

Extraction of Oligonucleotides From Plasma Samples Across Multiple Species Using OligoWorks™ SPE Microplate Kit

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Abstract

Oligonucleotide therapeutics are a key focus area for many drug developers today given their powerful ability to address disease biology at the level of gene transcription and translation with high target specificity and low toxicity. They are designed with specific types of modifications that enhance protein binding, half-life, and cellular uptake. Given their diversity, developing robust bioanalytical methods for oligonucleotide analysis can be quite challenging, and sample preparation is often its most challenging aspect due to the tight oligonucleotide-to-protein binding that occurs in biomatrices, often leading to low extraction recoveries and poor sensitivity. This work demonstrates use of the OligoWorks SPE Microplate Kit with RapiZyme™ Proteinase K Digestion to effectively extract oligonucleotides from plasma samples across multiple preclinical animal species including human. Using the starting protocol with no adjustments, oligonucleotide recoveries between 56–115% were obtained.

Experimental

Sample Preparation and Extraction

OligoWorks sample preparation and SPE extraction protocol

RapiZyme Proteinase K digestion sample pretreatment

Sample pretreatment
100 μ L biological sample, 20 μ L GuHCl (denaturation)
+ 10 μ L TCEP (Reduction) + 50 μ L RapiZyme Proteinase K (digestion)

Incubate 60 min, 55 $^{\circ}$ C, 600 rpm



OligoWorks SPE Microplate (2 mg/well)

Load
Pretreated Digested Plasma Sample (~180 μ L) to
SPE plate containing 180 μ L 50 mM NH_4OAc pH 5.5

Wash
Wash 1: 1 \times 200 μ L in 50 mM NH_4OAc pH 5.5
Wash 2: 1 \times 200 μ L in 10% MeOH



Elute
2 \times 25 μ L OligoWorks eluent
Dilute with 50 μ L Water (Optional)

Figure 1. Graphical representation of the OligoWorks SPE Microplate Kit (p/n: 186010614) sample preparation protocol, with plasma sample digestion pretreatment using RapiZyme Proteinase K, followed by WAX SPE using the OligoWorks SPE microplate. LC-MS analysis of plasma sample extracts was performed with an ACQUITY™ UPLC™ I-Class PLUS (FL) coupled to a Waters Xevo™ TQ-XS Tandem Quadrupole Mass Spectrometer using multiple-reaction monitoring mode (MRM). Chromatographic separation was achieved ACQUITY UPLC Premier Oligonucleotide C₁₈ Column, 130 Å, 1.7 μ m, 2.1 x 50 mm (p/n: 186009484).

Results

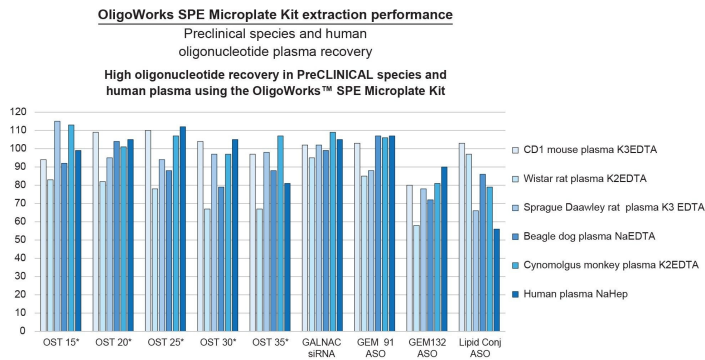


Figure 2. OligoWorks SPE Microplate Kit extraction performance (no internal standard correction) demonstrating high plasma recoveries (1-hour RapiZyme Proteinase K Digestion, 55 °C), for a diversity of oligonucleotides. *A 1:1 water dilution of digested plasma sample was applied prior to SPE loading. The 1:1 dilution minimized oligonucleotide loss (break-through) on SPE sample load, ensuring high SPE recovery.*

Ordering Information

| Description | P/N |
|--|-----------|
| OligoWorks SPE Microplate Kit | 186010614 |
| ACQUITY Premier Oligonucleotide C ₁₈ Column, 130Å, 1.7 µm 2.1 × 50 mm | 186009484 |
| QuanRecovery™ with MaxPeak, 700 µL plate | 186009184 |
| Polypropylene cap mat round well for 96-well | 186009452 |

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Xevo TQ-XS Triple Quadrupole Mass Spectrometer <

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RapiZyme Proteinase K Digestion Module <

<https://www.waters.com/nextgen/global/products/standards-and-reagents/rapizyme-proteinase-k-digestion-module.html>>

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