UHPLC/MS detection of yohimbine in food supplements.

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Abstract

UHPLC in combination with electrospray ionization mass spectrometric detection in the positive ion mode has been used successfully to screen the yohimbine content in food supplements. The chromatographic behaviour of yohimbine is examined using Accela UHPLC with mass spectral detection HRMS “Q-Exative” with H-ESI-interface (“Thermo Fisher Scientific”, Waltham, MA, USA) and columns “Kynetex” RP C8 50 × 3 mm × 2.6 μm type “core-shell” and “Poroshell” RP CE18 150 × 3 mm × 2.7 mm. We have used mobile phase containing: acetonitrile, methanol water, ammonium formate. We have made series of chromatograms in positive mode of ionization and in negative mode of ionization. We have found that the ionization in positive mode offers a significantly higher sensitivity and also higher selectivity than the negative ion mode. This analytical method could be successfully applied for analytical control of food supplements containing yohimbine.

Keywords: Yohimbe, Yohimbine, Pausinystalia yohimbe, Food supplement, HPLC.

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Introduction

Yohimbine is an alkaloid derived from the African plant Pausinystalia yohimbe (Rubiaceae). Pausinystalia yohimbe is a tree native to the coastal forests of Central Africa and is distributed from South East Nigeria to the Congolese Mayombe [1]. Its bark contains up to 6% of a mixture of alkaloids. The bark of the plant has been traditionally used in folk medicine as general tonic, performance enhancer and as an aphrodisiac [2-6]. The pharmacological researches on yohimbine have been started since 1949 [7,8]. It has been found that Yohimbine antagonizes α-2 adrenergic receptors. Nowadays yohimbine is used as an active ingredient in many food supplements. These products are promoted to improve sexual and athletic performance and also for weight loss. The daily intake of yohimbine can cause many adverse effects like headache, nausea, increased urinary urge; insomnia, anxiety, restlessness, irritability, increase of blood pressure and pulse rate, palpitation, dizziness, vomiting, anorexia, gastric complaints, flush, sweating, shivering, allergic reactions, nervousness, hypotension, tremor, bronchospasm, dysuria, decreased urge, genital pains, exanthema. For this reason in several European countries (e.g. United Kingdom, Ireland, Netherlands, Belgium, Denmark, Czech Republic) as well as in Canada, Australia and New Zealand the inclusion of yohimbine in food supplements or foods is prohibited [9]. In the most part of World yohimbine products are distributed as food supplements.

For manufacturing process yohimbine can either be obtained from the bark or chemically synthesised. Dried bark from the twigs and trunk of the Pausinystalia yohimbe is used in its entirety, cut up or ground into powder to be used as pharmaceutical products. A real problem for the consumers’s safety is that there is no information on standardisation of the extracts for use in food supplements with regard to the ratio of extracted material to starting material, extraction solvent, and content of biologically active ingredients [9]. In many food supplements the qualitative and quantitative composition of their components is not declared. According European Pharmacopoeia yohimbine hydrochloride should be kept in air tight containers, protected from light. In fact no data are available on the stability of Yohimbe bark or preparations such as extracts thereof in food supplements. The strong analytical control of these products is essential for customer safety. Unfortunately the analytical control is not obligatory for the food supplements. A variety of techniques have been used successfully to measure yohimbine including: GC-MS, HPLC with UV, fluorescence or amperometric detection [10-14] and also TLC determination but there are no official inter-
laboratory validated methods for determining the levels of yohimbine and accompanying alkaloids in food supplements.

Materials and Methods

Materials

I) Reference standard: Yohimbine with purity > 98% (Sigma Aldrich).

II) Reagents: Acetonitrile for HPLC ≥ 99.93% (Sigma Aldrich); Methanol for HPLC ≥ 99.9% (Sigma Aldrich); Ammonium formate for HPLC ≥ 99.0% (Sigma Aldrich), distilled water.

III) Samples: Samples were purchased from dietary supplement stores and pharmacies.

Sample 1-capsules 500 mg. Content: yohimbine 500 mg.

Sample 2-capsules 800 mg. Content: L-arginine 250 mg, Yohimbe extract 50 mg (10% extract=5 mg yohimbine), ginseng 150 mg, Tribulus terrestris 150 mg.

Sample 3-capsules 650 mg. Content: Yohimbe 50 mg (10% extract=5 mg yohimbine); Ginko biloba extract-40 mg; Vitamin E-40 mg; Vitamin B6-10 mg.

Methods

UHPLC/MS method: I) Instrumentation: “Accela UHPLC” with mass spectral detection HRMS “Q-Exactive” with H-ESI-interface (“Thermo Fisher Scientific”, Waltham, MA, USA) and columns “Kynetex” RP C8 50 × 3 mm × 2.6 μm type “core-shell” and “Poroshell” RP CE18 150 × 3 mm × 2.7 μm type “core-shell”; ultrasonic bath (Branson Wilmington, NC, USA); apparatus for ultra-pure water: “Milli-Q”, “Milipore” (Bedford, MA, USA) and “Elga” (VWR International, Randor, PA, USA).

