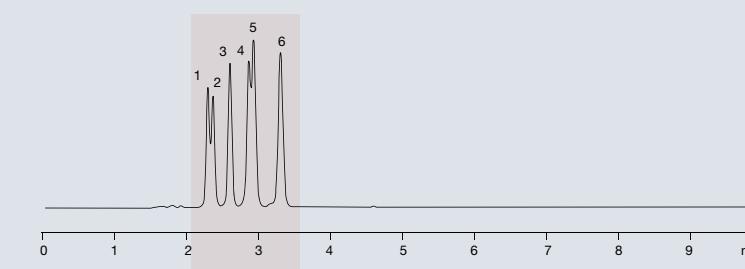
Phenomenex® Luna® 5 μm CN



Waters® Spherisorb® 5 μm CN



Agilent Technologies® Zorbax® 5 μm SB-CN



Conditions same for all columns

Dimension: 150 x 4.6 mm
Mobile Phase: A: Hexane, B: Methylene chloride/Methanol (80:20), A/B (99:1)
Flow Rate: 1.0 mL/min
Detection: UV @ 254 nm
Temperature: Ambient
Sample: 1. Di-n-octyl phthalate
2. Bis (2-Ethylhexyl) phthalate
3. Butylbenzyl phthalate
4. Di-n-butyl phthalate
5. Diethyl phthalate
6. Dimethyl phthalate

** The comparative data presented here may not be representative for all applications.

Luna (NH₂) Amino

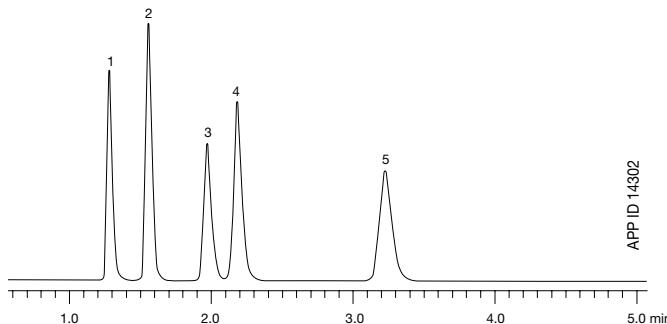
Developed for Ruggedness

Luna NH₂ columns were developed to provide improved amino column lifetime. Column life for most amino columns can be problematic as the amino bonding easily strips off the silica. Luna NH₂ columns, however, show good bonded phase stability under both normal and reversed phase modes and across a pH range of 1.5 to 11.0. Such a broad pH range indicates the bonded phase ruggedness and the density of the bonded phase coverage.

The result:

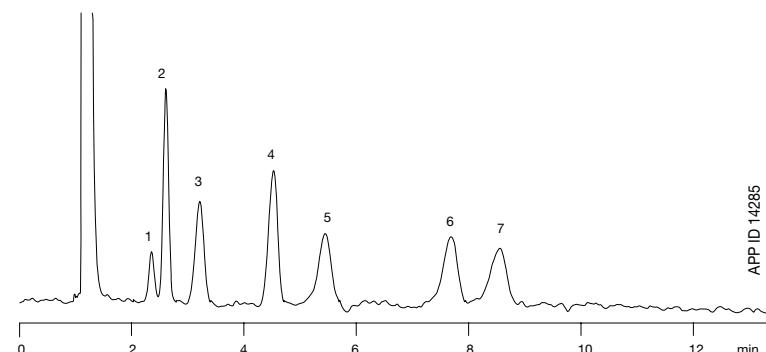
- » Long lifetimes and low phase bleed for more reproducible methods
- » Excellent retention of simple sugars, complex sugars, sugar alcohols by reverse phase conditions and hydrogen bonding compounds under normal phase conditions
- » pH stable from 1.5 to 11.0
- » Stable in 100 % aqueous mobile phases

