



Application Note AN-C-185

Potassium in potassium bicarbonate and potassium chloride effervescent tablets for oral solution

Method validation according to the U.S. Pharmacopoeia

Potassium bicarbonate and potassium chloride effervescent tablets for oral solution are used to prevent hypokalemia (low levels of potassium in blood) [1]. Pharmaceutical manufacturers and laboratories must use approved quality monitoring techniques for drugs and formulations as stipulated by U.S. Pharmacopoeia (USP) monographs.

As an alternative to flame photometry, ion chromatography with non-suppressed conductivity detection has been approved by the USP as a validated method to quantify potassium content in

potassium bicarbonate and potassium chloride effervescent tablets for oral solution [2].

The Metrosep C 6 - 150/4.0 column ([L76](#)) provides the required separation of potassium and magnesium. All acceptance criteria from the USP monograph «Potassium Bicarbonate and Potassium Chloride Effervescent Tablets for Oral Solution» are fulfilled [2]. The present IC method has been validated according to USP General Chapter <621> Chromatography, system suitability [3].

SAMPLE AND SAMPLE PREPARATION

Sample solutions are prepared from commercially available potassium bicarbonate and potassium chloride effervescent tablets for oral solution.

Standard analyses are performed with a solution of USP Potassium Chloride RS. No additional sample preparation is required.

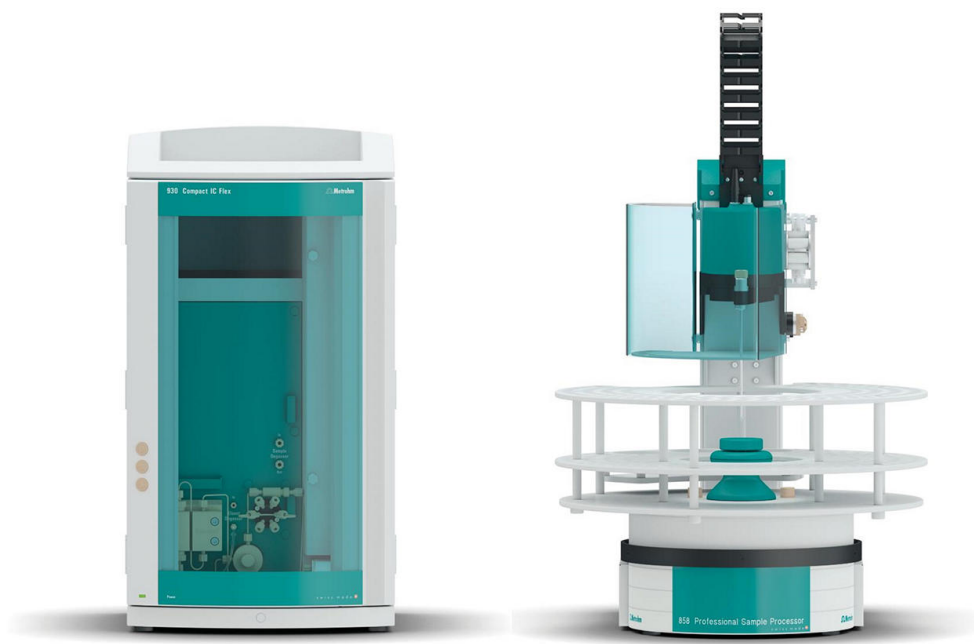


Figure 1. Instrumental setup including a 930 Compact IC Flex Oven and an 858 Professional Sample Processor.

EXPERIMENTAL

The sample stock solution is prepared through the addition of 25 g finely powdered potassium bicarbonate and potassium chloride effervescent tablets for oral solution to a 1000 mL volumetric flask. The powder is dissolved in 200 mL ultrapure water. After effervescence ceases, the volumetric flask is filled up to the mark. This stock solution contains nominally 4809.48 mg/L potassium. A 1.533 mL aliquot of this stock solution is transferred to a 500 mL volumetric flask and is filled up to the mark with ultrapure water. This final sample solution nominally contains 15.0 µg/mL potassium.

A working standard solution of 15 µg/mL potassium chloride is prepared from the respective USP reference standards.

All solutions (i.e., samples and standards) are injected directly into the ion chromatograph (**Figure 1**) using an 858 Professional Sample Processor. Potassium is separated from all other cations using the Metrosep C 6 - 150/4.0 column (L76).

The calibration is performed by using a 6-point linear calibration curve in the concentration range of 3.75–22.5 µg/mL potassium. The sample is then analyzed in duplicate.

RESULTS

The IC assay for potassium content was validated according to USP Monograph «Potassium Bicarbonate and Potassium Chloride Effervescent Tablets for Oral Solution» [2]. The accuracy of the potassium determination was calculated as 105% (Table 2 and Figure 2).

All acceptance criteria were fulfilled, e.g., the correlation coefficient for potassium was 0.9999, the resolution of adjacent peaks, and the relative standard deviation of the standard solutions was <0.15% (n = 6) (Table 2).

