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HIRANUMA APPLICATION DATA		Karl Fischer Titrator	Data No.	KF18	Nov.29, 2018
Water contents	Validation test with water standard KF Coulometry – Heat evaporation method				

1. Abstract

A heat evaporation method using Karl Fischer titrator combined with solid evaporator is suitable for insoluble sample in anode solution, such as plastics and inorganic compounds. The measurement of with the heat evaporation method is carried out by heating the sample while a carrier gas is blown into the evaporation chamber and introducing the evaporated moisture together with the carrier gas into the electrolysis cell.

This application introduces an example of measurement of solid water standard for validation of titrator system with solid evaporator.

2. Apparatus and Reagents

(1) Apparatus for heat evaporation method

- Titrator : HIRANUMA Karl Fischer Coulometric titrator AQ-series or MOICO-A19
- Evaporator : Hiranuma Solid Evaporator EV-2000
- Electrolytic cell : Standard Cell
Fritless Cell (No cathode solution required)

(2) Reagents

- Anode solution : HYDRANAL Coulomat AG
- Cathode solution : HYDRANAL Coulomat CG
- Dehydrated methanol : HYDRANAL Methanol
- Carrier gas : Nitrogen gas (Dew Point : -65 °C or less)

(3) Water standard material

(i) HYDRANAL- Standard sodium tartrate dihydrate (Honeywell)

CoA : 15.64 %(*1), Permissible range of recovery rate : ± 5 %(*2), Heat temperature : 160 °C

(ii) HYDRANAL- Water Standard KF-Oven, 140-160°C (Honeywell)

CoA : 5.04 %(*1), Permissible range of recovery rate : ± 4 %, Heat temperature : 140 – 160 °C

(iii) Water Standard Oven 1 % (Merck)

CoA : 0.99 %(*1), Permissible range of recovery rate : ± 3 %, Heat temperature : 140 – 400 °C

(*1) CoA values described in the test report are different with each production lot.

(*2) Permissible range of standard material (i) is set to our own criterion for inspection of evaporator, not described on the document of this standard.

3. Procedure

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(3) Flow N_2 carrier gas with flow rate 200 ml/min.

(4) Set the formed aluminum foil into the evaporation chamber of EV-2000 (Fig.3.2).

(5) Connect electrolytic cell and evaporation chamber with tube (Fig.3.3).

(6) Blanking become stable and then start pre-heating at set temperature.

(7) After the pre-heating procedure finished, confirm blanking stable with carrier gas flowing.

(8) Make formed aluminum foil to use balance dish.

(9) Weigh the sample on the aluminum foil. Weighing amounts are approximately 0.05 g for standard material (i), 0.1 g for standard material (ii) and 0.2 g for standard material (iii).

(10) Put the sample with aluminum foil into evaporation chamber from stopper of the chamber.

(11) Start titration. Measurement parameter is shown in Table 4.1.

(12) Blank measurement is also performed with same procedure except for sample addition.

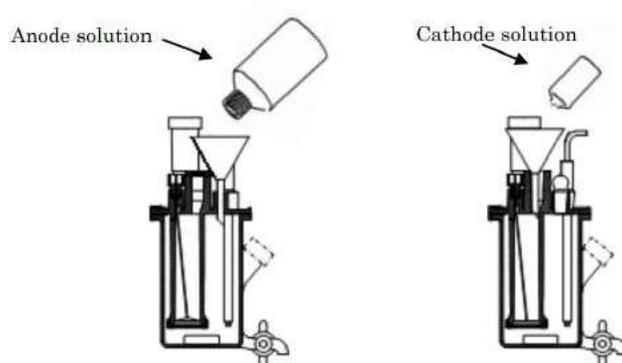


Fig.3.1. Preparation of the reagents. (Standard Cell)