NUCLEIC ACID BASES



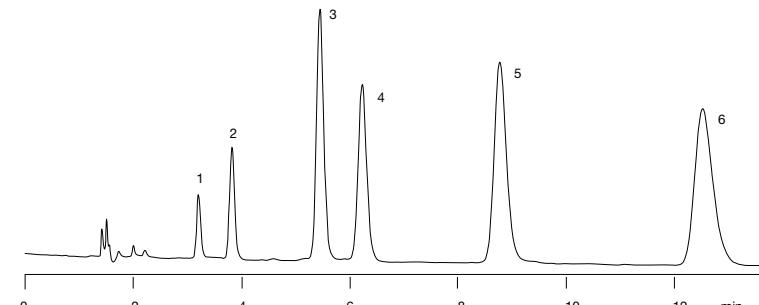
Column: Luna 5 μm NH₂
Dimension: 150 x 4.6 mm
Part No.: 00F-4378-E0
Mobile Phase: Acetonitrile/Water (80:20)
Flow Rate: 1.0 mL/min
Detection: UV @ 254 nm
Temperature: 40 °C
Sample: 1. Thymine
2. Uracil
3. Cytosine
4. Adenine
5. Guanosine

SIMPLE SUGARS



Column: Luna 5 μm NH₂
Dimension: 250 x 4.6 mm
Part No.: 00G-4378-E0
Mobile Phase: Acetonitrile/Water (80:20)
Flow Rate: 3.0 mL/min
Detection: RI
Temperature: 40 °C
Sample: 1. Xylose
2. Fructose
3. Glucose
4. Sucrose
5. Maltose
6. Melibiose
7. Raffinose

STEROIDS



Column: Luna 5 μm NH₂
Dimension: 250 x 4.6 mm
Part No.: 00G-4378-E0
Mobile Phase: Hexane/Ethanol (85:15)
Flow Rate: 2.0 mL/min
Detection: UV @ 240 nm
Temperature: 22 °C
Sample: 1. 11-Ketoprogesterone
2. 11-Hydroxyprogesterone
3. Cortisone acetate
4. Prednisolone 21-acetate
5. Cortisone
6. Prednisolone

Luna HILIC

Increase MS-Sensitivity and Retention for Polar Compounds

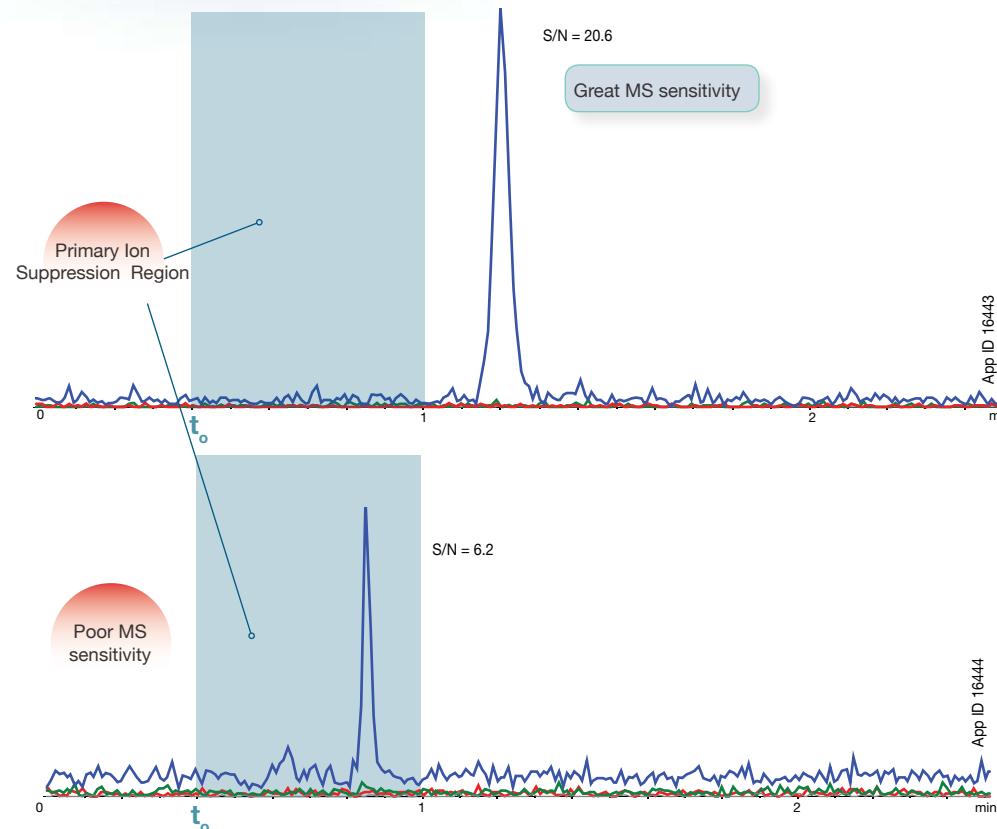
Luna HILIC columns retain a water-enriched layer on the surface of the silica. This water layer facilitates the transfer of polar compounds into the stationary phase for increased retention.

