

EXHIBIT 1012

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Drug Details

Drug Name(s)	LOESTRIN FE 1/20
FDA Application No.	(NDA) 017354
Active Ingredient(s)	ETHINYL ESTRADIOL; NORETHINDRONE ACETATE
Company	WARNER CHILCOTT
Original Approval or Tentative Approval Date	April 30, 1973
Chemical Type	3 New dosage form
Review Classification	P Priority review drug

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Products on Application (NDA) #017354

Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLDTE Code
LOESTRIN FE 1/20	ETHINYL ESTRADIOL; NORETHINDRONE ACETATE	0.02MG; 1MG	TABLET;ORAL-28	Prescription No	AB

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