SLEEP-AID- diphenhydramine hcl tablet ARMY AND AIR FORCE EXCHANGE SERVICE

Exchange Select 44-189

Active ingredient (in each tablet)

Diphenhydramine HCl 25 mg

Purpose

Nighttime sleep-aid

Uses

- for relief of occasional sleeplessness
- reduces time to fall asleep if you have difficulty falling asleep

Warnings

Do not use

- for children under 12 years of age
- with any other product containing diphenhydramine, even one used on skin

Ask a doctor before use if you have

- glaucoma
- a breathing problem such as emphysema or chronic bronchitis
- difficulty in urination due to enlargement of the prostate gland

Ask a doctor or pharmacist before use if you are

taking sedatives or tranquilizers.

When using this product

avoid alcoholic beverages.

Stop use and ask a doctor if

sleeplessness persists continuously for more than 2 weeks. Insomnia may be a symptom of a serious underlying medical illness.

If pregnant or breast-feeding,

ask a health professional before use.

Keep out of reach of children.

In case of overdose, get medical help or contact a Poison Control Center right away.

Directions

- do not take more than directed
- adults and children 12 years and over: take 2 tablets at bedtime if needed or as directed by a doctor
- children under 12 years: do not use

Other information

- each tablet contains: calcium 60 mg
- TAMPER EVIDENT: DO NOT USE IF OUTER PACKAGE IS OPENED OR BLISTER IS TORN OR BROKEN
- protect from moisture
- store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)
- see end flap for expiration date and lot number

Inactive ingredients

croscarmellose sodium, dibasic calcium phosphate dihydrate, FD&C blue #1 aluminum lake, magnesium stearate, microcrystalline cellulose, silicon dioxide, stearic acid

Questions or comments?

1-800-426-9391

Principal Display Panel

exchange**√select**™

Compare to The Active Ingredient of Original Formula Sominex®*

Sleep Aid

Diphenhydramine HCl 25 mg Nighttime Sleep-Aid

Actual Size

72 TABLETS

√guality value

*This product is not manufactured or distributed by Medtech Products Inc., owner of the registered trademark Original Formula Sominex[®].

50844 REV1019K18923

JOUTH NEVIOLDRIOSES

"SATISFACTION GUARANTEED OR YOUR MONEY BACK"

Manufactured For Your Military Exchanges Distributed by: LNK International, Inc.

Hauppauge, NY 11788 1-800-426-9391

TAMPER EVIDENT: DO NOT USE IF PACKAGE IS OPENED OR IF BLISTER UNIT IS TORN, BROKEN OR SHOWS ANY SIGNS OF TAMPERING



diphenhydramine hcl tablet

| Product Information | | | |
|----------------------------|----------------|--------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:55301-189 |
| Route of Administration | ORAL | | |

| Active Ingredient/Active Moiety | | | | |
|---|----------------------------------|----------|--|--|
| Ingredient Name | Basis of Strength | Strength | | |
| DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M) | DIPHENHYDRAMINE HYDROCHLORIDE | 25 mg | | |

| Inactive Ingredients | |
|--|----------|
| Ingredient Name | Strength |
| CROSCARMELLOSE SODIUM (UNII: M28OL1HH48) | |
| DIBASIC CALCIUM PHOSPHATE DIHYDRATE (UNII: O7TSZ97GEP) | |
| FD&C BLUE NO. 1 ALUMINUM LAKE (UNII: J9EQA3S2JM) | |
| MAGNESIUM STEARATE (UNII: 70097M6I30) | |
| MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U) | |
| SILICON DIOXIDE (UNII: ETJ7Z6XBU4) | |
| STEARIC ACID (UNII: 4ELV7Z65AP) | |

| Product Characteristics | | | |
|-------------------------|-------|--------------|----------|
| Color | blue | Score | no score |
| Shape | ROUND | Size | 10mm |
| Flavor | | Imprint Code | 44;189 |
| Contains | | | |

| Packaging | | | | | |
|------------------------|--|-------------------------|-----------------------|--|--|
| # Item Code | Package Description | Marketing Start Date | Marketing End Date | | |
| 1 NDC:55301-189- 23 | 9 in 1 CARTON | 04/10/1990 | | | |
| 1 | 8 in 1 BLISTER PACK; Type 0: Not a Combination Product | | | | |

| Marketing Information | | | | |
|-----------------------|---|-------------------------|-----------------------|--|
| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date | |
| OTC Monograph Drug | M010 | 04/10/1990 | | |
| | | | | |

| Establishment | | | |
|-------------------------|---------|-----------|---------------------|
| Name | Address | ID/FEI | Business Operations |
| LNK International, Inc. | | 038154464 | pack(55301-189) |

| Establishment | | | |
|-------------------------|---------|-----------|--|
| Name | Address | ID/FEI | Business Operations |
| LNK International, Inc. | | 832867837 | manufacture(55301-189) , pack(55301-189) |

| Establishment | | | |
|-------------------------|---------|-----------|----------------------------|
| Name | Address | ID/FEI | Business Operations |
| LNK International, Inc. | | 967626305 | pack(55301-189) |

| Establishment | | | |
|-------------------------|---------|-----------|----------------------------|
| Name | Address | ID/FEI | Business Operations |
| LNK International, Inc. | | 117025878 | manufacture(55301-189) |

Revised: 4/2024 ARMY AND AIR FORCE EXCHANGE SERVICE