The result:

- » Superior retention of polar compounds
- » Improve mass spec sensitivity
- » Increased laboratory throughput and productivity

Improve Mass Spec Sensitivity

Luna HILIC columns allow low level polar metabolites to be retained on column past the critical ion suppression zone, allowing: Increased MS sensitivity and Higher signal-to-noise ratio (S/N).



Separation is achieved through the partitioning of polar solutes from the high concentration, water-miscible, organic mobile phase into the hydrophilic surface environment. Polar solutes exhibit increased retention, and elute in the order of increasing hydrophilicity.

POLAR COMPOUND IN HILC MODE

Column: Luna 3 μ m HILIC
Dimension: 100 x 2.0 mm
Part No.: 00D-4449-B0
Mobile Phase: Acetonitrile / 100 mM Ammonium Formate, pH 3.2 (90:10)
Flow Rate: 0.4 mL/min
Detection: Mass Spectrometer (MS)
Temperature: Ambient
Sample: Bamethan

POLAR COMPOUND IN C18 REVERSED PHASE

Column: Gemini 3 μ m C18
Dimension: 100 x 2.0 mm
Part No.: 00D-4435-B0
Mobile Phase: 0.1 % Formic Acid / Acetonitrile (97:3)
Flow Rate: 0.4 mL/min
Detection: Mass Spectrometer (MS)
Temperature: Ambient
Sample: Bamethan

Ion Suppression Region is from 0.5-1.0 min

$$t_o \approx 0.5 \text{ min}$$

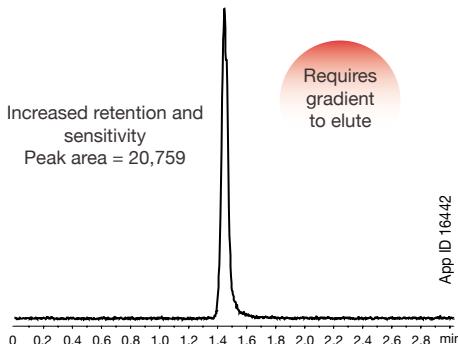
$$k' = 1 = \frac{t_R - t_o}{t_o} = \frac{1 - 0.5}{0.5}$$

Retain Polar Compounds

Highly polar compounds such as ribavirin may be poorly retained on reversed phase columns. HILIC techniques will increase polar compound retention and sensitivity.

RIBAVIRIN ON LUNA HILIC

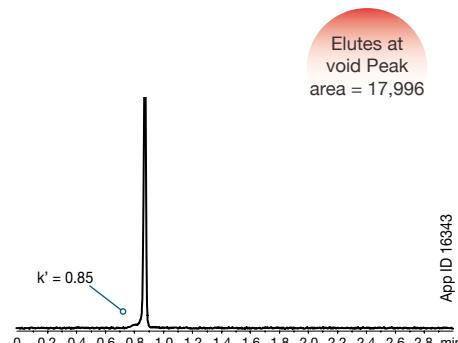
0.5 ng on column



Column: Luna 3 μ m HILIC
Dimension: 100 x 2.0 mm
Part No.: 00D-4449-80
Mobile Phase: A: Acetonitrile/100 mM Ammonium Formate, pH 3.2 (90:10)
B: Acetonitrile/20 mM Ammonium Formate, pH 3.2 (50:50)
Gradient: 100 % A for 3 min, then 100 % B to 4.5 min, switch to 100 % A for 10 min
Flow Rate: 0.4 mL/min
Detection: Mass Spectrometer (MS)
Temperature: Ambient
Sample: 1. Ribavirin (MRM: 245.2/113.2)

