LIQUICAL PLUS- multivitamin liquid
PureTek Corporation

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.

LiquiCal Plus

Liquid Multivitamin

DRUG DESCRIPTION:

LiquiCal™ Plus
Rx Only
Full Prescribing Information

Active Ingredients:
Each 5 mL Contains:
Vitamin D3 (as Cholecalciferol) ..... 10 mcg (400 IU)
Calcium (as Calcium Citrate) ..... 84 mg
Magnesium (as Magnesium Citrate) ..... 24 mg
Zinc (as Zinc Citrate) ..... 0.7 mg

Each 15 mL Contains:
Vitamin D3 (as Cholecalciferol) ..... 30 mcg (1200 IU)
Calcium (as Calcium Citrate) ..... 252 mg
Magnesium (as Magnesium Citrate) ..... 72 mg
Zinc (as Zinc Citrate) ..... 2.1 mg

Other Ingredients: Agave Tequilana (Tequila Agave) Stem Extract, Aqua (Purified Water), Citric Acid, Orange Flavor, Potassium Sorbate, Xanthan Gum.

INDICATIONS:

LiquiCal™ Plus is indicated for dietary management of patients with unique nutritional needs requiring increased Vitamin D3 and essential minerals, including Calcium, Magnesium, and Zinc in a convenient liquid form for easy administration.

CLINICAL PHARMACOLOGY

Vitamin D3 is essential for calcium absorption in the gut and maintaining adequate serum calcium and phosphate concentrations, supporting bone mineralization and immune function.

Calcium is crucial for the development and maintenance of healthy bones and teeth, as well as muscle function.

Magnesium is important for muscle and nerve function, blood glucose control, and bone health.

Zinc supports normal growth and development, immune function, and wound healing.

CONTRAINDICATIONS:

This product is contraindicated in patients with a known hypersensitivity to any of the ingredients. **LiquiCal™ Plus** is contraindicated in patients with hypercalcemia, malabsorption syndrome, abnormal sensitivity to the toxic effects of vitamin D, and hypervitaminosis D.

WARNINGS AND PRECAUTIONS:

Tell your doctor if you have: kidney problems, thyroid disease. This medication should be used as directed during pregnancy or while breastfeeding. Consult your doctor.

ADVERSE REACTIONS:

Adverse reactions are rare but may include gastrointestinal disturbances such as nausea or constipation. If adverse reactions occur, discontinue use and consult a healthcare professional.

You should call your doctor for medical advice about serious adverse events. To report adverse side effects or to obtain product information, contact **PureTek Corporation**, at **1-877-921-7873** or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch.

DOSAGE AND ADMINISTRATION:

Shake well before use.
For Children 1-3 years: 1-2 teaspoons (5-10 mL) daily
For Children 4+ years: 1 tablespoon (15 mL) up to 2 times daily

LiquiCal™ Plus should be administered under the supervision of a licensed healthcare practitioner.

HOW SUPPLIED:

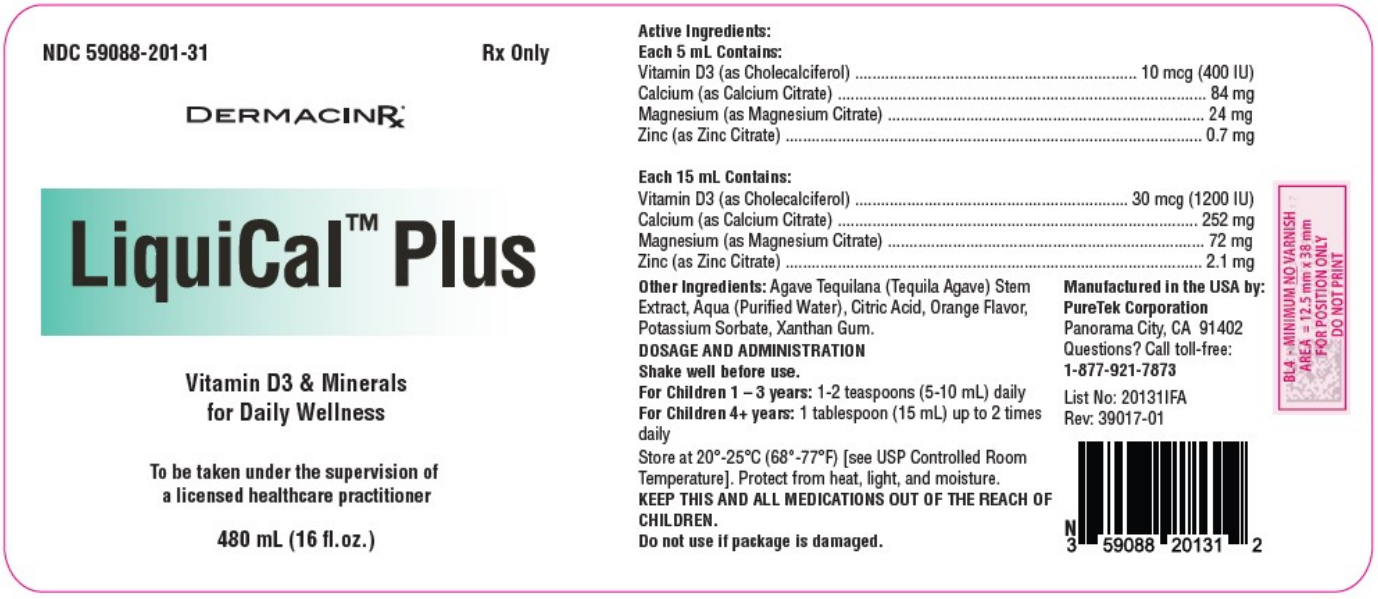
LiquiCal™ Plus is available in a 480 mL white HDPE bottle with NDC 59088-201-31.

STORAGE:

KEEP THIS AND ALL MEDICATIONS OUT OF THE REACH OF CHILDREN.
Store at 20°-25°C (68°-77°F) [see USP Controlled Room Temperature].
Protect from heat, light, and moisture.
Tamper Evident: Do not use if the seal is broken or missing

LiquiCal™ Plus

Manufactured in the USA by:
PureTek Corporation
Panorama City, CA 91402
For questions or information call toll-free: **877-921-7873**



LIQUICAL PLUS
multivitamin liquid

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:59088-201
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
CHOLECALCIFEROL (UNII: 1C6V77QF41) (CHOLECALCIFEROL - UNII:1C6V77QF41)	CHOLECALCIFEROL	1200 [iU] in 1 mL
ZINC CITRATE (UNII: K72I3DEX9B) (ZINC CATION - UNII:13S1S8SF37)	ZINC CATION	2.1 mg in 1 mL
MAGNESIUM CITRATE (UNII: RHO26O1T9V) (MAGNESIUM CATION - UNII:T6V3LHY838)	MAGNESIUM CITRATE	72 mg in 1 mL
CALCIUM CITRATE (UNII: MLM29U2X85) (CALCIUM CATION - UNII:2M83C4R6ZB)	CALCIUM CITRATE	252 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
WATER (UNII: 059QF0KO0R)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
XANTHAN GUM (UNII: TTV12P4NEE)	
AGAVE TEQUILANA STEM (UNII: J026JA743Y)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59088-201-31	480 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/12/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		06/12/2024	

Labeler - PureTek Corporation (785961046)

Revised: 9/2024 PureTek Corporation