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Drug Details

| | |
|---|-------------------------------|
| Drug Name(s) | BROMDAY |
| FDA Application No. | (NDA) 021664 |
| Active Ingredient(s) | BROMFENAC SODIUM |
| Company | BAUSCH AND LOMB INC |
| Original Approval or Tentative Approval Date | March 24, 2005 |
| Chemical Type | 3 New dosage form |
| Review Classification | S Standard review drug |

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Products on Application (NDA) #021664 Click on a column header to re-sort the table:

| Drug Name | Active Ingredients | Strength | Dosage Form/Route | Marketing Status | RLD TE Code |
|---------------------------|------------------------------------|--------------------------|-----------------------------------|----------------------------------|-----------------------------|
| BROMDAY | BROMFENAC SODIUM | EQ 0.09% ACID | SOLUTION/DROPS;OPHTHALMIC | Prescription | Yes AT2 |
| XIBROM | BROMFENAC SODIUM | EQ 0.09% ACID | SOLUTION/DROPS;OPHTHALMIC | Discontinued | No None |

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