# EDA Home<sup>3</sup> Drug Databases<sup>4</sup> Orange Book<sup>5</sup> Products with Therapeutic Equivalence Grange Book: Approved Drug Products with Therapeutic Equivalence Evaluations Search results from the "OB\_Rx" table for query on "021862."

Active Ingredient: NEPAFENAC

Dosage Form; Route: SUSPENSION/DROPS; OPHTHALMIC

Proprietary Name: NEVANAC

Applicant: ALCON PHARMS LTD

Strength: 0.1%
Application Number: N021862
Product Number: 001

Approval Date: Aug 19, 2005

Reference Listed Drug Yes RX/OTC/DISCN: RX

TE Code:

Patent and Exclusivity Info for this product: View

## Return to Electronic Orange Book Home Page<sup>6</sup>

FDA/Center for Drug Evaluation and Research

Office of Generic Drugs

Division of Labeling and Program Support

**Update Frequency:** 

Orange Book Data - Monthly

Generic Drug Product Information & Patent Information - Daily

Orange Book Data Updated Through February 2015

Patent and Generic Drug Product Data Last Updated April 06, 2015

### Links on this page:

- 1. http://www.addthis.com/bookmark.php? u508=true&v=152&username=fdamain
- 2. http://www.addthis.com/bookmark.php
- 3. http://www.fda.gov/default.htm
- 4. http://www.fda.gov/Drugs/InformationOnDrugs/default.htm
- 5. ../default.cfm
- 6. ../default.cfm

Note: If you need help accessing information in different file formats, see Instructions for Downloading Viewers and Players.

Accessibility Contact FDA Careers FDA Basics FOIA No Fear Act Site Map Transparency Website Policies





U.S. Food and Drug Administration 10903 New Hampshire Avenue Silver Spring, MD 20993 Ph. 1-888-INFO-FDA (1-888-463-6332) **Email FDA** 













Combination Products Advisory Committees Science & Research Regulatory Information Safety Emergency Preparedness International Programs News & Events Training and Continuing Education Inspections/Compliance State & Local Officials Consumers Industry Health Professionals FDA Archive



U.S. Department of Health & Human Services

Links on this page:



FDA Home<sup>3</sup> Drug Databases<sup>4</sup> Orange Book<sup>5</sup>

# Orange Book: Approved Drug Products with Therapeutic Equivalence Evaluations Patent and Exclusivity Search Results from query on Appl No 021862 Product 001 in the OB\_RX list.

#### **Patent Data**

| Appl No | Prod No | Patent No | Patent<br>Expiration | Drug Substance<br>Claim | Drug Product<br>Claim | Patent Use<br>Code | Delist<br>Requested |
|---------|---------|-----------|----------------------|-------------------------|-----------------------|--------------------|---------------------|
| N021862 | 001     | 7834059   | Jan 31, 2027         |                         |                       | U - 1095           |                     |
| N021862 | 001     | 8071648   | Dec 2, 2025          |                         | Υ                     |                    |                     |
| N021862 | 001     | 8324281   | Dec 2, 2025          |                         | Υ                     |                    |                     |

#### **Exclusivity Data**

There is no unexpired exclusivity for this product.

#### Additional information:

- 1. Patents are published upon receipt by the Orange Book Staff and may not reflect the official receipt date as described in 21 CFR 314.53(d)(5).
- 2. Patents listed prior to August 18, 2003 are flagged with method of use claims only as applicable and submitted by the sponsor. These patents may not be flagged with respect to other claims which may apply

View a list of all patent use codes View a list of all exclusivity codes

Return to Electronic Orange Book Home Page<sup>6</sup>

FDA/Center for Drug Evaluation and Research Office of Generic Drugs Division of Labeling and Program Support Update Frequency: Orange Book Data - Monthly Generic Drug Product Information & Patent Information - Daily Orange Book Data Updated Through February 2015 Patent and Generic Drug Product Data Last Updated April 06, 2015

#### Links on this page:

- 1. http://www.addthis.com/bookmark.php?u508=true&v=152&username=fdamain
- 2. http://www.addthis.com/bookmark.php
- 3. http://www.fda.gov/default.htm
- 4. http://www.fda.gov/Drugs/InformationOnDrugs/default.htm
- 5. ../default.cfm
- 6. ../default.cfm

Note: If you need help accessing information in different file formats, see Instructions for Downloading Viewers and Players.

Accessibility Contact FDA Careers FDA Basics FOIA No Fear Act Site Map Transparency Website Policies



U.S. Food and Drug Administration 10903 New Hampshire Avenue Silver Spring, MD 20993 Ph. 1-888-INFO-FDA (1-888-463-6332)





For Government For Press

Combination Products Advisory Committees Science & Research Regulatory Information Safety Emergency Preparedness International Programs News & Events Training and Continuing Education Inspections/Compliance State & Local Officials Consumers Industry Health Professionals FDA Archive



U.S. Department of Health & Human Services

Links on this page:

