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Product Details for NDA 020142

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CATAFLAM (DICLOFENAC POTASSIUM)
25MG **Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons**
Marketing Status: Discontinued

Active Ingredient: DICLOFENAC POTASSIUM
Proprietary Name: CATAFLAM
Dosage Form; Route of Administration: TABLET; ORAL
Strength: 25MG **Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons**
Reference Listed Drug: Yes
Reference Standard: No
TE Code:
Application Number: N020142
Product Number: 001
Approval Date: Nov 24, 1993
Applicant Holder Full Name: NOVARTIS PHARMACEUTICALS CORP
Marketing Status: Discontinued
Patent and Exclusivity Information (patent_info.cfm?Product_No=001&Appl_No=020142&Appl_type=N)

CATAFLAM (DICLOFENAC POTASSIUM)
50MG **Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons**
Marketing Status: Discontinued

Active Ingredient: DICLOFENAC POTASSIUM
Proprietary Name: CATAFLAM

Dosage Form; Route of Administration: TABLET; ORAL
Strength: 50MG **Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons**
Reference Listed Drug: Yes
Reference Standard: No
TE Code:
Application Number: N020142
Product Number: 002
Approval Date: Nov 24, 1993
Applicant Holder Full Name: NOVARTIS PHARMACEUTICALS CORP
Marketing Status: Discontinued
Patent and Exclusivity Information (patent_info.cfm?Product_No=002&Appl_No=020142&Appl_type=N)