

ANALYTICAL HPLC COLUMNS

50 Essential Selectivities for small organic molecules
Identification, Quantification & Purification
Ideal for Analysts & Chemists

[Uptisphere® CS Evolution](#)

[Uptisphere® Strategy™](#)

[Uptisphere® 120Å](#)

[puriFlash®](#)



 **Advion Interchim**
scientific

OUR SILICA TECHNOLOGIES

All of our Uptisphere® silicas (120Å, CS Evolution, Strategy™, puriFlash® & puriFlash® Bio) follow rigorous and innovative manufacturing processes. Base silicas are produced in ceramic reactors from standard particles for purification or totally free of all traces of metals for analysis. Each of the different synthesis steps is strictly controlled.

This rigor leads to extremely mechanically stable particles. The particle size and porosity distributions, as well as the specific surface areas, are perfectly defined and reproducible.

Our puriFlash® & puriFlash® Bio silicas are specifically designed to meet the requirements of preparative liquid chromatography. They combine quality and cost effectiveness.

Our silicas have three major advantages:

1. A perfect control of the surface state.

We physically or chemically modify the surface of the silica to choose the type, the amount of silanols or the overall surface energy according to the objective to be achieved.

2. Cylindrical pores.

The quantity of free silanols and their excellent accessibility allows to obtain a homogeneous and particularly dense functionalization (grafting). This results in a very good loading capacity and a good stability of these stationary phases under aggressive mobile phase conditions such as basic buffers.

3. High mechanical stability.

Our stationary phases can withstand multiple packings and de-packings without damaging the integrity of the substrate. They are the tool of choice for preparative chromatography.

Modified silicas:

The Laboratoire d'Étude des Techniques et des Instruments d'Analyse Moléculaire (LETIAM), a constituent unit of the analytical chemistry group of Paris Sud located at the IUT of Orsay, played a fundamental role in our reflection which led to the development of our stationary phases.

The laboratory of Sciences and Separative Methods - (SMS) of the Institute of Research in Fine Organic Chemistry - (IRCOF) has concretized part of our ideas by developing innovative synthesis schemes for the modification of our «Core-Shell» silicas Uptisphere® CS Evolution.

Today, we offer about 50 selectivities to meet the needs of analysts and chemists for the identification, quantification and purification of small organic molecules, peptides and proteins.

SILICAS FOR (U)HPLC, PREP-LC & FLASH COLUMNS

Uptisphere® CS Evolution

Core Shell columns for fast & highly efficient identification & quantification of small molecules.

Uptisphere® Strategy™

(U)HPLC, Analytical & prep LC columns with high Surface Area for identification, quantification & purification of small molecules & pharma compounds.

Uptisphere® 120Å

HPLC & prep LC columns for the identification, quantification of small molecules & pharma compounds.

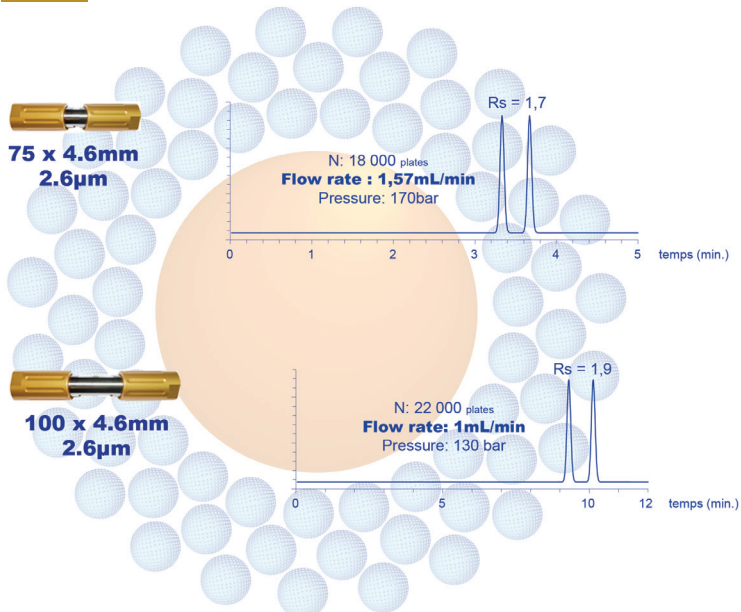
puriFlash®

Analytical, prep-LC & Flash columns with High Load-ability for routine analysis & easy transfer to purification of small organic molecules in pharma applications.

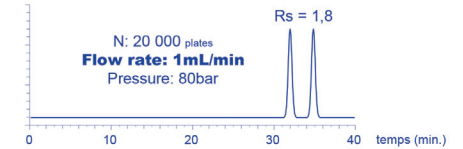
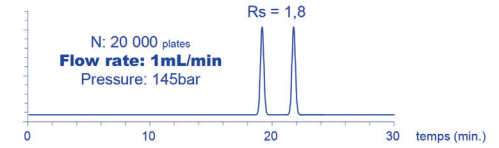
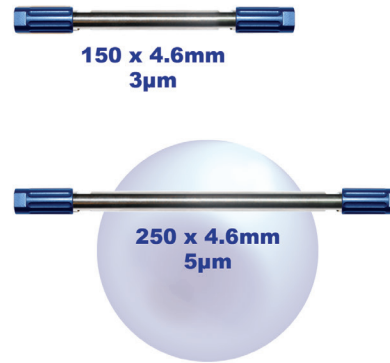


