

**POSITIVE SKIN TEST CONTROL - HISTAMINE- histamine injection**  
**Jubilant HollisterStier LLC**

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**Histamine 366140-H05 Initial Upload to Intagras SPL Portal**

**POSITIVE SKIN TEST CONTROL - HISTAMINE**

For Percutaneous (Scratch, Prick or Puncture) Administration

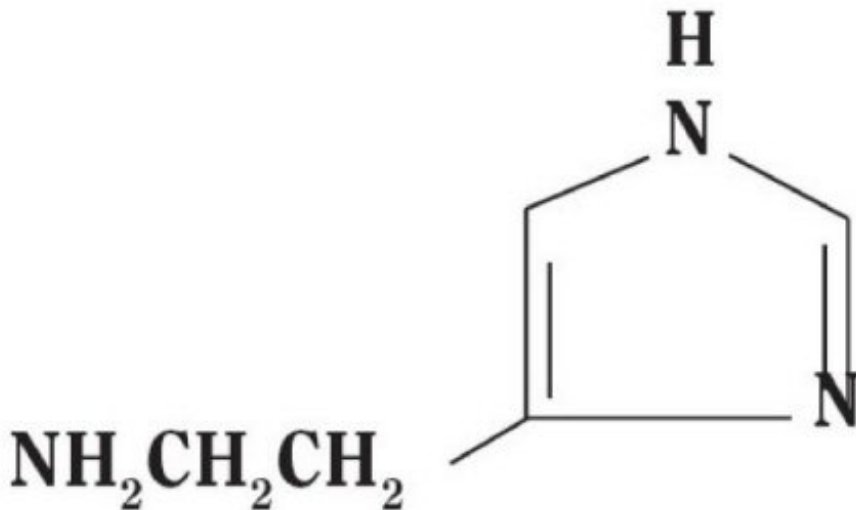
Histamine Base: 6mg/mL

(Histamine Dihydrochloride: 10mg/mL)

**This product is to be used by a physician or under the supervision of a physician.**

**DESCRIPTION**

Histamine Dihydrochloride contains histamine, a potent vasodilator having the chemical name 2-(4-Imidazolyl) ethylamine. Histamine has an empirical formula of  $C_5H_9N_3$ , a molecular weight of 111.15, and the following chemical structure:



Histamine Dihydrochloride is available in the following strengths:

**1. SCRATCH, PRICK or PUNCTURE TEST CONTROL:**

Positive Skin Test Control - Histamine contains 6.0 mg/mL Histamine Base and is a clear, colorless, sterile solution. It consists of Histamine Dihydrochloride 10mg/mL, Sodium Chloride 0.5%, Sodium Bicarbonate 0.275%, and Glycerin 50.0% (v/v) as a preservative.

**CLINICAL PHARMACOLOGY**

Pharmacological actions of histamine include the increase of capillary and post capillary venular permeability. This vascular change leads to the wheal-flare response. Large reactions can cause an amplification of a reaction to a nearby skin test. Histamine is

degraded either by oxidative deamination or by methylation and oxidative deamination so that the principal excretion products are imidazoleacetic acid-riboside and 1-methyl imidazoleacetic acid respectively.<sup>2</sup>

10 atopic subjects and 10 non-atopics were tested with Positive Skin Test Control - Histamine Solutions, and two negative control solutions. Thirteen females (35-49 years old; mean age 36.5) and 7 males (23-45 years old; mean age 36.4) were tested. This study included 19 Caucasian subjects and 1 African American subject. Different skin test devices produce different skin test responses. (14,15) Study results are summarized in Table 1 and Table 2. Atopic and non-atopic subject information was combined, as there were no significant differences in their skin responses.

Summation of wheal ( $\Sigma W$ ) or summation of erythema ( $\Sigma E$ ) was determined by measuring the longest diameter and the mid-point orthogonal diameter of the wheal or erythema response and summing the two measurements.

