

CANARY FEATHERS- serinus canaria solution
CATTLE EPITHELIA- bos taurus solution
CHICKEN FEATHERS- gallus gallus solution
DOG EPITHELIA- canis lupus familiaris solution
DUCK FEATHERS- anas platyrhynchos solution
GUINEA PIG EPITHELIA- cavia porcellus solution
GOOSE FEATHERS- anser anser solution
GERBIL EPITHELIA- meriones unguiculatus solution
RAT EPITHELIA- rattus norvegicus solution
MOUSE EPITHELIA- mus musculus solution
HORSE EPITHELIA- equus caballus solution
MIXED FEATHERS- gallus gallus, anas platyrhynchos and anser anser solution
GOAT EPITHELIA- capra hircus solution
RABBIT EPITHELIA- oryctolagus cuniculus solution
SILK WORM- bombyx mori solution
HAMSTER EPITHELIA- mesocricetus auratus solution
HOG EPITHELIA- sus scrofa solution
PARAKEET FEATHERS- melopsitticus undulatus solution
Greer Laboratories, Inc.

Non Standardized Allergenic Extracts
Pollens, Molds, Epidermals, Insects, Dusts, Foods, and Miscellaneous
Inhalants

HIGHLIGHTS OF PRESCRIBING INFORMATION

These highlights do not include all the information needed to use Non-Standardized Allergenic Extracts (Pollens, Molds, Epidermals, Insects, Foods and Miscellaneous Inhalants) safely and effectively. See full prescribing information for Non-Standardized Allergenic Extracts.

Non-Standardized Allergenic Extracts (Pollens, Molds, Epidermals, Insects, Foods, and Miscellaneous Inhalants)

Solutions for percutaneous, intradermal or subcutaneous administration.

Initial U.S. Approval: 1968

WARNING: SEVERE ALLERGIC REACTIONS

See full prescribing information for complete boxed warning.

- **Non-Standardized Allergenic Extracts can cause severe life-threatening systemic reactions, including anaphylaxis. (5.1)**
- **Do not administer these products to patients with severe, unstable or uncontrolled asthma. (4)**
- **Observe patients in the office for at least 30 minutes following treatment. Emergency measures and personnel trained in their use must be available immediately in the event of a life-threatening reaction. (5.1)**
- **Patients with extreme sensitivity to these products, on an accelerated immunotherapy build-up, switching to another lot, receiving high doses of**

these products, and patients exposed to similar allergens may be at increased risk of a severe allergic reaction. (5.1)

- **These products may not be suitable for patients with certain underlying medical conditions that may reduce their ability to survive a systemic allergic reaction, and for patients receiving medications such as beta-blockers that may make them unresponsive to epinephrine or inhaled bronchodilators. (5.1, 5.2)**
-

RECENT MAJOR CHANGES

Warning and Precautions, Anaphylaxis Following False Negative Food Allergen Skin Test Results (5.3) 01/2023

INDICATIONS AND USAGE

Non-Standardized Allergenic Extracts are indicated for:

- Skin test diagnosis of patients with a clinical history of allergies to one or more of the specific allergens. (1)
- Immunotherapy for reduction of allergen-induced allergic symptoms confirmed by appropriate positive skin tests or *in vitro* testing for allergen-specific IgE antibodies. (1)

Food extracts have not been proven safe or effective in allergen immunotherapy.

DOSAGE AND ADMINISTRATION

For percutaneous, intradermal or subcutaneous use only.

The extracts are diluted with sterile diluents when used for intradermal testing or subcutaneous immunotherapy. For percutaneous testing, the extracts are diluted with sterile diluents in patients expected to be at greater risk for systemic allergic reaction. Dosages vary by mode of administration and by individual response. See full prescribing information for instructions on preparation, administration, and adjustments of dose. (2.1)

DOSAGE FORMS AND STRENGTHS

Non-Standardized Allergenic Extracts are labeled in weight/volume and/or protein nitrogen units (PNU)/milliliter (a measure of total protein), and are supplied as sterile aqueous stock concentrates at up to 1:10 weight/volume or 40,000 PNU/milliliter, or 50% glycerin stock concentrates at up to 1:20 weight/volume. (3)

CONTRAINDICATIONS

- Severe, unstable or uncontrolled asthma. (4)
- History of any severe systemic or local allergic reaction to an allergen extract. (4)

WARNINGS AND PRECAUTIONS

Severe allergic reactions have occurred following the administration of other allergenic extracts and may occur in individuals following the administration of Non-Standardized Allergenic Extracts in the following situations:

- Extreme sensitivity to Non-Standardized Allergenic Extracts, receipt of high doses of Non-Standardized Allergenic Extracts, or concomitant exposure to similar

environmental allergens. (5.1)

- Receiving an accelerated immunotherapy build-up schedule (e.g., “rush” immunotherapy), or changing from one allergenic lot to another. (5.1)

ADVERSE REACTIONS

The most common adverse reactions, occurring in 26 to 82% of all patients who receive subcutaneous immunotherapy, are local adverse reactions at the injection site (e.g., erythema, itching, swelling, tenderness, pain). (6)

Systemic adverse reactions, occurring in $\leq 7\%$ of patients who receive subcutaneous immunotherapy, include generalized skin erythema, urticaria, pruritus, angioedema, rhinitis, wheezing, laryngeal edema, and hypotension. These can be fatal. (6)

To report SUSPECTED ADVERSE REACTIONS, contact GREER Laboratories, Inc. at 1-855-274-1322 or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch.

DRUG INTERACTIONS

- Antihistamines and other medications that suppress histamine, including topical corticosteroids, topical anesthetics and tricyclic antidepressants can interfere with skin test results. (7)

See 17 for PATIENT COUNSELING INFORMATION

FULL PRESCRIBING INFORMATION: CONTENTS*

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FULL PRESCRIBING INFORMATION

WARNING: SEVERE ALLERGIC REACTIONS

- **Non-Standardized Allergenic Extracts can cause severe life-threatening systemic reactions, including anaphylaxis. (5.1)**
- **Do not administer these products to patients with severe, unstable or uncontrolled asthma. (4)**
- **Observe patients in the office for at least 30 minutes following treatment. Emergency measures and personnel trained in their use must be available immediately in the event of a life-threatening reaction. (5.1)**
- **Patients with extreme sensitivity to these products, those on an accelerated immunotherapy build-up schedule, those switching to another allergenic lot, those receiving high doses of Non-Standardized Allergenic Extracts, or those also exposed to similar allergens may be at increased risk of a severe allergic reaction. (5.1)**
- **These products may not be suitable for patients with certain underlying medical conditions that may reduce their ability to survive a serious allergic reaction. (5.1)**
- **These products may not be suitable for patients who may be unresponsive to epinephrine or inhaled bronchodilators, such as those taking beta-blockers. (5.2)**

1 INDICATIONS AND USAGE

Non-Standardized Allergenic Extracts are indicated for:

- Skin test diagnosis of patients with a clinical history of allergies to one or more of the specific non-standardized allergens.
- Immunotherapy for the reduction of allergen-induced allergic symptoms confirmed by appropriate positive skin tests or by *in vitro* testing for allergen-specific IgE antibodies.

Food extracts have not been proven safe or effective in allergen immunotherapy.

2 DOSAGE AND ADMINISTRATION

For percutaneous, intradermal or subcutaneous use only.

The extracts are diluted with sterile diluents when used for intradermal testing or subcutaneous immunotherapy. For percutaneous testing, the extracts are diluted with sterile diluents in patients expected to be at greater risk for systemic allergic reaction. Dosages vary by mode of administration and by individual response.

2.1 Preparation for Administration

Parenteral drug products should be inspected visually for particulate matter and discoloration prior to administration, whenever solution and container permit. Discard solution if either of these conditions exist.

The extracts are diluted with sterile diluents when used for percutaneous and intradermal testing, or for subcutaneous immunotherapy.

Extracts labeled “For Diagnostic Use Only” are intended for percutaneous and intradermal testing only. These extracts have not been shown by adequate data to be safe and effective for therapeutic use. The extracts labeled For Diagnostic Use Only are the foods Barley, Coffee, Oat, Pineapple, Rye, Spinach, Wheat, the insects Flea, House Fly, Mosquito, and the plant and plant parts Cottonseed and Flax.

Undiluted 50% glycerin stock concentrate is used for percutaneous testing. To prepare 10-fold dilutions for percutaneous testing in patients suspected to be at greater risk for systemic allergic reaction, start with the stock concentrate. Proceed as shown in Table 1. The 10-fold dilution series uses 0.5 milliliters of concentrate added to 4.5 milliliters of sterile 50% glycerin diluent. Subsequent dilutions are made in a similar manner.

To prepare 10-fold dilutions for intradermal testing and immunotherapy, start with a 1:10 weight/volume, 1:20 weight/volume, or up to a 40,000 PNU/milliliter stock concentrate. Proceed as shown in Table 1. The 10-fold dilution series uses 0.5 milliliter of concentrate added to 4.5 milliliters of sterile diluent (glycerin-free diluents for intradermal testing, glycerin-free or 10% glycerin-saline diluents for immunotherapy). Subsequent dilutions are made in a similar manner.

Table 1: 10-fold Dilution Series*

Dilution	Extract	Milliliters of Diluent	Dilution Strength (w/v)	Dilution Strength (w/v)	Dilution Strength (PNU/milliliter)
0	Concentrate		1:10	1:20	20,000
1	0.5 mL Concentrate	4.5	1:100	1:200	2,000
2	0.5 mL Dilution 1	4.5	1:1,000	1:2,000	200
3	0.5 mL Dilution 2	4.5	1:10,000	1:20,000	20
4	0.5 mL Dilution 3	4.5	1:100,000	1:200,000	2
5	0.5 mL Dilution 4	4.5	1:1,000,000	1:2,000,000	0.2
6	0.5 mL Dilution 5	4.5	1:10,000,000	1:20,000,000	0.02

*There is no direct potency correlation across the table between PNU/milliliter and w/v.

Undiluted 50% glycerin stock concentrate is used for percutaneous testing. To prepare 5-fold dilutions for percutaneous testing in patients suspected to be at greater risk for systemic allergic reaction, start with the stock concentrate. Proceed as shown in Table 2. The 5-fold dilution series uses 1 milliliter of concentrate added to 4 milliliters of sterile 50% glycerin diluent. Subsequent dilutions are made in a similar manner.

To prepare 5-fold dilutions for intradermal testing or immunotherapy, start with a 1:10 weight/volume, 1:20 weight/volume, or up to a 40,000 PNU/milliliter stock concentrate. Proceed as shown in Table 2. The 5-fold dilution series uses 1 milliliter of concentrate added to 4 milliliters of sterile diluent (glycerin-free diluents for intradermal testing, glycerin-free or 10% glycerin-saline diluents for immunotherapy). Subsequent dilutions are made in a similar manner.

Table 2: 5-fold Dilution Series*

Dilution	Extract	Milliliters of Diluent	Dilution of Strength (w/v)	Dilution of Strength (w/v)	Dilution of Strength (PNU/milliliter)
0	Concentrate		1:10	1:20	20,000
1	1 mL Concentrate	4	1:50	1:100	4,000
2	1 mL Dilution 1	4	1:250	1:500	800
3	1 mL Dilution 2	4	1:1,250	1:2,500	160
4	1 mL Dilution 3	4	1:6,250	1:12,500	32
5	1 mL Dilution 4	4	1:31,250	1:62,500	6.4
6	1 mL Dilution 5	4	1:156,250	1:312,500	1.28

*There is no direct potency correlation across the table between PNU/milliliter and w/v.

2.2 Diagnostic Testing

Diagnostic testing can be performed via percutaneous or intradermal administration of the Non-Standardized Allergenic Extracts. A positive skin test reaction should be interpreted in relation to the patient's history and known exposure to the specific allergen(s).

Percutaneous Skin Testing

Preparation and Dose

For percutaneous testing (prick or puncture), use glycerinated extract; use the extracts at the highest available stock concentration. In patients suspected to be at greater risk for systemic allergic reaction, use 10-fold or 5-fold dilutions of the concentrate.

Prick test: Place one drop of extract with appropriate controls on the skin and with a skin test device, pierce through the drop into the skin with a slight lifting motion. Alternatively, use skin test devices loaded with the extract from the storage trays in a

similar manner or in accordance with the device manufacturer's recommendations.

Puncture test: Place one drop of extract or control on the skin and pierce the skin through the drop with a skin test device perpendicular to the skin. Alternatively, use skin test devices loaded with the extract from the storage trays in a similar manner or in accordance with the device manufacturer's recommendations.

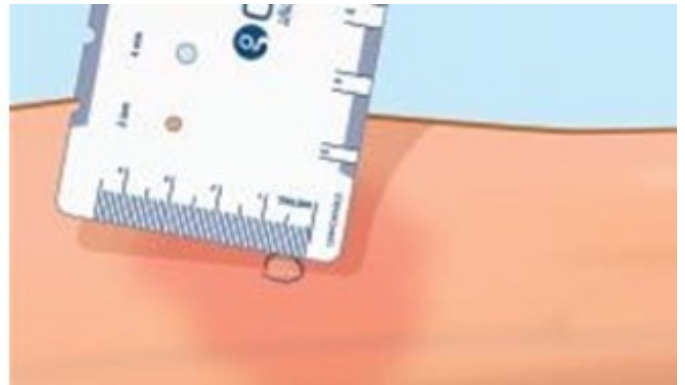
Interpreting Results

When using percutaneous skin test devices, follow the directions provided with the test devices. A glycerinated histamine control solution (6 milligrams/milliliter or 1 milligram/milliliter histamine base) may be used as the positive control. A 50% glycerin-saline solution may be used as the negative control.