II) Chromatographic conditions: Mobile phase A: 10% acetonitrile/5% methanol/75% water

Mobile phase B: 30% acetonitrile/20% methanol/50% water

Mobile phase C: 0.2 M ammonium formate, pH=6.4

Gradient: from 95% A/5% B to 95% C/5% B over 10 min linear gradient;

15 min 95% B/5% C, isocratic;

5 min 95% A/5% C, isocratic.

III) Sample preparation: Standard solution of yohimbine: 5.0 mg of referent substance yohimbine with purity > 98% (Sigma Aldrich) was weighed in a volumetric flask (10 ml). The substance was dissolved in 5 ml of the component “B” of the mobile phase and sonificated in an ultrasonic bath for 5 min. We have added to the solution 5 ml of component “A” of the mobile phase. The obtained standard solution has a concentration of 500 μg/ml and is stable upon storage in a refrigerator (4 ± 8°C) for more than 10 days. From the basic standard solution by dilution with component “A” of the mobile phase we have obtained working standard solutions with a concentration of 200 μg/ml, 100 μg/ml, 20 μg/ml, 10 μg/ml, 2 μg/ml, 1 μg/ml, 0.2 μg/ml and 0.02 μg/ml.

Sample 1 solution: 20 mg of the contents of the capsule are dissolved in 250 ml acetonitrile/water (the obtained solution has a concentration of 80 μg/ml);

Sample 2 solution: 37 mg of the contents of the capsule are dissolved in 250 ml acetonitrile/water (the solution has a concentration of 148 μg/ml).

Sample 3 solution: 82 mg of the content of the tablet is dissolved in 250 ml acetonitrile/water (the solution has a concentration of 328 μg/ml).

Results and Discussion

For optimization of the conditions of ionization we have obtained chromatograms of standard solution of yohimbine in positive ion mode and in negative ion mode (Figure 1). In the both modes of ionization we have observed formation of molecular ions, respectively (M-H)- with m/z 353.19 and (M +H)+ at m/z 355.20. Positive ion and negative ion mode were compared. We have found that ionization in positive mode offers a significantly higher sensitivity, with also higher selectivity. We have performed the analyses of the samples in positive ion mode with monitoring of the molecular ion of yohimbine with m/z 355.20 (Figure 2). We have found that the tree dietary supplements do not contain yohimbine in the concentration specified on the label (Table 1). These products invariably display incorrect or incomplete information on their packaging, increasing risks to consumers.

We have performed a validation of the described method.
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Figure 2. Chromatograms of sample 1, 2, 3 obtained with UHPLC-mass spectral detection in positive ion mode.

Figure 3. Chromatogram of standard solutions of yohimbine in concentration 0.2 μg/ml in negative ion mode.

Figure 4. Chromatogram of standard solutions of yohimbine in concentration 0.02 μg/ml in positive ion mode.

Selectivity of the method
Figures 3 and 4 show chromatograms of standard solutions of yohimbine at a concentration 0.2 μg/ml and 0.02 μg/ml, respectively resulting in standby negative ionization and in positive ionization mode.

Table 1. Total content of yohimbine in the analysed food supplements.

<table>
<thead>
<tr>
<th>Sample</th>
<th>Content of yohimbine (Declared on the label)</th>
<th>Content of yohimbine</th>
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<td>60 mg/1 capsule</td>
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<tr>
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<td>5 mg/1 capsule</td>
<td>6.1 mg/1 capsule</td>
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<td>Sample 3</td>
<td>5 mg/1 capsule</td>
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Table 2. Reproducibility of the method.

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<th>m</th>
<th>σ</th>
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</table>

Figure 5. Series of chromatograms of standard solutions of yohimbine (in concentrations from 0.02 to 200 μg/ml). Mass spectral detection in negative ion mode.

Linearity of the method
We have obtained chromatograms of a series of standard solutions of yohimbine (concentration from 0.02 to 200 μg/ml) in the monitoring of the molecular ion of yohimbine in the negative ionization mode and in the positive ionization mode (Figures 5 and 6). In both modes standard graphiques show...
linearity only in the concentration range up to 50 μg/ml. Figures 7 and 8 represent series chromatograms of standard solutions of yohimbine obtained with mass spectral detection in the positive ionization mode. Standard graphiques obtained with mass spectral detector show linearity in the concentration range 0-50 μg/ml.

Reproducibility was determined by introducing a 6-fold under the described chromatographic conditions and monitoring of the molecular ion of yohimbine in mode monitoring positive ions. Peak area of the chromatograms (S), the average value (x̄), standard deviation (σ) and the relative standard deviation (RS) are presented in Table 2.

Figure 7. Series of chromatograms of standard solutions of yohimbine (in concentrations from 0.02 to 200 μg/ml). Mass spectral detection in positive ion mode.

Conclusion
The development and validation of a rapid qualitative and quantitative method based on an UHPLC/MS allowed the identification of the target compound-yohimbine in mono and multi-botanical preparations, distributed as food supplements. The described method shows high sensitivity and selectivity. We have found that in the mode of positive ion monitoring the sensitivity and selectivity are higher in comparison with detection at monitoring of negative ions. The modified method was applied for the determination of yohimbine in food supplements. The analyses showed that in some of the food supplements the real content of yohimbine is different from the declared that indicate the importance of the analytical control of food supplements.

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References

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