**Table 1: Positive Skin Test Control Solutions (n = 20)**

	Skin Test Technique (Skin Test Device)					
	Scratch <sup>†</sup> (Scarifier)		Puncture <sup>†</sup> (Prick Lancetter)		Puncture <sup>†</sup> (Bifurcated Vaccinating Needle)	
Histamine Base Concentration	6mg/mL		6mg/mL		6mg/mL	
Summation of Wheal/ Summation of Erythema	$\Sigma W$	$\Sigma E$	$\Sigma W$	$\Sigma E$	$\Sigma W$	$\Sigma E$
Skin Response (mm) <sup>§</sup>	20	76	12	49	13	56
Range	11-36	55-103	8-16	37-66	9-18	43-78
Standard Deviation	5.6	11.9	2.1	6.8	2.2	8.7

<sup>†</sup>Dose – 1 drop topically

<sup>§</sup>Negative control values were not subtracted from response measurements

**Table 2: Negative Skin Test Control Solutions (n = 20)**

	Skin Test Technique (Skin Test Device)					
	Scratch <sup>†</sup> (Scarifier)		Puncture <sup>†</sup> (Prick Lancetter)		Puncture <sup>†</sup> (Bifurcated Vaccinating Needle)	
Control Solution	Sterile 50% (v/v) Glycerin		Sterile 50% (v/v) Glycerin		Sterile 50% (v/v) Glycerin	
Summation of Wheal/ Summation of Erythema	ΣW	ΣE	ΣW	ΣE	ΣW	ΣE
Skin Response (mm)	8	7	3	4	5	5
Range	0-11	0-11	0-10	0-10	3-8	2-8
Standard Deviation	2.4	3.0	2.8	2.2	1.3	1.5

<sup>†</sup>Dose – 1 drop topically

Malling tested 25 subjects using a straight needle prick technique with Histamine Dihydrochloride 10mg/mL and found wheal single diameters of 5.4-8.5mm (mean 7 mm). 13

Histamine Dihydrochloride at 10mg/mL has been shown to elicit fewer false negative and false positive reactions than Histamine Dihydrochloride at 1mg/mL. 12 10mg/mL Histamine Dihydrochloride has been shown to be the appropriate strength to perform biological standardization using skin reactivity to histamine as a standard of comparison.11,13

## INDICATIONS AND USAGE

Positive Skin Test Control - Histamine is indicated as an adjunct in allergy skin test for diagnosis, as a positive control to test wheal-flare response of skin for evaluation of skin test response to allergenic extracts.

## CONTRAINDICATIONS

Positive Skin Test Control - Histamine is contraindicated in patients with a history of hypersensitivity to histamine products, and in patients with hypotension, severe hypertension, vasomotor instability, severe cardiac, pulmonary or renal disease.

## WARNINGS

Attacks of severe asthma or other serious allergic conditions may be precipitated by the administration of Histamine Dihydrochloride in patients with bronchial disease. Caution is advised in using histamine in such patients and in those with a history of bronchial asthma. Histamine Dihydrochloride has not been approved for unlabeled use as gastric acid stimulus or for detecting bronchial hyperactivity.

## PRECAUTIONS

**(1) General:** It is necessary that scarifiers, syringes, and needles be properly sterilized before use on each patient to prevent the possibility of accidental transfer of serum hepatitis and other infectious agents from one person to another. Disposable products may also be used.

Always have injectable epinephrine and a tourniquet available when any skin tests are being made. (See ADVERSE REACTIONS Section)

Patients should be observed in the office for 30 minutes after administration of the test and instructed to return to the office promptly if symptoms of an allergic reaction or shock occur.

**(2) Drug Interactions:** Certain medications may lessen the skin test wheal and erythema responses elicited by histamine for varying time periods. Conventional antihistamines should be discontinued at least 5 days before skin testing. Long acting antihistamines should be discontinued for at least 3 weeks prior to skin testing.<sup>7</sup> Topical steroids should be discontinued at the skin test site for at least 2-3 weeks before skin testing.<sup>7,8</sup>

Tricyclic antidepressants such as Doxepin should be withheld for at least 7 days before skin testing.<sup>9</sup> The physician must determine whether the risk of severe depression occurring in patients who discontinue their medication outweighs the benefits that could be obtained from skin testing. Topical local anesthetics may suppress the flare responses and should be avoided in skin test sites.<sup>10</sup>

**(3) Carcinogenesis, Mutagenesis, Impairment of Fertility:** Studies have not been performed on Positive Skin Test Control-Histamine.

**(4) Pregnancy:** Positive Skin Test Control-Histamine. Animal reproduction studies have not been conducted on Histamine Dihydrochloride. It is also not known whether Histamine Dihydrochloride can cause fetal harm when administered to a pregnant woman or can affect reproduction capacity. Histamine Dihydrochloride should be given to a pregnant woman only if clearly needed.

**(5) Nursing Mothers:** It is unknown whether Histamine Dihydrochloride affects lactation or is excreted in human milk. Because many drugs are excreted in human milk, caution should be exercised when Positive Skin Test Control-Histamine is administered to a nursing woman.