TECHNICAL DATA

Core-shell silica



Totally Porous silica

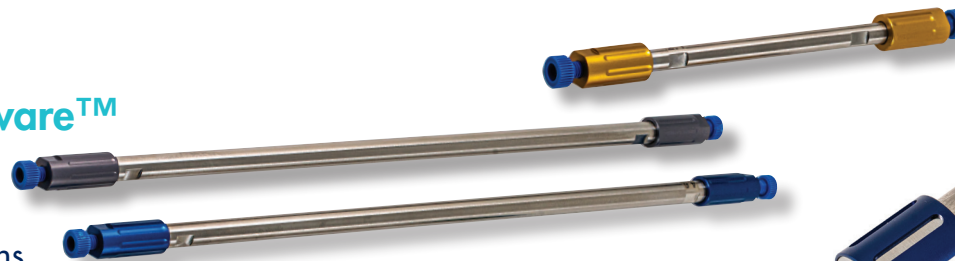


TECHNICAL DATA

High Performance Hardware™

- 1200 bar maximum pressure
- Used for HPLC & UHPLC columns
- High performance

(2.1 - 3.0 - 4.0 - 4.6 - 10.0 - 21.2 - 30.0 - 50.8) mm x (25 - 50 - 100 - 125 - 150 - 250) mm
 Easy transfert from any columns to "ready-to-go" & secured Analytical, or Gradient Purification, Method



Protects all 1.7 up to 5µm silicas columns

- 900 bar maximum pressure
- Low dead volume
- High performance

Core Shell Guard cartridges	Reverse Phase	Hilic Mode	Normal Phase
2.6µm - 5 x 2.1mm - 3u	CS-RP-2.6-005/021	CS-HILIC-2.6-005/021	---
2.6µm - 5 x 4.0mm - 3u	CS-RP-2.6-005/046	CS-HILIC-2.6-005/046	---

Guard cartridges	Reverse Phase	Hilic Mode	Normal Phase
1.7µm - 5 x 2.1mm - 3u	UP-RP-1.7-005/021	---	---
1.7µm - 5 x 4.0mm - 3u	UP-RP-1.7-005/046	---	---
2.2µm - 5 x 2.1mm - 3u	UP-RP-2.2-005/021	UP-HILIC-2.2-005/021	UP-NP-2.2-005/021
2.2µm - 5 x 4.0mm - 3u	UP-RP-2.2-005/046	UP-HILIC-2.2-005/046	UP-NP-2.2-005/046
3µm - 5 x 2.1mm - 3u	UP-RP-3-005/021	UP-HILIC-3-005/021	UP-NP-3-005/021
3µm - 5 x 4.0mm - 3u	UP-RP-3-005/046	UP-HILIC-3-005/046	UP-NP-3-005/046
5µm - 5 x 2.1mm - 3u	UP-RP-5-005/021	UP-HILIC-5-005/021	UP-NP-5-005/021
5µm - 5 x 4.0mm - 3u	UP-RP-5-005/046	UP-HILIC-5-005/046	UP-NP-5-005/046



Guard cartridge




UPTISPHERE® CS EVOLUTION STATIONARY PHASES



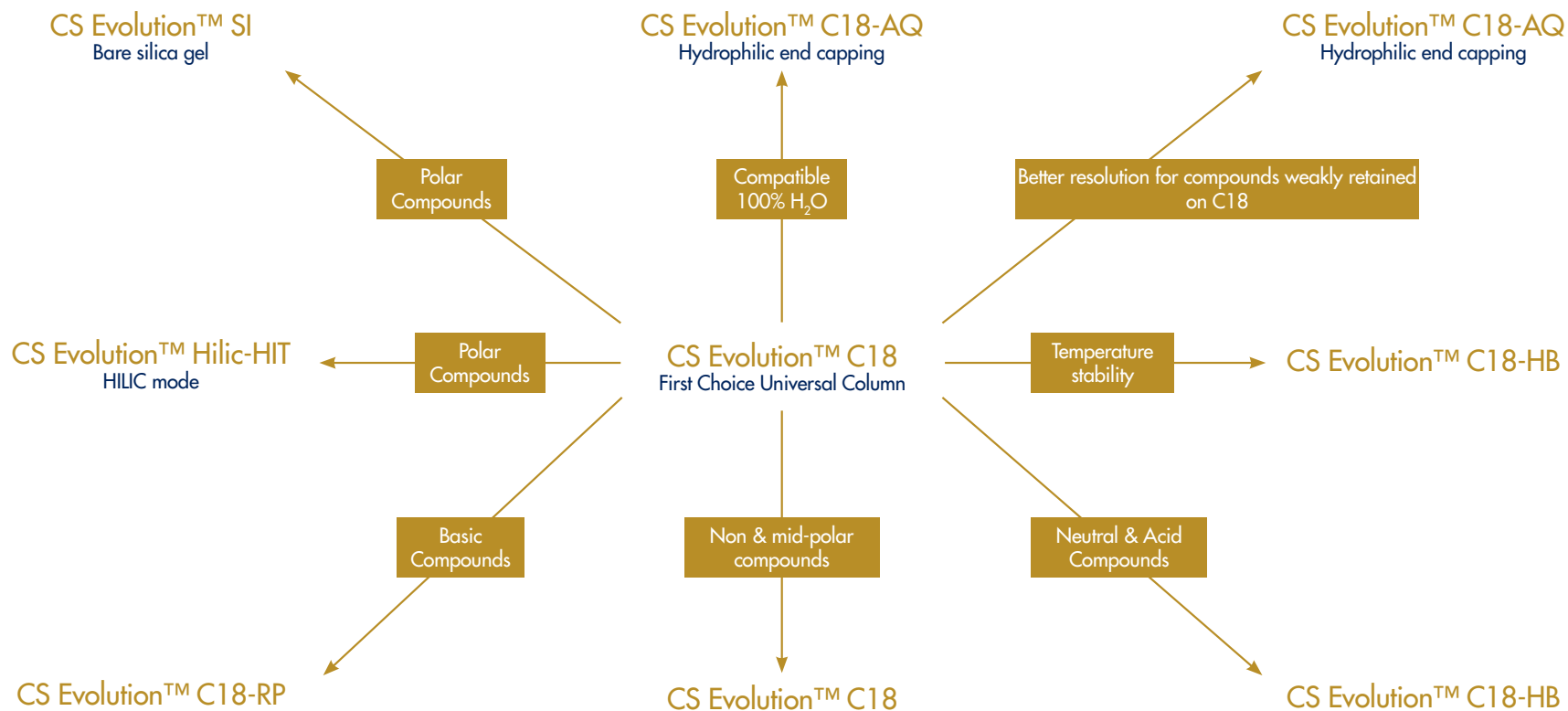
(MOST* Grade)

