Survey on the Quality of the Top-Selling European and American Botanical Dietary Supplements Containing Boswellic Acids

Authors

Jürgen Meins1, Christian Artaria2, Antonella Riva2, Paolo Morazzoni2, Manfred Schubert-Zsilavecz1, Mona Abdel-Tawab1

Affiliations

1 Central Laboratory of German Pharmacists, Eschborn, Germany
2 Indena S.p.A., Milano, Italy
3 Department of Pharmaceutical Chemistry, Goethe-University Frankfurt, Frankfurt am Main, Germany

Abstract

In consideration of the increasing popularity of frankincense and the widely published quality problems associated with botanical dietary supplements, a survey was conducted for the first time on the quality of frankincense containing botanical dietary supplements. Six US products representing 78% of the units sold and 70% of the market value, and 11 European products representing 30% of the units sold and 40% of the market value were tested for their boswellic acid composition profile, label compliance, and claimed health benefits. Special focus was also set on the statements made with regard to the frankincense applied. Only five products out of seventeen disclosed all relevant information for the Boswellia extract, mentioning the species, the part of plant used, and the boswellic acid content. Whereas all products but one claimed to use Boswellia serrata, three products did not mention the resin as the part applied and 10 products did not declare the boswellic acid content. Apart from the different boswellic acid composition determined with a sensitive LC/MS method, 41% of the products did not comply with the label declaration. Hence, one product from Italy did not contain any of the six characteristic boswellic acids (KBA, AKBA, αBA, AαBA, AβBA) and another US product contained only traces, suggesting the absence of frankincense or the use of Boswellia frereana instead of B. serrata. In another product, the ratios of the individual boswellic acids were different from B. serrata gum resin, indicating the use of another species such as Boswellia sacra or Boswellia carterii. Furthermore, two products revealed different boswellic acid contents from those declared on the label. Further, two products did not declare the use of manipulated Boswellia gum resin extract being enriched in acetyl-11-keto-boswellic acid content reaching up to 66%. In addition, consumers could be misled by outdated literature or references to in vitro studies performed at dosages that can never be achieved in humans following oral administration.

In summary, this survey reveals that in spite of increased regulations on botanical dietary supplements, the problem of mislabeling still exists and needs to be addressed by the manufacturers, so that consumers get greater confidence in the botanical dietary supplements they use.

Abbreviations

α-BA: alpha-boswellic acid
AαBA: acetyl- alpha-boswellic acid
AβBA: acetyl- beta-boswellic acid
AKBA: acetyl-11-keto-boswellic acid
β-BA: beta-boswellic acid
BAs: boswellic acids
catG: cathepsin G
IL: interleukin
KBA: 11-keto-boswellic acid
mPGES-1: microsomal prostaglandin E synthase-1
SIM: single ion mode

Introduction

With millions of people using medicinal plants worldwide, the market of herbal supplements is witnessing a steady growth. Recent data indicate that in 2012, 17.9% of all US adults used botanical supplements [1]. In Germany, 90% of the people use natural medicines at some time during their life and over 50% of the population has done so in other European countries [2, 3]. Thus, the glob-
al market for herbal dietary supplements or phytomedicines, estimated at approximately US$ 60 billion in 2000, is expected to increase dramatically, reaching US$107 billion by the year 2017 [4].

At the same time, episodes of contamination (with insecticides, pesticides, synthetic drugs, heavy metals) or adulteration (substituting one plant for another either purposefully or through misidentification) have been frequently reported, resulting in increased concerns about the safety, effectiveness, and quality of herbal products [5]. A study on selected commercial ginseng products marketed as botanical supplement in North America showed that the ginsenoside contents of 232 Panax ginseng C.A. Mey. (Araliaceae) and 81 Panax quinquefolius L. supplements ranged from 0.00% to 13.54% and from 0.009% to 8.00%, respectively, and that 26% of these products did not meet label claims [6]. Studies on the quality of St. John’s wort (Hypericum perforatum L., Hypericaceae) products showed hypericin content ranging from 22% to 140% of the label claim [7]. Similarly, silymarin, an extract from the seeds of milk thistle [Silybum marianum (L.) Gaertn., Asteraceae], was detected at 58–116% of the labeled claim [8].

