

**DUOLAQ- salicylic acid gel**  
**METHODIQ INC.**

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**Duolaq**

**Active ingredients**

Salicylic Acid 2%

**Purpose**

Salicylic Acid 2%.....Acne Treatment

**Uses**

- For the management / treatment of acne.
- Clears acne blemishes and blackheads.
- Helps prevent the development of new acne blemishes and blackheads

**Warnings**

For external use only

**When using this product**

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.

**Keep out of reach of children.**

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

**Directions**

- 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.

**Inactive ingredients**

Water, Glycerin, Ammonium Polyacryloyldimethyl Taurate, Propanediol, Sodium Polyacrylate, Butylene Glycol, Pentylene Glycol, Glycolic Acid, Dimethicone, Propylene Glycol, Dimethyl Isosorbide, 4-t-Butylcyclohexanol, Phenoxyethanol, Niacinamide, Xylitol, Caprylic Acid, Caprylyl Glycol, Allantoin, Hexylene Glycol, Ethylhexylglycerin, Chamomilla Recutita (Matricaria) Flower Extract, Hydrolyzed Sodium Hyaluronate, Citric Acid, Glycyrrhiza Glabra (Licorice) Root Powder, Fomes Officinalis (Mushroom) Extract,

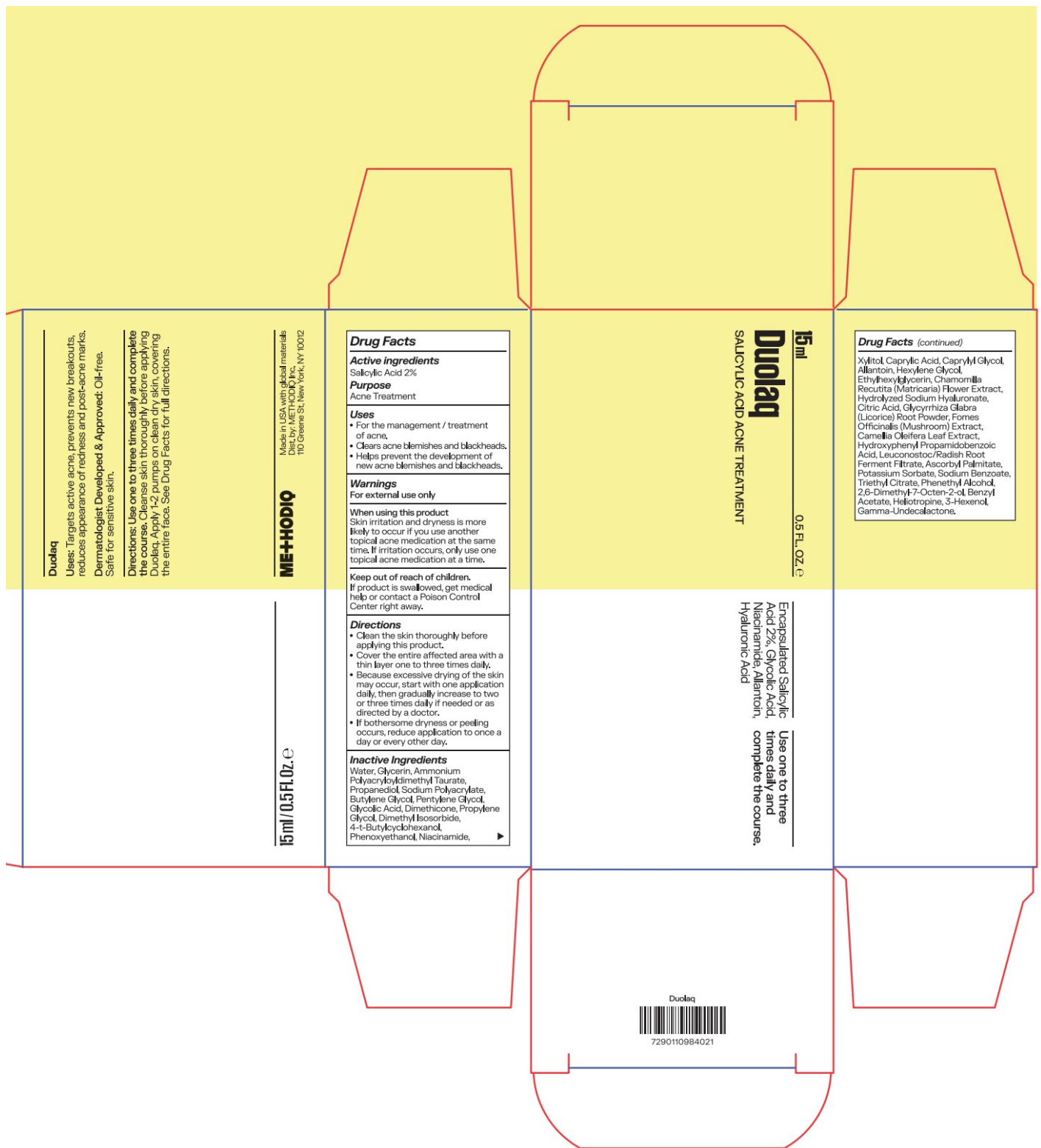
Camellia Oleifera Leaf Extract, Hydroxyphenyl Propamidobenzoic Acid, Leuconostoc/Radish Root Ferment Filtrate, Ascorbyl Palmitate, Potassium Sorbate, Sodium Benzoate, Triethyl Citrate, Phenethyl Alcohol, 2,6-Dimethyl-7-Octen-2-ol, Benzyl Acetate, Heliotropine, 3-Hexenol, Gamma-Undecalactone.

