

IMLYGIC- talimogene laherparepvec injection, suspension

Amgen Inc

HIGHLIGHTS OF PRESCRIBING INFORMATION

These highlights do not include all the information needed to use IMLYGIC safely and effectively. See full prescribing information for IMLYGIC.

IMLYGIC® (talimogene laherparepvec)

Suspension for intralesional injection

Initial U.S. Approval: 2015

RECENT MAJOR CHANGES

Warnings and Precautions (5.2)	2/2023
Warnings and Precautions (5.7)	6/2022

INDICATIONS AND USAGE

IMLYGIC is a genetically modified oncolytic viral therapy indicated for the local treatment of unresectable cutaneous, subcutaneous, and nodal lesions in patients with melanoma recurrent after initial surgery.

Limitations of use: IMLYGIC has not been shown to improve overall survival or have an effect on visceral metastases. (1)

DOSAGE AND ADMINISTRATION

- Administer IMLYGIC by injection into cutaneous, subcutaneous, and/or nodal lesions. (2.1)
- Recommended starting dose is up to a maximum of 4 mL of IMLYGIC at a concentration of 10^6 (1 million) plaque-forming units (PFU) per mL. Subsequent doses should be administered up to 4 mL of IMLYGIC at a concentration of 10^8 (100 million) PFU per mL. (2.1)

DOSAGE FORMS AND STRENGTHS

Injection: 10^6 (1 million) PFU per mL, 10^8 (100 million) PFU per mL in single-dose vials (3)

CONTRAINDICATIONS

- Immunocompromised Patients (4.1)
- Pregnant Patients (4.2)

WARNINGS AND PRECAUTIONS

- Accidental Exposure to IMLYGIC: Accidental exposure may lead to transmission of IMLYGIC and herpetic infection. Healthcare providers and close contacts should avoid direct contact with injected lesions, dressings, or body fluids of treated patients.
Accidental Exposure to IMLYGIC continued: Healthcare providers who are immunocompromised or pregnant should not prepare or administer IMLYGIC. If accidental exposure occurs, exposed individuals should clean the affected area. (5.1)
- Herpetic Infection: Patients who develop herpetic infections should be advised to follow standard hygienic practices to prevent viral transmission. (5.2)
- Injection Site Complications: Consider the risks and benefits before continuing IMLYGIC treatment if persistent infection or delayed healing develops. (5.3)
- Immune-Mediated Events: Consider the risks and benefits of IMLYGIC before initiating treatment in patients who have underlying autoimmune disease or before continuing treatment in patients who develop immune-mediated events. (5.4)
- Plasmacytoma at the Injection Site: Consider the risks and benefits in patients with multiple myeloma or in whom plasmacytoma develops during treatment. (5.5)
- Obstructive Airway Disorder: Use caution when injecting lesions close to major airways. (5.6)
- Hepatic Hemorrhage from Transcutaneous Intrahepatic Route of Administration: IMLYGIC is not indicated for treatment via transcutaneous intrahepatic route of administration. (5.7)

ADVERSE REACTIONS

The most commonly reported adverse drug reactions ($\geq 25\%$) in IMLYGIC-treated patients were fatigue, chills, pyrexia, nausea, influenza-like illness, and injection site pain. (6)

To report SUSPECTED ADVERSE REACTIONS, contact Amgen at 1-855-IMLYGIC (1-855-465-9442) or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch.

----- **USE IN SPECIFIC POPULATIONS** -----

- Pregnancy: Women of childbearing potential should be advised to use an effective method of contraception to prevent pregnancy during treatment with IMLYGIC. (8.1)
- Lactation: Discontinue drug or nursing. (8.2)

See 17 for PATIENT COUNSELING INFORMATION and Medication Guide.

Revised: 11/2024

FULL PRESCRIBING INFORMATION: CONTENTS*

1 INDICATIONS AND USAGE
2 DOSAGE AND ADMINISTRATION

- 2.1 Dose
- 2.2 Preparation and Handling
- 2.3 Administration

3 DOSAGE FORMS AND STRENGTHS

4 CONTRAINDICATIONS

- 4.1 Immunocompromised Patients
- 4.2 Pregnant Patients

5 WARNINGS AND PRECAUTIONS

- 5.1 Accidental Exposure to IMLYGIC
- 5.2 Herpetic Infection
- 5.3 Injection Site Complications
- 5.4 Immune-Mediated Events
- 5.5 Plasmacytoma at the Injection Site
- 5.6 Obstructive Airway Disorder
- 5.7 Hepatic Hemorrhage from Transcutaneous Intrahepatic Route of Administration

6 ADVERSE REACTIONS

- 6.1 Clinical Trials Experience
- 6.2 Postmarketing Experience

7 DRUG INTERACTIONS

8 USE IN SPECIFIC POPULATIONS

- 8.1 Pregnancy
- 8.2 Lactation
- 8.3 Females and Males of Reproductive Potential
- 8.4 Pediatric Use
- 8.5 Geriatric Use
- 8.6 Renal Impairment
- 8.7 Hepatic Impairment

10 OVERDOSAGE

11 DESCRIPTION

12 CLINICAL PHARMACOLOGY

- 12.1 Mechanism of Action
- 12.3 Pharmacokinetics

13 NONCLINICAL TOXICOLOGY

13.2 Animal Toxicology and/or Pharmacology

14 CLINICAL STUDIES**16 HOW SUPPLIED/STORAGE AND HANDLING****17 PATIENT COUNSELING INFORMATION**

* Sections or subsections omitted from the full prescribing information are not listed.

FULL PRESCRIBING INFORMATION**1 INDICATIONS AND USAGE**

IMLYGIC is a genetically modified oncolytic viral therapy indicated for the local treatment of unresectable cutaneous, subcutaneous, and nodal lesions in patients with melanoma recurrent after initial surgery.

Limitations of use: IMLYGIC has not been shown to improve overall survival or have an effect on visceral metastases.

2 DOSAGE AND ADMINISTRATION

For intralesional injection only. Do not administer intravenously.

2.1 Dose

Administer IMLYGIC by injection into cutaneous, subcutaneous, and/or nodal lesions that are visible, palpable, or detectable by ultrasound guidance.

IMLYGIC is provided in single-dose vials of 1 mL each in two different dose strengths:

- 10^6 (1 million) plaque-forming units (PFU) per mL (light green cap) – for initial dose only
- 10^8 (100 million) PFU per mL (royal blue cap) – for all subsequent doses

Recommended Dose and Schedule

The total injection volume for each treatment visit should not exceed 4 mL for all injected lesions combined. It may not be possible to inject all lesions at each treatment visit or over the full course of treatment. Previously injected and/or uninjected lesion(s) may be injected at subsequent treatment visits. The initial recommended dose is up to 4 mL of IMLYGIC at a concentration of 10^6 (1 million) PFU per mL. The recommended dose for subsequent administrations is up to 4 mL of IMLYGIC at a concentration of 10^8 (100 million) PFU per mL. The recommended dosing schedule for IMLYGIC is shown in Table 1.

