

**YOUMI ALCOHOL PREP PAD 70 ISOPROPYL ALCOHOL- isopropyl alcohol liquid  
PT YOUMI MEDIKA INDUSTRI**

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**Youmi Alcohol Prep Pad 70% Isopropyl Alcohol**

**Drug Facts**

**Active ingredient**

Isopropyl Alcohol 70%

**Purpose**

Antiseptic Cleanser

**Use**

for preparation of the skin prior to an injection.

**Warnings**

For External Use Only.  
Flammable, keep away from fire or flame.

**Do not use**

- with electrocautery procedures.
- in the eyes. If contact occurs, flush eyes with water.

**Stop use and ask a doctor if**

irritation and redness develop.

**Keep out of reach of children.**

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

**Directions:**

Wipe injection site vigorously and discard.

**Other Information:**

Store at room temperature.

**Inactive Ingredient:**

Purified Water.

**Package Labeling:**

<div style="text-align: center;">  <p>NDC XXXXXXXXXX REF XXXXXXXXXX</p> <p><b>ALCOHOL PREP PAD</b></p> <p>For External Use Only</p> <p>Saturated with 70% Isopropyl Alcohol Sterility guaranteed unless package is damaged or opened</p> <p><b>Sterile R</b>    NOT MADE WITH natural rubber latex    Single Use Only</p> <p><b>1/pouch</b> Made in Indonesia</p> <p>MANUFACTURED BY PT YOU MI MEDIKA INDUSTRI Jalan Akasia, Kedawung, Kec. Banyuputih, Kab. Batang, Central Java, Indonesia, P.C. 51271</p> <p style="text-align: right;">Rev.A</p> </div>	<table border="1" style="width: 100%; border-collapse: collapse;"> <tr> <th colspan="2" style="text-align: left;">Drug Facts</th> </tr> <tr> <th style="text-align: left;">Active ingredient</th> <th style="text-align: left;">Purpose</th> </tr> <tr> <td>Isopropyl Alcohol 70%.</td> <td>Antiseptic Cleanser</td> </tr> <tr> <th colspan="2">Warnings</th> </tr> <tr> <td colspan="2">For External Use Only. Flammable, keep away from fire or flame.</td> </tr> <tr> <td colspan="2">Use for preparation of the skin prior to an injection.</td> </tr> <tr> <th colspan="2">Do not use</th> </tr> <tr> <td colspan="2"> <ul style="list-style-type: none"> <li>■ with electrocautery procedures.</li> <li>■ in the eyes. If contact occurs, flush eyes with water.</li> </ul> </td> </tr> <tr> <td colspan="2">Stop use and ask a doctor if irritation and redness develop.</td> </tr> <tr> <th colspan="2">Keep out of reach of children</th> </tr> <tr> <td colspan="2">If swallowed, get medical help or contact a Poison Control Center right away.</td> </tr> <tr> <td colspan="2">Direction: Wipe injection site vigorously and discard.</td> </tr> <tr> <td colspan="2">Other Information: Store at room temperature.</td> </tr> <tr> <td colspan="2">Inactive Ingredient: Purified Water.</td> </tr> <tr> <td style="text-align: center;">Lot: XXXXXX</td> <td style="text-align: center;">Exp: XXXX-XX</td> </tr> </table>	Drug Facts		Active ingredient	Purpose	Isopropyl Alcohol 70%.	Antiseptic Cleanser	Warnings		For External Use Only. Flammable, keep away from fire or flame.		Use for preparation of the skin prior to an injection.		Do not use		<ul style="list-style-type: none"> <li>■ with electrocautery procedures.</li> <li>■ in the eyes. If contact occurs, flush eyes with water.</li> </ul>		Stop use and ask a doctor if irritation and redness develop.		Keep out of reach of children		If swallowed, get medical help or contact a Poison Control Center right away.		Direction: Wipe injection site vigorously and discard.		Other Information: Store at room temperature.		Inactive Ingredient: Purified Water.		Lot: XXXXXX	Exp: XXXX-XX
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YOU MI ALCOHOL PREP PAD 70 ISOPROPYL ALCOHOL				
isopropyl alcohol liquid				
Product Information				
<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:87401-000	
<b>Route of Administration</b>	TOPICAL			
Active Ingredient/Active Moiety				
<b>Ingredient Name</b>		<b>Basis of Strength</b>	<b>Strength</b>	
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)		ISOPROPYL ALCOHOL	70 mL in 100 mL	
Inactive Ingredients				
<b>Ingredient Name</b>			<b>Strength</b>	
WATER (UNII: 059QF0KO0R)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:87401-000-	0.34 mL in 1 POUCH; Type 0: Not a Combination	02/05/2026	

00	Product	02/05/2026	
<b>Marketing Information</b>			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	02/05/2026	

**Labeler** - PT YOUMI MEDIKA INDUSTRI (780789890)

**Registrant** - PT YOUMI MEDIKA INDUSTRI (780789890)

**Establishment**

Name	Address	ID/FEI	Business Operations
PT YOUMI MEDIKA INDUSTRI		780789890	manufacture(87401-000)

Revised: 2/2026

PT YOUMI MEDIKA INDUSTRI