Determination of Potassium Iodate in Edible Salt by Potassium Iodide-Iodine-Starch System

Xinrong Wen^{1,2,*} and Changqing Tu^{1,2}

¹School of Chemistry and Environment, Jiaying University, Meizhou, Guangdong 514015, China ²Guangdong Provincial Key Laboratory of Conservation and Precision Utilization of Characteristic Agricultural Resources in Mountainous Areas, Jiaying University, Meizhou, Guangdong 514015, China

Keywords: Spectrophotometry, Potassium Iodate, Potassium Iodide, Edible Salt.

Abstract: In acidic medium, ki can react with kio₃ to form i₂, then i₂ and starch form i₂-starch blue complex with a maximum absorption wavelength of 596 nm. Beer's law is obeyed between the kio₃ content and the absorbance of i₂-starch blue complex. Base on this, the kio₃ content can be determined indirectly. A novel method for the determination of potassium iodate in edible salt by potassium iodide-iodine-starch system has been established. The various effect factors on the determination of iodide by potassium iodide-iodine-starch system are investigated in detail. Under optimal conditions, when the mass concentration of kio₃ is 0.4000~1.280 µg/ml, the linear regression equation is a=-0.0649+0.04702c (µg/ml) with the linear correlation coefficient is 0.9992. this proposed method had been successfully applied to determinate kio₃ in edible salt, and the results agree well with those by standard method.

1 INTRODUCTION

Iodine is one of the essential microelements for humans. It can enhance the basic metabolism and promote the growth and development of human body. Both iodine deficiency and iodine excess do harm to the human body. Iodine deficiency can cause an endemic goiter and potential damage to children's intellectual growth, and iodine excess can lead to hypothyroidism, thyroid enlargement and other clinical manifestations. Eating iodized salt is the most important and effective way to prevent iodine deficiency disease. Potassium iodiate is usually added to the edible salt, which is the iodized salt. Eating iodized salt can achieve the effect of iodine supplement. Thus, the determination of iodine content in iodized salt has great practical significance. So far, the determination methods for potassium iodinate in salt are mainly included titration (Mohammad, 2020), spectrophotometry (Gavrilenko, 2019), emission spectrometry (Yu, 2013), flow-injection (Kuznetsov, 2007), ICP-OES (Sager, 2019), CE-ICP-MS (Chen, 2007) HPLC (Manju, 2010) and so on.

In this paper, a novel method for the determination of potassium iodate in edible salt by

potassium iodide-iodine-starch system is reported. In acidic medium, I reacts with KIO₃ to form I₂, then I₂ and starch form I₂-starch blue complex with a maximum absorption wavelength of 596 nm. There is a good linear relationship between the absorbance of I₂-starch blue complex and the KIO₃ dosage, the linear equation is A=-0.0649+0.04702C(μ g/mL) within the range of 0.4000~1.280 μ g/mL KIO₃ concentration. So, by measuring the absorbance of I₂starch blue complex, the content of KIO₃ can be determined indirectly. This proposed method has been applied to determinate of KIO₃ in edible salt with satisfactory result.

2 EXPERIMENTAL

2.1 Equipment and Reagents

UV-2401 UV-visible spectrophotometer (The Shimadzu Corporation, japan); 723S spectrophotometer (Shanghai Precision & Scientific Instrument Co,. Ltd).

 KIO_3 solution: 10.00 $\mu g \cdot m L^{\text{-1}},$ a 1.000 $m g \cdot m L^{\text{-1}}$ potassium iodiate solution is prepared and then

Wen, X. and Tu, C.

Determination of Potassium lodate in Edible Salt by Potassium lodide-lodine-Starch System.

DOI: 10.5220/0012012100003633 In Proceedings of the 4th International Conference on Biotechnology and Biomedicine (ICBB 2022), pages 21-26 ISBN: 978-989-758-637-8

Copyright © 2023 by SCITEPRESS - Science and Technology Publications, Lda. Under CC license (CC BY-NC-ND 4.0)

diluted to 10.00 μ g·mL⁻¹. KI solution: 1.000 g·L⁻¹. H₃PO₄ solution: 5.0 mol·L⁻¹. Starch solution:5.0 g·L⁻¹.

