CALCAREA FLUORICA- calcarea fluorica spray
Ratis, LLC

Disclaimer: This homeopathic product has not been evaluated by the Food and Drug Administration for safety or efficacy. FDA is not aware of scientific evidence to support homeopathy as effective.

DRUG FACTS:

ACTIVE INGREDIENT:

(in each spray) Calcarea Fluorica 12X 100%.

PURPOSES:

Calcarea Fluorica - elasticity*

USES:

Supports the health of bones, teeth, skin, joints, muscles, and veins.*

*Claims based on traditional homeopathic practice, not accepted medical evidence. Not FDA evaluated.

WARNINGS:

For oral use only.

If pregnant or breast-feeding, ask a health care professional before use.

Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.

DO NOT USE IF TAMPER EVIDENT SEAL IS BROKEN OR MISSING

KEEP OUT OF REACH OF CHILDREN:

In case of overdose, get medical help or contact a Poison Control Center right away.

DIRECTIONS:

• Adults: 2 sprays 3 times a day in mouth or half a cup of water.

• Children under 12: 1 spray as above.

• Consult a physician for use in children under 12 years of age.

Store in a cool, dry place.

INACTIVE INGREDIENTS:

Demineralized water, 20% organic ethanol

QUESTIONS:

www.HomeopathyStore.com

(888) 405-7551

Dist. by Ratis, LLC

1201 N Orange

Ste 7594

Wilmington, DE 19801

PACKAGE LABEL DISPLAY:

