

LEMTRADA REMS Education Program for Prescribers

This education program includes information about:

- > The LEMTRADA REMS Program requirements
- > Serious risks of autoimmune conditions, infusion reactions, and malignancies
- > Counseling and management of your patient


LEMTRADA[®]
alemtuzumab iv^{12mg}

What Is the LEMTRADA REMS Program?

A REMS (Risk Evaluation and Mitigation Strategy) is a strategy to manage known or potential risks associated with a drug, and is required by the FDA to ensure that the benefits of the drug outweigh its risks. Due to serious risks of autoimmune conditions, infusion reactions and malignancy, LEMTRADA is only available through a restricted program called the LEMTRADA REMS Program.

This brochure has been developed as part of the LEMTRADA REMS Program to help educate prescribers about the risks associated with LEMTRADA and how to help mitigate these risks through periodic monitoring for, and prompt identification of, signs and symptoms of these events.

- > **Prescribers** must be enrolled in the LEMTRADA REMS Program to be able to prescribe LEMTRADA.
- > **Pharmacies** must be enrolled in the LEMTRADA REMS Program to be able to dispense LEMTRADA.
- > **Healthcare Facilities** must be enrolled in the LEMTRADA REMS Program to be able to dispense and administer LEMTRADA.
- > **Patients** must be enrolled and authorized in the LEMTRADA REMS Program in order to receive LEMTRADA.

Steps for Prescriber Certification and Enrollment in the LEMTRADA REMS Program

1. Complete the training program, which includes reviewing the following materials:
 - > LEMTRADA Prescribing Information
 - > LEMTRADA REMS Program Overview
 - > LEMTRADA REMS Education Program for Prescribers
2. Successfully complete the 8-question Knowledge Assessment
3. Enroll in the program by completing a LEMTRADA REMS Prescriber Enrollment Form
4. Submit the completed and signed forms to the LEMTRADA REMS Program

The LEMTRADA REMS Program Overview, Knowledge Assessment, LEMTRADA Prescribing Information, and other REMS materials are available online at www.LemtradaREMS.com or by contacting the LEMTRADA REMS Program at 1-855-676-6326.

To enroll in the LEMTRADA REMS Program, call 1-855-676-6326 or go to www.LemtradaREMS.com

Genzyme will send confirmation of a prescriber's enrollment in the LEMTRADA REMS Program, including the prescriber's assigned LEMTRADA REMS Program identification number.

You will not be able to prescribe LEMTRADA without completing your certification in the LEMTRADA REMS Program.

You should understand that if you fail to comply with the Program requirements, you may no longer be able to participate in the Program.

Overview of Important Safety Information

INDICATION

LEMTRADA is indicated for the treatment of patients with relapsing forms of multiple sclerosis (MS). Because of its safety profile, the use of LEMTRADA should generally be reserved for patients who have had an inadequate response to two or more drugs indicated for the treatment of MS.

The Prescribing Information includes a **boxed WARNING** for LEMTRADA.

Please see the Prescribing Information for complete safety information, including boxed WARNING.

SERIOUS RISKS ASSOCIATED WITH LEMTRADA

Autoimmune Conditions

LEMTRADA has been associated with risk of autoimmune conditions, including immune thrombocytopenia, other cytopenias (including neutropenia, hemolytic anemia, and pancytopenia), thyroid disorders and glomerular nephropathies, which may occur many years after treatment and may be serious or life-threatening. Early detection can help to improve the outcomes of patients experiencing these events.

Please review the sections that follow to gain a better understanding of the risks of autoimmune conditions.

Immune Thrombocytopenia (ITP)

Immune thrombocytopenia occurred in 2% of LEMTRADA-treated patients in clinical studies in MS. Immune thrombocytopenia is an autoimmune disorder usually associated with anti-platelet antibodies. Platelet depletion reduces the ability of the blood to clot. Symptoms of ITP could include (but are not limited to) easy bruising, petechiae, spontaneous mucocutaneous bleeding (epistaxis, hemoptysis), and heavier than normal or irregular menstrual bleeding. These clinical signs of ITP may be apparent before serious bleeding develops.

