Extraction of Phosphatidylethanol (PEth) Species from Whole Blood Using ISOLUTE[®] SLE+ Prior to HPLC-MS/MS Analysis

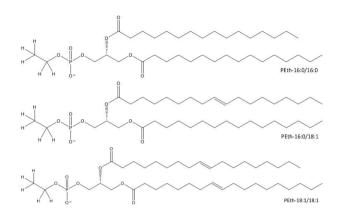


Figure 1. Chemical structures of three common PEth species.

Introduction

Phosphatidylethanol is an alcohol biomarker with a high degree of specificity; blood concentration of PEth correlates to the amount of alcohol consumed. This application note describes the extraction of 3 common species of PEth from whole blood using ISOLUTE® SLE+ supported liquid extraction prior to HPLC-MS/MS analysis.

ISOLUTE SLE+ supported liquid extraction fixed-well plates offer an efficient alternative to traditional liquid-liquid extraction (LLE) for bioanalytical sample preparation, providing high analyte recoveries, no emulsion formation, and significantly reduced sample preparation time.

This simple sample preparation procedure produces clean extracts, good recoveries with low RSD, and LOQ from 20 ng mL⁻¹. This method can be automated using Biotage[®] ExtraheraTM, see appendix for details.



Analytes

PEth is a group of phospholipids comprising a phosphoethanol head group and 2 fatty acid tails of varying length and saturation. This application note covers 3 commonly occurring species:

- » 1,2-dipalmitoyl-sn-glycero-3-phosphoethanol (PEth-16:0/16:0)
- » 1-palmitoyl-2-oleoyl-sn-glycero-3-phosphoethanol (PEth-16:0/18:1)
- » 1,2-dioleoyl-sn-glycero-3-phosphoethanol (PEth-18:1/18:1)
- » D_5 -1-palmitoyl-2-oleoyl-*sn*-glycero-3-phosphoethanol (D_5 -PEth-16:0/18:1) was used as an internal standard.

See Figure 1 for chemical structures.

Sample Preparation Procedure

Format

ISOLUTE[®] SLE+ 200 µL supported liquid extraction plate, part number 820-0200-P01.

Sample Pretreatment

To 20 μ L of whole blood, add 300 μ L of 6.25% (v/v) aqueous ammonium hydroxide in 30% aqueous methanol. Add an appropriate amount of internal standard separately or mix internal standard into the ammonium hydroxide pre-treatment solution prior to adding to sample. Mix thoroughly and allow to equilibrate.

Sample Loading

Load 140 μ L of the pre-treated whole blood into each well of the ISOLUTE SLE+ plate (equivalent to 8.75 μ L whole blood). Ensure the surface of the well frit is completely covered by the pre-treated sample. Using a Biotage[®] PRESSURE+96 Positive Pressure Manifold, apply 2–5 psi of pressure to load samples onto the sorbent. Wait 5 minutes for the sample to equilibrate on the sorbent.

Analyte Extraction

Apply 750 μ L of ethyl acetate and allow to flow under gravity for 5 minutes. Apply pressure (5–10 seconds) to remove any remaining extraction solvent.

Post Elution and Reconstitution

Dry the extract in a stream of air or nitrogen using a Biotage[®] SPE Dry 96 (40 °C at 60 L min⁻¹) or TurboVap 96 (40 °C at 1.0 bar). Reconstitute the extracts with 150 μ L mobile phase A:B (85:15 v/v). Mix thoroughly.



HPLC Conditions

Instrument

Waters Alliance 2795 HPLC with a 20 μ L loop.

Column

Agilent Poroshell 120 EC-C8 2.1 x 50 mm, 2.7 μ m analytical column; and Poroshell 120 EC-C8 2.1 x 5 mm, 2.7 μ m UHPLC guard column.

Mobile Phase

A: acetonitrile : 2 mM ammonium acetate (aq) 80:20 v/v;

B: propan-2-ol.

Flow Rate

0.25 mL min⁻¹.

Table 1. Gradient Conditions.

Time	% A	% В	Curve
0.00	85	15	1
0.40	85	15	1
3.50	15	85	6
3.51	5	95	6
5.50	5	95	6
6.00	85	15	6
8.90	85	15	1

Injection Volume

10 µL (partial loop)

Sample Temperature

12 °C

Column Temperature

Room temperature

Mass Spectrometry Conditions

Instrument

Waters Ultima Pt triple quadrupole mass spectrometer using electrospray ionization.

Capillary Voltage

3.2 kV

Desolvation Temperature

350 °C

Ion Source Temperature

100 °C

Negative ions acquired in multiple reaction monitoring (MRM) mode:

Table 2. MRM Conditions.

