





Fortis Philosophy

Here at Fortis Technologies we take pride in understanding the needs of our customers, levels of technical service are of the highest standard and all of our products are developed in close consultation with customers to ensure that their needs are met.

We believe in making method development as simple as possible, so our innovative phase chemistries offer the ability for resolution, efficiency, speed and sensitivity; all of the variables that are needed in today's high speed evolving world.

If you have any questions please feel free to contact us at : info@fortis-technologies.com

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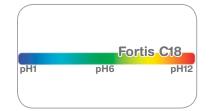
UHPLC Columns

- Optimised for Ultra High Pressure LC (UHPLC)
- Available C18, Diphenyl, HILIC, Cyano
- Operate at 18,000psi
- Fully Scaleable analytical to prep



1.7µm Fortis[™] C18

- Superior Peak Shape
- Low, Mid and High pH 1-12
- Based upon Ultra Pure Silica
- Fully Scaleable



1.7µm Fortis Diphenyl

- Unique diphenyl selectivity
- Separate Positional Isomers
- Metabolite separations
- Enhanced Polar Retention



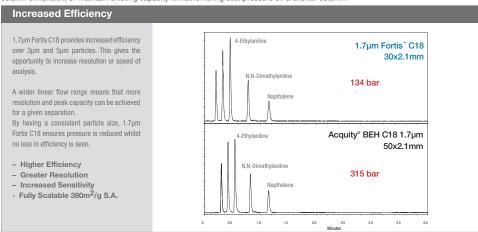
1.7µm UHPLC Columns

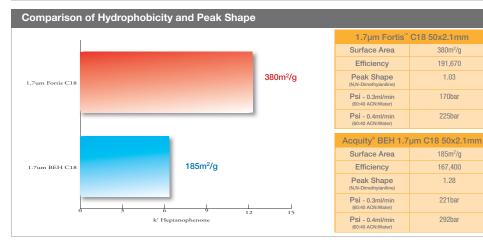
- 380m2/g Surface Area Provides Increased Peak Capacity
- High Efficiency small particles
- Robust Reproducible UHPLC columns
- Operate to 18,000psi

UHPLC

- Fully Scalable to Analytical and Prep Size

1.7um Fortis particles are designed to provide characteristics, which will aid in increased productivity within ultra high pressure chromatography (UHPLC). Designed to be robust, reproducible and fully scalable with 3µm, 5µm and 10µm particles. 1.7µm Fortis particles will operate upto 1200bar providing high linear velocities, increased efficiency, and allowing speed and sensitivity to be achieved on all the latest UHPLC systems. By choosing a high surface area (S.A.) UHPLC phase the analyst can increase peak capacity using their existing column dimension, or maintain existing capacity whilst lowering backpressure on a shorter column.







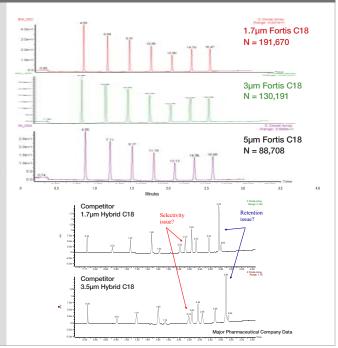
Fully Scaleable

Critically important to the analyst is the ability to have a fully scalable separation. Fortis C18 can be scaled from 1.7µm all the way through analytical 3 and 5µm particles to prep size without any change in retention profile.

By combining the same surface area, pore size characteristics with the identical bonding the analyst can be ensured of having the ability to either scale up methods to 'traditional' LC systems, or to be confident that a method can be transferred to another laboratory with the same selectivity being achieved.

If a small particle used in UHPLC is not the same as its larger 3um and 5um particle then changes in resolution and retention can occur, both of which can cause problems in method validity.

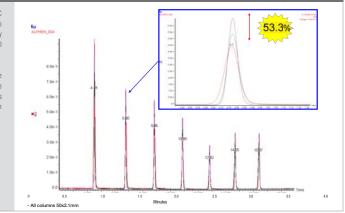
1.7µm Fortis C18 will alleviate all these potential issues, leaving the analyst confident in method transfer.



Sensitivity Gains

Critically peak height increases in UHPLC mode due to the rise in efficiency (N) from the smaller particle, but also it is also inversely proportional to peak width, so symmetrical peaks will lead to increased sensitivity.

By moving from a 3µm Fortis C18 particle size to 1.7µm Fortis C18 sensitivity can be increased. In this example peak height goes up by over 27%. The increase from 5um particles is even greater.



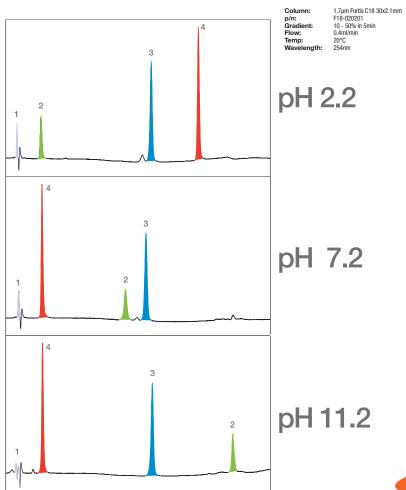
1. Uracil

4. 3-Nitrobenzoic acid

1.7µm Fortis C18 pH options

- pH selectivity for method development
- pH stable 1-12
- Gives high speed of equilibration

1.7µm Fortis C18 will operate across the pH spectrum giving the analyst the ability to optimise the correct pH region for their separation. Quickly equilibrating from formic acid to ammonium acetate through to ammonia allows pH, as a method variable, to be rapidly evaluated. Resolution of compounds can be changed radically by altering pH to optimise separation between compound classes.

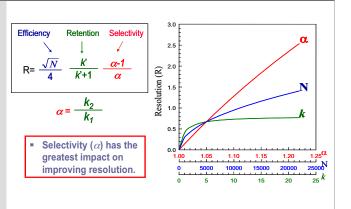


Fortis C18 page 14

Phase Chemistry Selectivity

Resolution vs Efficiency vs Selectivity

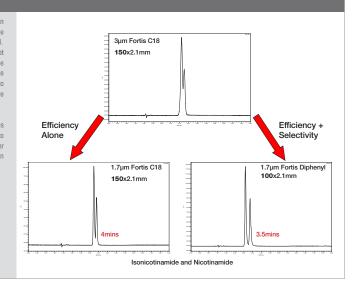
- 1.7µm Fortis C18 will provide Hydrophobic selectivity which is suitable for many compounds. However as we can see from the resolution equation having multiple phase chemistries available is a definite advantage. Selectivity can then be used in conjunction with efficiency of the small particle.
- 1.7µm Fortis UHPLC columns are also available
- 1.7µm Fortis Diphenyl
- 1.7µm Fortis Cyano
- 1.7µm Fortis HILIC



Improve Selectivity

If we are scaling a method and hoping that an increase in efficiency alone will provide the necessary resolution we can be disappointed. Scaling from 3µm to 1.7µm C18 has not provided baseline resolution between the compounds. Adding selectivity by phase chemistry into the equation has allowed us to go faster on a shorter column and now achieve

In this instance 1.7µm Fortis Diphenyl provides more resolution than C18. This then leads to the ability to increase speed by use of shorter columns, higher temperature, change in organic or a combination of the above.



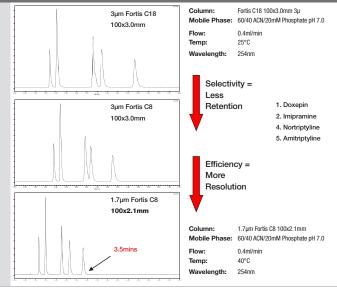
Method Development

Transfer from HPLC to UHPLC

Optimising resolution is one of the main reasons that we wish to move from HPLC to UHPLC, along with increased speed.

If we can change the hydrophobicity of the phase chemistry then we have the potential to reduce the retention time.

If we can then move to a smaller particle and increase resolution between our critical pairs we can improve the speed of analysis still



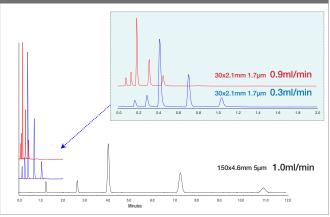
Transfer from HPLC to UHPLC

Greater speed can be achieved by moving to smaller particles, whilst still obtaining full baseline resolution.

1.7µm Fortis C18 will operate at higher linear flow rates whilst providing lower backpressure than other sub 2µm particles.

To move to smaller particles and smaller columns, without affecting selectivity, use some simple scaling equations to optimise your UHPLC parameters: e-mail us for the

technicalsupport@fortis-technologies.com



UHPLC Sample Filter



- Filter for all UHPLC columns
- No backpressure increase
- Increase lifetime of UHPLC columns
- Low volume in-line filter
- Change over time is seconds not minutes

Fortis UHPLC in-line filters are direct connect design, fitting in between the UHPLC column and the conventional fitting to filter out particulate matter. They contain low dead volume and pressure. In-line filters are ideal for 1.7µm Fortis columns where extra packed bed from a guard would be detrimental. UHPLC in-line filters are manufactured to withstand 20,000psi.

