DRY IDEA ANTIPERSPIRANT- dry idea clear gel antiperspirant - powder fresh gel
Henkel Corporation

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.

Dry Idea Antiperspirant Gel - Powder Fresh

Active Ingredient

Aluminum Zirconium Octachlorohydrex GLY 16.4%

Purpose

Antiperspirant

Use

• Reduces underarm perspiration • Extra effective

Warnings

For external use only. Do not use on broken skin.

Stop use and ask a doctor

if rash or irritation occurs.

Keep out of reach of children.

If swallowed, get medical help or contact a poison control center right away.

Ask a doctor before use if

you have kidney disease.

Directions

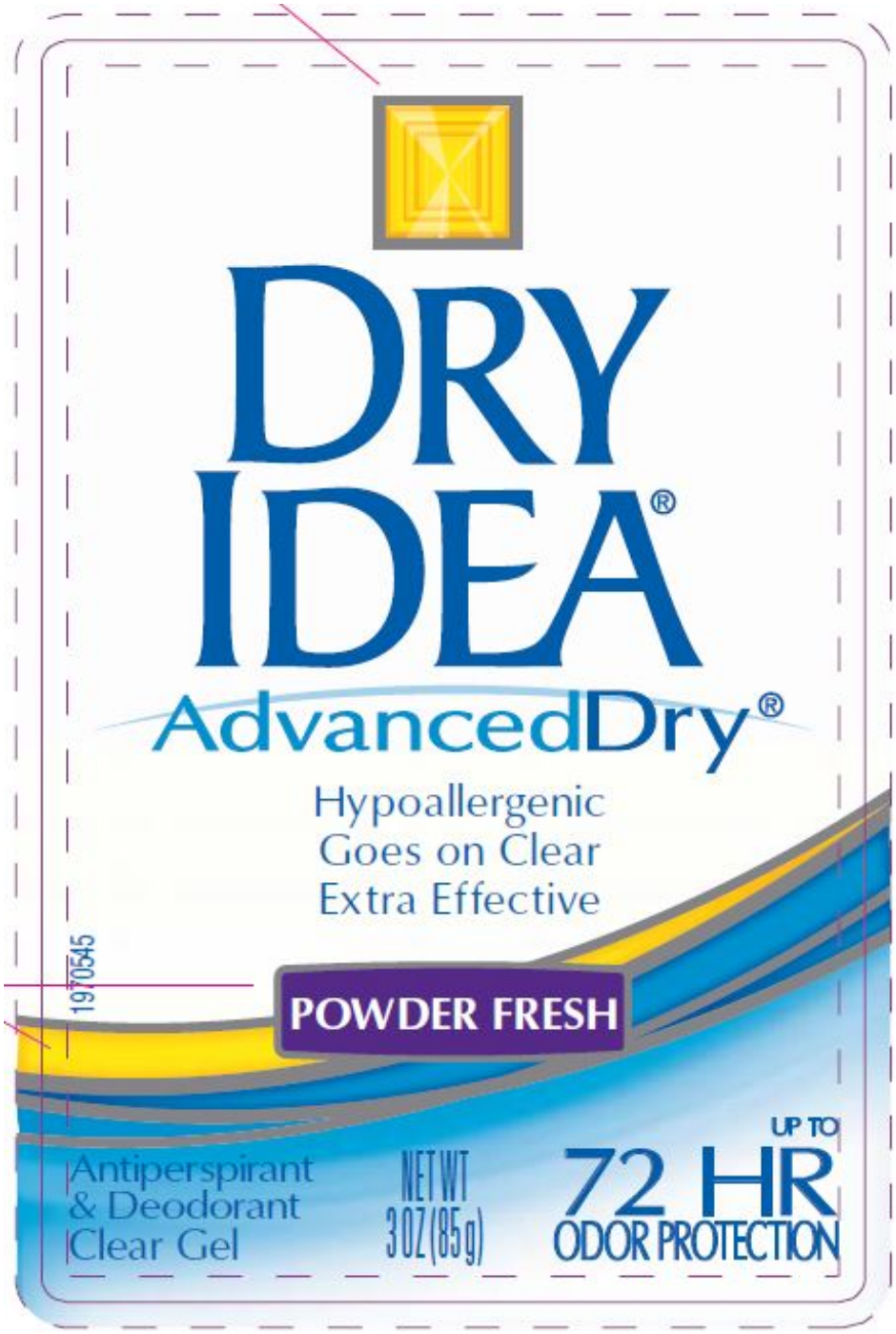
Apply to underarms only.

Inactive Ingredients

Aqua (Water, Eau) • Aluminum Zirconium Octachlorohydrex GLY • Alcohol Denat. • Cyclomethicone • Propylene Glycol • Dimethicone • Calcium Chloride • Trisiloxane • PEG/PPG-18/18 Dimethicone • TButyl Alcohol

Do not freeze or store above 105□F

Questions? 1-800-258-DIAL


DRY IDEA®
AdvancedDry®
Hypoallergenic
Goes on Clear
Extra Effective
1970545
POWDER FRESH
Antiperspirant
& Deodorant
Clear Gel
NET WT
3 OZ (85 g)
UP TO
72 HR
ODOR PROTECTION

DRY IDEA ANTIPERSPIRANT

dry idea clear gel antiperspirant - powder fresh gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:54340-291
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALUMINUM ZIRCONIUM OCTACHLOROHYDREX GLY (UNII: P9D3YP29MY) (ALUMINUM ZIRCONIUM OCTACHLOROHYDREX GLY - UNII:P9D3YP29MY)	ALUMINUM ZIRCONIUM OCTACHLOROHYDREX GLY	16.4 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	46.9 g in 100 g

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:54340-291-54	85 g in 1 CANISTER; Type 0: Not a Combination Product	01/01/2015	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part350	01/01/2015	

Labeler - Henkel Corporation (080887708)

Registrant - Henkel Corporation (080887708)

Establishment

Name	Address	ID/FEI	Business Operations
VVF ILLINOIS SERVICES LLC		024177178	manufacture(54340-291)

Revised: 2/2020 Henkel Corporation