

# A GREEN ANALYTICAL METHOD FOR THE DETERMINATION OF HYDROXYETHOXYPHENYL BUTANONE IN COSMETIC PRODUCTS

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**HEPB** 

Hydroxyetoxyphenyl

butanone

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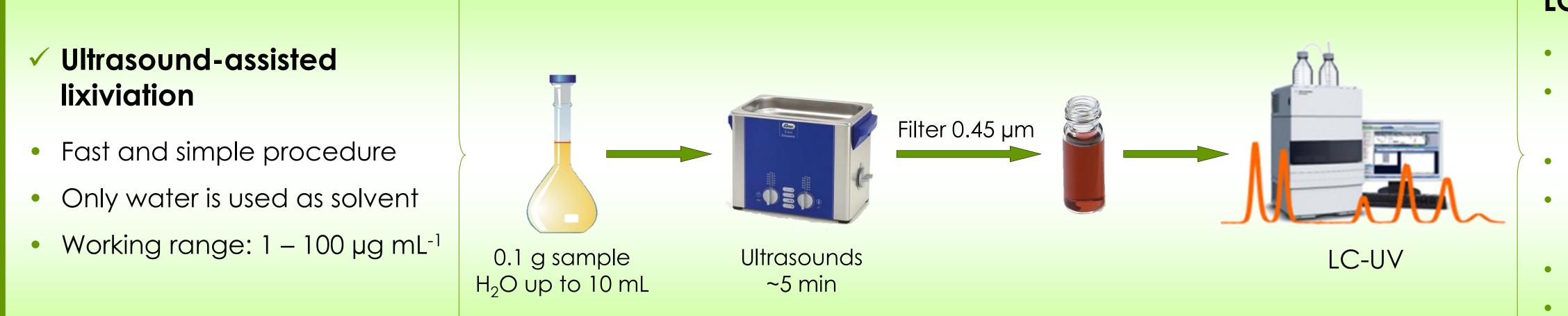
# INTRODUCTION



- HEPB is not regulated yet by the current European Regulation on Cosmetic Products
- However, according to the Scientific Committee on Consumer Safety, only a maximum concentration of 0.7 % can be considered safe due to potential toxicity for repeated exposure
- There are no published or official methods to quantify HEPB in cosmetic samples

The **aim** of this work is to develop and validate an accurate and fast analytical method to determine HEPB in different types of cosmetic products, with good analytical features and environmentally-friendly characteristics [1]

### EXPERIMENTAL



### **LC-UV** Conditions

- Injection volume: 20 µL
- Column: LiChrospher<sup>®</sup> 100 RP-18 (250x4 mm, 5 µm)
- Column temperature: 35 °C
- Mobile phase: EtOH : 1 % AcOH solution (23:77)
- Flow rate: 0.8 mL min<sup>-1</sup>
- Detection wavelength: 279 nm