RIBAVIRIN ON C18

0.5 ng on column



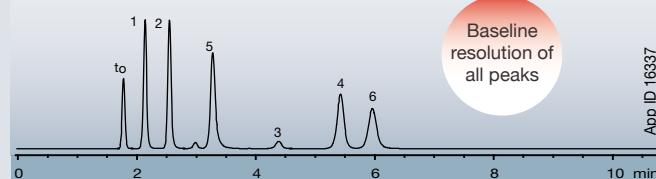
Column: Gemini 5 μ m C18
Dimension: 100 x 2.0 mm
Part No.: 00D-4435-B0
Mobile Phase: Acetonitrile with 0.1 % v/v Formic Acid/
Water with 0.1 % v/v % Formic Acid (3:97)
Flow Rate: 0.4 mL/min
Detection: Mass Spectrometer (MS) (ambient)
Temperature: Ambient
Sample: 1. Ribavirin (MRM: 245.2/113.2)

Unique HILIC Selectivity

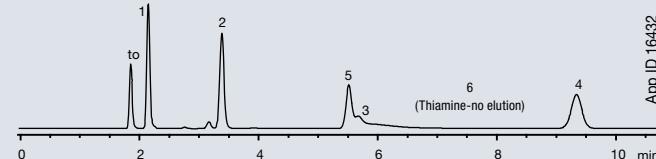
Not all HILIC columns are alike. Luna HILIC columns deliver on the exacting standards you've come to trust from the Luna product line.

HILIC COLUMN COMPARISON**

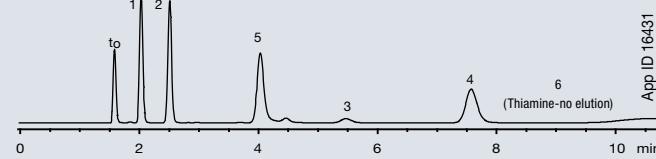
Phenomenex® Luna® 5 μ m



Waters® Atlantis® 5 μ m HILIC



SeQuant 5 μ m ZIC®-HILIC



Conditions same for all columns:

Column: As noted
Dimension: 150 x 4.6 mm
Mobile Phase: Acetonitrile/100 mM Ammonium Formate, pH 3.2 (90:10)
Flow Rate: 1.0 mL/min
Detection: UV @ 260 nm
Sample: 1. PABA
2. Nicotinamide
3. Riboflavin
4. Nicotinic Acid
5. Pyridoxine
6. Thiamine

** The comparative data presented here may not be representative for all applications.

Luna SCX

Develop Robust Strong Cation Exchange Methods

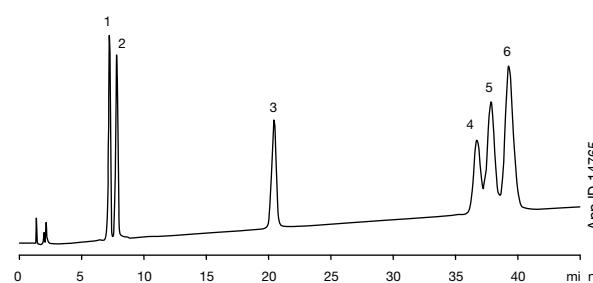
Luna SCX columns provide excellent resolution and peak shape of basic, cationic compounds. However, most SCX columns show poor peak shape and bad resolution causing many chromatographers to ignore this important phase for small molecule method development, until now.

