IMAGE SKINCARE CLEAR CELL CLARIFYING ACNE SPOT TREATMENT- salicylic acid gel
Natural Technology, LLC

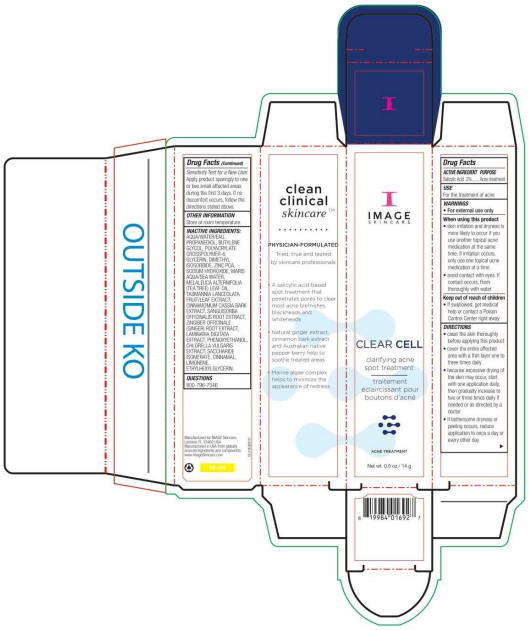
Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Salicylic Acid 2%. Purpose: Acne Treatment

CLEAR CELL CLARIFYING
ACNE SPOT TREATMENT
AND CLEAR CELL CLARIFYING
SALICYLIC BLEMISH GEL

Assigned Color Swatches

BLACK 288 C WHITE SILVER FOIL EMBOSSED



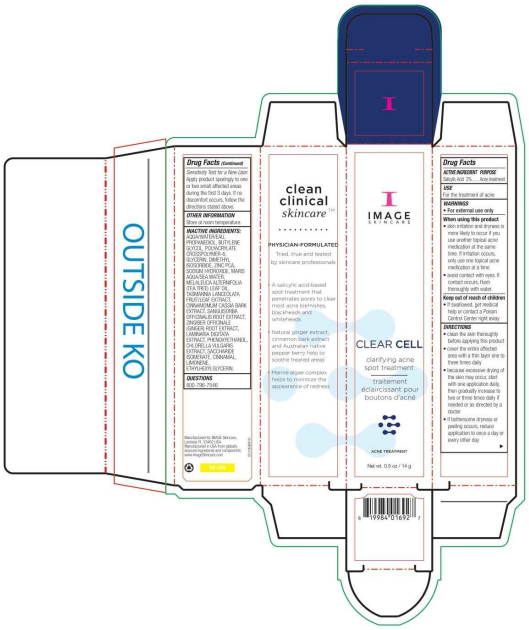
14g. NDC: 79458-002-00

Purpose: Acne Treatment

CLEAR CELL CLARIFYING
ACNE SPOT TREATMENT
AND CLEAR CELL CLARIFYING
SALICYLIC BLEMISH GEL

Assigned Color Swatches

BLACK | 288 C | WHITE | SILVER FOIL | EMBOSSED



14 g. NDC: 79458-002-00

For external use only

CLEAR CELL CLARIFYING
ACNE SPOT TREATMENT
AND CLEAR CELL CLARIFYING
SALICYLIC BLEMISH GEL

Assigned Color Swatches

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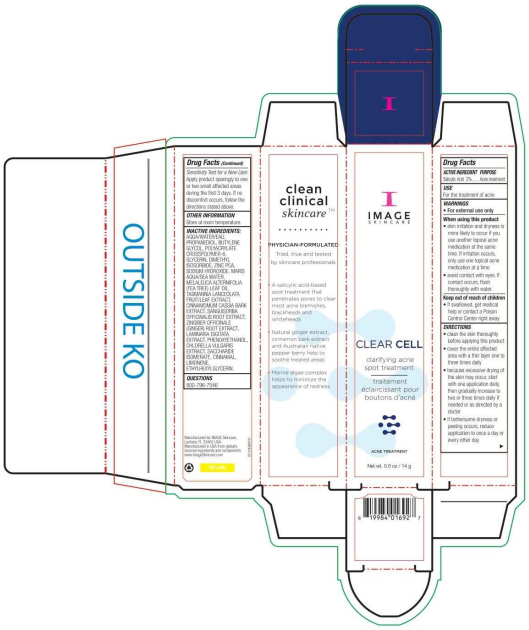

