



MOBITIL JOURNAL

12 Pages



Issue date : Aug 2011

VOL.NO. 1



Medical Union Pharmaceuticals, Abu-Sultan, Ismailia, Egypt.

THE NSAIDs REPORT

More and more patients become unsatisfied with ordinary NSAIDs treatment results

Condition: Rheumatoid Arthritis & Joint pain

Drug: Ibuprofen

Reviewer: Female, 45, on treatment for 3 months

Comment: Never had a blood pressure problem until I began taking this drug on a regular basis to control pain/inflammation. This NSAID is definitely not for me!

Condition: Tennis elbow

Drug: Diclofenac Potassium

Reviewer: Male, 50, on treatment for less than one month

Comment: No relief noticed after 8 days at 3 tablets per day... Very irritating...



Condition: Osteoarthritis

neck and knee

Drug: Piroxicam

Reviewer: Female, 67, on treatment for 10 years

Comment: It worked great as pain killer. But eventually my BP started rising only due to Piroxicam.



Condition: Menstrual cramps

Drug: Ibuprofen

Reviewer: Female, 21, on treatment for 1 day

Comment: Triggered my asthma within a day. Severe shortness of breath and constant stabbing pain when taking a slightly deep breath for 3 days after that. Is good for cramps but not so good for people with asthma as it triggers it.





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One Step above



Tablet formulation containing meloxicam and beta-cyclodextrin: mechanical characterization and bioavailability evaluation.

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Abstract

The purpose of this research was to evaluate beta-cyclodextrin (beta-CD) as a vehicle, either singly or in blends with lactose (spray-dried or monohydrate), for preparing a meloxicam tablet. Aqueous solubility of meloxicam in presence of beta-CD was investigated. The tablets were prepared by direct compression and wet granulation techniques. The powder blends and the granules were evaluated for angle of repose, bulk density, compressibility index, total porosity, and drug content. The tablets were subjected to thickness, diameter, weight variation test, drug content, hardness, friability, disintegration time, and in vitro dissolution studies. The effect of beta-CD on the bioavailability of meloxicam was also investigated in human volunteers using a balanced 2-way crossover study. Phase-solubility studies indicated an A(L)-type diagram with inclusion complex of 1:1 molar ratio. The powder blends and granules of all formulations showed satisfactory flow properties, compressibility, and drug content. All tablet formulations prepared by direct compression or wet granulation showed acceptable mechanical properties. The dissolution rate of meloxicam was significantly enhanced by inclusion of beta-CD in the formulations up to 30%. The mean pharmacokinetic parameters (C(max), K(e), and area under the curve [AUC](0-infinity)) were significantly increased in presence of beta-CD. These results suggest that beta-CD would facilitate the preparation of meloxicam tablets with acceptable mechanical properties using the direct compression technique as there is no important difference between tablets prepared by direct compression and those prepared by wet granulation. Also, beta-CD is particularly useful for improving the oral bioavailability of meloxicam.

MOBITIL improved
Meloxicam and β -Cyclodextrin
oral bioavailability

Tablet Formulation Containing Meloxicam and β -Cyclodextrin: Mechanical Characterization and Bioavailability Evaluation

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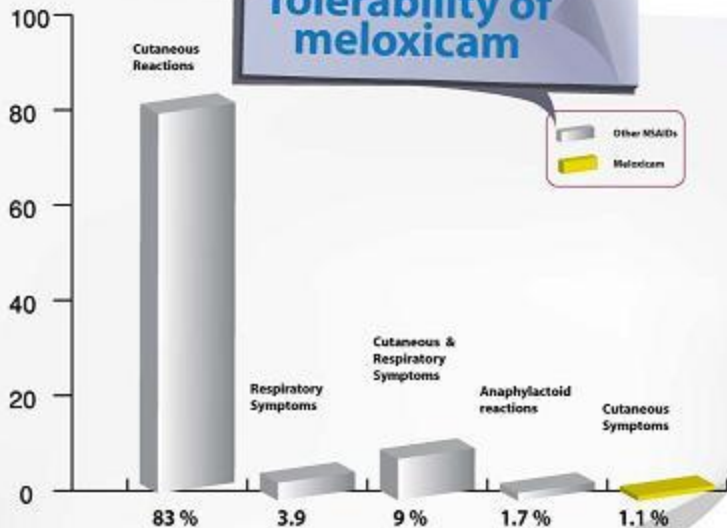
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Submitted: March 22, 2004; Accepted: July 27, 2004.

In conclusion, inclusion complexation of MX in β -CD results not only in an improvement of the bioavailability of the drug but also in the rapid plasma appearance of the drug observed for the complex, which would be highly advantageous for the use of this form in oral MX therapy. Furthermore, the use of β -CD would facilitate the pharmaceutical preparation of the tablets, particularly from the viewpoint of enhancement of the dissolution rate and compression behavior, which give good mechanical properties when used as a direct compression vehicle.

Tolerability of meloxicam

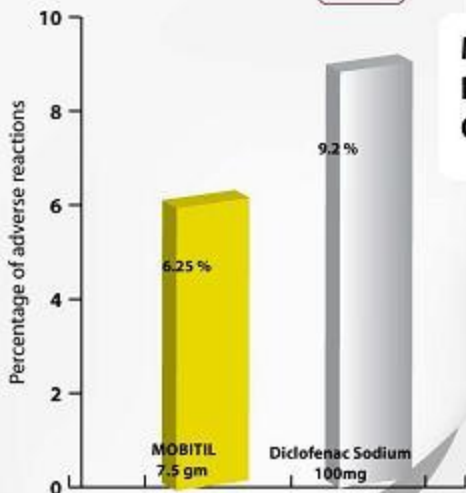


Tolerability of meloxicam in patients with histories of adverse reactions to nonsteroidal anti-inflammatory drugs

Meloxicam seems to be well tolerated by NSAID-sensitive subjects whose reactions are manifested by urticaria/angioedema. Additional study is needed for a more complete assessment of its tolerability in patients with aspirin-induced asthma and other severe manifestations of NSAID sensitivity.

MOBITIL Comparable Efficacy & Safety

n = 336



Meloxicam 7.5mg has the same efficacy of Diclofenac Sodium 100mg in treatment of Osteoarthritis with a better safety profile.

In December 2009, the FDA issued a warning about the **potential for liver damage during treatment with all diclofenac products, including the gel.** The smallest effective amount should be applied for the shortest time possible to minimize the potential for an adverse liver-related event.

Meloxicam in Osteoarthritis: A 6-Month, Double-Blind Comparison with Diclofenac Sodium

Oxford Journals Medicine Rheumatology Volume35, Issue suppl1Pp.39-43