SHIMADZU APPLICATION NEWS

SPECTROPHOTOMETRIC ANALYSIS



Measurement by Flame Micro Sampling Method

Introduction

The AA-6300 combined with an autosampler ennables the automated flame micro sampling method. In the flame micro sampling method, flame atomic absorption analysis is conducted with small sample quantities (2 - 100μ L), while in the conventilnal flame method (hereafter "flame continuous suction method"), the sample is continuously aspirated so rather large amount of sample is necessary.

Table 1 shows the comparison of the flame micro sampling method with the flame continuous suction method. The flame micro sampling method has sevral advantages over the flame continuous suction method: analysis is possible with a small amount of sample, and when an autosampler is used, automatic dilution of the sample and automatic addition of interference inhibitor are possible. Moreover, since only a small amount of sample is introduced, the flame micro sampling method is effective for analysis of high matrix samples which may cause clogging of the burner by the flame continuous suction method.

We introduce here an example of analysis of Na and Mg in the standard river water sample, using the flame micro sampling method. (The standard riverwater; JAC0031, unspiked, cetified values for Na;4.2±0.1mg/L and Mg;2.83±0.06mg/L, provided by the Japan Society of Analytical Chemistry)

Table 1 Comaprison of flame continuous suction method and flame micro sampling method

| | Measure-ment time | Volume of Sample Consumption (per analysis) | Sensitivity (Absorbance) | Coefficient of Variation (%) (Simple repetition) | Auto dilution | |
|---------------------------|-------------------|--|-------------------------------|---|-----------------------------|--|
| Continuous Suction method | 10 – 20sec | 1 – 2mL | - | Approx. 1% | Not possible | |
| Micro sampling method | Approx. 100sec | 2 – 100µL | Approx. 80% of suction method | 2 – 3% | Possible (50 times max.) | |

The maximum dilution factor refers to maximum value when the mixing port is not used (2µL/100µL).

Instrument and Analytical Conditions

Table 2 indicates the instrument and main analytical conditions. For both elements, the conditions were set so that if the measured value exceeded the 1mg/L

upper limit concentration of the calibration curve, the sample amount injected would be automatically decreased and the sample reanalyzed.

Table 2 Instrument and Analytical Conditions

| Instrument | Atomic Absorption Spectrophotometer AA-6300 | | | | | |
|--|---|--------------------|--|--|--|--|
| Autosampler | ASC-6100F, ASK-6100, ASK-6300, Micro sampling kit | | | | | |
| Measurement element | Na | Mg | | | | |
| Measurement wavelength | 589.0nm | 285.2nm | | | | |
| Ignition mode | NON-BGC | BGC-D ₂ | | | | |
| Flame type | Air / acetylene | | | | | |
| Burner angle | 0° | | | | | |
| Sampling time | 30 sec. | | | | | |
| Interference inhibitor | None | Lanthanum (La) | | | | |
| Standard solution concentration (mg/L) | 0.2, 0.5, 1.0 | | | | | |
| Max. conc. (mg/L) for auto dilution/reanalysis | 1.0 | | | | | |

Measurement Results

The measurement results for Na and Mg are shown in Table 3 and 4, respectively. Measurements were repeated twice for the standard sample and three times for the actual sample. The calibration curves were generated for both Na and Mg by injecting different volumes of 1mg/L solutions using an autosampler. For the Mg, lanthanum was also added as the interference inhibitor using the autosampler. In the analysis of the actual samples, since the first measurement value exceeded the set upper limit concentration, dilution and reanalysis were automatically conducted with smaller volumes of samples. The obtained values agreed well with certified values for both elements.

Table 3 Measurement result for sodium (Na)

| Operation | Sample ID | Exclude | Set concentration (mg/L) | Concentration (mg/L) | Abs. | Position | Sample amount (µL) | Diluent (μL) | Total volume (μL) | Auto dilution | Actual concentration (mg/L) | CV (%) |
|-----------|--------------|---------|-----------------------------|-------------------------|--------|----------|-----------------------|-----------------|----------------------|------------------|--------------------------------|--------|
| STD-AV | 0.2mg/L | | 0.2 | | 0.1184 | R2 | 20 | 80 | 100 | | | 1.31 |
| STD-AV | 0.5mg/L | | 0.5 | | 0.2954 | R2 | 50 | 50 | 100 | | | 0.55 |
| STD-AV | 1.0mg/L | | 1.0 | | 0.5652 | R2 | 100 | 0 | 100 | | | 1.58 |
| UNK1 | JAC0031 | X | | 3.3821 | 1.8933 | 2 | 100 | 0 | 100 | | 3.38 | |
| UNK2-1 | JAC0031 | | | 0.5972 | 0.3433 | 2 | 14 | 86 | 100 | 7.143 | 4.27 | |
| UNK2-2 | JAC0031 | | | 0.6121 | 0.3516 | 2 | 14 | 86 | 100 | 7.143 | 4.37 | |
| UNK2-3 | JAC0031 | | | 0.6074 | 0.3490 | 2 | 14 | 86 | 100 | 7.143 | 4.34 | |
| UNK2-AV | JAC0031 | | | 0.6056 | 0.3480 | 2 | 14 | 86 | 100 | 7.143 | 4.33 | 1.22 |

Table 4 Measurement result for magnesium (Mg)

| Operation | Sample ID | Exclude | Set concentration (mg/L) | Concentration (mg/L) | Abs. | Position | Sample amount (µL) | Diluent (μL) | La5% (μL) | Total volume (µL) | Auto dilution | Actual concentration (mg/L) | CV (%) |
|-----------|--------------|---------|-----------------------------|-------------------------|--------|----------|-----------------------|-----------------|--------------|----------------------|------------------|--------------------------------|--------|
| STD-AV | 0.2mg/L | | 0.2 | | 0.1946 | R2 | 16 | 64 | 20 | 100 | | | 3.05 |
| STD-AV | 0.5mg/L | | 0.5 | | 0.4750 | R2 | 40 | 40 | 20 | 100 | | | 0.61 |
| STD-AV | 1.0mg/L | | 1.0 | | 0.9388 | R2 | 80 | 0 | 20 | 100 | | | 1.25 |
| UNK1 | JAC0031 | X | | 2.0887 | 1.9516 | 2 | 80 | 0 | 20 | 100 | | 2.09 | |
| UNK2-1 | JAC0031 | | | 0.7446 | 0.7016 | 2 | 20 | 60 | 20 | 100 | 4 | 2.98 | |
| UNK2-2 | JAC0031 | | | 0.7272 | 0.6854 | 2 | 20 | 60 | 20 | 100 | 4 | 2.91 | |
| UNK2-3 | JAC0031 | | | 0.7377 | 0.6952 | 2 | 20 | 60 | 20 | 100 | 4 | 2.95 | |
| UNK2-AV | JAC0031 | | | 0.7365 | 0.6941 | 2 | 20 | 60 | 20 | 100 | 4 | 2.95 | 1.18 |



Fig.1 Examples of peak profiles for sodium (Na)



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