# SCHUESSLER TISSUE SALTS COMB D SKIN DISORDERS- kali muriaticum, kali sulphuricum, calcarea sulphurica, silicea tablet, chewable Martin & Pleasance Pty Ltd

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.

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## Active Ingredients Purpose

Calcarea Sulphurica 6X HPUS	skin disorders, acne
Kali Muriaticum 6X HPUS	minor skin eruptions
Kali Sulphuricum 6X HPUSdry	, scaly & flaky skin, minor skin eruptions
Silicea 6X HPUS	skin disorders, minor skin eruptions

<sup>&</sup>quot; HPUS" is the Homeopathic Pharmacopoeia of the United States

#### Uses

Temporarily relieves symptoms associated with

- Skin disorders
- Minor skin eruptions
- Acne
- Dry, scaly & flaky skin

#### Warnings

#### Stop use and ask a doctor if:

- Symptoms persist for more than 7 days or worsen
- Inflammation, fever or infection is present or develops
- Fever gets worse or lasts more than 3 days
- If skin eruptions are severe, or painful, seek the advice of a doctor.

If pregnant or breastfeeding, ask a health professional before use. Keep out of reach of children.

#### **Directions**

- Use as directed
- Take 30 minutes away from food.

#### Adults:

- Acute dose: Chew 1 tablet every 30 minutes until symptoms subside, up to 6 times per day.
- Maintenance dose: Chew 1 tablet, 4 times daily.

#### Children 6 years and older:

- Acute dose: Crush ½ tablet, dissolve in water, and take every 30 minutes until symptoms subside, up to 6 times per day.
- Maintenance dose: Crush ½ tablet, dissolve in water, 4 times per day.

#### Other Information

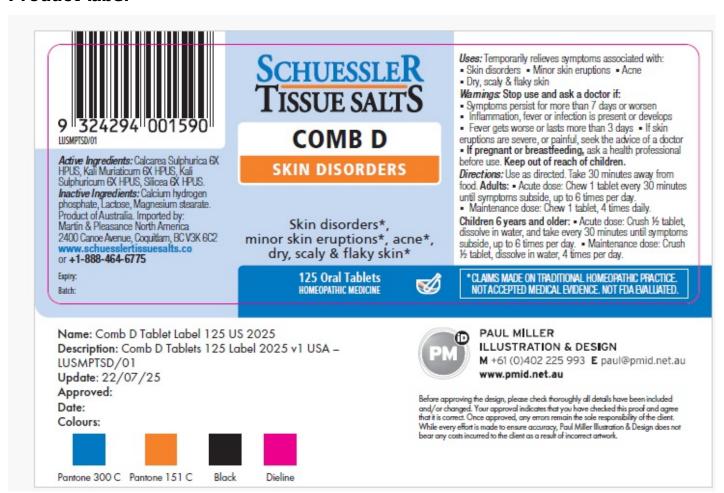
Store below 86°F

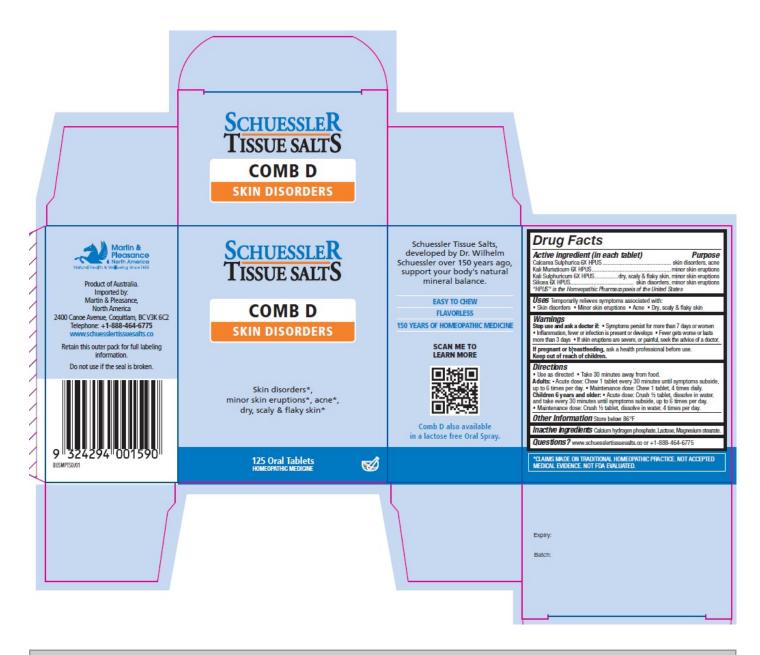
#### **Inactive Ingredients**

Calcium hydrogen phosphate, Lactose, Magnesium stearate.

Questions? www.schuesslertissuesalts.co or + 1-888-464-6775

#### Product label





#### SCHUESSLER TISSUE SALTS COMB D SKIN DISORDERS

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Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:84999-026	
Route of Administration	ORAL			

Active Ingredient/Active Moiety			
Ingredient Name	<b>Basis of Strength</b>	Strength	
POTASSIUM CHLORIDE (UNII: 660YQ98I10) (POTASSIUM CATION - UNII:295053K152)	POTASSIUM CATION	6 [hp_X]	
POTASSIUM SULFATE (UNII: 1K573LC5TV) (POTASSIUM CATION - UNII:295053K152)	POTASSIUM SULFATE	6 [hp_X]	
CALCIUM SULFATE DIHYDRATE (UNII: 4846Q921YM) (POTASSIUM CATION - UNII:295053K152)	CALCIUM SULFATE DIHYDRATE	6 [hp_X]	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4) (SAND - UNII:ETJ7Z6XBU4)	SILICON DIOXIDE	6 [hp_X]	

Inactive Ingredients			
Ingredient Name	Strength		
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)			
MAGNESIUM STEARATE (UNII: 70097M6I30)			
ANHYDROUS DIBASIC CALCIUM PHOSPHATE (UNII: L11K75P92J)			

Product Characteristics			
Color	white	Score	no score
Shape	OVAL	Size	12mm
Flavor		Imprint Code	
Contains			

Packaging				
# Item Code	Package Description	Marketing Start Date	Marketing End Date	
NDC:84999-026-	1 in 1 CARTON	08/01/2025		
1	125 in 1 BOTTLE; Type 0: Not a Combination Product			

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		08/01/2025	

### Labeler - Martin & Pleasance Pty Ltd (752869859)

Revised: 8/2025 Martin & Pleasance Pty Ltd