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离子色谱法同时测定化妆品中的铵和6种烷基胺

钟志雄1*,李攻科2,朱炳辉1,罗志彬1,吴西梅1

1. 广东省疾病预防控制中心, 广东 广州 510300; 2. 中山大学化学与化学工程学院, 广东 广州 510275

Simultaneous determination of ammonium and six alkylamine chromatography

ZHONG Zhixiong1*, LI Gongke2, ZHU Binghui1, LUO Zhibin1, WU Ximei1

 Center for Disease Control and Prevention of Guangdong Province, Guangzhou 510300, China;
School of Sen University, Guangzhou 510275, China

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摘要 建立同时测定化妆品中铵和6种烷基胺的离子色谱(IC)分析方法。优化了色谱条件和样品前处理方法,样品经100 mmol/L乙酸乙腈溶液浸提,固相萃取(SPE)柱去除阴离子、中和氢离子后进样测定。考察了提取溶液的pH、有机溶剂和共存离子对测定结果的影的线性范围为0.3~15 mg/L,检出限为2.1~7.9 mg/kg,定量限为7~26 mg/kg。采用建立的分析方法测定了清洗、柔肤、祛斑、发、染发和育发类化妆品的加标回收率的范围在80.2%~109.2%之间,相对标准偏差(RSD)的范围为0.5%~3.1%。方法选择性灯 抗干扰强,用于实际样品测定结果准确。

关键词: 离子色谱法 铵 烷基胺 化妆品

Abstract: A method for the simultaneous determination of ammonium and six alkylamines in cosmetic product chromatography (IC) was developed. The sample pretreatment process and the separation resolution of chromatography were investigated. The samples were extracted by 100 mmol/L acetic acid-20%(v/v) acetonit solution at room temperature, and then solid phase extraction (SPE) column was used to eliminate the interfee The influences of pH value, organic solvent and coexisted ions were investigated. The separation was carried IonPac CS17 (250 mm×4 mm)analytical column and IonPac CG17 (50 mm×4 mm)guard column using $1.5 \sim 10^{-2}$ methanesulfonic acid and $0.5\% \sim 5\% (v/v)$ acetonitrile gradient elution at a flow rate of 1.0 mL/min at 24%, consuppressed conductivity detector. Under the optimum conditions, a measurement could be completed less that The linearity ranged from 0.3 to 15 mg/L, the detection limits and the quantification limits were in the ranges of mg/kg and $7\sim 26$ mg/kg, respectively. The method was successfully employed for the determination of ammonimethylamine, dimethylamine, trimethylamine, ethylamine, propylamine and butylamine in samples including clebody lotion, skin-bleaching, sun block, marcel, hair dye and pilatory cosmetics with the recoveries of $80.2\% \sim 1$ the relative standard deviations (RSDs) of $0.5\% \sim 3.1\%$. The method offered high selectivity, sensitivity, and gasatisfactory results for real sample analysis.

Keywords: ion chromatography (IC) ammonium alkylamine cosmetic products

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Corresponding Authors: 钟志雄

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