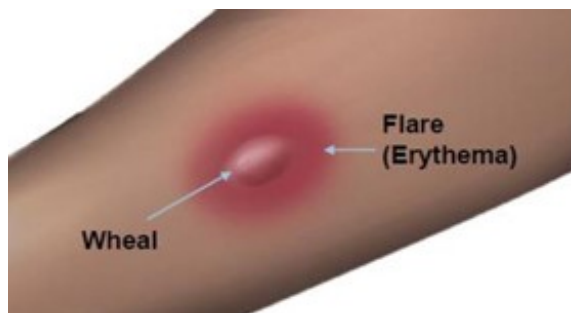
Read and record skin test responses 15 to 20 minutes after exposure. Individual patient reactivity can vary with time, allergen potency, and/or immunotherapy, as well as testing technique. The most reliable method of recording a skin test reaction is to measure the largest diameter of both wheal and erythema. While some correlation exists between the size of the skin test reaction and the degree of sensitivity, other factors should be considered in the diagnosis of allergy to specific allergens (see Figure 1 below).

Figure 1: Measurement of Wheal and Flare

Use a paper or plastic millimeter skin reaction guide as shown below.



Fifteen minutes after application of the skin test, measure the length and midpoint orthogonal width of each flare and wheal from the inner edge of the reaction.

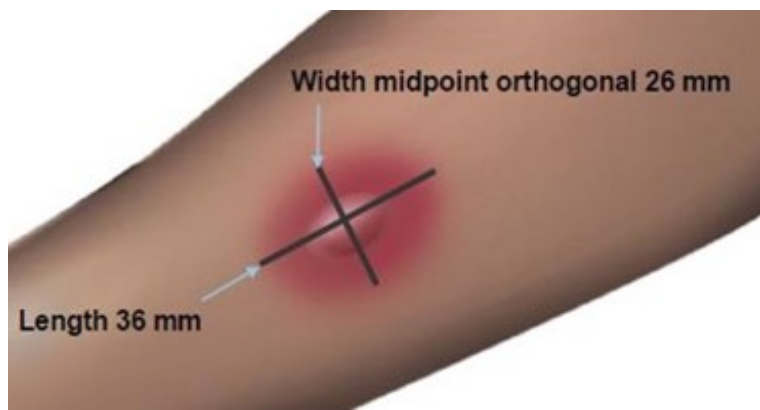


The wheal is a smooth, slightly elevated area which is redder or paler than the surrounding skin.

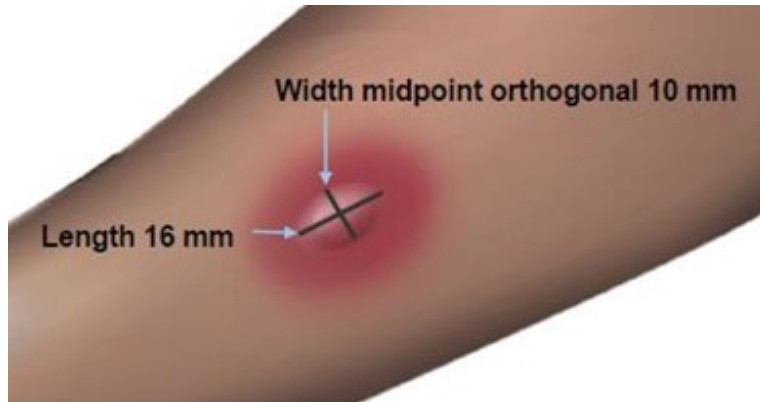
The flare is the red outermost zone of a wheal reaction.

The length of the skin test is defined as the largest diameter and the width of the skin test is defined as the diameter perpendicular to the length at its midpoint. Consider the wheal and flare as separate entities. First, measure the flare and then independently measure the wheal.

Measuring the Flare



Measuring the Wheal



The average diameter measurement in the example above of the flare is $(26 \text{ mm} + 36 \text{ mm})/2 = 31 \text{ mm}$ and the average diameter of the wheal is $(10 \text{ mm} + 16 \text{ mm})/2 = 13 \text{ mm}$.

Responses to positive controls should be at least 3 millimeters larger than responses to the negative controls.

Negative controls should elicit no reaction or only reactions of small diameter (less than 2 millimeters wheal, less than 5 millimeters erythema).

If either the positive or negative control response does not meet the above criteria, results for the allergenic extracts tested at the same time should be considered invalid and be repeated.

Intradermal Skin Testing

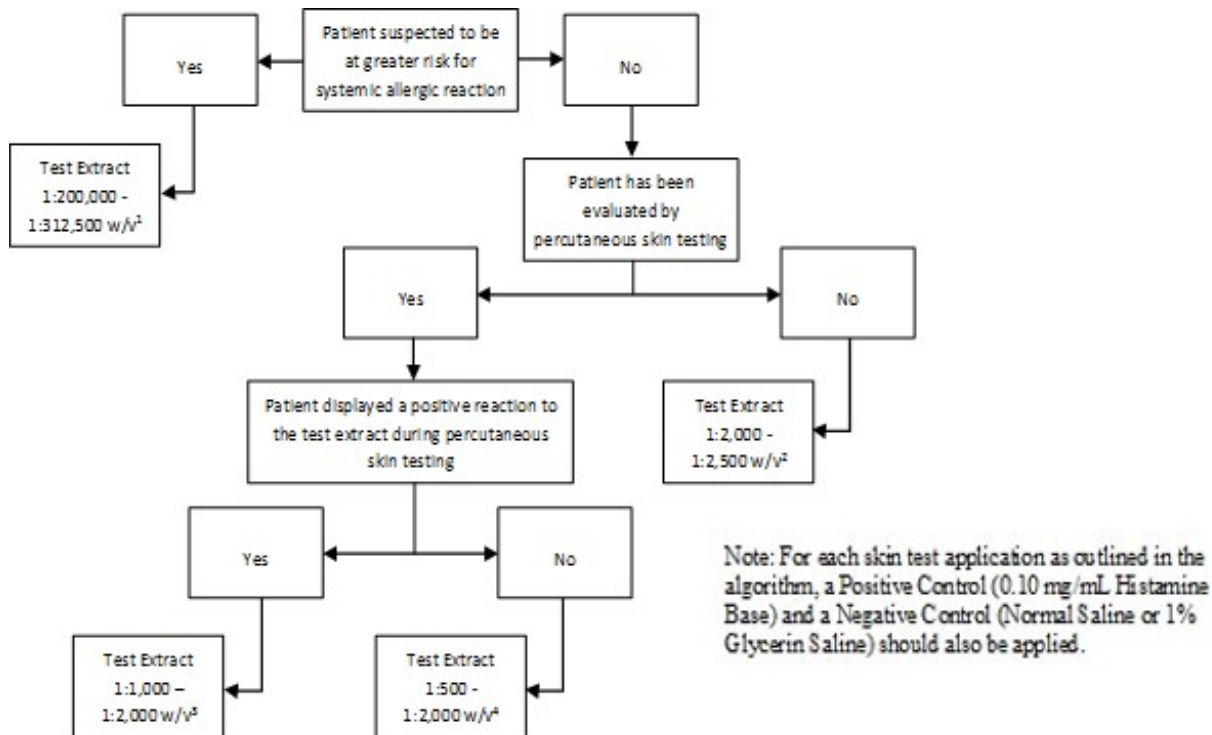
Preparation and Dose

For intradermal testing, dilute stock concentrate to 1:100 to 1:1000 volume to volume of Non-Standardized Allergenic Extracts stock concentrate solution. Dilute the stock concentrate solution with sterile diluent [see *Dosage and Administration* (2.1)]. Use normal or buffered saline or normal saline with human serum albumin (HSA) diluent. If the result from the initial test dose is negative, subsequent intradermal tests using increasingly stronger doses may be performed up to the maximum recommended strength of 1:25 volume to volume dilution of the extract concentrate solution.

Inject 0.02 milliliters of the extract solution intradermally according to the algorithms shown in Figure 2.

Figure 2: Algorithm for Dilution of Stock Concentrate Solution of Non-

Standardized Allergenic Extracts for Intradermal Skin Testing



¹ Corresponds to 1:10,000 - 1:15,625 volume to volume dilution of 1:20 weight/volume glycerinated extract concentrates

² Corresponds to 1:100 - 1:125 volume to volume dilution of 1:20 weight/volume glycerinated extract concentrates

³ Corresponds to 1:50 - 1:100 volume to volume dilution of 1:20 weight/volume glycerinated extract concentrates

⁴ Corresponds to 1:25 - 1:100 volume to volume dilution of 1:20 weight/volume glycerinated extract concentrates

2.3 Immunotherapy

For subcutaneous administration only.

Preparation and Dose

Stock concentrates of Non-Standardized Allergenic Extracts are available in aqueous (up to 1:10 weight/volume or 40,000 PNU/milliliter) and 50% glycerin (up to 1:20 weight/volume) strengths for immunotherapy. Stock concentrates are diluted in normal saline, buffered saline, HSA-saline, or 10% glycerin-saline, depending on the patient's reactivity to the diluent. See Table 1 and Table 2 for dilution preparation.

Administration of Immunotherapy

Administer immunotherapy by subcutaneous injection in the lateral aspect of the upper arm or thigh. Avoid injection directly into any blood vessel.

The optimal interval between doses of allergenic extract varies among individuals. Injections are usually given 1 to 2 times per week until the maintenance dose is reached, at which time the injection interval is increased to 2, then 3, and finally 4 weeks. Dosages vary by mode of administration, and by clinical response and tolerance. The minimum course of treatment may be three to five years, depending on the clinical response.

Guidelines for Immunotherapy

The initial dose of the extract should be based on the skin test reactivity. In patients suspected to be at greater risk for systemic allergic reaction by history and skin test, the initial dose of the extract should be 0.05 milliliter of a 1:20,000,000 to 1:2,000,000 weight/volume extract dilution. Patients not suspected to be at greater risk for systemic allergic reaction may be started at 0.1 milliliter of a 1:200,000 to 1:20,000 weight/volume extract dilution.

The dose of allergenic extract is increased at each injection by no more than 50% of the previous dose, and the next increment is governed by the response to the last injection.

Select the maximum tolerated maintenance dose based on the patient's clinical response and tolerance. Doses larger than 0.2 milliliter of the stock concentrate are rarely administered because an extract in 50% glycerin diluent can cause discomfort upon injection.

Dosage Modification Guidelines for Immunotherapy

The following conditions may indicate a need to withhold or reduce the dosage of immunotherapy.

- Symptoms of rhinitis and/or asthma
- Infection accompanied by fever
- Exposure to excessive amounts of clinically relevant environmental allergen prior to a scheduled injection
- Large local reactions that persist for longer than 24 hours can be an indication for repeating the previous dose or reducing the dose at the next administration

Any evidence of a systemic reaction is an indication for a significant reduction (at least 75%) in the subsequent dose. Repeated systemic adverse reactions are sufficient reason for the cessation of further attempts to increase the dose.

Local adverse reactions require a decrease in the next dose by at least 50%. Proceed cautiously in subsequent dosing. In situations prompting dose reduction, once the reduced dose is tolerated, a cautious increase in dosage can be attempted.

Changing extract to a different lot or from a different manufacturer: When switching to a different lot of extract, or from another manufacturer's extract, decrease the starting dose. Because manufacturing processes and sources of raw materials differ among manufacturers, the interchangeability of extracts from different manufacturers cannot be assured. In general, a dose reduction of 50 to 75% of the previous dose should be adequate, but each situation must be evaluated separately, considering the patient's history of sensitivity, tolerance of previous injections, and other factors. Dose intervals should not exceed one week when rebuilding the dose.

Unscheduled gaps between treatments: Patients can lose tolerance to allergen injections during prolonged periods between doses, which increases their risk for an adverse reaction. The duration of tolerance between injections varies from patient to patient.

During the build-up phase, when patients receive injections 1 to 2 times per week, repeat or reduce the extract dosage if there has been a substantial time interval between injections. This depends on: 1) the concentration of allergen immunotherapy extract that

is to be administered; 2) a previous history of systemic reactions; and 3) the degree of variation from the prescribed

interval of time, with longer intervals since the last injection leading to greater reductions in the dose to be administered.

This suggested approach to dose modification, due to unscheduled gaps between treatments during the build-up phase, is not based on published evidence. The individual physician should use this or a similar protocol for the specific clinical setting.

Similarly, if unscheduled gaps occur during maintenance therapy, it may be necessary to reduce the dosage and bring the patient up to maintenance dosing using an established build-up protocol.

Changing from non-stabilized to human serum albumin (HSA) stabilized diluents: Allergenic extracts prepared with diluents containing HSA and 0.4% phenol are more stable than those prepared with diluents that do not contain stabilizers. When switching from a non-stabilized to an HSA-stabilized diluent, consider lowering the dose for immunotherapy.

3 DOSAGE FORMS AND STRENGTHS

Non-Standardized Allergenic Extracts are labeled in weight/volume and/or protein nitrogen units (PNU)/milliliter (a measure of total protein), and are supplied as sterile aqueous stock concentrates at up to 1:10 weight/volume or 40,000 PNU/milliliter, or 50% glycerin stock concentrates at up to 1:20 weight/volume.

