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## Solubility Enhanced Itraconazole Anti Fungal Gel

**design and optimization of itraconazole tablet employing ...** - solubility and dissolution rate of itraconazole, a highly hydrophobic antifungal agent. materials and methods: the objective was strived through spray drying technique amalgamated with experiment design approach. **solubility enhancement of poorly hydrophilic drugs by ...** - international journal of therapeutic applications, volume 6, 2012, 8 - 13 **8 solubility enhancement of poorly hydrophilic drugs by using different newer techniques: a review research article enhancement of dissolution of poorly ...** - solubility of poorly water soluble drug, by means of solubilising agent. in case of poorly water soluble drug, dissolution may be the in case of poorly water soluble drug, dissolution may be the rate limiting step in the process of absorption. **pegylated nanoarchitecture mediated solubility ...** - delivery of anti-inflammatory agents. polyethylene glycol (peg) conjugation or linking with the dendritic system is called pegylation, which improves water solubility and non-statistical ... **2 produced via quasi nanoemulsion precipitation** - itraconazole is an anti- 37 infective drug that suffers from the above drawbacks, however some attempts involving 38 fabrication in the nanoscale have been found to boost its bioavailability 2, 3 . **a novel formulations for solubility enhancement of ...** - sporanox is a trade name for itraconazole, which has antifungal activity and low solubility. mellaerts et al. (2007) reported that an in vitro study confirmed an enhancement in the release of the **emerging liquisolid compact technology for solubility ...** - emerging liquisolid compact technology for solubility enhancement of bcs class-ii drug p gowree manogar1, bn vedha hari2, d ramya devi2\* 1department of pharmaceutical technology, school of chemical and bio-technology, sastra university, **journal of chemical and pharmaceutical research, 2016, 8(3 ...** - journal of chemical and pharmaceutical research, 2016, 8(3):58-63 ... the drug class known as anti-fungal. it is used for the inhibition of fungal cytochrome p450 enzyme "lanosterol 4 demethylase", used in the conversion of lanosterol to ergosterol, which is a main sterol in fungal cell membrane, thus inhibits replication and promotes cell death in case of the yeast cells transformation ... **research article enhancement of dissolution and anti ...** - meloxicam, a non-steroidal anti-inflammatory drug is widely used in the treatment of rheumatoid arthritis, ankylosing spondylitis and osteoarthritis. one of the major problems with this drug is its low solubility in biological fluids, which results into poor bioavailability after oral ... **company overview - pdfpocket** - enhanced solubility dissolve the hydrophobic drug in water drug (mg/ml) itraconazole paclitaxel water 50 solubility [micelle/water] 2000 times or more 500 times or more plasma level toxic level . 9 abc transporter (e.g. p-glycoprotein) mechanism of overcoming multi-drug resistance micellar nanoparticle . advantages of micellar nanoparticle anti-cancer agents 10 ... **international journal of pharmaceutical research and bio ...** - meloxicam is a nonsteroidal anti-inflammatory drug (nsaid), belongs to bcs ii category (low solubility, high permeability), it's low aqueous dissolution leads to slow onset, thus solubility enhancement could accelerate this process.

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