1.7µm UHPLC part numbers

| 1.7 | um Fortis C18 |
|------------|----------------------------|
| F18-020701 | 1.7µm Fortis C18 150x2.1mm |
| F18-020501 | 1.7µm Fortis C18 100x2.1mm |
| F18-020301 | 1.7µm Fortis C18 50x2.1mm |
| F18-020201 | 1.7µm Fortis C18 30x2.1mm |
| F18-020101 | 1.7µm Fortis C18 20x2.1mm |
| F18-030501 | 1.7µm Fortis C18 100x3.0mm |
| F18-030301 | 1.7µm Fortis C18 50x3.0mm |
| F18-030201 | 1.7µm Fortis C18 30x3.0mm |
| F18-050301 | 1.7µm Fortis C18 50x4.6mm |

| 1.7um Fortis Cyano | | | | |
|--------------------|------------------------------|--|--|--|
| FCN-020701 | 1.7µm Fortis Cyano 150x2.1mm | | | |
| FCN-020501 | 1.7µm Fortis Cyano 100x2.1mm | | | |
| FCN-020301 | 1.7µm Fortis Cyano 50x2.1mm | | | |
| FCN-020201 | 1.7µm Fortis Cyano 30x2.1mm | | | |
| FCN-020101 | 1.7µm Fortis Cyano 20x2.1mm | | | |
| FCN-030501 | 1.7µm Fortis Cyano 100x3.0mm | | | |
| FCN-030301 | 1.7µm Fortis Cyano 50x3.0mm | | | |
| FCN-030201 | 1.7µm Fortis Cyano 30x3.0mm | | | |
| FCN-050301 | 1.7µm Fortis Cyano 50x4.6mm | | | |

| 1.7um Fortis Diphenyl | | | | | |
|-----------------------|---------------------------------|--|--|--|--|
| FPH-020701 | 1.7µm Fortis Diphenyl 150x2.1mm | | | | |
| FPH-020501 | 1.7µm Fortis Diphenyl 100x2.1mm | | | | |
| FPH-020301 | 1.7µm Fortis Diphenyl 50x2.1mm | | | | |
| FPH-020201 | 1.7µm Fortis Diphenyl 30x2.1mm | | | | |
| FPH-020101 | 1.7µm Fortis Diphenyl 20x2.1mm | | | | |
| FPH-030501 | 1.7µm Fortis Diphenyl 100x3.0mm | | | | |
| FPH-030301 | 1.7µm Fortis Diphenyl 50x3.0mm | | | | |
| FPH-030201 | 1.7µm Fortis Diphenyl 30x3.0mm | | | | |
| FPH-050301 | 1.7µm Fortis Diphenyl 50x4.6mm | | | | |
| | | | | | |

| 1.7um Fortis HILIC | | | | | |
|--------------------|------------------------------|--|--|--|--|
| FHI-020701 | 1.7µm Fortis HILIC 150x2.1mm | | | | |
| FHI-020501 | 1.7µm Fortis HILIC 100x2.1mm | | | | |
| FHI-020301 | 1.7µm Fortis HILIC 50x2.1mm | | | | |
| FHI-020201 | 1.7µm Fortis HILIC 30x2.1mm | | | | |
| FHI-020101 | 1.7µm Fortis HILIC 20x2.1mm | | | | |
| FHI-030501 | 1.7µm Fortis HILIC 100x3.0mm | | | | |
| FHI-030301 | 1.7µm Fortis HILIC 50x3.0mm | | | | |
| FHI-030201 | 1.7µm Fortis HILIC 30x3.0mm | | | | |
| FHI-050301 | 1.7µm Fortis HILIC 50x4.6mm | | | | |

High Performance 2.5µm particles

- Over 20% more efficiency than 3µm
- Lower backpressure than UHPLC columns
- Operate in 400bar or 1000bar systems
- Fully Scalable to analytical and prep size

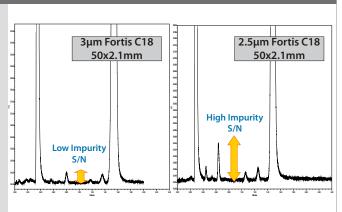
Fortis 2.5µm particles are designed to be the next step in improving both resolution and speed. Allowing the analyst to move towards ultra high pressure chromatography (UHPLC) whilst still operating on traditional 400bar LC systems. Can be combined with the companies innovative column designs 2.5µm particles offer speed and efficiency without compromising loadability and scaleability. The optimised C18 bonding maintains the phases ability to be stable from pH 1-12.

Optimised Resolution Resolution of closely related species can be Fortis C18 100x4.6mm 2.5µ Column: Tricyclic antidepressants achieved by the use of 2.5µm particles and F18-050502 the optimised peak shapes afforded by Fortis Mobile Phase: A - H₂0 + 0.1% Formic acid C18 stationary phase. B - ACN + 0.1% Formic acid Basic, Acidic and Neutral analyte performance Gradient: 25 - 40% in 10min is first class across the pH spectrum. Flow: - Higher Efficiencies - Greater Reproducibility - Resolution enhanced 1. Protriptyline 2. Nortriptyline 3. Amitriptyline 4. Trimipramine

Sensitivity Gain - Impurities

With sensitivity of degradents and impurities being an issue in pharmaceutical analysis, peak shapes and peak height need to be optimal in order to obtain low level LOD (limits

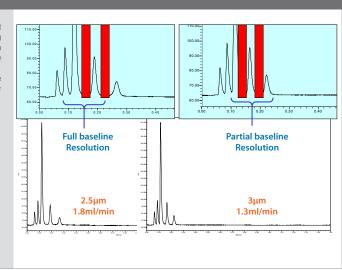
Fortis 2.5um particles allow higher sensitivity to be obtained than 3um particles, therefore lower LOD's.



Increased Peak Capacity

As we run faster and faster we get to the point where peak capacity starts to fall, by being able to increase efficiency by moving to a smaller particle we can increase the efficiency and therefore maintain peak capacity.

An improvement in baseline resolution can be seen by the use of 2.5um particles at higher linear velocities.



2.5µm Fortis C18 particles can be provided packed in hardware that is fully compatible with UHPLC systems so the low dead volume capability of these systems can be fully utilised:



To learn more about the stability and peak shapes of Fortis C18 turn to page 14.

| 2.5µm Fortis C18 | Column Length | | | | |
|---------------------|---------------|------------|------------|------------|--|
| | 30 | 50 | 100 | 150 | |
| 2.1 | F18-020202 | F18-020302 | F18-020502 | F18-020702 | |
| Column Diameter 3.0 | F18-030202 | F18-030302 | F18-030502 | - | |
| 4.6 | F18-050202 | F18-050302 | F18-050502 | - | |

| 2.5µm Fortis C18 in UHPLC | | Column | Length | |
|---------------------------|---------------|---------------|---------------|---------------|
| Hardware | 30 | 50 | 100 | 150 |
| 2.1 | F18-020202UHP | F18-020302UHP | F18-020502UHP | F18-020702UHP |
| Column Diameter 3.0 | F18-030202UHP | F18-030302UHP | F18-030502UHP | - |
| 4.6 | F18-050202UHP | F18-050302UHP | - | - |



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Fortis Method Development Options

- Choice of Stationary phase functionality
- Based on Ultra pure silica
- Reversed Phase (RP) and Normal Phase (NP) options

Fortis[™] H2o

- Polar endcapped C/18
- Increased polar retention
- Organic acids
- Catecholamines

Fortis[™] Diphenyl - Unique di-phenyl structure

- Separate Positional Isomers
- Metabolite profiling

Fortis[™] C18 - General HPLC use - Method dev. from pH 1-12 - Acids, Bases and Neutral

Fortis[™] Cyano

- Cyano functionality
- RP or NP use
- Explosives
- Pesticides

Fortis[™] HILIC

- High Polar Retention
- Highly Pure Silica
- Carboxylic acids
- Nucleotides

Fortis[™] C8

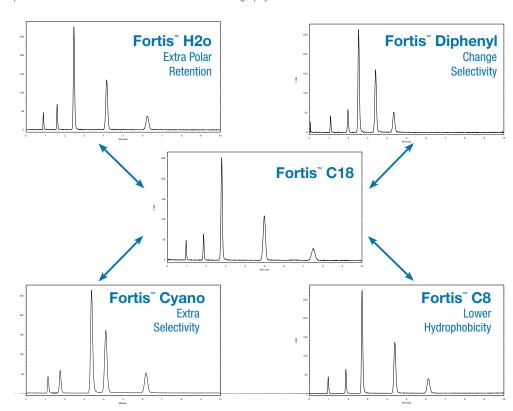
- Reduced Hydrophobicity
- Lipids
- Steroids

Getting Started:

Method development typically starts with a C18 or C8 column, both provide Hydrophobic retention with good peak shapes for neutral, acidic and basic analytes. Generally if retention of polar molecules is also needed then a polar endcapped stationary phase such as Fortis H2o is a good starting choice.

If selectivity is insufficient then Diphenyl or Cyano stationary phases are a good alternative, they will change selectivity and even elution order since they work on dipole characteristics as opposed to just hydrophobicity.

Fortis Cyano is good in normal phase (NP) conditions for polar analytes with COOH, NH2, NHR2 or NR2 groups. If small polar molecules still do not retain then HILIC chromatography is a suitable alternative.



Acidic, Neutral & basic analytes

- Fortis C18
- Fortis C8
- Fortis Diphenyl

Polar basic molecules

- Fortis C18 operated at high pH
- Fortis Diphenyl
- Fortis H2o

Polar acidic molecules

- Fortis H2o
- Fortis HILIC
- Fortis Cyano in NP mode

Alternate Selectivity

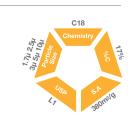
- Fortis Diphenyl
- Fortis Cyano

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Fortis[™] C18

- Superior Peak Shapes
- pH Range 1-12
- Based on Ultra Pure Silica
- Fully Scalable UHPLC to Prep

Fortis C18 is a pure silica based stationary phase with unique high and low pH performance. Whether carrying out simple compound screens or complex metabolite identification Fortis C18 will provide the best in peak shape, resolution and extended pH range for method development flexibility.



Optimised Peak Shape

Whatever the compound functionality the optimised hydrophobic bonding of Fortis C18 leads to peak symmetries being near perfect whatever the analyte type.

Basic, Acidic and Neutral analyte performance is first class across the pH spectrum.