4 CONTRAINDICATIONS

Non-Standardized Allergenic Extracts are contraindicated in patients with:

- Severe, unstable or uncontrolled asthma
- History of any severe systemic or local allergic reaction to an allergen extract

5 WARNINGS AND PRECAUTIONS

5.1 Serious Systemic Adverse Reactions

Serious systemic adverse reactions have occurred following the administration of other allergenic extracts and may occur in individuals following the administration of Non-Standardized Allergenic Extracts in the following situations:

- Extreme sensitivity to the specific allergen(s)
- Receipt of an accelerated immunotherapy build-up schedule (e.g., “rush” immunotherapy)
- Receipt of high doses of allergenic extracts or concomitant exposure to similar environmental allergens
- Change from one allergenic extract lot to another allergenic extract lot

High-risk patients have had fatal reactions. Consider using more dilute preparations in patients suspected to be at greater risk of systemic allergic reaction [see *Dosage and Administration* (2.1)].

Administer Non-Standardized Allergenic Extracts in a healthcare setting under the supervision of a physician prepared to manage a severe systemic or a severe local allergic reaction. Observe patients in the office for at least 30 minutes following administration. ¹

5.2 Epinephrine

Non-Standardized Allergenic Extracts may not be suitable for patients with certain medical conditions that may reduce the ability to survive a serious allergic reaction or increase the risk of adverse reactions after epinephrine administration. Examples of these medical conditions include but are not limited to: markedly compromised lung function (either chronic or acute), unstable angina, recent myocardial infarction, significant arrhythmia, and uncontrolled hypertension.

These products may not be suitable for patients who are taking medications that can potentiate or inhibit the effect of epinephrine. These medications include:

Beta-adrenergic blockers: Patients taking beta-adrenergic blockers may be unresponsive to the usual doses of epinephrine used to treat serious systemic reactions, including anaphylaxis. Specifically, beta-adrenergic blockers antagonize the cardiostimulating and bronchodilating effects of epinephrine.

Alpha-adrenergic blockers, ergot alkaloids: Patients taking alpha-adrenergic blockers may be unresponsive to the usual doses of epinephrine used to treat serious systemic reactions, including anaphylaxis. Specifically, alpha-adrenergic blockers antagonize the vasoconstricting and hypertensive effects of epinephrine. Similarly, ergot alkaloids may reverse the pressor effects of epinephrine.

Tricyclic antidepressants, levothyroxine sodium, monoamine oxidase inhibitors, and certain antihistamines: The adverse effects of epinephrine may be potentiated in patients taking tricyclic antidepressants, levothyroxine sodium, monoamine oxidase inhibitors, and the antihistamines chlorpheniramine, and diphenhydramine.

Cardiac glycosides, diuretics: Patients who receive epinephrine while taking cardiac glycosides or diuretics should be observed carefully for the development of cardiac arrhythmias.

5.3 Anaphylaxis Following False Negative Food Allergen Skin Test Results

False negative skin test results associated with anaphylaxis from subsequent exposure to the allergen have been reported during postmarketing diagnostic use of some food allergenic extracts. Based on the patient's clinical history and index of suspicion, healthcare providers should consider confirming negative skin testing with serologic testing by measuring specific serum IgE or with a medically-supervised oral food challenge.

5.4 Cross-Reactions and Dose Sensitivity

When determining the final dose of an allergen mixture for immunotherapy, consider cross-reactivity among component extracts.

Determine the initial dilution of allergenic extract, starting dose, and progression of dosage based on the patient's history and results of skin tests ² [see *Dosage and Administration* (2.1)]. Strongly positive skin tests can be indicators for potential adverse reactions.

6 ADVERSE REACTIONS

The most common adverse reactions, occurring in 26 to 82% of all patients who receive subcutaneous immunotherapy are local adverse reactions at the injection site (e.g., erythema, itching, swelling, tenderness, pain).¹ Systemic adverse reactions, occurring in $\leq 7\%$ of patients who receive subcutaneous immunotherapy,³ include generalized skin erythema, urticaria, pruritus, angioedema, rhinitis, wheezing, laryngeal edema, and hypotension. These adverse reactions can be fatal.¹

The allergenic extracts labeled “For Diagnostic Use Only” that contain sodium formaldehyde sulfoxylate (SFS) can cause slight discoloration of the skin at the site of administration. This discoloration can remain for extended amounts of time.

7 DRUG INTERACTIONS

7.1 Antihistamines

Do not perform skin testing with allergenic extracts within 3 to 10 days of use of first-generation H₁-histamine receptor blockers (e.g., clemastine, diphenhydramine) and second-generation antihistamines (e.g., loratadine, cetirizine). These products suppress histamine skin test reactions and could mask a positive response.²

7.2 Topical Corticosteroids and Topical Anesthetics

Topical corticosteroids can suppress skin reactivity; therefore, discontinue use at the skin test site for 2 to 3 weeks before skin testing. Avoid use of topical local anesthetics at skin test sites as they can suppress flare responses.²

7.3 Tricyclic Antidepressants

Tricyclic antidepressants can have potent antihistamine effects that can affect skin testing. If tricyclic medication has been recently discontinued, allow 7 to 14 days before initiating skin testing.²

8 USE IN SPECIFIC POPULATIONS

8.1 Pregnancy

Risk Summary

All pregnancies have a risk of birth defect, loss, or other adverse outcomes. In the U.S. general population, the estimated background risk of major birth defects and miscarriage in clinically recognized pregnancies is 2 to 4% and 15 to 20%, respectively. There are no human or animal data to establish the presence or absence of Non-Standardized Allergenic Extracts-associated risks during pregnancy.

8.2 Lactation

Risk Summary

It is not known whether Non-Standardized Allergenic Extracts are present in human milk. Data are not available to assess the effects of these extracts on the breastfed child or on milk production/excretion. The developmental and health benefits of breastfeeding should be considered along with the mother’s clinical need for Non-Standardized

Allergenic Extracts and any potential adverse effects on the breastfed child from the extracts or from the underlying maternal condition.

8.4 Pediatric Use

For use of these products in children younger than 5 years of age, consideration should be given to the patient's ability to comply and cooperate with allergen immunotherapy and the potential for difficulty in communicating with the child regarding systemic reactions. ¹

8.5 Geriatric Use

Data are not available to determine if subjects 65 years of age and older respond differently to allergen immunotherapy than younger subjects.

11 DESCRIPTION

Non-Standardized Allergenic Extracts are sterile solutions used for percutaneous testing, intradermal testing, or subcutaneous immunotherapy. Aqueous extracts contain the soluble extractants of the source material in water for injection, 0.5% sodium chloride, 0.54% sodium bicarbonate, and 0.4% phenol. Glycerinated extracts contain the soluble extractants of the source material in water for injection and 50% glycerin, 0.25% sodium chloride, 0.27% sodium bicarbonate, and 0.2% phenol. The pH of the extracts range from 6 to 9.

Certain food extracts (Barley, Oat, Pineapple, Rye, Spinach, and Wheat), labeled "For Diagnostic Use Only", contain 0.1% sodium formaldehyde sulfoxylate as an antioxidant.

Source materials used in the manufacture of allergenic extracts are collected from natural sources or from laboratory cultures.

Non-Standardized Allergenic Extracts appear as clear and colorless to dark brown solutions that should be free of particulate matter.

Extracts are labeled either as weight-to-volume based on the weight of the source material to the volume of the extracting fluid, or as PNU/milliliter with one PNU representing 0.00001 mg of protein nitrogen per milliliter.

12 CLINICAL PHARMACOLOGY

12.1 Mechanism of Action

The skin test reaction results from interaction of the introduced allergen and allergen-specific IgE antibodies bound to mast cells, leading to mast cell degranulation and release of histamine, tryptase and other mediators, which results in the formation of the wheal and flare.

The precise mechanisms of action of allergen immunotherapy are not known. Immunologic responses to immunotherapy include changes in allergen-specific IgE levels, allergen-specific IgG levels, and regulatory T cell responses. ¹

14 CLINICAL STUDIES

Specific immunotherapy with allergenic extracts is helpful in reducing symptoms

associated with exposure to the offending allergens. A summary of effectiveness by the Panel on Review of Allergenic Extracts, an advisory committee to the U.S. Food and Drug Administration, has been published. ⁴

15 REFERENCES

1. Cox LJ, Nelson H, Lockey R. Allergen immunotherapy: A practice parameter third update. *J Allergy Clin Immunol.* 2011;127:(1)S1-55.
2. Bernstein IL, Li JT, Bernstein DI, et al. Allergy diagnostic testing: an updated practice parameter. *Ann Allergy Asthma Immunol.* 2008;100:S1-148.
3. Greenberg MA, Kaufman CR, Gonzalez GE, et al. Late and immediate systemic-allergic reactions to inhalant allergen immunotherapy. *J Allergy Clin Immunol.* 1986;77:865-870.
4. Federal Register Proposed Rule: Biological Products: Implementation of Efficacy Review, Allergenic Extracts, Federal Register 1985;50: 3082-3288.

16 HOW SUPPLIED/STORAGE AND HANDLING

Non-Standardized Allergenic Extracts and mixes may be supplied as aqueous stock concentrates of up to 1:10 weight/volume or 40,000 PNU/milliliter for intradermal and subcutaneous testing. The extracts may also be supplied as 50% glycerin stock concentrates of up to 1:20 weight/volume for use in percutaneous skin testing and subcutaneous immunotherapy. Non-Standardized Allergenic Extracts are labeled in weight/volume and/or PNU/milliliter and may be provided in 5, 10, and 50 milliliter vials. Glycerinated extracts are also supplied in 5 milliliter dropper vials for prick or puncture testing.

Non-Standardized Allergenic Extracts available are as follows:

Pollens - Grasses

Bahia Grass, *Paspalum notatum*

Brome, Smooth, *Bromus inermis*

Canarygrass, Reed, *Phalaris arundinacea*

Johnson Grass, *Sorghum halepense*

Quack (Couch) Grass, *Elymus repens*

Ryegrass, Giant Wild, *Leymus condensatus*

Ryegrass, Italian, *Lolium multiflorum*

Velvetgrass, *Holcus lanatus*

Wheatgrass, Western, *Pascopyrum smithii*

Pollens - Trees

Acacia, *Acacia dealbata*

Alder, Hazel, *Alnus serrulata*

Alder, Red, *Alnus rubra*

Alder, White, *Alnus rhombifolia*

Ash, Arizona (Velvet), *Fraxinus velutina*

Ash, Green, *Fraxinus pennsylvanica*

Ash Mix (Equal parts *Fraxinus pennsylvanica*, *Fraxinus americana*)

Ash, Oregon, *Fraxinus latifolia*

Ash, White, *Fraxinus americana*
Aspen, *Populus tremuloides*
Beech, American, *Fagus grandifolia*
Birch, Black-Sweet, *Betula lenta*
Birch, Mix (Equal parts *Betula lenta*, *Betula nigra*, *Betula populifolia*)
Birch, River, *Betula nigra*
Birch, Spring, *Betula occidentalis*
Birch, White, *Betula populifolia*
Box Elder, *Acer negundo*
Cedar, Mountain, *Juniperus ashei*
Cedar, Red, *Juniperus virginiana*
Cedar, Salt (Tamarisk), *Tamarix gallica*
Central/Eastern 4 Tree Mix (Equal parts *Ulmus americana*, *Acer negundo*, *Carya illinoensis*, *Quercus virginiana*)
Cottonwood, Arizona (Fremont), *Populus fremontii*
Cottonwood, Black, *Populus trichocarpa*
Cottonwood, Eastern, *Populus deltoides*
Cottonwood, Western, *Populus deltoides* ssp. *monilifera*
Cypress, Arizona, *Callitropsis arizonica*
Cypress, Bald, *Taxodium distichum*
Eastern Oak Mix (Equal parts *Quercus velutina*, *Quercus rubra*, *Quercus alba*)
Eastern 6 Tree Mix (Equal parts *Fagus grandifolia*, *Populus deltoides*, *Quercus rubra*, *Betula nigra*, *Carya ovata*, *Fraxinus americana*)
Eastern 7 Tree Mix (Equal parts *Ulmus americana*, *Fagus grandifolia*, *Populus deltoides*, *Quercus rubra*, *Betula nigra*, *Carya ovata*, *Fraxinus americana*)
Eastern 8 Tree Mix (Equal parts *Ulmus americana*, *Fagus grandifolia*, *Populus deltoides*, *Quercus rubra*, *Betula nigra*, *Carya ovata*, *Fraxinus americana*, *Acer saccharum*)
Eastern 10 Tree Mix (Equal parts *Platanus occidentalis*, *Ulmus americana*, *Fagus grandifolia*, *Populus deltoides*, *Quercus rubra*, *Betula nigra*, *Carya ovata*, *Fraxinus americana*, *Acer saccharum*, *Liquidambar styraciflua*)
Elm, American, *Ulmus americana*
Elm, Cedar, *Ulmus crassifolia*
Elm Mix (Equal parts *Ulmus americana*, *Ulmus pumila*)
Elm, Siberian, *Ulmus pumila*
Eucalyptus, Bluegum, *Eucalyptus globulus*
Hackberry, *Celtis occidentalis*
Hazelnut, American, *Corylus americana*
Hickory Mix (Equal parts *Carya glabra*, *Carya ovata*, *Carya laciniosa*, *Carya tomentosa*)
Hickory-Pecan Mix (Equal parts *Carya illinoensis*, *Carya ovata*)
Hickory, Shagbark, *Carya ovata*
Hickory, Shellbark, *Carya laciniosa*
Hickory, White, *Carya tomentosa*
Juniper Mix (Equal parts *Juniperus monosperma*, *Juniperus scopulorum*)
Juniper, Oneseed, *Juniperus monosperma*
Juniper, Pinchot, *Juniperus pinchotii*
Juniper, Rocky Mountain, *Juniperus scopulorum*