Compound	MRM Transition	Cone Voltage (V)	Collision Energy (eV)
PEth-16:0/16:0	675.4 > 255.2	35	31
PEth-16:0/18:1	701.4 > 281.2	35	34
PEth-18:1/18:1	727.5 > 281.2	35	32
D ₅ -PEth-16:0/18:1 (IS)	706.5 > 281.2	35	32

Results and Discussion

Chromatography

PEth species were chromatographed with a superficially porous C8 column using dilute ammonium acetate (aq) / acetonitrile and propan-2-ol. The overlaid extracted ion chromatogram (EIC) in Figure 2 demonstrates partial separation of the three PEth species was achieved. No inter-species contributions were observed in the EICs, demonstrating the HPLC-MS/MS method is capable of distinguishing between common PEth species.

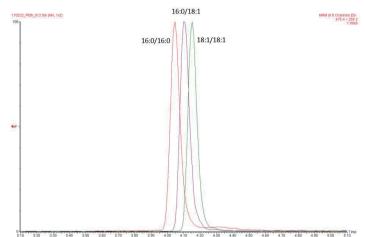


Figure 2. PEth Standards in Solvent Equivalent to 200 ng mL^1 in Whole Blood.

Recovery and Repeatability

Method performance was assessed by spiking whole blood with three PEth species plus internal standard at 200 ng mL⁻¹, equivalent to 1.75 ng when extracting 140 μ L of pre-treated whole blood. Recovery was determined relative to fortified blanks containing the same amount of extracted matrix. Recoveries for three PEth species were between 84% and 89% using an optimized ISOLUTE[®] SLE+ 200 μ L protocol (Figure 3). Extraction RSDs were between 5% and 6% (Figure 3). Representative EICs of 3 PEth species extracted from 20 μ L whole blood spiked at 200 ng mL⁻¹ are free from interfering peaks (Figure 4).

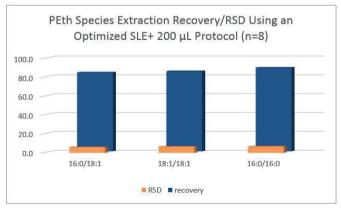


Figure 3. Percentage Recovery and RSD of PEth Species.



Compound name: 16:0/18:1 7014

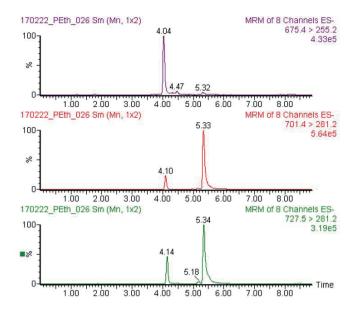


Figure 4. Extracted Ion Chromatograms of PEth Species (16:0/16:0, 16:0/18:1, 18:1/18:1) 20 μL Whole Blood Spiked at 200 ng mL $^{-1}$

Calibration Curves

Calibration curves were constructed by spiking whole blood from 20 ng mL⁻¹ to 20 μ g mL⁻¹ for each PEth species prior to extraction; the internal standard was spiked at 2 μ g mL⁻¹. Coefficient of determination (r²) values are demonstrated greater than 0.990 for each PEth species using the optimized extraction protocol. Representative curves are shown in Figure 5.

LOQ was estimated from the calibration curves where the signal/noise ratio was greater than 10:1. S/N values, estimated LOQ, coefficients of determination and precision RSDs are presented in Table 3.

 $\label{eq:table_to_state} \begin{array}{l} \textbf{Table 3.} \ \text{Lower Limits of Quantitation (LLOQ) using} \\ \text{optimized ISOLUTE}^* \ \text{SLE+ extraction protocol.} \end{array}$

PEth Species	LOQ, ng mL ⁻¹ (S/N)	LOQ, nmol L ⁻¹	r²	Precision % RSD (n=8)
16:0/16:0	20 (20)	29.5	0.995	6.1
16:0/18:1	20 (12)	28.4	0.994	6.6
18:1/18:1	20 (14)	27.4	0.994	6.8

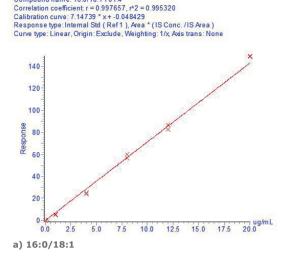
Additional Notes

Unless specified, all reagents and solvents are HPLC-grade.

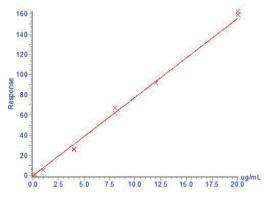
6.25% ammonium hydroxide (aq) / 30% methanol (aq): add 6.25 mL of concentrated ammonium hydroxide (28–30%) and 30 mL of methanol to 63.75 mL 18.2 M Ω cm water.

