VIVARIN- caffeine tablet
Navajo Manufacturing Company Inc.

Vivarin

Drug Facts

Active ingredient (in each tablet)

Caffeine 200mg

Purpose

Alertness aid

Use

Helps restore mental alertness or wakefulness when experiencing fatigue or drowsiness

Warnings

Do not use

- as a substitute for sleep. This product is intended for occasional use only.
- in children under 12 years of age

When using this product

limit the use of caffeine-containing medications, foods, or beverages because too much caffeine may cause nervousness, irritability, sleeplessness, and occasionally, rapid heartbeat. The recommended dose of this product contains about as much caffeine as a cup of coffee.

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 of age and over: take 1 tablet not more often than every 3 to 4 hours.

Other information

- **each tablet contains calcium 60 mg**
- store in a dry place at 15° - 30°C (59° - 86°F)
- avoid excessive heat (greater than 100°F) or humidity

Inactive ingredients

carnauba wax, colloidal silicon dioxide, croscarmellose sodium, D&C yellow #10 aluminum lake, dicalcium phosphate dihydrate, FD&C yellow #6 aluminum lake, hypromellose, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polysorbate 80, starch, titanium dioxide

Questions or comments?

Call toll-free **1-855-874-0970** weekdays

PACKAGE LABEL-PRINCIPAL DISPLAY PANEL

Made in India

VIVARIN

caffeine tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:67751-226(NDC:71179-018)
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
CARNAUBA WAX (UNII: R12CBM0EIZ)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
D&C YELLOW NO. 10 ALUMINUM LAKE (UNII: CQ3XH3DET6)	
DIBASIC CALCIUM PHOSPHATE DIHYDRATE (UNII: O7TSZ97GEP)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
STARCH, CORN (UNII: O8232NY3SJ)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	yellow	**Score**	no score
Shape	ROUND	**Size**	11mm
Flavor		**Imprint Code**	V;V
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67751-226-04	1 in 1 CARTON	06/19/2023	
1		4 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	06/19/2023	

Labeler - Navajo Manufacturing Company Inc. (091917799)

Registrant - Navajo Manufacturing Company Inc. (136941411)

Establishment

Name	Address	ID/FEI	Business Operations
Navajo Manufacturing Company Inc.		136941411	relabel(67751-226) , repack(67751-226)

Revised: 4/2024

Navajo Manufacturing Company Inc.