

## N-to-C 4.0 and C-to-N 4.0 Protein Sequencer Methods

## **Technical Note**

HP 241 Protein Sequencer (N+C)

The N-to-C 4.0 and C-to-N 4.0 sequencer methods enable the switching between the N-terminal and C-terminal sequencing chemistry methods on the Hewlett-Packard column-based protein sequencer.

The N-to-C 4.0 method is run when switching from the N-terminal to the C-terminal sequencing chemistry. The column method pauses the column, while the flask method prepares a coordinated injection with the on-line HPLC to place the proper HPLC solvents in the HPLC binary pumps for the C-terminal chromatography. The C-to-N 4.0 method is run when switching from the C-terminal to the N-terminal sequencing chemistry. The column method washes the column, while the flask method, in addition to washing the flask, prepares a coordinated injection with the on-line HPLC to place the proper HPLC solvents in the HPLC binary pumps for the N-terminal chromatography. These methods are generally run with a Zitex strip in an empty RP/SAX biphasic column, but any column configuration is applicable. No column preparation is required with this method.

The methods are:

- N-to-C 4.0 (column)
- N-to-C 4.0 (flask)
- Column wash 4.0
- Flask wash 4.0

The sequence program controls all of the column and flask methods, and well as any cycle exception methods. The N-to-C 4.0 and C-to-N 4.0 sequencer methods do not use a exception method.

The Routine 4.0 sequencer methods require the bottle configuration shown. The		
Bottle Diagram X	reagents and solvents are purchased from Hewlett-Packard. These methods produce	
Couple	less that	i mu/cycle of ilquid waste.
SZA 1R R1 2R R2	S2A	Neat ethyl acetate
	18	Dinhenvlnhosphorylisothiocyanate (DPP-ITC) in toluene/hentane (23·27:50)
	R1	Phenylisothiocyanate (PITC) in hentane (3.97)
Extract Cleave	2R	Pyridine in ethylacetate (2:98)
SOS HEA SH SS HS	R2	Dijsopropylethylamine (DIEA) in 1-propanol/water (1:3:6)
	S2/3	Acetonitrile/toluene (23:77)
<b>R2A</b> Ocylamine in heptane (3:97)		
L1 PTH TH B4 55	3R	Potassium trimethylsilanolate (0.1M) in methanol/t-butanol (50:50)
	<b>S</b> 3	Acetonitrile/toluene (15:85)
	R3	Neat trifluoroacetic acid (TFA)
L3 Acetic acid in methanol/water (1:74:25)		
LCA LCB LCC	PTH	Mixture of PTH amino acids (10 pmol/100 µl) in acetonitrile with DPTU
		as a marker
	TH	Mixture of TH amino acids (50 pmol/100 µl) in acetonitrile
<b>R4</b> Trifluoroacetic acid (TFA) in water (1:3)		
	S5	Phosphate buffer (pH 2.9), 0.3% ion pairing reagent

#### **Method Details**

The steps for the N-to-C 4.0 and C-to-N 4.0 methods are described below. The steps indicate the ranges in volume that are appropriate for the various reagents/solvent deliveries. All sequencer methods are accessed by choosing the top menu item Edit/Method in the Protein Sequencer window. The sequencer column configuration used with the N-to-C 4.0 and C-to-N 4.0 methods consists of an empty reverse-phase (RP) sample column (top) mated with an empty strong anion exchange (SAX) column (bottom).

# Method (Column): Column wash 4.0

Total time: 38.8 min

All metering steps deliver the specified reagent or solvent to waste.

**Step 3:** The delivery of R2A should completely wet the membrane.

**Step 6, 9:** The volume of R2 delivered during this step should completely fill the column.

**Step 11, 42:** "Flush with S2/3 and S2A" flushes the delivery lines and should not wet the column

**Steps 13, 15, 16, 17, 18, 19, 20, 21:** The delivery of S2A should completely fill the column, followed by drying.

**Steps 23, 24, 36, 37:** The delivery of S3 should completely fill the column, followed by drying.

**Steps 26, 27, 33, 34:** The delivery of S2/3 should completely fill the column, followed by drying.

**Step 28, 30, 31:** The volume of R3 (approximately 1/2 column volume) delivered during this step should completely wet the membrane.