* MOST : Maximum Operational Surface Technology

	USP Code	Ø Pore	Surface	2.6µm	Bonding	Function-ization	% Carbon	End-Capping	pH stability	Use Mode	Application	
	C18	L1	85Å	130m ² /g	x	C18 - octadecyl	mono-functional	9.0%	One step	1.5 - 7.5	RP	Serves a broad-scope of analytical & prep LC requirements for separating non polar compounds.
	C18-HB	L1	85Å	130m ² /g	x	C18 - octadecyl	mono-functional	8.0%	One step	1.5 - 8.0	RP	Suitable for non polar compounds separation. Exhibits a very hydrophobic surface. HB shows excellent stability under high temperature.
	C18-RP	L1	85Å	130m ² /g	x	C18 - octadecyl	mono-functional	6.0%	Mixte	1.5 - 8.0	RP	Suitable for mid & non polar compounds separation. RP shows excellent mechanical stability that make it an excellent tool for analysis under acidic or basic conditions.
	 C18-AQ	L1	85Å	130m ² /g	x	C18 - octadecyl	mono-functional	6.5%	Mixte	1.5 - 7.0	RP	Suitable for mid & non polar compound separation. Shows excellent stability under 100% aqueous mobile phase condition.
	HIT	L3	85Å	130m ² /g	x	Proprietary	Proprietary	---	Proprietary	1.5 - 7.0	Hilic	Aqueous normal phase separation (ANP) of water-soluble compounds . Typical mobile phase: water / ACN (> 70%) ANP is an excellent alternative to RP purification for highly polar compounds.

SELECTION GUIDE

Uptisphere® (U)HPLC Core Shell Silicas

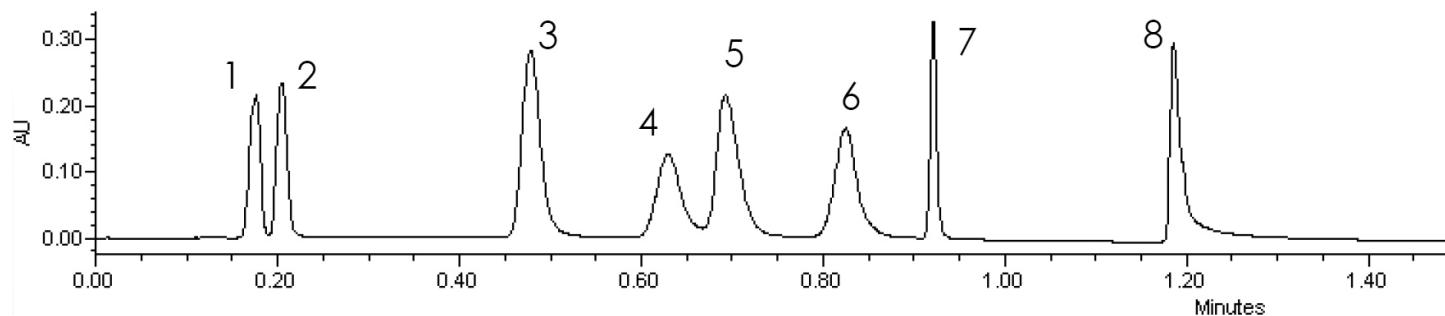


MOST : Maximum Operational Surface Technology



APPLICATION

Nucleobases, Nucleosides & Nucleotides



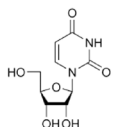
Column Uptisphere® CS Evolution™ 2.6µm HIT 50 x 2,1mm

Gradient: ACN / 20mM formiate d'ammonium pH 3,
Flow rate: 1 mL/min, Temp : 30°C, UV 254nm

	pKa _{acid}	pKa _{basic}	logP
Uracil	9.80-13.80	---	-0.86
Uridine	9.70-12.60	---	-2.42
Adenine	9.90	5.20	-0.53
Adenosine	12.45	4.99	-2.09
Adenosine monophosphate	---	---	-5.19
Cytosine	9.98	2.35	-1.24
Cytidine	12.80-13.60	---	-2.80
Guanosine	10.20	1.8	-2.71



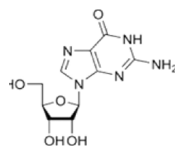
Uracil (1)



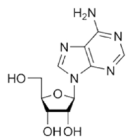
Uridine (2)



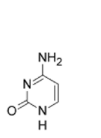
Adenine (3)



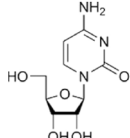
Guanosine (4)



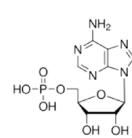
Adenosine (5)



Cytidine (6)



Cytosine (7)



AMP(8)

APPLICATIONS IN ENVIRONMENT

Automatic, simultaneous and rapid analysis of pesticides in surface and underground water by online SPE and UHPLC-MS/MS

Doriane Toinon , Alban Huteau, SHIMADZU France, le Lizard II, bat A, Bd Allende, Noisiel, 77448 Marne La Vallée cedex 2, France.