14 g. NDC: 79458-002-00

- Skin irritation and dryness is more likely to occur if you use another topical acne medication at the same time. If irritation occurs, only use one topical acne medication at a time.
- Avoid contact with eyes. If contact occurs, flush thoroughly with water.

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SALICYLIC BLEMISH GEL

Assigned Color Swatches

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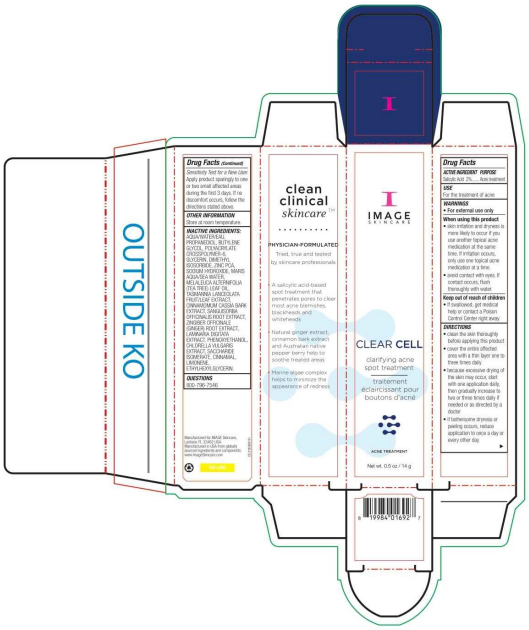

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If swallowed, get medical help or contact Poison Control Center right away.

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ACNE SPOT TREATMENT
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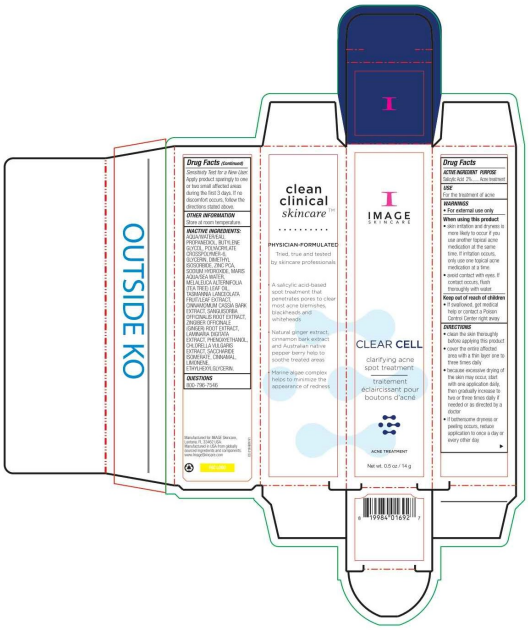

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- Clean the skin thoroughly before applying this product
- Cover the entire affected area with a thin layer one to three times daily
- Because excessive drying of the skin may occur, start with one application daily, then gradually increase to two or three times daily if needed or as directed by a doctor
- If bothersome dryness or peeling occurs, reduce application to once a day or every other day

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ACNE SPOT TREATMENT
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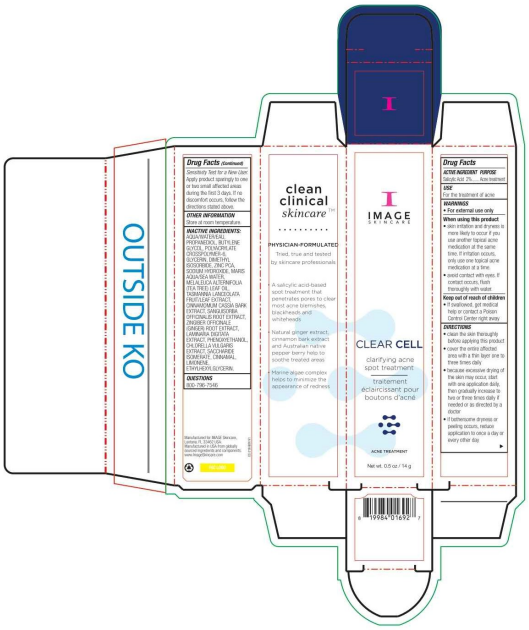