Table 1. Recommended Dose and Schedule for IMLYGIC

Treatment	Treatment	Maximum Injection Volume per	Dose	Prioritization of Lesions to be
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Treatment	Interval	Treatment Visit (all lesions combined)	Strength	Injected
Initial	-	4 mL	10^6 (1 million) PFU per mL	<ul style="list-style-type: none"> Inject largest lesion(s) first. Prioritize injection of remaining lesion(s) based on lesion size until maximum injection volume is reached or until all injectable lesion(s) have been treated.
Second	3 weeks after initial treatment	4 mL	10^8 (100 million) PFU per mL	<ul style="list-style-type: none"> Inject any new lesion(s) (lesions that have developed since initial treatment) first. Prioritize injection of remaining lesion(s) based on lesion size until maximum injection volume is reached or until all injectable lesion(s) have been treated.
All subsequent treatments (including reinitiation)	2 weeks after previous treatment	4 mL	10^8 (100 million) PFU per mL	<ul style="list-style-type: none"> Inject any new lesion(s) (lesions that have developed since previous treatment) first. Prioritize injection of remaining lesion(s) based on lesion size until maximum injection volume is reached or until all injectable lesion(s) have been treated.

Dose Volume Determination (per Lesion)

Use Table 2 to determine the volume of IMLYGIC injection for each lesion.

Table 2. Determination of IMLYGIC Injection Volume Based on Lesion Size

Lesion Size (longest dimension)	Injection Volume
> 5 cm	up to 4 mL
> 2.5 cm to 5 cm	up to 2 mL
> 1.5 cm to 2.5 cm	up to 1 mL
> 0.5 cm to 1.5 cm	up to 0.5 mL
≤ 0.5 cm	up to 0.1 mL

When lesions are clustered together, inject them as a single lesion according to Table 2.

Continue IMLYGIC treatment for at least 6 months unless other treatment is required or until there are no injectable lesions to treat.

Reinitiate IMLYGIC treatment if new unresectable cutaneous, subcutaneous, or nodal lesions appear after a complete response.

2.2 Preparation and Handling

Healthcare providers who are immunocompromised or pregnant should not prepare or administer IMLYGIC and should not come into direct contact with the IMLYGIC injection sites, dressings, or body fluids of treated patients [see *Warnings and Precautions* (5.1)].

Avoid accidental exposure to IMLYGIC and follow below instructions for preparation, administration, and handling of IMLYGIC:

- Wear personal protective equipment (protective gown or laboratory coat, safety glasses or face shield, and gloves) while preparing or administering IMLYGIC.
- Avoid accidental exposure to IMLYGIC, especially contact with skin, eyes, and mucous membranes.
 - Cover any exposed wounds before handling.
 - In the event of an accidental occupational exposure (e.g., through a splash to the eyes or mucous membranes), flush with clean water for at least 15 minutes.
 - In the event of exposure to broken skin or needle stick, clean the affected area thoroughly with soap and water and/or a disinfectant.
- Clean all surfaces that may have come in contact with IMLYGIC and treat all IMLYGIC spills with a virucidal agent such as 1% sodium hypochlorite or 70% isopropyl alcohol and blot using absorbent materials.
- Dispose of all materials that may have come in contact with IMLYGIC (e.g., vial, syringe, needle, cotton gauze, gloves, masks, or dressings) as biohazardous waste.
- Advise patients to place used dressings and cleaning materials into a sealed plastic bag and dispose in household waste.

Thawing IMLYGIC Vials

1. Determine the total volume required for injection, up to 4 mL [see *Dosage and Administration* (2.1)].
2. Thaw frozen IMLYGIC vials at room temperature [20°C to 25°C (68°F to 77°F)] until IMLYGIC is liquid. The time to achieve complete vial thaw is expected to be 30 to 70 minutes, depending on the ambient temperature. Do not expose the vial to higher temperatures. Keep the vial in original carton during thawing.
3. Swirl gently. Do NOT shake.
4. After thawing, administer IMLYGIC immediately or store in its original vial and carton, as follows:
 - Thawed IMLYGIC is stable when stored at temperatures of 2°C (36°F) up to 25°C (77°F) protected from light in its original vial and carton for the storage times specified in Table 3. Do not exceed the storage times specified in Table 3.
 - IMLYGIC must not be refrozen once it is thawed. Discard any thawed IMLYGIC in the vial stored longer than the specified times in Table 3.

Table 3. Maximum Storage Time for Thawed Imlygic in Vial

	10^6 (1 million) PFU/mL	10^8 (100 million) PFU/mL
2°C to 8°C (36°F to 46°F)	24 hours	1 week (7 days)
up to 25°C (77°F)	12 hours	24 hours

5. Prepare sterile syringes and needles. A detachable 18-26 G needle or nondetachable 22-26 G staked needle (which minimizes hold up volume) may be used for IMLYGIC withdrawal and a detachable or nondetachable staked needle of 22-26 G may be used for injection. Small unit syringes (e.g., 0.5 mL insulin syringes) are recommended for better injection control.
6. Using aseptic technique, remove the vial cap and withdraw the product from the vial into the syringe(s), noting the total volume. Avoid generating aerosols when loading syringes with product and use a biologic safety cabinet if available.

2.3 Administration

Follow the steps below to administer IMLYGIC to patients:

Pre-Injection

1. Clean the lesion and surrounding areas with an alcohol swab and let dry.
2. Treat the injection site with a topical or local anesthetic agent, if necessary. Do not inject anesthetic agent directly into the lesion. Inject anesthetic agent around the periphery of the lesion.

Injection

1. Inject IMLYGIC intralesionally into cutaneous, subcutaneous, and/or nodal lesions that are visible, palpable, or detectable by ultrasound guidance. Using a single insertion point, inject IMLYGIC along multiple tracks as far as the radial reach of the needle allows within the lesion to achieve even and complete dispersion. Multiple insertion points may be used if a lesion is larger than the radial reach of the needle.