- Superior Peak Shapes
- Higher Efficiencies
- Excellent Reproducibility

Column:

Fortis™ C18 150x4.6mm 5µ Luna® C18(2) 150x4.6mm 5µ

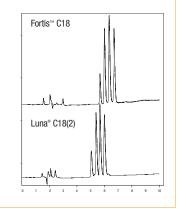
Mobile Phase: A - H₂0 + 0.1% Formic acid B - ACN + 0.1% Formic acid

Gradient:

25 - 40% in 10min

Flow:

- 1. Protriptyline
- 2. Nortriptyline
- 3. Amitriptyline
- 4. Trimipramine

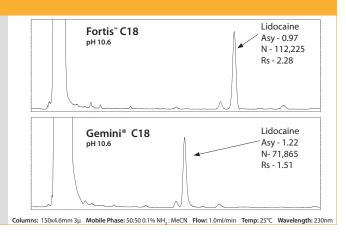


Extreme pH range

Fortis C18 has the ability to not only operate at low pH like other silica based phases, but also to operate at high pH like hybrid phases to aid with basic analyte retention and performance.

The ability to quickly equilibrate from formic acid or TFA into ammonia or bicarbonate aids in method development. Mass transfer, loadability and precision of a silica matrix are all maintained.

- Higher Efficiency than Hybrids
- Excellent Reproducibility
- Retain Polar Basic Analytes

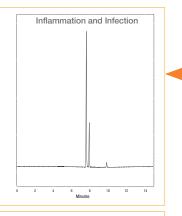


Extended operating pH range

Mobile Phase: A - H₂0 + 0.1% Formic acid B - ACN + 0.1% Formic acid 10 - 50% in 10min

Flow: 20°C

- 2. Neomycin Sulphate
- 3. Acetic acid

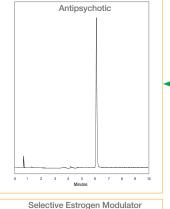


Fortis C18 50x4.6mm 5µ F18-050305

A - 50mM NH₄OAc Mobile Phase: B - ACN

10 - 40% in 10min Flow: 20°C





Column: Fortis C18 50x3.0mm 3µ

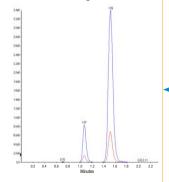
Mobile Phase: 30:70 H₂0 + 10mM ammonium bicarbonate : MeOH

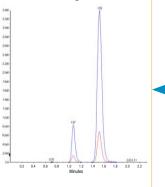
0.4ml/min

MS Detection

Raloxifene Glucuronides

Data Courtesy of : Pharmaceutical company, USA



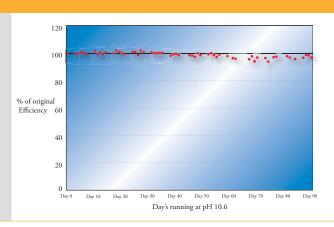


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The unique bonding of Fortis C18 enables stability at extremes of pH to be maintained.

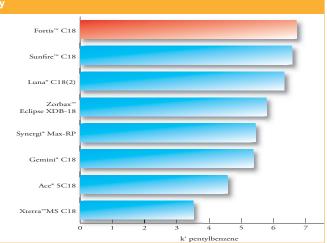
Run continuously in 0.1% ammonia Fortis C18 shows no deterioration in efficiency over a 90 day period.



Fortis C18 high surface area combined with the optimised C18 ligand bonding provides high retention for compounds.

This is advantageous in a number of ways:

- Higher retention of analytes, more organic modifier can be used to elute, therefore greater MS sensitivity.
- Higher retention of analytes, more organic leads to shorter 'dry-down' in fraction collection.
- Higher retention of analytes, more chance of resolution



Fortis[™] C18

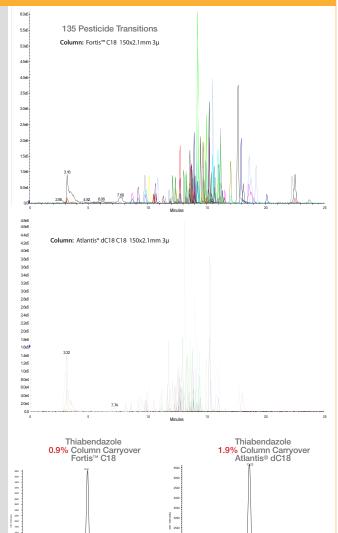
Optimised Resolution

Only by optimising all factors of stationary phase design can the analyst be assured of the best possible chromatography.

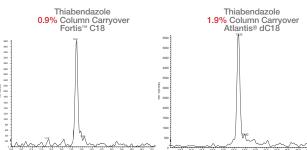
Fortis C18's unique bonded character ensures that not only is reproducibility and robustness assured, but also that resolution is of the highest level. Only by obtaining sharp peak shapes for many analyte types both polar and non polar can this sort of resolution be achieved.

Analysed here are 135 transitions of pesticide residue from an apple matrix. Good LC resolution leads to excellent sensitivity in MS detection.

Polar organophosphates such as Acephate and Methamidophos are retained well due to the high surface area of the Fortis C18 phase.



Thiabendazole can be bound on the column from one gradient cycle to the next, the optimised hydrophobicity of Fortis C18 means that carryover on column is greatly reduced since there is no secondary silanol activity to bind with analytes.



Data Courtesy of : Central Science Laboratories, UK

Fortis[™] C18

Analyte Loading

Based on a silica template Fortis C18 has high loading capability for those wishing either scale up to preparative separation or needing to load in order to correctly identity low level components.

Having a 380m2/g surface area means that the phase chemistry will not overload causing poor peak shapes. This can be especially important in biological work where a high concentration of matrix interference is also often present

Smaller surface area phases and solid-core-shell particles can suffer from lower loading capability and potentially higher backpressure.

Overload can be viewed as loss of efficiency and/or peak shape.

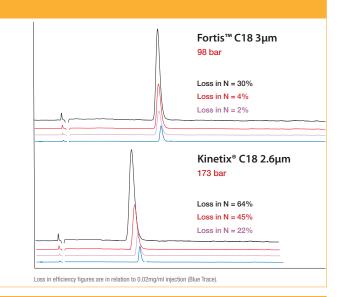
Column: 50x3.0mm

Mobile Phase: H₂0 + 0.01% formic acid : ACN

Flow: 0.6ml/min

Temp: 30°C

Diphenhydramine 0.02, 0.2, 0.5 & 1mg/ml



Selectivity of C18 - Plant Hormones

All C18 chemistries are capable of providing different selectivity. Selectivity can be just as important as efficiency, here we see radically different peak shapes and resolution regardless of C18 particle size for some plant hormones.

Column: 50x2.1mm

Mobile Phase: A - $H_2O + 0.01\%$ formic acid

B - MeOH + 0.1% formic acid 10-40% in 30min

Gradient:

Temp: 30°C
Wavelength: MS Detection

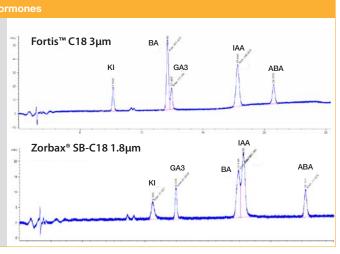
KI = Kinetin

BA = Benzyladenine

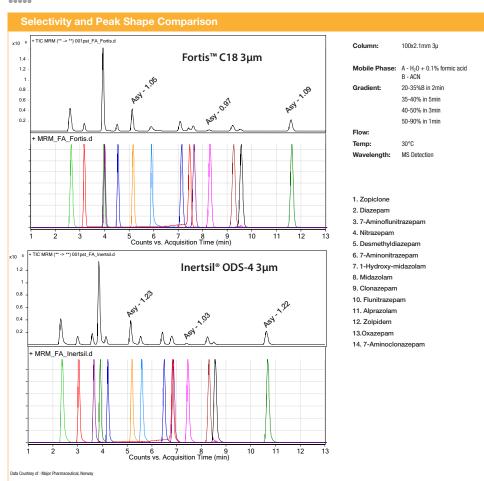
IAA = Indol-3-yl acetate

ABA = Abscisic acid GA3 = Gibberellin acid

Data Courtesy of : Kings College, UK



Fortis[™] C18



| Fortis C18 | | | Column | Length | |
|-----------------|-----|------------|------------|------------|------------|
| | | 50 | 100 | 150 | 250 |
| | 2.1 | F18-0203xx | F18-0205xx | F18-0207xx | - |
| Column Diameter | 3.0 | F18-0303xx | F18-0305xx | F18-0307xx | - |
| | 4.6 | F18-0503xx | F18-0505xx | F18-0507xx | F18-0509xx |

| Fortis C18 Guards | | Length |
|-------------------|-----|--------------|
| | | 10 |
| Column Diameter | 2.1 | DC18-0200xxG |
| | 4.6 | DC18-0500xxG |

Replace xx -01 for 1.7μm - 02 for 2.5μm - 03 for 3μm - 05 for 5μm - 10 for 10μm

Fortis[™] Diphenyl

- Unique Selectivity
- Separate Positional Isomers
- No "MS bleed", Stable Hydrophobic Ligand
- Enhanced Polar Retention

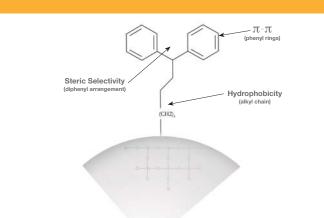
Fortis Diphenyl is designed to provide characteristics which will enhance selectivity. It provides the analyst with extra retention of compounds containing aromatic functionality. Extra selectivity and retention can be found for polar substrates, along with metabolite profiling. Fortis Diphenyl is now available in 1.7µm particle size for UHPLC.



Fortis Diphenyl is based upon a unique di-phenyl functionality. Three controlled mechanisms of interaction can occur.

This allows for unique resolution of closely related species, and metabolites. No complex mobile phases are necessary simplifying method development.

