

INSTRUCTION MANUAL FOR COLUMNS CHIRALPAK® IA-U / IB-U / IC-U

<Reverse Phase>

Please read this instruction sheet completely before using these columns

**These columns can also be used in normal phase mode.
Please refer to the corresponding instruction sheet for details.**

General recommendations

To switch from reversed phase mode to normal phase mode, and vice versa, column should be carefully flushed with miscible solvent.

It is highly recommended:

- to use a UHPLC system to preserve the best separation performance of the column.
- to apply the **regeneration procedure** described in the instruction sheet for normal phase mode. Before applying this protocol, any traces of salts should be removed by flushing with a mobile phase that does not contain any salts / buffers.
- to adjust the flow rate to uphold the column pressure < 700 bar.

Method Development / Reversed Phase

A - Mobile phases / For both UV and Mass detections

		ACIDIC (AMPHOTERIC) Compounds ④	NEUTRAL Compounds ④	BASIC Compounds ④
CHIRALPAK® IA-U CHIRALPAK® IB-U CHIRALPAK® IC-U	Aqueous solution ①	HCOOH aq. pH 2.0	Water	20mM NH ₄ HCO ₃ aq. pH 9.0 adjusted with a basic additive ①
	Organic modifier ②	CH ₃ CN or MeOH or EtOH or IPA or THF		
	Typical starting conditions ③	Aqueous solutions CH ₃ CN	60% 40% ⑤	

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

B - Complementary aqueous and buffer solutions / For UV detection

		ACIDIC (AMPHOTERIC) Compounds	NEUTRAL Compounds	BASIC Compounds ④
CHIRALPAK® IA-U CHIRALPAK® IB-U CHIRALPAK® IC-U	Aqueous solution ①	50mM Phosphate Buffer pH 2.0 OR H ₃ PO ₄ aq. pH 2.0 OR 100mM KPF ₆ (or NaPF ₆) aq. pH 2.0 adjusted with H ₃ PO ₄	Water	20mM Borate Buffer pH 9.0 OR 20mM Phosphate Buffer pH 8.0 ⑥ OR 100mM KPF ₆ (or NaPF ₆) aq.

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

- ❶ Refer to **section C** for preparation of aqueous solution and choice of basic additives.
- ❷ It is recommended to use CH₃CN to start the investigation
 - The elution power of organic modifiers for these columns is in the descending order of CH₃CN > EtOH > MeOH: 50%CH₃CN ≈ 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 CH₃CN due to their high viscosity in mixtures with water.
- ❸ Retention can be adjusted by changing the proportion of CH₃CN. Retention may be very sensitive to the amount of CH₃CN present into 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 the column life, it is essential to inject filtered clean sample solutions.**
 - 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.**
- ❺ High percentages of organic modifier in the mobile phase **may precipitate the buffering salt** from the solution, and lead to consequent 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

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

C – Buffer preparation – Examples

➤ Preparation of pH 2 Phosphate buffer:

Solution A: 50mM potassium dihydrogenphosphate
3.40g KH₂PO₄ / FW 136.09, make up the volume to 500ml with HPLC grade water

Solution B: phosphoric acid (H₃PO₄ 85% by weight)
Adjust the pH of solution A to a value of 2.0 using solution B.

➤ Preparation of pH 2 KPF₆ (NaPF₆) solution:

Solution A: 100mM potassium (sodium) hexafluorophosphate
9.20g KPF₆ / FW 184.06 or 8.40g NaPF₆ / FW 167.95, make up the volume to 500ml with HPLC grade water

Solution B: phosphoric acid (H₃PO₄ 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: 20mM ammonium bicarbonate
0.78g NH₄HCO₃ / FW 78.05, make up the volume to 500ml with HPLC grade water

Solution B: Basic additive such as diethylamine (DEA), triethylamine (TEA), ammonia (NH₃) and so on.
* *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: 20mM potassium hydrogenophosphate
1.74g of K₂HPO₄ / FW 174.18, make up the volume to 500ml with HPLC grade water

Solution B: 20mM potassium dihydrogenophosphate
1.36g KH₂PO₄ / FW 136.09, make up the volume to 500ml 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: 20mM sodium tetraborate decahydrate
3.81g of Na₂B₄O₇·10H₂O / FW 381.37, make up the volume to 500ml with HPLC grade water

Solution B: 20mM boric acid
0.62g H₃BO₃ / FW 61.83, make up the volume to 500ml with HPLC grade water

Adjust the pH of solution A to a value of 9.0 using solution B.

Column care / Maintenance

- Any traces of salts should be removed before column storage and /or before switching to 100% organic solvent (use Water/CH₃CN 60:40 (v/v) for instance)

For column care/maintenance, refer to Instruction Manual for normal phase

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

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