

Orange Book: Approved Drug Products with Therapeutic Equivalence Evaluations

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Product Details for NDA 007073

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AZULFIDINE (SULFASALAZINE)
500MG

Marketing Status: Prescription

Active Ingredient: SULFASALAZINE

Proprietary Name: AZULFIDINE

Dosage Form; Route of Administration: TABLET; ORAL

Strength: 500MG

Reference Listed Drug: Yes

Reference Standard: Yes

TE Code: AB

Application Number: N007073

Product Number: 001

Approval Date: Approved Prior to Jan 1, 1982

Applicant Holder Full Name: PHARMACIA AND UPJOHN CO

Marketing Status: Prescription

Patent and Exclusivity Information (patent_info.cfm?

Product No=001&Appl No=007073&Appl type=N)

AZULFIDINE EN-TABS (SULFASALAZINE)
500MG

Marketing Status: Prescription