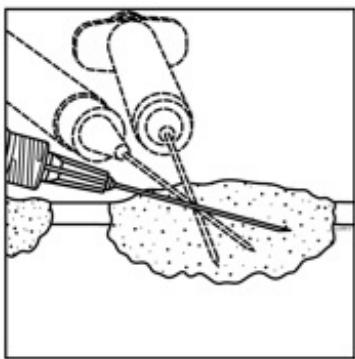


Figure 1: Injection administration for cutaneous lesions

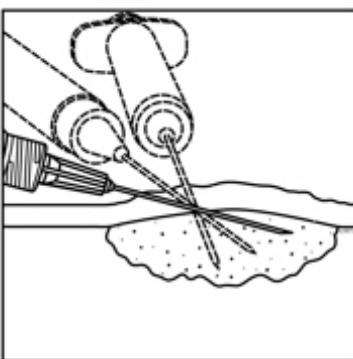


Figure 2: Injection administration for subcutaneous lesions

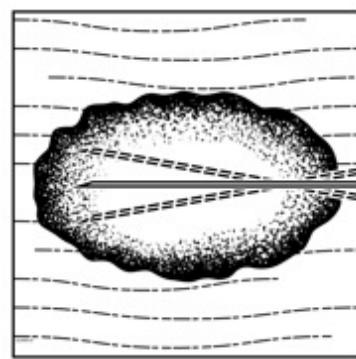


Figure 3: Injection administration for nodal lesions

2. Inject IMLYGIC evenly and completely within the lesion by pulling the needle back without exiting the lesion. Redirect the needle as many times as necessary while injecting the remainder of the dose of IMLYGIC. Continue until the full dose is evenly and completely dispersed.
3. When removing the needle, withdraw it from the lesion slowly to avoid leakage of IMLYGIC at the insertion point.
4. Repeat steps 1-2 under pre-injection and steps 1-3 under injection for other lesions to be injected.
5. Use a new needle any time the needle is completely removed from a lesion and each

time a different lesion is injected.

Post-Injection

1. Apply pressure to the injection site(s) with sterile gauze for at least 30 seconds.
2. Swab the injection site(s) and surrounding area with alcohol.
3. Change gloves and cover the injected lesion(s) with an absorbent pad and dry occlusive dressing.
4. Wipe the exterior of occlusive dressing with alcohol.
5. Advise patients to:
 - Keep the injection site(s) covered for at least the first week after each treatment visit or longer if the injection site is weeping or oozing.
 - Replace the dressing if it falls off.

3 DOSAGE FORMS AND STRENGTHS

- Initial dose only: 10^6 (1 million) PFU per mL solution in 1 mL single-dose vial (light green cap)
- Subsequent doses: 10^8 (100 million) PFU per mL solution in 1 mL single-dose vial (royal blue cap)

4 CONTRAINDICATIONS

4.1 Immunocompromised Patients

IMLYGIC is a live, attenuated herpes simplex virus and may cause life-threatening disseminated herpetic infection in patients who are immunocompromised. Do not administer IMLYGIC to immunocompromised patients, including those with a history of primary or acquired immunodeficient states, leukemia, lymphoma, AIDS or other clinical manifestations of infection with human immunodeficiency viruses, and those on immunosuppressive therapy [see *Nonclinical Toxicology* (13.2)].

4.2 Pregnant Patients

Do not administer IMLYGIC to pregnant patients.

5 WARNINGS AND PRECAUTIONS

5.1 Accidental Exposure to IMLYGIC

Accidental exposure may lead to transmission of IMLYGIC and herpetic infection. Accidental needle stick and flashback to the eyes have been reported in healthcare providers during preparation and administration of IMLYGIC.

Healthcare providers, close contacts (household members, caregivers, sex partners, or persons sharing the same bed), pregnant women, and newborns should avoid direct contact with injected lesions, dressings, or body fluids of treated patients [see *Dosage and Administration* (2.2)]. Healthcare providers who are immunocompromised or pregnant should not prepare or administer IMLYGIC.

Caregivers should wear protective gloves when assisting patients in applying or

changing occlusive dressings and observe safety precautions for disposal of used dressings, gloves, and cleaning materials [see *Dosage and Administration* (2.2)].

In the event of an accidental exposure to IMLYGIC, exposed individuals should clean the affected area thoroughly with soap and water and/or a disinfectant. If signs or symptoms of herpetic infection develop, the exposed individuals should contact their healthcare provider for appropriate treatment [see *Warnings and Precautions* (5.2)].

Patients should avoid touching or scratching injection sites or their occlusive dressings, as doing so could lead to inadvertent transfer of IMLYGIC to other areas of the body.

5.2 Herpetic Infection

Herpetic infections (including but not limited to cold sores and herpetic keratitis) and serious cases of disseminated herpetic infections have been reported in patients treated with IMLYGIC, including fatal disseminated herpetic infection in the immunocompromised patient population [see *Clinical Trials Experience* (6.1) and *Postmarketing Experience* (6.2)]. Immunocompromised patients may be at increased risk of life-threatening disseminated herpetic infection [see *Contraindications* (4.1)].

Patients who develop suspicious herpes-like lesions should follow standard hygienic practices to prevent viral transmission. Patients or close contacts with suspected herpetic infections should also contact their healthcare provider to evaluate the lesions. Suspected herpetic lesions should be reported to Amgen at 1-855-IMLYGIC (1-855-465-9442); patients or close contacts have the option of follow-up testing for further characterization of the infection.

IMLYGIC is sensitive to acyclovir. Acyclovir or other antiviral agents may interfere with the effectiveness of IMLYGIC. Therefore, consider the risks and benefits of IMLYGIC treatment before administering antiviral agents to manage herpetic infection.

5.3 Injection Site Complications

Necrosis or ulceration of tumor tissue may occur during IMLYGIC treatment. Cellulitis and systemic bacterial infection have been reported in clinical studies. Careful wound care and infection precautions are recommended, particularly if tissue necrosis results in open wounds.

In clinical studies, impaired healing at the injection site has been reported. IMLYGIC may increase the risk of impaired healing in patients with underlying risk factors (e.g., previous radiation at the injection site or lesions in poorly vascularized areas). One patient had an amputation of a lower extremity 6 months after IMLYGIC injection due to an infected non-healing wound. This wound area had been treated with surgery and radiation prior to IMLYGIC treatment and had previous wound complications.

If there is persistent infection or delayed healing of the injection site(s), consider the risks and benefits of IMLYGIC before continuing treatment with IMLYGIC.

5.4 Immune-Mediated Events

IMLYGIC may result in immune-mediated events. In clinical studies, immune-mediated events, including glomerulonephritis, vasculitis, pneumonitis, worsening psoriasis, and vitiligo have been reported in patients treated with IMLYGIC.

Consider the risks and benefits of IMLYGIC before initiating treatment in patients who

have underlying autoimmune disease or before continuing treatment in patients who develop immune-mediated events.

5.5 Plasmacytoma at the Injection Site

In a clinical study, a plasmacytoma has been reported in proximity to the injection site after administration of IMLYGIC in a patient with smoldering multiple myeloma.

Consider the risks and benefits of IMLYGIC in patients with multiple myeloma or in whom plasmacytoma develops during treatment.

5.6 Obstructive Airway Disorder

Obstructive airway disorder has been reported following IMLYGIC treatment. Use caution when injecting lesions close to major airways.

5.7 Hepatic Hemorrhage from Transcutaneous Intrahepatic Route of Administration

IMLYGIC is not indicated for transcutaneous intrahepatic route of administration. In clinical studies, cases of hepatic hemorrhage resulting in hospitalization and death have been reported in patients receiving transcutaneous intrahepatic IMLYGIC injections.

