## **Determination of Purity of Titanocene Dichloride**

## 1 Method Summary

The sample was dissolved in acetonitrile and tested with acetonitrile and water as the mobile phase. The stainless-steel column with C18 as the filler and variable ultraviolet detector are used to separate and determine the purity of Titanocene Dichloride with area normalization.

## 2 Reagents and Solutions

- 2.1 Acetonitrile: chromatographically pure
- 2.2 New steamed double distilled water

#### 3 Instruments

- 3.1 High pressure liquid chromatography; variable UV wavelength detector; chromatographic data processor or chromatography workstation
- 3.2 Column: 250mm × 4.6mm (i.d) stainless steel column, filled with ODS (C18) filler, particle size 10μm
- 3.3 Micro Injector: 50 µL

#### 4 Operating Conditions

- 4.1 Column temperature: room temperature
- 4.2 Mobile phase: acetonitrile + water = 70+30
- 4.3 Flow rate: 1mL/min
- 4.4 Detection wavelength: 220nm
- 4.5 Injection volume: 20μL
- 4.6 Retention time: About 2.16min

The above operating parameters are typical, and the given parameters can be appropriately adjusted according to the characteristics of different instruments to obtain the best results.

#### 5 Determination Steps

Approximately 0.5 g (accurate to 0.0002 g) of the sample was weighed into a 500 mL volumetric flask, dissolved in acetonitrile, and diluted to the mark with a mobile phase. Under the above operating conditions, after the baseline of the instrument was stabilized, 20  $\mu$ L of the sample solution was injected and determine the purity by area normalization.

# 6. Allowable Difference

| The difference between two parallel determination results shall not be greater than | n 1.0% | greater than | not be | esults shall | parallel determination | vo paralle | between tw | difference | The |
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