



Formulation and Bioequivalence Evaluation of Extended Release Solid Drug Delivery System for Metronidazole Using Eudragit NM30D and Methocel Premium K4M as Retardant Material

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ABSTRACT

Conventional indefinite quantity style of antiprotozoal drug create a drug blood level time profile which does not maintain inside the therapeutic vary for extended amount of your time. within the gift study, a shot was finished to develop Associate in Nursing extended unharness antiprotozoal drug composition which may be capable of delivering acceptable bioavailability for up to twenty four hours. Complete unharness formulations of antiprotozoal drug were ready employing a type of proportions of Eudragit NM30D and Methocel Premium K4M, on an individual basis through wet granulation. Eudragit NM30D and Methocel Premium K4M were use as agent material. 3 formulations (U-1 to U-3) were developed mistreatment Eudragit NM30D and whereas a brand new 5 (M-1 to M-5) mistreatment Methocel Premium K4M. The grain for tableting were evaluated for wetness content, squeezability index and angle of repose etc. Tablets were subjected to thickness, stiffness, breakableness and in vitro unharness studies. Dissolution study of the developed pill matrices were administered in zero.1 N Hydrochloric acids for twenty-four hours stage. Dissolution profiles were then value with innovator's drug metronidazole ER in name only of distinction issue (f1) and similarity issue (f2). it absolutely was experimental that, formulation U-1, U-2, M-2, M-3, M-4 meet the need of bioequibalance with metronidazole ER. Among them, M-3 showed the easiest similarity and lowest distinction issue. it's evident from the study that, formulation M-3 posses all the desired characteristics to provide Associate in Nursing extended unharness antiprotozoal drug composition which can be stable enough and capable of delivering acceptable bioavailability for up to twenty four hours. Metronidazole (2-methyl-5-nitroimi-dazole-1-ethanol) is Associate in Nursing oral synthetic nitroimidazole antibiotic medication used for the treatment of infections caused by anaerobic bacterium and protozoa. it's an antibiotic, amebicide, and antiprotozoal drug.

In the past many matrix-type and polymeric-coated antiprotozoal drug formulations has been wide investigated as a result of it is a temporary plasma elimination [*fr1] life ranging from 6 to seven hours. a good vary of controlled unharness matrices are developed supported hydrophilic polymers such as hydroxypropylmethyl polysaccharide (HPMC), hydroxypropyl polysaccharide (HPC), sodium alginate, chitosan and xanthan gum. within the current study, a shot was created to develop Associate in Nursing extended unharness Metronidazole composition which may be capable of delivering adequate bioavailability for up to twenty four hours. For this reason Eudragit NM30D and Methocel Premium K4M were used as agent material. mistreatment numerous proportions of Eudragit NM30D and Methocel Premium K4M, complete unharness formulations of antiprotozoal drug were ready on an individual basis through wet granulation. The technology of chemical compound drug delivery has been studied in details over the past thirty years and numerous wonderful reviews area unit obtainable. The 3 key blessings that chemical compound drug delivery product can give area unit localized, extended delivery and stabilization of the drug. many reviews have been given on the mechanisms and therefore the mathematical aspects of unharness of medication from compound matrices. For a given drug, the discharge mechanics from the compound matrix is ruled preponderantly by 3 factors the polymer sort, polymer morphology and therefore the excipients gift in the system. The mechanisms of drug release from numerous chemical compound matrix systems are extensively mentioned. The diffusion of drug molecules to & from the matrix across the hydraulics diffusion layer could also be treated as one-dimensional diffusion to a plane surface. Eudragit NM30D offers time controlled unharness and pH scale free dissolution, makes it appropriate for extended unharness antiprotozoal drug pill. On the opposite

hand, Methocel Premium K4M showed higher unharass agent impact for controlled unharass of pill. The objective of this study was to arrange Associate in Nursing extended unharass antiprotozoal drug pill that will sustain bioavailability of the drug for up to 24 hours.

The present study was designed to develop extended unharass tablets of Metronidazole by mistreatment Eudragit NM30D and Methocel Premium K4M as rate retarding factor on an individual basis by wet granulation methodology. Eudragit NM30D was employed in the planned formulations (U-1 to U-3) and Methocel Premium K4M in (M-1 to M-5) so as to evaluate the quantity of compound needed to produce desired unharass rate for twenty-four hours amount. The granules of planned formulations were evaluated for moisture content, angle of repose; compressibility index and drug content. Moisture content of the granules for formulation U-1 to U-3 ranged from two.32 to 2.87 and for formulation M-1 to M-5 ranged from one.63% to 1.85%. Angles of repose The results of angles of repose of planned formulations ranged from 19.64 ± 0.02 (°) to twenty three. 15 ± 0.03 (°) (Table 4). Compressibility index (%) The results of squeezability index (%) of planned formulations ranged from 14.55 ± 0.02 to 20.34 ± 0.04 (Table 4). Drug content The drug content in a very weighed amount of all formulations ranged from ninety nine.14% to 100.21%. of these results indicate that the granules possess satisfactory flow properties, compressibility and drug content (Table 4). Evaluation of tablet The tablets of the proposed formulations (U-1 to U-3 and M-1 to M-5) were subjected to varied analysis tests such as thickness, hardness, weight variation test and breakableness take a look at and drug content. Keywords: Retardant material, Eudragit NM30D, Methocel Premium K4M, Metronidazole

