Spectrophotometric determination of hesperidin in supplements and orange juices

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INTRODUCTION

Flavonoids as the biggest group of plant polyphenols are widespread, but mostly can be found in angiosperms. Actually, they are pigments responsible for the flowers, fruits and leaves colors, but they also take parts in numerous metabolic and physiologic processes in plants.

Availability of citrus fruits all around the world nowadays, during the whole year, leads to numerous studies focused on the effects of hesperidin (Hesp), a flavonoid that consists of aglycone hesperitin and sugar rutinoside (Figure 1).

Hesperidin is flavanone mainly stored in the inner peel of citrus fruits, called albedo. Hesperidin can be used alone or in combination with other bioflavonoids and/or vitamin C as dietary supplements.

Positive effects of hesperidin are mainly attributed to its strong antioxidant activity, increasing its ability to scavenge free radicals [1]. Some of the most often reported positive effects of hesperidin are anti-inflammatory, antiallergenic, antiviral and anticarcinogenic effect [2]. Hesperidin contributes to strength of the blood vessels walls, leads to the bruise preventing and some conditions related to certain vascular disorders [3]. Further, even promising effects of orange juice daily consumption have been reported as the results of performed studies, such as lowering of diastolic blood pressure [4]. Further, significant radioprotective effect of hesperidin has been reported [5].

As a very important fact in hesperidin usage, no signs of toxicity have been reported with the normal intake of hesperidin or related compounds. Further, both hesperidin and its aglycone hesperetin have been observed to possess a wide range of pharmacological properties.

Abstract

Hesperidin is the flavonoid heteroside most found in citrus species, such as grapefruit, lemon and orange. It is applied in the treatment of numerous cardiovascular diseases, with many products containing hesperidin in combination with other flavonoid heterosides or vitamin C. The aim of this study was to develop and validate a simple, rapid and sensitive and low-cost method for determination of hesperidin in tablets and orange juices. The developed spectrophotometric method is based on the formation of the Zn(II)-hesperidin complex in 70 v/v% methanol at pH 3.12, with max of absorption at λmax= 283 nm. The method of variation of equimolar solutions confirmed the stoichiometry of the complex as hesperidin: zinc= 2:1. The stability constant of the complex was obtained by the Bjerrum method and it is logβ2 = 17.01 at pH 3.12. Linearity range for the developed method was defined as 0.61-7.32 μg/mL, (confirmed by coefficient of determination, r² = 0.9882), while limits of detection and quantification were calculated as LOD = 0.15 μg/mL and LOQ = 0.45 μg/mL. Fairly high precision of the method is indicated by low values of CV, varying in the range 0.26-1.23%. The proposed method fulfills all aimed requirements, and it was successfully applied for determination of hesperidin in tablets with recovery value 92.2 % and commercially available orange juices.

Key words: hesperidin, orange juice, spectrophotometry, zinc complex.
Hesperidin possesses the strong ability to form complexes with transition metal ions, helping the plant defense mechanism against the pathogens and plant-eaters. The same ability of metal ions complexation is used as the quality test for the presence of flavonoids in extracts and biological specimens.

Hesperidin can be determined alone and/or with other flavonoids by many methods, such as HPLC [6,7], adsorptive-stripping voltammetry [8], spectrophotometry [9,10], and pulse perturbation of the oscillatory reaction system [11]. The very sensitive method for hesperidin determination in plasma has been reported to be liquid chromatography tandem mass spectrometry (LC-MS/MS) [12].

Considerable numbers of the reported methods are based on the analysis of flavonoid-metal complexes with aim to follow the quality control of dietary supplements and food products.

Previously, we have developed the methods for the determination of hesperidin in human plasma and pharmaceutical forms based on the fluorescence properties of the aluminium-hesperidin complex in the presence of zwitterionic surfactant sulfobetaine – SB 12 [13], and without presence of surfactant [14]. The methods have great sensitivity but a spectrophotometer is still a too expensive instrument for regular daily control of the products.

In this work we have proposed a spectrophotometric method based on the absorption of the zinc(II) – hesperidin complex. We have tested and chosen the optimal conditions for the formation of zinc(II) – hesperidin complex with aim to apply the developed method for the quantification of hesperidin in tablets and several samples of orange juices.

**MATERIAL AND METHODS**

**Reagents and Materials**

Hesperidin (Fluka AG), methanol, NaOH, CH₃COOH, CH₃COONa, zinc-chloride, (Merck), all p.a. grade, were used. The stock solution of zinc-chloride (1.0 × 10⁻⁴ mol/L) was prepared by dissolving ZnCl₂ in doubly distilled water. The stock solution of hesperidin (1.0 × 10⁻⁴ mol/L) was prepared by dissolving hesperidin in methanol (70 % v/v) and was stored in a refrigerator.