Juniper, Utah, *Juniperus osteosperma*
Juniper, Western, *Juniperus occidentalis*
Locust Blossom, Black, *Robinia pseudoacacia*
Mango Blossom, *Mangifera indica*
Maple-Box Elder Mix (Equal parts *Acer saccharum*, *Acer negundo*)
2 Maple Mix (Equal parts *Acer rubrum*, *Acer saccharum*)
3 Maple Mix (Equal parts *Acer rubrum*, *Acer saccharinum*, *Acer saccharum*)
Maple, Red, *Acer rubrum*
Maple, Silver/Soft, *Acer saccharinum*
Maple, Sugar/Hard, *Acer saccharum*
Melaleuca, *Melaleuca quinquenervia*
Mesquite, Velvet, *Prosopis velutina*
Mulberry, Paper, *Broussonetia papyrifera*
Mulberry, Red, *Morus rubra*
Mulberry, White, *Morus alba*
Oak, Arizona (Gambel), *Quercus gambelii*
Oak, Black, *Quercus velutina*
Oak, Bur, *Quercus macrocarpa*
Oak, California Black, *Quercus kelloggii*
Oak, California Live, *Quercus agrifolia*
Oak, California White, *Quercus lobata*
Oak, Post, *Quercus stellata*
Oak, Red, *Quercus rubra*
Oak, Virginia Live, *Quercus virginiana*
Oak, Water, *Quercus nigra*
Oak, Western White, *Quercus garryana*
Oak, White, *Quercus alba*
Olive, *Olea europaea*
Olive, Russian, *Elaeagnus angustifolia*
Orange Pollen, *Citrus X sinensis*
Palm, Queen, *Syagrus romanzoffiana*
Pecan, *Carya illinoensis*
Peppertree Mix (Equal parts *Schinus molle*, *Schinus terebinthifolius*)
Pine, Australian (Beefwood), *Casuarina equisetifolia*
Pine, Loblolly, *Pinus taeda*
Pine, Longleaf, *Pinus palustris*
Pine Mix (Equal parts *Pinus taeda*, *Pinus strobus*, *Pinus echinata*)
Pine, Ponderosa, *Pinus ponderosa*
Pine, Virginia Scrub, *Pinus virginiana*
Pine, White (Eastern), *Pinus strobus*
Pine, White (Western), *Pinus monticola*
Pine, Yellow, *Pinus echinata*
Poplar, Lombardy's, *Populus nigra*
Poplar, White, *Populus alba*
Privet, *Ligustrum vulgare*

Sweetgum, *Liquidambar styraciflua*
Sycamore, American, *Platanus occidentalis*
Sycamore, California (Western), *Platanus racemosa*
11 Tree Mix (Equal parts *Fagus grandifolia*, *Platanus occidentalis*, *Ulmus americana*,
Juglans nigra, *Salix nigra*, *Populus deltoides*, *Quercus rubra*, *Betula nigra*, *Carya ovata*,
Acer saccharum, *Fraxinus americana*)
Walnut, Black, *Juglans nigra*
Walnut, California Black, *Juglans californica*
Walnut, English, *Juglans regia*
Wax Myrtle, *Morella cerifera*
Western Oak Mix (Equal parts *Quercus kelloggii*, *Quercus agrifolia*, *Quercus garryana*)
Western 3 Tree Mix (Equal parts *Olea europaea*, *Ulmus pumila*, *Platanus racemosa*)
Western 10 Tree Mix (Equal parts *Acacia dealbata*, *Acer negundo*, *Populus fremontii*,
Olea europaea, *Ulmus pumila*, *Betula occidentalis*, *Juniperus occidentalis*, *Platanus*
racemosa, *Quercus garryana*, *Morus alba*)
Western Walnut Mix (Equal parts *Juglans californica*, *Juglans regia*)
Willow, Arroyo, *Salix lasiolepis*
Willow, Black, *Salix nigra*

Pollens - Weeds and Garden Plants

Allscale, *Atriplex polycarpa*
Amaranth, Green, *Amaranthus hybridus*
Baccharis Mix (Equal parts *Baccharis sarothroides*, *Baccharis halimifolia*)
Burningbush (Kochia), *Kochia scoparia* spp. *scoparia*
Burrobrush, *Ambrosia salsola*
Central/Western Weed Mix (Equal parts *Kochia scoparia* ssp. *scoparia*, *Chenopodium*
album, *Salsola kali*)
Cocklebur, *Xanthium strumarium*
Common Weed Mix (Equal parts *Xanthium strumarium*, *Plantago lanceolata*,
Chenopodium album, *Amaranthus retroflexus*, *Salsola kali*)
Dock-Sorrel Mix (Equal parts *Rumex acetosella*, *Rumex crispus*)
Dock, Yellow (Curly), *Rumex crispus*
Dogfennel, *Eupatorium capillifolium*
Goldenrod, *Solidago canadensis*
Iodinebush, *Allenrolfea occidentalis*
Lamb's Quarters, *Chenopodium album*
Lenscale (Quailbrush), *Atriplex lentiformis*
Marsh Elder, True (Rough), *Iva annua*
Marshelder, Burweed (Giant Poverty), *Cyclachaena xanthiifolia*
Mixed Amaranths (Equal parts *Amaranthus hybridus*, *Amaranthus palmeri*, *Amaranthus*
retroflexus)
Mugwort, Common, *Artemisia vulgaris*
National Weed Mix (Equal parts *Xanthium strumarium*, *Ambrosia trifida*, *Chenopodium*
album, *Amaranthus retroflexus*, *Ambrosia artemisiifolia*)
Nettle, *Urtica dioica*
Palmer's Amaranth, *Amaranthus palmeri*
Pigweed, Rough Redroot, *Amaranthus retroflexus*

Pigweed, Spiny, *Amaranthus spinosus*
Plantain, English, *Plantago lanceolata*
Plantain-Sorrel Mix (Equal parts *Plantago lanceolata*, *Rumex acetosella*)
Rabbit Bush, *Ambrosia deltoidea*
Ragweed, Desert, *Ambrosia dumosa*
Ragweed, False, *Ambrosia acanthicarpa*
Ragweed, Giant (Tall), *Ambrosia trifida*
Ragweed, Lanceleaf, *Ambrosia bidentata*
Ragweed, Slender, *Ambrosia confertiflora*
Ragweed, Western, *Ambrosia psilostachya*
Russian Thistle, *Salsola kali*
Sagebrush, Common, *Artemisia tridentata*
Sage Mix (Equal parts *Artemisia tridentata*, *Artemisia ludoviciana*)
Sage, Prairie, *Artemisia ludoviciana*
Saltbush, Annual, *Atriplex wrightii*
Scale/Atriplex Mix (Equal parts *Atriplex polycarpa*, *Atriplex lentiformis*, *Atriplex canescens*)
Sorrel, Sheep (Red), *Rumex acetosella*
Waterhemp, Tall, *Amaranthus tuberculatus*
3 Weed Mix (Equal parts *Xanthium strumarium*, *Chenopodium album*, *Amaranthus retroflexus*)
Western Ragweed Mix (Equal parts *Ambrosia acanthicarpa*, *Ambrosia psilostachya*)
Wingscale, *Atriplex canescens*

Plants and Plant Parts

Cotton Linters, *Gossypium hirsutum*
Cottonseed, *Gossypium hirsutum* (For Diagnostic Use Only)
Flax, *Linum usitatissimum* (For Diagnostic Use Only)
Gum, Arabic, *Acacia senegal*
Gum, Karaya, *Sterculia urens*
Gum, Tragacanth, *Astragalus gummifer*
Kapok, *Ceiba pentandra*
Orris Root, *Iris germanica*
Pyrethrum, *Chrysanthemum cinerariifolium*
Tobacco, Cultivated, Leaf, *Nicotiana tabacum*

Pollens - Cultivated Farm Plants

Alfalfa, *Medicago sativa*
Beet, Sugar, *Beta vulgaris*
Corn, Cultivated, *Zea mays*
Oat, Cultivated, *Avena sativa*
Rape (Mustard), *Brassica napus*
Red Clover, *Trifolium pratense*
Rye, Cultivated, *Secale cereale*
Wheat, Cultivated, *Triticum aestivum*

Pollens - Flowers

Daisy, *Leucanthemum vulgare*
Dandelion, *Taraxacum officinale*

Sunflower, *Helianthus annuus*

Molds, Rusts and Smuts

AHH Mold Mix (Equal parts *Alternaria alternata*, *Bipolaris sorokiniana*, *Cladosporium sphaerospermum*)

Alternaria alternata

Alternaria/Hormodendrum Mix (Equal parts *Alternaria alternata*, *Cladosporium sphaerospermum*)

Aspergillus amstelodami

Aspergillus flavus

Aspergillus fumigatus

Aspergillus Mix (Equal parts *Aspergillus amstelodami*, *Aspergillus flavus*, *Aspergillus fumigatus*, *Aspergillus nidulans*, *Aspergillus niger*)

Aspergillus nidulans

Aspergillus niger

Aureobasidium pullulans

Bermuda Grass Smut, *Ustilago cynodontis*

Bipolaris sorokiniana

Botrytis cinerea

Candida albicans

Chaetomium globosum

Cladosporium herbarum

Cladosporium sphaerospermum

Corn Smut, *Ustilago maydis*

Curvularia spicifera

Dematiaceae Mix (Equal parts *Alternaria alternata*, *Aureobasidium pullulans*, *Bipolaris sorokiniana*, *Cladosporium herbarum*, *Curvularia spicifera*, *Helminthosporium solani*)

Epicoccum nigrum

Epidermophyton floccosum

Fusarium Mix (Equal parts *Gibberella fujikuroi*, *Fusarium solani*)

Fusarium solani

Geotrichum candidum

Gibberella fujikuroi

Gliocladium viride

Grain Smut Mix (Equal parts *Ustilago maydis*, *Ustilago tritici*, *Ustilago nuda*, *Ustilago avenae*)

Grass Smut Mix (Equal parts *Ustilago cynodontis*, *Sporisorium cruentum*)

Helminthosporium solani

Hypomyces perniciosus

Loose Kernel Smut, *Sporisorium cruentum*

Loose Smut, Wheat, *Ustilago tritici*

Microsporum canis

Mold Mix #1 (Equal parts *Alternaria alternata*, *Aspergillus niger*, *Bipolaris sorokiniana*, *Cladosporium sphaerospermum*, *Penicillium chrysogenum* var. *chrysogenum*)

Mold Mix #2 (Equal parts *Aureobasidium pullulans*, *Curvularia spicifera*, *Gibberella fujikuroi*, *Mucor plumbeus*, *Rhizopus stolonifer*)

Mold Mix #3 (Equal parts *Alternaria alternata*, *Aspergillus niger*, *Cladosporium*

sphaerospermum, *Penicillium chrysogenum* var. *chrysogenum*)
Monilia Mix (Equal parts *Candida albicans*, *Neurospora intermedia*)
Mucor circinelloides f. *circinelloides*
Mucor circinelloides f. *lusitanicus*
Mucor Mix (Equal parts *Mucor circinelloides* f. *lusitanicus*, *Mucor plumbeus*)
Mucor plumbeus
Neurospora intermedia
New Stock Fungi Mix (Equal parts *Sarocladium strictum*, *Alternaria alternata*, *Aspergillus niger*, *Aureobasidium pullulans*, *Bipolaris sorokiniana*, *Botrytis cinerea*, *Candida albicans*, *Chaetomium globosum*, *Cladosporium sphaerospermum*, *Epicoccum nigrum*, *Gibberella fujikuroi*, *Mucor plumbeus*, *Penicillium chrysogenum* var. *chrysogenum*, *Phoma betae*, *Rhizopus stolonifer*, *Trichophyton mentagrophytes*)
Oat Smut, *Ustilago avenae*
Paecilomyces variotii
Penicillium chrysogenum var. *chrysogenum*
Penicillium digitatum
Penicillium Mix (Equal parts *Penicillium camemberti*, *Penicillium chrysogenum*, *Penicillium digitatum*, *Penicillium chrysogenum* var. *chrysogenum*, *Penicillium roqueforti*)
Phoma betae
Phycomycetes Mix (Equal parts *Mucor circinelloides* f. *lusitanicus*, *Rhizopus stolonifer*)
Rhizopus arrhizus
Rhizopus Mix (Equal parts *Rhizopus stolonifer*, *Rhizopus arrhizus*)
Rhizopus stolonifer
Rhodotorula mucilaginosa
Saccharomyces cerevisiae
Sarocladium strictum
Stemphylium solani
Trichoderma harzianum
Trichophyton mentagrophytes
Trichophyton rubrum
Trichothecium roseum
Animal Allergens
Canary Feathers, *Serinus canaria*
Cattle Epithelia, *Bos taurus*
Chicken Feathers, *Gallus gallus*
Dog Epithelia, *Canis lupus familiaris*
Duck Feathers, *Anas platyrhynchos*
Gerbil Epithelia, *Meriones unguiculatus*
Goat Epithelia, *Capra hircus*
Goose Feathers, *Anser anser*
Guinea Pig Epithelia, *Cavia porcellus*
Hamster Epithelia, *Mesocricetus auratus*
Hog Epithelia, *Sus scrofa*
Horse Epithelia, *Equus caballus*
Mixed Feathers (Equal parts *Gallus gallus*, *Anas platyrhynchos*, *Anser anser*)
Mouse Epithelia, *Mus musculus*