**(6) Pediatric Use:** Indications and dosage for the pediatric population are the same as for adults; however, the reactions to histamine are smaller. Studies have shown that the use of histamine solutions for skin test is safe in infants and young children.<sup>4,5,6</sup>

## ADVERSE REACTIONS

Large doses of histamine may precipitate systemic reactions. These reactions may include flushing, dizziness, headache, bronchial constriction, urticaria, asthma, marked hypotension or hypertension, abdominal cramps, vomiting, metallic taste, local or

generalized allergic manifestations.

An antihistamine preparation may be given orally, I.M. or I.V. to prevent or ameliorate systemic reactions to the drug.

**If a systemic or anaphylactic reaction does occur, apply a tourniquet above the site of injection and inject 1:1000 epinephrine-hydrochloride intramuscularly or subcutaneously into the opposite arm. Loosen the tourniquet at least every 10 minutes. Do not obstruct arterial blood flow with the tourniquet.**

Epinephrine Dosage:

ADULT: 0.3 to 0.5 mL should be injected. Repeat in 5 to 10 minutes if necessary.

PEDIATRIC: The usual initial dose is 0.01 mg (mL) per kg body weight or 0.3 mg(mL) per square meter of body surface area. Suggested dosage for infants to 2 years of age is 0.05 mL to 0.1 mL; for children 2 to 6 years, 0.15 mL; and children 6 to 12 years, 0.2 mL. Single pediatric doses should not exceed 0.3 mg (mL). Doses may be repeated as frequently as every 20 minutes, depending on the severity of the condition and the response of the patient.

After administration of epinephrine, profound shock or vasomotor collapse should be treated with intravenous fluids, and possibly vasoactive drugs. Oxygen should be given by mask. Aminophylline or adrenal corticosteroids may be used if necessary after adequate epinephrine and circulatory support has been given.

Emergency resuscitation measures and personnel trained in their use should be available immediately in the event of a serious systemic or anaphylactic reaction not responsive to the above measures (Ref. J. ALLERGY AND CLINICAL IMMUNOLOGY 77s (2): p. 271-273, 1986).<sup>16</sup>

Rarely are all of the above measures necessary, the tourniquet and epinephrine usually producing prompt responses. However, the physician should be prepared in advance for all contingencies. Promptness in beginning emergency treatment measures is of utmost importance.

### **Adverse Event Reporting**

Report all adverse events to Jubilant HollisterStier LLC Customer Technical Services Department at 1 (800) 992-1120. A voluntary adverse event reporting system for health professionals is available through the FDA MEDWATCH program. Preprinted forms (FDA Form 3500) are available from the FDA by calling 1 (800) FDA-1088. Completed forms should be mailed to MEDWATCH, 5600 Fisher Lane, Rockville, MD 20852-9787 or Fax to: 1 (800) FDA-0178.

### **OVERDOSAGE**

Overdosage may cause severe symptoms, including circulatory collapse, shock, and even death. Intravenous administration of histamine in normal volunteers at doses of up to 1.0 µg/kg/min produced flushing, headaches, tachycardia and decreased diastolic blood pressure. 1,2,3

See ADVERSE REACTIONS Section for emergency treatment steps.

### **DOSAGE AND ADMINISTRATION**

## (1) General

Parenteral drug products should be inspected visually for particulate matter and discoloration prior to administration, whenever solution and container permit. Prior to testing, clean the skin area with ether or alcohol and allow to dry. The back or the volar surface of the arms are the most satisfactory sites for testing.

Skin of the posterior thighs or abdomen may be used if necessary. Avoid very hairy areas where possible, since the reactions will be smaller and more difficult to interpret there. The most satisfactory areas of the back are from the posterior axillary fold to 2.5 cm from the spinal column, and from the top of the scapula to the lower rib margins. The best areas of the arms are the volar surfaces from the axilla to 2.5 or 5 cm above the wrist, skipping the anti-cubital space.

The histamine test should be applied in the same test area as other allergenic extracts tests, but spaced no closer than 4 to 5 cm from adjacent test sites. Use the same technique or procedure that you use for allergen testing.

The negative control is the diluent used in the extract to be tested (e.g. 50% glycerin, Sterile Albumin Saline with Phenol, Sterile Buffered Saline with Phenol).

With each skin testing method, in order for the reaction to Positive Skin Test Control-Histamine (6 mg/mL Histamine Base) to be considered valid, erythema must be present which exceeds the respective negative control by 4 mm ( $\Sigma E$ ). A wheal reaction does not have to be elicited unless there is a wheal reaction to the respective negative control. In this case, the wheal of the positive control must exceed the negative control by 4 mm ( $\Sigma W$ ) in order to be considered appropriate. Record measurement of erythema and wheal diameters.