- π - π High Selectivity
- Resolution Enhanced
- Sharp Peak Shapes
- Highly Stable Diphenyl Ligand

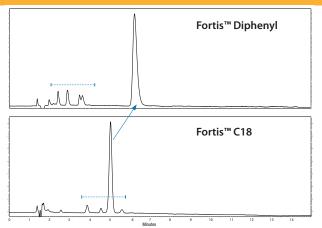


Diphenyl vs C18 Selectivity

Selectivity of the Fortis Diphenyl is radically different to that of a C18 stationary phase.

In this pharmaceutical mixture we can see an increase in retention of the parent drug, whilst the degradents are all eluted quickly, removing them from co-elution with the parent.

Selectivity such as this can be extremely useful, combined with the ability to separate closely related species such as metabolites and positional isomers.



Data Courtesy of : Major Pharmaceutical company, USA

Metabolite Profiling

Fortis Diphenyl's extended selectivity leads to its ability to discriminate between very closely related species, such as those often associated as metabolites or excipiants. The stationary phase's three modes of interaction allow subtle changes in positional spacing, loss or gain of an atom or functional group to be differentiated and separation to be achieved.

Separate Positional Isomers

Selectivity of compounds normally difficult to resolve on a hydrophobic alkyl chain stationary phase is simplified by the π - π interactions provided by the phenyl functionality.

In this application two hydroxyestradiol steroids exhibit resolution from each other, which is not achievable on alkyl chain phases. No complex mobile phases are necessary.

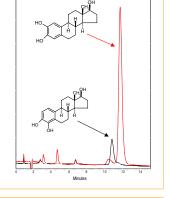
- Isomer Selectivity
- Metabolite Resolution
- Alternate Selectivity

Column: Fortis Diphenyl 150x4.6mm 5µ FPH-050705

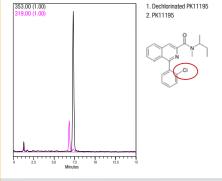
p/n: Mobile Phase: 40:60 H₂0: MeOH

1ml/min 210nm

- 1. 4-Hydroxyestradiol (mw=288.38)
- 2. 2-Hydroxyestradiol (mw=288.38)

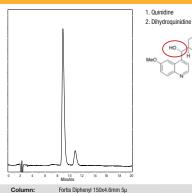


PET Tracer - PK11195



Column: Fortis Diphenyl 150x4.6mm 5µ FPH-050705 40:60 H₂0:ACN

Flow 1ml/min Temp: 25°C MS Detection



FPH-050705 70:30 H₂0 + 0.1% formic acid

1ml/min

25°C



1.7µm Fortis Diphenyl page 7

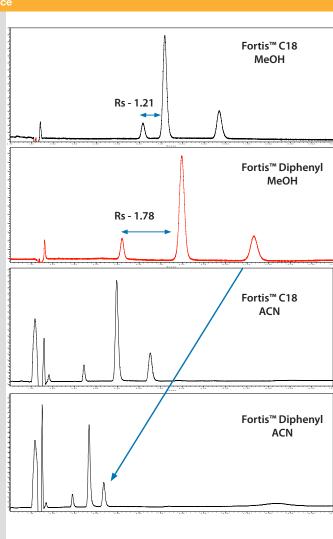
Fortis Diphenyl

Effect of Mobile phase choice

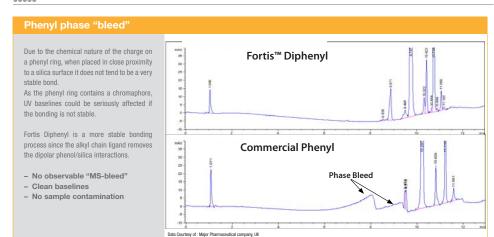
Choice of mobile phase can be very important in a running a phenyl column. Whilst many people have standardised upon ACN as the organic modifier of choice, MeOH is a better choice in order to let the $\pi\text{-}\pi$ interactions occur on the phenyl rings.

Using ACN can not only suppress retention but also selectivity.

It can be seen how maximum retention and resolution is obtained on Fortis Diphenyl in MeOH mobile phase, even greater than C18. Once the organic modifier is substituted for ACN not only is resolution reduced but also a large amount of retention is lost in relation to that lost on a C18.



Fortis[™] Diphenyl



| Fortis Diphenyl | | Column Length | | | |
|-----------------|-----|---------------|------------|------------|------------|
| | | 50 | 100 | 150 | 250 |
| | 2.1 | FPH-0203xx | FPH-0205xx | FPH-0207xx | - |
| Column Diameter | 3.0 | FPH-0303xx | FPH-0305xx | FPH-0307xx | - |
| | 4.6 | FPH-0503xx | FPH-0505xx | FPH-0507xx | FPH-0509xx |

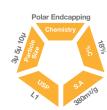
| Fortis Diphenyl Guards | | Length |
|------------------------|-----|--------------|
| | | 10 |
| Column Diameter 2.1 | | DCPH-0200xxG |
| | 4.6 | DCPH-0500xxG |

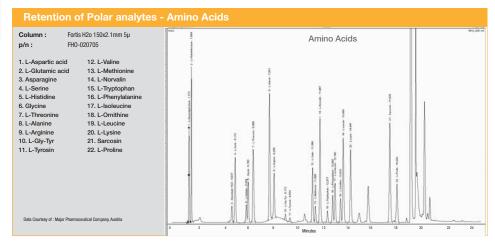
Replace xx - 01 for 1.7μm - 02 for 2.5μm - 03 for 3μm - 05 for 5μm - 10 for 10μm

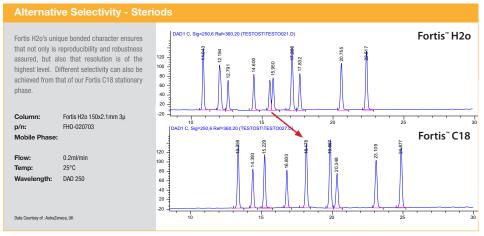
Fortis[™] H2o

- Retention of Polars by Polar Endcapping Group
- Enhanced Resolution
- 100% Aqueous Compatible
- Fully Scalable

Fortis H2o is designed to aid in the separation and retention of polar analytes. Complex mobile phase systems can be bypassed if sufficient retention can be provided by the stationary phase chemistry. Fortis H2o is designed to supply additional interaction with polar molecules which allows their successful retention



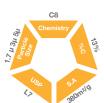




Fortis[™] C8

- Reduced Hydrophobicity over C18
- Excellent Peak Shapes
- Fully Scalable

Fortis C8 is designed to provide characteristics similar to Fortis C18 but specifically for situations where less hydrophobicity is required. The same gains in peak shape, efficiency, resolution and scaleability are available providing increased productivity to the analyst.



Optimised Peak Shape

Fortis C8 is optimised to provide the best possible peak shapes and efficiency.

Basic, Acidic and Neutral analyte performance is first class.

- Higher Efficiencies
- Greater Reproducibility
- Symmetrical peak shapes
- Lower Hydrophobicity

Column: Fortis C8 150x4.6mm 5μ **p/n:** F08-050705

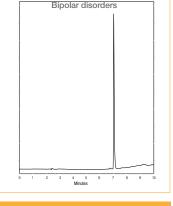
Mobile Phase: A - H₂O + 0.1% Formic acid B - MeOH + 0.1% Formic acid

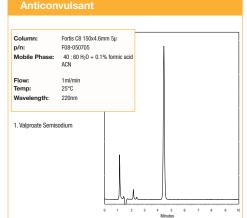
Gradient: 10 - 90% in 10min **Flow:** 1ml/min

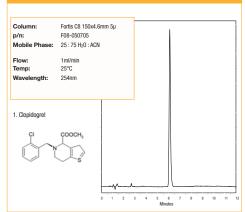
Temp: 25°C Wavelength: 254nn

Lamotrigine









Fortis[™] Cyano

- Retention of Polars
- Alternative Selectivity
- Normal Phase or Reverse Phase system
- Rapid Equilibration

Fortis Cyano allows the use of aqueous reversed phase conditions to provide less retention for compounds too heavily retained on C18 functionality. However, it can also be used in normal phase solvent systems to retain and separate polar analyte species.

Cyano columns are particularly useful for polar species. Fortis Cyano is now also available in $1.7\mu m$ particle size for UHPLC work.



Herhicides

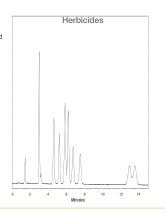
Fortis Cyano is optimised not only to help retain and resolve polar analytes, but also to be complimentary in resolution to other Fortis phases.

- Normal phase as well as Reversed phase use
- Alternative Selectivity
- Rapid Equilibration

- Column : Fortis Cyano 50x2.1mm 3μ p/n : FCN-020303
- Mobile Phase: 80:20 H₂0 : ACN + 0.2% Acetic acid
- Flow: 0.2ml/min

 Temp: 20°C

 Wavelength: 280nm
- 1. Banvel
- 2. Internal Std 3. 2,4-D
- 4. MCPA
- 5. PCOC 6. 2,4-DCP
- 7. 2,4-DP
- 8. CMPP
- 9. 2.4-DB
- 10. MCPB



To see more applications on Fortis Cyano turn to page 37. To learn more about Fortis Cyano 1.7µm see page 7.

| Fortis Cyano | Column Length | | | | |
|---------------------|---------------|------------|------------|------------|--|
| | 50 | 100 | 150 | 250 | |
| 2.1 | FCN-0203xx | FCN-0205xx | FCN-0207xx | - | |
| Column Diameter 3.0 | FCN-0303xx | FCN-0305xx | FCN-0307xx | - | |
| 4.6 | FCN-0503xx | FCN-0505xx | FCN-0507xx | F18-0509xx | |

Replace xx $\,$ - 01 for 1.7 μm $\,$ - 03 for 3 μm $\,$ - 05 for 5 μm

| Fortis Cyano Guards | | Length |
|---------------------|-----|--------------|
| | | 10 |
| Column Diameter | 2.1 | DCCN-0200xxG |
| | 4.6 | DCCN-0500xxG |

Fortis HILIC

- Retention of Polar Compounds
- Increased MS Sensitivity
- Alternate Selectivity
- Reduced Extraction (SPE) and Dry Down Times.