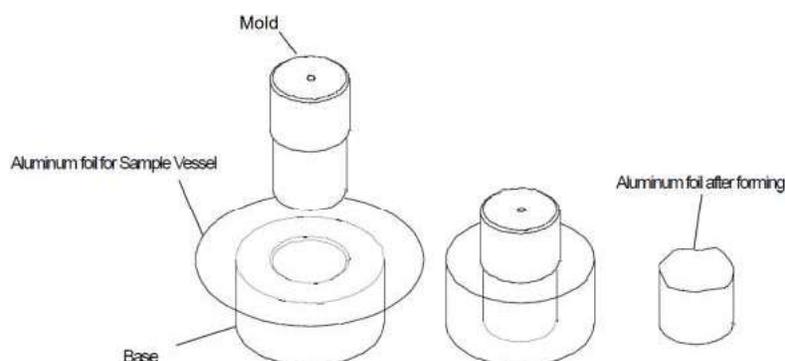
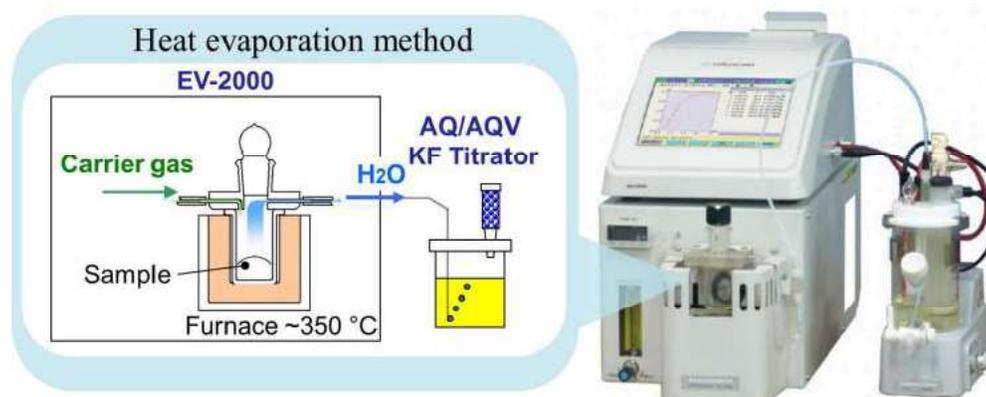


Fig.3.2. Forming the aluminum foil.



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4. parameters and results

Table 4.1 Parameters for heat evaporation method.

Condition File		EV.File	
Cal Mode	o:Sample weight(net) X=(H ₂ O-BLANK)/SIZE	* Step 1 Temp.	160 °C
Interval Time	30 sec	Time	0 min
Current	MEDIUM	Step 2 Temp.	0 °C
Min.Timer	5 min	Time	0 min
Blank Value	0 µg	Step 3 Temp.	0 °C
Unit Mode	AUTO	Time	0 min
Auto Interval	0 g	Pre. Heat End Time	5 min
Minimum Count	5 µg	End B.G.	0 µg
Back Ground	ON	Cooling Time	5 min
Sample Size Input	Every Time	B.G. Count	60
*Cell Type	Stadard/Fritless	Back Purge Time	5 sec

(* Set the electrolytic cell and heating temperature appropriately.)

Table 4.3 Results of water content measurement of water standard.

Water Standard	Temp. (°C)	Cell	No.	Sample Size (g)	Water (µg)	Water Content(%)	Statistics Results		Recovery rate(%)
15.64%	160	Standard	1	0.0529	8149.4	15.4053	Avg.	15.418 %	98.6
			2	0.0615	9506.1	15.4571	SD	0.035 %	
			3	0.0527	8110.6	15.3901	RSD	0.23 %	
		Fritless	1	0.0216	3327.3	15.4042	Avg.	15.436 %	
			2	0.0378	5853.4	15.4852	SD	0.043 %	
			3	0.0577	8896.3	15.4182	RSD	0.28 %	
5.04%	150	Standard	1	0.1426	7242.6	5.0790	Avg.	5.083 %	100.8
			2	0.1406	7181.3	5.1076	SD	0.023 %	
			3	0.1180	5973.0	5.0619	RSD	0.45 %	
		Fritless	1	0.1430	7294.1	5.1008	Avg.	5.088 %	
			2	0.1011	5143.9	5.0879	SD	0.013 %	
			3	0.1080	5479.9	5.0740	RSD	0.26 %	
0.99%	300	Standard	1	0.1236	1240.1	1.0033	Avg.	1.007 %	101.7
			2	0.1223	1230.4	1.0061	SD	0.004 %	
			3	0.1523	1539.0	1.0105	RSD	0.36 %	

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	3	0.2419	2431.8	1.0053	RSD 0.26 %		

5. Note

- (1) Validation test of titrator with solid evaporator is evaluated from the recovery rate of the result with respect to the CoA value written in the test report of the water standard. Permissible range of recovery rates differed depending on the type of water standard materials. The recovery rate of standards (i), (ii) and (iii) should fall within the range of ± 5 %, ± 4 % and ± 3 % respectively.
- (2) Select the type of water standard that is close to the water content or heating temperature of your actual sample.
- (3) The optimized electrolysis control for fritless cell of AQ series released after 2009 improves the measurement accuracy of fritless cell. It can be used with the evaporator as well. Suitable reagent for fritless cell is required. For example, Hydranal coulomat AG and AG-Oven are compatible with fritless cell.
- (4) When nitrogen gas for carrier gas is not available, optional a dry air pump for EV-2000 can be used to supply air carrier gas. When using air as a carrier gas, pay attention to the following points.
 - i) Background moisture tends to be higher than nitrogen gas.
 - ii) When the sample has tendency of thermal decomposition, it wouldl be promoted in the air.

Keywords: Karl Fischer, Coulometric titration, Heat evaporation, Water standard, Validation, Fritless cell

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