Working solutions of the zinc(II)-hesperidin complex was prepared by dilution of the stock solutions of zinc(II) (2.5 × 10⁻⁵ mol/L ZnCl₂ and hesperidin (5.0 × 10⁻⁷ to 1.0 × 10⁻⁵ mol/L).

Acetate buffers (in 70 wt% methanol and pH 3.12), previously prepared according to Perrin [15], were used for all spectrophotometric measurements.

**Instruments**

Spectrophotometric measurements were performed on a Beckman DU-650 spectrophotometer, using 1 cm of quartz cells.

Measurements of pH were carried out using a Mettler Toledo mp 120 pH meter, equipped with a combination electrode.

**Sample preparation**

**Procedure for analysis of hesperidin in pharmaceutical preparations**

Chosen tablets Helopyrin (Rosch & Handel, Vienna, Austria) were prepared for analysis according the following procedure. Ten tablets were weighed and powdered using pestle and mortar. A portion of the powder, equivalent to weight of one tablet, was dissolved in 100 mL 70 v/v % of methanol and kept in an ultrasonic bath at 25 °C for 30 min, after which the solution was filtered through Millipore membrane filter.

Appropriate volumes of each filtrate and a 3.0 mL portion of 1.0 × 10⁻⁴ mol/L ZnCl₂, were mixed in a volumetric flask of 10 mL and diluted to the mark with 70 v/v% methanol. The expected concentration of hesperidin in prepared solutions was approximately 5.0 × 10⁻⁶ mol/L. Values of pH of these solutions were adjusted to 3.12. The absorbance of prepared solutions was measured at λ max = 283 nm against 70 v/v % methanol at pH 3.12 as blank.

All measurements were performed in triplicate. Origin 7 software was used for all necessary calculations and to obtain appropriate analytical parameters.

**Procedure for analysis of hesperidin in orange juices**

To the aliquot of 0.25 mL of tested orange juice, add 2.5 mL ZnCl₂ of concentration 2.5 × 10⁻⁵ mol/L and 7.25 mL of acetate buffer pH 3.12 in methanol and centrifuged at 900 rpm for 5 min. Measured absorbance of transparent supernatant at λ max = 283 nm with 70 v/v % methanol at pH 3.12 as blank was used with previously obtained calibration curve to calculate the hesperidin content in tested samples.

**RESULTS AND DISCUSSION**

**Complex formation between hesperidin and zinc(II)-ion**

Hesperidin is poorly soluble in water, but it dissolves in methanol, where the water and methanol ratio strongly affects the solubility of hesperidine. It was found that the most favorable ratio of methanol to water is 70:30, since that is the optimum condition for the formation of the Zn²⁺ -hesperidin complex.

Absorption spectra of hesperidin, ZnCl₂, and Zn-hesperidin complex solutions were recorded in the range 250 nm – 400 nm against 70 v/v% methanol as blank. The complex is stable in a wide range of pH values, from pH 1 to pH 7. It is confirmed that complex Zn²⁺ – Hesp shows maximum of absorbance at λ max = 283 nm, at pH 3.12, while the solution of ZnCl₂ does
not exhibit considerably significant absorbance in the same range, 250 – 400 nm.

The ratio of Zn^{2+} and hesperidin in their complex were determined by the method of variations of equimolar solutions [16], prepared in acetate buffer pH 3.12 in 70% methanol; as a result, the complex was defined as ZnHesp_2. The composition of the zinc – hesperidin complex was also followed by the mole ratio method [17], confirming the zinc–hesperidin ratio 1:2 for the complex formed at pH 3.12.

The dependence of absorbance intensity on pH was tested in the acetate buffers (in 70 v/v % methanol) of different pH values, prepared according to Perrin [15]. Figure 2 represents the absorbance intensity as function of solution pH, where the strong pH dependence can be noticed.

Figure 2. Dependence intensity of absorbance on the pH

Spectrophotometric determination of hesperidin

For the spectrophotometric determination of hesperidin based on absorbance depending on the concentration of formed Zn^{2+}-hesperidin complex, the calibration curve method was applied. Eight solutions of hesperidin were prepared, with hesperidin concentrations increasing from 1.0 x 10^{-6} mol/L to 1.2 x 10^{-5} mol/L, while Zn^{2+} ion was in excess, c = 2.5 x 10^{-5} mol/L. All used solutions were prepared in acetate buffer, pH 3.12. Solutions were prepared 0.5 h prior to spectra recording at \lambda = 283 nm against 70%v/v methanol as blank.