Aside from ginseng, St. John’s wort, and milk thistle, frankincense is counted among the well-established botanical dietary supplements. This is mainly attributed to the anti-inflammatory properties of frankincense [9] and the growing prevalence of symptoms like joint pain and stiffness in a progressively ageing Western population [10]. Hence, it was shown that a number of pivotal enzymes in inflammation like 5-LO, catG, and mPGES-1 as well as NF-xB and several cytokines like TNFα, IL-1β, and IL-6 are inhibited by BAs, the main active ingredients of frankincense (Fig. 1). Whereas in the past the anti-inflammatory effects were mainly attributed to the inhibition of 5-LO by AKBA [11], recent research revealed the relevance of the whole fraction of triterpenoid acids and the involvement of much more molecular targets than only 5-LO, especially catG and several cytokines like TNFα, IL-1β, and IL-6 are inhibited by BAs, the main active ingredients of frankincense (Fig. 1). Whereas in the past the anti-inflammatory effects were mainly attributed to the inhibition of 5-LO by AKBA [11], recent research revealed the relevance of the whole fraction of triterpenoid acids and the involvement of much more molecular targets than only 5-LO, especially catG and several cytokines like TNFα, IL-1β, and IL-6 are inhibited by BAs, the main active ingredients of frankincense (Fig. 1). Whereas in the past the anti-inflammatory effects were mainly attributed to the inhibition of 5-LO by AKBA [11], recent research revealed the relevance of the whole fraction of triterpenoid acids and the involvement of much more molecular targets than only 5-LO, especially catG and several cytokines like TNFα, IL-1β, and IL-6 are inhibited by BAs, the main active ingredients of frankincense (Fig. 1). Whereas in the past the anti-inflammatory effects were mainly attributed to the inhibition of 5-LO by AKBA [11], recent research revealed the relevance of the whole fraction of triterpenoid acids and the involvement of much more molecular targets than only 5-LO, especially catG and several cytokines like TNFα, IL-1β, and IL-6 are inhibited by BAs, the main active ingredients of frankincense (Fig. 1). Whereas in the past the anti-inflammatory effects were mainly attributed to the inhibition of 5-LO by AKBA [11], recent research revealed the relevance of the whole fraction of triterpenoid acids and the involvement of much more molecular targets than only 5-LO, especially catG and several cytokines like TNFα, IL-1β, and IL-6 are inhibited by BAs, the main active ingredients of frankincense (Fig. 1). Whereas in the past the anti-inflammatory effects were mainly attributed to the inhibition of 5-LO by AKBA [11], recent research revealed the relevance of the whole fraction of triterpenoid acids and the involvement of much more molecular targets than only 5-LO, especially catG and several cytokines like TNFα, IL-1β, and IL-6 are inhibited by BAs, the main active ingredients of frankincense (Fig. 1). Whereas in the past the anti-inflammatory effects were mainly attributed to the inhibition of 5-LO by AKBA [11], recent research revealed the relevance of the whole fraction of triterpenoid acids and the involvement of much more molecular targets than only 5-LO, especially catG and several cytokines like TNFα, IL-1β, and IL-6 are inhibited by BAs, the main active ingredients of frankincense (Fig. 1). Whereas in the past the anti-inflammatory effects were mainly attributed to the inhibition of 5-LO by AKBA [11], recent research revealed the relevance of the whole fraction of triterpenoid acids and the involvement of much more molecular targets than only 5-LO, especially catG and several cytokines like TNFα, IL-1β, and IL-6 are inhibited by BAs, the main active ingredients of frankincense (Fig. 1).

Results and Discussion

As BAs represent the major pharmacologically active ingredients of frankincense, 17 of the most popular botanical dietary supplements in the American and European supermarkets/outlets claiming to contain frankincense have been analyzed for their BA composition profile using a sensitive LC/MS method. Special focus was also set on the statements made on the label or on the leaflet with regard to the frankincense applied. Fourteen products contained a mixture of Boswellia extract with up to 10 other ingredients comprising vitamins, minerals, glucosamine, methylsulfonylmethane (MSM), collagen, condroitin sulphate, or other plant extracts. Only three products were solely composed of Boswellia extract.

