



Application Note AN-H-127

Thermometric analysis of aluminum by back-titration

Fast and robust technique for aluminum determination

A thermometric complexometric titration procedure has been adapted to the determination of aluminum in solutions, where the direct titration with fluoride is not practicable because of the interference of silica (e.g., from digestion of clays, zeolites, or other alumino-silicate-containing substances).

The new method involves the use of a thermometric indicator (hydrogen peroxide) to give a sharp temperature change at the endpoint. When all the excess EDTA has reacted with the copper(II) titrant, the first trace of free Cu^{2+} ions causes the H_2O_2 to

decompose very rapidly, causing a sudden increase in the temperature of the solution. The heat of reaction ΔH_f for $\text{H}_2\text{O}_2 \rightarrow \text{H}_2\text{O} + [\text{O}]$ is approximately -98 kJ/mol, or twice the heat created during the reaction of a strong acid with a strong base. This makes the technique very robust.

Additionally thermometric titrations have very short titration durations as the titrant is added continuously while monitoring the temperature. Results are usually obtained within 2–3 minutes.

SAMPLE AND SAMPLE PREPARATION

Aluminum sulfate and potassium alum salts are used as samples. To an Erlenmeyer flask containing the aluminum salts, EDTA solution in excess and ammonia

solution is added. Then the obtained solution is stirred for five minutes while boiling to facilitate the complexation reaction between aluminum and EDTA.

EXPERIMENTAL

After allowing to cool down to room temperature, an aliquot of the solution is used for titration. Ammonia buffer and hydrogen peroxide are added subsequently. The excess of EDTA is back titrated with Cu^{2+} solution.

The thermometric titration is carried out automatically with the *tiamo*TM software in combination with an 859 Titrotherm and a Thermoprobe.



Figure 1. 859 Titrotherm equipped with a Thermoprobe and tiamo. Example setup for the analysis of aluminum.

RESULTS

The analysis of aluminum is very reproducible. Relative standard deviations < 0.3% are obtained with this

method.

Table 1. Results of the aluminum determination in aluminum sulfate ($\text{Al}_2(\text{SO}_4)_3 \cdot 16 \text{H}_2\text{O}$) and potassium alum ($\text{AlK}(\text{SO}_4)_2 \cdot 12 \text{H}_2\text{O}$).

	Assay Al in $\text{Al}_2(\text{SO}_4)_3 \cdot 16 \text{H}_2\text{O}$ / %	Assay Al in $\text{AlK}(\text{SO}_4)_2 \cdot 12 \text{H}_2\text{O}$ / %
n	8	10
Mean	7.87	5.11
SD(abs)	0.02	0.01
SD(rel)	0.25	0.20

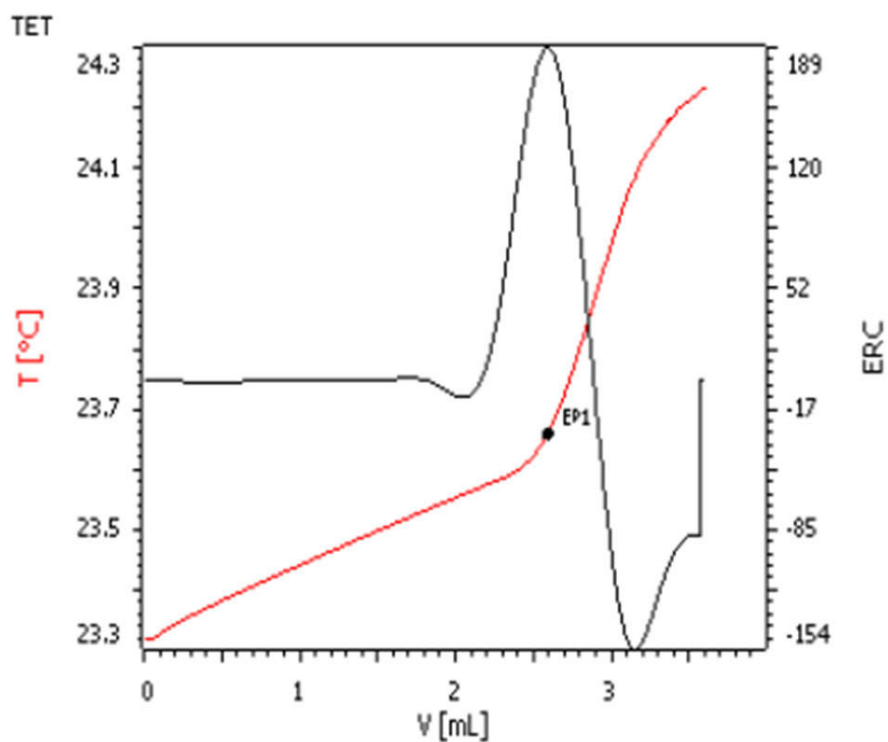


Figure 2. Example curve of thermometric determination of aluminum in aluminum sulfate.

CONCLUSION

This application shows a fast alternative method to the potentiometric titration of aluminum which also can be used in the presence of silicates.

Thermometric titration is a very fast and maintenance-

free technique, which leads to reliable and precise results. The addition of peroxide enhances the reaction enthalpy therefore additionally increases the reproducibility.

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