SCHUESSLER CELL SALTS

LACTOSE FREE

ANNA

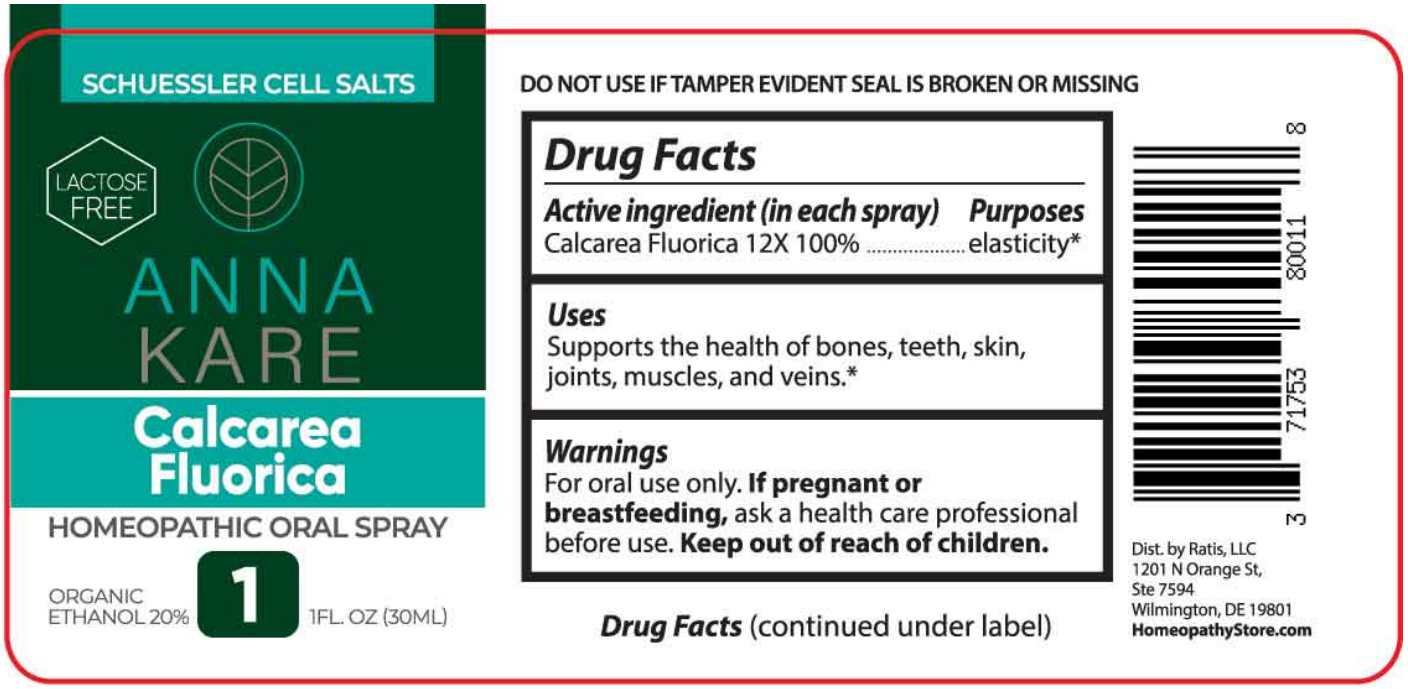
KARE

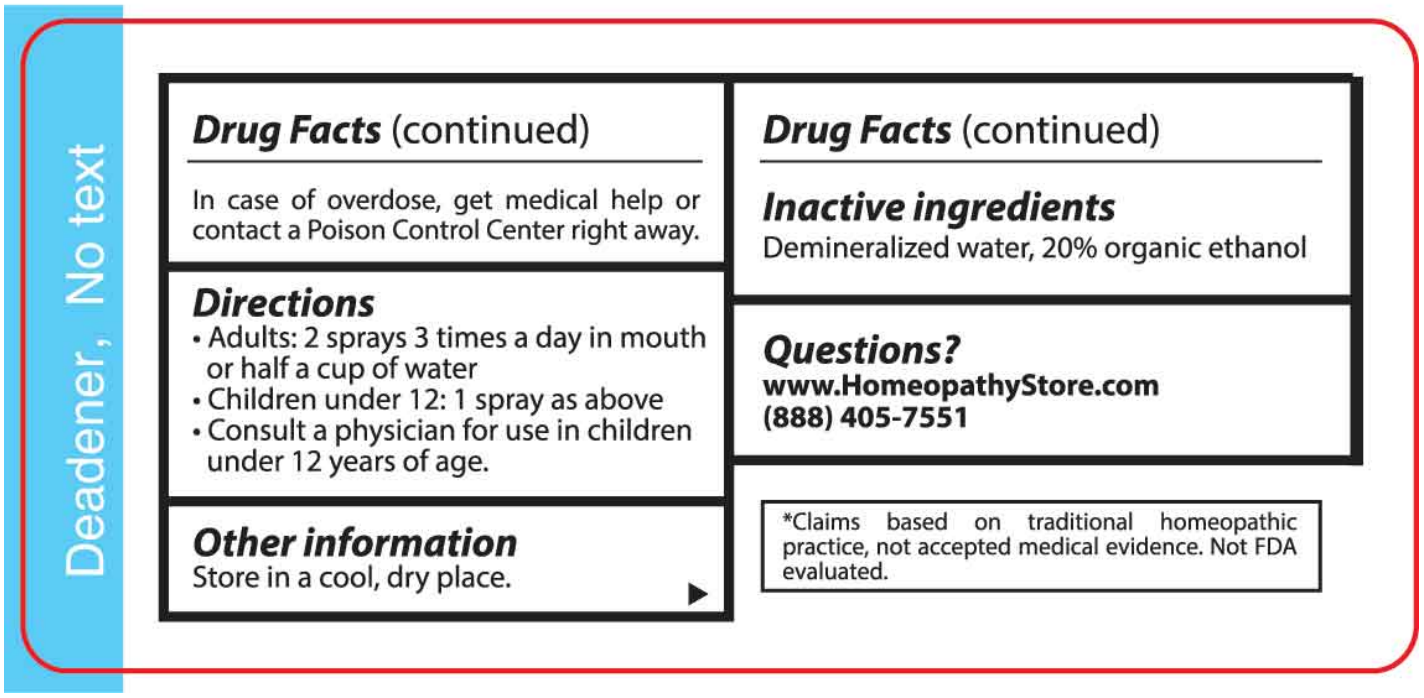
Calcarea

Fluorica

HOMEOPATHIC ORAL SPRAY

1 FL. OZ (30ML)





CALCAREA FLUORICA

calcarea fluorica spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71753-8001
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**

CALCIUM FLUORIDE (UNII: O3B55K4YKI) (FLUORIDE ION - UNII:Q80VPU408O)	CALCIUM FLUORIDE	12 [hp_X] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71753-8001-1	30 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	09/30/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		09/30/2020	

Labeler - Ratis, LLC (964594324)

Revised: 7/2023 Ratis, LLC