

International Conference and Exhibition on

# Pharmaceutical Development and Technology

April 24-26, 2017 Dubai, UAE

## Formation of solid dispersions famotidine with HPMC E5LV and mannitol with co-grinding technique

Robby Kurniawan, Erizal Zaini, Lili Fitriani and Sherly Ramadhani  
Andalas University, Indonesia

**Background & Aim:** Solid dispersion has attracted considerable interest as an efficient means of improving the solubility and the dissolution rate of poorly water-soluble drug. The aim of this study was to prepare solid dispersions of famotidine with HPMC E5LV and mannitol as carrier to improve its solubility and its dissolution rate.

**Methods:** Co-grinding techniques by using ball milling was used. 18 formulae with 3 different ratios to HPMC and mannitol (1:1, 1:2, 2:1) and 3 different grinding times (30', 60', 90') were prepared. Characterization of solid dispersion was analyzed with scanning electron microscopy analysis (SEM), X-ray diffraction, Fourier transform infrared (FTIR), optilab microscope camera, solubility test and dissolution test. The solid state interaction of co-ground and physical mixture was evaluated by X-ray powder diffraction and SEM. The dissolution studies were conducted in USP type II apparatus.

**Results:** The result of X-ray powder diffraction analysis showed that the co-ground of famotidine with HPMC E5LV and mannitol decreased the drug crystallinity. X-ray powder diffraction showed the transformation of crystalline state of famotidine to amorphous by co-grinding with HPMC E5LV and mannitol.

**Conclusion:** SEM results showed the co-ground mixture with HPMC E5LV had smaller size and co-ground mixture with mannitol showed agglomerate form. The highest in solubility and dissolution rate was observed for famotidine-HPMC and E5LV showed in 1:1 ratio with 90' grinding time and famotidine-mannitol showed in 1:2 ratio with 30' grinding time compared to the intact famotidine and its physical mixture.

[robbyk142@gmail.com](mailto:robbyk142@gmail.com)

## Potential healing effects of *Hibiscus sabdariffa* L. flowers on arthritis

Zena M Fahmi Qaragholi  
University of Baghdad, Iraq

The present study has been focused to assess the anti-inflammatory activity and healing effects of the aqueous extract of *Hibiscus sabdariffa* L. flowers on induced arthritis in mice and compare it with meloxicam (Mobic®), one of the most conventional drugs used to treat arthritis. The water extract of *Hibiscus sabdariffa* was administered orally at a dose of 300 mg/kg, 400 mg/kg and 500 mg/kg body weight for 14 days after induction of arthritis with incomplete Freund's adjuvant. (T<sub>1</sub>, T<sub>2</sub> and T<sub>3</sub>) showed a significant increase in body weight when compared with T<sub>4</sub>, negative and positive control groups. A significant decrease in the levels of RBC and Hb was observed in all groups subjected to arthritic (T<sub>1</sub>, T<sub>2</sub>, T<sub>3</sub>, T<sub>4</sub> and T<sub>5</sub>) when compared to the negative group (T<sub>6</sub>). The administration of the aqueous extract of *Hibiscus sabdariffa* L. flowers to arthritic mice in (T<sub>1</sub>, T<sub>2</sub> and T<sub>3</sub>) improved the levels of Hb and RBC to near normal. A significant reduction (P≤0.01) in spleen weight, WBC, ESR, CPR and serum copper level was found at all treatment groups with the water extract of *Hibiscus sabdariffa* in comparison with the groups treated with meloxicam, the positive and the negative (T<sub>1</sub>, T<sub>2</sub> and T<sub>3</sub>) revealed a significant (P≤0.01) reduction of the inflammation in comparison with the other treatment groups (T<sub>4</sub>, T<sub>5</sub>). A better activity was observed at 500 mg/kg body weight in mice.

[zenaqaragholi@yahoo.com](mailto:zenaqaragholi@yahoo.com)