## **RESULTS AND DISCUSSION**

Analytical features of the proposed method

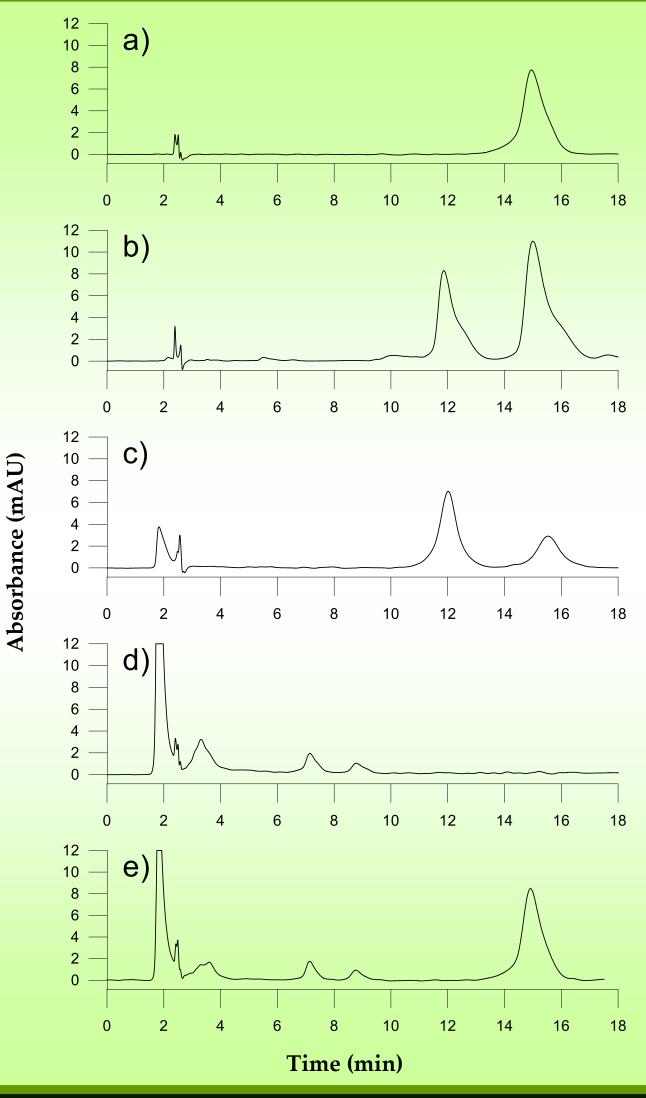
Analysis of commercial cosmetic samples

 $\checkmark$  High level of linearity (R<sup>2</sup> > 0.9997) at least to 100 µg mL<sup>-1</sup>

 $\checkmark$  Limit of detection (3 S/N): 0.3 µg mL<sup>-1</sup> (30 µg g<sup>-1</sup> in sample) Limit of quantification (10 S/N): 0.9  $\mu$ g mL<sup>-1</sup> (90  $\mu$ g g<sup>-1</sup> in sample)

proposed method ✓ The Was satisfactorily applied 10 7 commercial cosmetic samples

✓ Different types of cosmetics:



- $\checkmark$  Intra-day and inter-day repeatability, expressed as RSD (%):
  - 2.5 and 1.2 % at 5  $\mu$ g mL<sup>-1</sup>
  - 0.2 and 1.2 % at 25  $\mu$ g mL<sup>-1</sup>
  - 0.9 and 4.7 % at 50  $\mu$ g mL<sup>-1</sup>
- ✓ **Relative error** below 5.6 % in the analysis of a laboratory-made cosmetic cream sample
- $\checkmark$  Quantitative **recovery values**, 86 103 %, in the analysis of spiked samples

✓ ACCURACY

PRECISION

- - 2 liquid hand soaps
  - 2 sunscreens
  - 1 shampoo
  - 1 moisturizing cream
  - 1 make-up
- ✓ HEPB found the WQS In moisturizing (0.083 ± cream 0.002 % w/w) according to its label.



The method is rapid and provides a good analytical performance, allowing the quality control of cosmetic products containing HEPB as alternative preservative agent.

The method is in accordance with the principles of the so-called Green Analytical Chemistry, as it is harmless to the operator and the environment.

Cromatograms obtained applying the proposed LC-UV method to: a) standard solution (25  $\mu$ g mL<sup>-1</sup>), b) laboratory-made cream, c) moisturizing cream, d) sunscreen, e) sunscreen spiked with HEPB (25  $\mu$ g mL<sup>-1</sup>).



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### Consulta otras comunicaciones presentadas por el GICAPC en la XXII **Reunión de la Sociedad Española de Química Analítica:**

Determination of nitro musks in environmental waters by stir bar sorptive dispersive microextraction followed by thermal desorption-gas chromatography-mass spectrometry. J.L. Benedé, A. Chisvert, A. Salvador. Flash Communication. Ver comunicación.

Stir bar sorptive-dispersive microextraction mediated by a magnetic

nanoparticles-metal organic framework composite for the determination of *n-nitrosamines in cosmetic products*. P. Miralles, I. Van Gemert, A. Chisvert, A. Salvador. Flash Communication. Ver comunicación.

Development of an analytical method for the determination of acrylamide in cosmetic products based on dispersive liquid-liquid microextraction. L. Schettino, J.L. Benedé, A. Chisvert, A. Salvador. Flash Communication. Ver comunicación.

Determination of hydroxylated ingredients with preservative activity in cosmetic products by gas chromatography-mass spectrometry. C. Azorín, J.L. Benedé, A. Chisvert, A. Salvador. Ver comunicación.

A green analytical method for the determination of hydroxyethoxyphenyl butanone in cosmetic products. P. Miralles, J.L. Benedé, A. Mata-Martín, A. Chisvert, A. Salvador. Ver comunicación.

Determination of polycyclic aromatic hydrocarbons in cosmetics by stir bar sorptive dispersive microextraction and gas chromatography-mass spectrometry. Vállez-Gomis, J. Grau, J.L. Benedé, A. Chisvert, A. Salvador. Ver comunicación.

Reversed-phase dispersive liquid-liquid microextraction prior to liquid chromatography-tandem mass spectrometry for the determination of acrylamide in cosmetic products. L. Fernández, J.L. Benedé, A. Chisvert, A. Salvador. Ver comunicación.

Development of dispersive liquid-solid microextraction: application to the determination of cortisone and cortisol in human saliva. J. Grau, J.L. Benedé, A. Chisvert, A. Salvador. Ver comunicación.