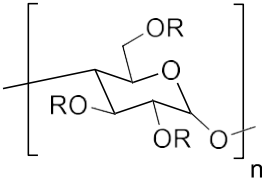
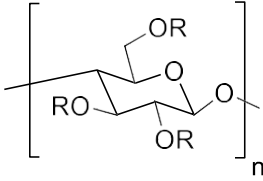
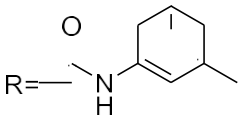
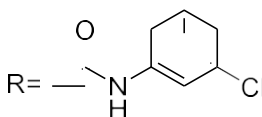
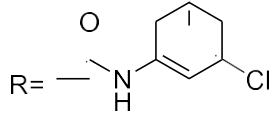
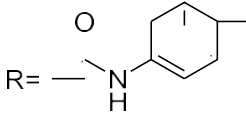
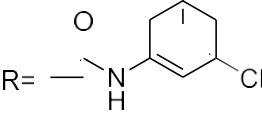
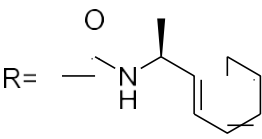
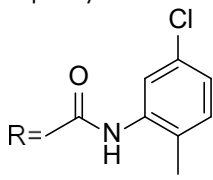
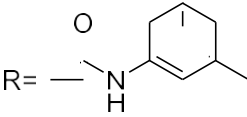
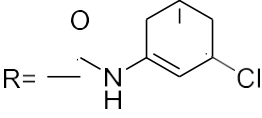
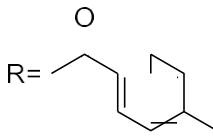
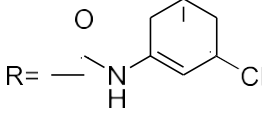
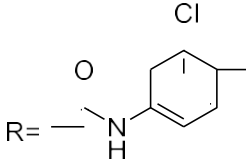


## INSTRUCTION MANUAL FOR CHIRALPAK® IA, IB, IB N-5, IC, ID, IE, IF, IG, IH, IJ, IK, IM, and IN

### <Reversed-Phase>

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

These columns can also be used in normal phase and SFC. Please refer to the corresponding instruction manual for details.

AMYLOSE-BASED  Immobilized on 5 µm silica gel		CELLULOSE-BASED  Immobilized on 5 µm silica gel	
<b>CHIRALPAK® IA</b> Amylose tris (3,5-dimethyl-phenylcarbamate) 	<b>CHIRALPAK® ID</b> Amylose tris (3-chloro-phenylcarbamate) 	<b>CHIRALPAK® IE</b> Amylose tris(3,5-dichlorophenylcarbamate) 	<b>CHIRALPAK® IF</b> Amylose tris(3-chloro-4-methylphenylcarbamate) 
<b>CHIRALPAK® IG</b> Amylose tris(3-chloro-5-methylphenylcarbamate) 	<b>CHIRALPAK® IH</b> Amylose tris[(S)-α-methylbenzylcarbamate] 		<b>CHIRALPAK® IN</b> Amylose tris(5-chloro-2-methylphenylcarbamate) 
<b>CHIRALPAK® IB</b> <b>CHIRALPAK® IB N-5</b> Cellulose tris(3,5-dimethylphenylcarbamate) 	<b>CHIRALPAK® IC</b> Cellulose tris(3,5-dichlorophenylcarbamate) 	<b>CHIRALPAK® IJ</b> Cellulose tris(4-methylbenzoate) 	
<b>CHIRALPAK® IK</b> Cellulose tris(3-chloro-5-methylphenylcarbamate) 		<b>CHIRALPAK® IM</b> Cellulose tris(3-chloro-4-methylphenylcarbamate) 	

#### Shipping Solvent:

1. Hexane/Isopropanol (IPA) = 90:10 (v/v) for analytical columns (2.1 mm i.d. x 150 mm, 4.6 mm i.d. x 150 and 250 mm), guard, and semi-prep and prep columns (only applicable to HPLC columns).
2. 100% Methanol for analytical (4.6 mm i.d. x 50 and 100 mm), guard, and semi-prep and prep columns (only applicable to SFC columns).