**Step 38:** After step 38, the volume of S5 delivered to the flask should be  $50 \mu l (+/-5 \mu l)$ 

Step	Description	Primary Time	Temperature
1.	Wash: Dry column DOWN	30.0	55
2.	Wash: Meter R2A	4.8	55
ζ.	Wash: Deliver R2A DOWN (closed)	20.0	55
Δ·	Wash: Dry column DOWN	30.0	55
5.	Wash: Meter R2 DOWN	120.0	55
6 <sup>.</sup>	Wash: Deliver R2 DOWN	30.0	55
7.	Wash: Dry column DOWN	30.0	55
8.	Wash: Meter R2 UP	120.0	55
g.	Wash: Deliver R2 LIP	30.0	55
10.	Wash: Dry column UP	30.0	55
11:	Wash: Flush with S2/3 and S2A	15.0	55
12:	Wash: Meter S2A UP	60.0	55
13:	Wash: Deliver and DRY UP (closed)	10.0	55
14:	Wash: Meter S2A DOWN	60.0	55
15:	Wash: Deliver and DRY DOWN (closed)	10.0	55
16:	Wash: Flush with S2A UP	14.0	55
17:	Wash: Flush with S2A DOWN	14.0	55
18:	Wash: Flush with-S2A UP	14.0	55
19:	Wash: Flush with S2A DOWN	14.0	55
20:	Wash: Flush with S2A DOWN to flask	14.0	55
21:	Wash: Flush with S2A UP to flask	14.0	55
22:	Wash: Dry column DOWN	30.0	55
23:	Wash: Flush with S3 UP	15.0	55
24:	Wash: Flush with S3 DOWN	15.0	55
25:	Wash: Dry column UP	30.0	55
26:	Wash: Flush with S2/3 DOWN	15.0	55
27:	Wash: Flush with S2/3 UP	15.0	55
28:	Wash: Flush with R3 UP	0.0	55
29:	Wash: Dry column UP	30.0	55
30:	Wash: Flush with R3 DOWN	30.0	55
31:	Wash: Flush with R3 DOWN to flask	30.0	55
32:	Wash: Dry column DOWN to flask	60.0	55
33:	Wash: Flush with S2/3 DOWN to flask	15.0	55
34:	Wash: Flush with S2/3 UP to flask	15.0	55
35:	Wash: Dry column DOWN	30.0	55
36:	Wash: Flush with S3 UP to flask	15.0	55
37:	Wash: Flush with S3 DOWN to flask	15.0	55
38:	!Wash: Deliver S5 to flask	7.0	55
39:	!Wash: Extract with S3 DOWN to flask	15.0	55
40:	!Wash: Evaporate flask	50.0	55
41:	Wash: Dry column DOWN	30.0	55
42:	Wash: Flush with S2/3 and S2A	15.0	55
43:	Wash: Dry column UP	60.0	55

#### Method (Column): N-to-C 4.0 Total time: 20.7 min

The column pauses during this method while the N-to-C 4.0 flask method runs

Step	Description	Primary Time	Temperature
1:	wait	20.0	30
2:	!wait system	20.0	30
3:	wait	600.0	30
4:	wait	600.0	30

#### Method (Flask): Flask wash 4.0 Total time: 17.0 min

**Step 4:** After Step 4, the volume of S5 in the flask should be 75-80 µl.

### **TEST Delivery protocol:**

To check the delivery volume of S5, copy step 4 to the clipboard and run from the clipboard. At the end of Step 4, remove the flask from the heating chamber. Carefully unscrew the flask cap and measure the delivered volume using a syringe. Be careful that some of the delivered volume is not retained on the tube sides or top of the flask.

**Step 7:** Before injection the injector loop should be filled, leaving the solvent visible in both the inlet and outlet lines of the injector loop. The time for Step 7 may have to be adjusted in order to optimize the delivery.

Step	Description	Primary Time	Temperature
1: W	/ash: Flush with L3	85.0	50
2: W	/ash: Mix	20.0	50
3: W	/ash: Empty flask	30.0	50
4: W	ash: Deliver S5 dripwise	12.0	50
5: Wash: Mix and equilibrate		15.0	50
6: Wash: Solubilize		20.0	50
7: W	ash: Fill loop and inject	2.0	50
8: W	ash: Empty flask	30.0	50
9:	Wash: Deliver R4 wash	40.0	50
10:	Wash: Empty flask	30.0	50
11:	Wash: Flush with L3	85.0	50
12:	Wash: Mix	20.0	50
13:	Wash: Empty flask	30.0	50
14:	Wash: Flush with L3	85.0	50
15:	Wash: Mix	20.0	50
16:	Wash: Empty flask	30.0	70
17:	Wash: Dry waste line	100.0	70
18:	Wash: Dry vent line	60.0	70
18:	Wash: Dry vent line	60.0	70



#### Method (Flask): N-to-C 4.0 Total time: 9.6 min

**Step 2:** After Step 2, the volume of S5 in the flask should be 75-80 µl.

#### TEST delivery protocol:

To check the delivery volume of S5, copy step 2 to the clipboard and run from the clipboard. At the end of Step 2, remove the flask from the heating chamber. Carefully unscrew the flask cap and measure the delivered volume using a syringe. Be careful that some of the delivered volume is not retained on the tube sides or top of the flask

Step	Description	Primary Time	Temperature
1:	Wash: Empty flask	30.0	50
2:	Wash: Deliver S5 dripwise	12.0	50
3:	Wash: Mix and equilibrate	15.0	50
4:	Wash: Solubilize	20.0	50
5:	Wash: Fill loop and inject	2.0	50
6:	Wash: Empty flask	30.0	50
7:	Wash: Flush with L3	85.0	50
8:	Wash: Mix	20.0	50
9:	Wash: Empty flask	30.0	50
10:	Wash: Dry waste line	100.0	50
11:	Wash: Dry vent line	60.0	50

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