6 ADVERSE REACTIONS

The most commonly reported adverse drug reactions ($\geq 25\%$) in IMLYGIC-treated patients were fatigue, chills, pyrexia, nausea, influenza-like illness, and injection site pain.

The following adverse reactions are discussed in greater detail in another section of the label:

- Herpetic Infection [see *Warnings and Precautions (5.2)*]
- Injection Site Complications [see *Warnings and Precautions (5.3)*]

6.1 Clinical Trials Experience

Because clinical trials are conducted under widely varying conditions, adverse reaction rates observed in the clinical trials of a drug cannot be directly compared to rates in the clinical trials of another drug and may not reflect the rates observed in practice.

The safety of IMLYGIC was evaluated in 419 patients who received at least 1 dose of either IMLYGIC ($n = 292$) or subcutaneously administered granulocyte-macrophage colony-stimulating factor (GM-CSF) ($n = 127$) in an open-label, randomized clinical study of patients with stage IIIB, IIIC, and IV melanoma that was not considered to be surgically resectable [see *Clinical Studies (14)*]. The median duration of exposure to IMLYGIC was 23 weeks (5.3 months). Twenty-six patients were exposed to IMLYGIC for at least 1 year.

Most adverse reactions reported were mild or moderate in severity and generally resolved within 72 hours. The most common grade 3 or higher adverse reaction was cellulitis [see *Warnings and Precautions (5.3)*].

Pyrexia, chills, and influenza-like illness can occur any time during IMLYGIC treatment but were more frequent during the first 3 months of treatment.

Table 4 below lists adverse reactions with a 5% or greater incidence in the IMLYGIC arm compared to the GM-CSF arm in the clinical study [see *Clinical Studies* (14)].

Table 4. Adverse Reactions Reported with At Least a 5% Greater Incidence in Patients Treated with IMLYGIC Compared to GM-CSF

Adverse Reactions	IMLYGIC (n = 292)		GM-CSF (n = 127)	
	Any Grade n (%)	Grade 3 n (%)	Any Grade n (%)	Grade 3 n (%)
General disorders and administration site conditions				
Fatigue	147 (50.3)	6 (2.1)	46 (36.2)	1 (< 1)
Chills	142 (48.6)		11 (8.7)	
Pyrexia	125 (42.8)		11 (8.7)	
Influenza-like illness	89 (30.5)	2 (< 1)	19 (15.0)	
Injection site pain	81 (27.7)	2 (< 1)	8 (6.3)	
Gastrointestinal disorders				
Nausea	104 (35.6)	1 (< 1)	25 (19.7)	
Vomiting	62 (21.2)	5 (1.7)	12 (9.5)	
Diarrhea	55 (18.8)	1 (< 1)	14 (11.0)	
Constipation	34 (11.6)		8 (6.3)	1 (< 1)
Abdominal pain	26 (8.9)	2 (< 1)	3 (2.4)	
Musculoskeletal and connective tissue disorders				
Myalgia	51 (17.5)	1 (< 1)	7 (5.5)	
Arthralgia	50 (17.1)	2 (< 1)	11 (8.7)	
Pain in extremity	48 (16.4)	4 (1.4)	12 (9.5)	1 (< 1)
Nervous system disorders				
Headache	55 (18.8)	2 (< 1)	12 (9.5)	
Dizziness	28 (9.6)		4 (3.2)	
Respiratory, thoracic, and mediastinal disorders				
Oropharyngeal pain	17 (5.8)		1 (< 1)	
Investigations				
Weight decreased	17 (5.8)	1 (< 1)	1 (< 1)	

Other adverse reactions associated with IMLYGIC in the open-label, randomized study include rash, dermatitis, glomerulonephritis, vitiligo, worsening psoriasis, cellulitis, pneumonitis, vasculitis, herpetic keratitis, obstructive airway disorder, plasmacytoma at the injection site, deep vein thrombosis, and oral herpes.

6.2 Postmarketing Experience

The following adverse reactions have been identified during postapproval use of IMLYGIC. Because these reactions are reported voluntarily from a population of uncertain size, it is not always possible to reliably estimate their frequency or establish a causal relationship to drug exposure.

- Herpetic infections including disseminated herpetic infections [see *Warnings and Precautions* (5.2)]. Serious including fatal disseminated herpetic infections in

immunocompromised patient population [see *Contraindications* (4.1) and *Warnings and Precautions* (5.2)].

7 DRUG INTERACTIONS

IMLYGIC is sensitive to acyclovir. Acyclovir or other antiherpetic viral agents may interfere with the effectiveness of IMLYGIC. No drug interaction studies have been conducted with IMLYGIC.

8 USE IN SPECIFIC POPULATIONS

8.1 Pregnancy

Risk Summary

Adequate and well-controlled studies with IMLYGIC have not been conducted in pregnant women. No effects on embryo-fetal development have been observed in a study conducted in pregnant mice. The design of the study limits application of the animal data to humans [see *Data*].

In the U.S. general population, the estimated background risk of major birth defects and miscarriage in clinically recognized pregnancies is 2-4% and 15-20%, respectively.

Clinical Considerations

If the patient becomes pregnant while taking IMLYGIC, the patient should be apprised of the potential hazards to the fetus and neonate. Women of childbearing potential should be advised to use an effective method of contraception to prevent pregnancy during treatment with IMLYGIC.

If a pregnant woman has an infection with wild-type Herpes Simplex Virus Type 1 (HSV-1) (primary or reactivation), there is potential for the virus to cross the placental barrier and also a risk of transmission during birth due to viral shedding. Infections with wild-type HSV-1 have been associated with serious adverse effects, including multi-organ failure and death, if a fetus or neonate contracts the wild-type herpes infection. While there are no clinical data to date on IMLYGIC infections in pregnant women, there could be a risk to the fetus or neonate if IMLYGIC were to act in the same manner.

Data

Animal Data

No effects on embryo-fetal development were observed when IMLYGIC was intravenously administered during organogenesis to immunocompetent pregnant mice at doses up to 4×10^8 (400 million) PFU per kg (60-fold higher, on a PFU per kg basis, compared to the maximum clinical dose). Levels of IMLYGIC DNA in pooled fetal blood were at or below the assay detection level. Study design limitations included: 1) administration of IMLYGIC expressing human granulocyte-macrophage colony-stimulating factor (huGM-CSF), which is not biologically active in mice; 2) unknown transplacental kinetics of IMLYGIC following intravenous administration in pregnant mice; and 3) unknown significance of IMLYGIC dose extrapolation from animal to human based on body weight.

8.2 Lactation

Risk Summary

There is no information regarding the presence of IMLYGIC in human milk, the effects on the breastfed infant, or the effects on milk production. The developmental and health benefits of breastfeeding should be considered along with the mother's clinical need for IMLYGIC and any potential adverse effects on the breastfed infant from IMLYGIC or from the underlying maternal condition.

Clinical Considerations

Because medicinal products can be found in human milk, a decision should be made whether to discontinue nursing or to discontinue IMLYGIC while nursing.