Uptisphere® CS Evolution 2.6µm C18-AQ 150x3.0mm

A: water + 0.002% formic acid

+ 2mM ammonium formate

B: 50/50 acetonitrile/Methanol

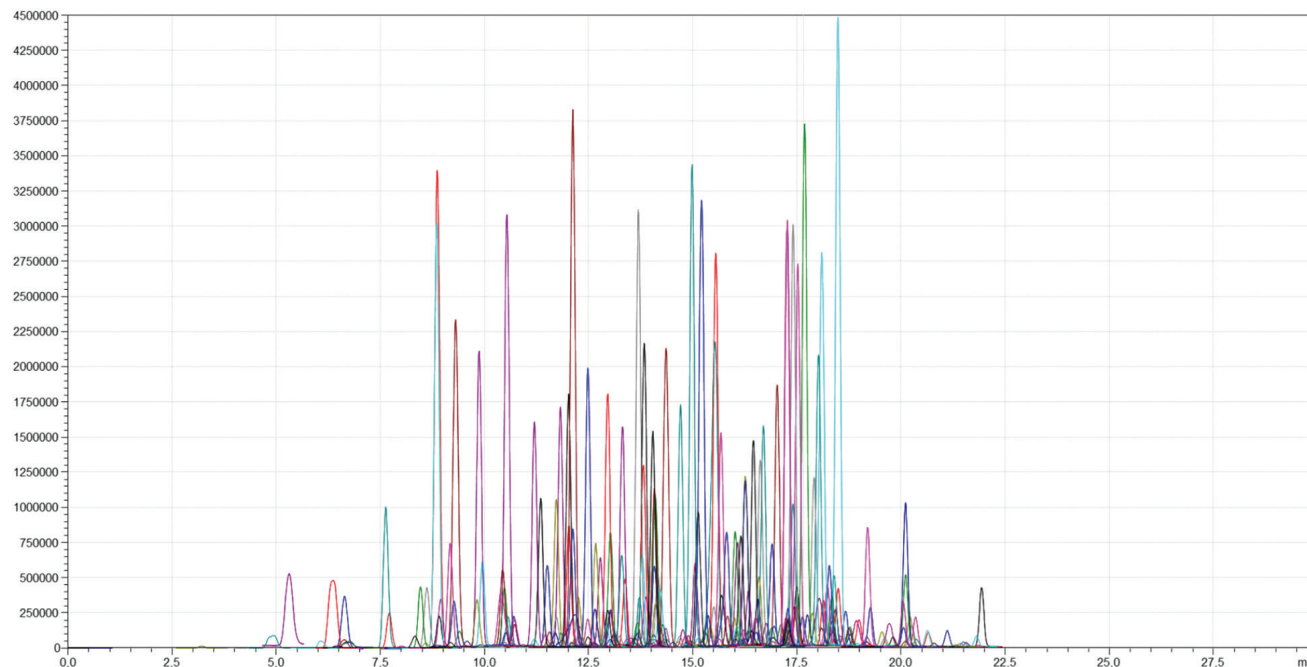
+ 0.002% formic acid

+ 2mM ammonium formate

Flow: 0.7mL/min, 40°C

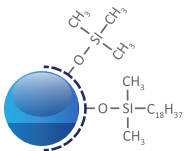
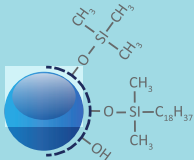
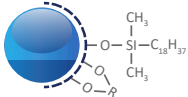
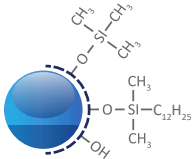
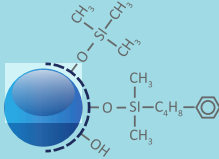
System: Shimadzu Nexera X2

Method validated
with LOQ up to 1ng/L
for 272 pesticides
on water matrix




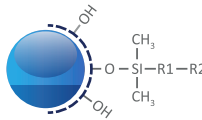

UPTISPHERE® STRATEGY™ STATIONARY PHASES



	USP Code	Ø Pore	Surface	1.7µm	2.2µm	3µm	5µm	Bonding	Functionalization	% Carbon	End-Capping	pH stability	Use Mode	Application	
	C18-3	L1	100Å	425m ² /g	---	---	x	x	C18 - octadecyl	mono-functional	22.0%	Multi step	1.0 - 12	RP	The high bonding density of C18-3 facilitates a strong separation of non polar compounds. Multi step bonding technology guarantees a fully end-capped phase, stable under basic pH conditions. C18-3 is an excellent phase for the separation of basic drugs at up to pH: 12.
	C18-HQ	L1	100Å	425m ² /g	x	x	x	x	C18 - octadecyl	mono-functional	19.0%	Multi step	1.0 - 10	RP	This utility phase serves many pharmaceutical applications. Its 425m²/g surface area providing excellent loading capacity.
	C18-RP	L1	100Å	425m ² /g	---	x	x	x	C18 - octadecyl	mono-functional	16.0%	Multi step Mixte	1.5 - 8.0	RP	Suitable for mid & non polar compounds separation. RP shows excellent mechanical stability that make it an excellent tool for purification under acidic or basic conditions.
	C12	---	100Å	425m ² /g	---	x	---	x	C12 - dodecyl	mono-functional	16.0%	One step	1.5 - 8.0	RP	Non polar compounds. Less retentive than C18 with greater capacity.
	PHC4	L11	100Å	300m ² /g	---	x	x	x	Phenyl - Butyl	mono-functional	12.0%	One step	1.5 - 7.5	RP	Very selective for compounds with aromatic cycles and mid-polar compounds.

