

**INSTRUCTION MANUAL FOR SFC OPTIMIZED (3.0 mm i.d.)  
CHIRALPAK® IA-3, IB-3, IB N-3, IC-3, ID-3, IE-3, IF-3, IG-3, IH-3, and IJ-3**

**<Supercritical Fluid Chromatography (SFC)>**

**Please read this instruction sheet completely before using these columns.**

These columns can also be used in reversed-phase and normal phase. Please refer to the corresponding instruction sheet for details.

**Column Description**

| <p><b>AMYLOSE-BASED</b></p> <p><b>Immobilized on 3 μm silica gel</b></p>            |   | <p><b>CELLULOSE-BASED</b></p> <p><b>Immobilized on 3 μm silica gel</b></p>                                    |
|---|---|---|
| <p><b>CHIRALPAK® IA-3</b></p> <p>Amylose tris(3,5-dimethylphenylcarbamate)</p>      | <p><b>CHIRALPAK® ID-3</b></p> <p>Amylose tris(3-chlorophenylcarbamate)</p>          | <p><b>CHIRALPAK® IB-3</b><br/><b>CHIRALPAK® IB N-3</b></p> <p>Cellulose tris(3,5-dimethylphenylcarbamate)</p> |
| <p><b>CHIRALPAK® IE-3</b></p> <p>Amylose tris(3,5-dichlorophenylcarbamate)</p>      | <p><b>CHIRALPAK® IF-3</b></p> <p>Amylose tris(3-chloro-4-methylphenylcarbamate)</p> | <p><b>CHIRALPAK® IC-3</b></p> <p>Cellulose tris(3,5-dichlorophenylcarbamate)</p>                              |
| <p><b>CHIRALPAK® IG-3</b></p> <p>Amylose tris(3-chloro-5-methylphenylcarbamate)</p> | <p><b>CHIRALPAK® IH-3</b></p> <p>Amylose tris[(S)-α-methylbenzylcarbamate]</p>      | <p><b>CHIRALPAK® IJ-3</b></p> <p>Cellulose tris(4-methylbenzoate)</p>   |

Shipping Solvent:

**Methanol = 100%**

All columns have been pre-tested before packaging. Test parameters and results, as well as the Column Lot Number, were included with the column when purchased.

\*Because different columns, including custom columns, can be shipped in different solvents, we recommend flushing them with 100% Ethanol or Isopropanol, at the typical flow rate listed below, before their first use to avoid any damage (see column transfer conditions between LC and SFC on page 5).\*

**THIS INSTRUCTION MANUAL IS NOT APPLICABLE TO ANY OTHER DAICEL COLUMNS**

## Operating Instructions

|                                  | 50 x 3.0 mm i.d.<br>100 x 3.0 mm i.d.<br>150 x 3.0 mm i.d.<br>Analytical Columns                             |
|----------------------------------|--|
| Flow Rate Direction              | As indicated on the column label   |
| Typical Flow Rate <b>in SFC</b>  | ~ 0.5 - 4.0 ml/min   |
| Pressure Limitation <sup>①</sup> | < 300 bar (4350 psi) for maximum column life<br><b>Typical CO<sub>2</sub> backpressure (BPr) 110-150 bar</b> |
| Temperature                      | 0 to 40°C  |
| Column Fitting                   | Please contact <a href="#">Technical Support</a> for details   |

① The relevant pressure value is the one generated by the column itself (pressure drop). The pressure drop is the difference between the inlet pressure ( $P_{inlet}$ ) and the outlet pressure ( $P_{outlet}$ ) in the system. The pressure drop generated by the system alone (without any column) has to be subtracted from the total value (system + column).

The column can be operated up to 300 Bar (pressure drop). However, it is necessary to check if the SFC system has been designed to withstand these conditions. The flow rate has to be adapted considering the pressure drop in the column (this pressure being dependent upon flow rate as well as the amount and type of modifier in the mobile phase).

## Method Development / SFC

### A - Mobile Phases

CHIRALPAK® IA-3, IB-3, IB N-3, IC-3, ID-3, IE-3, IF-3, IG-3, IH-3, and IJ-3 can be used **with all ranges of organic miscible solvents as co-solvent combined with the carbon dioxide (CO<sub>2</sub>)**, progressing from the traditional solvents (mixtures of CO<sub>2</sub> with alcohols or acetonitrile (ACN)), to mobile phases containing CO<sub>2</sub> with tetrahydrofuran (THF), dichloromethane (DCM), chloroform (CHCl<sub>3</sub>), ethyl acetate (EtOAc), and methyl *tert*-butyl ether (MtBE), among others.

### B - Method Development - Screening

When developing methods, we would recommend a screening approach.

1. The conditions described in Table 1 should be used as Primary Screening.
2. If the compound or compound series are not soluble in any of these mobile phases, we recommend to try the Primary Screening with the product dissolved in a stronger solvent (DCM/alcohol...).

**Table 1. Immobilized Primary Screening Solvents**

| Primary Solvent Mixtures    | CO <sub>2</sub> / MeOH | CO <sub>2</sub> / EtOH | CO <sub>2</sub> / 2-PrOH | CO <sub>2</sub> / ACN <sup>①</sup> |
|-----------------------------|------------------------|------------------------|--------------------------|------------------------------------|
| Typical Starting Conditions | 80:20                  | 80:20                  | 80:20                    | 70:30 <sup>①</sup>                 |
| Advised Optimization Range  | 99:1<br>to 40:60       | 99:1<br>to 40:60       | 99:1<br>to 40:60         | 99:1<br>to 40:60 <sup>①</sup>      |

① Alcohols can be added into ACN to enhance the eluting strength for strongly retained compounds.

