A variety of biologically active compounds have been found in raspberry leaves, including flavonoids, phenolic acids, organic acids, ellagitannins, gallotannins and proanthocyanidins [1, 2]. There is a rather high interest in determining the amount of organic acids in raspberry leaves in literary sources [3, 4]. According to these studies, the total amount of organic acids is relatively high and ranges from 1.1 to 2.3%; it makes raspberry leaves a promising raw material of organic acids [5, 6].

Organic acids are a group of biologically active compounds with a wide spectrum of the pharmacological activity. They reveal vitamin properties, possess the choleretic effect, and normalize the activity of the digestive system. Organic acids regulate the secretion of the bile and pancreatic juice, improve appetite, possess bactericidal properties and reduce preteractive processes in the body [7].

Nowadays there are various analytical methods for determining organic acids such as titration, ion chromatography [8], high-performance liquid chromatography-ultraviolet detection (HPLC-UV) [9], capillary electrophoresis [10], gas chromatography (GC) [11] and thin-layer chromatography [12].
Although HPLC-UV and GC have been proposed for the determination of organic acids before, they require high skillful analysts and expensive instruments. In addition, most of the procedures previously described require expensive instrumental setup. That is why in the current work the method of alkalimetry with potentiometric detection of the end-point was chosen for determining organic acids in raspberry leaves as it is simple, reliable, rapid and economical. Thus, the aim of the study was to validate the method proposed for the quantitative determination of free organic acids in raspberry leaves.

**Experimental part**

The study object was raspberry leaves collected in the Kharkiv region during the period of full ripening, according to the rules of harvesting this type of plant. Drying of leaves was carried out at a temperature of 60°C to a residual humidity of not more than 20%. A Hanna 2550 pH meter with HI 1131P potentiometric electrode was used in the study. All titrations were carried out manually. Free organic acids were titrated using a microburette with Class A accuracy. Weighing was carried out using an AN100 digital analytical balance (AXIS, Ukraine) with \( d = 0.0001 \) g.

Citric acid was purchased from Sigma Aldrich (≥ 98%), NaOH was of analytical grade. In order to prepare NaOH solution with the concentration of 0.05 M, 1.0 g of NaOH was dissolved in distilled water. The solution was diluted to 250 mL with the same water and standardized.

The procedure of the quantitative determination of free organic acids in raspberry leaves. Place 2.0 g (accurate weight) of the crushed raw material in a 100 mL flask with a ground glass joint, then pour 40 mL of distilled water in the flask, equip it with a condenser and keep the flask for 2 h on a boiling water bath. After cooling quantitatively transfer the solution into a 50 mL volumetric flask and dilute to the volume (solution A).

Place 5.00 mL of the solution A in a 100 mL flask, and add 45.0 mL of distilled water with the subsequent titration of the solution by 0.05 M sodium hydroxide. After adding each portion of 0.05 M sodium hydroxide mix the solution thoroughly, and record the electrode potential.

The blank experiment was also performed. According to it, the blank volume of 0.05 M sodium hydroxide was 0.03 mL.

The content of organic acids in the solution was calculated from the value of the equivalent volume of the titrant. The equivalent volume of the titrant was determined by a differential curve constructed in the \( \Delta E/\Delta V – V \) coordinates. The equivalence point was fixed at the maximum of the differential curve. The perpendicular line was dropped to the horizontal axis (the volume of the titrant) through the maximum, and the volume of the titrant spent on titration was determined (Fig. 1).

The content of free organic acids (X, %) with reference to citric acid in the completely dry raw material was calculated by the following formula:

\[
X = \frac{(V_x - V_{eq}) \cdot 0.0032 \cdot 50 \cdot 100 \cdot K}{m \cdot 5 \cdot (100 - W)}
\]

where: 0.0032 – is the amount of citric acid equivalent to 1 mL of sodium hydroxide solution (0.05 mol/L), g; \( V_x \) – is the volume of sodium hydroxide solution (0.05 mol/L) used for titration, mL; \( V_{eq} \) – is volume of sodium hydroxide solution (0.05 mol/L) spent for titration in the blank experiment, mL; m – is the mass of the raw material used, g; K – is the correction coefficient for 0.05 mol/L sodium hydroxide solution; W – is the loss on drying of the raw material, %.

![Fig. 1. The potentiometric titration curve of determination total free organic acids in raspberry leaves](image-url)
Validation

Validation of the alkalimetric method for the quantitative determination of the amount of free organic acids in raspberry leaves by potentiometric titration was performed according to the International Conference on Harmonization (ICH). The titrimetric method proposed was validated by the following parameters: specificity, accuracy, linearity, repeatability, intermediate precision, robustness.

The specificity of the method was studied by potentiometric titration of the solvent.