Fortis HILIC (Hydrophilic Interaction Chromatography) is designed to aid in the separation and retention of very polar analytes. Extended retention is afforded by the partitioning, ion-exchange and hydrogen bonding that can occur on a HILIC stationary phase. Fortis HILIC can increase sensitivity in MS analysis and provide alternate selectivity to that achieved with reversed phase C18. Fortis HILIC is now also available in 1.7µm particle size for UHPLC work.



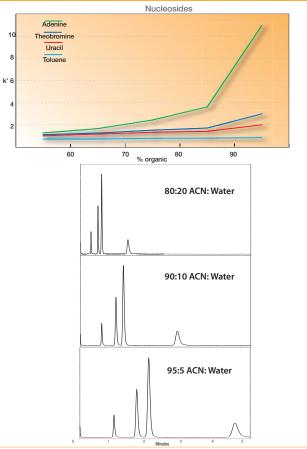
Polar retention in HILIC mode

Fortis HILIC is optimised to help retain and resolve polar analytes. By use of high concentrations of organic solvent polar analytes partition with the stationary phase.

- Polar Retention
- Alternative Selectivity
- Rapid Equilibration

Hydrophilic Interaction Chromatography (HILIC) works in a similar way to normal phase chromatography. A polar surface combined with a non-polar mobile phase, typically ACN, allows for partition of the polar analytes and hence retention and separation. Water is used in low concentration as the strong solvent in order to elute the compounds.

Usually no more than 20%-30% water is needed in order to elute most analyte species.





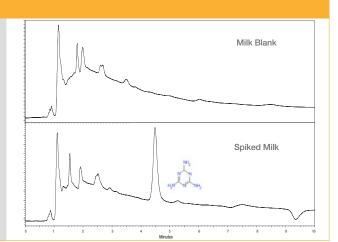
Melamine Contamination

Melamine has been adulterated into many products, but most importantly into baby milk in order to increase the apparent protein content. Due to its highly polar organic nature, 1,3,5-Triazine structure, it can be very difficult to retain in HPLC. HILIC provides a simple method in order to quickly quantitate melamine.

Column : Fortis HILIC 100x2.1mm 3μ **p/n :** FHI-020503

Mobile Phase: 90:10 ACN: 20mM NH₃OAc

Flow: 0.2ml/min
Temp: 20°C



Nucleosides

Nucleosides are typically difficult to retain due to the ribose or deoxyribose sugar that forms part of their structure. Fortis HILIC provides a good tool to retain and separate these polar analytes in simple mobile phase conditions.

Mobile Phase: 95:5 ACN:100mM NH₃0Ac

 Flow :
 1ml/min

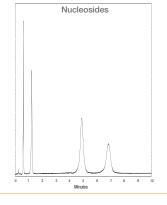
 Temp :
 20°C

 Wavelength:
 254nm

1. Uracil

2. Uridine

Cytosine
 Guanosine



| Fortis HILIC | Column Length | | | |
|---------------------|---------------|------------|------------|------------|
| | 50 | 100 | 150 | 250 |
| 2.1 | FHI-0203xx | FHI-0205xx | FHI-0207xx | = |
| Column Diameter 3.0 | FHI-0303xx | FHI-0305xx | FHI-0307xx | - |
| 4.6 | FHI-0503xx | FHI-0505xx | FHI-0507xx | FHI-0509xx |

Replace xx $\,$ - 01 for 1.7 μm $\,$ - 03 for 3 μm - 05 for 5 μm - 10 for 10 μm

| Fortis HILIC Guards | | Length | |
|---------------------|-----|--------------|--|
| | | 10 | |
| Column Diameter | 2.1 | DCHI-0200xxG | |
| | 4.6 | DCHI-0500xxG | |

- LC/MS Optimised Column Hardware

- 20mm and 30mm Column Lengths
- High Throughput
- High Efficiency and Resolution

Fortis Pace™ columns are designed with High Throughput Screening (HTS) applications in mind. Optimised for use in LC/MS to provide greatest sensitivity by achieving sharp peak shapes combined with excellent resolution and retention. Any Fortis stationary phase and particle sizes can be supplied in this hardware.



Optimised Hardware

Fortis Pace column hardware is specifically designed for HTS, whether isocratic or by ballistic gradients.

Optimised packing density in this low volume hardware leads to ultra sharp peak shapes combined with maximum efficiency.

By combining a low volume flow path with an optimised frit Fortis PACE columns provide improved efficiency, Asymmetry and pressure.

- Reduced peak widths
- Higher Efficiency
- Eliminated dead volume

| 0.5μ through hole | 0.005" through hole |
|-------------------|---------------------|

| Hardware Comparison | | | | | |
|----------------------------|---------|-------|----|--|--|
| plates/m Sym Bar | | | | | |
| Standard Hardware | 93,100 | 1.173 | 48 | | |
| PACE [™] Hardware | 100,176 | 1.113 | 38 | | |
| % Change +7.6% -5.4% -21% | | | | | |

Complimentary Stationary Phases

Fortis stationary phases have been proven to exhibit excellent peak shapes and efficiency, packed in Pace hardware allows speed and resolution to be achieved without the need for UHPLC systems.

Providing highly retentive and selective phases allows strong retention properties, enabling high concentrations of organic modifier to be utilised optimising the MS ionisation process.

Gains are also made in:

- Reduced analysis time
- Increased productivity
- Lower solvent consumption

Column: Fortis Pace C18 30x2.1mm 5μ p/n: F18-020205

Mobile Phase: A - H₂0 + 0.1% Formic acid B - ACN + 0.1% Formic acid

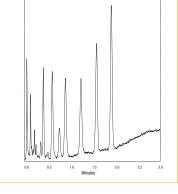
 Gradient:
 60 - 90% in 2min

 Flow:
 1ml/min

 Temp:
 25°€

Wavelength: 254nm

- 1. Uracil
- 2. Benzene
- 3. Ethylbenzene
- 4. Propylbenzene
- 5. Butylbenzene
- 6. Pentylbenzene
- 7. Hexylbenzene
- 8. Heptylbenzene



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www.fortis-technologies.com Tel: +44 151 336 2266



- 5µm and 10µm particles
- High Loadability
- Optimised Packing Efficiency
- Narrow peak profile, High Efficiency and Resolution

Fortis Prep columns are designed for high sample loading, high throughput applications. The optimised packed bed (OPB) process ensures excellent peak shapes and efficiency, whilst the lifetime of the column is increased.



Columns & Bulk

Fortis Prep columns come in sizes from 30mm to 250mm in length and from 10mm in diameter all the way up to 50mm $(2^n i.d.)$.

Pre-packed columns are advised for < 2" i.d. after this Bulk material can be supplied for those wishing to pack DAC (Dynamic Axial Compression) columns.

If preparative columns are packed with the identical media to their analytical counterpart then the ability to scale up with the theoretical calculations will be accurate.

- 10mm, 21.2mm and 30mm i.d.
- 5µm and 10µm particles
- 100g to multi Kg bulk available

Contact us for more information on availability of prep options/bulk packings, or to discuss your application and the ability to scale up. Our technical experts will be happy to discuss your needs with you.



Filter & Guard Options

- Guard system for all 3µm, 5µm and 10µm phases
- Low volume in-line filters for LC and UHPLC
- Maintain chromatographic integrity

Fortis Guards and filters are designed to ensure that erroneous materials do not find there way onto the more important and expensive analytical column. Guards are available in sizes to match all analytical and preparative column dimensions. Filters are particularly suitable for short fast LC/MS (PaceTM) columns and UHPLC columns.



- Direct connect guard system for all 3µm, 5µm and 10µm phases
- Quick replacement cartridges
- Highly Cost Effective



- Preparative guard system 10mm & 21.2mm
- Quick replacement cartridges
- Highly Cost Effective
- Reduced volume coupler available



- In-line Filter for all LC columns
- Low volume in-line filters
- Change over time is seconds not minutes

Fortis in-line filters are fingertight direct connect design, fitting in between the column and the conventional peek fitting to filter out particulate matter, it contains low dead volume and pressure. In-line filters are ideal for very short fast columns such as Fortis Pace LC/MS columns where extra packed bed from a quard would be detrimental.

In-line filters are also available in UHPLC format, capable of withstanding the elevated pressures involved.



- Filter for all UHPLC columns
- No backpressure increase
- Increase lifetime of UHPLC columns
- Low volume in-line filter
- Change over time is seconds not minutes

| In-line Filters | | | |
|-----------------|---------------------------|--|--|
| 2-SAV5 | 2μm In-line filter pk 5 | | |
| 2-SAV10 | 2μm In-line filter pk 10 | | |
| UHPSAV2 | UHPLC In-line filter pk 2 | | |
| UHPSAV4 | UHPLC In-line filter pk 4 | | |





Surface Area -175m²/g

Pore Size 140Å

% Carbon 11%C

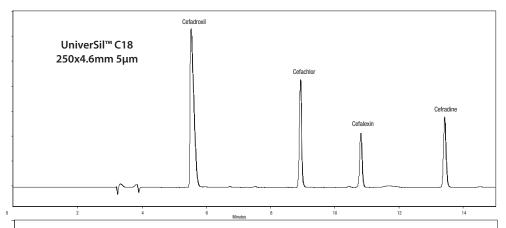
UniverSil HPLC columns are a suitable economical alternative to many of the older brands on the marketplace from Alltech, Macherey-Nagel, Hypersil and Waters. Many of the phases have very similar physical characteristics meaning that UniverSil will provide similar selectivity and resolution as your current chromatography. Use UniverSil to upgrade your method in terms of peak shape, robustness and reproducibility, or just to provide a economical back-up to your current column. UniverSil silica is a new Type B silica meaning improved peak shape and improved lifetime.