All reagents are of analytical reagent grade. Bidistilled water is used.

2.2 Method

KI solution 3.00 mL, starch solution 3.00 mL, H₃PO₄ solution 1.50 mL, a certain volume of KIO₃ solution or edible salt sample solution are added into a 25 mL volumetric flask. The solution is diluted to the mark with bidistilled water, mixed well and placed at room temperature for 40 minutes in the dark. The

absorbance of I_2 -starch blue complex is measured at 596 nm against the reagent blank.

3 RESULTS AND DISCUSSION

3.1 Maximum Absorption Wavelength

In 500 ~ 700 nm, the absorption spectrum of I₂-starch blue complex is obtained using UV-2401 UV-visible spectrophotometer (Fig. 1). Fig. 1 show that the maximum absorption wavelength of I₂-starch blue complex is 596 nm.



3.2 Reaction Temperature

the increase of reaction temperature. Hereby, the room temperature is used.

The effect of reaction temperature is seen in table 1. We can sen from table 1 that the absorbance of I_{2} -starch blue complex keep constantly decreasing with

Temperature /°C	room temperature	30	35	40
Absorbance	0.380	0.375	0.374	0.372
Temperature /°C	45	50	55	60
Absorbance	0.368	0.364	0.353	0.342

Table 1: The effect of reaction temperature on the absorbance.

Experimental conditions: KIO₃:2.00 mL; KI:2.00 mL; H₃PO₄:3.00 mL; starch:2.00 mL; reaction time:15 min.

3.3 Reaction Time

The effect of the reaction time is showed in Fig. 2. It is found that the absorbance of I_2 -starch blue complex gradually increased with the reaction time, and the absorbance of I_2 -starch blue complex reaches greatest when the reaction time is 35 minutes or more. So, 40 minute is selected.



KI:2.00mL; H₃PO₄:3.00mL; KIO₃:2.00mL; starch:2.00mL.

Figure 2: Effect of the reaction time.

3.4 KI Solution Dosage

The effect of KI solution dosage can be seen in fig. 3. The results showed that as the amount of KI increases, the absorbance of I_2 -starch blue complex also

gradually increases. The absorbance of I₂-starch blue complex reaches the maximum value when KI solution dosage is 2.50 mL. Thereafter, the absorbance of I₂-starch blue is basicly stable as KI dosage increases. Thus, 3.00 mL KI solution is chosed.



H₃PO₄:3.00mL; KIO₃:2.00mL; starch:2.00mL; reaction time:40 min.

Figure 3: Effect of KI solution dosage.

3.5 H₃PO₄ Solution Dosage

The effect of H_3PO_4 solution dosage is showed in Fig. 4. The results show that the absorbance of I_2 -starch blue complex gradually increases with the amount of H_3PO_4 increases. The absorbances of I_2 -starch blue complex are essentially constant when the H_3PO_4 solution dosage is $1.00 \sim 2.00$ mL. Hence, 1.50 mL H_3PO_4 solution is used. ICBB 2022 - International Conference on Biotechnology and Biomedicine



V(H₃PO₄)/mL

KI:3.00mL; KIO₃:2.00mL; starch:2.00mL; reaction time:40 min.

Figure 4: Effect of H3PO4 solution dosage.

3.6 Starch Solution Dosage

The effect of starch solution dosage is showed in Table 2. The experimental results show that the absorbance of I_2 -starch blue complex increase with the

increase of starch solution dosage. The absorbance of I_2 -starch blue complex is maintained at stable values when the starch solution dosage is 2.50 mL or more. Therefore, the starch solution dosage is chosen as 3.00 mL.

Table 2: The effect of starch solution dosage on the absorbance.



Experimental conditions: KI:3.00 mL; H₃PO₄:1.50 mL; KIO₃:2.00mL; reaction time:40 min.

3.7 Calibration Curve

Under the optimum conditions, a series of determination solutions with different KIO₃ concentrations are prepared, then the absorbances of

these solutions are measured at 596 nm against the reagent blank. Using concentration as the abscissa and absorbance as the ordinate, the calibration curve (Fig. 5) is obtained. In the range of 0.4000-1.280 μ g/mL KIO₃, a good linear relationship between the KIO₃ concentration and the absorbance of I₂-starch blue complex, the linear equation is A=-0.0649+0.4702C(μ g/mL) and the correlation coefficient is 0.9992.



