

PLUS BRASEPT ALCOCOL GEL- alcohol gel
NEWDROP NORTH AMERICA LLC

Plus Brasept Alcolol Gel 6605 Drug Facts and Label

Drug Facts Box OTC-Active Ingredient Section

Ethyl Alcohol 70% b/v

Drug Facts Box OTC-Purpose Section

Antiseptic

Drug Facts Box OTC-Indications & Usage Section

for hand-washing to decrease bacteria on the skin, only when water is not available

Drug Facts Box OTC-Warnings Section

FLAMMABLE, keep away from fire and flames

For external use only

Drug Facts Box OTC-When Using Section

do not get into eyes

if contact occurs, rinse eyes thoroughly with water

Drug Facts Box OTC-Stop Use Section

irritation and redness develop

Drug Facts Box OTC-Keep Out of Reach of Children Section

if swallowed, get medical help or contact a Poison Control Center right away

Drug Facts Box OTC-Dosage & Administration Section

wet hands thoroughly with product and allow to dry without wiping

Drug Facts Box OTC-Inactive Ingredient Section

water, DMD hydantoin, diisopropylamine, carbomer, propylene glycol, tocopheryl acetate, albe barbadensis, fragrance

Plus Brasept Alcocol Gel 6605 Drug Facts and Label

PLUS BRASEPT
Alcohol Gel
Instant Hand Sanitizer

Contains
VITAMIN E
and
ALOE VERA

This product is designed exclusively for industrial and institutional use by trained personnel. This product is sold 'as is' and the manufacturer makes no warranty, expressed or implied, of merchantability, and/or fitness for a particular purpose or otherwise.

PLUS BRASEPT

Alcohol Gel

Instant Hand Sanitizer

NEWDROP
USA

DANGER: FLAMMABLE
KEEP OUT OF REACH OF CHILDREN
KEEP AWAY FROM FIRE OR FLAME. FOR EXTERNAL USE ONLY.
Si usted no entiende la etiqueta, busque a alguien para que se la explique a usted en detalle.
(If you do not understand this label, find someone to explain it to you in detail.)
NET CONTENTS: 1 GALLON (3.8 L)

| Active Ingredient | Purpose |
|---|------------|
| Ethyl Alcohol 70% v/v | Antiseptic |
| Use for hand-washing to decrease bacteria on the skin, only when water is not available. | |
| Warnings Flammable, keep away from fire and flames For external use only. | |
| When using this product | |
| <ul style="list-style-type: none"> ■ do not get into eyes ■ if contact occurs, rinse eyes thoroughly with water ■ stop use and ask a doctor if ■ irritation and redness develop | |
| Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away. | |
| Directions ■ wet hands thoroughly with product and allow to dry without wiping | |
| Inactive Ingredients water, DMDM hydantoin, diisopropylamine, carbomer, propylene glycol, tocopheryl acetate, aloe barbadensis | |

NEWDROP
USA

Sold by: **Newdrop North America LLC**
3480 Oakcreek Rd, Suite D2, Doraville, GA, 30340
(470) 275-3196

6605413000000000

PLUS BRASEPT ALCOCOL GEL

alcohol gel

Product Information

| | | | |
|--------------------------------|----------------|---------------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:83318-605 |
| Route of Administration | TOPICAL | | |

Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|---|-------------------|----------------|
| ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M) | ALCOHOL | 0.7 mL in 1 mL |

Inactive Ingredients

| Ingredient Name | Strength |
|---|----------|
| WATER (UNII: 059QF0KO0R) | |
| CARBOMER 934 (UNII: Z135WT9208) | |
| PROPYLENE GLYCOL (UNII: 6DC9Q167V3) | |
| ALOE VERA LEAF (UNII: ZY81Z83H0X) | |
| .ALPHA.-TOCOPHEROL ACETATE, DL- (UNII: WR1WPI7EW8) | |
| DIISOPROPYLAMINE (UNII: BR9JLI40NO) | |
| DMDM HYDANTOIN (UNII: BYR0546TOW) | |

Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|------------------|---|----------------------|--------------------|
| 1 | NDC:83318-605-14 | 3785 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product | 03/24/2023 | 11/20/2025 |
| 2 | NDC:83318-605-08 | 208200 mL in 1 DRUM; Type 0: Not a Combination Product | 06/25/2024 | |

Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|--------------------|--|----------------------|--------------------|
| OTC Monograph Drug | 505G(a)(3) | 03/24/2023 | |

Labeler - NEWDROP NORTH AMERICA LLC (094392481)

Registrant - ABC Compounding Co., Inc. (003284353)

Establishment

| Name | Address | ID/FEI | Business Operations |
|---------------------------|---------|-----------|------------------------|
| ABC Compounding Co., Inc. | | 003284353 | manufacture(83318-605) |

Revised: 11/2025

NEWDROP NORTH AMERICA LLC