Orange Book: Approved Drug Products with Therapeutic Equivalence Evaluations

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Product Details for NDA 018932

REVIA (NALTREXONE HYDROCHLORIDE)
50MG Marketing Status: Discontinued

Active Ingredient: NALTREXONE HYDROCHLORIDE
Proprietary Name: REVIA
Dosage Form; Route of Administration: TABLET; ORAL
Strength: 50MG
Reference Listed Drug: No
Reference Standard: No
TE Code:
Application Number: N018932
Product Number: 001
Approval Date: Nov 20, 1984
Applicant Holder Full Name: TEVA WOMENS HEALTH INC
Marketing Status: Discontinued
Patent and Exclusivity Information (patent_info.cfm?Product_No=001&Appl_No=018932&Appl_type=N)