

**HYDROCORTISONE- hydrocortisone cream**  
**J&A Digital Inc.**

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**Hydrocortisone**

***Drug Facts***

***Active ingredient***

Hydrocortisone 1%

***Purpose***

Anti-itch

***Uses***

- for the temporary relief of itching associated with minor irritations and rashes
- other uses of this product should be only under the advice and supervision of a doctor.

***Warnings***

**For external use only.**

***Do not use***

for the treatment of diaper rash. Consult a doctor.

***Stop use and ask a doctor if***

- condition worsens or lasts more than 7 days, or clears up and occurs again within a few days.
- you begin use of any other hydrocortisone product
- bleeding occurs

**Keep out of reach of children.**

If swallowed contact a Poison Control Center right away.

***Directions***

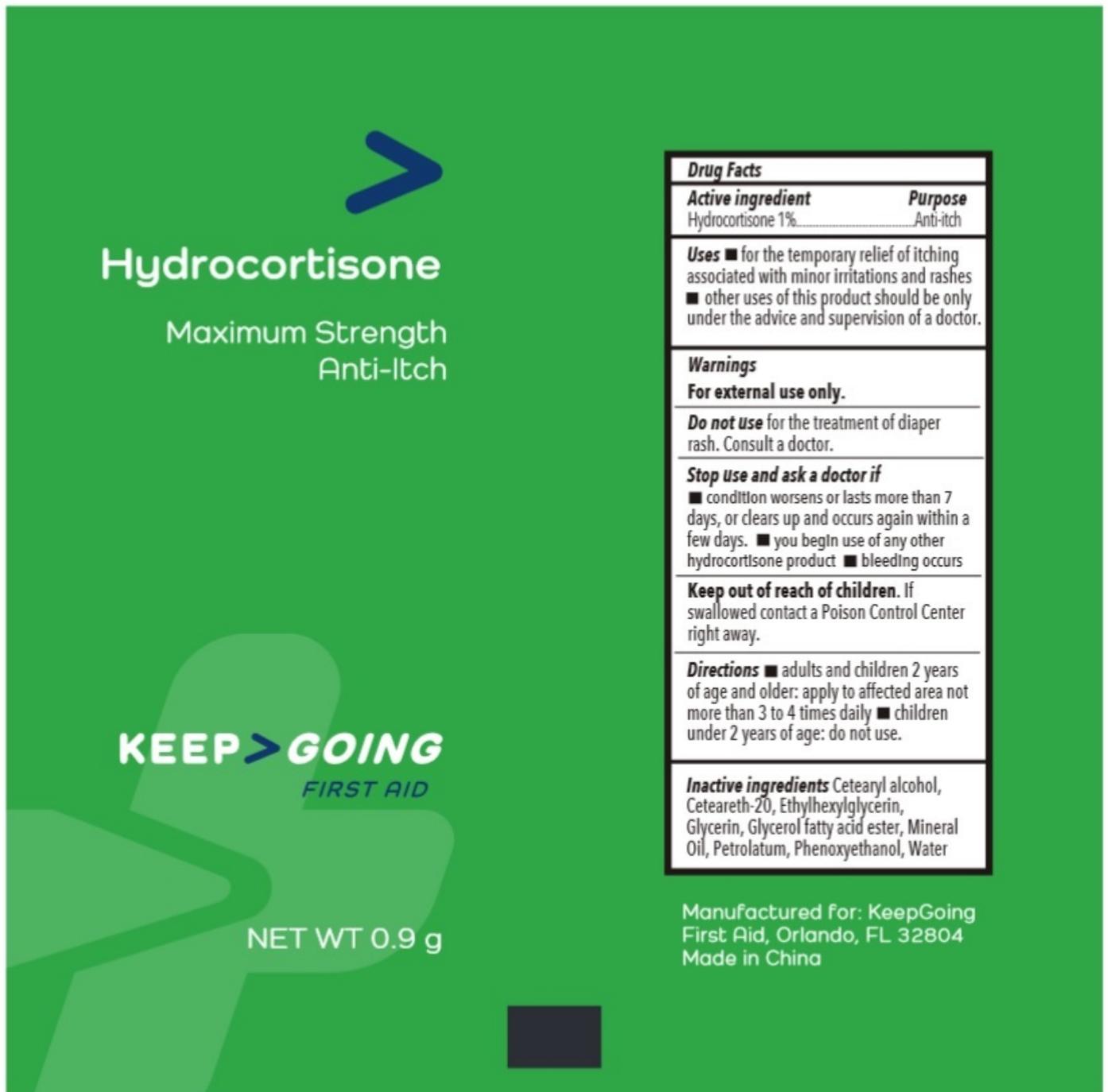
- adults and children 2 years of age and older: apply to affected area not more than 3 to 4 times daily
- children under 2 years of age: do not use.

***Inactive ingredients***

Cetearyl alcohol, Cetareth-20, Ethylhexylglycerin, Glycerin, Glycerol fatty acid ester,

Mineral Oil, Petrolatum, Phenoxyethanol, Water

### Package Labeling:



<b>HYDROCORTISONE</b>			
hydrocortisone cream			
<b>Product Information</b>			
<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:82942-1003
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
<b>HYDROCORTISONE</b> (UNII: W4X0X7BPJ) (HYDROCORTISONE - UNII:W4X0X7BPJ)	HYDROCORTISONE	10 mg in 1 g

**Inactive Ingredients**

<b>Ingredient Name</b>	<b>Strength</b>
<b>CETOSTEARYL ALCOHOL</b> (UNII: 2DMT128M1S)	
<b>POLYOXYL 20 CETOSTEARYL ETHER</b> (UNII: YRC528SWUY)	
<b>ETHYLHEXYLGLYCERIN</b> (UNII: 147D247K3P)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>MINERAL OIL</b> (UNII: T5L8T28FGP)	
<b>PETROLATUM</b> (UNII: 4T6H12BN9U)	
<b>PHENOXYETHANOL</b> (UNII: HIE492ZZ3T)	
<b>WATER</b> (UNII: 059QF0KO0R)	

**Packaging**

<b>#</b>	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
1	NDC:82942-1003-1	0.9 g in 1 PACKET; Type 0: Not a Combination Product	05/08/2023	

**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	M017	05/08/2023	

**Labeler** - J&A Digital Inc. (040268672)

Revised: 12/2024

J&amp;A Digital Inc.