UPTISPHERE® STRATEGY™ STATIONARY PHASES



	USP Code	Ø Pore	Surface	2.2µm	3µm	5µm	Bonding	Functiona- lization	End- Capping	pH stability	Use Mode	Application
 HILIC-HIT	L3	100Å	425m ² /g	x	x	x	Proprietary	Proprietary	Proprietary	1.5 - 7.0	Hilic	Aqueous normal phase separation (ANP) of water-soluble compounds. Typical mobile phase: water / ACN (> 70%). ANP is an excellent alternative to RP purification for highly polar compounds.
 HILIC-HIA	L3	100Å	300m ² /g	x	x	x	Proprietary	Proprietary	Proprietary	2.0 - 7.0	Hilic	Aqueous normal phase separation (ANP) of water-soluble compounds. Typical mobile phase: water / ACN (> 70%). ANP is an excellent alternative to RP purification for highly polar compounds.
 SI	L3	100Å	425m ² /g	x	x	x	Ultra pure silica	---	none	1.5 - 7.0	NP	Non-ionic, polar organic compounds.



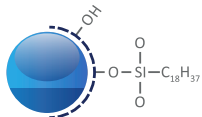
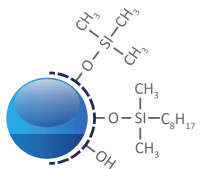
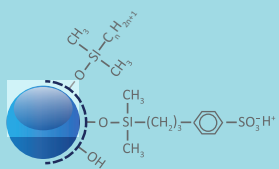
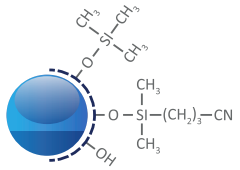
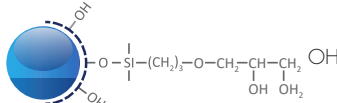
UPTISPHERE® 120Å STATIONARY PHASES



	USP Code	Ø Pore	Surface	2.2µm	3µm	5µm	Bonding	Functionalization	% Carbon	End-Capping	pH stability	Use Mode	Application	
	C18-HSC	L1	60Å	---	---	x	x	C18 - octadecyl	mono-functional	20.0%	Multi step	1.5 - 8.0	RP	Non-polar compounds.
	C18-ODB	L1	120Å	320m ² /g	x	x	x	C18 - octadecyl	mono-functional	18.0%	One step	1.5 - 7.0	RP	Serves a broad-ship of analytical & prep LC requirements for separating non polar compounds.
	C18-HDO	L1	120Å	320m ² /g	x	x	x	C18 - octadecyl	mono-functional	17.0%	Mixte	1.5 - 7.0	RP	Suitable for mid & non polar compound separation. Shows excellent stability under 100% aqueous mobile phase condition.
	C18-NEC	L1	120Å	320m ² /g	x	x	x	C18 - octadecyl	mono-functional	16.0%	none	1.5 - 6.5	RP	NEC strongly retains the polar and mid-polar compounds. It overcome peak tailing with compounds that contains chains and / or carbon cycles combined with numerous polar groups and/or basic in character.

UPTISPHERE® 120Å STATIONARY PHASES

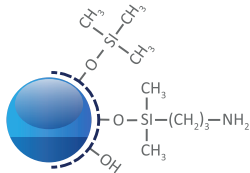
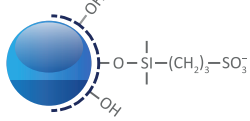
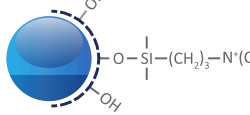


	USP Code	Ø Pore	Surface	3µm	5µm	Bonding	Functiona- lization	% Carbon	End- Capping	pH stability	Use Mode	Application	
	C18-TF	L1	---	---	x	C18 - octadecyl	poly-functional	14.0%	One step	1.5 - 8.0	RP	Alternative selectivity for challenging separations & for - but not limited to - aromatic, polyphenol, PAHs etc.	
	C8	L7	120Å	320m ² /g	(x)	x	C8 - octyl	mono-functional	11.0%	One step	2.0 - 7.0	RP	Mid-polar and non polar compounds. Less retentive than C18.
	MM1	L44	120Å	320m ² /g	---	x	C8 / SCX	mono-functional	---	One step	2.0 - 6.5	RP / EI	Ion exchange and hydrophobic chains are bonded onto the surface of silica providing unique selectivity. Compounds that possess basic functionality are retained by ion exchange functionality. An organic solvent will elute hydrophobic compounds.
	CN	L10	120Å	320m ² /g	x	x	CN - cyano	mono-functional	8.0%	One step	2.0 - 7.0	RP / NP	CN functional groups can be used either in normal phase to purify polar compounds or in reversed phase for mid-polar compounds.
	L20	120Å	320m ² /g	---	(x)	OH - diol	mono-functional	6.0%	none	1.5 - 6.5	RP / NP	The diol function provide globally a neutral surface onto the silica. It leads to greater separation of basic compounds by normal phase vs. regular silica.	



