

**Formulation and Characterization of Film Coated Matrix
Tablets of Anti-Anginal Drug**

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Abstract

The purpose of the present study was to design and evaluate once-daily sustained release film coated matrix tablets of anti-anginal drug Ivabradine hydrochloride. The tablets were prepared by direct compression method using hydroxyl propyl methylcellulose (HPMC) K4M and K30M, xanthan gum, carbopol 934 and lactose as a channeling agent. Prepared matrix tablets were evaluated for various parameters like hardness, thickness, weight variation, friability, drug content, *in vitro* drug release, drug release kinetics and matrix tablets were film coated using HPMCK4M as a polymer. The optimized matrix tablet had a drug to polymer ratio 1:3:3 for xanthan gum and carbopol 934 offered the required *in-vitro* drug release among the all nine batch and that follow non-fickin diffusion. The evaluation parameters hardness, thickness, weight variation, friability and drug content were found within the limits. This optimized formulation was film coat with 5% HPMC K4M and *in vitro* studies showed that tablets released the drug over a period of 24 hr and 5% fim coating did not change the drug release profile. Hence it can be conclude that a combination of release retarding polymer xanthan gum, release modifying agent carbopol

934 and film coating polymer HPMC K4M can effectively control the drug release for freely water soluble drugs over a period of 24 h.

Keywords: Ivabradine hydrochloride, direct compression, film coated matrix tablet, sustained release.