

Cece Furwanti:

We have reached a decision regarding your submission to Media Pharmaceutica Indonesiana (MPI), "Simultaneously HPLC determination of Lidocaine Hydrochloride and Hexachlorophene in Suppositories, and its Validation According USP 41".

Our decision is to: Accept Submission

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Media Pharmaceutica Indonesiana (MPI)

## Simultaneous HPLC determination of Lidocaine Hydrochloride and Hexachlorophene in a Suppository Product

## Abstract

Introduction: Antihemoroid® suppository has been produced commercially by PT Kimia Farma, Indonesia. For QC purposes, a separated densitometric method for analysis of its active ingredients, lidocaine hydrochloride and hexachlorophene, was applied. Objective: The objective of this study was obtaining more efficient analysis method of LH and HC, therefore an HPLC procedure has been developed for the determination of both compounds simultaneously. Methods: AYMC-Triart C18 column was used with a gradient mobile phase consisting of acetonitrile and phosphate buffer 0.05 M (pH 6.0). Quantitative evaluation was performed at 220 nm. Method validation was performed according to the new methods of USP 41. Result: Showed that the HPLC method was simple, accurate, precise, and robust. Conclusion: The HPLC method can be applied in simultaneous determination of LH and HC in suppositories as a QC tool in the pharmaceutical industries.

**Key words:** hexachlorophene, HPLC, lidocaine HCl, method validation, suppository

## 1. Introduction

PT. Kimia Farma (Persero) Tbk., Plant Watudakon, Jombang, Indonesia, produced antihemoroid®suppository which have indication as anti-hemorrhoids;