An overview on the products included in this survey is given in Table 1 and the BAs composition profile of the individual products is presented in Fig. 2. Given that for the production of an extract several Boswellia species may be used, the Boswellia species and the part of the plant applied as well as the content of BAs is of upmost importance. In fact, five products out of seventeen (3, 5, 10, 12, 16) disclosed all this information. Whereas all products but one (7) claimed to use B. serrata, two products (6, 8) did not declare the part of the plant applied nor the BAs content, eight products (1, 2, 4, 9, 11, 13, 14, 15) did not make declarations on the BAs content, but on the part used, and one product (17) did not declare the part of the plant applied, but the BAs content. Moreover, the composition profile revealed that the BAs differed greatly, suggesting the use of other Boswellia sources than that declared on the package in the case of seven products. According to a previously carried out analysis of different authenticated gum resins of Boswellia species by Frank and Unger, B. serrata contains the complete spectrum of the characteristic six BAs, whereas B. frereana does not contain appreciable amounts of the six BAs at all. On the other hand, the ratio of the signal intensities of the non-acetylated (α-BA, β-BA) to the acetylated BAs (AcαBA and AcβBA) is < 1 in the case of frankincense extracts from B. carterii and B. sacra, but is > 1 in the case of frankincense extracts from B. serrata [16]. Based on that background, one product from Italy (11) not containing any of the six BAs (KBA, AKBA, α-BA, β-BA, AcαBA, and AcβBA) at all and another product from USA (2)
Table 1  Overview of the top-selling American and European botanical dietary supplements included in the survey

<table>
<thead>
<tr>
<th>Product</th>
<th>Country Manufacturer/Distributor</th>
<th>Claimed health benefits</th>
<th>Dosing</th>
<th>Boswellia extract [mg] per dosage form</th>
<th>Labeled Boswellia extract</th>
<th>Determined total boswellic acid content [%]</th>
<th>Statement on Boswellia extract</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>USA</td>
<td>Healthy joints and cartilage improved joint mobility**</td>
<td>2 tablets daily</td>
<td>50</td>
<td>5-Loxin® Advanced B. serrata Extract (resin)</td>
<td>31.43 (AKBA: 28.51)</td>
<td>+ + –</td>
</tr>
<tr>
<td>2</td>
<td>USA</td>
<td>Support of the immune system’s balanced response to normal internal metabolic stress as well as to typical wear and tear of everyday life**</td>
<td>2 capsules daily</td>
<td>250</td>
<td>Boswellia extract (B. serrata) (Gum resin) (min. 65% Organic acids including BAs)</td>
<td>0.81</td>
<td>+ + –</td>
</tr>
<tr>
<td>3</td>
<td>India/USA</td>
<td>Support of normal joint function, normal flexibility of the body and the body’s natural inflammation response**</td>
<td>2 × 2 capsules daily</td>
<td>125</td>
<td>Indian Frankincense standardized extract (oleo gum resin) (B. serrata Roxb.) (60% BAs)</td>
<td>40.0</td>
<td>+ + +</td>
</tr>
<tr>
<td>4</td>
<td>USA</td>
<td>Increase of mobility, flexibility, and range of motion in insensitive joints; improvement in joint comfort*</td>
<td>1 capsule daily</td>
<td>100</td>
<td>5-Loxin® Advanced (B. serrata Extract [resin])</td>
<td>34.0 (AKBA: 29.6)</td>
<td>+ + –</td>
</tr>
<tr>
<td>5</td>
<td>USA</td>
<td>No indication</td>
<td>1 capsule two times a day</td>
<td>300</td>
<td>Boswellia (B. serrata) (resin extract) (Guaranteed 65% BAs)</td>
<td>43.5</td>
<td>+ + +</td>
</tr>
<tr>
<td>6</td>
<td>Spain</td>
<td>Contribution to good flexibility and well-being of joints</td>
<td>1–2 capsules daily</td>
<td>100</td>
<td>Boswellia extract (B. serrata)</td>
<td>91.1 (AKBA: 66.5)</td>
<td>+ – –</td>
</tr>
<tr>
<td>7</td>
<td>USA</td>
<td>Support of joint health and mobility**</td>
<td>1 tablet three times daily</td>
<td>307</td>
<td>Boswellia extract (tree resin) 65% BAs</td>
<td>49.7</td>
<td>– + +</td>
</tr>
<tr>
<td>8</td>
<td>Germany</td>
<td>No indication</td>
<td>1 capsule daily</td>
<td>350</td>
<td>B. serrata extract (standardized to 85% total acids)</td>
<td>44.6</td>
<td>+ – –</td>
</tr>
<tr>
<td>9</td>
<td>Italy</td>
<td>Dietary supplementation</td>
<td>1–2 sachets daily</td>
<td>400</td>
<td>Boswellia dried resin extract (B. serrata Roxb.) Fitosoma® (Casperome®)</td>
<td>16.80</td>
<td>+ + +</td>
</tr>
<tr>
<td>10</td>
<td>Italy</td>
<td>Physiological comfort of joints</td>
<td>1 sachet daily</td>
<td>200</td>
<td>B. serrata e. s. (B. serrata Roxb., resin) Total boswellic acid content 70 mg</td>
<td>70.1</td>
<td>+ + +</td>
</tr>
<tr>
<td>11</td>
<td>Poland/Italy</td>
<td>Physiological effects against states of localized tension</td>
<td>1 soft capsule daily</td>
<td>36</td>
<td>B. serrata Roxb. (B. serrata) resin dried extract</td>
<td>0.0</td>
<td>+ + –</td>
</tr>
<tr>
<td>12</td>
<td>Italy</td>
<td>Support of normal joint function</td>
<td>1 sachet daily</td>
<td>200</td>
<td>Boswellia dried extract 30% AKBA (B. serrata Roxb. resin)</td>
<td>66.2 (AKBA: 49.5)</td>
<td>+ + +</td>
</tr>
<tr>
<td>13</td>
<td>Italy</td>
<td>No indication</td>
<td>1 tablet two times daily</td>
<td>100</td>
<td>Casperome® (B. serrata Roxb. ex. Co-lebr. Extract, oleoresin)</td>
<td>38.5</td>
<td>+ + –</td>
</tr>
<tr>
<td>14</td>
<td>Italy</td>
<td>Against states of localized tension</td>
<td>2–6 tablets daily</td>
<td>25</td>
<td>B. serrata Roxb. titrated resin dried extract Titrated alpha keto boswellic acid</td>
<td>50.6</td>
<td>+ + –</td>
</tr>
<tr>
<td>15</td>
<td>Italy</td>
<td>No indication</td>
<td>1 tablet three times daily</td>
<td>75</td>
<td>B. serrata E.S., resin</td>
<td>33.1</td>
<td>+ + –</td>
</tr>
<tr>
<td>16</td>
<td>Italy</td>
<td>Inhibition of inflammatory mediators</td>
<td>2 tablets daily</td>
<td>100</td>
<td>AKBAMAX™ (B. serrata Roxb. gum) dried extract corresponding to 10 mg acetyl-keto-β-boswellic acid (AKBA)</td>
<td>85.0 (AKBA: 17.1)</td>
<td>+ + +</td>
</tr>
<tr>
<td>17</td>
<td>Romania</td>
<td>Support of elasticity and flexibility of joints</td>
<td>3 tablets daily</td>
<td>50</td>
<td>Boswellin® (B. serrata extract; min. 70% BAs min. 20% acetyl-β-boswellic acid</td>
<td>41.7 (acetyl-β-boswellic acid: 7.7)</td>
<td>+ – +</td>
</tr>
</tbody>
</table>

