

Society for Research Development in Health Sciences (RDHS), Sponsored



2nd International Conference

Organized By

Ambe Durga Education Society's

Dadasaheb Balpande College of Pharmacy

(Degree and Diploma), Near Swami Samarth Dham Mandir, Besa, Nagpur-440037, Maharashtra, India.



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FORMULATION AND EVALUATION VALSARTAN FAST DISINTEGRATING TABLET

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ABSTRACT

To develop orodispersible tablet of Valsartan by using natural and synthetic superdisintegrant. Cordia dichotoma fruits were collected from Maharashtra region (India) in the month of June. Isolation of gum from Cordia dichotoma fruit, The identification characteristics of drug are performed and all the parameters were observed satisfactory. Direct compression technique was used, microcrystalline cellulose PH 102(MCC) was used as a diluent, sodium starch glycolate (SSG), and crospovidone(CP) were used as synthetic superdisintegrants, Cordia dichotoma gum was used as natural superdisintegrant, talc was used as flow promoter magnesium stearate was used as lubricant, aspartate as sweetener and vanilla flavour was used to improve mouth feel. The results obtained by FTIR and DSC studies revealed that there was no chemical interaction between the pure drug and excipients. Direct compression method was employed to formulate the tablets, the pre-compression and post compression parameter study was determined formulations. The final optimized formulations of Valsartan tablets containing 7.5% Cordia dichotoma gum was compared with marked conventional tablets of valsartan (Valtan) and the results revealed that formulated fast dissolving tablets of Valsartan were effective and better to meet with official compliances. The concept of formulating orodispersible tablets containing Valsartan offers a suitable, practical approach to achieve fast release of the drug. Gives fast release and meets its aim and objectives.

Keywords Valsartan, Orodispersible tablets, Chordiadichotoma

FORMULATION AND EVALUATION OF THEOPHYLLINE FLOATING TABLET Bamba Sakshi ¹, Sharma Ravindrakumar ², Kawitkwar P.S. ¹, Thakare Vinod ¹, Patil V. R. ²

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ABSTRACT

To develop gastric floating drug delivery system containing Theophylline Anhydrous and evaluation of the same. The floating behavior, which showed floating lag time 1- 2 min and total floating time more than 12 hrs. In-vitro drug release study was performed in1.2 pH buffer. In formulation FT1-FT4, this contained carbopol, as concentration of polymer (HPMC K15 and HPMC K100) increase simultaneous weight of tablet and hence increases in swelling percentage. In this study the maximum percentage observed in FT3 and formulation FT5-FT8 showed minimum swelling percentage. All the formulations showed increase in swelling indices for 9-12 hrs. Formulation without carbopol showed drug release in 9-11 hrs. Formulation with high polymer content (FT6) extended drug release up to 11 hrs. The above result demonstrate that the theophylline anhydrous reducing dosing frequency and thus increasing patient compliance.

Keywords Theophylline, Floating tablet, Carbopol, HPMC K15, HPMC K100, Drug release