C/mg.mL⁻¹

KI:3.00 mL; H₃PO₄:1.50 mL; starch:3.00 mL; reaction time:40 min.

Figure 5: Calibration curve.

3.8 Sample Analysis

25.0000 g edible salt sample is weighed and dissolved in proper amounts of bidistilled water, then it is transferred into a 250 mL volumetric flask and diluted to the mark with bidistilled water, shaked well. This is the edible salt sample solution. According to the experimental method, 5.00 mL edible salt sample solution is added, then the absorbance of I₂-starch blue complex is determined, and the content of KIO₃ is calculated. Meanwhile, the recovery tests of standard addition are performed and the content of KIO₃ is determined by standard method. The results as show in Table 3.

Table 3: The content of KIO3 in edible salt.

	Proposed method (µg·g ⁻¹)	RSD (%)	Standard method (GB 26402- 2011) (μg·g ⁻¹)	Added (μg∙mL⁻¹)	Recovered (μg∙mL⁻¹)	Recovery yield (%)	
Natural sea salt	37.61	0.2	38.82	$0.08000 \\ 0.1600$	0.07614 0.1642	95.2 102.6	-
Low sodium salt	36.06	0.6	37.15	$0.08000 \\ 0.1600$	0.07571 0.1510	94.6 94.4	
Well cooked salt	34.40	0.3	35.45	$0.08000 \\ 0.1600$	0.07512 0.1540	93.9 96.2	

From Table 3, we can seen that the content of KIO₃ in edible salt by this proposed method is consistent with the standard method, and the recovery yields are $93.9\% \sim 102.6\%$.

4 CONCLUSION

A novel method for the determination of KIO_3 in edible salt by potassium iodide-iodine-starch system has been reported in this paper. This method has been successfully applied to the determination the content of KIO_3 in different edible salt with satisfactory results. It is obvious that the determination the content of KIO_3 by potassium iodide-iodine-starch system has certain practical significance and foreground of application.

REFERENCES

- Chen J H, Wang K E, Jiang S J. (2007) Determination of Iodine and Bromine Compounds in Foodstuffs by CE-Inductively Coupled Plasma MS[J]. Electrophoresis., 28(22): 4227-4232.
- Gavrilenko N A, Fedan D A, Saranchina N V, et al. (2019) Solid Phase Colorimetric Determination of Iodine in Food Grade Salt using Polymethacrylate Matrix[J]. Food Chemistry., 280: 15-19.
- Kuznetsov V V, Ermolenko Y V, Seffar L. (2007) Amylose and Amylopectin as Reagents for the Flow-Injection

ICBB 2022 - International Conference on Biotechnology and Biomedicine

Determination of Elemental Iodine [J]. Journal of Analytical Chemistry., 62(5): 479-485.

- Manju G, Aradhana K.K.V. P, Amrita S, et al. (2010) Salt-Assisted Liquid–Liquid Microextraction for the Determination of Iodine in Table Salt by High-Performance Liquid Chromatography-Diode Array Detection[J]. Food Chemistry.,124(4):1741-1746.
- Mohammad R A, Moumita D, Md K I, et al. (2020) Determination of Iodine Content of Commercially Available Table Salts at the Retailer Level in Selected Areas of Bangladesh[J]. European Journal of Nutrition & Food Safety., 284-288.
- Sager M. (2019) Determination of High Iodine Levels by ICP-OES after Separation from Excess Phosphate by Co-precipitation[J], Journal of Food Science and Engineering.,9(2):74-80.
- The Minister of Health of the People's Republic of China. (2011) National Food Safety Standards (GB 26402-2011): Food Additives Potassium Iodate[S]. Standards Press of China: Beijing. (In Chinese)
- Yu Y L, Dou S, Chen M L, et al. (2013) Iodine Excitation in a Dielectric Barrier Discharge Micro-Plasma and its Determination by Optical Emission Spectrometry[J]. Analyst.,138(6): 1719-1725.