The accuracy was verified by the method of additives in a triplicate analysis of three levels of concentration of organic acids corresponding to 40, 60, 80% of the working concentrations of organic acids. The standard solution of citric acid was prepared as follows: 0.076 g (accurate weight) of citric acid was placed in a 200.0 mL volumetric flask, and the solution was diluted to the volume with distilled water. Then an aliquot of the resulting standard solution of 2.00, 3.00, 4.00 mL was taken and placed in a 100.0 mL flask. After that 5.00 mL of the extract obtained from raspberry leaves was added to the flask, then 45.0 mL of distilled water was added, and the solution was titrated. The evaluation criterion in determining the accuracy was the value of the relative standard deviation (RSD), which according to the requirements should be not more than 2%, and the percentage of recovery should be from 95 to 105%.

The linearity of the method was studied at 9 concentration levels (40, 60, 80, 100, 120, 140, 160, 180, 200%) of the theoretical content of the total amount of free organic acids (calculated with reference to citric acid, %) in raspberry leaves. In order to evaluate linearity of the method, different aliquots of the extract were taken (2.00; 3.00; 4.00; 5.00; 6.00; 7.00; 8.00; 9.00; 10.00 mL). After that each aliquot was placed in a 100.0 mL flask and diluted to the volume with distilled water. The quantitative content of the total amount of free organic acids (calculated with reference to citric acid, %) in raspberry leaves in the solutions obtained was then determined according to the alkalimetry method of titration. The linearity was assessed by a linear relationship between the concentration of the total amount of free organic acids and the equivalent volume of the titrant, as well as the correlation coefficient calculated. The linear regression was calculated by the method of least squares to obtain the regression equation and determine the correlation coefficient (r²). According to the requirements of ICH, the value of the correlation coefficient when studying the linearity of the analytical method for determining the quantitative content of the active substance should be ≥0.999.

The repeatability of the method was checked by preparing an aqueous extract of raspberry leaves from 6 portions of the raw material within a short period of time using the same set of reagents and with the participation of the same analyst. The intermediate precision was determined as described above in the same laboratory, but in different days. The acceptance criterion is expressed by the value of the relative standard deviation, which should not exceed 2%.

The robustness of the potentiometric procedure was tested at 100% concentration of free organic acids with the participation of two different analysts and two different burettes. The acceptance criterion is expressed by the value of the relative standard deviation, which should not exceed 2%.

The statistical processing of experimental data obtained was performed in accordance with the monograph «Statistical analysis of the results of a chemical experiment» of the State Pharmacopeia of Ukraine.

Results and discussion

When studying the specificity of the method, it was shown that the solvent used in the samples preparation and the probable impurities did not affect the result of the quantification of the amount of free organic acids in raspberry leaves (Table 1).

Linearity was proven in the concentration range from 40% to 200%. A calibration curve of the dependence of the equivalent volume of the titrant values on the volume of aliquots of the extract from raspberry leaves was plotted (Fig. 2). The regression equation of the curve had the following form: \( y = 0.1667x - 0.2722 \). The value of the correlation coefficient (r²) was equal to 0.9991 (Fig. 2).

When testing the method for the quantitative determination of the amount of free organic acids by linearity parameters it was found that the correlation coefficient (r²) was 0.9991. Therefore, there is a direct linear relation between the equivalent volume of the titrant and the concentration of organic acids.

| \( V_{\text{t}_0} \), mL | Content of organic acids, % | Statistical analysis |
|-------------------------|-----------------------------|----------------------|
| Blank experiment (titration of distilled water) | | 0.045 ± 0.02% \( s_x = 0.0014 \) |
| 0.03 | 0.05 | |
| 0.02 | 0.035 | |
| 0.03 | 0.05 | |
| Results of titration of the extract from raspberry leaves | | 1.00 ± 0.02% \( s_x = 0.0033 \) |
| 0.57 | 1.00 | |
| 0.56 | 0.99 | |
| 0.57 | 1.00 | |
The accuracy of the method was assessed using the percentage of recovery and the relative standard deviation. The percentage of recovery was found to be in the range from 98.77 to 102.48%, and its average value was 100.45%, the value of the relative standard deviation when assessing the correctness of the method was 1.45% and did not exceed 2% (Table 2).

The precision of the method was confirmed by repeatability and intermediate precision. The values of RSD for repeatability and intermediate precision were 1.58 and 1.74%, respectively. The RSD values were less than 2%. It proves that the method is precise (Tables 3, 4).

