

Where design
meets
performance ...



IFC PAL Applications

Valve Selection and Flow Rates
Analytical and Prep Applications
Safety Issues

...CTC ANALYTICS

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❖ Valve Selection

- ✓ Bore Size 0.75 mm : Flow Rates 5 to approx.100 ml/min
- ✓ Bore Size 0.40 mm : Flow Rates 0.5 to approx. 20 ml/min
- ✓ Bore Size 0.25 mm : Flow Rates 10 to 500 µl/min
- ✓ Bore Size 0.15 mm : Flow Rates nl – Range to low µl Range ¹
- ✓ Bore Size 0.10 mm : Flow Rates nl – Range to low µl Range ²

1: no vertical port. Valve mounted on holder with 60 ° angle, used for injection

2: no vertical port, valve mounted on holder with 60 ° angle,
used for column switching

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❖ Tubing ID

- Analytical System:
OD 1/16 inch, ID 0.8 mm
- Preparative System:
OD 1/16 inch, ID 1.0 mm

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❖ **Syringe on Injector Side:**
5ml HPLC Syringe Needle Gauge 19
Gauge 19: OD 1.07 , ID 0.69 mm

❖ **Consequences:**

- ✓ Injection Unit: Needle Guides
- ✓ Valve: Needle Guide, Needle Seal (Teflon, blue color)
- ✓ FastWashStation: Glass Inserts

Details: see Service Note # 07/2004

IFC PAL Analytical Version

- ❖ Example of typical Application for Drug Discovery:
 - Biological Screening, MT-Plates with „hits“
 - Hit is checked if a single or several compounds are present. Chromatographic separation.
 - Flow rate: 400 ul/min, UV detection
 - Time based fraction collection, 6 sec/well
 - Collected into MT384
 - Collected Fraction are sent back to bio-screening

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IFC Preparative Version

- ❖ Example of Typical Application for Drug Discovery and Purification:
 - Purification or Chiral Separation of a new drug for further studies, like tox or clinical trials, mg to gram range
 - Flow Rates: 5 ml to 50 ml/min
 - Often normal phase separation
 - Column Bed pressure sensitive

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IFC Preparative Version

- ❖ Example of Typical Application for Drug Discovery and Purification continued:
 - Avoid any backpressure, tubing, Valve, Injection Syringe, SP-Syringe
 - Detection by UV-, Chiral- or MS-Detector
 - Peak-Detection Mode
 - Collection into Reagent-Glass-Tubes
 - Safety Measures have to be taken!!!
 - Solvent evaporation: various approaches

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❖ Safety Measures:

- ✓ Grounding
- ✓ Grounding
- ✓ Grounding

especially for IFC Prep-Version.....

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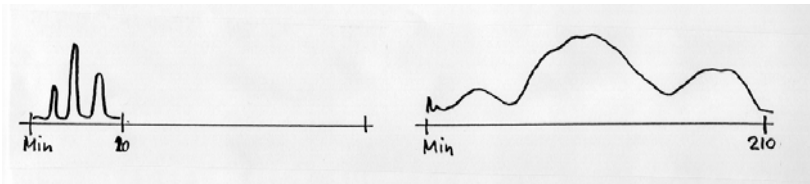
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❖ Sample Re-Injection

- Re-Injection Directly from Collection Tube
 - Needle Length
 - Layer of Solvent/Sample
- Evaporation of Solvent
 - Sample Stability

IFC PAL | Scale-up

❖ Scale-up Analytical to Preparative Scale:



Particles
Length
ID
Flow
Runtime
Total Solvent

10 μm
25 cm
4.6 mm
1 ml/min
20 min
20 ml

40 - 150 μm
65 cm
25 mm
20 ml/min
210 min
4200 ml

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Scaleable and Absolute Factors

- It is important to normalize all of the scaleable factors by the appropriate linear scale-up factor. Do **not** change any absolute factors.

