NODOZ ALERTNESS AID MAXIMUM STRENGTH- caffeine tablet, film coated Lil' Drug Store Products, 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.

NoDoz® Alertness Aid

Maximum Strength

Drug Facts

Active ingredient (in each caplet)

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 heartbeat.

Do not use

- in children under 12 years of age
- as 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 of age and over: take 1/2 to 1 caplet not more often than every 3 to 4 hours

Other information

• store at controlled room temperature of 20-25°C (68-77°F)

Inactive ingredients

carnauba wax, corn starch, FD&C blue #1, hypromellose, microcrystalline cellulose, povidone K30, propylene glycol, stearic acid, sucrose, titanium dioxide

Questions or comments?

call toll-free 1-877-507-6516 (M-F 8AM-4:30PM CST) or www.nodoz.com

PRINCIPAL DISPLAY PANEL - 200 mg Caplet Pouch Carton

Max Strength • Fast Acting NoDoz® Alertness Aid 200 mg CAFFEINE EACH 4 CAPLETS 2 POUCHES, 2 CAPLETS PER POUCH Lil' Drug Store ®



NODOZ ALERTNESS AID MAXIMUM STRENGTH

caffeine tablet, f	film coated						
Duraharitat							
Product Info	rmation						
Product Type		HUMAN OTC DRUG	HUMAN OTC DRUG Item Code		NDC:66	:66715-9799	
Route of Admin	nistration	ORAL					
Active Ingred	lient/Active	Moietv					
.		ient Name		Basis of Stre	nath	Strength	
CAFFEINE (UNII: 3	-	FFEINE - UNII:3G6A5W338E)		CAFFEINE		200 mg	
Inactive Ingr	edients						
Ingredient Name				Strength			
CARNAUBA WAX							
STARCH, CORN (UNII: O8232NY3S	J)					
FD&C BLUE NO.	1 (UNII: H3R47K3	TBD)					
HYPROMELLOSE,	, UNSPECIFIED	(UNII: 3NXW29V3WO)					
MICROCRYSTALL	INE CELLULOS	E (UNII: OP1R32D61U)					
POVIDONE, UNSI	PECIFIED (UNII:	FZ989GH94E)					
PROPYLENE GLY	COL (UNII: 6DC9	Q167V3)					
STEARIC ACID (U	NII: 4ELV7Z65AP						
SUCROSE (UNII: C	C151H8M554)						
TITANIUM DIOXII	DE (UNII: 15FIX9V	'2JP)					
Product Chai	racteristics						
Color white			Score		2 pieces		
Shape	OVAL (C	Caplet)	Size		15mm		
Flavor			Imprint Cod	int Code		NoDoz	
Contains					nobe		
contains							
Packaging							
# Item Code	Pa	ckage Description	1	Marketing Start Date	Mar	keting End Date	
1 NDC:66715- 9799-4	2 in 1 CARTON		12	2/07/2017		Date	
1	10 in 1 BLISTER PACK; Type 0: Not a Co Product		mbination				
2 NDC:66715- 9799-3	3 in 1 CARTON		12	2/07/2017			
	2 in 1 POUCH; Type 0: Not a Combinatio		n Product				
2		rype of Not a Combinatio					
2 NDC:66715- 9799-6	1 in 1 CARTON	rype 0. Not a combinatio		2/07/2017			

3		Combination Product					
4	NDC:66715- 9799-2	2 in 1 CARTON	06/01/2018	12/30/2023			
4		2 in 1 POUCH; Type 0: Not a Combination Product					
5		10 in 1 VIAL, PLASTIC; Type 0: Not a Combination Product	04/01/2019	4/01/2019			
Marketing Information							
	Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
01	FC monograph fin	al part340	12/07/2017				

Labeler - Lil' Drug Store Products, Inc. (093103646)

Revised: 4/2022

Lil' Drug Store Products, Inc.