ITP can be a serious condition leading to morbidity and mortality, and may occur several years after dosing. It is important to monitor all patients for ITP as follows:

- > Complete blood counts with differential should be obtained ≤ 30 days prior to initiation of treatment and at monthly intervals thereafter until 48 months after the patient's last infusion of LEMTRADA. After this period of time, testing should be performed based on clinical findings suggestive of ITP.
- > Check the patient for clinical symptoms of ITP.
- > Counsel the patient on the importance of complying with monthly monitoring of their blood and the need to continue for 48 months after their last infusion.
- > Educate the patient on how to recognize ITP-related symptoms, and emphasize the need to remain vigilant for them.
- > If ITP is suspected, appropriate medical intervention should be promptly initiated, including immediate referral to a specialist. Severe or widespread bleeding is life-threatening and demands immediate care.

The potential risk associated with retreatment with LEMTRADA following the occurrence of ITP is unknown.



Potential Clinical Presentations of ITP

Note: These pictures are only a guide in order to show examples of bruises or petechiae. The patient may have a less severe type of bruise or petechiae than these pictures and still have ITP.



This is an example of a leg with petechiae.

Petechiae are small, scattered, “pinprick” spots under the skin that are red, pink or purple.

Petechiae can occur anywhere on the patient’s body, not just the legs.



This is an example of easy or excessive bruising.

This could occur anywhere on the patient’s body.



This is an example of purpura under the tongue.

Purpura could occur on any mucous membrane, including anywhere in the mouth (under the tongue, roof of the mouth, inner cheeks, tongue, gums).

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Other Autoimmune Cytopenias

Autoimmune cytopenias such as neutropenia, hemolytic anemia, and pancytopenia have been infrequently reported in clinical studies in MS. Symptoms of autoimmune hemolytic anemia may include weakness, chest pain, jaundice, dark urine, and tachycardia. Use the monthly CBC results to monitor for cytopenias. If a cytopenia is confirmed, appropriate medical intervention should be promptly initiated.

Glomerular Nephropathies

Glomerular nephropathies, including anti-glomerular basement membrane (anti-GBM) disease, have been reported after treatment with LEMTRADA in multiple sclerosis patients in clinical trials. Cases of anti-GBM disease have been diagnosed up to 40 months after the last dose of LEMTRADA.

Anti-GBM disease is life-threatening if not treated and therefore demands immediate care. Without prompt treatment, patients can rapidly develop renal failure requiring dialysis and/or kidney transplantation, and this may lead to death.

Clinical manifestations of nephropathy may include elevation in serum creatinine, hematuria, and/or proteinuria.

While not observed in clinical trials, alveolar hemorrhage manifested as hemoptysis may occur with anti-GBM disease. Since patients may be asymptomatic, it is important that the monthly tests are conducted.

- > Serum creatinine levels should be obtained ≤ 30 days prior to the first infusion of LEMTRADA and at monthly intervals thereafter until 48 months after the patient's last infusion. After this period of time, testing should be performed based on clinical findings suggestive of nephropathies.
- > Urinalysis with urine cell counts should be obtained ≤ 30 days prior to the first infusion of LEMTRADA and at monthly intervals thereafter until 48 months after the last infusion. After this period of time, testing should be performed based on clinical findings suggestive of nephropathies. In menstruating females, consider the timing of urinalysis to avoid false positives. The observation of clinically significant changes from baseline in serum creatinine, unexplained hematuria, and/or proteinuria should prompt further evaluation for nephropathies, including referral to a specialist.
- > Early detection and treatment of nephropathies may decrease the risk of poor outcomes.
- > Immediate referral to a specialist for further assessment for patients with suspected nephropathy is strongly recommended.

Thyroid Disorders

During clinical trials, autoimmune thyroid disorders including Graves' disease, hyperthyroidism, and hypothyroidism were reported. Autoimmune thyroid disorders occurred in 34% of LEMTRADA-treated patients in clinical studies.

Newly diagnosed thyroid disorders occurred throughout the uncontrolled clinical study follow-up period, more than 7 years after the first LEMTRADA dose. Serious thyroid events occurred in 2% of patients. Most thyroid disorders were managed with conventional medical therapy; some patients required surgical intervention.

