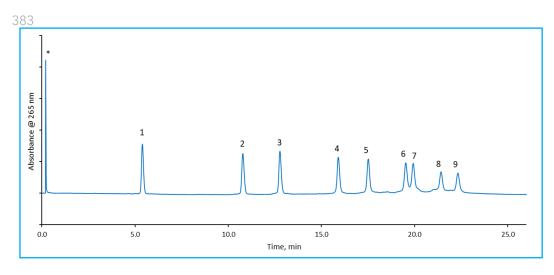


BIOPHARMACEUTICALS



Separation of Large ssDNA Oligomers on HALO® OLIGO C18



PEAK IDENTITIES

- 1. 20 mer
- 2. 30 mer
- 3. 40 mer
- 4. 50 mer
- 5. 60 mer
- 6. 70 mer
- 7. 80 mer
- 8. 90 mer
- 9. 100 mer
- * Tris HCI/EDTA

TEST CONDITIONS:

Column: HALO 120 Å OLIGO C18, 2.7 µm, 2.1 x 50 mm

Part Number: P2A62-402

Mobile Phase A: 100mM TEAA, pH 7

Mobile Phase B: Acetonitrile

Gradient:	Time	%B
	0.0	6.5
	30.0	11
	31.0	11
	31.1	6.5
	35.0	6.5

Flow Rate: 0.5 mL/min Back Pressure: 144 bar Temperature: 60 °C Injection: 2.0 µL

Sample Solvent: 10mM Tris HCI/1mM EDTA pH=8.0

Wavelength: PDA, 265 nm

Flow Cell: 1 µL Data Rate: 40 Hz

Response Time: 0.05 sec.

LC System: Shimadzu Nexera X2

The OLIGO C18 column with a 120 Å pore size separates a ladder of larger ssDNA nucleotides. Ranging from 20 to 100 bases, each oligonucleotide is separated under IPRP conditions. The oligonucleotide size that can be separated on the HALO® OLIGO C18 depends upon the requirements of the separation. While the OLIGO C18 column excels with smaller oligonucleotides like, antisense strands , it also has the ability to separate larger oligomers like this ssDNA ladder. This increases the diversity of samples that can be analyzed on the HALO® OLIGO C18 column.