Parakeet Feathers, *Melopsittacus undulatus*

Rabbit Epithelia, *Oryctolagus cuniculus*

Rat Epithelia, *Rattus norvegicus*

Silk Worm Cocoon, *Bombyx mori*

Insects (Whole Body)

Ant, Black Carpenter, *Camponotus pennsylvanicus*

Ant, Fire, *Solenopsis invicta*

Ant, Fire, *Solenopsis richteri*

Cockroach, American, *Periplaneta americana*

Cockroach, German, *Blattella germanica*

2 Cockroach Mix (Equal parts *Periplaneta americana*, *Blattella germanica*)

Deer Fly, *Chrysops vittatus*

Flea, *Ctenocephalis felis* (For Diagnostic Use Only)

House Fly, *Musca domestica* (For Diagnostic Use Only)

Mosquito, *Aedes taeniorhynchus* (For Diagnostic Use Only)

Food - Animal Products and Poultry Products

Beef, *Bos taurus*

Chicken Meat, *Gallus gallus*

Egg, White, Chicken, *Gallus gallus*

Egg, Whole, Chicken, *Gallus gallus*

Egg, Yolk, Chicken, *Gallus gallus*

Lamb, *Ovis aries*

Pork, *Sus scrofa*

Turkey Meat, *Meleagris gallopavo*

Food - Dairy Products

Milk, Cow, *Bos taurus*

Food - Fish and Shellfish

Bass, Black, *Centropomus striata*

Catfish, Channel, *Ictalurus punctatus*

Clam, Northern Quahog, *Mercenaria mercenaria*

Cod, Atlantic, *Gadus morhua*

Crab, Blue, *Callinectes sapidus*

Fish Mix (Equal parts *Gadus morhua*, *Paralichthys lethostigma*, *Hippoglossus hippoglossus*, *Scomber scombrus*, *Thunnus albacares*)

Flounder, Southern, *Paralichthys lethostigma*

Lobster, American, *Homarus americanus*

Mackerel, King/Atlantic, *Scomber scombrus*

Oyster, Atlantic/Eastern, *Crassostrea virginica*

Perch, Ocean, *Sebastes alutus*

Salmon, Atlantic, *Salmo salar*

Scallops, Sea, *Placopecten magellanicus*

Shellfish Mix (Equal parts *Mercenaria mercenaria*, *Callinectes sapidus*, *Crassostrea virginica*, *Placopecten magellanicus*, *Farfantepenaeus aztecus*)

Shrimp, Brown, *Farfantepenaeus aztecus*

Trout, Rainbow, *Oncorhynchus mykiss*

Tuna, Yellowfin, *Thunnus albacares*

Food - Plant Source

Almond, *Prunus dulcis*

Apple, *Malus pumila*

Apricot, *Prunus armeniaca*

Banana, *Musa acuminata*

Barley, Whole Grain, *Hordeum vulgare* (For Diagnostic Use Only, Contains SFS*)

Bean, Lima, *Phaseolus lunatus*

Bean, Navy, *Phaseolus vulgaris*

Bean, String Green, *Phaseolus vulgaris*

Blueberry, Velvetleaf, *Vaccinium myrtilloides*

Brazil Nut, *Bertholletia excelsa*

Broccoli, *Brassica oleracea* var. *botrytis*

Buckwheat, *Fagopyrum esculentum*

Cabbage, *Brassica oleracea* var. *capitata*

Cacao Bean, *Theobroma cacao*

Cantaloupe, *Cucumis melo*

Carrot, *Daucus carota*

Cashew Nut, *Anacardium occidentale*

Cauliflower, *Brassica oleracea* var. *botrytis*

Celery, *Apium graveolens* var. *dulce*

Cherry, Sweet, *Prunus avium*

Cinnamon, *Cinnamomum verum*

Coconut, *Cocos nucifera*

Coffee, *Coffea arabica* (For Diagnostic Use Only)

Corn, *Zea mays*

Cranberry, *Vaccinium macrocarpon*

Cucumber, *Cucumis sativus*

Garlic, *Allium sativum*

Ginger, *Zingiber officinale*

Grape, White Seedless, *Vitis vinifera*

Grapefruit, *Citrus X paradisi*

Hazelnut (Filbert), *Corylus americana*

Hops, *Humulus lupulus*

Lemon, *Citrus X limon*

Lettuce, *Lactuca sativa*

Malt (Barley), *Hordeum vulgare*

Mushroom, *Agaricus campestris*

Mustard Seed, *Sinapis alba*

Nutmeg, *Myristica fragrans*

Oat, *Avena sativa* (For Diagnostic Use Only, Contains SFS*)

Olive, Green, *Olea europaea*

Onion, *Allium cepa*

Orange, *Citrus X sinensis*

Pea, Green or English, *Pisum sativum*

Peach, *Prunus persica*

Peanut, *Arachis hypogaea*
Pear, *Pyrus communis*
Pecan, *Carya illinoensis*
Pepper, Black, *Piper nigrum*
Pepper, Green, *Capsicum annuum*
Pineapple, *Ananas comosus* (For Diagnostic Use Only, Contains SFS*)
Potato, Sweet, *Ipomoea batatas*
Potato, White, *Solanum tuberosum*
Raspberry, Red, *Rubus idaeus*
Rice, *Oryza sativa*
Rye, *Secale cereale* (For Diagnostic Use Only, Contains SFS*)
Sesame Seed, *Sesamum indicum*
Soybean, *Glycine max*
Spinach, *Spinacia oleracea* (For Diagnostic Use Only, Contains SFS*)
Squash, Yellow Summer, *Cucurbita pepo* var. *ovifera*
Strawberry, *Fragaria X ananassa*
Tomato, *Solanum lycopersicum*
Vanilla, *Vanilla planifolia*
Walnut, Black, *Juglans nigra*
Walnut, English, *Juglans regia*
Watermelon, *Citrullus lanatus*
Wheat, Whole, *Triticum aestivum* (For Diagnostic Use Only, Contains SFS*)

*SFS – Sodium Formadehyde Sulfoxylate

16.2 Storage and Handling

Maintain at 2 to 8°C (36 to 46°F) during storage and use.

Dilutions of concentrated extracts that result in a glycerin content of less than 50% can reduce extract stability. Extract dilutions at 1:100 volume to volume should be kept no longer than a month, and more dilute solutions no more than a week. The potency of a dilution can be checked by skin test comparison to a fresh dilution of the extract on a known allergic patient.

17 PATIENT COUNSELING INFORMATION

Instruct patient to remain under observation in the office for 30 minutes or longer after an injection.

Caution patient that reactions can occur more than 30 minutes after skin testing or an injection.

Instruct patient to recognize the following symptoms as adverse reactions and to immediately return to the office or immediately seek other medical attention if any of these symptoms occur following skin testing or an injection:

- Unusual swelling and/or tenderness at the injection site
- Hives or itching of the skin
- Swelling of the face and/or mouth

- Sneezing, coughing or wheezing
- Shortness of breath
- Nausea
- Dizziness or faintness

Manufacturer:

U.S. License No. 308

Greer Laboratories, Inc.

Lenoir, NC 28645 U.S.A

Sterile Multiple Dose Vial U.S. Rx Only Store at 2-8°C

ALLERGENIC EXTRACT

DOG EPITHELIA

Canis lupus familiaris

Item: E7A05 10 mL 20,000 PNU/mL

Preservative 0.4% Phenol.

No U.S. Standard of Potency. See Package Insert for Contents, Dose and Directions for Use.



GTIN (01) 00322840110627
 S/N (21) 000000000000
 LOT (10) SAMPLE
 EXP (17) 01 Jan 2025



GREER Laboratories, Inc. Lenoir, NC 28645 U.S. License 308

Skin Test Only Vial U.S. Rx Only Store at 2-8°C

ALLERGENIC EXTRACT

HORSE EPITHELIA

Equus caballus

Item: GE17A01 5 mL 1:20 W/V

Preservative 0.2% Phenol.

Contains 50% v/v Glycerin. No U.S. Standard of Potency. See Package Insert for Contents, Dose and Directions for Use.



GTIN (01) 00322840510656
 S/N (21) 000000000000
 LOT (10) SAMPLE
 EXP (17) 01 Jan 2025



GREER Laboratories, Inc. Lenoir, NC 28645 U.S. License 308

Sterile Multiple Dose Vial U.S. Rx Only Store at 2-8°C

ALLERGENIC EXTRACT

MIXED FEATHERS

Equal Parts *G. gallus*, *A. platyrhynchos*, *A. anser*

Item: GEO1A04 50 mL 1:20 W/V

Preservative 0.2% Phenol.

Contains 50% v/v Glycerin. No U.S. Standard of Potency. See Package Insert for Contents, Dose and Directions for Use.



GTIN (01) 00322840910746
S/N (21) 000000000000
LOT (10) SAMPLE
EXP (17) 01 Jan 2025



GREER Laboratories, Inc. Lenoir, NC 28645 U.S. License 308

CANARY FEATHERS

serinus canaria solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1150
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SERINUS CANARIA FEATHER (UNII: 9EL3384IQY) (SERINUS CANARIA FEATHER - UNII:9EL3384IQY)	SERINUS CANARIA FEATHER	10000 [PNU] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1150-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

CATTLE EPITHELIA

bos taurus solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-5100
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BOS TAURUS SKIN (UNII: 7J12CD6O9L) (BOS TAURUS SKIN - UNII:7J12CD6O9L)	BOS TAURUS SKIN	0.05 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-5100-5	5 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		
2	NDC:22840-5100-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		
3	NDC:22840-5100-4	50 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		
4	NDC:22840-5100-3	30 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

CATTLE EPITHELIA

bos taurus solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1102
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BOS TAURUS SKIN (UNII: 7J12CD609L) (BOS TAURUS SKIN - UNII:7J12CD609L)	BOS TAURUS SKIN	0.05 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1102-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

CATTLE EPITHELIA

bos taurus solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1100
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BOS TAURUS SKIN (UNII: 7J12CD609L) (BOS TAURUS SKIN - UNII:7J12CD609L)	BOS TAURUS SKIN	0.1 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1100-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

CHICKEN FEATHERS

gallus gallus solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1154
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
GALLUS GALLUS FEATHER (UNII: 1FCM16V0FV) (GALLUS GALLUS FEATHER - UNII:1FCM16V0FV)	GALLUS GALLUS FEATHER	20000 [PNU] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
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#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1154-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

DOG EPITHELIA				
canis lupus familiaris solution				
Product Information				
Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1105	
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS			
Active Ingredient/Active Moiety				
	Ingredient Name	Basis of Strength	Strength	
	CANIS LUPUS FAMILIARIS SKIN (UNII: X2W7CLE97T) (CANIS LUPUS FAMILIARIS SKIN - UNII:X2W7CLE97T)	CANIS LUPUS FAMILIARIS SKIN	0.01 g in 1 mL	
Inactive Ingredients				
	Ingredient Name	Strength		
	SODIUM BICARBONATE (UNII: 8MDF5V39QO)			
	PHENOL (UNII: 339NCG44TV)			
	SODIUM CHLORIDE (UNII: 451W47IQ8X)			
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1105-4	50 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		
2	NDC:22840-1105-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
BLA	BLA101833	09/15/1981		

DOG EPITHELIA

canis lupus familiaris solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1107
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CANIS LUPUS FAMILIARIS SKIN (UNII: X2W7CLE97T) (CANIS LUPUS FAMILIARIS SKIN - UNII:X2W7CLE97T)	CANIS LUPUS FAMILIARIS SKIN	10000 [PNU] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1107-4	50 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		
2	NDC:22840-1107-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

DUCK FEATHERS

anas platyrhynchos solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-5112
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ANAS PLATYRHYNCHOS FEATHER (UNII: 83B65P4796) (ANAS PLATYRHYNCHOS FEATHER - UNII:83B65P4796)	ANAS PLATYRHYNCHOS FEATHER	0.05 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-5112-5	5 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		
2	NDC:22840-5112-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

CANARY FEATHERS

serinus canaria solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1151
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SERINUS CANARIA FEATHER (UNII: 9EL3384IQY) (SERINUS CANARIA FEATHER - UNII:9EL3384IQY)	SERINUS CANARIA FEATHER	0.001 g in 1 mL

Inactive Ingredients

Ingredient Name		Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)		
PHENOL (UNII: 339NCG44TV)		
SODIUM CHLORIDE (UNII: 451W47IQ8X)		

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1151-1	5 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

GUINEA PIG EPITHELIA				
cavia porcellus solution				
Product Information				
Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1168	
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS			
Active Ingredient/Active Moiety				
Ingredient Name		Basis of Strength	Strength	
CAVIA PORCELLUS SKIN (UNII: GM3H4U6QS8) (CAVIA PORCELLUS SKIN - UNII:GM3H4U6QS8)		CAVIA PORCELLUS SKIN	0.05 g in 1 mL	
Inactive Ingredients				
Ingredient Name		Strength		
SODIUM BICARBONATE (UNII: 8MDF5V39QO)				
PHENOL (UNII: 339NCG44TV)				
SODIUM CHLORIDE (UNII: 451W47IQ8X)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1168-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

