# STAY AWAKE- caffeine tablet CHAIN DRUG MARKETING ASSOCIATION INC

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**Quality Choice 44-226** 

## Active ingredient (in each tablet)

Caffeine 200 mg

## Purpose

Alertness aid

#### Use

helps restore mental alertness or wakefulness when experiencing fatigue or drowsiness

# Warnings

## For occasional use only

**Caffeine warning:** The recommended dose of this product contains about as much caffeine as a cup of coffee. Limit the use of caffeine-containing medications, foods, or beverages while taking this product because too much caffeine may cause nervousness, irritability, sleeplessness, and, occasionally, rapid heart beat.

#### Do not use

- for children under 12 years of age
- as a substitute for sleep

# Stop use and ask a doctor if

fatigue or drowsiness persists or continues to recur.

# 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**

- adults and children 12 years and over: take 1 tablet not more often than every 3 to 4 hours
- children under 12 years: do not use

#### Other information

- each tablet contains: calcium 35 mg
- TAMPER EVIDENT: DO NOT USE IF OUTER PACKAGE IS OPENED OR BLISTER IS TORN OR BROKEN
- 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

corn starch, D&C yellow #10 aluminum lake, dextrates hydrated, dibasic calcium phosphate dihydrate, FD&C yellow #6 aluminum lake, magnesium stearate, microcrystalline cellulose, silicon dioxide

Questions or comments?

1-800-426-9391

Principal Display Panel

NDC 63868-114-16

**QC**® QUALITY CHOICE

\*Compare to the Active ingredient in VIVARIN®

Stay Awake

Caffeine 200 mg Alertness Aid Equal to About a Cup of Coffee

actual size

**16** Tablets

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

\*This product is not manufactured or distributed by Meda AB, owner of the registered trademark Vivarin®. 50844 REV1219B22621

SATISFACTION GUARANTEED 100%

QC

Distributed by C.D.M.A., Inc© 43157 W 9 Mile Rd Novi, MI 48375 www.qualitychoice.com Questions:800-935-2362





# **Stay Awake**

\*Compare to the Active Ingredient in VIVARIN®

Caffeine 200 mg



NDC 63868-114-16

\*Compare to the Active Ingredient in **VIVARIN®** 

# Stay Awake

# Caffeine 200 mg

**Alertness Aid** 

actual

Equal to About a Cup of Coffee

**16** Tablets



magnesium stearate, microcrystalline cellulose, silicon phosphate dihydrate, PD &C yellow #6 aluminum lake, #10 aluminum lake, dextrates hydrated, dibasic calcium Inactive ingredients corn starch, D&C yellow

**Drug Facts** (continued)

B-0220-226-21 REV1219B22621

Unestions of comments? 1-800-426-9391

**Drug Facts** (continued)

■ see end flap for expiration date and lot number 12,-30,C (26,-86,E) ■ store at 25°C (77°F); excursions permitted between

**BBOKEN** PACKAGE IS OPENED OR BLISTER IS TORN OR ■ TAMPER EVIDENT: DO NOT USE IF OUTER ■ each tablet contains: calcium 35 mg

Uther intormation

■ children under 12 years: do not use not more often than every 3 to 4 hours ■ adults and children 12 years and over: take 1 tablet

Directions

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when experiencing tatigue or drowsiness **N26** helps restore mental alertness or wakefulness

Caffeine 200 mg. . Aletinessaid (in each tablet)

Drug Facts

\*This product is not manufactured or distributed by Meda AB, owner of the registered trademark Vivarin®. 50844 REV1219B22621 Marnings Active ingredient

No Print / No Varnish

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PRODUCT INFORMATION ANY SIGNS OF TAMPERING KEEP OUTER PACKAGE FOR COMPLETE