Aqueous mobile phase (A): dissolve 280 mg LC-MS grade ammonium acetate in 200 mL 18.2 M Ω cm water, add to 800 mL LC-MS grade acetonitrile and mix thoroughly.

Organic Mobile Phase (B): Use an appropriate volume of LC-MS grade propan-2-ol.

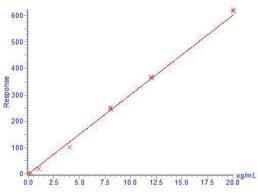


 $\label{eq:compound_name: 18:1/18:1 727.5} Correlation coefficient r = 0.997486, r^2 = 0.994978 Calibration curve: 7.78183 * x + 0.0543766 Response type: Internal Std (Ref1), Area * (IS Conc. /IS Area) Curve type: Linear, Origin: Exclude, Weighting: 1/x, Avis trans: None$





 $\label{eq:compound_name: 16:0/16:0 675.4} Correlation coefficient: r = 0.996908, r^2 = 0.993826 Calibration curve: 30.1208 * x + -0.438999 Response type: Internal Std (Ref 1), Area * (IS Conc. /IS Area) Curve type: Linear, Origin: Exclude, Weighting: 1/x, Avis trans: None$



c) 16:0/16:0

Figure 4. Representative PEth Species Calibration Curves from 20 ng mL $^{\rm 1}$ to 20 μg mL $^{\rm 1}$



Ordering Information

Part Number	Description	Quantity
820-0200-P01	ISOLUTE° SLE+ 200 μL Supported Liquid Extraction Plate	1
For Manual Process	ing	
PPM-96	Biotage [®] PRESSURE+ 96 Positive Pressure Manifold	1
For Automated Pro	cessing	
414001	Biotage® Extrahera	1
Rack and Reservoir	Options	
413991SP	Solvent Rack (25 mL)	1
414045SP	Solvent Reservoir (25 mL)	1
415560SP	Solvent Rack (100 mL)	1
414214SP	Solvent Reservoir (100 mL)	1
Evaporation		
SD-9600-DHS-EU	Biotage [®] SPE Dry 96 Sample Evaporator 220/240V	1
SD-9600-DHS-NA	Biotage® SPE Dry 96 Sample Evaporator 100/120V	1
C103199	TurboVap [®] LV Evaporator	1



Appendix Biotage[®] Extrahera[™] Settings

The method described in this application note was automated on the Biotage[®] Extrahera[™], using ISOLUTE[®] SLE+ 200 µL plates. Method performance was comparable to manual processing: recovery 73% to 77%, RSD 4.0% to 4.4%, r² 0.995 to 0.997. This appendix contains the software settings required to configure Extrahera to run this method.

Using this automated procedure, 96 samples can be processed in 22 minutes 33 secs.

An importable electronic copy of this method for Extrahera can be downloaded from www.biotage.com

Method Name:	TBD
Sample Plate/Rack:	2 mL Sample Plate, 96
Extraction Media:	PEth SLE 200





Settings

"Sample" Tab	
Sample Type:	PEth Sample
Starting Sample Volume (µL)	20
Reuse sample tips?	No
Method comment:	

- 1. If needed, centrifuge the sample plate briefly to collect the sample at the bottom of the well.
- 2. Prepare internal standard at an appropriate concentration in 6.25% ammonia 30% methanol. We recommend using Solvent Reservoir 6 in Accessory Position 6. An additional Solvent Rack and Solvent Reservoir will be needed**.

Alternatively, use a Solvent Reservoir from position 1 to 5. For this option, a larger volume of solvent + internal standard is needed and the Solvent Inlet Lines will need flushing after use.

** see ordering information on page 4 for suitable solvent rack and reservoir options.



Screenshot

thod name	100		SI	ample plate/rack		Extraction media	
PEth Whole B	lood			2 mL Sample Plate, 96	-	PEth SLE 200	-
etreatment	Sample	Pretreatment	Load	Elution			
On ad On tion	Number o 1 1 5 olvent 6,25%	• NH4OH 30% met				Pause after last step? No	Dispose solvent after each step? NC
	300 Mix numbe 0 Wait time (r of times Mix volume	+ (pL)				
	0						

Settings

0	
Pre-treatment	Activated
No. of steps	1
Pause after last step	No
Dispose tips after last step	No

	Solvent				
1	6.25% NH₄OH 30	% methano	l in water		
2					
3					
4					
		1	2	3	4
Volur	ne (µL)	300			
Mix n	umber of times	0			
		÷			
Mix v	olume (µL)	0			

tethod name	9	Sample plate/rack		Extraction media	
PEth Whole I	Blood	2 mL Sample Plat	e, 96 👻	PEth SLE 200	-
retreatment	Sample Pretreatmen	t Load Elution			
On	Volume (µL)	Air push time (s)	Walt time (mi	n)	
ad	140	2	5		
On	Premix? Numbe	r of times			
ution	Yes 3	÷			
On	Pause after each				
		in position			

