Drugs@FDA
FDA Approved Drug Products

Label and Approval History

Drug Name(s)	**LUNELLE**
FDA Application No.	**(NDA) 020874**
Active Ingredient(s)	**ESTRADIOL CYPIONATE; MEDROXYPROGESTERONE ACETATE**
Company	**PHARMACIA AND UPJOHN**

Go to Approval History

Label Information

What information does a label include?
Note: Not all labels are available in electronic format from FDA.

The supplement type of the 04/26/2002 approval does not usually require new labeling.

View the label approved on 10/05/2000 (PDF) for **NDA no. 020874**

- To see if other previously-approved labels are available on this site, go to the "Approval History" section of this page. **Older labels are for historical information only and should not be used for clinical purposes.**

Approval History
NDA 020874

Note: Not all reviews are available in electronic format from FDA.
Older labels are for historical information only, and should not be used for clinical purposes.
Approval dates can only be verified from 1984 to the present.

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

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Action Date	Supplement Number	Approval Type	Letters, Reviews, Labels, Patient Package Insert	Note
04/26/2002	002	Control Supplement		This supplement type does not usually require new labeling.
06/07/2001	001	Control Supplement		This supplement type does not usually require new labeling.
10/05/2000	000	Approval	Label (PDF) Letter (PDF) Review	

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

- **There are no Therapeutic Equivalents**