

**W.SKIN TREEMON CLEANSING WATER- citric acid liquid**  
**J&J COMPANY**

*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.*

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**J&J COMPANY - STOP-AGING PEPTIDE TONER**

citric acid

Water, Dipropylene Glycol, PEG-6 Caprylic/Capric Glycerides, PEG-7 Glyceryl Cocoate, Phenoxyethanol, Sodium Citrate, Ethylhexylglycerin, Disodium EDTA, Citric Acid, Butylene Glycol, Polyglyceryl-10 Myristate, Hydroxyacetophenone, Pentylene Glycol, Caprylyl Glycol, Lavandula Angustifolia (Lavender) Extract, Melissa Officinalis Leaf Extract, Mentha Piperita (Peppermint) Leaf Extract, Monarda Didyma Leaf Extract, 1,2-Hexanediol, Cetyl Ethylhexanoate, Caprylic/Capric Triglyceride, Ethylhexyl Palmitate, Sodium Hyaluronate, Camellia Japonica Flower Extract, Melaleuca Alternifolia (Tea Tree) Leaf Extract, Salvia Officinalis (Sage) Leaf Extract

Maintaining balance between the t-zone. Dry skin or peeling, cleaning and purification of the skin of acne. Peeling or soft cleaning without simulation and water supply

keep out of reach of the children

Apply twice a day with a cotton pad. Do not need to rinse.

Avoid contact with eyes. In case of direct contact with the eyes, rinse thoroughly. Do not ingest. For external use only. Do not to be used for children under 3years of age.Keep out of reach of children

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PRODUCT NAME: TREEMON CLEANSING WATER

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INGREDIENTS:

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## W.SKIN TREEMON CLEANSING WATER

citric acid liquid

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:74997-0022
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>CITRIC ACID MONOHYDRATE</b> (UNII: 2968PHW8QP) (ANHYDROUS CITRIC ACID - UNII:XF417D3PSL)	ANHYDROUS CITRIC ACID	0.03 g in 100 mL

### Inactive Ingredients

Ingredient Name	Strength
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**HYALURONATE SODIUM** (UNII: YSE9PPT4TH)

**WATER** (UNII: 059QF0KO0R)

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:74997-0022-1	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/08/2021	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		05/08/2021	

**Labeler** - J&J COMPANY (688995236)

**Registrant** - J&J COMPANY (688995236)

### Establishment

Name	Address	ID/FEI	Business Operations
J&J COMPANY		688995236	manufacture(74997-0022) , label(74997-0022) , pack(74997-0022)

Revised: 5/2021

J&J COMPANY