8.3 Females and Males of Reproductive Potential

No nonclinical or clinical studies were performed to evaluate the effect of IMLYGIC on fertility.

8.4 Pediatric Use

Safety and effectiveness of IMLYGIC have not been established in pediatric patients.

8.5 Geriatric Use

In clinical studies, no overall differences in safety or efficacy were observed between geriatric patients (≥ 65 years old) and younger patients.

8.6 Renal Impairment

No clinical studies have been conducted to evaluate the effect of renal impairment on the pharmacokinetics of IMLYGIC.

8.7 Hepatic Impairment

No clinical studies have been conducted to evaluate the effect of hepatic impairment on the pharmacokinetics of IMLYGIC.

10 OVERDOSAGE

There is no clinical experience with an overdose with IMLYGIC. Doses up to 4 mL at dose strength of 10^8 (100 million) PFU per mL every 2 weeks (maximum cumulative dose of 222.5×10^8 PFU) have been administered in clinical studies, with no evidence of dose-limiting toxicity. The maximum dose of IMLYGIC that can be safely administered has not been determined. In the event of a suspected overdose, the patient should be treated symptomatically and supportive measures instituted as required [see *Warnings and Precautions (5)*].

11 DESCRIPTION

IMLYGIC (talimogene laherparepvec) is a sterile suspension for intralesional injection. IMLYGIC is a live, attenuated HSV-1 that has been genetically

modified to express huGM-CSF. The parental virus for IMLYGIC was a primary isolate, which was subsequently altered using recombinant methods to result in gene deletions and insertions.

Each vial contains 1 mL deliverable volume of IMLYGIC at either 1×10^6 (1 million) PFU per mL or 1×10^8 (100 million) PFU per mL concentrations and the following excipients: di-sodium hydrogen phosphate dihydrate (15.4 mg), sodium dihydrogen phosphate dihydrate (2.44 mg), sodium chloride (8.5 mg), myo-inositol (40 mg), sorbitol (20 mg), and water for injection.

The 10^6 (1 million) PFU per mL vial of IMLYGIC contains a clear to semi-translucent liquid following thaw from its frozen state. The 10^8 (100 million) PFU per mL vial of IMLYGIC contains a semi-translucent to opaque liquid following thaw from its frozen state. The liquid in each vial may contain white, visible, variously shaped, virus-containing particles.

Each vial of IMLYGIC may also contain residual components of VERO cells including DNA and protein and trace quantities of fetal bovine serum.

The product contains no preservative.

12 CLINICAL PHARMACOLOGY

12.1 Mechanism of Action

IMLYGIC has been genetically modified to replicate within tumors and to produce the immune stimulatory protein GM-CSF. IMLYGIC causes lysis of tumors, followed by release of tumor-derived antigens, which together with virally derived GM-CSF may promote an antitumor immune response. However, the exact mechanism of action is unknown.

12.3 Pharmacokinetics

Biodistribution (within the body) and Viral Shedding (excretion/secretion)

IMLYGIC viral DNA levels in various tissues and secretions were determined using a quantitative polymerase chain reaction (qPCR) assay. Infectious IMLYGIC at the injection sites and at some potential herpetic lesions was also quantified using viral infectivity assays.

Nonclinical data

Following repeat intratumoral administration in mice, IMLYGIC DNA was primarily detected in the tumor, blood, spleen, lymph node, liver, heart, and kidney. IMLYGIC DNA was not detected in bone marrow, eyes, lachrymal glands, nasal mucosa, or feces. The highest level of IMLYGIC DNA was found in the injected tumor. IMLYGIC DNA was found in the injected tumor through 84 days and in blood samples through 14 days after the last administration of IMLYGIC.

Clinical data

The biodistribution and shedding of intralesionally administered IMLYGIC were investigated in a clinical study that measured IMLYGIC DNA in blood, urine, injection site, occlusive dressings, oral mucosa, anogenital area, and suspected herpetic lesions. Occlusive dressings samples were collected during treatment. Blood and urine samples

were collected during treatment and for up to 30 days after the end of treatment. Injection site, oral mucosa, and anogenital area samples were collected during treatment and for up to 60 days after the end of treatment. Suspected herpetic lesion samples were collected any time a patient experienced lesions of suspected herpetic origin. If the qPCR testing for IMLYGIC DNA was positive, then a TCID₅₀ assay was performed to measure viral infectivity. In the 60 patients with melanoma who received IMLYGIC intralesional injection at a dose and schedule same as the clinical study [see *Clinical Studies (14)*], data indicate that IMLYGIC DNA was present in all sites during the study (see Table 5).

Table 5. Patients with Detectable DNA During Treatment

Body fluid/Site	Patients with detectable DNA during treatment (n = 60)
Injection site	60 (100%)
Blood	59 (98%)
Urine	19 (32%)
Injection site	60 (100%)
Occlusive dressing	48 (80%)
Oral mucosa	8 (13%)
Anogenital area	5 (out of 26)*

* For the anogenital area, 26 patients were tested.

The proportion of samples and subjects with IMLYGIC DNA was highest during cycle 2 of treatment for the blood, urine, injection site, and occlusive dressings; highest in cycle 1 of treatment for the oral mucosa; and highest in cycles 1 and 2 for the anogenital area. Among patients with detectable IMLYGIC DNA in the blood, urine, oral mucosa and anogenital area, no samples had detectable IMLYGIC DNA 30 days after the end of treatment. For patients with detectable DNA in injected lesions, no samples had detectable IMLYGIC DNA 60 days after the end of treatment.

Overall 3 of 19 patients with lesions of suspected herpetic origin had IMLYGIC DNA present at any time during the study. Of the three subjects who had lesions of suspected herpetic origin that had IMLYGIC DNA detected during the study, none of the suspected lesions were injected with IMLYGIC. None of the subjects with the lesions of the suspected herpetic origin with IMLYGIC DNA had disseminated lesions or received antiviral therapy and these events were non-serious, lasting from 16 days to 4 months. The data suggest that IMLYGIC may cause herpetic lesions in treated patients.

In addition to the presence of IMLYGIC DNA, viral activity was measured in samples that were positive for IMLYGIC DNA from the injection site, occlusive dressings, oral mucosa, anogenital area, and suspected herpetic lesions. No viral activity was detected in samples of the occlusive dressings, oral mucosa, anogenital area, and suspected herpetic lesions. Infectious IMLYGIC virus was detected at the site of injection in 7 (11%) patients at multiple time points in the study; no samples were positive for viral infectivity after cycle 2 or after the end of treatment.

13 NONCLINICAL TOXICOLOGY

13.2 Animal Toxicology and/or Pharmacology

Repeated intratumoral administration at 2×10^8 (200 million) PFU per kg (30-fold maximum proposed clinical dose, extrapolated based on body weight) did not demonstrate any adverse effects in immunocompetent mice. Severe combined immunodeficient (SCID) mice administered repeat intratumoral injections of IMLYGIC at a dose of 30-fold maximum proposed clinical dose developed systemic viral infection (viral inclusion bodies or necrosis in enteric neurons in the gastrointestinal tract, adrenal gland, skin, pancreatic islet cells, eye, pineal gland, and brain).