Purpose



ONIT IS TORN, BROKEN OR SHOWS PACKAGE IS OPENED OR IF BLISTER TAMPER EVIDENT: DO NOT USE IF

# **STAY AWAKE**

caffeine tablet

| Product  | Intorm | ation |
|----------|--------|-------|
| I I OGGC |        | чи    |

| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:63868-114 |
|--------------|----------------|--------------------|---------------|
|--------------|----------------|--------------------|---------------|

Route of Administration ORAL

# **Active Ingredient/Active Moiety**

| Ingredient Name  | Basis of Strength | Strength |
|--|-------------------|----------|
| CATTERNS (UNIV. 20045) (CATTERNS - UNIV. 20045) (CATTERNS - UNIV. 20045) | CAFFEINE          | 200      |

CAFFEINE (UNII: 3G6A5W338E) (CAFFEINE - UNII:3G6A5W338E) CAFFEINE 200 mg

# **Inactive Ingredients**

| Ingredient Name                                    | Strength |
|--|----------|
| STARCH, CORN (UNII: O8232NY3SJ)                    |          |
| D&C YELLOW NO. 10 ALUMINUM LAKE (UNII: CQ3XH3DET6) |          |

DEXTROSE MONOHYDRATE (UNII: LX22YL083G)

DIBASIC CALCIUM PHOSPHATE DIHYDRATE (UNII: O7TSZ97GEP)

FD&C YELLOW NO. 6 (UNII: H77VEI93A8)
MAGNESIUM STEARATE (UNII: 70097M6I30)

MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)

SILICON DIOXIDE (UNII: ETJ7Z6XBU4)

# **Product Characteristics**

| Color    | yellow | Score        | no score |
|----------|--------|--------------|----------|
| Shape    | ROUND  | Size         | 11mm     |
| Flavor   |        | Imprint Code | 44;226   |
| Contains |        |              |          |

Packaging

|   | ackaging             |  |                         |                       |
|---|----------------------|--|-------------------------|-----------------------|
| # | Item Code            | Package Description                                    | Marketing Start<br>Date | Marketing End<br>Date |
| 1 | NDC:63868-114-<br>16 | 2 in 1 CARTON  | 11/21/1996              |                       |
| 1 |                      | 8 in 1 BLISTER PACK; Type 0: Not a Combination Product |                         |                       |
| 2 | NDC:63868-114-<br>40 | 5 in 1 CARTON  | 11/21/1996              | 03/25/2023            |
| 2 |                      | 8 in 1 BLISTER PACK; Type 0: Not a Combination Product |                         |                       |
|   |                      |  |                         |                       |

| Marketing Information |   |                         |                       |  |
|-----------------------|---|-------------------------|-----------------------|--|
| Marketing<br>Category | Application Number or Monograph<br>Citation | Marketing Start<br>Date | Marketing End<br>Date |  |
| OTC Monograph Drug    | M011  | 11/21/1996              |                       |  |
|                       |   |                         |                       |  |

# Labeler - CHAIN DRUG MARKETING ASSOCIATION INC (011920774)

| Establishment           |         |           |                            |
|-------------------------|---------|-----------|----------------------------|
| Name                    | Address | ID/FEI    | <b>Business Operations</b> |
| LNK International, Inc. |         | 038154464 | pack(63868-114)            |

| <b>Establishment</b>    |         |           |  |
|-------------------------|---------|-----------|--|
| Name                    | Address | ID/FEI    | Business Operations                      |
| LNK International, Inc. |         | 832867837 | manufacture(63868-114) , pack(63868-114) |

| Establishment           |         |           |                            |
|-------------------------|---------|-----------|----------------------------|
| Name                    | Address | ID/FEI    | <b>Business Operations</b> |
| LNK International, Inc. |         | 967626305 | pack(63868-114)            |

| Establishment           |         |           |                        |
|-------------------------|---------|-----------|------------------------|
| Name                    | Address | ID/FEI    | Business Operations    |
| LNK International, Inc. |         | 117025878 | manufacture(63868-114) |

Revised: 6/2023 CHAIN DRUG MARKETING ASSOCIATION INC