

This label may not be the latest approved by FDA.

For current labeling information, please visit <https://www.fda.gov/drugsatfda>

PEDIATRIC USE ONLY

45 mg for 6-month administration

↑
PULL TAB HERE
↓

Lupron Depot-PED® NDC 0074-3575-01
45 mg for 6-month administration

LOT

LOT

EXP.

EXP.



2 007 8807

20078807

SN

Includes:

- One prefilled dual-chamber syringe containing 23 gauge 1 1/2 inch needle with LuproLoc® safety device
- One plunger
- Two alcohol swabs

Not made with natural rubber latex.

Do not remove from clamshell until ready to use.

Recommended dosage:

After mixing, a **healthcare provider** should immediately administer entire contents of syringe by intramuscular injection every 6 months. See prescribing information.

Only Activate Safety Device Post-Injection.

Store at 20°C to 25°C (68°F to 77°F); excursions 15-30°C (59-86°F)

Manufactured for: AbbVie Inc.
North Chicago, IL 60064
by: Takeda Pharmaceutical
Company Limited
Osaka, Japan 540-8645
Product of Japan.

Rx only



(01) 00300743575019



NDC 0074-3575-01
Single Dose Administration Kit
with prefilled dual-chamber syringe.

LUPRON DEPOT-PED®
(Leuprolide Acetate for
Depot Suspension)

Dispense the accompanying Medication Guide to each patient.

**45 mg
for 6-month administration
FOR INTRAMUSCULAR INJECTION**

The front chamber contains: leuprolide acetate 45 mg • polylactic acid 169.9 mg • D-mannitol 39.7 mg • stearic acid 10.1 mg

The second chamber contains: carboxymethylcellulose sodium 7.5 mg • D-mannitol 75.0 mg • polysorbate 80 1.5 mg • water for injection, USP, and glacial acetic acid, USP to control pH

to be the latest a
n, please visit h

NDC 0074-3575-01

SINGLE DOSE

(Prefilled Dual-
Chamber Syringe)

LUPRON

DEPOT-PED[®]

(Leuprolide Acetate
for Depot Suspension)

45 mg

for 6-month

administration

**FOR INTRAMUSCULAR
INJECTION**

Each 1.5 mL (when mixed)
contains: leuprolide
acetate 45 mg

20078808



EXP.

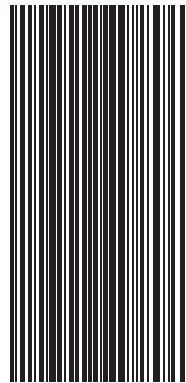
LOT

This label may not be the latest approved by FDA.
For current labeling information, please visit <https://www.fda.gov/drugsatfda>

PEDIATRIC USE ONLY

45 mg for **6-month** administration

20078806



20078806

This label may not be the latest approved by FDA.
For current labeling information, please visit <https://www.fda.gov/drugsatfda>



Hasmukh
Patel

Digitally signed by Hasmukh Patel
Date: 12/04/2023 11:13:15AM
GUID: 508da71e00029e3864a0ad90c32d1866
Comments: Signed for Ramesh Raghavachari, Ph.D.