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Sensitivity Test for a New User.
Apply product sparingly to one or two small affected areas during the first three days. If no discomfort occurs, follow the directions stated above.

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ACNE SPOT TREATMENT
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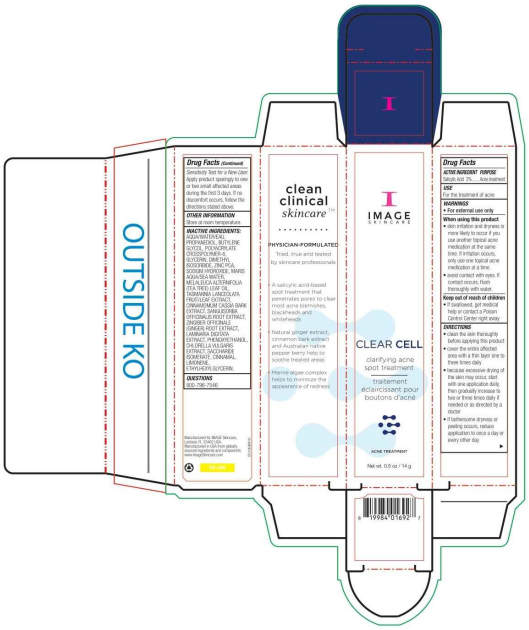

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Store at room temperature.

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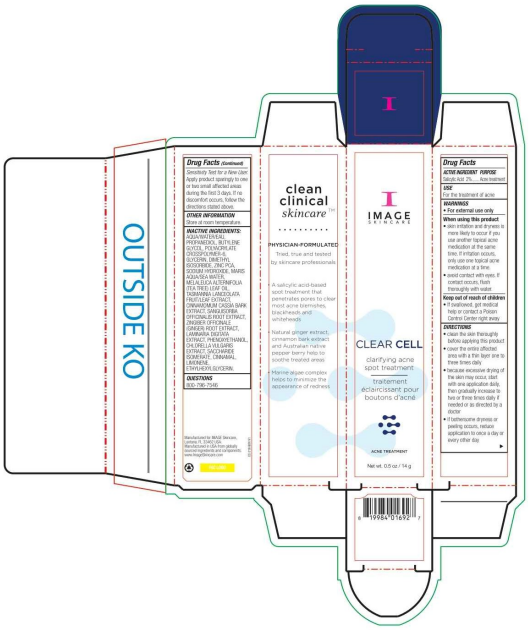

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Aqua/Water/EAU, Propanediol, Butylene Glycol, Polyacrylate Crosspolymer-6, Glycerin, Dimethyl Isosorbide, Zinc PCA, Sodium Hydroxide, Maris Aqua/Sea Water, Melaleuca Alternifolia (Tea Tree) Leaf Oil, Tasmannia Lanceolata Fruit/Leaf Extract, Cinnamomum Cassia Bark Extract, Sanguisorba Officinalis Root Extract, Zingiber Officinale (Ginger) Root Extract, Laminaria Digitata Extract, Phenoxyethanol, Chlorella Vulgaris Extract, Saccharide Isomerate, Cinnamal, Limonene, Ethylhexylglycerin.