* Mentioned in the supplement facts box or anywhere else on the package or in the leaflet; ** Indication on US products that these statements have not been evaluated by the Food and Drug Administration and that this product is not licensed to diagnose, treat, cure, or prevent diseases.
containing only traces suggests, in contradiction to what was declared in the label, the absence of *B. serrata* or the use of another *Boswellia* species such as *B. frereana*. Another product (10) displaying all basic label information revealed a ratio of non-acetylated to acetylated BAs < 1, indicating the use of *B. carterii* or *B. sacra* [16]. In addition, based on the applied LC/MS analysis, two products (16, 17) revealed different BA contents from those declared on the label. Thus, only 7.7% AFBA was determined in one product (17), corresponding to half of the amount mentioned on the label. The other product (16) claimed to contain 10 mg AKBA, whereas an almost twofold higher amount of 17.1 mg AKBA could be determined. Furthermore, two products (6, 12) did not declare the use of manipulated *Boswellia* gum resin extract being enriched in AKBA content reaching up to 66%. However, it should be noted in this context that the observed deviations from the declaration on the label may not always be attributed to deliberate adulteration, but in many cases results from the confusing characterization of *Boswellia* gum resin extracts and/or insufficient quality control measures applied by manufacturers. Thus, some *Boswellia* gum resin extracts are characterized by the organic acid content, others by the total acid, and again others by the BA content. As the BAs represent only a minor portion of the total acid content, a product claiming to contain 65% total acids or organic acids is not equivalent to a product claiming 60% BAs. Moreover, different analytical methods may result in different boswellic BA content as demonstrated in the product information for AKBAmax™ [17], a commercially available *Boswellia* extract enriched in AKBA. Hence, the BA content was determined to be 70% when quantified with a titration method, but did not exceed 35–45% when quantified with HPLC. In the first place, this may be attributed to the insufficient specificity of titration methods compared to chromatographic methods. Moreover, the varying spectrum of analytes covered by different analytical methods represents another reason for varying BA contents. This is also the reason why the total BA content determined in the present study turned out to be generally lower than the BA content labeled on the different products. It may be generally assumed that the declaration on the label is based on HPLC analysis, including more BAs than the six major BAs determined by the LC/MS method in this survey, leading, in consequence, to higher BA contents. Because of this diversity in characterizing *Boswellia* extracts and the varying analytical spectrum of the applied methods for quantification, it is very important for the manufacturer to get detailed information on the analytical methods and to ensure that a valid method is used to characterize the *Boswellia* extract in order to be able to properly evaluate its quality.