- **Scaleable Factors**

- **Flow Rate**
- **Tubing Area**
- **Sample Injection Volume**
- **Detector Flow Cell Size**
- **Solvent Consumption**
- **Throughput**
- **Dead Volume**
- **Sample Loop Size**
- **Fraction Volumes**
-

- **Absolute Factors**

- **Packing Material / Column Length**
- **Back Pressure**
- **Sample Concentration**
- **Peak Height and Area**
- **Percent Recovery**
- **Purity**
- **Gradient Slope**
- **Run Time**
- **Column Performance**
- **Temperature**

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IFC PAL | Scale-up

- ✓ Make your first “preparative injection” on a small diameter “analytical” column.
Optimize the separation.
- ✓ Increase all of the scaleable elements of the system linearly.
Do not change any of the “absolute” elements.
- ✓ Perform the preparative separation.

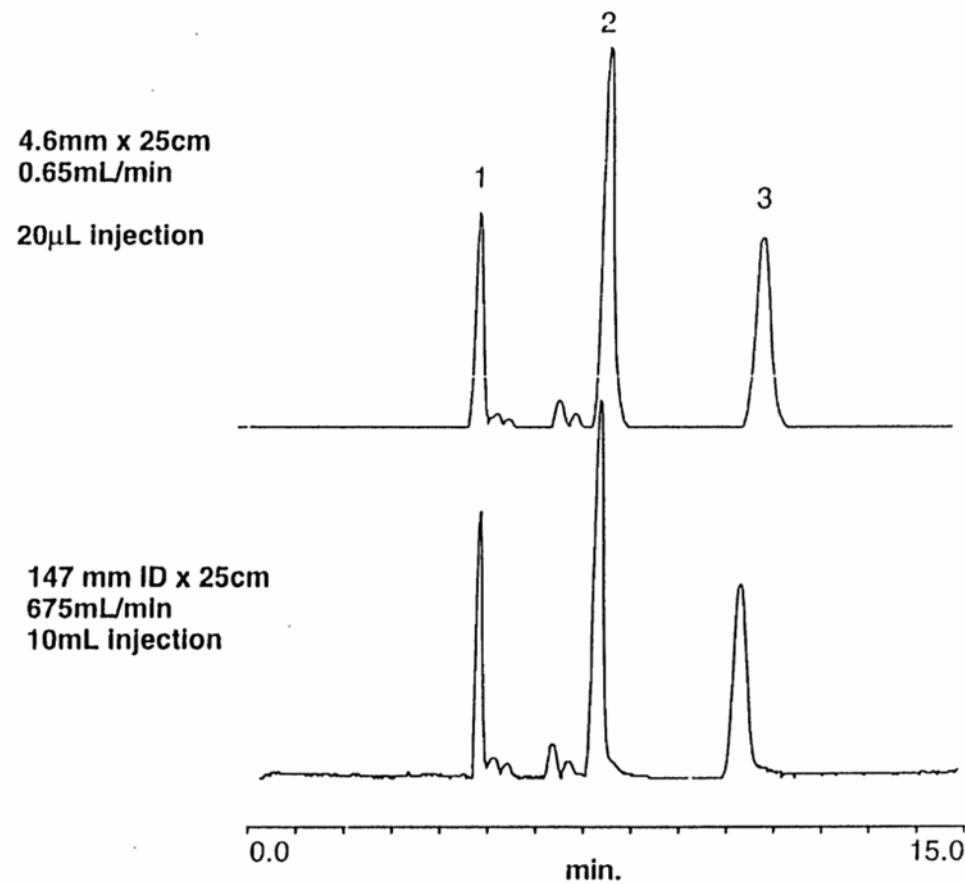
IFC PAL | Scale-up

Use A Linear Scale-Up Factor

❖ A linear scale-up factor is simply the ratio of the cross sectional areas of the analytical column and the intended preparative column.

$$\begin{aligned}\text{❖ Scale-Up Factor} &= P r^2_{\text{prep}} / P r^2_{\text{analytical}} \\ &= r^2_{\text{prep}} / r^2_{\text{analytical}}\end{aligned}$$

IFC PAL | Scale-up



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