UPTISPHERE® 120Å STATIONARY PHASES



	USP Code	Ø Pore	Surface	2.2µm	3µm	5µm	Bonding	Functiona- lization	% Carbon	End-Capping	pH stability	Use Mode	Application
	NH2	L8	120Å	320m ² /g	x	x	x	NH2 - amino	mono-functional	5.0%	none	2.0 - 6.5	RP / NP / EI Can be either weak anion exchangers for strong acids, or polar media that can interact with OH, NH, SH...
	SCX	L50	120Å	320m ² /g	---		x	Strong Cation Exchanger	mono-functional	---	none	1.0 - 7.5	EI Strong cation exchange (SCX) contains sulfonic acid used to analyze weak basic compounds which have one or more positive charges.
	SAX	L14	120Å	320m ² /g	---		x	Strong Anion Exchanger	mono-functional	---	none	1.0 - 7.5	EI Strong anion exchange (SAX) contains quaternary amine used to analyze weak acid compounds which have one or more negative charged, nucleotides, nucleosides, organic acids...

PURIFLASH® STATIONARY PHASES

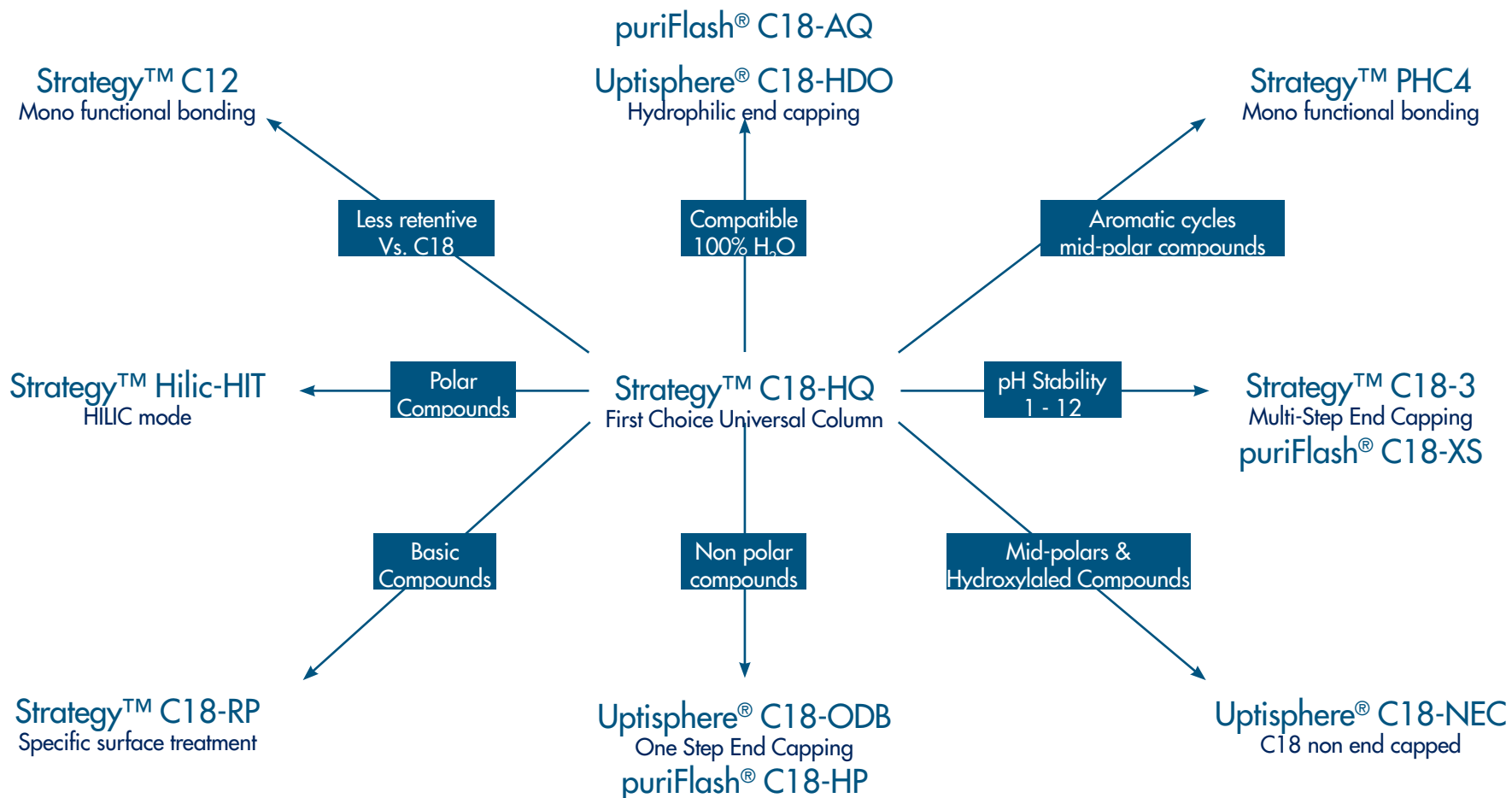


	USP Code	Ø Pore	Surface	5µm	Bonding	Functionalization	% Carbon	End-Capping	pH stability	Use Mode	Application	
	C18-XS	L1	100Å	300m ² /g	x	C18 - octadecyl	mono-functional	17.0%	Multi-step	1.0 - 10.0	RP	The proprietary multi-step bonding technology guarantees a fully end-capped phase, stable under basic pH conditions up to pH: 10. It's an excellent phase for the integral purification of basic drugs.
	C18-HP	L1	100Å	300m ² /g	x	C18 - octadecyl	mono-functional	16.5%	One-step	1.5 - 7.5	RP	Serves many pharmaceutical applications. Excellent choice for routine purification in reverse phase mode.