Tables 1 and 2 summarize skin testing results with histamine base and controls in atopic and non-atopic subjects using four different devices and methods.

(2) Scratch, Prick or Puncture Test: Positive Skin Test Control-Histamine, 6.0 mg/mL Histamine Base.

The Scratch, Prick or Puncture test should be read in 15 minutes. If a large wheal reaction occurs before that time, wipe excess histamine solution from test site.

a) Use a sterile scarifier for each patient.

Hold the scarifier between the thumb and index finger, press the sharp edge of the instrument against the skin and twirl instrument rapidly. The scratch should disrupt the outer layers of epidermis down to the germinal layer, but should not produce immediate oozing of blood. The amount of pressure needed to produce a satisfactory scratch will vary between patients according to the thickness or fragility of their skin.

Apply one drop of Positive Skin Test Control-Histamine to the scratch test site.

b) Prick or Puncture Test: Prick tests are performed by placing a drop of extract on the skin and piercing through the drop into the skin with a slight lifting motion. Puncture tests are performed by placing a drop of extract on the skin and piercing through the drop perpendicular to the skin with a device such as a Prick Lancetter. After about 1 minute the extract may be wiped away with a dry sponge.

## HOW SUPPLIED

SCRATCH, PRICK or PUNCTURE TEST: Positive Skin Test Control-Histamine (6.0 mg/mL Histamine Base), 5 mL sterile amber vial with dropper assembly.

## STORAGE

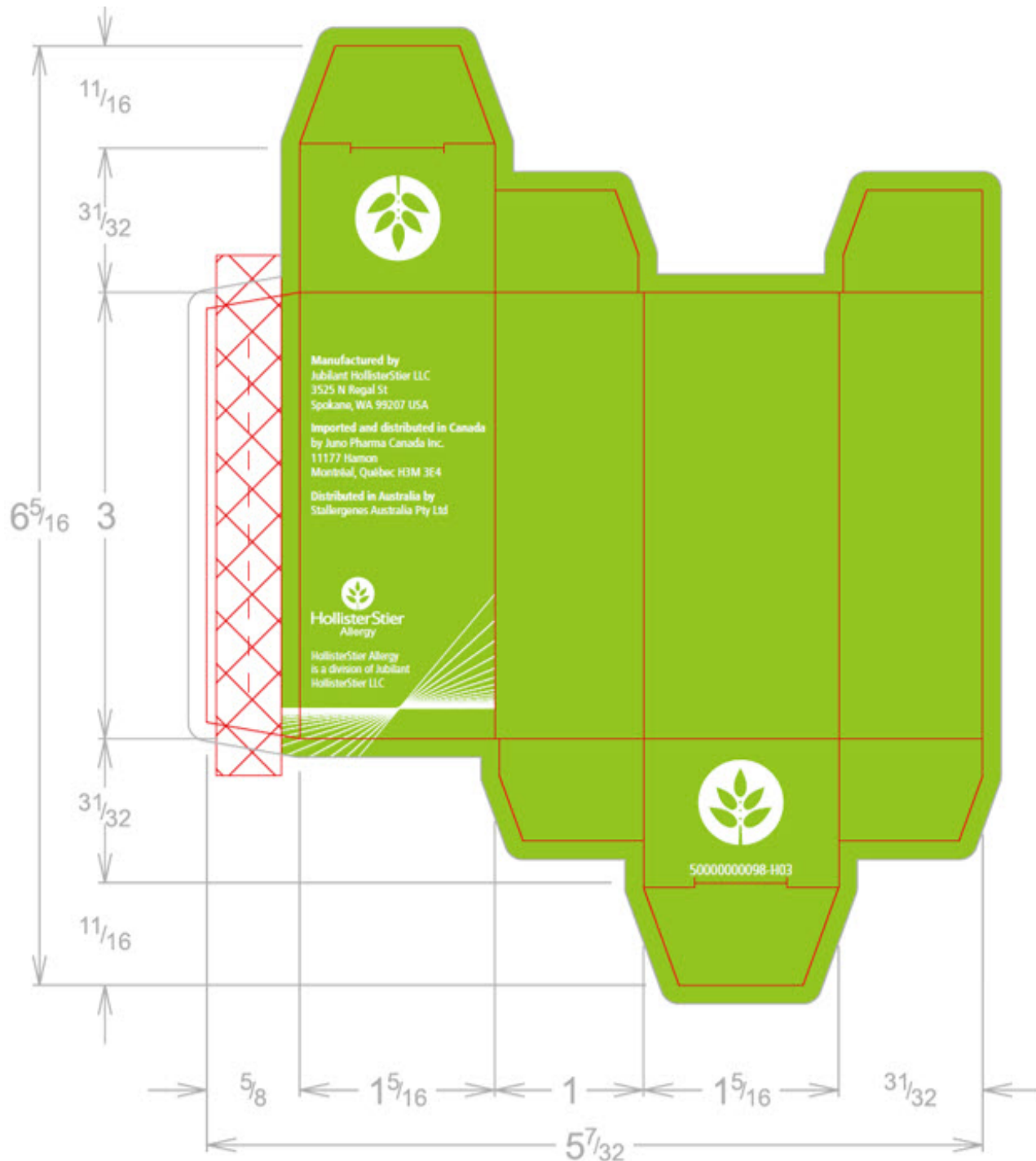
Store at 2°- 8°C. Protect from light when not in use.