The result:

- » Resolving power and sharp peak shape to separate complex cationic/basic and nitrogen containing compounds
- » Benzene sulfonic acid ligand provides mixed-mode interaction improving separation for 2D peptide applications
- » Excellent first dimension of 2D LC applications

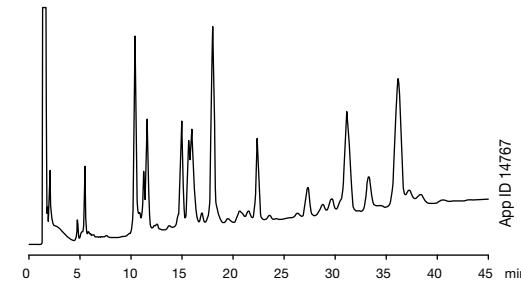
Luna SCX columns contain a benzene sulfonic acid ligand providing ion exchange reversed phase, and aromatic interactions. In combination with the ultra pure silica, Luna SCX columns are a stable, robust phase for strong cation exchange chromatography.

PEPTIDES



Column: Luna 5 μ m SCX
Dimension: 150 x 4.6 mm
Part No.: 00F-4398-E0
Mobile Phase: A: 20 mM Potassium Phosphate, 25 % Acetonitrile, pH 2.5
B: 20 mM Potassium Phosphate, 25 % Acetonitrile, 400 mM Potassium Chloride, pH 2.5
Gradient: A/B (95:5) to A/B (10:90) in 45 minute
Flow Rate: 1 mL/min
Temperature: 35 °C
Detection: UV @ 215 nm
Injection Volume: 2 μ L (5 μ g on column)
Sample: Peptide Mixture - Substance P
1. Fragment 5-11 (+1)
2. Fragment 4-11 (+1)
3. Fragment 2-11 (+2)
4. Fragment 1-9 (+3)
5. Intact (+3)
6. (ARG-PHE-TRP-LEU) (+3)

TRYPTIC DIGEST OF BOVINE CYTOCHROME C



Column: Luna 5 μ m SCX
Dimension: 150 x 4.6 mm
Part No.: 00F-4398-E0
Mobile Phase: A: 20 mM Potassium Phosphate, pH 2.5 /25 % Acetonitrile
B: 20 mM Potassium Phosphate, pH 2.5 /25 % Acetonitrile / 350 mM Potassium Chloride
Gradient: 100 % A to 100 % B in 50 minutes
Flow Rate: 1 mL/min
Temperature: 35 °C
Detection: UV @ 215 nm
Injection Volume: 50 μ L (20 μ g on column)
Sample: Bovine Cytochrome C trypsin digest



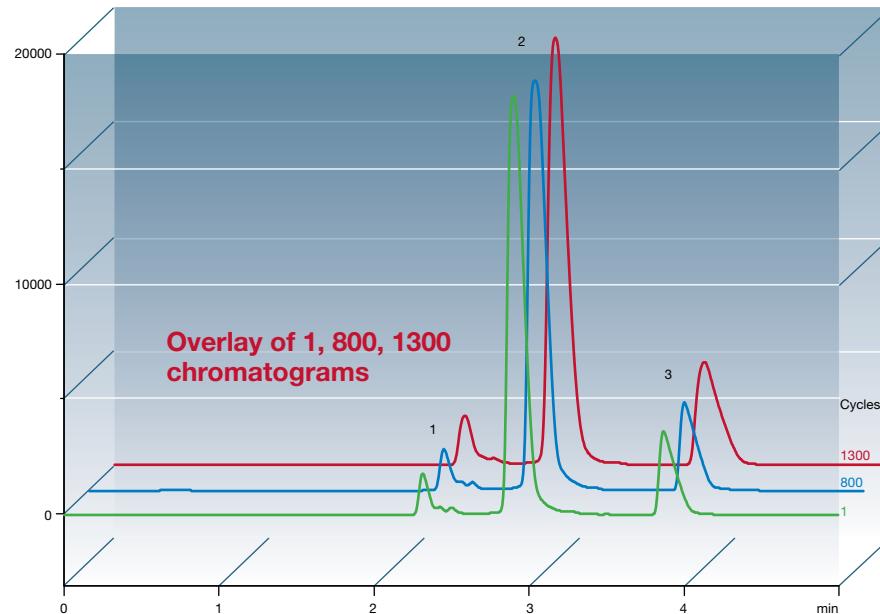
AXIA

Revolutionize Lab-Scale Purification

An advanced column packing and hardware design, Axia™ columns incorporate patent-pending Hydraulic Piston Compression technology to eliminate bed collapse as a source of failure in preparative columns. Using a single, controlled hydraulic compression, the piston assembly is locked in place without allowing the media to decompress or "relax," thus maintaining media and column bed integrity.

