

Review on Formulation and Evaluation of Aceclofenac Tablet

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Abstract: In 1843, the British painter and the inventor was William Brockedon is granted a patent for a machine capable of such "Shaping Pills, Lozenges and Black Lead by the Pressure in that Dies". The device was capable to be compressing powder into the tablet without use of an adhesive Aceclofenac it is an oral non-steroidal anti-inflammatory drug (NSAID) which having anti-inflammatory and analgesic properties. Although there are various differences in the authorized indications between countries, aceclofenac is mainly to recommended for the treatment of inflammatory and painful processes, such as the low back pain (LBP), scapulothoracic per arthrosis, extraarticular rheumatism, odontalgia, and the osteo arthritis (O A), rheumatoid arthritis (RA), and the enclosing spondylitis (AS). The purpose that of study was to develop fast and rapid dissolving tablets of the Aceclofenac using different concentration superdisintegrants. Fast dissolving tablets of the Aceclofenac were prepared by wet granulation technique using the sodium starch glycol ate together with Polyplasdone xl-10 as super disintegrants. The porous granules where that compressed in to tablets. These tablets are were evaluated for drug content, weight variation, friability, hardness and wetting time and Dispersion time. All the formulations are showed at low weight variation with the dispersion time less than 90 seconds and the fast in vitro dissolution. These types of drug content of all the formulations was within the acceptable limits. The optimized formulation is showed good release of profile with maximum drug being released at prolong time intervals. That was concluded that fast dissolving tablets with improved the Aceclofenac dissolution could be prepared by wet granulation of tablet. The dispersion time and the dissolution parameter (t50% and t80%) decreased with increasing the concentration of Polyplasdone xl-10 advertisement.

Keywords: Aceclofenac

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