

OXALIS MONTANA 6X- oxalis montana 6x globule
True Botanica, 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.

Oxalis Montana 6X

Oxalis Montana 6X, HPUS

The letters HPUS indicate that these ingredients are officially included in the Homeopathic Pharmacopoeia of the United States.

For temporary relief of fatigue. Helps to overcome metabolic sluggishness. May also be used for standard homeopathic indications or as directed by your physician.

Use:

For temporary relief of fatigue. Helps to overcome metabolic sluggishness. May also be used for standard homeopathic indications or as directed by your physician.

Warnings:

Consult your health care provider if symptoms persist more than 5 days or worsen.

If pregnant or breastfeeding ask a health professional before use.

Keep out of the reach of children.

TAMPER EVIDENT: Do not use if safety seal is broken before first use.

Directions for Use:

Place globules under tongue for 30 seconds. Adults ~3-5 globules 3 times a day.

Children 2-12 years ~ 1-3 globules 3 times a day. Under 2 years, consult your doctor.

Use until symptoms are relieved or as directed by your health care provider.

Other Ingredients:

Sucrose

Developed and Made by

True Botanica, LLC

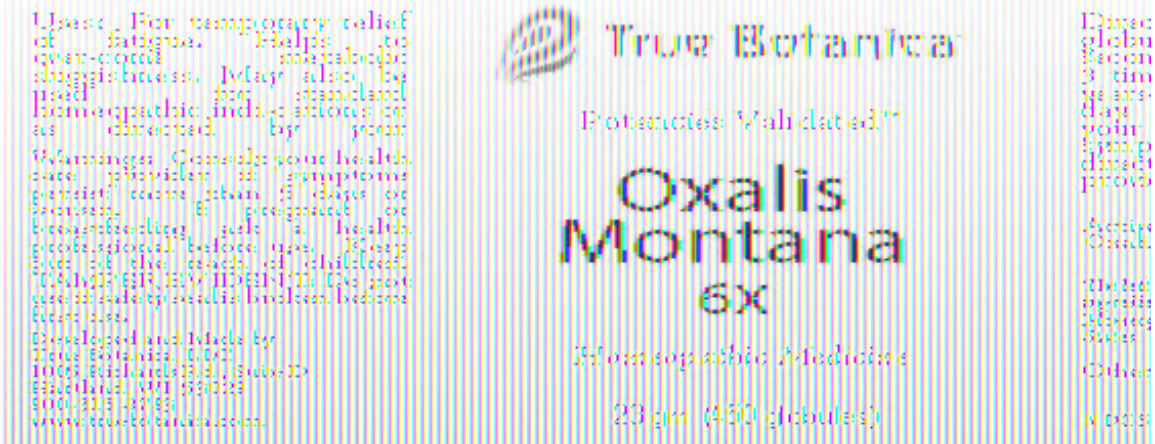
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Principal Display Panel



OXALIS MONTANA 6X

oxalis montana 6x globule

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:53645-1950
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
OXALIS ACETOSELLA WHOLE (UNII: GON5VU1D0Z) (OXALIS ACETOSELLA WHOLE - UNII:GON5VU1D0Z)	OXALIS ACETOSELLA WHOLE	6 [hp_X] in 23 g

Inactive Ingredients

Ingredient Name	Strength
SUCROSE (UNII: C151H8M554)	

Product Characteristics

Color	white	Score	no score
Shape	ROUND	Size	4mm
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:53645-1950-2	23 g in 1 BOTTLE; Type 0: Not a Combination Product	06/02/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		06/02/2023	

Labeler - True Botanica, LLC (182071519)

Establishment

Name	Address	ID/FEI	Business Operations
True Botanica, LLC		182071519	manufacture(53645-1950)

Revised: 6/2023

True Botanica, LLC