

STAY AWAKE- caffeine tablet
Geiss, Destin & Dunn Inc.

GoodSense 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°C-30°C (59°F-86°F)
- see end flap for expiration date and lot number

Inactive ingredients

corn starch, D&C yellow #10 aluminum lake, dextrans 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

GOODSENSE®

Caffeine for Mental Alertness

NDC 50804-262-21

Stay Awake

**Caffeine
Alertness Aid**

16 Tablets - 200 mg Each

actual size

***Compare to the** *active ingredient of*
Vivarin® 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® Tablets.

50844 REV1219B22621

Distributed by:
Perrigo Direct, Inc., Peachtree City, GA 30269
www.PerrigoDirect.com (1-800-426-9391)
GoodSense® is a registered trademark of
L. Perrigo Company.

GOODSENSE®

NDC 50804-262-21

Caffeine for Mental Alertness

Stay Awake

GOODSENSE®

NDC 50804-262-21

Stay Awake



actual size

Caffeine
Alertness Aid

**Compare to the active ingredient of Vivarin® Tablets*

16 Tablets - 200 mg Each

Caffeine for Mental Alertness

E11025

| | |
|--|---|
| Drug Facts (continued) calcium phosphate dihydrate, FD&C yellow #6 aluminum lake, magnesium stearate, microcrystalline cellulose, silicon dioxide | Drug Facts (continued) #10 aluminum lake, dextrans hydrated, dibasic com starch, D&C yellow |
| Questions or comments? 1-800-426-9391 | ■ see end flap for expiration date and lot number |

REV 12/14/11
262262-0565-B

| | |
|---|---|
| Drug Facts (continued) 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. | Drug Facts KEEP OUTER PACKAGE FOR COMPLETE PRODUCT INFORMATION Active ingredient (in each tablet) Caffeine 200 mg.....Alertness aid Purpose Alertness aid |
| 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 | 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 ■ as a substitute for sleep ■ for children under 12 years of age Stop use and ask a doctor if fatigue or drowsiness |
| Other information ■ each tablet contains: calcium 35 mg ■ TAMPER EVIDENT: DO NOT USE IF OUTER OR BROKEN PACKAGE IS OPENED OR BLISTER IS TORN OR | Use helps restore mental alertness or wakefulness when experiencing fatigue or drowsiness |
| ■ store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F) | |

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

no print/no varnish area
lot no. & exp date

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STAY AWAKE

caffeine tablet

Product Information

| | | | |
|--------------------------------|----------------|---------------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:50804-262 |
| Route of Administration | ORAL | | |

Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|--|-------------------|----------|
| 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

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|------------------|--|----------------------|--------------------|
| 1 | NDC:50804-262-10 | 5 in 1 CARTON | 07/01/2020 | |
| 1 | | 8 in 1 BLISTER PACK; Type 0: Not a Combination Product | | |
| 2 | NDC:50804-262-21 | 2 in 1 CARTON | 07/01/2020 | |
| 2 | | 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 | M011 | 07/01/2020 | |

Labeler - Geiss, Destin & Dunn Inc. (076059836)

Establishment

| Name | Address | ID/FEI | Business Operations |
|-------------------------|---------|-----------|---------------------|
| LNK International, Inc. | | 038154464 | pack(50804-262) |

Establishment

| Name | Address | ID/FEI | Business Operations |
|-------------------------|---------|-----------|--|
| LNK International, Inc. | | 832867837 | manufacture(50804-262) , pack(50804-262) |

Establishment

| Name | Address | ID/FEI | Business Operations |
|-------------------------|---------|-----------|---------------------|
| LNK International, Inc. | | 967626305 | pack(50804-262) |

Establishment

| Name | Address | ID/FEI | Business Operations |
|-------------------------|---------|-----------|------------------------|
| LNK International, Inc. | | 117025878 | manufacture(50804-262) |

Revised: 12/2023

Geiss, Destin & Dunn Inc.