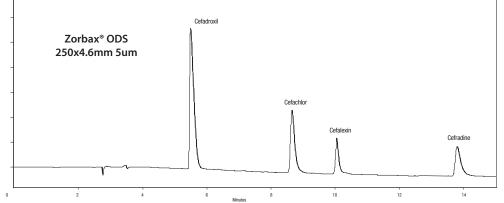
| UniverSil [™] C18 is an alternative to: | | | | |
|--|-----------------|---------|--|--|
| Alltech® ODS2 | Alltech | 220m²/g | | |
| Alltech® Exsil® C18 | Alltech | 220m²/g | | |
| Hypersil® ODS | Thermo Electron | 170m²/g | | |
| Hypersil® BDS | Thermo Electron | 170m²/g | | |
| Nucleosil® 120 C18 | Macherey-Nagel | 200m²/g | | |
| Pinnacle® C18 | Restek | 170m²/g | | |
| Spherisorb® ODS1 | Waters | 200m²/g | | |
| Spherisorb® ODS2 | Waters | 200m²/g | | |
| Supelcosil® LC18 | Supelco | 170m²/g | | |
| Ultrasphere® C18 | Beckman | 200m²/g | | |
| YMC [®] ODS-A | YMC | 170m²/g | | |
| Zorbax [®] Rx-C18 | Agilent | 180m²/g | | |

UniverSil™ C18 is a trademark of Fortis™ Technologies Ltd.

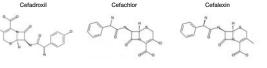
For the contract of the contra

Fortis is not associated with these companies. Comparative separations may not be representative of all applications. All columns are original manufacturers own.





| Column: | UniverSil [™] C18 250x4.6mm 5µ | |
|---------------|---|------------|
| p/n: | U18-050905 | Cefadroxil |
| Mobile Phase: | A: Phosphate pH3 B: ACN | They C |
| Flow: | 1.0ml/min | |
| Temp: | 25°C | h. |



| iverSil [™] C18 | | | Column | Length | | UniverSil C |
|--------------------------|-----|------------|------------|------------|------------|-------------|
| | | 50 | 100 | 150 | 250 | |
| | 2.1 | U18-0203xx | U18-0205xx | U18-0207xx | - | Column Di |
| lumn Diameter | 3.0 | U18-0303xx | U18-0305xx | U18-0307xx | - | |
| | 4.6 | U18-0503xx | U18-0505xx | U18-0507xx | U18-0509xx | |

| Universil C16 Guards | | Length |
|----------------------|-----|---------------|
| | | 10 |
| Column Diameter | 2.1 | DCU18-0200xxG |
| | 4.6 | DCU18-0500xxG |

Replace xx - 03 for 3µm - 05 for 5µm

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Wavelength: 254nm





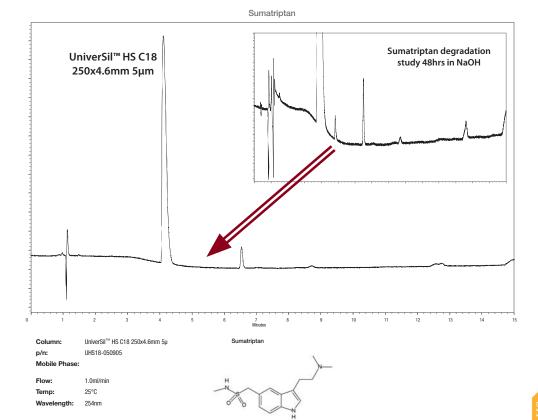
Surface Area - 325m²/g

Pore Size - 100Å

% Carbon - 16%C

UniverSil HPLC columns are a suitable economical alternative to many of the older brands on the marketplace from Kromasil, Macherey Nagel, Hypersil and Merck. Many of the phases have very similar physical characteristics meaning that UniverSil will provide similar selectivity and resolution as your current chromatography. Use UniverSil to upgrade your method in terms of peak shape, robustness and reproducibility, or just to provide a economical back-up to your current column. UniverSil silica is a new Type B silica meaning improved peak shape and improved lifetime.

| UniverSil [™] HS C18 is an alternative to: | | | |
|---|--------------------------|---------|--|
| Alltech® Alltima | Alltech® Alltima Alltech | | |
| Betasil® C18 | Thermo Electron | 300m²/g | |
| Cosmosil® 18 | Nacalai Tesque | 330m²/g | |
| Genesis® ODS | Jones | 300m²/g | |
| Hypersil® HS | Thermo Electron | 300m²/g | |
| Inertsil® ODS2 | GL Sciences | 320m²/g | |
| Kromasil® C18 | Akzo Nobel | 340m²/g | |
| Lichrosphere® RP18 | Merck | 350m²/g | |
| Nucleosil® 100 C18 | Macherey Nagel | 350m²/g | |
| Symmetry® C18 | Waters | 335m²/g | |
| YMC [®] Pack ODS AL | YMC | 300m²/g | |
| Zorbax® C18 | Agilent | 300m²/g | |



| UniverSil™ HS C18 | | Column Length | | | |
|-------------------|-----|---------------|--------------|--------------|--------------|
| | | 50 | 100 | 150 | 250 |
| | 2.1 | UHS18-0203xx | UHS18-0205xx | UHS18-0207xx | - |
| Column Diameter | 3.0 | UHS18-0303xx | UHS18-0305xx | UHS18-0307xx | - |
| | 4.6 | UHS18-0503xx | UHS18-0505xx | UHS18-0507xx | UHS18-0509xx |

| UniverSil HS C18 Gua | Length | |
|----------------------|--------|-----------------|
| | | 10 |
| Column Diameter | 2.1 | DCUHS18-0200xxG |
| | 4.6 | DCUHS18-0500xxG |

Replace xx - 03 for 3µm - 05 for 5µm

UniverSII** HS C18 is a trademark of Fortis** Technologies Ltd.

Symmetry* is a registered trademark of Vilates Corporation. Alllech* is a registered trademark of Alltech. Zorbax* is a registered trademarks of Agilent Technologies. YMC* is a registered trademark of YMC. Lichrosphere* is a registered trademark of Hermo Electron. Genesis* is a registered trademark of Jones. Kromasil* is a registered trademark of Alzon Nobel. Inertsil* is a registered trademark of Nobel. Inertsil* is a registered trademark of Sciences. Cosmosil* is a registered trademark of Nobel. Inertsil* is a registered trademark of Sciences.

Fortis is not associated with these companies. Comparative separations may not be representative of all applications. All columns are original manufacturers own.

Column Reproducibility

- Robust Column Bondings
- Assured Peak Shapes
- 20% Lower Asymmetry Specification
- 10% Higher Efficiency

Fortis HPLC columns are tested using the industries most rigorous QC test, utilising basic analyte probes as well as neutral efficiency markers ensures that the column reproducibility is first class. Fortis columns are also subject to a 20% lower peak shape specification than other manufacturers columns.

QC Test

Fortis stationary phases have been proven to exhibit excellent peak shapes and efficiency for the full range of analyte species.

By employing a QC mix that accurately probes silanol activity (the measure of good peak shape) the analyst can be assured of quality time and time again.

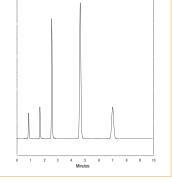
Gains are also made in:

- Sample carry over
- Increased Resolution
- Increased Sensitivity

Column: Fortis C18 100x4.6mm 5µ p/n: F18-050505 Mobile Phase: 60:40 ACN:H₂0

Temp: Wavelength: 254nm

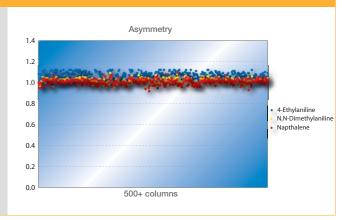
- 1. Uracil
- 3. 4-Ethylaniline
- 4. N.N-Dimethylaniline
- 5. Napthalene



Fortis columns are subject to tight specification using basic analytes in an unbuffered mobile phase system.

If there were residual uncovered hydroxyl groups present then these basic probes would highlight this fact.

Fortis Technologies unique bondings combined with the ultra pure silica matrix ensure that the peak shapes achieved are first class.



Applications



Column: 1.7µm Fortis C18 100x2.1mm

F18-020501

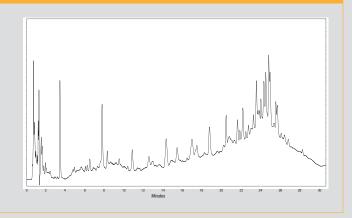
Mobile Phase: A - H₂0 + 0.1% Formic acid B - ACN + 0.1% Formic acid

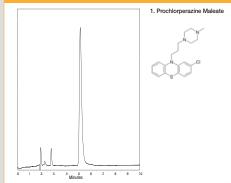
Omin 5% B

5min 5% B 30min 20% B 35min 50% B

0.25 ml/min Flow: 35°C 200nm

1. Casein



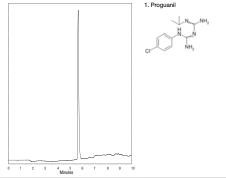


Column: Fortis Diphenyl 150x4.6mm 5u FPH-050705

: MeOH 25°C Temp

1ml/min 254nm

35:65 H₂O + 0.1% Formic acid



Column Fortis Diphenyl 50x4.6mm 5u FPH-050305 A: H₂O + 0.1% formic acid Mobile Phase: B: MeOH + 0.1% formic acid 10 - 90% in 10mins 1ml/min

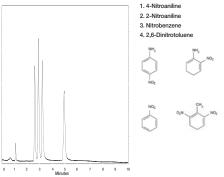
25°C



Fortis Diphenyl 150x4.6mm 5µ FPH-050705

30:60 H₂0 + 0.1% Formic acid 1ml/min

Flow

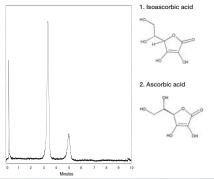


Fortis Cyano 50x2.1mm 3µ FCN-020303 A: 80:20 H₂0 : MeOH B: ACN 90:10 Isocratio 0.2ml/min Ambient