With Axia technology, the correct slurry amount and packing pressure are automated to give not only higher efficiency and sharper peaks, but also drastically reduced column-to-column variability. This will help improve longer column lifetime, column-to-column reproducibility, and recover higher compound purity with analytical like efficiency.

Axia™ Gradient Lifetime Study

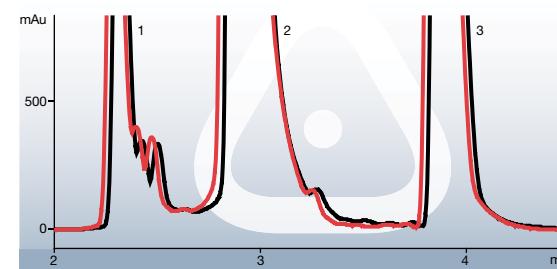


Columns: Luna® 5 µm C18(2) Axia™ Packed
Dimension: 50 x 21.2mm
Part No.: 00B-4252-P0-AX
Mobile Phase: A: 0.5% TFA in Water
 B: 0.5% TFA in Acetonitrile
Gradient: Linear 95:5 to 5:95 (A/B) over 7 min, hold 3 min
Injection Volume: 500 µL
Flow Rate: 30 mL/min
Temperature: Ambient
Detection: UV @ 254nm
Sample: 1. Triprolidine 1.6 mg
 2. Methacycline 16 mg
 3. Amitriptyline 5.25 mg

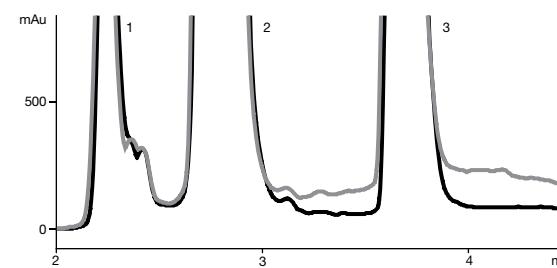


2006 R&D 100 Award Recipient

Compare Lifetime**



VS.



same for both columns except where noted

Column: Luna 5 µm C18(2) Axia™ Packed; Waters® Xterra® 5 µm Prep MS C18 OBD™
Dimension: 50 x 21.2 mm (Luna); 50 x 19 mm (Xterra®)
Mobile Phase: A: 0.5 % TFA in Water B: 0.5 % TFA in Acetonitrile
Gradient: Linear 95:5 (A/B) to 5:95 (A/B) over 7 min, hold 3 min
Flow Rate: 30 mL/min (Luna) 24 mL/min (Xterra®)
Temperature: Ambient
Detection: UV @ 254 nm
Sample: 1. Triprolidine 1.6 mg
 2. Methacycline 16 mg
 3. Amitriptyline 5.25 mg
† Same linear velocity

** The comparative data presented here may not be representative for all applications.

Fast LC

When you want Fast LC, you need BALANCE

The ever-increasing demand for high-throughput analysis of drug candidates during the early stages of drug discovery has generated an acute need for rapid methods of analysis.

3 Balanced Solutions to:

Balance Your Speed, Pressure, Efficiency and Selectivity

	Speed	Pressure	Efficiency	Selectivity	
1	High Speed Technology (HST) Columns	Fast	< 400 Bar	Highest	Several phases
2	MercuryMS Columns and Cartridges	Fast	< 400 Bar	High	Most phases
3	Monolithic Columns	Fastest	< 200 Bar	Good	Some phases

Developing ultra-fast and efficient methods for potential drugs has become a constant challenge for analysts. Use the chart above to determine the HPLC column that meets your performance needs.