Load	Activated
Pause after each load	No
Volume (µL)	140
Collect in position	D
Air Push Time (s)	2
Wait time (min)	5
Premix	Yes
Number of times	3



thod name			Sample plate/rack	Extraction media	
PEth Whole B	lood		2 mL Sample Plate, 96 🛛 👻	PEth SLE 200	-
etreatment	Sample Pro	etreatment Load	Elution		
On	Number of steps		Air push after last elution? Air push time Yes 10	(s)	Dispose solvent ti after each step? No
On	1 Solvent				NO
tion On	Ethyl Aceta	te 🔶 🗸			
	Volume (μL) 750	Collect in position			
	Wait time (min)	Advanced pressure settings			
	5 Repeat (number of	Edit			
	times)	step?			
	1004				

Elution	Activated
No. of steps	1
Air push after last elution	Yes
Air push time (s)	10
Dispose tips after each step	No

	Solvent				
1	Ethyl Acetate				
2	MTBE				
3					
4					
_		_	_	_	_
		1	2	3	4
Volu	me (µL)	750			
Colle	Collect in position				
Wait	time (min)	5			
Repe	at	1			
Paus	e	No			

'Advanced Settings'

Solvent Properties

	Solvent Description
1	6.25% NH4OH 30% methanol in water
2	Ethyl Acetate
3	
4	
5	
6	
7	
8	
9	
10	

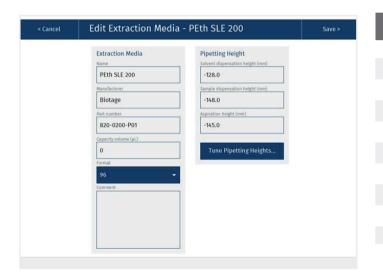


Solvent	1	2	3	4	5	6	7	8	9	10
Reservoir Type		Refi	lable				N	on Refillab	le	
Capacity	N/A	N/A								
Aspiration flow rate (mL/min)	10	10								
Dispense flow rate (mL/min)	20	10								
Lower air gap flow rate (mL/min)	20	20								
Lower air gap volume (µL)	5	5								
Upper air gap flow rate (mL/min)	120	120								
Upper air gap volume (µL)	100	100								
Upper air gap dispense pause	300	300								
Conditioning?	Yes	Yes								
Conditioning number of times	3	2								
Conditioning flow rate (mL/min)	20	10								
Conditioning volume (%)	100	100								
Aspirate post dispense	Yes	Yes								
Chlorinated	No	Yes								
Serial dispense	No	No								



Air Gap Lower air gap flow rate (mL/min) 20 Lower air gap volume (pL) 5 Upper air gap flow rate (mL/min) 10	Aspirate Aspirate post disperse? No
Lower air gap volume (pL) 5 Upper air gap flow rate (mL/min)	No
5 Upper air gap flow rate (mL/min)	
Contraction and Advancement of the second statement of the	
Upper air gap volume (µL)	
Upper air gap dispense pause (ms)	
0	
	500

"Sample" Screen	
Sample name	PEth Sample
Sample description	For 1 mL SLE columns
Aspiration flow rate (mL/min)	10
Dispense flow rate (mL/min)	5
Lower air gap flow rate (mL/min)	20
Lower air gap volume (µL)	5
Upper air gap flow rate (mL/min)	100
Upper air gap volume (µL)	500
Upper air gap dispense pause	0
Aspirate post dispense	No



"Extraction Media" Screen	
Name	PEth SLE 200
Manufacturer	Biotage
Part number	820-0200-P01
Capacity volume (µL)	0
Format	96
Solvent dispensation height (mm)	-128.0
Sample dispensation height (mm)	-148.0
Aspiration height (mm)	-145.0
Comment	

Sample Plate/Rack	Pipetting Height	
2 mL Sample Plate, 96	Aspiration height (mm) -162.0	
Capacity volume (µL)	Pretreatment dispensation height (mm)	
1800	-128.0	
Format		
96 👻	Tune Pipetting Heights	

"Sample Plate/Rack" Screen	
Name	2 mL Sample Plate, 96
Capacity volume (µL)	1800
Format	96
Aspiration height (mm)	-162.0
Pre-treatment dispensation height (mm)	-128.0



Pipette Tip	
1000 µL Biotage tip	
Manufacturer	
Biotage	
Part number	
414141	
Capacity (pl.)	
1000	
Length (mm)	
95	

"Pipette tip" Screen	
Name	1000 µL Biotage Tip
Manufacturer	Biotage
Part number	414141
Capacity (µL)	1000
Length (mm)	95

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