The robustness of the method was determined by changing the analyst and burette. It was found that the RSD values of inter-analyst and inter-burette were

---

**Table 2**

Recovery studies by the procedure of standard additions

| The amount present, g | The amount of citric acid added, g | The amount of organic acids taken, g | The amount recovered, g | Recovery, % | SD, % | RSD, % |
|----------------------|-----------------------------------|-----------------------------------|------------------------|------------|-------|--------|
| 0.019                | 0.0052                            | 0.0242                            | 0.024                  | 99.17      |       |        |
| 0.019                | 0.0134                            | 0.0324                            | 0.032                  | 98.77      | 1.46  | 1.45   |
| 0.019                | 0.0184                            | 0.0374                            | 0.037                  | 98.90      |       |        |
|                      |                                   |                                   | 0.0372                 | 101.07     |       |        |

**Table 3**

Repeatability of organic acids in raspberry leaves

| Number of samples | Content of free organic acids, % | SD              | Confidence interval (P = 95%), % | RSD, % |
|-------------------|----------------------------------|-----------------|----------------------------------|-------|
| 1                 | 0.95                             |                 |                                  |       |
| 2                 | 0.97                             |                 |                                  |       |
| 3                 | 0.93                             |                 |                                  |       |
| 4                 | 0.95                             |                 |                                  |       |
| 5                 | 0.95                             |                 |                                  |       |
| 6                 | 0.95                             |                 |                                  |       |
| Mean, %           | 0.95                             |                 |                                  |       |
| SD                | 0.0151                           |                 | 0.0120                           | 1.58  |
1.52 and 1.37%, respectively. The RSD values were less than 2%, showing that minor changes in conditions have little effect on the results (Table 5).

Conclusions

The alkalimetric method for the quantitative determination of free organic acids in the raspberry leaves has been developed and validated according to the following parameters: specificity, linearity, accuracy, repeatability, intermediate precision, robustness. It has been confirmed that the method is simple, reliable, accurate and cost-effective.

Conflict of Interests: the authors have no conflict of interests to declare.

References

1. Padmanabhan, P.; Correa-Betanzo, J.; Paliyath, G. Berries and Related Fruits. In Encyclopedia of Food and Health; Caballero, B.; Finglas, P. M.; Toldrà, F., Eds. Academic Press: Oxford, 2016; pp 364–371. https://doi.org/10.1016/B978-0-12-384947-2.00060-X.

2. Sariburun, E.; Şahin, S.; Demir, C.; Türkben, C.; Uylaşer, V. Phenolic Content and Antioxidant Activity of Raspberry and Blackberry Cultivars. J. Food Sci. 2010, 75 (4), C328–C335. https://doi.org/10.1111/j.1750-3841.2010.01571.x.

3. Velichko, V. V.; Makarova, D. L. Comparative pharmacognostic analysis of leaves and fruit of raspberry ordinary. Medicine and Education in Siberia 2015, 4.

4. Дергачева, Ж. М.; Гурина, Н. С.; Мушкина, О. В. Фитохимический анализ листьев малины обыкновенной (Rubi Idaeus Folia). Рецепт 2015, 6, 64–74. https://doi.org/10.3390/antiox5020017.

5. Bobinaitė, R.; Viškelis, P.; Alarcón, S. H. Multivariate analysis of organic acids in fermented food from reversed-phase high-performance liquid chromatography data. Talanta 2018, 178, 15–23. https://doi.org/10.1016/j.talanta.2017.09.005.
10. Nogueira, T.; Lago, C. L. d. Determination of Ca, K, Mg, Na, sulfate, phosphate, formate, acetate, propionate, and glycerol in biodiesel by capillary electrophoresis with capacitively coupled contactless conductivity detection. *Microchem. J.* 2011, 99 (2), 267–272. https://doi.org/10.1016/j.microc.2011.05.014.

11. Zhilkina, V. Yu.; Marakhova, A. I.; Stanishevskiy, Ya. M. Qualitative and quantitative analysis of organic acids in mixture of multivitamin raw material. *Drug development & registration* 2016, 1, 156–159.

12. Umarov, U. A.; Maskov, O. Y.; Kolisnyk, S. V.; Fathullaeva, M. Development and Validation of The Conductometric Titration Method of Quantitative Determination of Free Organic Acids in The Anise Fruits. *European Journal of Molecular & Clinical Medicine* 2020, 7 (3), 3874–3883.

13. Sochorova, L.; Torokova, L.; Baron, M.; Sochor, J. Electrochemical and others techniques for the determination of malic acid and tartaric acid in must and wine. *Int. J. Electrochem. Sci.* 2018, 13 (9), 9145–9165. https://doi.org/10.20964/2018.09.20.

14. Strömberg, N.; Sahlin, E. Determination of the short-chain fatty acid pattern in biodiesel using high throughput syringe solvent extraction and ion exclusion chromatography. *Fuel* 2012, 97, 531–535. https://doi.org/10.1016/j.fuel.2012.01.032.

15. The International Council for Harmonisation. Quality Guidelines. https://www.ich.org/page/quality-guidelines (accessed Jan 10, 2021), Validation of analytical procedures: text and methodology Q2(R1).

16. Державна фармакопея України: в 3 т., 2-е вид.; Державне підприємство «Український науковий фармакопейний центр якості лікарських засобів»: Харків, 2015; Т. 1.