## REFERENCES

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2. Samter, Max, K. Frank Austin, ed.. Histamine and other mediators of allergic reactions. Immunological Diseases. Little, Brown and Company, Boston, pp. 332-335, 1971.
3. Summers, Richard, Robert Sigler, James H. Schelhamer, and Michael Kaliner. Effects of infused histamine on asthmatic and normal subjects: comparison of skin test responses. J. Allergy Clin. Immunol. 67 (6): 456-464, June 1981.
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9. Rao, Kamineni S., et al. Duration of suppressive effect of tricyclic anti-depressants on histamine induced wheal and flare reactions on human skin. J. Allergy Clinical Immunology. 82: 752-757, November 1988.
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## PRINCIPAL DISPLAY PANEL



**5mL Positive Skin Test Control-Histamine, Carton**

**POSITIVE SKIN TEST  
CONTROL - HISTAMINE**

Preservative:  
50% Glycerin v/v

Inactive Ingredients:  
0.5% Sodium  
chloride  
0.275% Sodium  
bicarbonate

*Strength: 6mg/mL  
Histamine Base (10 mg/mL  
Histamine Dihydrochloride)*



(01)00365044999810  
(17)260102  
(10)XXXXXXXXXX  
(21)012345678910

Item: 7099ED  
Lot: XXXXXXXX  
Exp: 2026Jan02

Dose/Route: 1 drop topically  
Protect from light

DIN: 02439514  
NDC: 65044-9998-1  
U.S. License No. 1272

5 mL                      Item: 7099ED

Store at 2-8°C

**50000004970 - H01**

Non-Returnable

Rx Only - Sterile Until Opened

**7099ED - 5mL 10 mg/mL Positive Skin Test Control-Histamine, Carton Label**

**POSITIVE SKIN TEST  
CONTROL - HISTAMINE**

Strength: 6mg/mL Histamine Base  
(10 mg/mL Histamine Dihydrochloride)



50000004523-H02

5 mL vial  
Dose / Route: 1 drop topically  
Protect from light

Rx Only - Sterile  
Until Opened  
Store at 2-8°C  
Item: 7099ED  
Lot: S1234567  
Exp:2025Jan13

U.S. License No. 1272

Jubilant HollisterStier LLC Spokane, WA 99207

**7099ED - 5mL 10 mg/mL Positive Skin Test Control-Histamine, Container Label**

**POSITIVE SKIN TEST CONTROL - HISTAMINE**

histamine injection

**Product Information**

<b>Product Type</b>	HUMAN PRESCRIPTION DRUG	<b>Item Code (Source)</b>	NDC:65044-9998
<b>Route of Administration</b>	PERCUTANEOUS		

**Active Ingredient/Active Moiety**

<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
<b>HISTAMINE</b> (UNII: 820484N8I3) (HISTAMINE - UNII:820484N8I3)	HISTAMINE	6 mg in 1 mL

**Inactive Ingredients**

<b>Ingredient Name</b>	<b>Strength</b>
<b>SODIUM CHLORIDE</b> (UNII: 451W47IQ8X)	
<b>SODIUM BICARBONATE</b> (UNII: 8MDF5V39QO)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	

**Packaging**

<b>#</b>	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
1	NDC:65044-9998-1	5 mL in 1 VIAL; Type 0: Not a Combination Product	06/15/1995	

**Marketing Information**

<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
BLA	BLA103891	06/15/1995	

**Labeler** - Jubilant HollisterStier LLC (069263643)**Registrant** - Jubilant HollisterStier LLC - HollisterStier Allergy (119241112)

Revised: 4/2026

Jubilant HollisterStier LLC