

TOPICAL ANALGESIC & SKIN PROTECTANT · OTC DRUG

I.C.® IVY RELIEF

Roll-on stick · Diphenhydramine HCl 2% + Zinc Acetate 0.1% · 0.68 oz. (20 mL)

PRODUCT DESCRIPTION

I.C. Ivy Relief is a 0.68 oz. (20 mL) roll-on stick for outdoor crews who work around poison ivy, oak and sumac. It is an OTC drug with two actives: diphenhydramine HCl 2%, a topical analgesic, and zinc acetate 0.1%, a skin protectant. It temporarily relieves pain and itching associated with rashes due to poison ivy, poison oak and poison sumac, insect bites, minor burns, minor cuts, sunburn, minor skin irritations and scrapes, and dries the oozing and weeping of poison ivy, oak and sumac. Alcohol-free and steroid-free.

DIRECTIONS FOR USE (LABEL)

Do not use more than directed · Hold stick straight down over infected area · press tip of stick repeatedly on affect skin area until liquid flows, then dab sparingly · adults and children 2 years of age and older: apply to affected area not more then 3 to 4 times daily · children under 2 years of age: ask a doctor

USES (LABEL)

Temporary relieves pain and itching associated with:

- Insect Bites · Minor Burns · Minor Cuts
- Sunburn · Minor Skin Irritations · Scrapes
- Dries oozing and weeping of Poison Ivy, Poison Oak, and Poison Sumac
- Rashes due to Poison Ivy, Poison Oak, and Poison Sumac

APPLICATIONS

Outdoor Workers, Municipalities, Utilities, Power Gen, Telecommunications, Arborists, Construction, Military, First Responders.

FEATURES & BENEFITS

TOPICAL ANALGESIC	Diphenhydramine HCl 2% temporarily relieves pain and itching
SKIN PROTECTANT	Zinc acetate 0.1% dries the oozing and weeping of poison ivy, oak and sumac
POISON IVY RELIEF	Temporarily relieves pain and itching associated with rashes due to poison ivy, oak and sumac
BITES & MINOR IRRITATIONS	Also labeled for insect bites, minor burns, minor cuts, scrapes, sunburn and minor skin irritations
EASY APPLICATION	One-handed roll-on stick: press the tip until the liquid flows, then dab sparingly
ALCOHOL-FREE	Contains no alcohol, which can dry skin
STEROID-FREE	Contains no steroids
POCKET SIZE	A 20 mL stick for the shirt pocket, truck or first-aid kit
BLOCK, CLEANSE, RELIEF	The third step after I.C. Ivy Block and I.C. Ivy Cleanse



FREE OF

ALCOHOL-FREE	STEROID-FREE
PARABEN-FREE	NO PRESERVATIVES
NO BAKING SODA	NO ANIMAL TESTING

ACTIVE INGREDIENTS

- Diphenhydramine HCl 2% · Topical Analgesic
- Zinc Acetate 0.1% · Skin Protectant

INACTIVE INGREDIENTS

Propylene Glycol, DI Water, PVP K-30, Xanthan Gum

OTHER INFORMATION

Protect from excessive heat (40°C/104°F) (label). Questions or Comments? 1-888-860-7424

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DRUG FACTS WARNINGS (LABEL)

For external use only. Do not use on large areas of the body • with any other product containing diphenhydramine, even one taken by mouth. **Ask doctor before use** on chicken pox • on measles. **When using this product** avoid contact with eyes. **Stop use and ask doctor if** conditions worsen, or do not improve within 7 days • symptoms persist for more than 7 days or clear up and occur again within a few days. **Keep out of reach of children.** If swallowed, get medical help or contact Poison Control Center right away.

[CONFIRM: Drug Facts are copied verbatim from the current stick label (LN0166_A), including its wording slips: "Temporary relieves", "infected area", "affect skin", "not more than". Fix on the next label revision, then here. The label prints PN# ICR-STICK; master, web and GSA use ICVR-STICK.]

