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Niosomal Carriers Enhance Oral Bioavailability Of carvedilol: effects of bile salt-enriched vesicles and carrier surface charge *Int J Nanomedicine* . 2015 Jul 29;10:4797-813. doi: 10.2147/IJN.S84703.

Niosomal Carriers Enhance Oral Bioavailability Of carvedilol: effects of bile salt-enriched vesicles and carrier surface charge. Abstract: Carvedilol (CRV) is an antihypertensive drug with both alpha and beta receptor blocking activity used to preclude angina and cardiac arrhythmias.

Intestinal and hepatic first-pass metabolism can also restrict oral bioavailability to a significant extent.^{4,5} In the efforts to enhance oral bioavailability, various approaches have been employed, such as solid dispersions,⁶ salt forms,⁷ nanosizing, and micronization.⁸ In the last two decades, there has been increased interest in studying colloidal particulate carriers such as liposomes ...

1/1/2015 · Carvedilol is an antihypertensive drug with both alpha and beta receptor blocking activity used to preclude angina and cardiac arrhythmias. To overcome the low, variable oral bioavailability of

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CRV, niosomal formulations were prepared and characterized: plain niosomes (without bile salts), bile salt-enriched niosomes (bilosomes containing various percentages of sodium cholate or sodium ...

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Niosomal Carriers Enhance Oral Bioavailability Of carvedilol: effects of bile salt-enriched vesicles and carrier surface charge gelareh arzani¹ azadeh haeri¹ Marjan Daeihamed¹ hamid Bakhtiari-Kaboutaraki¹ simin Dadashzadeh^{1,2} ¹Department of Pharmaceutics, Faculty of Pharmacy, ²Pharmaceutical sciences research center, shahid

Carvedilol is an antihypertensive drug with both alpha and beta receptor blocking activity used to preclude angina and cardiac arrhythmias. To overcome the low, variable oral bioavailability of CRV, niosomal formulations were prepared and characterized: plain niosomes (without bile salts), bile salt-enriched niosomes (bilosomes containing various percentages of sodium cholate or sodium ...

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The in vivo study revealed that the niosomal dispersion significantly improved the oral bioavailability of griseofulvin in albino rats after a single oral dose. The maximum concentration (C_{max}) achieved in case of niosomal formulation was approximately double (2.98 µg/ml) as compared to free drug (1.54 µg/ml).

Second and third groups were treated with a single oral dose of 40 mg/kg of AJPER October-

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December 2014, Vol 3, Issue 4 (122-133) Srivastava et al. Enhancement of Oral Bioavailability of Irbesartan by Niosomal Formulation free Irbesartan solution and the freshly prepared unpurified niosomal dispersion containing both the free (89%) and the entrapped drug (11%) by oral route.

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BCS classification. RLX has oral bioavailability of only 2% owing to extensive first pass metabolism. Therefore, it is necessary to increase the solubility and dissolution rate of RLX which lead to improvement in oral bioavailability [3]. Enhancement in oral bioavailability can be achieved by

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reducing the hepatic first pass metabolism.

31/5/2020 · nanomaterials Article Oral Bioavailability Enhancement of Raloxifene with Nanostructured Lipid Carriers Aditya Murthy 1,2, Punna Rao Ravi 2,* , Himanshu Kathuria 2,3 and Shrinivas Malekar 2 1 Differentiated Formulations, Strides Pharma Science Ltd., R & D Centre, J.P. Nagar 2nd Phase, Bangalore 560083, Karnataka, India; adityamurthy1212@gmail.com

Nanostructured lipid (NLCs) carriers as a bioavailability enhancement tool for oral administration Bharti Gaba¹, Mohammad Fazil¹, Asgar Ali¹, Sanjula Baboota¹, Jasjeet K. Sahni², and Javed Ali¹ ¹Department of Pharmaceutics, Faculty of Pharmacy, Jamia Hamdard, Hamdard Nagar, New Delhi, India and ²Khalsa College of Pharmacy, Amritsar, Punjab, India.

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Amisulpride (AMS), a second generation antipsychotic, suffers from low oral bioavailability (48%). This might be due to its pH-dependent solubility or being a substrate of P-glycoprotein efflux pump.

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