

**SULFADERM- sulfur ointment**  
**Pharmadel LLC**

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**Sulfaderm Acne Treatment (WP)(DA)**

***Drug Facts***

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***Active ingredient & Purpose***

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***Active ingredient***

Sulfur 10% .....

***Purpose***

Acne treatment

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***Uses***

For the treatment of acne. Clears up acne:

- blemishes
- pimples
- blackheads
- whiteheads

***Warnings***

**For external use only. Avoid contact with the eyes.**

***Do not use on***

- broken skin
- large areas of the skin

***When using this product***

- skin irritation and dryness are more likely to occur if you use another topical acne medication at the same time. If irritation occurs, only use one topical acne medication at a time.
- apply only to areas with acne

***If pregnant or breast feeding,***

ask a health professional before use

***Keep out of the reach of children.***

If swallowed, get medical help or contact a Poison Control Center right away.

***Directions***

- cleanse the skin thoroughly before applying this product
- cover the entire affected area with a thin layer one to three times daily
- because excessive drying of the skin may occur, start with one application daily, then gradually increase to two or three times daily if needed or as directed by a doctor
- if bothersome dryness or peeling occurs, reduce application to once a day or every other day

**Other information**

- store between (59-86°F) 15-30°C
- don't use if clear seal over cap is broken, torn, or missing

**Inactive ingredients**

cetyostearyl alcohol, glycerin, glyceryl monostearate, methylparaben, mineral oil, paraffin, petrolatum, propylparaben, polyethylene glycol, sorbitan monooleate, water

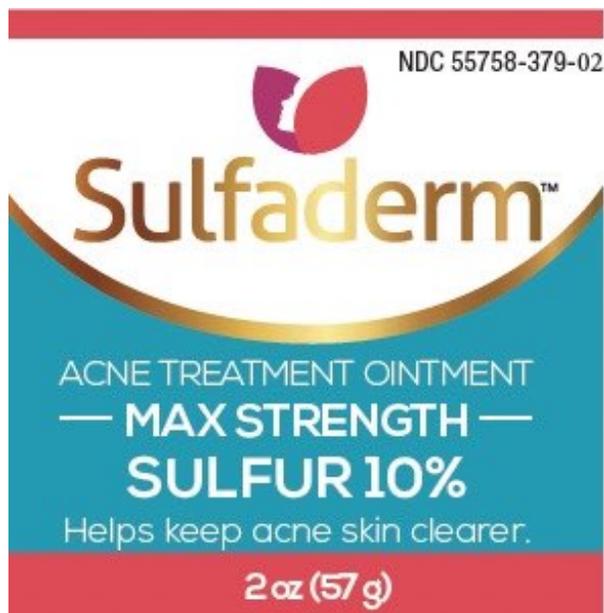
**Distributed by:**

**Pharmadel LLC**

New Castle, DE 19720

**+1-866-359-3478**

**Principal Display Panel**



**SULFADERM**

sulfur ointment

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:55758-379
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
<b>SULFUR</b> (UNII: 70FD1KFU70) (SULFUR - UNII:70FD1KFU70)	SULFUR	10 g in 100 g

**Inactive Ingredients**

<b>Ingredient Name</b>	<b>Strength</b>
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>CETOSTEARYL ALCOHOL</b> (UNII: 2DMT128M1S)	
<b>GLYCERYL MONOSTEARATE</b> (UNII: 230OU9XXE4)	
<b>METHYLPARABEN</b> (UNII: A2I8C7HI9T)	
<b>MINERAL OIL</b> (UNII: T5L8T28FGP)	
<b>PARAFFIN</b> (UNII: I9O0E3H2ZE)	
<b>PETROLATUM</b> (UNII: 4T6H12BN9U)	
<b>POLYETHYLENE GLYCOL, UNSPECIFIED</b> (UNII: 3WJQ0SDW1A)	
<b>PROPYLPARABEN</b> (UNII: Z8IX2SC1OH)	
<b>SORBITAN MONOOLEATE</b> (UNII: 06XEA2VD56)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	

**Product Characteristics**

<b>Color</b>	yellow	<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>		<b>Imprint Code</b>	
<b>Contains</b>			

**Packaging**

<b>#</b>	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
1	NDC:55758-379-02	57 g in 1 JAR; Type 0: Not a Combination Product	06/10/2023	04/01/2026

**Marketing Information**

<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
OTC Monograph Drug	M006	06/10/2023	04/01/2026

**Labeler** - PharmadeI LLC (030129680)