14 CLINICAL STUDIES

The safety and efficacy of intralesional injections of IMLYGIC compared with subcutaneously administered GM-CSF was evaluated in a multicenter, open-label, randomized clinical study in patients with stage IIIB, IIIC, and IV melanoma that was considered to be not surgically resectable. IMLYGIC was injected into cutaneous, subcutaneous, or nodal melanoma lesions and was not injected into visceral lesions. Previous systemic treatment for melanoma was allowed. Patients with active cerebral metastases, bony metastases, extensive visceral disease, primary ocular or mucosal melanoma, evidence of immunosuppression, or receiving treatment with a systemic antiherpetic agent were excluded from the study.

The study included 250 (57%) men and 186 (43%) women. The mean age was 63 (range: 22 to 94) years. Most patients (98%) were white. Seventy percent (70%) of patients had baseline Eastern Cooperative Oncology Group (ECOG) performance status of zero. Seventy percent (70%) of patients had stage IV disease (27% M1a; 21% M1b; and 22% M1c), and 30% had stage III disease. Fifty-three percent (53%) of patients had received prior therapy for melanoma (other than or in addition to surgery, adjuvant therapy, or radiation), and 58% were seropositive for wild-type HSV-1 at baseline.

A total of 436 patients were randomized to receive either IMLYGIC ($n = 295$) or GM-CSF ($n = 141$). IMLYGIC was administered by intralesional injection at an initial concentration of 10^6 (1 million) PFU per mL on Day 1, followed by a concentration of 10^8 (100 million) PFU per mL on Day 21 and every 2 weeks thereafter, at a dose of up to 4 mL per visit. GM-CSF was administered subcutaneously in 28-day cycles, i.e., 125 µg/m² daily for 14 days followed by 14 days without GM-CSF administration.

Patients were to be treated for at least 6 months or until there were no injectable lesions. During this period, treatment could continue despite an increase in size in existing lesion(s) and/or development of new lesion(s), unless the patient developed intolerable toxicity or the investigator believed that it was in the best interest of the patient to stop treatment or to be given other therapy for melanoma. After 6 months of treatment, patients were to continue treatment until clinically relevant disease progression (i.e., disease progression associated with a decline in performance status and/or alternative therapy was required in the opinion of the investigator), up to 12 months. Patients experiencing a response at 12 months after the start of treatment could continue treatment for up to an additional 6 months, unless there were no remaining injectable lesions or disease progression. All patients were to be followed for survival status for at least 36 months.

The major efficacy outcome was durable response rate (DRR), defined as the percent of patients with complete response (CR) or partial response (PR) maintained continuously

for a minimum of 6 months. Tumor responses were determined according to World Health Organization (WHO) response criteria modified to allow patients who developed new lesions or disease progression of existing lesions to continue the treatment and be evaluated later for tumor response.

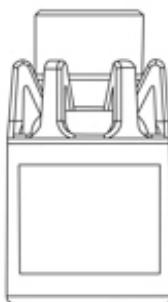
The DRR was 16.3% in the IMLYGIC arm and 2.1% in the GM-CSF arm in the overall study population. The unadjusted relative risk was 7.6 (95% CI: 2.4, 24.1), with a p-value < 0.0001. The median time to response was 4.1 (range: 1.2 to 16.7) months in the IMLYGIC arm.

There was no statistically significant difference in overall survival (OS) between the IMLYGIC and the GM-CSF arms. The median OS in the overall study population was 22.9 months in the IMLYGIC arm and 19.0 months in the GM-CSF arm ($p = 0.116$).

16 HOW SUPPLIED/STORAGE AND HANDLING

How Supplied

- IMLYGIC is provided as a sterile frozen suspension in a single-dose, cyclic olefin polymer (COP) plastic resin vial with a chlorobutyl elastomer stopper, aluminum seal, and polypropylene cap in two different presentations:
 - Figure 4: Single-dose vial permanently inserted into a clear copolyester plastic sleeve



OR

- Figure 5: Single-dose vial without a clear plastic sleeve



- Each vial contains a retrievable minimal volume of 1 mL.
- The vial cap is color coded:
 - 10^6 (1 million) PFU per mL is light green (NDC 55513-078-01).
 - 10^8 (100 million) PFU per mL is royal blue (NDC 55513-079-01).

Storage and Handling

- Store and transport IMLYGIC at –90°C to –70°C (–130°F to –94°F).
- Protect IMLYGIC from light.
- Store IMLYGIC in the carton until use.
- Thaw IMLYGIC immediately prior to administration [see *Dosage and Administration* (2.2)].
- Do not draw IMLYGIC into a syringe until immediately prior to administration [see *Dosage and Administration* (2.2)].

17 PATIENT COUNSELING INFORMATION

Advise patients and/or close contacts to:

- **Read the FDA-approved patient labeling (Medication Guide).**
- Follow instructions below to prevent viral transmission [see *Warnings and Precautions* (5.1)]:
 - Avoid direct contact with injection sites, dressings, or body fluids of patients.
 - Wear gloves when changing dressing.
 - Avoid touching or scratching injection sites.
 - Keep injection sites covered for at least the first week after each treatment visit or longer if the injection site is weeping or oozing. Replace dressing if it falls off.
 - Dispose of used dressings and cleaning materials in household waste in a sealed plastic bag.
- Female patients of childbearing potential should use an effective method of contraception to prevent pregnancy during treatment with IMLYGIC [see *Contraindications* (4.2) and *Use in Specific Populations* (8.1)].
- Close contacts who are pregnant or immunocompromised should not change dressings or clean injection sites [see *Warnings and Precautions* (5.1)].
- In case of accidental exposure to IMLYGIC, clean the exposed area with soap and water and/or a disinfectant. Patients or close contacts with suspected herpetic infections should contact their healthcare provider to evaluate the lesions. Suspected herpetic lesions should be reported to Amgen at 1-855-IMLYGIC (1-855-465-9442); patients or close contacts have the option of follow-up testing for further characterization of the infection [see *Warnings and Precautions* (5.1) and (5.2)].

IMLYGIC® (talimogene laherparepvec)

Manufactured by:

BioVex LLC, a subsidiary of Amgen Inc.
One Amgen Center Drive
Thousand Oaks, California 91320-1799

Patent: <http://pat.amgen.com/Imlygic/>

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v9

Medication Guide

IMLYGIC® (imm-LY-jik)

(talimogene laherparepvec)

Read the Medication Guide before you start treatment with IMLYGIC and before each IMLYGIC treatment. There may be new information. This Medication Guide does not tell you everything about IMLYGIC. Talk with your healthcare provider if you have any questions about treatment with IMLYGIC.

What is IMLYGIC?