If a suitable chiral separation is not found using the Primary Screening strategy, we recommend a Secondary Screening to be applied using the following conditions:

**Table 2. Immobilized Secondary Screening Solvents**

| Secondary Solvent Mixtures  | CO <sub>2</sub> / THF | CO <sub>2</sub> / (DCM+MeOH 90:10) | CO <sub>2</sub> / (EtOAc+MeOH 90:10) | CO <sub>2</sub> / (MtBE+MeOH 80:20) |
|-----------------------------|-----------------------|------------------------------------|--------------------------------------|-------------------------------------|
| Typical Starting Conditions | 75:25                 | 80:20                              | 80:20                                | 75:25                               |
| Advised Optimization Range  | 99:1<br>to 40:60      | 99:1<br>to 40:60                   | 99:1<br>to 40:60                     | 99:1<br>to 40:60                    |

Notes: The alcohol content and type (MeOH, EtOH and 2-PrOH) can be used to modulate retention and recognition. THF can be added into DCM and EtOAc to enhance the eluting strength for strongly retained compounds.

All solvent proportions indicated in this manual are by volume.

### C – General Comments

- ⇒ Only immobilized CHIRALPAK® IA-3, IB-3, IB N-3, IC-3, ID-3, IE-3, IF-3, IG-3, IH-3, and IJ-3 are suitable for the Secondary Screening.
- ⇒ Additional modifiers such as CHCl<sub>3</sub>, 1,4-Dioxane, Toluene, or Acetone can also be investigated with CHIRALPAK® IA-3, IB-3, IB N-3, IC-3, ID-3, IE-3, IF-3, IG-3, IH-3, and IJ-3 columns.
- ⇒ The typical starting conditions consist in mobile phases of upper middle eluting strength. Under such conditions, most of the analytes can be eluted within a reasonable time range with a good probability of full resolution of the enantiomers.
- ⇒ It is also important to ensure your SFC system (seals...) is compatible with all types of solvents and to take into account UV cut-off of certain solvents, in order to avoid detection issues. Detection with a regular UV detector may become difficult depending on a combination of sample and mobile phase (e.g. EtOAc, high percentages of DCM).

## D – Additives

For basic samples, it is necessary to incorporate an additive into the mobile phase in order to optimize the chiral separation.

Acidic samples **do not always** require the presence of an additive. In fact, the acidic properties of the carbon dioxide (CO<sub>2</sub>) are sometimes enough to elute the product properly.

- In practice, 1% of the additive is incorporated with the modifier. The total amount of additive into the mobile phase will be dependent upon the percentage of modifier. For example, if the mobile phase is CO<sub>2</sub> / EtOH = 90: 10, with EtOH containing 1% of additive, then the mobile phase composition will be CO<sub>2</sub> / EtOH / additive = 90: 10: 0.1).

| Basic Samples require Basic additives ❶   | Acidic Samples require Acidic additives❷                 |
|---|--|
| Diethylamine (DEA)<br>Triethylamine (TEA) | Trifluoroacetic acid (TFA)<br>Acetic acid<br>Formic acid |

⇒ **STRONGLY BASIC solvent additives or sample solutions MUST BE AVOIDED, because they are likely to damage the silica gel used in this column**

## Column Care / Maintenance

- ❑ The use of a guard cartridge is highly recommended for maximum column life.
- ❑ Samples should preferably be dissolved in the modifier.
- ❑ Sample solutions should be filtered through a membrane filter of approximately 0.5µm porosity to ensure that there is no precipitate before use.

Following extensive use of the column in multiple solvents, there may be a change in separation reproducibility. In order to ensure consistent performance, a regeneration method may be implemented to eliminate any change in chiral recognition due to the history of the column (mobile phases, additives...).

**For detailed Regeneration Procedures, please [click here](#)**

### ↻ Column transfer between modes:

#### From LC to SFC

- Flush with 100% EtOH at 0.2 ml/min<sup>(\*)</sup> for 45 min
- Flush with 100% CO<sub>2</sub> or CO<sub>2</sub>+co-solvent at 0.25 ml/min<sup>(\*)</sup> for 45 min

#### From SFC to LC

- Flush with 100% EtOH at 0.2 ml/min<sup>(\*)</sup> for 45 min
- Flush with the mobile phase at 0.25 ml/min<sup>(\*)</sup> for 45 min

<sup>(\*)</sup> *Recommended flow rate for analytical columns (3.0mm i.d.).*

## Column Storage

- ❑ For column storage, remove the acidic or basic additives by flushing the column with several column volumes of 100% 2-PrOH or 100% methanol, without additives.
- ❑ Columns can be stored with ends capped in the additive-free mobile phase, or the shipping solvent, at room temperature.

***Operating these columns in accordance with the guidelines outlined here will result in a long column life.***

⇒ If you have any questions about the use of these columns, or encounter a problem, contact:

In the USA: [questions@cti.daicel.com](mailto:questions@cti.daicel.com) or call 800-6-CHIRAL

In the EU: [cte@cte.daicel.com](mailto:cte@cte.daicel.com) or call +33 (0) 3 88 79 52 00

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