

**STAY AWAKE- caffeine tablet**  
**CHAIN DRUG MARKETING ASSOCIATION INC**

*Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.*

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**Quality Choice 44-226**

***Active ingredient (in each tablet)***

Caffeine 200 mg

***Purpose***

Alertness aid

***Uses***

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, 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**

**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](http://www.qualitychoice.com)

Questions:800-935-2362

Equal to About a Cup of Coffee



# Stay Awake

Caffeine 200 mg

\*Compare to the Active Ingredient in VIVARIN®

E1025



NDC 63868-114-16

\*Compare to the Active Ingredient in VIVARIN®

# Stay Awake

Caffeine 200 mg

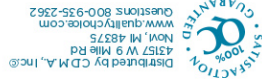
Alertness Aid

Equal to About a Cup of Coffee

actual size



16 Tablets



**Drug Facts (continued)**  
**Questions or comments?** 1-800-426-9391

**Drug Facts (continued)**  
**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

**Drug Facts (continued)**  
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■ each tablet contains: calcium 35 mg  
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■ store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)  
■ see end flap for expiration date and lot number

**Drug Facts**  
**Active ingredient** (in each tablet) Caffeine 200 mg. Alertness aid  
**Purpose**  
Use helps restore mental alertness or wakefulness when experiencing fatigue or drowsiness  
**Warnings**  
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.

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No Print / No Varnish Lot no. & Exp. date

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

KEEP OUTER PACKAGE FOR COMPLETE PRODUCT INFORMATION

B-0220-226-21  
REV1219B22621



Quality Choice 44-226

## STAY AWAKE

caffeine tablet

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:63868-114
<b>Route of Administration</b>	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 (UNII: 35SW5USQ3G)	
DEXTROSE MONOHYDRATE (UNII: LX22YL083G)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
ANHYDROUS DIBASIC CALCIUM PHOSPHATE (UNII: L11K75P92J)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	

### Product Characteristics

<b>Color</b>	YELLOW	<b>Score</b>	no score
<b>Shape</b>	ROUND	<b>Size</b>	11mm
<b>Flavor</b>		<b>Imprint Code</b>	44;226
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63868-114-40	5 in 1 CARTON	11/21/1996	
1		8 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:63868-114-16	2 in 1 CARTON	11/21/1996	
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 FINAL	part340	11/21/1996	

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**Labeler - CHAIN DRUG MARKET ING ASSOCIATION INC (011920774)****Establishment**

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

**Establishment**

<b>Name</b>	<b>Address</b>	<b>ID/FEI</b>	<b>Business Operations</b>
LNK International, Inc.		832867894	MANUFACTURE(63868-114)

**Establishment**

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

**Establishment**

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

**Establishment**

<b>Name</b>	<b>Address</b>	<b>ID/FEI</b>	<b>Business Operations</b>
LNK International, Inc.		868734088	PACK(63868-114)

Revised: 4/2020

CHAIN DRUG MARKETING ASSOCIATION INC