15ml

0.5 Fl. Oz.

Duolaq

Salicylic Acid Acne Treatment



DUOLAQ			
salicylic acid gel			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:87110-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name		Basis of Strength	Strength
SALICYLIC ACID (UNII: 0414PZ4LPZ) (SALICYLIC ACID - UNII:0414PZ4LPZ)		SALICYLIC ACID	20 mg in 1 mL
Inactive Ingredients			
Ingredient Name			Strength
LARICIFOMES OFFICINALIS FRUITING BODY (UNII: 7IFM8431X3)			
PENTYLENE GLYCOL (UNII: 50C1307PZG)			
DIMETHYL ISOSORBIDE (UNII: SA6A6V432S)			
4-T-BUTYLCYCLOHEXANOL (UNII: K0H1405S9C)			
TRIETHYL CITRATE (UNII: 8Z96QXD6UM)			
DIMETHICONE (UNII: 92RU3N3Y1O)			
ALLANTOIN (UNII: 344S277G0Z)			
NIACINAMIDE (UNII: 25X51I8RD4)			
GAMMA-UNDECALACTONE (UNII: QB1T0AG2YL)			
GLYCOLIC ACID (UNII: 0WT12SX38S)			
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)			
CAPRYLIC ACID (UNII: OBL58JN025)			
CAPRYLYL GLYCOL (UNII: 00YIU5438U)			
CHAMOMILE (UNII: FGL3685T2X)			
SODIUM HYALURONATE (UNII: YSE9PPT4TH)			
CITRIC ACID (UNII: 2968PHW8QP)			
HYDROXYPHENYL PROPAMIDOBENZOIC ACID (UNII: 25KRT26H77)			
ASCORBYL PALMITATE (UNII: QN83US2B0N)			
HELIOTROPINE (UNII: KE109YAK00)			
WATER (UNII: 059QF0KO0R)			
GLYCERIN (UNII: PDC6A3C0OX)			
PHENETHYL ALCOHOL (UNII: ML9LGA7468)			
GLYCYRRHIZA GLABRA (LICORICE) ROOT POWDER (UNII: 2788Z9758H)			
CAMELLIA OLEIFERA LEAF (UNII: 5077EL0C60)			
SODIUM BENZOATE (UNII: OJ245FE5EU)			
PROPANEDIOL (UNII: 5965N8W85T)			
2,6-DIMETHYL-7-OCTEN-2-OL (UNII: 46L1B02ND9)			
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)			
BENZYL ACETATE (UNII: 0ECG3V79ZJ)			
XYLITOL (UNII: VCQ006KQ1E)			
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)			
LEUCONOSTOC/RADISH ROOT FERMENT FILTRATE (UNII: D2QHA03458)			
3-HEXENOL (UNII: V14F8G75P4)			
AMMONIUM POLYACRYLOYLDIMETHYL TAURATE (UNII: F01RIY4371)			
SODIUM POLYACRYLATE (8000 MW) (UNII: 285CYO341L)			
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)			
PHENOXYETHANOL (UNII: HIE492ZZ3T)			
HEXYLENE GLYCOL (UNII: KEH0A3F75J)			
Packaging			
		Marketing Start	Marketing End

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:87110-001-01	1 in 1 CARTON	11/18/2025	
1		15 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M006	11/18/2025	

**Labeler -**
METHODIQ INC. (119377051)