TENSION HEADACHE RELIEF- acetaminophen, caffeine tablet, film coated Walgreen Company

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.

Walgreens 44-428

Active ingredients (in each caplet)

Acetaminophen 500 mg (formulated with 65 mg caffeine)

Purpose

Pain reliever

Uses

- temporarily relieves minor aches and pains due to:
 - headache
 - muscular aches

Warnings

Liver warning: This product contains acetaminophen. Severe liver damage may occur if you take

- more than 4,000 mg of acetaminophen in 24 hours
- with other drugs containing acetaminophen
- 3 or more alcoholic drinks every day while using this product

Allergy alert: Acetaminophen may cause severe skin reactions. Symptoms may include:

- blisters
- rash
- skin reddening

If a skin reaction occurs, stop use and seek medical help right away.

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

- with any other drug containing acetaminophen (prescription or nonprescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist.
- if you are allergic to acetaminophen

Ask a doctor before use if you have

liver disease.

Ask a doctor or pharmacist before use if you are

taking the blood thinning drug warfarin.

Stop use and ask a doctor if

- new symptoms occur
- painful area is red or swollen
- pain gets worse or lasts more than 10 days
- fever gets worse or lasts more than 3 days

These could be signs of a serious condition.

If pregnant or breast-feeding,

ask a health professional before use.

Keep out of reach of children.

In case of accidental overdose, get medical help or contact a Poison Control Center (1-800-222-1222) right away. Prompt medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

Directions

- do not use more than directed
- adults and children 12 years and over: take 2 caplets every 6 hours. Do not take more than 6 caplets in 24 hours.
- children under 12 years: ask a doctor

Other information

- 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, crospovidone, D&C red #27 aluminum lake, FD&C blue #2 aluminum lake, FD&C yellow #6 aluminum lake, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, povidone, sodium starch glycolate, stearic acid, talc, titanium dioxide

Questions or comments?

1-800-426-9391

Principal Display Panel

Walgreens

Compare to Excedrin® Tension Headache Caplets active ingredients^{††}

NDC 0363-0428-12

Tension Headache Relief

ACETAMINOPHEN / CAFFEINE / PAIN RELIEVER

ASPIRIN FREE

CAPLETS

100 CAPLETS

ACTUAL SIZE

TAMPER EVIDENT: DO NOT USE IF IMPRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING

Walgreens Pharmacist Recommended Walgreens Pharmacist Survey ^{††}This product is not manufactured or distributed by GSK group of companies, owner of the registered trademark Excedrin® Tension Headache Caplets.

50844 ORG041742857

DISTRIBUTED BY: WALGREEN CO. 200 WILMOT RD., DEERFIELD, IL 60015 *Walgreens* **100% SATISFACTION GUARANTEED** Walgreens.com ©2018 Walgreen Co.



TENSION HEADACHE RELIEF										
acetaminophen, caffeine tablet, film coated										
Pr	oduct Informa	ation								
Pr	oduct T ype		H	HUMAN OTC E	DRUG	Item Code	(Source)	1	NDC:0363-	0428
Ro	ute of Administr	ation	C	ORAL						
Ac	tive Ingredie	nt/Active	Moiet	ty						
	0			, redient Nam	ie			Basis o	f Strength	Strength
AC	ETAMINO PHEN (UNII: 36209	-			O9ITL9D)			-	500 mg
CA	FFEINE (UNII: 3G6	6A5W338E)	(CAFFE	INE - UNII:3G6	A5W338E)			CAFFEIN	E	65 mg
Ina	active Ingredi	ents								
				Ingred	ient Name					Strength
D&	C RED NO. 27 AL	LUMINUM I	L AKE (U	NII: ZK64F7XS	STX)					
TA	LC (UNII: 7SEV7J4	4R1U)								
FDa	&C YELLOW NO	.6 (UNII: H7	77VEI93/	A8)						
CR	O SPO VIDO NE, U	NSPECIFIE	E D (UNII:	2S7830E561)						
	ARCH, CORN (UN									
	GNESIUM STEAF									
	CROCRYSTALLI				61U)					
	VIDONE, UNSPEC			GH94E)						
	EARIC ACID (UNI ANIUM DIO XIDE		,							
	&C BLUE NO. 2				C584)					
	LYETHYLENE GI			, -	,					
	LYVINYL ALCOI									
	DIUM STARCH G				,	42)				
						,				
Pr	oduct Charac	teristics								
Color RED Score no score										
Shape			OVAL		Size				17mm	
Flavor				Imprint Code			44;428			
Co	Contains									
Pa	ckaging									
#	Item Code		1	Package Des	crintion			ting Sta		eting End
				achage Des	-iption			Date		Date
	NDC:0363-0428- 2	0428- 1 in 1 CARTON 01/17/2007			7					
1		100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product				ina tio n				

	1 in 1 CAF	TON	01/17/2007	07/15/2018		
	24 in 1 BC Product	TTLE, PLASTIC; Type 0: Not a Combination				
	1 in 1 CAF	TON	0 1/17/20 0 7	02/07/2021		
	125 in 1 B Product	OTTLE, PLASTIC; Type 0: Not a Combination				
Marketing Information						
<u> </u>						
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC MONOGRAPH NOT FINAL		nart343	01/17/2007			
1	08 NDC:0363-0428- 57 Iarketing Inf Marketing Cate	08 I III I CAR 24 in 1 BO Product 1 in 1 CAR 57 I in 1 CAR 125 in 1 BO Product 125 in 1 BO Product	08If it CARION0824 in 1 BOTLE, PLASTIC; Type 0: Not a Combination ProductNDC:0363-0428- 571 in 1 CARTON125 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination ProductProductInternet Combination ProductInternet Combination Product<	081 In T CARION0 I/1/20070824 in 1 B ∪ TLE, PLASTIC; Type 0: Not a Combination Product0 1/17/2007NDC:0363-0428- 571 in 1 C ARION0 1/17/20071025 in 1 B ∪ TLE, PLASTIC; Type 0: Not a Combination Product0 1/17/2007Image: Specific Strippe 0: Not a Combination Product10 1/17/2007Image: Specific Strippe 0: Not a Combination ProductNation Number or Monograph Citation Marketing Start DateImage: Specific Strippe 0: Not a Combination ProductMarketing Start Date		

Labeler - Walgreen Company (008965063)

Establishment						
Name	Address	ID/FEI	Business Operations			
LNK International, Inc.		038154464	PACK(0363-0428)			

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867894	MANUFACTURE(0363-0428)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		967626305	PACK(0363-0428)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		868734088	PACK(0363-0428)

Revised: 2/2020

Walgreen Company