

# Application Data Sheet

## No.6

### GCMS

Gas Chromatograph Mass Spectrometer

## Analysis of Adipose Fatty Acids in Blood Utilizing a GC × GC-MS System

When fatty acids are analyzed with a GC-MS, an enormous number of components are detected due to differences in the number of double bonds and isomers of each component. It is impossible to completely separate these using conventional GC/MS analysis.

A GCMS-QP2010 Ultra, equipped with a GC × GC modulator, is capable of separating and detecting components that cannot be separated with a conventional GC-MS.

This application datasheet investigates the separation of the C18 fatty acids: oleic acid and gamma-linolenic acid. These substances could not be separated using only a 1-dimensional column, but could be separated 2-dimensionally using a GC × GC-MS system.

#### Analysis Conditions

GC × GC modulator	: ZX1-GC × GC modulator		
GC-MS	: GCMS-QP2010 Ultra		
[GC × GC]			
Column	: 1 <sup>st</sup> DB-5ms (30 mL × 0.25 mm I.D., 0.25 μm)	[MS]	
	: 2 <sup>nd</sup> BPX50 (2.5 mL × 0.1 mm I.D., 0.1 μm)	Interface temperature	: 240°C
Injection quantity	: 1.0 μL	Ion source temperature	: 200°C
Injection mode	: Split (split ratio 100)	Solvent elution time	: 15.5 min
Vaporization chamber temperature	: 250°C	Data sampling time	: 16 to 80 min
Column oven temperature	: 40 °C (2 min) -> (30 °C/min) -> 160 °C (2 °C/min)	Measurement mode	: Scan
	: -> 300 °C (5 min)	Mass range	: <i>m/z</i> 45-330
Control mode	: Constant pressure (150 kPa)	Scan speed	: 20,000 <i>u</i> /sec
Modulation time	: 8 sec		
Hot pulse time	: 0.5 sec (325°C)		