GOOSE FEATHERS

anser anser solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1160
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ANSER ANSER FEATHER (UNII: 15XI414745) (ANSER ANSER FEATHER - UNII:15XI414745)	ANSER ANSER FEATHER	0.001 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1160-1	5 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

GUINEA PIG EPITHELIA

cavia porcellus solution

Product Information				
Product Type		NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-5115
Route of Administration		INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		
Active Ingredient/Active Moiety				
Ingredient Name			Basis of Strength	Strength
CAVIA PORCELLUS SKIN (UNII: GM3H4U6QS8) (CAVIA PORCELLUS SKIN - UNII:GM3H4U6QS8)			CAVIA PORCELLUS SKIN	0.05 g in 1 mL
Inactive Ingredients				
Ingredient Name				Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)				
PHENOL (UNII: 339NCG44TV)				
SODIUM CHLORIDE (UNII: 451W47IQ8X)				
GLYCERIN (UNII: PDC6A3C0OX)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-5115-5	5 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		
2	NDC:22840-5115-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		
Marketing Information				
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA		BLA101833	09/15/1981	

GERBIL EPITHELIA				
meriones unguiculatus solution				
Product Information				
Product Type		NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1113
Route of Administration		INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		
Active Ingredient/Active Moiety				

Ingredient Name		Basis of Strength	Strength	
MERIONES UNGUICULATUS SKIN (UNII: 9WN2H714TG) (MERIONES UNGUICULATUS SKIN - UNII:9WN2H714TG)		MERIONES UNGUICULATUS SKIN	0.001 g in 1 mL	
Inactive Ingredients				
Ingredient Name		Strength		
SODIUM BICARBONATE (UNII: 8MDF5V39QO)				
PHENOL (UNII: 339NCG44TV)				
SODIUM CHLORIDE (UNII: 451W47IQ8X)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1113-1	5 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
BLA	BLA101833	09/15/1981		

RAT EPITHELIA

rattus norvegicus solution

Product Information			
Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1145
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		
Active Ingredient/Active Moiety			
Ingredient Name		Basis of Strength	Strength
RATTUS NORVEGICUS SKIN (UNII: Y69HPD48AI) (RATTUS NORVEGICUS SKIN - UNII:Y69HPD48AI)		RATTUS NORVEGICUS SKIN	0.05 g in 1 mL
Inactive Ingredients			
Ingredient Name		Strength	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)			
PHENOL (UNII: 339NCG44TV)			
SODIUM CHLORIDE (UNII: 451W47IQ8X)			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1145-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

RAT EPITHELIA

rattus norvegicus solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1146
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
RATTUS NORVEGICUS SKIN (UNII: Y69HPD48AI) (RATTUS NORVEGICUS SKIN - UNII:Y69HPD48AI)	RATTUS NORVEGICUS SKIN	0.1 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1146-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

RAT EPITHELIA

rattus norvegicus solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1147
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
RATTUS NORVEGICUS SKIN (UNII: Y69HPD48AI) (RATTUS NORVEGICUS SKIN - UNII:Y69HPD48AI)	RATTUS NORVEGICUS SKIN	0.001 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1147-1	5 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

MOUSE EPITHELIA

mus musculus solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1135
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety				
Ingredient Name			Basis of Strength	Strength
MUS MUSCULUS SKIN (UNII: 390AN9GB09) (MUS MUSCULUS SKIN - UNII:390AN9GB09)			MUS MUSCULUS SKIN	0.05 g in 1 mL

Inactive Ingredients	
Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1135-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

MOUSE EPITHELIA			
mus musculus solution			
Product Information			
Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1136
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
MUS MUSCULUS SKIN (UNII: 390AN9GB09) (MUS MUSCULUS SKIN - UNII:390AN9GB09)	MUS MUSCULUS SKIN	0.001 g in 1 mL

Inactive Ingredients	
Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	

PHENOL (UNII: 339NCG44TV)

SODIUM CHLORIDE (UNII: 451W47IQ8X)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1136-1	5 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

HORSE EPITHELIA

equus caballus solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1128
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
EQUUS CABALLUS SKIN (UNII: 88VZV9HGT4) (EQUUS CABALLUS SKIN - UNII:88VZV9HGT4)	EQUUS CABALLUS SKIN	10000 [PNU] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1128-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

MIXED FEATHERS

gallus gallus, anas platyrhynchos and anser anser solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-9104
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
GALLUS GALLUS FEATHER (UNII: 1FCM16V0FV) (GALLUS GALLUS FEATHER - UNII:1FCM16V0FV)	GALLUS GALLUS FEATHER	0.0006666 g in 1 mL
ANAS PLATYRHYNCHOS FEATHER (UNII: 83B65P4796) (ANAS PLATYRHYNCHOS FEATHER - UNII:83B65P4796)	ANAS PLATYRHYNCHOS FEATHER	0.0006666 g in 1 mL
ANSER ANSER FEATHER (UNII: 15XI414745) (ANSER ANSER FEATHER - UNII:15XI414745)	ANSER ANSER FEATHER	0.0006666 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-9104-1	5 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

MIXED FEATHERS

gallus gallus, anas platyrhynchos and anser anser solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-9103
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
GALLUS GALLUS FEATHER (UNII: 1FCM16V0FV) (GALLUS GALLUS FEATHER - UNII:1FCM16V0FV)	GALLUS GALLUS FEATHER	0.0166 g in 1 mL
ANAS PLATYRHYNCHOS FEATHER (UNII: 83B65P4796) (ANAS PLATYRHYNCHOS FEATHER - UNII:83B65P4796)	ANAS PLATYRHYNCHOS FEATHER	0.0166 g in 1 mL
ANSER ANSER FEATHER (UNII: 15XI414745) (ANSER ANSER FEATHER - UNII:15XI414745)	ANSER ANSER FEATHER	0.0166 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-9103-4	50 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

GERBIL EPITHELIA

meriones unguiculatus solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1111
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MERIONES UNGUICULATUS SKIN (UNII: 9WN2H714TG) (MERIONES UNGUICULATUS SKIN - UNII:9WN2H714TG)	MERIONES UNGUICULATUS SKIN	0.05 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1111-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

GOAT EPITHELIA

capra hircus solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1173
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CAPRA HIRCUS SKIN (UNII: JLG9853E2P) (CAPRA HIRCUS SKIN - UNII:JLG9853E2P)	CAPRA HIRCUS SKIN	0.1 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1173-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

HORSE EPITHELIA

equus caballus solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1127
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
EQUUS CABALLUS SKIN (UNII: 88VZV9HGT4) (EQUUS CABALLUS SKIN - UNII:88VZV9HGT4)	EQUUS CABALLUS SKIN	20000 [PNU] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1127-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		
2	NDC:22840-1127-4	50 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

HORSE EPITHELIA

equus caballus solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-5106
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
EQUUS CABALLUS SKIN (UNII: 88VZV9HGT4) (EQUUS CABALLUS SKIN - UNII:88VZV9HGT4)	EQUUS CABALLUS SKIN	0.05 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-5106-5	5 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		
2	NDC:22840-5106-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		
3	NDC:22840-5106-4	50 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

MIXED FEATHERS

gallus gallus, anas platyrhynchos and anser anser solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-9105
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
GALLUS GALLUS FEATHER (UNII: 1FCM16V0FV) (GALLUS GALLUS FEATHER - UNII:1FCM16V0FV)	GALLUS GALLUS FEATHER	0.0003333 g in 1 mL
ANAS PLATYRHYNCHOS FEATHER (UNII: 83B65P4796) (ANAS PLATYRHYNCHOS FEATHER - UNII:83B65P4796)	ANAS PLATYRHYNCHOS FEATHER	0.0003333 g in 1 mL
ANSER ANSER FEATHER (UNII: 15XI414745) (ANSER ANSER FEATHER - UNII:15XI414745)	ANSER ANSER FEATHER	0.0003333 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-9105-1	5 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

RABBIT EPITHELIA

oryctolagus cuniculus solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1138
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ORYCTOLAGUS CUNICULUS SKIN (UNII: Z91W4U43WC) (ORYCTOLAGUS CUNICULUS SKIN - UNII:Z91W4U43WC)	ORYCTOLAGUS CUNICULUS SKIN	0.1 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1138-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

RABBIT EPITHELIA

oryctolagus cuniculus solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1139
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ORYCTOLAGUS CUNICULUS SKIN (UNII: Z91W4U43WC) (ORYCTOLAGUS CUNICULUS SKIN - UNII:Z91W4U43WC)	ORYCTOLAGUS CUNICULUS SKIN	20000 [PNU] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	

SODIUM CHLORIDE (UNII: 451W47IQ8X)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1139-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

RABBIT EPITHELIA

oryctolagus cuniculus solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1140
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ORYCTOLAGUS CUNICULUS SKIN (UNII: Z91W4U43WC) (ORYCTOLAGUS CUNICULUS SKIN - UNII:Z91W4U43WC)	ORYCTOLAGUS CUNICULUS SKIN	0.05 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1140-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

RABBIT EPITHELIA

oryctolagus cuniculus solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-5108
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ORYCTOLAGUS CUNICULUS SKIN (UNII: Z91W4U43WC) (ORYCTOLAGUS CUNICULUS SKIN - UNII:Z91W4U43WC)	ORYCTOLAGUS CUNICULUS SKIN	0.05 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-5108-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		
2	NDC:22840-5108-5	5 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		
3	NDC:22840-5108-4	50 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

RAT EPITHELIA

rattus norvegicus solution

Product Information				
Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1144	
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS			
Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
RATTUS NORVEGICUS SKIN (UNII: Y69HPD48AI) (RATTUS NORVEGICUS SKIN - UNII:Y69HPD48AI)	RATTUS NORVEGICUS SKIN	10000 [PNU] in 1 mL		
Inactive Ingredients				
Ingredient Name	Strength			
SODIUM BICARBONATE (UNII: 8MDF5V39QO)				
PHENOL (UNII: 339NCG44TV)				
SODIUM CHLORIDE (UNII: 451W47IQ8X)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1144-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
BLA	BLA101833	09/15/1981		

HORSE EPITHELIA			
equus caballus solution			
Product Information			
Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1131
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		
Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
EQUUS CABALLUS SKIN (UNII: 88VZV9HGT4) (EQUUS CABALLUS SKIN -	EQUUS CABALLUS	0.001 g	

UNII:88VZV9HGT4)		SKIN	in 1 mL	
Inactive Ingredients				
Ingredient Name			Strength	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)				
PHENOL (UNII: 339NCG44TV)				
SODIUM CHLORIDE (UNII: 451W47IQ8X)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1131-1	5 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
BLA	BLA101833	09/15/1981		

RABBIT EPITHELIA				
oryctolagus cuniculus solution				
Product Information				
Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1142	
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS			
Active Ingredient/Active Moiety				
Ingredient Name		Basis of Strength	Strength	
ORYCTOLAGUS CUNICULUS SKIN (UNII: Z91W4U43WC) (ORYCTOLAGUS CUNICULUS SKIN - UNII:Z91W4U43WC)		ORYCTOLAGUS CUNICULUS SKIN	0.001 g in 1 mL	
Inactive Ingredients				
Ingredient Name			Strength	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)				
PHENOL (UNII: 339NCG44TV)				
SODIUM CHLORIDE (UNII: 451W47IQ8X)				
Packaging				
			Marketing Start	Marketing End

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1142-1	5 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

SILK WORM				
bombyx mori solution				
Product Information				
Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-5517	
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS			
Active Ingredient/Active Moiety				
	Ingredient Name	Basis of Strength	Strength	
	BOMBYX MORI FIBER (UNII: 6LK42KUV6W) (BOMBYX MORI FIBER - UNII:6LK42KUV6W)	BOMBYX MORI FIBER	0.05 g in 1 mL	
Inactive Ingredients				
	Ingredient Name	Strength		
	SODIUM CHLORIDE (UNII: 451W47IQ8X)			
	GLYCERIN (UNII: PDC6A3C0OX)			
	SODIUM BICARBONATE (UNII: 8MDF5V39QO)			
	PHENOL (UNII: 339NCG44TV)			
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-5517-5	5 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		
2	NDC:22840-5517-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	

BLA	BLA101833	09/15/1981	
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HORSE EPITHELIA
equus caballus solution

Product Information			
Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1129
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
EQUUS CABALLUS SKIN (UNII: 88VZV9HGT4) (EQUUS CABALLUS SKIN - UNII:88VZV9HGT4)	EQUUS CABALLUS SKIN	0.05 g in 1 mL

Inactive Ingredients	
Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1129-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		
2	NDC:22840-1129-4	50 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

HORSE EPITHELIA
equus caballus solution

Product Information			
Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1130

Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS			
Active Ingredient/Active Moiety				
	Ingredient Name	Basis of Strength	Strength	
	EQUUS CABALLUS SKIN (UNII: 88VZV9HGT4) (EQUUS CABALLUS SKIN - UNII:88VZV9HGT4)	EQUUS CABALLUS SKIN	0.002 g in 1 mL	
Inactive Ingredients				
	Ingredient Name	Strength		
	SODIUM BICARBONATE (UNII: 8MDF5V39QO)			
	PHENOL (UNII: 339NCG44TV)			
	SODIUM CHLORIDE (UNII: 451W47IQ8X)			
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1130-1	5 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
BLA	BLA101833	09/15/1981		