Linearity

The high value of the stability constant of the zinc(II) – hesperidin complex guarantees the quantitative determination of hesperidin based on the absorbance characteristics of the complex. The calibration curve method was used, requiring solutions containing constant concentration of ZnCl_2 and different concentrations of hesperidin in acetate buffer pH 3.12 (70 v/v % methanol as solvent) and acetate buffer pH 3.12 as blank. Linear dependence of the absorbance of the complex on the concentration of hesperidin was obtained in the interval as 0.60 – 7.32 μg/mL. The regression equation; 

\[ A = 0.0079 + 0.0186 \cdot c \]

was calculated with the aid of Origin v. 7 software, where A is absorbance (\lambda = 283 nm) and c is concentration in μg/mL. The good linearity of the calibration curve and small scatter of experimental points resulted in a high coefficient of determination, \( r^2 = 0.9882 \).

LOD (Limit of Detection) and LOQ (Limit of Quantification)

The limit of detection (LOD) was calculated according to the formula: LOD = 3.3 s_b/a, while the limit of quantification (LOQ) was determined by using the formula: LOQ = 10 s_b/a. where s_b is the standard deviation in the intercept and a is the slope of the calibration line [18, 19]. It was found that LOD, establishing the minimum level at which hesperidin can be detected, is 0.15 μg/mL, while hesperidin can be quantified at a concentration of 0.45 μg/mL.

Precision

The repeatability of the method was determined for four different hesperidin concentrations (Table 1). The accuracy and repeatability of the method are satisfied, as indicated by good recovery and low values of CV.

The quantitative spectrophotometric determination of hesperidin

The formation of a stable zinc-hesperidin complex in methanolic solution with enhanced absorbance can be utilized for quantitative determination of hesperidin in both tablets and orange juices in considerable amounts.

We have applied the standard addition method to test the matrix effect of the tested samples. Those preliminary tests showed that nonspecific absorption from juice matrix is eliminated by high dilution of tested juices and there was no need to apply the standard addition method. The same preliminary test (the standard

| Table 1. The spectrophotometric determination of hesperidin in aqueous – methanolic solutions. |
|-------------------------------------------------------------|
| Taken (μg/mL) | Found (μg/mL) | Recovery (%) | SD            | CV(%)         |
|----------------|----------------|--------------|----------------|---------------|
| 0.80           | 0.79           | 98.75        | 9.75 x 10^{-3} | 1.23          |
| 2.40           | 2.38           | 99.17        | 1.66 x 10^{-2} | 0.70          |
| 5.00           | 5.01           | 100.20       | 1.32 x 10^{-2} | 0.26          |
| n=5            |                |              |                |               |
addition method) was performed prior to application of the proposed spectrophotometric determination of hesperidin in tested tablets.

**Hesperidin content in dietary supplement tablets**

As a result, by the proposed spectrophotometric method, 18.44 mg of declared 20 mg was found in tested supplement tablets, with recovery value of 92.2 % and coefficient of variation 0.70 %.

**Hesperidin content in orange juices**

Taking into account that citrus and citrus juices are naturally abundant sources of hesperidin, it is necessary to develop a simple, accurate and precise method for its determination in such samples. We found that the method we have proposed has more than satisfactory sensitivity for the routine determination of hesperidin in citrus juices.

The results of hesperidin spectrophotometric determination based on the zinc complex for four commercially available orange juices available on the Serbian market are given in Table 2. The recovery values are not given as the tested orange juices do not have declared content of hesperidin.

Table 2. Hesperidin content in tested orange juices.

| Juices                  | Hesperidin content, mg/L |
|-------------------------|--------------------------|
| Sample 1 – pulpy        | 67.8                     |
| Sample 2 – nectar       | 281.9                    |
| Sample 3 – 100 % fruit  | 443.9                    |
| Sample 4 – 100 % fruit  | 423.2                    |

Without doubt, the reported spectrofluorimetric method [13] leads to better sensitivity and precision of determination, necessary for the quantification in biological fluids. However, having in mind the expected quantity of hesperidin in orange juices, the spectrophotometric method is more than satisfactory.

The linearity range for the spectrophotometric determination of hesperidin reported in literature was from $2.5 \times 10^{-5}$ mol/L to $1.75 \times 10^{-4}$ mol/L [10], while the here developed method may be successfully applied in the range from $1.0 \times 10^{-6}$ mol/L to $1.2 \times 10^{-5}$ mol/L. Linearity in such low concentration range provides the use of very diluted samples of juices, which leads to avoiding of matrix effects, as we have confirmed by the standard addition method in our preliminary experiments to verify the method applied to juice samples.

As it was mentioned before, the highest concentration of hesperidin in citrus species fruits are present in the albedo. That is the explanation why citrus juices obtained by squeezing the fruits do not have high concentration of this flavonoid, comparing to industrial produced orange juices, as the most represented citrus juice on the market. These juices are produced by grinding whole fruits, explaining their very high concentration of hesperidin.