**\*Columns should be flushed with 100% Ethanol or IPA before connecting to a reversed phase system to ensure all Hexane is removed before introducing an aqueous mobile phase to the column.\***

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.\***

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

### Operating Instructions

	150 x 2.1 mm i.d. Analytical Column	50 x 4.6 mm i.d. 100 x 4.6 mm i.d. 150 x 4.6 mm i.d. 250 x 4.6 mm i.d. Analytical Columns	250 x 10 mm i.d.① 250 x 20 mm i.d.① 250 x 30 mm i.d.① 250 x 50 mm i.d.① Semi-Prep Columns
Guard	//	10 x 4.0 mm i.d. Guard Cartridge	20 x 10 mm i.d. 50 x 20 mm i.d. 50 x 30 mm i.d. Guard Column
Flow Rate Direction	As indicated on the column label		
Typical Flow Rate	0.1-0.5 ml/min	0.5-2.5 ml/min	5 ml/min (10 mm i.d.) 20 ml/min (20 mm i.d.) 42 ml/min (30 mm i.d.) 118 ml/min (50 mm i.d.)
Pressure Limitation②	Should be maintained < 300 Bar (4350 psi) for maximum column life <a href="#">Adapt flow rates to column size.</a>		
Temperature	0 to 40°C		
Column Fitting	Please contact <a href="#">Technical Support</a> for details		

① When using a semi-preparative column, it is highly recommended to discard at least the first 150 ml (for 250 x 10 mm i.d.) or 500 ml (for 250 x 20 mm i.d.) of eluent at the beginning of each preparative work.

② The column pressure is the total pressure minus the system pressure. At a given temperature, the column back pressure is linearly proportional to the flow rate.

### Switching Between RP and NP or SFC

To switch from reversed-phase to normal phase or SFC, and vice versa, the column should be carefully flushed with miscible solvent, such as ethanol or isopropanol. The column should be flushed in a similar manner with ethanol or isopropanol when initially received after purchase, before first used in reversed-phase, as it could contain a hexane/alcohol mixture.

It is highly recommended that the **regeneration procedure** described in the instruction manual for normal phase (located in the column care and maintenance section) be used when switching from reversed-phase to normal phase or SFC. Before applying this procedure, any traces of salts should be removed by flushing with a mobile phase that does not contain any salts / buffers, for example Water/ACN = 60/40, and then flushing with ethanol or isopropanol.

## A - Mobile Phases / For Both UV and Mass Detections

		ACIDIC (AMPHOTERIC) Compounds	NEUTRAL Compounds	BASIC Compounds ❶
<b>CHIRALPAK® IA</b> <b>CHIRALPAK® ID</b> <b>CHIRALPAK® IE</b> <b>CHIRALPAK® IF</b> <b>CHIRALPAK® IG</b> <b>CHIRALPAK® IH</b> <b>CHIRALPAK® IN</b>  <b>CHIRALPAK® IB</b> <b>CHIRALPAK® IB N-5</b> <b>CHIRALPAK® IC</b> <b>CHIRALPAK® IJ</b> <b>CHIRALPAK® IK</b> <b>CHIRALPAK® IM</b>	Aqueous Solution ❶	HCOOH aq. pH 2.0	Water	20 mM NH <sub>4</sub> HCO <sub>3</sub> aq. pH 9.0 adjusted with a basic additive ❶
	Organic Modifier ❷	ACN or MeOH or EtOH or IPA or THF		
	Typical Starting Conditions ❸	Aqueous solutions      60% ACN      40% ❹		

NOTE 1: If you cannot achieve sufficient resolution, try the complementary aqueous solutions

## B – Complementary Aqueous and Buffer Solutions / For UV Detection Only

		ACIDIC (AMPHOTERIC) Compounds	NEUTRAL Compounds	BASIC Compounds ❶
<b>CHIRALPAK® IA</b> <b>CHIRALPAK® ID</b> <b>CHIRALPAK® IE</b> <b>CHIRALPAK® IF</b> <b>CHIRALPAK® IG</b> <b>CHIRALPAK® IH</b> <b>CHIRALPAK® IN</b>  <b>CHIRALPAK® IB</b> <b>CHIRALPAK® IB N-5</b> <b>CHIRALPAK® IC</b> <b>CHIRALPAK® IJ</b> <b>CHIRALPAK® IK</b> <b>CHIRALPAK® IM</b>	Aqueous Solution ❶	50 mM Phosphate Buffer pH 2.0 OR H <sub>3</sub> PO <sub>4</sub> aq. pH 2.0 OR 100 mM KPF <sub>6</sub> (or NaPF <sub>6</sub> ) aq. pH 2.0 adjusted with H <sub>3</sub> PO <sub>4</sub>	Water	20 mM Borate Buffer pH 9.0 OR 20 mM Phosphate Buffer pH 8.0 ❷ OR 100 mM KPF <sub>6</sub> (or NaPF <sub>6</sub> ) aq.

NOTE 2: The concentration of all the buffering salt should be less than 500mM.