	C18-AQ	L1	100Å	300m ² /g	x	C18 - octadecyl	mono-functional	14.0%	Mixte	2.0 - 7.5	RP	The bonding chemistry allow to start gradient with 100% of water. Suitable for the purification of mid and non polar compounds.
	RP-AQ	L7	60Å	500m ² /g	x	RP-alkyl	mono-functional	6.0%	Mixte	2.0 - 7.5	RP	The bonding chemistry allow to start gradient with 100% of water. Suitable for the purification of high and mid polar compounds. Compare to C18, peaks are elutes earlier from the beginning of the gradient.



SELECTION GUIDE

(U)HPLC Fully Porous Silicas



APPLICATION

Characterization of Furosemid within rat plasma & broncho-alveolar washing

Uptisphere® Strategy™ 1.7µm C18-HQ,
50 x 2.1mm

Conditions:

Agilent 1200 RRLC + Qtrap 4000
Solvent A: 0.1% CH₃COOH
Solvent B: ACN + 0.1% CH₃COOH
Isocratic (50/50)
Flow rate: 600µL/min @ 500bar (7000psi)
LD: 2.14µg/L
Injection: 10µL
MS: 329 to 285
System: Shimadzu Nexera X2

Comments:

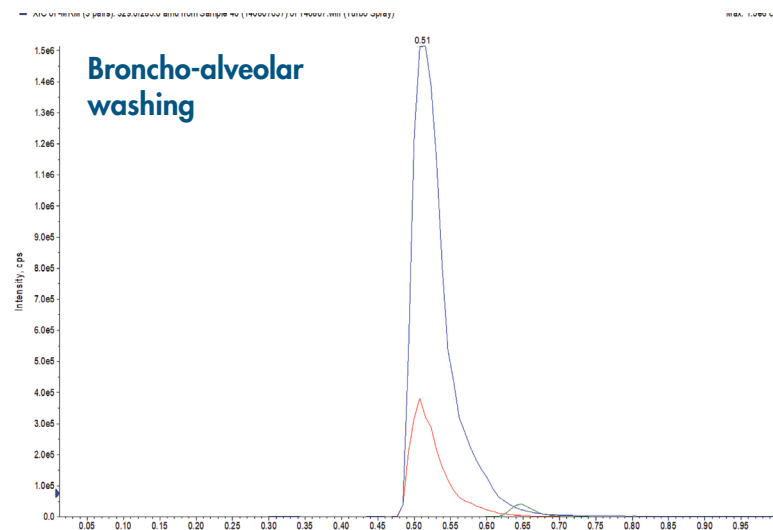
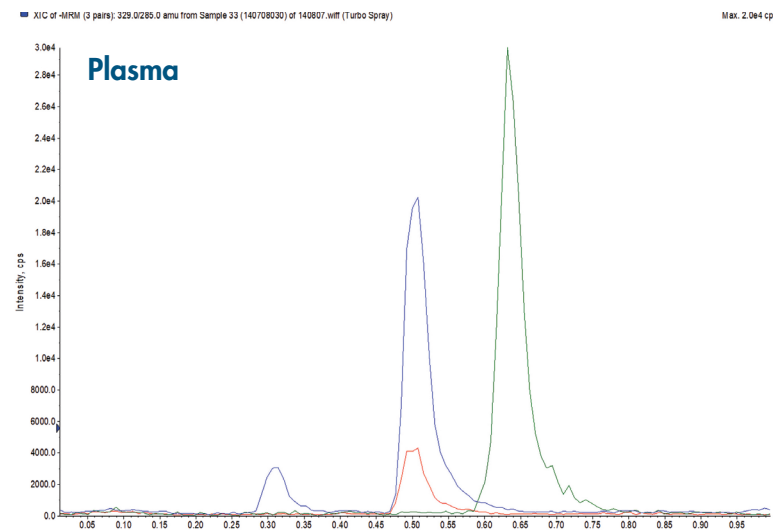
This study will be published.

We can see a big peak within the plasma and nothing except the Internal Standard (IS) within the broncho-alveolar washing.

IS run time: 0.5min

Furosemid run time: 0.64min - a "supposed" metabolite is elute within the plasma extract at 0.3min.

Total run time: 1 min.



APPLICATION

Analysis of Furocoumarins in Citrus Essential Oils

Uptisphere® 3µm C18-ODB, 150 x 2.1mm

Conditions:

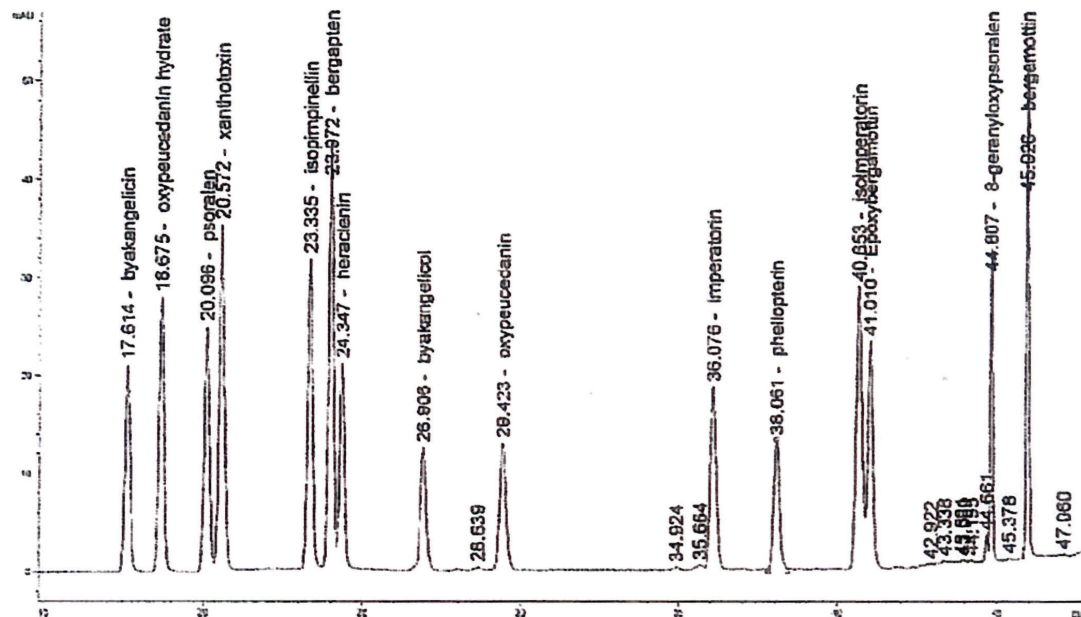
Solvent A: water - acetonitrile - THF (85/10/5)

Solvent B: acetonitrile - methanol - THF (65/30/5)

Flow rate: 0.3mL/min

DAD: 310nm

Time min	0 to 5	5 to 20	20 to 24	24 to 38	38 to 40	Rinse	Equilibration
% B	0	0-32	32	32-55	55-90	90	0
	isocratic	linear	isocratic	linear	linear	10min	10min



APPLICATION

Urea from diesel particles extracts

**Uptisphere® Strategy™ 5µm Hilic-HIT,
250 x 4.6mm**

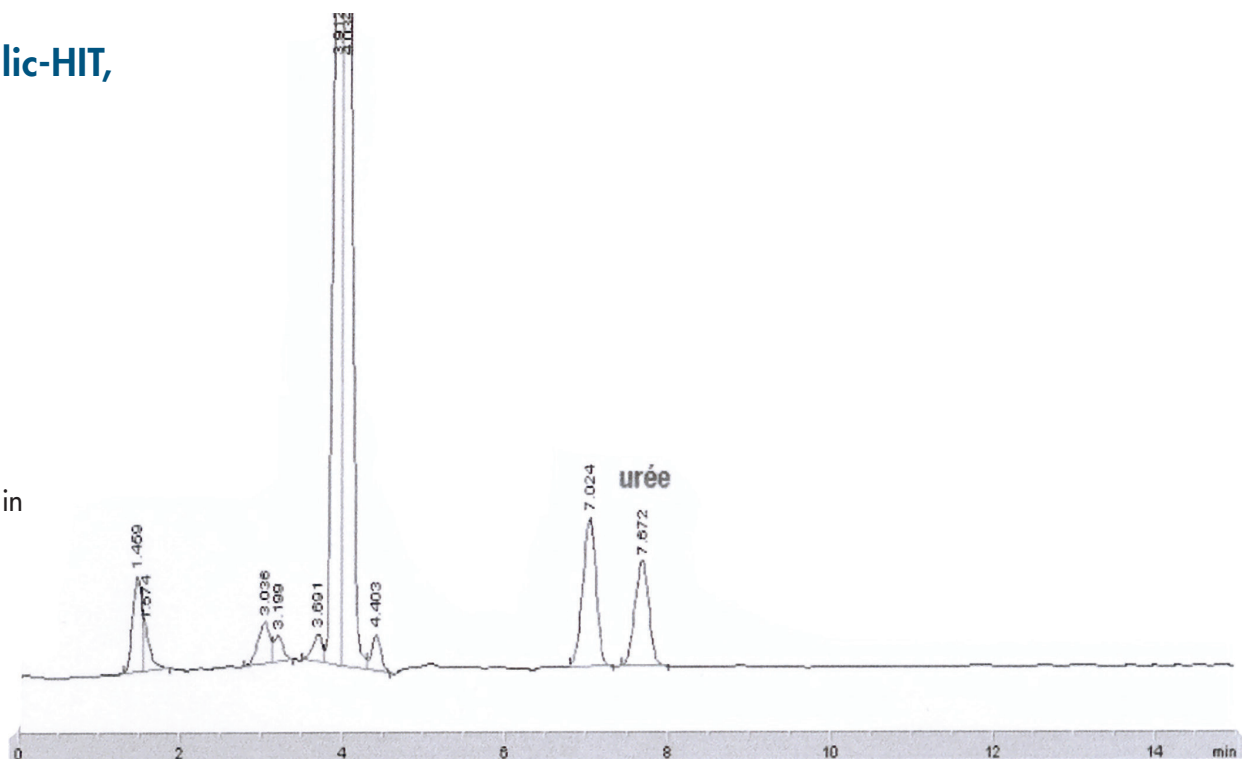
Conditions:

Acetonitrile 100%
Flow rate: 1 mL/min
Temperature: 20°C
DAD: 195nm
Injection: 10µL

Comments:

The column provides the following advantage in comparison to the previous one

- Better Urea retention
- Deletion of interferences
- Symmetric Urea peak



Uptisphere® Strategy™ 5µm Hilic-HIT, 250 x 4.6mm





Advion Interchim scientific

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