IMLYGIC is a prescription medicine used to treat a type of cancer called melanoma when it is on your skin or in your lymph glands. IMLYGIC is a weakened form of Herpes Simplex Virus Type 1, which is commonly called the cold sore virus. Your healthcare provider will inject IMLYGIC directly into your tumor(s).

IMLYGIC may not help you live longer and may not shrink cancer in your organs (for example, lung or liver).

Who should not get IMLYGIC?

You should not get IMLYGIC if you are pregnant or have a weakened immune system (for example, an immune deficiency, blood or bone marrow cancer, steroid use, or HIV/AIDS).

What should I tell my healthcare provider before I get IMLYGIC?

Before getting IMLYGIC, tell your healthcare provider if you:

- Are taking steroids or other medicines that suppress your immune system.
- Are taking antiviral medicines to treat or prevent herpes, such as acyclovir.
- Have or ever had medical conditions such as:
 - HIV infection or AIDS.
 - Blood or bone marrow cancer.
 - Autoimmune disease.
 - Other medical conditions that can weaken your immune system.
- Have close contact with someone who has a weakened immune system or is pregnant.
- Are pregnant or plan to become pregnant.
 - IMLYGIC may harm your unborn baby.
 - You should not become pregnant during treatment with IMLYGIC.
 - Talk to your healthcare provider about effective birth control methods.
- Are breastfeeding or plan to breastfeed.

Tell your healthcare provider about all the medicines you take, including prescription and over-the-counter medicines, vitamins, and herbal supplements.

IMLYGIC may affect the way other medicines work and other medicines may affect how IMLYGIC works.

How is IMLYGIC given?

Your healthcare provider will inject IMLYGIC directly into your tumor(s) with a needle and syringe. You will get a second treatment 3 weeks after the first treatment. After that, you will get treatments every 2 weeks for as long as you have tumor(s). You can get treated for 6 months or longer.

Your healthcare provider will decide which tumor(s) to inject and may not inject every one.

It is important to care for the treatment sites properly so that IMLYGIC does not spread

to other people. Your healthcare provider will show you how to do this.

What should I avoid while getting IMLYGIC?

IMLYGIC virus can spread to other areas of your body or to your close contacts (household members, caregivers, sex partners, or persons sharing the same bed).

Do the following to avoid spreading IMLYGIC to other areas of your body or to your close contacts:

- Avoid direct contact between your treatment sites, dressings, or body fluids and close contacts (for example, use condoms when engaging in sexual activity, avoid kissing close contacts if either has an open mouth sore).
- Wear gloves while putting on or changing your dressings.
- Keep treatment sites covered with airtight and watertight dressings for at least 1 week after each treatment (or longer if the treatment site is weeping or oozing).
- If the dressing comes loose or falls off, replace it right away with a clean dressing.
- Place all used dressings and cleaning materials in a sealed plastic bag and throw them away in the garbage.
- Do not touch or scratch the treatment sites.

What are possible side effects of IMLYGIC?

The most common side effects of IMLYGIC include:

- Tiredness
- Chills
- Fever
- Nausea
- Flu-like symptoms
- Pain at treatment site

Tell your doctor right away if you get any of the signs and symptoms of herpes infection, including but not limited to:

- Pain, burning, tingling, or blister formation around the mouth, genitals, or any part of the body, near or far from the injection site
- Eye pain, light sensitivity, discharge from the eyes, or blurry vision
- Weakness in arms or legs
- Extreme drowsiness (feeling sleepy)
- Mental confusion

If you think you have a herpes infection, inform your healthcare provider. Life threatening-herpes infection and herpes infection spreading to any part of the body far from the injection site (disseminated herpetic infection) may occur in patients treated with IMLYGIC. If you have any new or worsening symptoms, call your healthcare provider right away. You or your healthcare provider should call Amgen at 1-855-IMLYGIC (1-855-465-9442) for follow-up testing if needed.

These are not all the possible side effects of IMLYGIC. Your healthcare provider can give you more detailed information. Tell your healthcare provider if you have any side effects that bother you or that do not go away. You may report side effects to FDA at 1-800-FDA-1088.

What are the ingredients in IMLYGIC?

Active ingredient: talimogene laherparepvec

Inactive ingredients: di-sodium hydrogen phosphate dihydrate, sodium dihydrogen phosphate dihydrate, sodium chloride, myo-inositol, sorbitol, and water for injection

General information about the safe and effective use of IMLYGIC

This Medication Guide summarizes the most important information about IMLYGIC. Medicines are sometimes prescribed for purposes other than those listed in a Medication Guide. If you would like more information, talk with your healthcare provider. You can ask your healthcare provider for information about IMLYGIC that was written for healthcare professionals.

This Medication Guide has been approved by the U.S. Food and Drug Administration.

IMLYGIC® (talimogene laherparepvec)

Manufactured by:

BioVex LLC, a subsidiary of Amgen Inc.
One Amgen Center Drive
Thousand Oaks, California 91320-1799