Luna: High Speed Technology(HST) columns

- » High efficiency 2.5 µm particles on ultra-pure silica
- » Ultra-high performance results on your current HPLC
- » Easy method transfer
- » Orthogonal selectivity options

Luna HST columns are manufactured in specific dimensions utilizing new, highly controlled and robust packing technologies. The technology allows for consistent, high performance results on newer and existing HPLC instrumentation. Get the benefit of increased speed and efficiency with standard HPLC system pressure capabilities!

Luna HST can be used with your current standard HPLC and newer high performance systems so that there will be no need for time consuming method revalidation.

Luna HST 2.5 µm columns allow the scientist to reduce analysis time by increasing flow rates without a loss in performance.

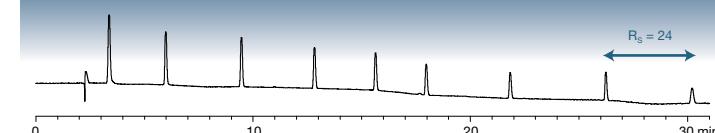


Speed Influence on Performance

HST Columns: 66 % Faster. No Loss in Resolution

Luna 5 µm C18(2) 250 x 4.6 mm

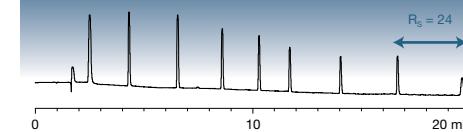
Flow Rate: 1.5 mL/min



App ID 16281

Luna 3 µm C18(2) 150 x 4.6 mm

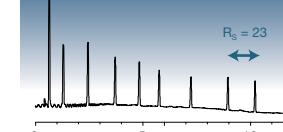
Flow Rate: 1.5 mL/min



App ID 16274

Luna 2.5 µm C18(2)-HST 100 x 2.0 mm

Flow Rate: 0.65 mL/min



Run time reduced
by 20 min
with virtually no effect
on resolution!

Column: Luna C18(2), particle size as noted

Dimension: as noted

Mobile Phase: A: Water B: Acetonitrile

Gradient: 90:10 (A/B) to 5:95 (A/B)