MIXED FEATHERS			
gallus gallus, anas platyrhynchos and anser anser solution			
Product Information			
Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-9107
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		
Active Ingredient/Active Moiety			
	Ingredient Name	Basis of Strength	Strength
	GALLUS GALLUS FEATHER (UNII: 1FCM16V0FV) (GALLUS GALLUS FEATHER - UNII:1FCM16V0FV)	GALLUS GALLUS FEATHER	0.0166 g in 1 mL
	ANAS PLATYRHYNCHOS FEATHER (UNII: 83B65P4796) (ANAS PLATYRHYNCHOS FEATHER - UNII:83B65P4796)	ANAS PLATYRHYNCHOS FEATHER	0.0166 g in 1 mL
	ANSER ANSER FEATHER (UNII: 15XI414745) (ANSER ANSER FEATHER - UNII:15XI414745)	ANSER ANSER FEATHER	0.0166 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-9107-4	50 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		
2	NDC:22840-9107-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		
3	NDC:22840-9107-5	5 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

MOUSE EPITHELIA

mus musculus solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1133
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MUS MUSCULUS SKIN (UNII: 390AN9GB09) (MUS MUSCULUS SKIN - UNII:390AN9GB09)	MUS MUSCULUS SKIN	0.1 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1133-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

MOUSE EPITHELIA

mus musculus solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1134
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MUS MUSCULUS SKIN (UNII: 390AN9GB09) (MUS MUSCULUS SKIN - UNII:390AN9GB09)	MUS MUSCULUS SKIN	20000 [PNU] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1134-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing	Application Number or Monograph	Marketing Start	Marketing End
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Category	Citation	Date	Date
BLA	BLA101833	09/15/1981	

GERBIL EPITHELIA

meriones unguiculatus solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1112
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MERIONES UNGUICULATUS SKIN (UNII: 9WN2H714TG) (MERIONES UNGUICULATUS SKIN - UNII:9WN2H714TG)	MERIONES UNGUICULATUS SKIN	10000 [PNU] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1112-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

GERBIL EPITHELIA

meriones unguiculatus solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-5102
Route of Administration	INTRADERMAL, PERCUTANEOUS,		

Route of Administration SUBCUTANEOUS

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MERIONES UNGUICULATUS SKIN (UNII: 9WN2H714TG) (MERIONES UNGUICULATUS SKIN - UNII:9WN2H714TG)	MERIONES UNGUICULATUS SKIN	0.05 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-5102-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		
2	NDC:22840-5102-5	5 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

HAMSTER EPITHELIA

mesocricetus auratus solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1119
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MESOCRICETUS AURATUS SKIN (UNII: 3K873H631W) (MESOCRICETUS AURATUS SKIN - UNII:3K873H631W)	MESOCRICETUS AURATUS SKIN	0.001 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1119-1	5 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

HAMSTER EPITHELIA

mesocricetus auratus solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-5104
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MESOCRICETUS AURATUS SKIN (UNII: 3K873H631W) (MESOCRICETUS AURATUS SKIN - UNII:3K873H631W)	MESOCRICETUS AURATUS SKIN	0.05 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	NDC:22840-	5 mL in 1 BOTTLE, DROPPER; Type 0: Not a		

1	NDC:22840-5104-5	5 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		
2	NDC:22840-5104-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

HOG EPITHELIA				
sus scrofa solution				
Product Information				
Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1122	
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS			
Active Ingredient/Active Moiety				
	Ingredient Name	Basis of Strength	Strength	
	SUS SCROFA SKIN (UNII: 3EM4VW6TQN) (SUS SCROFA SKIN - UNII:3EM4VW6TQN)	SUS SCROFA SKIN	20000 [PNU] in 1 mL	
Inactive Ingredients				
	Ingredient Name	Strength		
	SODIUM BICARBONATE (UNII: 8MDF5V39QO)			
	PHENOL (UNII: 339NCG44TV)			
	SODIUM CHLORIDE (UNII: 451W47IQ8X)			
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1122-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
BLA	BLA101833	09/15/1981		

HOG EPITHELIA

sus scrofa solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1174
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SUS SCROFA SKIN (UNII: 3EM4VW6TQN) (SUS SCROFA SKIN - UNII:3EM4VW6TQN)	SUS SCROFA SKIN	10000 [PNU] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1174-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

HORSE EPITHELIA

equus caballus solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1126
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety				
Ingredient Name		Basis of Strength	Strength	
EQUUS CABALLUS SKIN (UNII: 88VZV9HGT4) (EQUUS CABALLUS SKIN - UNII:88VZV9HGT4)		EQUUS CABALLUS SKIN	0.1 g in 1 mL	
Inactive Ingredients				
Ingredient Name		Strength		
SODIUM BICARBONATE (UNII: 8MDF5V39QO)				
PHENOL (UNII: 339NCG44TV)				
SODIUM CHLORIDE (UNII: 451W47IQ8X)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1126-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		
2	NDC:22840-1126-4	50 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
BLA	BLA101833	09/15/1981		

RAT EPITHELIA			
rattus norvegicus solution			
Product Information			
Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-5109
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		
Active Ingredient/Active Moiety			
Ingredient Name		Basis of Strength	Strength
RATTUS NORVEGICUS SKIN (UNII: Y69HPD48AI) (RATTUS NORVEGICUS SKIN - UNII:Y69HPD48AI)		RATTUS NORVEGICUS SKIN	0.05 g in 1 mL
Inactive Ingredients			
Ingredient Name		Strength	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)			

PHENOL (UNII: 339NCG44TV)				
SODIUM CHLORIDE (UNII: 451W47IQ8X)				
GLYCERIN (UNII: PDC6A3C0OX)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-5109-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		
2	NDC:22840-5109-5	5 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
BLA	BLA101833	09/15/1981		

SILK WORM				
bombyx mori solution				
Product Information				
Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1537	
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS			
Active Ingredient/Active Moiety				
	Ingredient Name	Basis of Strength	Strength	
	BOMBYX MORI FIBER (UNII: 6LK42KUV6W) (BOMBYX MORI FIBER - UNII:6LK42KUV6W)	BOMBYX MORI FIBER	0.1 g in 1 mL	
Inactive Ingredients				
	Ingredient Name	Strength		
	SODIUM BICARBONATE (UNII: 8MDF5V39QO)			
	PHENOL (UNII: 339NCG44TV)			
	SODIUM CHLORIDE (UNII: 451W47IQ8X)			
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1537-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

MIXED FEATHERS

gallus gallus, anas platyrhynchos and anser anser solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-9100
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
GALLUS GALLUS FEATHER (UNII: 1FCM16V0FV) (GALLUS GALLUS FEATHER - UNII:1FCM16V0FV)	GALLUS GALLUS FEATHER	0.033 g in 1 mL
ANAS PLATYRHYNCHOS FEATHER (UNII: 83B65P4796) (ANAS PLATYRHYNCHOS FEATHER - UNII:83B65P4796)	ANAS PLATYRHYNCHOS FEATHER	0.033 g in 1 mL
ANSER ANSER FEATHER (UNII: 15XI414745) (ANSER ANSER FEATHER - UNII:15XI414745)	ANSER ANSER FEATHER	0.033 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-9100-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		
2	NDC:22840-9100-4	50 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

MIXED FEATHERS

gallus gallus, anas platyrhynchos and anser anser solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-9101
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
GALLUS GALLUS FEATHER (UNII: 1FCM16V0FV) (GALLUS GALLUS FEATHER - UNII:1FCM16V0FV)	GALLUS GALLUS FEATHER	6666.6666 [PNU] in 1 mL
ANAS PLATYRHYNCHOS FEATHER (UNII: 83B65P4796) (ANAS PLATYRHYNCHOS FEATHER - UNII:83B65P4796)	ANAS PLATYRHYNCHOS FEATHER	6666.6666 [PNU] in 1 mL
ANSER ANSER FEATHER (UNII: 15XI414745) (ANSER ANSER FEATHER - UNII:15XI414745)	ANSER ANSER FEATHER	6666.6666 [PNU] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-9101-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		
2	NDC:22840-9101-4	50 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

MIXED FEATHERS

gallus gallus, anas platyrhynchos and anser anser solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-9102
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
GALLUS GALLUS FEATHER (UNII: 1FCM16V0FV) (GALLUS GALLUS FEATHER - UNII:1FCM16V0FV)	GALLUS GALLUS FEATHER	3333.3333 [PNU] in 1 mL
ANAS PLATYRHYNCHOS FEATHER (UNII: 83B65P4796) (ANAS PLATYRHYNCHOS FEATHER - UNII:83B65P4796)	ANAS PLATYRHYNCHOS FEATHER	3333.3333 [PNU] in 1 mL
ANSER ANSER FEATHER (UNII: 15XI414745) (ANSER ANSER FEATHER - UNII:15XI414745)	ANSER ANSER FEATHER	3333.3333 [PNU] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-9102-4	50 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

SILK WORM

bombyx mori solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1538
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety				
Ingredient Name		Basis of Strength	Strength	
BOMBYX MORI FIBER (UNII: 6LK42KUV6W) (BOMBYX MORI FIBER - UNII:6LK42KUV6W)		BOMBYX MORI FIBER	0.001 g in 1 mL	
Inactive Ingredients				
Ingredient Name		Strength		
SODIUM BICARBONATE (UNII: 8MDF5V39QO)				
PHENOL (UNII: 339NCG44TV)				
SODIUM CHLORIDE (UNII: 451W47IQ8X)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1538-1	5 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
BLA	BLA101833	09/15/1981		

RABBIT EPITHELIA			
oryctolagus cuniculus solution			
Product Information			
Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1141
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		
Active Ingredient/Active Moiety			
Ingredient Name		Basis of Strength	Strength
ORYCTOLAGUS CUNICULUS SKIN (UNII: Z91W4U43WC) (ORYCTOLAGUS CUNICULUS SKIN - UNII:Z91W4U43WC)		ORYCTOLAGUS CUNICULUS SKIN	10000 [PNU] in 1 mL
Inactive Ingredients			
Ingredient Name		Strength	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)			
PHENOL (UNII: 339NCG44TV)			
SODIUM CHLORIDE (UNII: 451W47IQ8X)			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1141-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

MOUSE EPITHELIA

mus musculus solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-5107
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MUS MUSCULUS SKIN (UNII: 390AN9GB09) (MUS MUSCULUS SKIN - UNII:390AN9GB09)	MUS MUSCULUS SKIN	0.05 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-5107-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		
2	NDC:22840-5107-5	5 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

HAMSTER EPITHELIA

mesocricetus auratus solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1117
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MESOCRICETUS AURATUS SKIN (UNII: 3K873H631W) (MESOCRICETUS AURATUS SKIN - UNII:3K873H631W)	MESOCRICETUS AURATUS SKIN	0.05 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1117-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

HAMSTER EPITHELIA

mesocricetus auratus solution

Product Information

Item Code NDC 22840

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1118	
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS			
Active Ingredient/Active Moiety				
	Ingredient Name	Basis of Strength	Strength	
	MESOCRICETUS AURATUS SKIN (UNII: 3K873H631W) (MESOCRICETUS AURATUS SKIN - UNII:3K873H631W)	MESOCRICETUS AURATUS SKIN	10000 [PNU] in 1 mL	
Inactive Ingredients				
	Ingredient Name		Strength	
	SODIUM BICARBONATE (UNII: 8MDF5V39QO)			
	PHENOL (UNII: 339NCG44TV)			
	SODIUM CHLORIDE (UNII: 451W47IQ8X)			
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1118-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
BLA	BLA101833	09/15/1981		

HOG EPITHELIA			
sus scrofa solution			
Product Information			
Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1121
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		
Active Ingredient/Active Moiety			
	Ingredient Name	Basis of Strength	Strength
	SUS SCROFA SKIN (UNII: 3EM4VW6TQN) (SUS SCROFA SKIN - UNII:3EM4VW6TQN)	SUS SCROFA SKIN	0.1 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1121-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

HOG EPITHELIA

sus scrofa solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1123
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SUS SCROFA SKIN (UNII: 3EM4VW6TQN) (SUS SCROFA SKIN - UNII:3EM4VW6TQN)	SUS SCROFA SKIN	0.002 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
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1	NDC:22840-1123-1	5 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		
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Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

HOG EPITHELIA

sus scrofa solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1125
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SUS SCROFA SKIN (UNII: 3EM4VW6TQN) (SUS SCROFA SKIN - UNII:3EM4VW6TQN)	SUS SCROFA SKIN	0.001 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1125-1	5 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

HOG EPITHELIA

sus scrofa solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-5105
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SUS SCROFA SKIN (UNII: 3EM4VW6TQN) (SUS SCROFA SKIN - UNII:3EM4VW6TQN)	SUS SCROFA SKIN	0.05 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-5105-5	5 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		
2	NDC:22840-5105-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

PARAKEET FEATHERS

melopsitticus undulatus solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1162
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MELOPSITTACUS UNDULATUS FEATHER (UNII: 1XY644QKWG) (MELOPSITTACUS UNDULATUS FEATHER - UNII:1XY644QKWG)	MELOPSITTACUS UNDULATUS FEATHER	0.05 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1162-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

PARAKEET FEATHERS

melopsitticus undulatus solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1163
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MELOPSITTACUS UNDULATUS FEATHER (UNII: 1XY644QKWG) (MELOPSITTACUS UNDULATUS FEATHER - UNII:1XY644QKWG)	MELOPSITTACUS UNDULATUS FEATHER	10000 [PNU] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1163-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