APPEARANCE & PHYSICAL DATA (2026 SDS)

PRODUCT TYPE	Clear liquid
ODOR	Characteristic
PH	Neutral
SPECIFIC GRAVITY	0.98–1.03 @ 25°C
BOILING / FREEZING POINT	100°C (212°F) / 26–30°F
SOLUBILITY IN WATER	Soluble
VISCOSITY	TBD – confirm

[CONFIRM: TBD – confirm: same formula as I.C. Itch & Pain Relief (F012), whose TDS prints slightly cloudy, odorless, pH 4.5–6.0 and viscosity 0–20 cPs. The SDS says odor "Characteristic" and pH "Neutral"; the web page says odorless. Use the ISIP values here?]

STORAGE TEMPERATURE & SHELF LIFE

Can be stored up to 2 years from date of manufacture (SDS). Store in a cool place in the original container and protect from direct sunlight; avoid temperatures >115°F (SDS). Protect from excessive heat (40°C/104°F) (label).

QUICK CHEMICAL REGULATORY REFERENCES

PROP 65	No warning required (2026 SDS)
TSCA	All ingredients listed or exempt
SARA TITLE III	No hazard category
REACH SVHC	None ≥ 0.1%
GHS / OSHA HCS	Not a dangerous substance; FD&C Act labeled, exempt from HCS labeling

SIZES & PART NUMBERS

PART NO.	SIZE	CASE	UPC
ICVR-STICK	Roll-on stick, 0.68 oz. (20 mL)	20	666080851217
ICVR-FOIL-100	1 mL single-use foil, box of 100	1	666080851194
ICVR-FOIL-200	1 mL single-use foil, box of 200	1	666080851200
ICVR-2	2 oz. (59 mL) bottle	–	666080851224

[CONFIRM: TBD – confirm: the foil boxes and the 2 oz are Active in the Part Number Master and have label art (foil ICR-FOIL, 1 mL; 2 oz ICR-2, LN143_A) but are on no channel (Woo, Amazon, GSA list the stick only). Still sold? Part nos. ICVR-FOIL-100/-200 and ICVR-2 are assumed; 2 oz case pack and foil NDC not on file. The 2 oz label (2021) also prints "Kills Germs" and stick directions – not used here.]

REGULATORY

This product is an OTC drug (external analgesic and skin protectant), manufactured and labeled in compliance with the Federal Food, Drug, and Cosmetic Act administered by the Food and Drug Administration (FDA). R&R Lotion is an FDA and EPA registered and audited facility.

FDA EST. ID	3004167955
EPA FACILITY	91651-AZ-001
NDC	59555-230-19 (stick, label) • -04 (2 oz, label)
GOV'T CONTRACT	GSA / VA / DOD 47QSMS24D0078

SAFETY DATA SHEETS

For safety, environmental, handling, first aid and disposal information, refer to the Safety Data Sheet (Ivy Relief Ointment, issued January 1, 2026) in the SDS library at rrlotion.com/msds/

QUALITY ASSURANCE

Facility standards. R&R Lotion facilities follow Current Good Manufacturing Practice (cGMP) and/or Cosmetic GMP requirements.

Material standards. Incoming raw materials must undergo quality inspection and are efficacy tested prior to being used for manufacturing.

Equipment standards. All equipment and tools utilized are cleaned and sanitized both before and after use once verified by QA using sterility testing equipment.

Formula testing. Formulas developed by R&R Lotion undergo a series of tests and approvals to verify stability, product efficacy and preservative effectiveness.

Finished product testing. Every batch produced by R&R Lotion must go through physical, chemical and microbiological testing prior to being approved for distribution. This includes in-house lab testing of appearance, color, odor, density, pH and viscosity, followed by third-party active ingredient efficacy and product sterility testing.

Product traceability. The collection and annual review of all raw materials, finished goods, process controls and formulas allows for consistent control and correction in good manufacturing and documentation procedures.

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This information and all further technical advice are based upon our present knowledge and experience. Customer is responsible for determining whether products and the information in this document are appropriate for Customer's use. The claims made may not have been approved for use in all countries. R&R Lotion Inc assumes no obligation or liability for the information in this document. No warranties are given; all implied warranties of merchantability or fitness for a particular purpose are expressly excluded. Dated: 10/2026 (supersedes no prior TDS; new sheet).