New spectrophotometric estimation of indomethacin capsules with niacinamide as hydrotropic solubilizing agent

Abstract

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Background: Hydrotropic solubilization process involves cooperative intermolecular interaction with several balancing molecular forces, rather than either a specific complexation event or a process dominated by a medium effect, such as co-solvency or salting-in. **Materials and Methods:** In the present investigation, hydrotropic solution of 2 M niacinamide was employed as the solubilizing agent to solubilize the poorly water-soluble drug, indomethacin, from the capsule dosage form for spectrophotometric determination in ultraviolet region. **Results:** Hydrotropic agent used did not interfere in the spectrophotometric analysis. In preliminary solubility studies, it was found that there was more than fivefold enhancement in the aqueous solubility of indomethacin (poorly water-soluble drug) in 2 M niacinamide solution as compared to its aqueous solubility at $28 \pm 1^{\circ}$ C. **Conclusion:** The proposed method is new, simple, safe, environmentally friendly, economic, accurate and cost-effective and can be successfully employed in routine analysis.

Key words: Environmentally safe, hydrotropic solubilization, indomethacin, niacinamide, spectrophotometric

INTRODUCTION

Hydrotropy is the term originally put forward by Neuberg to describe the increase in the solubility of a solute by the addition of fairly high concentrations of alkali metal salts of various organic acids.^[1] Hydrotropic solubilization process involves cooperative intermolecular interaction with several balancing molecular forces, rather than either a specific complexation event or a process dominated by a medium effect, such as co-solvency or salting-in. Hydrotropic agents have been observed to enhance the aqueous solubility of poorly water-soluble drugs.^[2-15]

It is a phenomenon where addition of large amount of second solute results in increase in aqueous solubility of another solute. Concentrated aqueous hydrotropic solutions of sodium benzoate, sodium salicylate, urea, nicotinamide, sodium citrate and sodium glycinate have been observed to enhance the aqueous solubilities of poorly water-soluble drugs. Hydrotropic solutions can be employed to replace organic solvents employed in analysis of poorly water-soluble drugs.^[15-22]

Mixed hydrotropic solubilization technique is the phenomenon to increase the solubility of poorly water-soluble drugs, using blends of hydrotropic agents.^[19-25] This technique can provide additive or synergistic effect on the solubility of poorly water-soluble drugs. Utilization of this method in the formulation of dosage forms made of water-insoluble drugs can also reduce the concentration of individual hydrotropic agents, in order to minimize the side effects (in place of using a large concentration of one hydrotrope, a blend of several hydrotropes can be employed in much smaller concentrations, reducing their individual toxicities).

The spectrophotometric analytical method available for indomethacin in literature is in United States Pharmacopeia, in which methanol and methylene chloride were used to solubilize indomethacin. In this method, approximately 200 ml of methylene chloride was used, which is a toxic and costlier organic solvent.^[26]

Therefore, the basic objective of the present investigation was to employ the use of hydrotropic solution to extract the drug from the dosage forms, excluding the use of costlier solvents. Costlier organic solvents are more often employed to solubilize the poorly water-soluble drugs for spectrophotometric analysis. Volatility, high cost, toxicity and pollution are the drawbacks of such solvents. In this investigation, a hydrotropic solution has been employed to solubilize the drug for its spectrophotometric analysis precluding the use of organic solvent.

MATERIALS AND METHODS

Materials

Indomethacin Capsules, Indocap[®] manufactured by Jagson Pharmaceuticals, Uttarakhand (B No JR09A009) as formulation I and Donica[®] manufactured by IPCA Pharmaceuticals, Uttarakhand (B No IP18D105) as formulation II were purchased from local market. Gift sample of bulk indomethacin drug was provided by Ranbaxy Laboratories Ltd., Dewas, India. Free gift sample of niacinamide was obtained from Alkem Laboratories Ltd., Mumbai, India.

Calibration curve

A Shimadzu[®] 1700, double-beam UV-visible spectrophotometer, with 10-mm matched silica cells was used for spectrophotometric analysis (software used UV Probe Ver 7.0). 2 M niacinamide was scanned

against water and no interference was found in 300–350 nm range, in which indomethacin is being analyzed [Figure 1]. Twenty milligrams of indomethacin was dissolved in 50 ml of methanol and the volume was made up to 100 ml with methanol. Further dilutions were made with water and analyzed against the corresponding reagent blank [Figure 2]. From Figure 2, the characteristic peak of indomethacin was found at 320 nm, peak 1, which is far out of the range of niacinamide peak. So, we can conclude that no interference was there due to the use of hydrotropic agent.

Accurately weighed 50 mg of indomethacin was transferred to 50 ml volumetric flask and 40 ml of 2 M niacinamide was added, the drug was solubilized by shaking and the volume made up to the mark with distilled water. The standard stock solution was diluted with distilled water to obtain various dilutions. The dilutions of 10, 20, 30, 40 and 50 µg/ml were used to plot the calibration curve by noting the absorbance at λ_{max} 320 nm against the corresponding reagent blank. Beer's law was obeyed in the concentration range of 10–50 µg/ml (R² = 0.999).

Table 1: Analytical data of indomethacin capsules with statistical evaluation $(n = 3)$				
Capsule formulation	Label claim per capsule (mg)	Percent label claim estimated* (Mean ± SD)	Percent coefficient of variation	Standard error
I	25	98.39 ± 2.113	2.147	1.220
II	25	98.04 ± 1.387	1.415	0.801

*Average of three determinations



Figure 1: 2 M niacinamide scanned against water from 200 to 400 nm

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