PARAKEET FEATHERS

melopsitticus undulatus solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1164
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MELOPSITTACUS UNDULATUS FEATHER (UNII: 1XY644QKWG) (MELOPSITTACUS UNDULATUS FEATHER - UNII:1XY644QKWG)	MELOPSITTACUS UNDULATUS FEATHER	0.001 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1164-1	5 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
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BLA	BLA101833	09/15/1981	
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GOOSE FEATHERS

anser anser solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-5113
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ANSER ANSER FEATHER (UNII: 15XI414745) (ANSER ANSER FEATHER - UNII:15XI414745)	ANSER ANSER FEATHER	0.05 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-5113-5	5 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		
2	NDC:22840-5113-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

PARAKEET FEATHERS

melopsitticus undulatus solution

Product Information

		Item Code	NDC:22840
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Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-5114	
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS			
Active Ingredient/Active Moiety				
Ingredient Name		Basis of Strength	Strength	
MELOPSITTACUS UNDULATUS FEATHER (UNII: 1XY644QKWG) (MELOPSITTACUS UNDULATUS FEATHER - UNII:1XY644QKWG)		MELOPSITTACUS UNDULATUS FEATHER	0.025 g in 1 mL	
Inactive Ingredients				
Ingredient Name		Strength		
SODIUM BICARBONATE (UNII: 8MDF5V39QO)				
PHENOL (UNII: 339NCG44TV)				
SODIUM CHLORIDE (UNII: 451W47IQ8X)				
GLYCERIN (UNII: PDC6A3C0OX)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-5114-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		
2	NDC:22840-5114-5	5 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
BLA	BLA101833	09/15/1981		

CANARY FEATHERS			
serinus canaria solution			
Product Information			
Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1149
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		
Active Ingredient/Active Moiety			
Ingredient Name		Basis of Strength	Strength
SERINUS CANARIA FEATHER (UNII: 9EL3384IQY) (SERINUS CANARIA FEATHER - UNII:9EL3384IQY)		SERINUS CANARIA FEATHER	0.05 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1149-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

DOG EPITHELIA

canis lupus familiaris solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1110
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CANIS LUPUS FAMILIARIS SKIN (UNII: X2W7CLE97T) (CANIS LUPUS FAMILIARIS SKIN - UNII:X2W7CLE97T)	CANIS LUPUS FAMILIARIS SKIN	0.001 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
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1	NDC:22840-1110-1	5 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product	
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Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

DUCK FEATHERS

anas platyrhynchos solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1176
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ANAS PLATYRHYNCHOS FEATHER (UNII: 83B65P4796) (ANAS PLATYRHYNCHOS FEATHER - UNII:83B65P4796)	ANAS PLATYRHYNCHOS FEATHER	0.1 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1176-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

DOG EPITHELIA

canis lupus familiaris solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-5101
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CANIS LUPUS FAMILIARIS SKIN (UNII: X2W7CLE97T) (CANIS LUPUS FAMILIARIS SKIN - UNII:X2W7CLE97T)	CANIS LUPUS FAMILIARIS SKIN	0.05 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-5101-4	50 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		
2	NDC:22840-5101-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		
3	NDC:22840-5101-5	5 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

DOG EPITHELIA

canis lupus familiaris solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1108
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CANIS LUPUS FAMILIARIS SKIN (UNII: X2W7CLE97T) (CANIS LUPUS FAMILIARIS SKIN - UNII:X2W7CLE97T)	CANIS LUPUS FAMILIARIS SKIN	0.05 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1108-4	50 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		
2	NDC:22840-1108-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

CANARY FEATHERS

serinus canaria solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-5110
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SERINUS CANARIA FEATHER (UNII: 9EL3384IQY) (SERINUS CANARIA FEATHER - UNII:9EL3384IQY)	SERINUS CANARIA FEATHER	0.05 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	

SODIUM BICARBONATE (UNII: 8MDF5V39QO)				
PHENOL (UNII: 339NCG44TV)				
SODIUM CHLORIDE (UNII: 451W47IQ8X)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-5110-5	5 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		
2	NDC:22840-5110-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
BLA	BLA101833	09/15/1981		

CATTLE EPITHELIA				
bos taurus solution				
Product Information				
Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1101	
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS			
Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
BOS TAURUS SKIN (UNII: 7J12CD6O9L) (BOS TAURUS SKIN - UNII:7J12CD6O9L)	BOS TAURUS SKIN	20000 [PNU] in 1 mL		
Inactive Ingredients				
Ingredient Name	Strength			
SODIUM BICARBONATE (UNII: 8MDF5V39QO)				
PHENOL (UNII: 339NCG44TV)				
SODIUM CHLORIDE (UNII: 451W47IQ8X)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a		

1101-2	Combination Product		
Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

CATTLE EPITHELIA

bos taurus solution

Product Information			
Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1103
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
BOS TAURUS SKIN (UNII: 7J12CD6O9L) (BOS TAURUS SKIN - UNII:7J12CD6O9L)	BOS TAURUS SKIN	0.001 g in 1 mL	

Inactive Ingredients		
Ingredient Name	Strength	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)		
PHENOL (UNII: 339NCG44TV)		
SODIUM CHLORIDE (UNII: 451W47IQ8X)		

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1103-1	5 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

CATTLE EPITHELIA

bos taurus solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1175
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BOS TAURUS SKIN (UNII: 7J12CD609L) (BOS TAURUS SKIN - UNII:7J12CD609L)	BOS TAURUS SKIN	10000 [PNU] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1175-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

CHICKEN FEATHERS

gallus gallus solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1153
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
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Ingredient Name		Basis of Strength	Strength	
GALLUS GALLUS FEATHER (UNII: 1FCM16V0FV) (GALLUS GALLUS FEATHER - UNII:1FCM16V0FV)		GALLUS GALLUS FEATHER	0.1 g in 1 mL	
Inactive Ingredients				
Ingredient Name		Strength		
SODIUM BICARBONATE (UNII: 8MDF5V39QO)				
PHENOL (UNII: 339NCG44TV)				
SODIUM CHLORIDE (UNII: 451W47IQ8X)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1153-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
BLA	BLA101833	09/15/1981		

CHICKEN FEATHERS

gallus gallus solution

Product Information			
Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1155
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		
Active Ingredient/Active Moiety			
Ingredient Name		Basis of Strength	Strength
GALLUS GALLUS FEATHER (UNII: 1FCM16V0FV) (GALLUS GALLUS FEATHER - UNII:1FCM16V0FV)		GALLUS GALLUS FEATHER	0.001 g in 1 mL
Inactive Ingredients			
Ingredient Name		Strength	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)			
PHENOL (UNII: 339NCG44TV)			
SODIUM CHLORIDE (UNII: 451W47IQ8X)			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1155-1	5 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

CHICKEN FEATHERS

gallus gallus solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-5111
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
GALLUS GALLUS FEATHER (UNII: 1FCM16V0FV) (GALLUS GALLUS FEATHER - UNII:1FCM16V0FV)	GALLUS GALLUS FEATHER	0.05 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-5111-5	5 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		
2	NDC:22840-5111-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

DOG EPITHELIA

canis lupus familiaris solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1106
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CANIS LUPUS FAMILIARIS SKIN (UNII: X2W7CLE97T) (CANIS LUPUS FAMILIARIS SKIN - UNII:X2W7CLE97T)	CANIS LUPUS FAMILIARIS SKIN	20000 [PNU] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1106-4	50 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		
2	NDC:22840-1106-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

GOAT EPITHELIA

capra hircus solution

Product Information				
Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-5103	
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS			
Active Ingredient/Active Moiety				
	Ingredient Name	Basis of Strength	Strength	
	CAPRA HIRCUS SKIN (UNII: JLG9853E2P) (CAPRA HIRCUS SKIN - UNII:JLG9853E2P)	CAPRA HIRCUS SKIN	0.05 g in 1 mL	
Inactive Ingredients				
	Ingredient Name		Strength	
	SODIUM BICARBONATE (UNII: 8MDF5V39QO)			
	PHENOL (UNII: 339NCG44TV)			
	SODIUM CHLORIDE (UNII: 451W47IQ8X)			
	GLYCERIN (UNII: PDC6A3C0OX)			
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-5103-5	5 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		
2	NDC:22840-5103-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
BLA	BLA101833	09/15/1981		

GOOSE FEATHERS			
anser anser solution			
Product Information			
Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1159
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		
Active Ingredient/Active Moiety			
	Ingredient Name	Basis of Strength	Strength

Ingredient Name		Basis of Strength	Strength	
ANSER ANSER FEATHER (UNII: 15XI414745) (ANSER ANSER FEATHER - UNII:15XI414745)		ANSER ANSER FEATHER	0.1 g in 1 mL	
Inactive Ingredients				
Ingredient Name		Strength		
SODIUM BICARBONATE (UNII: 8MDF5V39QO)				
PHENOL (UNII: 339NCG44TV)				
SODIUM CHLORIDE (UNII: 451W47IQ8X)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1159-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
BLA	BLA101833	09/15/1981		

DUCK FEATHERS			
anas platyrhynchos solution			
Product Information			
Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1157
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		
Active Ingredient/Active Moiety			
Ingredient Name		Basis of Strength	Strength
ANAS PLATYRHYNCHOS FEATHER (UNII: 83B65P4796) (ANAS PLATYRHYNCHOS FEATHER - UNII:83B65P4796)		ANAS PLATYRHYNCHOS FEATHER	0.05 g in 1 mL
Inactive Ingredients			
Ingredient Name		Strength	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)			
PHENOL (UNII: 339NCG44TV)			
SODIUM CHLORIDE (UNII: 451W47IQ8X)			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1157-1	5 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

GOAT EPITHELIA

capra hircus solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1116
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CAPRA HIRCUS SKIN (UNII: JLG9853E2P) (CAPRA HIRCUS SKIN - UNII:JLG9853E2P)	CAPRA HIRCUS SKIN	0.001 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1116-1	5 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

GUINEA PIG EPITHELIA

cavia porcellus solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1167
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CAVIA PORCELLUS SKIN (UNII: GM3H4U6QS8) (CAVIA PORCELLUS SKIN - UNII:GM3H4U6QS8)	CAVIA PORCELLUS SKIN	0.1 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W471Q8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1167-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

GUINEA PIG EPITHELIA

cavia porcellus solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1169
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CAVIA PORCELLUS SKIN (UNII: GM3H4U6QS8) (CAVIA PORCELLUS SKIN - UNII:GM3H4U6QS8)	CAVIA PORCELLUS SKIN	10000 [PNU] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1169-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

GUINEA PIG EPITHELIA

cavia porcellus solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1171
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CAVIA PORCELLUS SKIN (UNII: GM3H4U6QS8) (CAVIA PORCELLUS SKIN - UNII:GM3H4U6QS8)	CAVIA PORCELLUS SKIN	0.001 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	

PHENOL (UNII: 339NCG44TV)

SODIUM CHLORIDE (UNII: 451W47IQ8X)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1171-1	5 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

GUINEA PIG EPITHELIA

cavia porcellus solution

Product Information

Product Type	NON-STANDARDIZED ALLERGENIC	Item Code (Source)	NDC:22840-1170
Route of Administration	INTRADERMAL, PERCUTANEOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CAVIA PORCELLUS SKIN (UNII: GM3H4U6QS8) (CAVIA PORCELLUS SKIN - UNII:GM3H4U6QS8)	CAVIA PORCELLUS SKIN	20000 [PNU] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PHENOL (UNII: 339NCG44TV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22840-1170-2	10 mL in 1 VIAL, MULTI-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA101833	09/15/1981	

Labeler - Greer Laboratories, Inc. (024671414)

Registrant - Greer Laboratories, Inc. (024671414)

Establishment

Name	Address	ID/FEI	Business Operations
Greer Laboratories, Inc.		024671414	manufacture(22840-1100, 22840-1101, 22840-1102, 22840-1103, 22840-1105, 22840-1106, 22840-1107, 22840-1108, 22840-1110, 22840-1111, 22840-1112, 22840-1113, 22840-1116, 22840-1117, 22840-1118, 22840-1119, 22840-1121, 22840-1122, 22840-1123, 22840-1125, 22840-1126, 22840-1127, 22840-1128, 22840-1129, 22840-1130, 22840-1131, 22840-1133, 22840-1134, 22840-1135, 22840-1136, 22840-1138, 22840-1139, 22840-1140, 22840-1141, 22840-1142, 22840-1144, 22840-1145, 22840-1146, 22840-1147, 22840-1149, 22840-1150, 22840-1151, 22840-1153, 22840-1154, 22840-1155, 22840-1157, 22840-1159, 22840-1160, 22840-1162, 22840-1163, 22840-1164, 22840-1167, 22840-1168, 22840-1169, 22840-1170, 22840-1171, 22840-1173, 22840-1174, 22840-1175, 22840-1176, 22840-1537, 22840-1538, 22840-5100, 22840-5101, 22840-5102, 22840-5103, 22840-5104, 22840-5105, 22840-5106, 22840-5107, 22840-5108, 22840-5109, 22840-5110, 22840-5111, 22840-5112, 22840-5113, 22840-5114, 22840-5115, 22840-5517, 22840-9100, 22840-9101, 22840-9102, 22840-9103, 22840-9104, 22840-9105, 22840-9107)

Revised: 11/2023

Greer Laboratories, Inc.