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## Development of polar lipid profiling method by supercritical fluid chromatography/mass spectrometry

Data No. LL007-0000

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**Data source** : Poster  
**Year** : 2011

### Conditions

**Column** : Inertsil ODS-EP (5 $\mu$ m, 250 x 4.6 mm I.D.)  
**Column Cat. No.** : 5020-02646  
**Mobile Phase** : Carbon dioxide (CO<sub>2</sub>, 99.9% grade)  
**Modifier** : Methanol with 0.1% (w/w) HCOONH<sub>4</sub>  
**Flow rate** : 3 mL/min  
**Oven temperature** : 37 °C  
**Back pressure** : 10 MPa

**Sample** : Polar lipids  
**Analyte** : table below

**Detection** : SFC/MS/MS  
**Ionization method** : Electrospray ionization  
**Ion mode** : positive  
**Capillary voltage** : 3.00 kV  
**Desolvation temperature** : 450 °C  
**Desolvation gas flow** : 800 L/hr  
**Cone gas flow** : 60 L/hr  
**Collision gas flow** : 13.2 mL/hr  
**MS collision energy** : 6 V  
**Source temperature** : 150 °C  
**Extractor voltage** : 3 V

### Optimized MS/MS method by TMS derivatization

Polar lipids	SRM transitions	CV	CE	Number of adducted TMS
Phosphatidylglycerol (PG)	[M+H] <sup>+</sup> > [M-315]	20	30	2
Phosphatidic acid (PA)	[M+H] <sup>+</sup> > [M-169]	25	25	1
Phosphatidylinositol (PI)	[M+H] <sup>+</sup> > [M-619]	35	30	5
Lysophosphatidylcholine (LPC)	[M+H] <sup>+</sup> > 184	45	35	1
Lysophosphatidylethanolamine (LPE)	[M+H] <sup>+</sup> > [M-140]	30	15	1
Lysophosphatidylglycerol (LPG)	[M+H] <sup>+</sup> > [M-315]	30	20	3
Lysophosphatidic acid (LPA)	[M+H] <sup>+</sup> > [M-169]	30	15	2
Lysophosphatidylinositol (LPI)	[M+H] <sup>+</sup> > [M-547]	45	25	5
Sphingomyeline (SM)	[M+H] <sup>+</sup> > 184	45	30	1
Sphingosine-1-phosphate (S1P)	[M+H] <sup>+</sup> > 264	25	20	2

CV: Cone voltage, CE: MS/MS collision energy

### Profiling of 10 polar lipid standards