- ❶ Refer to **section C** for the preparation of an aqueous solution and choice of basic additives.
- ❷
  - ❑ It is recommended to use ACN to start the investigation.
  - ❑ The elution power of organic modifiers for these columns is in the descending order of ACN > EtOH > MeOH: 50%ACN ≈ 65-70%EtOH ≈ 75-80%MeOH.
  - ❑ The use of other organic solvents, **except THF**, has not been investigated and could be harmful to the columns.
  - ❑ The use of alcohols causes the back pressure to be significantly higher compared to ACN due to their higher viscosity in mixtures with water.
- ❸
  - ❑ Retention can be adjusted by changing the proportion of ACN. Retention may be very sensitive to the amount of ACN present in the mobile phase.
  - ❑ Lowering the column temperature may increase the retention time and the selectivity.
  - ❑ Increasing the column temperature and decreasing the flow rate may increase the resolution.
- ❹
  - ❑ To maximize column life, the use of a guard cartridge or guard column is essential when basic conditions are employed.
  - ❑ The use of strong basic conditions (> pH 9) must be avoided, as they are known to damage the silica gel matrix.

- ❑ When these columns are used at pH > 7, **the temperature should be maintained between 5°C and 25°C for maximum column life.**
- ⊗ High percentages of organic modifier in the mobile phase **may precipitate the buffering salt** from the solution, and lead to subsequent clogging of the column (refer to the table below).

Water / Organic Modifier	Buffer solution / Organic Modifier
90 / 10 to 0 / 100	90 / 10 to 15 / 85

- ⊗ Do not use a phosphate buffer for pH > 8. When pH 9 is necessary, the use of the ammonium bicarbonate solution or borate buffer is recommended for maximum column life.

## C – Buffer Preparation – Examples

- Preparation of pH 2 Phosphate buffer:  
**Solution A:** 50 mM potassium dihydrogenphosphate  
3.40g KH<sub>2</sub>PO<sub>4</sub> / FW 136.09, make up the volume to 500ml with HPLC grade water  
**Solution B:** phosphoric acid (H<sub>3</sub>PO<sub>4</sub> 85% by weight)  
Adjust the pH of solution A to a value of 2.0 using solution B.
- Preparation of pH 2 KPF<sub>6</sub> (NaPF<sub>6</sub>) solution:  
**Solution A:** 100 mM potassium (sodium) hexafluorophosphate  
9.20g KPF<sub>6</sub> / FW 184.06 or 8.40g NaPF<sub>6</sub> / FW 167.95, make up the volume to 500 ml with HPLC grade water  
**Solution B:** phosphoric acid (H<sub>3</sub>PO<sub>4</sub> 85% by weight)  
Adjust the pH of solution A to a value of 2.0 using solution B.
- Preparation of pH 9 Ammonium bicarbonate solution:  
**Solution A:** 20 mM ammonium bicarbonate  
0.78g NH<sub>4</sub>HCO<sub>3</sub> / FW 78.05, make up the volume to 500 ml with HPLC grade water  
**Solution B:** Basic additive such as diethylamine (DEA), triethylamine (TEA), ammonia (NH<sub>3</sub>) and soon.  
*\* DEA tends to give better peak shape than other bases.*  
Adjust the pH of solution A to a value of 9.0 using solution B.
- Preparation of pH 8 Phosphate buffer:  
**Solution A:** 20 mM potassium hydrogenophosphate  
1.74g of K<sub>2</sub>HPO<sub>4</sub> / FW 174.18, make up the volume to 500 ml with HPLC grade water  
**Solution B:** 20 mM potassium dihydrogenophosphate  
1.36g KH<sub>2</sub>PO<sub>4</sub> / FW 136.09, make up the volume to 500 ml with HPLC grade water.  
Adjust the pH of solution A to a value of 8.0 using solution B.
- Preparation of pH 9 Borate buffer:  
**Solution A:** 20 mM sodium tetraborate decahydrate  
3.81g of Na<sub>2</sub>B<sub>4</sub>O<sub>7</sub>·10H<sub>2</sub>O / FW 381.37, make up the volume to 500 ml with HPLC grade water  
**Solution B:** 20 mM boric acid  
0.62g H<sub>3</sub>BO<sub>3</sub> / FW 61.83, make up the volume to 500 ml with HPLC grade water  
Adjust the pH of solution A to a value of 9.0 using solution B.

## Column Care / Maintenance

- ❑ The use of a guard cartridge or guard column is highly recommended for maximum column life.
- ❑ Samples should preferably be dissolved in the mobile phase.
- ❑ The mobile phase and the sample solution should be filtered through a membrane filter of approximately 0.5 µm porosity to ensure that there is no precipitate before using.

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 Storage

- ❑ For column storage and/or switching to 100% organic solvent, any traces of salts should be removed by flushing the column with a mobile phase which does not contain any salts or buffers, for instance Water/ACN = 60/40 (v/v).
- ❑ 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

In India: [chiral@chiral.daicel.com](mailto:chiral@chiral.daicel.com) or call +91 84 1866 0700 & 703

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