

Chemistry Conference for Young Scientists Blankenberge, Belgium

— FEBRUARY 21-23 — — —

BOOK OF ABSTRACTS

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Analytical and Environmental Chemistry: Oral communications

Session 1 (21/02/2018 - 14:30)

Swati Chandrawanshi - Pt. Ravishankar Shukla University

Determination of nitrate by using ion-pair single-drop micro extraction (SDME) method with the attenuated total reflectance –Fourier transform infrared (ATR-FTIR) technique

Liselotte Neven - University of Antwerp

Detection of specific DNA sequences by photoelectrochemistry

Mathijs Baert - Ghent University

Investigating the potential for improved temperature responsive separations in liquid chromatography

Ravindra Hegade - Ghent University

Enhanced resolution of stereoisomers through Stationary phase optimized selectivity liquid and supercritical chromatography (SOS-LC and SOS-SFC)

Brecht Laforce - Ghent University

The Herakles 3D X-ray scanner: a novel tool for lab-based 3D analysis

Session 2 (22/02/2018 - 08:30)

Monica Hernandez Rodriguez - Universidad de Oriente

Evaluation of activation parameters of activated carbon from coffee and cocoa seed husk rests: carbon yields and Ni(II) adsorption capacity study

Myrthe Van Hal - University of Antwerp

Harvesting energy from air pollution with an un-biased gas phase photoelectrochemical cell

Camille Gaulier - Vrije Universiteit Brussel

The geochemical behaviour of trace metals in the surface water of the Belgian Coastal Zone

Adrian Frank Herbort - Universität Koblenz-Landau

Removal of Inert Organic Chemical Stressors (IOCS) from Wastewater by Adding Innovative Hybrid materials – Wasser 3.0

Session 3 (22/02/2018 - 10:10)

Saranya Thiruvottriyur Shanmugam - University of Antwerp

Wireless electrochemical sensor for on-site detection of ecstasy

Monika Strozynska - SAS Hagmann GmbH

A new derivatisation reaction for perfluorcarboxylic acids prior to GC/MS analysis

Ivan Bezruk - National University of Pharmacy

Using of the HPLC method for quantitative determination of active pharmaceutical ingredients in the process of the industrial synthesis



Using of the HPLC method for quantitative determination of active pharmaceutical ingredients in the process of the industrial synthesis

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Quality of medicines depends from many factors. One of the main and defining is quality of its components and first of all active pharmaceutical ingredients (API).

Manufacturers, who plan to develop API synthesis, have to carry out scientific researches, which would allow optimizing production process for providing quality of substances and the quality of the medicine as a whole.

Epilepsy is a neurological disease that affects about 9 people per thousand of population throughout the world. Nowadays, dibenzylamide of malonic acid (Dibamk) is one of the most promising substances that have anticonvulsant effect.

Developing of simple in using methods for analysis of the dibamk with needed effectiveness is necessary condition for possibility for medical application of the API. Also manufacturer should development methods of controlling the entry of starting substances (specifically benzyl amine and diethyl malonate) in the API for ensuring quality of the drug.

The "ProStar" Varian liquid chromatograph was used in the study. The chromatography was conducted in the following conditions: Waters XBridge® C18 column (150mm × 4.6 mm, 3.5 mm particle size); mobile phase A – phosphate buffer solution with pH 5.5 and mobile phase B [acetonitrile; the gradient program for chromatography was as follows: time (min) /%; mobile phase A: 0/90; $2/90 \rightarrow 35$; 5/35; $9/35 \rightarrow 90$; 12/90; flow rate – 1.2 ml/min; column temperature — 25 °C; detection was performed at the wavelength of 254 nm; injection size — 50μ L; run time — 15 min. In the specified conditions the retention time of main substance is about 8 minutes.

The HLPC method of quantitative determination of dibamk is a reliable and simple method with proper specificity. Therefore, it can be recommended for use in controlling of the yield of the product in the process of industrial synthesis as well as for the quantitative content of API in the substance.