BANOPHEN- diphenhydramine hcl tablet, film coated
Health Department, Oklahoma State

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

Diphenhydramine 25mg (Emergency Kits)

marked drowsiness may occur
avoid alcoholic beverages
alcohol, sedatives, and tranquilizers may increase drowsiness
use caution when driving a motor vehicle or operating machinery
excitability may occur, especially in children


diphenhydrAMINE HCl
25 MG TABS (PO)
NDC:00904555159
Labeler Code: 83112-5551-1
LOT: P136926
EXP: 11/30/25
OK State Dept of Health
405-426-8000

BANOPHEN

diphenhydramine hcl tablet, film coated

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:83112-555(NDC:0904-5551)
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	25 mg

Product Characteristics

Color	pink	**Score**	no score
Shape	OVAL	**Size**	11mm
Flavor		**Imprint Code**	44;329
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83112-555-59	1 in 1 PACKET; Type 0: Not a Combination Product	01/31/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		01/31/2025	

Labeler - Health Department, Oklahoma State (143673015)

Registrant - Health Department, Oklahoma State (143673015)

Establishment

Name	Address	ID/FEI	Business Operations
Health Department, Oklahoma State		143673015	repack(83112-555)

Revised: 1/2025 Health Department, Oklahoma State