Patent: <http://pat.amgen.com/lmlygic/>

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Revised: 11/2024

V5

PRINCIPAL DISPLAY PANEL - 10⁶ PFU/mL Vial Carton

10⁶ PFU/mL Single-Dose Vial

NDC 55513-078-01

AMGEN

talimogene laherparepvec
IMLYGIC®

10⁶ Plaque Forming Units (PFU)
per mL

10⁶
PFU/mL

For Intralesional Injection Only

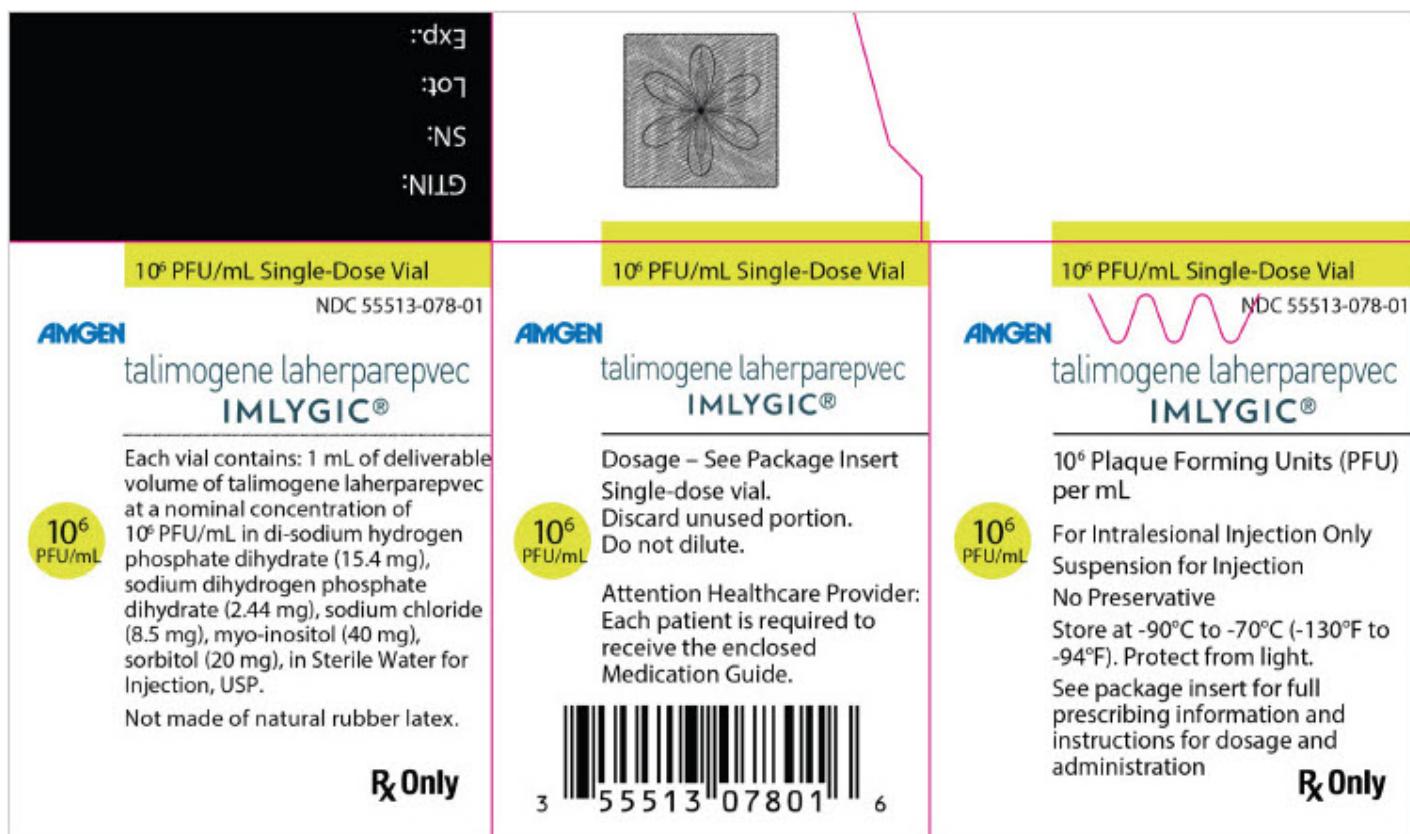
Suspension for Injection

No Preservative

Store at -90°C to -70°C (-130°F to
-94°F). Protect from light.

See package insert for full
prescribing information and
instructions for dosage and
administration

Rx Only



PRINCIPAL DISPLAY PANEL - 10⁸ PFU/mL Vial Carton

10⁸ PFU/mL Single-Dose Vial

NDC 55513-079-01

AMGEN

talimogene laherparepvec
IMLYGIC®

10⁸ Plaque Forming Units (PFU)
per mL

10⁸
PFU/mL

For Intralesional Injection Only

Suspension for Injection

No Preservative

Store at -90°C to -70°C (-130°F to -94°F). Protect from light.

See package insert for full prescribing information and instructions for dosage and administration

Rx Only

<p>Exp: Lot: SN: GTIN:</p> <p>10⁸ PFU/mL Single-Dose Vial NDC 55513-079-01</p> <p>AMGEN talimogene laherparepvec IMLYGIC®</p> <p>Each vial contains: 1 mL of deliverable volume of talimogene laherparepvec at a nominal concentration of 10⁸ PFU/mL in di-sodium hydrogen phosphate dihydrate (15.4 mg), sodium dihydrogen phosphate dihydrate (2.44 mg), sodium chloride (8.5 mg), myo-inositol (40 mg), sorbitol (20 mg), in Sterile Water for Injection, USP.</p> <p>Not made of natural rubber latex.</p> <p>10⁸ PFU/ml</p> <p>Rx Only</p>	 <p>10⁸ PFU/mL Single-Dose Vial</p> <p>AMGEN talimogene laherparepvec IMLYGIC®</p> <p>Dosage – See Package Insert Single-dose vial. Discard unused portion. Do not dilute.</p> <p>Attention Healthcare Provider: Each patient is required to receive the enclosed Medication Guide.</p> <p>10⁸ PFU/ml</p> <p>Rx Only</p>	<p>10⁸ PFU/mL Single-Dose Vial NDC 55513-079-01</p> <p>AMGEN talimogene laherparepvec IMLYGIC®</p> <p>10⁸ Plaque Forming Units (PFU) per mL</p> <p>For Intralesional Injection Only Suspension for Injection No Preservative Store at -90°C to -70°C (-130°F to -94°F). Protect from light. See package insert for full prescribing information and instructions for dosage and administration</p> <p>10⁸ PFU/ml</p> <p>Rx Only</p>
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IMLYGIC

talimogene laherparepvec injection, suspension

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:55513-078
Route of Administration	INTRALESIONAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
TALIMOGENE LAHERPAREPVEC (UNII: 07730V90L6) (TALIMOGENE LAHERPAREPVEC - UNII:07730V90L6)	TALIMOGENE LAHERPAREPVEC	1000000 [PFU] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM PHOSPHATE, DIBASIC, DIHYDRATE (UNII: 9425516E2T)	15.4 mg in 1 mL
SODIUM PHOSPHATE, MONOBASIC, DIHYDRATE (UNII: 5QWK665956)	2.44 mg in 1 mL
SODIUM CHLORIDE (UNII: 451W47IQ8X)	8.5 mg in 1 mL
INOSITOL (UNII: 4L6452S749)	40 mg in 1 mL
SORBITOL (UNII: 506T60A25R)	20 mg in 1 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55513-078-01	1 in 1 CARTON	11/02/2015	
1		1 mL in 1 VIAL; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA125518	11/02/2015	

IMLYGIC

talimogene laherparepvec injection, suspension

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:55513-079
Route of Administration	INTRALESIONAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
TALIMOGENE LAHERPAREPVEC (UNII: 07730V90L6) (TALIMOGENE LAHERPAREPVEC - UNII:07730V90L6)	TALIMOGENE LAHERPAREPVEC	100000000 [PFU] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM PHOSPHATE, DIBASIC, DIHYDRATE (UNII: 9425516E2T)	15.4 mg in 1 mL
SODIUM PHOSPHATE, MONOBASIC, DIHYDRATE (UNII: 5QWK665956)	2.44 mg in 1 mL
SODIUM CHLORIDE (UNII: 451W47IQ8X)	8.5 mg in 1 mL
INOSITOL (UNII: 4L6452S749)	40 mg in 1 mL
SORBITOL (UNII: 506T60A25R)	20 mg in 1 mL
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55513-079-01	1 in 1 CARTON	11/02/2015	

1	1 mL in 1 VIAL; 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	BLA125518	11/02/2015	

Labeler - Amgen Inc (039976196)

Establishment

Name	Address	ID/FEI	Business Operations
West Services and Solutions LLC		118935952	ANALYSIS(55513-078, 55513-079)

Revised: 4/2025

Amgen Inc