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Bile salts are endogenous detergents used extensively in drug delivery as

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permeability enhancers, facilitating drug penetration across biological barriers including skin,²⁰ the intestinal wall,²¹ the blood-brain barrier,²² nasal mucosa,²³ and the cornea.²⁴ Liposomes containing bile salts have been claimed to improve the oral bioavailability of some drugs and macromolecules.^{21,25,26} Although the

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exact mechanisms of this enhanced absorption have not been determined, it has been proposed that ...

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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

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SC or STC) bile salts to the optimal plain
niosomes while keeping the molar ratio of
Span 60 and

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bioavailability of carvedilol ... Niosomal carriers enhance oral bioavailability of carvedilol: eff | IJN. Figure 1 Morphology of plain (F1) (A), bile salt-enriched (F5) (B), cationic (F7) (C), and anionic (F10) (D) carvedilol-loaded niosomes by AFM.

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leads to its oral bioavailability achieved within short span of time, but with short half-life (Doodipala et al., 2011). The antibiotic therapy of Levofloxacin can be markedly enhanced by maintaining the therapeutic level of the drug for extended time in the biological system. An oral

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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

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oral dose. The maximum concentration (C_{max}) achieved in case of niosomal formulation was approximately double ($2.98 \mu\text{g/ml}$) as compared to free drug ($1.54 \mu\text{g/ml}$).

Enhanced Oral Bioavailability of
Griseofulvin via Niosomes ...

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(2016). Sugar-based novel niosomal nanocarrier system for enhanced oral bioavailability of levofloxacin. Drug Delivery: Vol. 23, No. 9, pp. 3653-3664.

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Abstract. Proniosomes (PN) are the dry

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water-soluble carrier systems that may enhance the oral bioavailability, stability, and topical permeability of therapeutic agents. The low solubility and low oral bioavailability due to extensive first pass metabolism make Pentazocine as an ideal candidate for oral and topical sustained release delivery.

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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

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