Flow Rate: As noted

Detection: UV @ 270 nm

Sample: Ketones C₃ to C₁₆

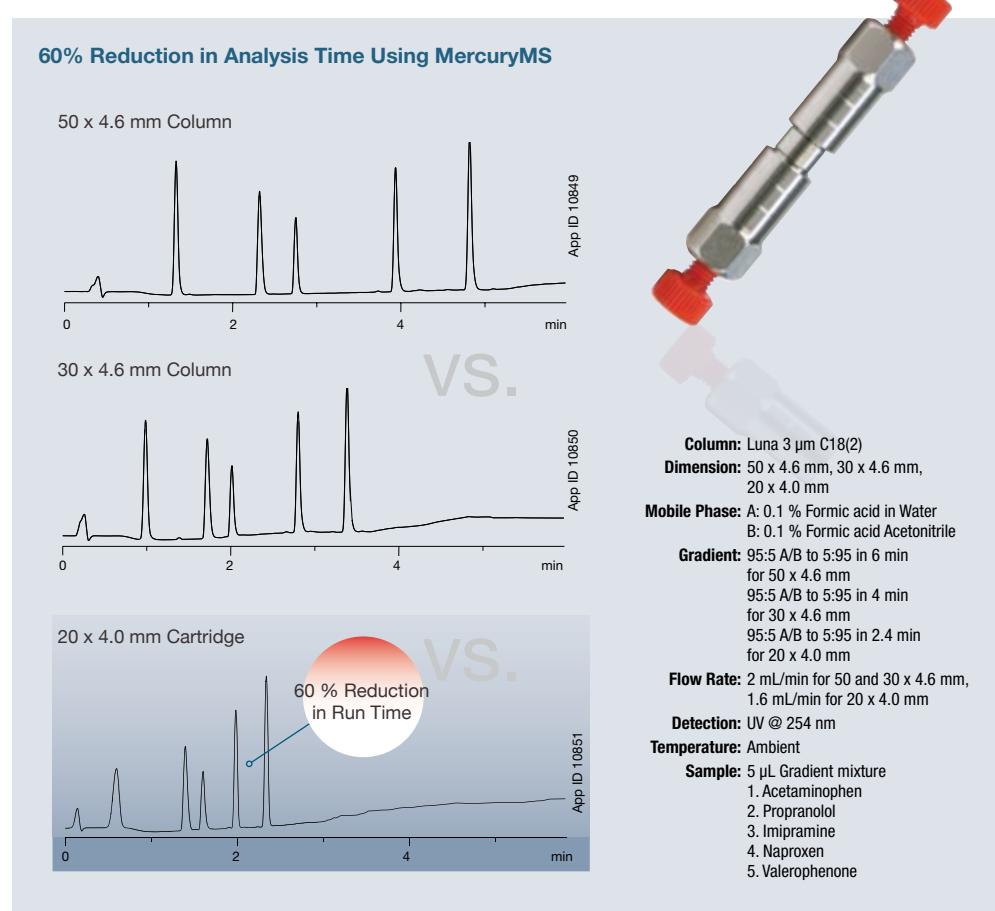
MercuryMS™: Columns and Cartridges

- Ultra-fast, low-cost analysis for high-throughput laboratories
- Packed with Luna®, Synergi™, and Gemini® material
- Short 10 and 20 mm cartridge formats use a new proprietary slurry packing process



Reduce Analysis Times by 60 %

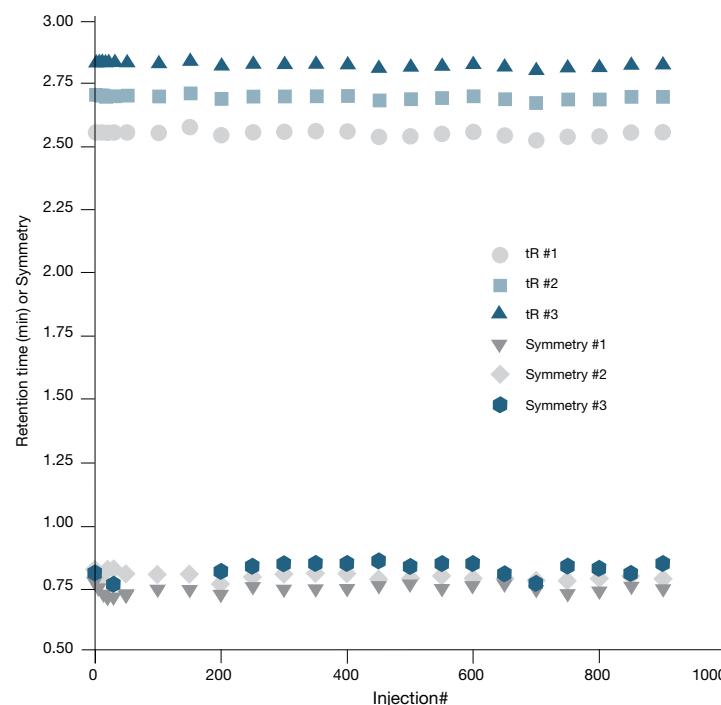
With the increasing emphasis on high sample throughput for screening combinatorial libraries as well as the need for overall faster cycle time, it has become necessary for the chromatographer to reduce analysis time while still maintaining acceptable resolution. As shown, retention times can be significantly reduced with a 20 x 4.0 mm MercuryMS cartridge column.



Rugged Durability for Over 1,000 Injections

When running long sequences, often involving hundreds of valuable samples, it is imperative that the column does not fail during the middle of the run. As shown, the Luna 3 µm C18(2) 20 x 2.0 mm MercuryMS cartridge was stable for over 1000 injections (over 20,000 column volumes of mobile phase) in this gradient assay, offering exceptional reliability for automated screening systems.

MercuryMS: Stable Over 1,000 Injections



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