

ALERTNESS AID- caffeine tablet
Chain Drug Consortium

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.

Premier Value 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

***Premier
Value®***

***COMPARE TO THE ACTIVE INGREDIENT
IN VIVARIN®**

Alertness Aid

Caffeine 200 mg
ALERTNESS AID

Equal to about a cup of coffee

40 Tablets

actual
size

**INDEPENDENTLY TESTED
PV
SATISFACTION GUARANTEED**

**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 REV1219A22610

Distributed By:

Pharmacy Value Alliance, LLC
407 East Lancaster Avenue,
Wayne, PA 19087

**If for any reason you are not satisfied with
this product, please return it to the store
where purchased for a full refund.**



Alertness Aid

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ALERTNESS AID

40 Tablets



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8 409886 03210 3

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ALERTNESS AID
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Drug Facts (continued)
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Drug Facts (continued)
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B-1590-226-10
REV1219A22610



44-226

ALERTNESS AID

caffeine tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68016-680
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:68016-680-40	5 in 1 CARTON	11/21/1996	
1		8 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:68016-680-16	2 in 1 CARTON	11/21/1996	05/17/2023
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	

Labeler - Chain Drug Consortium (101668460)**Establishment**

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		038154464	pack(68016-680)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	manufacture(68016-680) , pack(68016-680)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		967626305	pack(68016-680)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		117025878	manufacture(68016-680)

Revised: 3/2023

Chain Drug Consortium