254nm

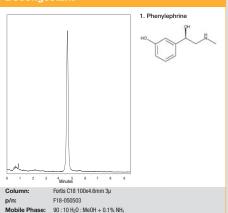
25°C

25°C



Fortis HILIC 50x4.6mm 5µ FHI-050305 ase: 90:10 ACN:NH3OAc

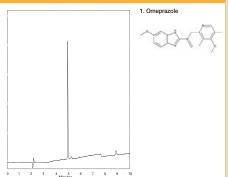
1ml/min 25°C 254nm



Applications

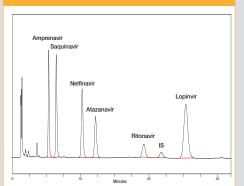






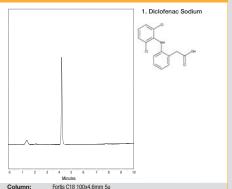
Column: Fortis C18 100x4.6mm 3µ F18-050503 A: H₂O + 0.1% Formic acid Mobile Phase: B: ACN + 0.1% Formic acid

Gradient: 10 - 100% in 10mins Flow: 1.0ml/min Temp: 25°C 254nm



Fortis C18 150x4 6mm 5u Column: F18-050705 A: H₂O + Buffer B: MeOH + ACN

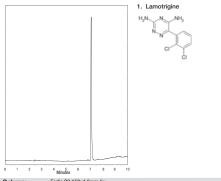
1.0ml/min 25°C 215nm



Column F18-050505 p/n: Mobile Ph Gradient 50 - 100% in 10mins

Temp:

A: H₂0 + 0.1% Formic acid B: ACN + 0.1% Formic acid Flow: 1.0ml/min Wavelength:



Fortis C8 150x4.6mm 5µ F08-050705 A: H₂0 + 0.1% formic acid B: MeOH + 0.1% formic acid 10 - 90% in 10mins 1.0ml/min

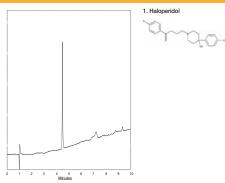
25°C



1. Loratadine

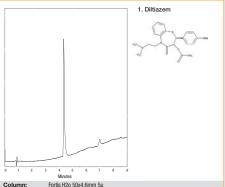
Fortis C18 100x4.6mm 5µ Column F18-050505 Mobile Phase: A: H₂0 + 0.1% Formic acid B: ACN + 0.1% Formic acid 10 - 100% in 10mins

1.0ml/min 254nm

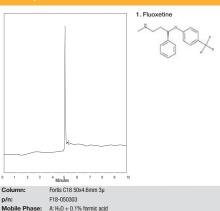


Fortis C18 50x4.6mm 5µ F18-050305 Mobile Phase: A: H₂0 + 0.1% formic acid B: ACN + 0.1% formic acid 10 - 100% in 5mins

25°C 254nm



FHO-050305 A: H₂0 + 0.1% formic acid B: ACN + 0.1% formic acid 1.0ml/min 25°C



B: ACN + 0.1% formic acid

20°C

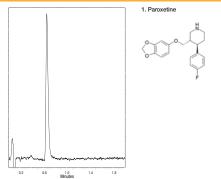
Temp:

Wavelength:

Applications



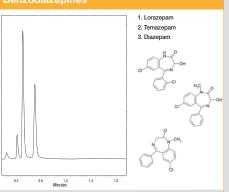




Column Fortis Pace H2o 30x2.1mm 3µ FH0-020203

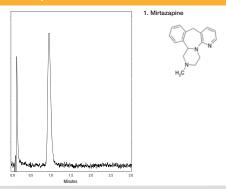
Mobile Phase: 70:30 H₂0 : ACN + 0.1% Formic acid

25°C



Fortis Pace H2o 30x2.1mm 3µ FH0-020203 Mobile Phase: 50:50 ACN : H₂0 0.7ml/mir

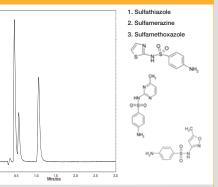
220nm



Column: Fortis Pace C18 30x2.1mm 5u F18-020205

Mobile Phase: 80:20 H₂0 : MeOH + 0.1% Formic acid

0.7ml/min Temp: 25°C



Fortis Pace H2o 30x2.1mm 3u Column: FH0-020203 70:30 H₂0 : ACN + 0.1% Formic acid

0.2ml/min 25°C

Applications



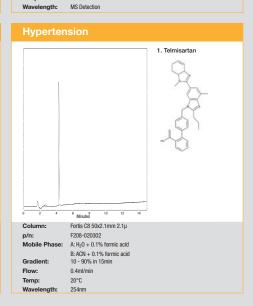
1. Rosuvastatin 2. Atorvastatin

Fortis Pace C18 30x2.1mm 5µ F18-020205 Mobile Phase: 75:25 H₂0:ACN + 0.1% Formic acid

0.4ml/min 254nm

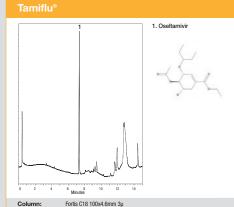
1. MDA (3,4-Methylenedioxyamphetamine) 2. Amphetamine 3. MDMA (Ecstasy) 4. Methaphetamine 5. MDEA (3,4-Methylenedioxy-N-ethylamphetamine) Fortis C18 150x3.0mm 5µ F18-030705 Mobile Phase: A: H₂0 + 0.1% ammonia B: ACN + 0.1% ammonia 20°C

1. Progesterone 2. 17αHydroxyprogesterone 3. 11 a Hydroxyprogesterone 4. Hydroxy-21-acetate 5. Cortisone 6. Prednisone 7. Prednisolone Fortis Cyano 50x2.1mm 1.7µ FCN-020301 Heptane: THF: MeOH 90:5:5 25°C Wavelength: 254nm

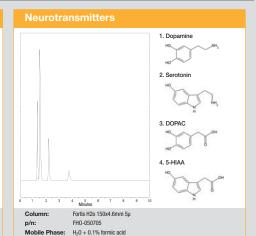


Applications





F18-050503 Mobile Phase: A: H₂0 + 0.1% ammonia B: ACN + 0.1% ammonia Gradient 10 - 100% in 15mins Flow: 1.0ml/min Temp: 20°C Wavelength: 254nm

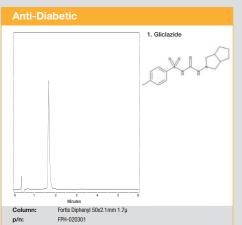


0.8ml/min

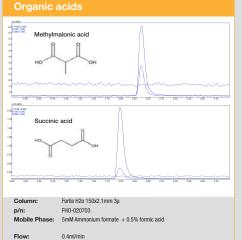
MS Detection

20°C

270nm



Mobile Phase: 80:20 H₂0 : ACN + 0.1% formic acid Flow: Temp: 25°C





1. Amiloride

Column Fortis C18 100x4.6mm 3µ F18-050503 Mobile Phase: 75:25 H₂0+ 0.1% NH₃:Me0H

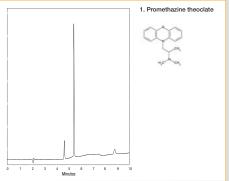
1.0ml/min 254nm

1. Tenofovir 2. Thymidine (IS)

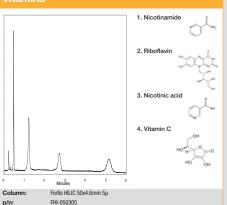
Fortis H2o 150x4.6mm 5µ FH0-030705

Mobile Phase: 95:5 10mM Ammonium formate pH6.5 : Me0H

20°C



Fortis C18 100x4.6mm 3µ F18-050503 se: A: H₂0 + 0.1% ammonia B: ACN + 0.1% ammonia 10 - 100% in 10mins 1.0ml/min Temp: 20°C



Mobile Phase: 90:10 ACN:100mM ammonium acetate

1.0ml/min

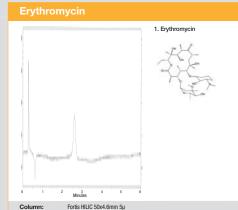
20°C

254nm

Temp:

Applications

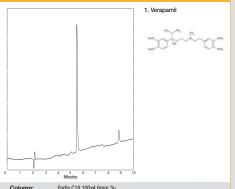




FHI-050305

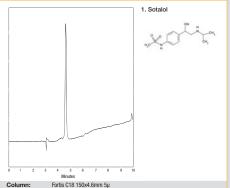
Mobile Phase: 90: 10 ACN : 100mM ammonium acetate

0.8ml/min 200nm



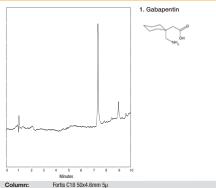
Fortis C18 100x4.6mm 3µ F18-050503 A: H₂O + 0.1% formic acid B: ACN + 0.1% formic acid 10 - 90% in 10min 1 Oml/min 20°C

