Actavis

Regulatory Affairs Department
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December 8, 2015

VIA FEDEX OVERNIGHT COURIER

Dockets Management Branch
Food and Drug Administration
Room 1061
5630 Fishers Lane
Rockvlle, MD 20852

CITIZEN PETITION FDA-2006-P-0279

Dear Sir or Madam:

Reference is made to the petition cited above which was originally submitted on October 30, 2006 in accordance with sections 21 CFR 10.20 and 21 CFR 10.30 of the regulations requesting the Commissioner of the Food and Drug Administration to determine that the drug product Oxycodone Hydrochloride Capsules, 5 mg is suitable for submission of an ANDA.

Actavis is not pursuing this product at this time. Therefore, this petition is moot and we request that it be withdrawn.

Respectfully submitted by
Actavis Elizabeth LLC

Madhulika Joshi

for Janak Jadeja, R.Ph
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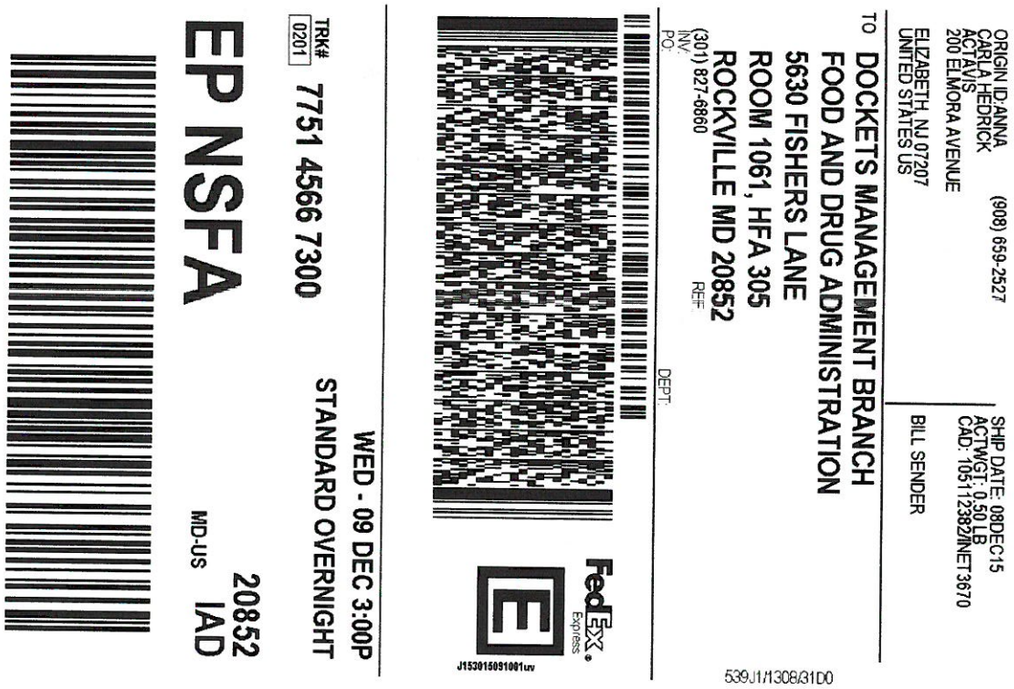

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UNITED STATES US

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