# EXHIBIT 1012

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## Drug Details

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|---|---|
| **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** |

- **Therapeutic Equivalents**
- **Approval History, Letters, Reviews, and Related Documents**
- **Labels are not available**

**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 | RLD | TE Code |
|---|---|---|---|---|---|---|
| LOESTRIN FE 1/20 | ETHINYL ESTRADIOL; NORETHINDRONE ACETATE | 0.02MG; 1MG | TABLET;ORAL-28 | Prescription | No | AB |

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