In the frame of misleading composition declarations, consumers find it increasingly difficult to actually estimate the quality and efficacy of dietary *Boswellia* products. Hence, they may correlate better efficacy with an increased acid content, although the declared acid content should not necessarily reflect the pharmacologically active BAs. They may be also misled by outdated literature or references to in vitro studies performed at dosages that can never be achieved in humans following oral administration. This was particularly evident for some studies cited [11, 18] in the case of product 16 using an AKBA-enriched extract. In fact, it was recently shown that AKBA is extensively bound to albumin in plasma, never approaching the concentrations needed to modulate the various targets of BAs even when large dosages of extracts were administered [9]. Furthermore, AKBA failed to inhibit 5-LO product formation in human whole blood, showing activity only in enzyme assays [19]. In addition, recent research revealed little relevance for 5-LO in the inflammatory response associated with osteoarthritis, the major clinical indication of *Boswellia* extracts [20, 21].

All products included in this survey are sold as botanical dietary supplements and are not intended to diagnose, treat, cure, or prevent any disease. Moreover, all US products clearly indicate that the statements made have not been evaluated by the Food and Drug Administration. Nevertheless, consumers should have access to reliable and accurately labeled botanical dietary supplements that do not claim effects that are hardly achieved. Surely many high-quality botanical dietary supplements are produced by reputable companies, but this cannot be always guaranteed, as demonstrated in the present survey. Thus, quality may still represent a paramount and complex issue when dealing with botanicals, as shown by the great differences in the chemical composition of BAs. Although all products but one declared the use of *B. serrata* extract, seven products out of seventeen have been identified where the BA content did not comply with the label claim, or other undeclared *Boswellia* extracts have been used instead of *B. serrata* or the *Boswellia* extract was even totally absent. Thus, in spite of increased regulations on botanical dietary supplements, the problem of mislabeling still exists and needs to be addressed by the manufacturers, so that consumers get greater confidence in the botanical dietary supplements they use.

**Materials and Methods**

**Botanical dietary supplements**

The six US top-selling *Boswellia* dietary products were selected on the basis of the data from the market research company SPINS. One product was purchased from a food, drug, and mass merchandiser store, four from Natural Health Food Stores, and two from the Internet at amazon.com. The 11 European top-selling products were selected based on the data of the market research company IMS OTC and were purchased in local pharmacies. All tested products were assigned voucher numbers and representative voucher specimens have been deposited in the Central Laboratory of German Pharmacists, Eschborn, Germany.
Chemicals and reagents
BAs (α-BA, β-BA, AαBA, AKBA, A®BA, KBA) (purity > 99%) were purchased from Phytoplan. Methanol of LC/MS quality was purchased from Carl Roth GmbH, ammonium formiate from Alfa Aesar GmbH, and water Emsure® p. a. from Merck KGaA.

Standard preparation
Stock standard solutions of each BA were prepared by weighing into a 20-mL volumetric flask 20 mg of each BA standard and diluting it with 20 mL methanol to yield a concentration of 1 mg/mL of each BA, respectively. Mixed spike solutions were prepared by mixing the appropriate amount of each boswellic stock standard solution with methanol to yield spike solutions K1 (4 µg/mL), K2 (12 µg/mL), and K3 (24 µg/mL).

