(12) PATENT APPLICATION PUBLICATION
(19) INDIA
(22) Date of filing of Application :22/12/2021

(21) Application No.202111060045 A
(43) Publication Date : 31/12/2021

(54) Title of the invention : DEVELOPMENT OF SIMVASTATIN NIOSOME TO INCREASE BIOAVAILABILTY

(51) International classification :A61K0031366000, A61K0009000000, C07D0309300000, A61K0009127000, A61K0009200000
(86) International Application No :NA
Filing Date :NA
(87) International Publication No : NA
(61) Patent of Addition to Application Number :NA
Filing Date :NA
(62) Divisional to Application Number :NA
Filing Date :NA

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(57) Abstract :
The present invention relates to preparation of simvastatin niosomes to enhance its solubility and bioavailability. Simvastatin is derived synthetically from fermentation products of Aspergillus terreus. It is used to treat hyperlipidaemia. Simvastatin when hydrolysed produces beta, delta, dihydroxy acid which is similar to HMG – CoA (Hydroxylmethyl glutaryl CoA) in structure. So hydrolysed simvastatin competes with HMG – CoA for HMG – CoA reductase. simvastatin niosomes were prepared by using hand shaking method. From the result of the experiment, it may be concluded that formulation F2 containing 2:1 (Span 60: Cholesterol) was found to be high % of entrapment efficiency and desired sustained release of simvastatin. The in-vivo study value it was found that the bioavailability of simvastatin niosome was greater than the plain simvastatin drug due to the decrease in particle size.

No. of Pages : 19 No. of Claims : 2

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