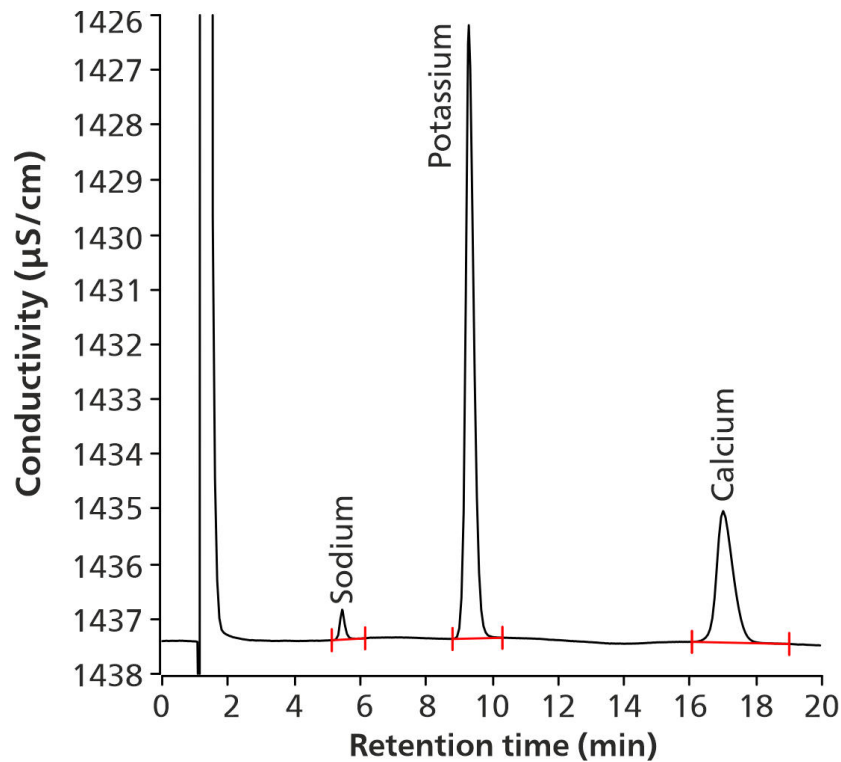


Figure 2. Chromatogram of the sample solution (105% recovery of the nominal concentration for potassium). Sodium and calcium were not quantified. The resolution was 11, for both pairs: sodium/potassium and potassium/calcium.

CONCLUSION

The presented IC method for the determination of potassium content in potassium bicarbonate and potassium chloride effervescent tablets for oral solution using the Metrosep C 6 column (packing material L76) for separation is officially included into the USP [2]. Robustness and reliability of the method

was demonstrated following the guidelines of the USP General Chapter <621> [3]. The presented setup is suitable to quantify potassium according to the USP requirements. Additional USP methods are summarized in the flyer «[Bring your USP methods up to date!](#)» [4].

REFERENCES

1. Kardalas, E.; Paschou, S. A.; Anagnostis, P.; et al. Hypokalemia: A Clinical Update. *Endocr Connect* **2018**, 7 (4), R135–R146. <https://doi.org/10.1530/EC-18-0109>.
2. *Potassium Bicarbonate and Potassium Chloride Effervescent Tablets for Oral Solution*; Monograph; U.S. Pharmacopeia/National Formulary: Rockville, MD. https://doi.org/10.31003/USPNF_M67253_02_01.
3. <621> *Chromatography, General Chapter*, U.S. Pharmacopeia/National Formulary: Rockville, MD. <https://www.uspnf.com/notices-gc-621-nitr-20220826>.
4. Metrohm AG. Bring Your USP Methods up to Date!, 2023. [8.000.5436EN](https://www.metrohm.com/asset/documents/8.000.5436EN)

CONTACT

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CONFIGURATION



Metrosep C 6 - 150/4.0

The high-capacity C 6 material makes the Metrosep C 6 - 150/4.0 separation column the optimum solution for separating standard cations with high differences in concentration in conjunction with reasonable retention times. Drinking water with low ammonium contents can be determined with this column.



Metrosep C 6 Guard/4.0

The Metrosep C 6 Guard/4.0 contains the C 6 column material and is used to protect against particles and contamination. This considerably lengthens the service life of the analytical separation column. The Metrosep C 6 Guard/4.0 works according to the "On Column Guard System" and is screwed directly onto the respective separation column with virtually no dead volume.



930 Compact IC Flex Deg

The 930 Compact IC Flex Deg is the intelligent Compact IC instrument **without suppression** with built-in **degasser**. The instrument can be used with any separation and detection methods.

Typical areas of application:

- Anion or cation determinations without suppression with conductivity detection
- Simple applications with UV/VIS or amperometric detection



858 Professional Sample Processor

The 858 Professional Sample Processor processes samples from 500 μL to 500 mL. The sample transfer takes place either by means of a peristaltic pump on the 850 Professional IC system or with an 800 Dosino.