# **AccuTOF**<sup>™</sup> Introduction of Dual ESI and Corona ESI Ion Sources

We have developed two new ion sources, the Dual ESI and Corona ESI, for the AccuTOF LC-TOF MS system that was introduced in the spring of 2002. The Dual ESI (Figure 1) provides two ESI (electrospray ionization) sprayers. This permits the introduction of a reference compound without disturbing or suppressing the spray of the analyte. In addition, the dual-probe system can be useful for high-throughput analysis.

The Corona ESI ion source (Figure 2) is equipped with both an ESI sprayer and a coronadischarge electrode. The combination of ESI and APCI (atmospheric pressure chemical ionization) in a single source is useful for rapid analysis of unknown elements, and for improved efficiency in determining optimal analysis conditions. In addition, the orthogonal spray ion sources feature long-term stability and easy maintenance. For further details about the orthogonal-spray API source, refer to applications note MS-021024B.

## Secondary Sprayer



Figure1. Dual ESI Ion Source



**ESI Sprayer** 

Electrode

Figure 2. Corona ESI Ion Source

## **Exact Mass Measurement Using Dual ESI Ion Source**

Exact mass measurement using two sprayers is one of the benefits of the Dual ESI Ion Source. We conducted an exact mass measurement of rhodamine as a test sample, which was introduced from the main sprayer. Reserpine was introduced from the secondary sprayer as standard reference (drift correction or "lock mass") for mass calibration.

Table 1 shows the ion source conditions. Measurement was conducted under three different conditions, by switching on and off the nebulizing gas. Condition 1 is used for the measurement of rhodamine by ionization using the main sprayer. Condition 2 is used for the measurement of reserpine (the drift-correction standard) by the secondary sprayer, and condition 3 is used for obtaining spectra of both, by using two sprayers simultaneously.

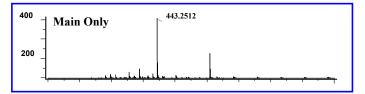
The results are shown in Figure 3. The measured value of rhodamine by using reserpine as a drift correction was 443.2323, and the deviation from the theoretical value was 1.13 millimass unit.

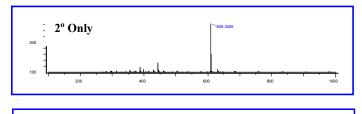
www.jeol.com/ms/accutof.html ms@jeol.com October 2002



	Main Sprayer On	$2^{\circ}$ Sprayer On	Both On
Main needle potential	2000 V	2000 V	2000 V
Nebulizing gas 1	ON	OFF	ON
2 <sup>°</sup> needle potential	2000 V	2000 V	2000 V
Nebulizing gas 2	OFF	ON	ON

Table 1.





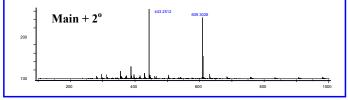


Figure 3. Rhodamine + Reservine

Ion Source Condition (ESI Mode)			
Needle Voltage	2000V		
Corona Electrode	0V		
Orifice Voltage	80V		
Ring Lens	10V		
Ion Source Condition (APCI Mode)			
Needle Voltage	0V		
Corona Electrode	5000V		
Orifice 1 Voltage	80V		
Ring Lens Voltage	10V		
Sample Introduction Condition			
Introduction Method	Infusion		
Flow Rate	0.2mL/min		

Table 3. Measurement condition of hydrocortisone

### Measurement Using Corona ESI Ion Source

Hydrocortisone is measured by using the Corona ESI Ion Source. Table 3 shows the measurement condition and Figure 4 shows the measured spectra. The sodium adduct is observed in ESI mode, and a proton adduct is observed in APCI mode as shown above.

### Summary

As mentioned above, the newly-developed Dual ESI ion source provides an easier exact mass measurement with excellent mass accuracy. In addition, the Corona ESI enables acquisition of information of ESI and APCI with a single ion source, and is therefore an extremely useful tool for rapid analysis of unknowns and for efficient optimization of analysis conditions.

Measured Value(m/z)	443.2323	
Theoretical Value(m/z)	443.2335	
Difference	-1.13 mmu	
	-2.55 ppm	

Table 2. Accurate mass measurement result

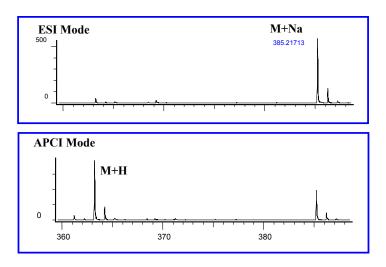


Figure 4. ESI Spectra and APCI Spectra of Hydrocortisone by ESI