Beta-Blockers



F18-050705 A: H₂0 + 0.1% formic acid B: ACN + 0.1% formic acid Flow: 1.0ml/min

Temp: 20°C



F18-050305 A: H₂O + 0.1% formic acid B: ACN + 0.1% formic acid 1.0ml/min 20°C



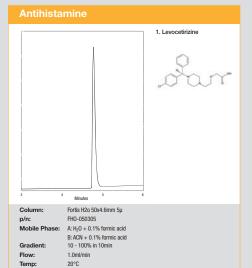
1. Folic acid

Fortis H2o 50x4.6mm 5µ

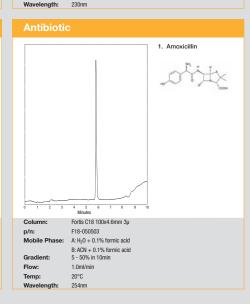
FH0-050305

1.0ml/min

Mobile Phase: 60:40 H₂0 : ACN

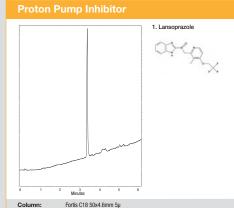


1. Citalopram Fortis C18 50x4.6mm 5µ F18-050305 A: 50mM ammonia acetate B: ACN Flow: 1.0ml/min 20°C

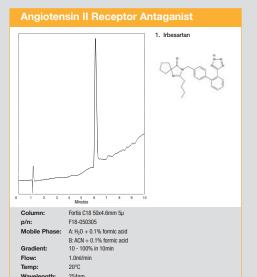


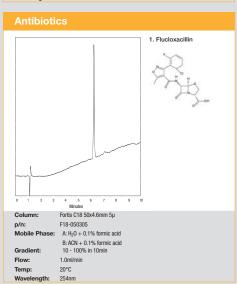
Applications

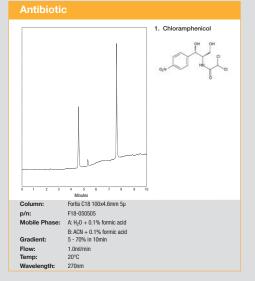




F18-050305 A: H₂O + 0.1% formic acid B: ACN + 0.1% formic acid 10 - 100% in 10min 1 Oml/min Flow: Wavelength:







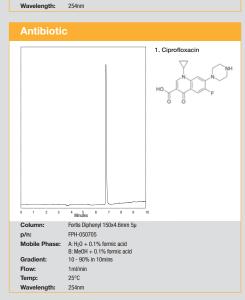


1. Tramadol 2.0 2.5 3.0 3.5 4.0 4.5 5.0 Minutes

Fortis C18 50x2.1mm 5µ F18-020305 A: H₂0 + B: ACN + 0.1% NH 10 - 100% in 5mins Ambient Temp:

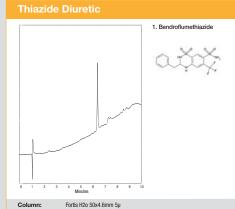
1. Phenoxymethylpenicilin Fortis C18 50x4.6mm 5µ F18-050305 A: H₂0 + 0.1% formic acid B: ACN + 0.1% formic acid 10 - 100% in 10mins 1ml/min Flow: 25°C

Fortis Diphenyl 150x4.6mm 5µ FPH-050705 Mobile Phase: 100% H₂0 + 0.1% Formic acid 0.8ml/min

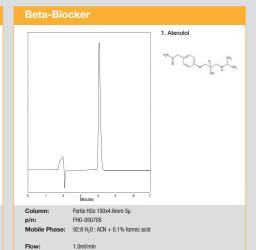


Applications

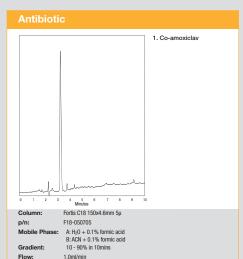


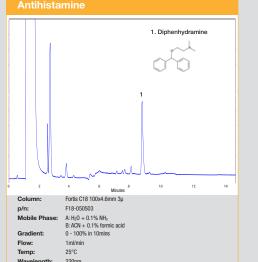


FH0-050305 A: H₂O + 0.1% formic acid B: ACN + 0.1% formic acid 10 - 100% in 10min 1 Oml/min Flow: Wavelength:



20°C





25°C

254nm

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Part number tables

| Fortis C18 | | Column Length | | | | | | |
|-----------------|-----|---------------|------------|------------|------------|------------|------------|--|
| | | 20 | 30 | 50 | 100 | 150 | 250 | |
| | 2.1 | F18-0201xx | F18-0202xx | F18-0203xx | F18-0205xx | F18-0207xx | - | |
| Column Diameter | 3.0 | - | F18-0302xx | F18-0303xx | F18-0305xx | F18-0307xx | - | |
| | 4.6 | - | F18-0502xx | F18-0503xx | F18-0505xx | F18-0507xx | F18-0509xx | |

Replace xx $\,$ -01 for 1.7 μm $\,$ - 02 for 2.5 μm $\,$ - 03 for 3 μm $\,$ - 05 for 5 μm $\,$ - 10 for 10 μm

| Fortis Diphenyl | | Column Length | | | | | | |
|-----------------|-----|---------------|------------|------------|------------|------------|------------|--|
| | | 20 | 30 | 50 | 100 | 150 | 250 | |
| | 2.1 | FPH-0201xx | FPH-0202xx | FPH-0203xx | FPH-0205xx | FPH-0207xx | - | |
| Column Diameter | 3.0 | - | FPH-0302xx | FPH-0303xx | FPH-0305xx | FPH-0307xx | - | |
| | 4.6 | - | FPH-0502xx | FPH-0503xx | FPH-0505xx | FPH-0507xx | FPH-0509xx | |

Replace xx - 01 for 1.7μm - 02 for 2.5μm - 03 for 3μm - 05 for 5μm - 10 for 10μm

| Fortis H2o | Column Length | | | | | | |
|-----------------|---------------|------------|------------|------------|------------|------------|------------|
| | | 20 | 30 | 50 | 100 | 150 | 250 |
| | 2.1 | FH0-0201xx | FH0-0202xx | FH0-0203xx | FH0-0205xx | FH0-0207xx | - |
| Column Diameter | 3.0 | - | FH0-0302xx | FH0-0303xx | FH0-0305xx | FH0-0307xx | - |
| | 4.6 | - | FH0-0502xx | FH0-0503xx | FH0-0505xx | FH0-0507xx | FH0-0509xx |

Replace xx - 03 for 3µm - 05 for 5µm

| Fortis C8 | | Column Length | | | | | |
|-----------------|-----|---------------|------------|------------|------------|------------|------------|
| | | 20 | 30 | 50 | 100 | 150 | 250 |
| | 2.1 | F08-0201xx | F08-0202xx | F08-0203xx | F08-0205xx | F08-0207xx | - |
| Column Diameter | 3.0 | - | F08-0302xx | F08-0303xx | F08-0305xx | F08-0307xx | - |
| | 4.6 | - | F08-0502xx | F08-0503xx | F08-0505xx | F08-0507xx | F08-0509xx |

Replace xx - 03 for 3µm - 05 for 5µm

| Fortis Cyano | Column Length | | | | | | |
|-----------------|---------------|------------|------------|------------|------------|------------|------------|
| | | 20 | 30 | 50 | 100 | 150 | 250 |
| | 2.1 | FCN-0201xx | FCN-0202xx | FCN-0203xx | FCN-0205xx | FCN-0207xx | - |
| Column Diameter | 3.0 | - | FCN-0302xx | FCN-0303xx | FCN-0305xx | FCN-0307xx | - |
| | 4.6 | - | FCN-0502xx | FCN-0503xx | FCN-0505xx | FCN-0507xx | F18-0509xx |

Replace xx $\,$ - 01 for 1.7 μm $\,$ - 03 for 3 μm $\,$ - 05 for 5 μm

| Fortis HILIC | | Column Length | | | | | | |
|-----------------|-----|---------------|------------|------------|------------|------------|------------|--|
| | | 20 | 30 | 50 | 100 | 150 | 250 | |
| | 2.1 | FHI-0201xx | FHI-0202xx | FHI-0203xx | FHI-0205xx | FHI-0207xx | - | |
| Column Diameter | 3.0 | | FHI-0302xx | FHI-0303xx | FHI-0305xx | FHI-0307xx | - | |
| | 4.6 | | FHI-0502xx | FHI-0503xx | FHI-0505xx | FHI-0507xx | FHI-0509xx | |

Replace xx $\,$ - 01 for 1.7 μm $\,$ - 03 for 3 μm $\,$ - 05 for 5 μm $\,$ - 10 for 10 μm

| 5μm F | 5µm Fortis Guard Cartridges | | | | | | |
|----------------|------------------------------|--|--|--|--|--|--|
| DCGUA-1 | Guard Cartridge Holder | | | | | | |
| DCxx-040005G/2 | 10x4mm Fortis 5µm Guard pk 2 | | | | | | |
| DCxx-040005G/4 | 10x4mm Fortis 5µm Guard pk 4 | | | | | | |
| DCxx-020005G/2 | 10x2mm Fortis 5µm Guard pk 2 | | | | | | |
| DCxx-020005G/4 | 10x2mm Fortis 5µm Guard pk 4 | | | | | | |

| 3μm Fortis Guard Cartridges | | | | | | | |
|-----------------------------|------------------------------|--|--|--|--|--|--|
| DCGUA-1 | Guard Cartridge Holder | | | | | | |
| DCxx-040003G/2 | 10x4mm Fortis 3µm Guard pk 2 | | | | | | |
| DCxx-040003G/4 | 10x4mm Fortis 3µm Guard pk 4 | | | | | | |
| DCxx-020003G/2 | 10x2mm Fortis 3µm Guard pk 2 | | | | | | |
| DCxx-020003G/4 | 10x2mm Fortis 3µm Guard pk 4 | | | | | | |

Replace xx 18 for Fortis C18 PH for Fortis Diphenyl HO for Fortis H2o 08 for Fortis C8 CN for Fortis Cyano HI for Fortis HILIC

| Analytical In-line Filters | | | | |
|----------------------------|--------------------------|--|--|--|
| 2-SAV5 | 2μm In-line filter pk 5 | | | |
| 2-SAV10 | 2μm In-line filter pk 10 | | | |

| UHPLC In-line Filters | | | | |
|-----------------------|---------------------------|--|--|--|
| UHPSAV2 | UHPLC In-line filter pk 2 | | | |
| UHPSAV4 | UHPLC In-line filter pk 4 | | | |

WORLDWIDE AVAILABILITY





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