Sample preparation
The contents of 10 tablets/capsules/sachets were pulverized and mixed well. An equivalent of 100 mg Boswellia extract was weighed in a 50-mL centrifuge tube (Eco, PP, Roth, Art. AN78.1) and shaken with 20 mL of methanol for 60 min at 200 rpm on a vertical shaker, followed by treatment in an ultrasonic bath for 30 min and centrifugation for 10 min at 2000 rpm. Four aliquots of 100 µL of the clear supernatant were then transferred into four 10 mL volumetric flasks, respectively. In order to overcome any possible matrix interferences, the standard addition method was applied for the quantification of the BAs. For that purpose, the three spike solutions, K1, K2, and K3, were added to the three aliquots of the clear supernatant, respectively, to yield three samples spiked with 4 µg/mL, 12 µg/mL, and 24 µg/mL, respectively. To one aliquot, no spike solution was added. Finally, 20 µL of each sample solution were injected into the chromatographic system.

Liquid chromatography-mass spectrometry analysis
Apart from several analytical methods described in the literature for the quantification of BAs in Boswellia extracts and plasma, one HPTLC and one HPLC method have been reported for the analysis of KBA and AKBA in market formulations [22, 23]. Moreover, only one HPLC analysis was devoted to the determination of the non-ketylated BAs, β-BA, α-BA, AβBA, and AαBA, besides KBA and A®BA in market formulations before [24]. In brief, however, the non-ketylated BAs do not possess a chromophore, the HPLC analysis of the complex extracts had to be conducted at a rather unsuitable wavelength of 210 nm. In this regard, the application of LC/MS is advantageous because all BAs, even those without a chromophore, may be clearly identified via their typical m/z values. Based on that background, the chromatographic and mass spectrometric parameters applied in a previously developed LC-MS method for the simultaneous determination of the six BAs in plasma have been applied for the analysis of BAs in the present study after its suitability had been assessed [25]. In brief, liquid chromatography was performed on an Agilent 1200 series equipped with a gradient pump with a vacuum degasser, an autosampler, and a column oven. A Hypersil™ BDS RP C18 column (100 × 4 mm; 3 µm; Thermo scientific) and an upstream Gemini SecurityGuard™ cartridge (Phenomenex; 4 × 3 mm) were used for chromatography. Separation was achieved using a gradient program starting with 90% mobile phase A (methanol: water 90:10, 400 mg/L ammonium formate) and 10% mobile phase B (methanol: water 80:20, 400 mg/L ammonium formate), changing to 100% mobile phase A within 20 min. This was kept constant for 14 min before returning to the initial conditions within 1 min. The total run time was 35 min at a flow rate of 0.4 mL/min. The column oven was set to 40°C and the autosampler was kept at room temperature. MS analysis was performed in the negative SIM on an Agilent Triple Quadrupole LC/MS 6410 series (Agilent Technologies) equipped with an ESI source. The monitoring mass was set at m/z 469.3 for KBA, 511.5 for AKBA, 455.5 for β-BA and α-BA, and 497.4 for AαBA and AαBA. The dwell time was chosen to be 200 ms. MassHunter® software was used for data acquisition and processing.

The fitness of the applied LC/MS method was assessed by judging the linearity, accuracy, and precision. The identity of the BAs was confirmed by their retention times and the standard addition approach. As can be seen exemplary in Fig. 3, no interfering peaks are found in the chromatogram of the tested products, indicating that the formulation excipients do not interfere in the estimation of BAs. The accuracy of the method was evaluated by determining the recovery of each added BA standard at each concentration level in every product tested in addition to the mean recovery of each BA extending over all test products. The precision was assessed through the repetitive use of the same analytical procedure over the different matrices of all products (n = 17) on two consecutive days. Taking into consideration that no placebo matrix was available, the LOD for each BA was calculated as three times the baseline noise over three consecutive days. Taking into consideration that no placebo matrix was available, the LOD for each BA was calculated as three
times the corresponding standard deviation of the mean intercept divided by the slope of the mean standard addition calibration curve. The LOQ for each BA was calculated as 10 times the corresponding standard deviation of the mean intercept divided by the slope of the mean standard addition calibration curve. The results are summarized in Table 2. Both the high accuracy reflected in the high recovery rates as well as the good precision at each concentration level for each BA in different matrices verify the ruggedness of the applied method and its independence from minor deviations in the experimental conditions, especially in the matrix composition of the different products tested.

Conflict of Interest

P.M. and A.R. are employed by Indena, the producer of Casperone®, C.A. was employed by Indena at the time of the study. The other authors have no conflict of interest.

References

5. Moraes CDF, Still DW, Lum MR, Hirsch AM. DNA-based authentication of botanicals and plant-derived dietary supplements: where have we been and where are we going? Planta Med 2015; 81: 687–695