According to the *Code of Practice for evaluation of quality and authenticity of fruit and vegetable juices*, published by the AIJN-European Fruit Juice Association [20], the content of hesperidin is one of the special requirements for quality of citrus based juices such as oranges, lemons and grapefruits, and that has to be 250-700 mg/L.

Results of hesperidin determination of the here proposed spectrophotometric method are within the defined range for all tested juices, except for one, Sample 1. Although the “pulpy” is part of the name of that product, according to very low content of hesperidin, 67.8 mg/L, the fact is that it is not fruit juice, but diluted fruit base with added pulp.

Sample 2 is labeled as nectar, and that is the product containing less than 50% of fruit content. If we compare the hesperidin content for Sample 2 (281.9 mg/L) with contents for samples 3 and 4 labeled as 100 % fruit (443.9 and 423.2 mg/L), it can be observed that two times more fruit content do not result in doubled flavonoid content, such as hesperidin in this particular case. It has been previously reported in literature that sometimes juices labelled as consisting of 50% fruit nectar content have almost the same quality as those declared as 100% fruit juice [21]. Although it has been observed a high correlation has been observed between the flavonoids content, the total phenol content, and antioxidant capacities of fruit products such as juices, fruit juice labels based only on fruit % could sometimes misinform consumers.

It is well known fact that the concentration of bioactive compounds (such as flavonoids) in fruits is a very complex multifactorial dependence, related to geographical region, climate, soil type, cultivar methods, growing season, harvest date, storage conditions, low-dose irradiation, and other conditions [22, 23], so like for other compounds, the content of hesperidin could vary in a wide range.

However, here represented results of hesperidin content for all tested samples are in agreement with the *Code of Practice for evaluation of quality and authenticity of fruit and vegetable juices*, and indicates just the origin, production method and thus the quality of final product are satisfactory.

Although the numerous beneficial effects of daily fruits and vegetables (and therefore flavonoids) intake are a wide spread fact, it should be noted that overlarge intake of hesperidin may trigger a number of side effects, including abdominal pain, diarrhea, and nausea [24]. Because of hesperidin effects, it is necessary to have in mind its possible interactions with certain
medications such as anticoagulants, blood pressure drugs, and calcium channel blockers. That is one more reason for having more precise data about the content of flavonoids, including hesperidin, and finding the appropriate low-cost method for routine products’ analysis, such as juices and dietary supplements.

CONCLUSION
In this work, a simple spectrophotometric determination of hesperidin in orange juice, after its complexation with zinc(II)-ion, provides good accuracy and precision and may be used for routine analysis for both tablets and orange juices. Tested orange juices fulfill the requirements considered for the content of hesperidin.

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Spektrofotometrijsko određivanje hesperidina u suplementima i sokovima od pomorandže

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Kratak sadržaj
Hesperidin je flavonoidni heterozid koji se najčešće nalazi u citrusnim vrstama, kao što su grejpfrut, limun i pomorandža. Primjenjuje se u lečenju brojnih kardiovaskularnih bolesti, sa mnogim proizvodima koji sadrže hesperidin u kombinaciji sa drugim flavonoidnim heterozidima ili vitaminom C. Cilj ove studije je razvijanje i primena jednostavne, brze i osetljive, a pri tome pristupačne metode za određivanje hesperidina u tabletama i sokovima od pomorandže. Razvijena spektrofotometrijska metoda zasniva se na formiranju Zn(II)-hesperidin kompleksa u 70 v/v% metanolu pri pH 3,12, sa maksimumom absorbancije na λ<sub>max</sub> = 283 nm. Metodom varijacije ekvimagazin rastvora utvrđena je stehiometrija kompleksa hesperidin:cink = 2:1. Konstanta stabilitnosti kompleksa dobijena Bjerumovom metodom iznosi logβ<sub>2</sub> = 17,01 pri pH 3,12. Linearnost razvijene metode utvrđena je u oblasti 0,61 – 7,32 μg/mL, (potvrđeno koeficijentom određivanja, r<sup>2</sup> = 0,9882), dok su izračunati limiti detekcije i kvantifikacije LOD = 0,15 μg/mL i LOQ = 0,45 μg/mL. Na visoku preciznost metode ukazuju niske vrednosti koeficijenata varijacije (CV je u oblasti 0,26 – 1,23 %). Predložena metoda ispunjava sve postavljene zahteve i sa uspehom je primenjena za određivanje hesperidina u tabletama uz recovery vrednost 92,2 % i komercijalno dostupnim sokovima od pomorandže.

Ključne reči: hesperidin, sok od pomorandže, spektrofotometrija, kompleks cinka.