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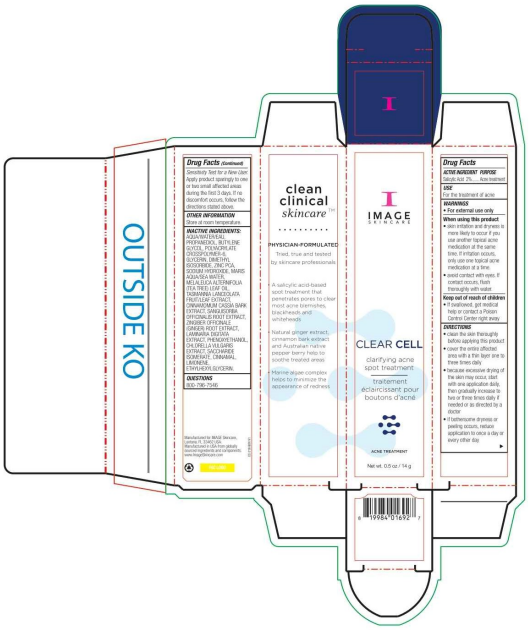

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Clear Cell clarifying acne spot treatment

CLEAR CELL CLARIFYING
ACNE SPOT TREATMENT
AND CLEAR CELL CLARIFYING
SALICYLIC BLEMISH GEL

Assigned Color Swatches

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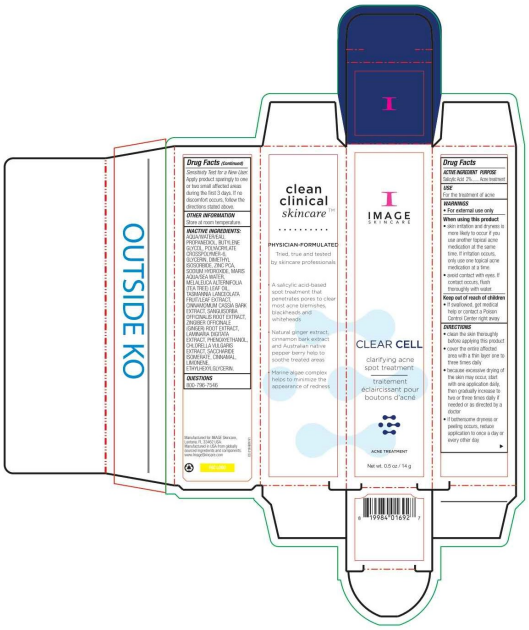

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IMAGE SKINCARE CLEAR CELL CLARIFYING ACNE SPOT TREATMENT

salicylic acid gel

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:79458-002
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	20 g in 1000 g

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
CHLORELLA VULGARIS (UNII: RYQ4R60M02)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
TASMANNIA LANCEOLATA LEAF (UNII: H8C2R3TXSF)	
WATER (UNII: 059QF0KO0R)	

AMMONIUM ACRYLOYLDIMETHYLTAURATE, DIMETHYLACRYLAMIDE, LAURYL METHACRYLATE AND LAURETH-4 METHACRYLATE COPOLYMER, TRIMETHYLOLPROPANE TRIACRYLATE CROSSLINKED (45000 MPA.S) (UNII: Q7UI015FF9)	
TEA TREE OIL (UNII: VIF565UC2G)	
TASMANNIA LANCEOLATA FRUIT (UNII: PNT2HDL13Q)	
CHINESE CINNAMON (UNII: WS4CQ062KM)	
SANGUISORBA OFFICINALIS ROOT (UNII: 4NYV2HT01X)	
GINGER (UNII: C5529G5JPQ)	
LAMINARIA DIGITATA (UNII: 15E7C67EE8)	
SACCHARIDE ISOMERATE (UNII: W8K377W98I)	
CINNAMALDEHYDE (UNII: SR60A3XG0F)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
PROPANEDIOL (UNII: 5965N8W85T)	
ZINC PIDOLATE (UNII: C32PQ86DH4)	
LIMONENE, (+)- (UNII: GFD7C86Q1W)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
DIMETHYL ISOSORBIDE (UNII: SA6A6V432S)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79458-002-01	1 in 1 BOX	03/17/2022	
1	NDC:79458-002-00	14 g in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333D	03/17/2022	

Labeler - Natural Technology, LLC (618561906)

Revised: 3/2022 Natural Technology, LLC