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<p>(51) International classification :A61K0009000000, A61K0009200000, A61K0047260000, A61K0031138000, A61K0047380000</p> <p>(86) International Application No :NA Filing Date :NA</p> <p>(87) International Publication No : NA</p> <p>(61) Patent of Addition to Application Number :NA Filing Date :NA</p> <p>(62) Divisional to Application Number :NA Filing Date :NA</p>	<p>(71)Name of Applicant : 1)Ms. Pooja Malik Address of Applicant :Assistant Professor, School of Pharmaceutical Sciences, IFTM University, Moradabad, Uttar Pradesh, Pin Code: 244102</p> <p>-----</p> <p>2)Ms. Payal Malik 3)Mr. Amit Kumar 4)Mr. Shivam 5)Ms. Mujeeba Rehman 6)Mr. Navneet Thakkar 7)Mr. Sandeep Singh Name of Applicant : NA Address of Applicant : NA</p> <p>(72)Name of Inventor : 1)Ms. Pooja Malik Address of Applicant :Assistant Professor, School of Pharmaceutical Sciences, IFTM University, Moradabad, Uttar Pradesh, Pin Code: 244102</p> <p>-----</p> <p>2)Ms. Payal Malik Address of Applicant :Student, School of Biotechnology, IFTM University, Moradabad, Uttar Pradesh, Pin Code: 244102 -----</p> <p>----</p> <p>3)Mr. Amit Kumar Address of Applicant :Assistant Professor, School of Pharmaceutical Sciences, IFTM University, Moradabad, Uttar Pradesh, Pin Code: 244102</p> <p>-----</p> <p>4)Mr. Shivam Address of Applicant :Assistant Professor, School of Pharmaceutical Sciences, IFTM University, Moradabad, Uttar Pradesh, Pin Code: 244102</p> <p>-----</p> <p>5)Ms. Mujeeba Rehman Address of Applicant :PhD Scholar, Babasaheb Bhim Rao Ambedkar University, Lucknow, Uttar Pradesh, Pin Code: 226025 -----</p> <p>-</p> <p>6)Mr. Navneet Thakkar Address of Applicant :Research Scholar, School of Pharmaceutical Sciences, IFTM University, Moradabad, Uttar Pradesh, Pin Code: 244102</p> <p>-----</p> <p>7)Mr. Sandeep Singh Address of Applicant :Research Scholar, NKBR College of Pharmacy & Research Centre, Hapur road, Meerut, Uttar Pradesh, Pin Code: 250004 --</p> <p>-----</p>
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(57) Abstract :

The present invention relates to the preparation of oral disintegrating tablets of Metoprolol tartrate, using various pharmaceutical compositions to enhance patient compliance. Metoprolol tartrate was formulated into orally disintegrating using the direct compression method and suitable excipients such as Carbopol 934P, HPMC, carboxy methyl cellulose, Mannitol and ethyl cellulose. The prepared tablets are evaluated for weight variation, thickness, hardness, friability, disintegration time, a simulated wetting test, and in-vitro dissolution. The drug content of all the formulations were in the range of 90.19±0.71%to 99.51±0.07%. Optimized formulation was selected by using the higher mucoadhesive time, mucoadhesive force and higher ex-vivo permeation. Out of all the prepared formulations formulation (F6) was considered as optimize formulations.

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