It is important to monitor all patients for thyroid disorders as follows:

- > Thyroid function tests such as thyroid-stimulating hormone (TSH) levels should be obtained ≤ 30 days prior to the first infusion of LEMTRADA and then every 3 months thereafter continuing for 48 months following the last infusion. Continue to test thyroid function after 48 months if clinically indicated.
- > Additionally, watch out for signs and symptoms of thyroid disorders, which may include excessive sweating, unexplained weight loss, eye swelling, nervousness and fast heartbeat (hyperthyroidism), or unexplained weight gain, feeling cold, worsening tiredness, and newly occurring constipation (hypothyroidism).

- > Thyroid disease poses special risks in women who become pregnant. Untreated thyroid disease can cause harm to the unborn and newborn baby. Special caution should be taken for pregnant women with Graves' disease, as maternal thyroid-stimulating hormone receptor antibodies can be transferred to a developing fetus and can cause transient neonatal Graves' disease. The HCP responsible for managing the patient's pregnancy must be made aware of the increased risk of thyroid disorders due to the patient's LEMTRADA treatment, and the need for these to be appropriately treated.

Strategies to Mitigate the Risk of Autoimmune Conditions

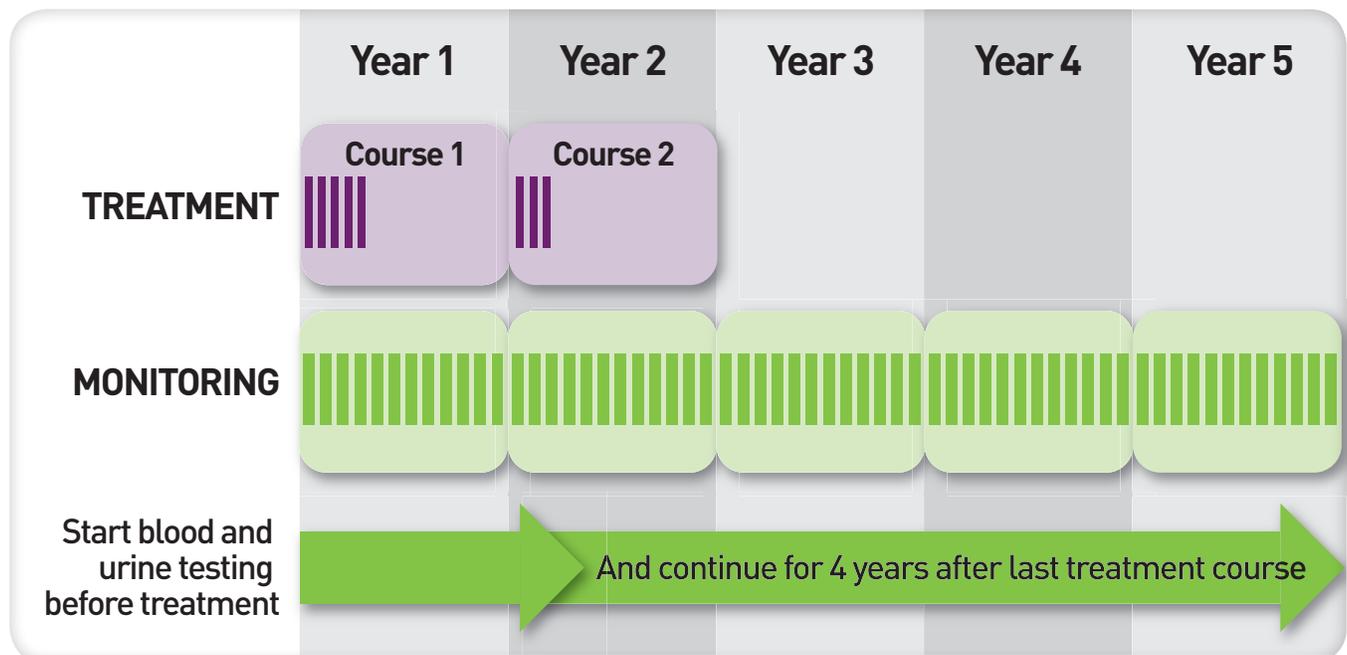
In order to minimize possible risks and side effects of LEMTRADA, prescribers and patients must commit to 48 months of follow-up after the last infusion of LEMTRADA. It is important that patients understand that they should continue with the monitoring, even if they are feeling well.

Creating a partnership between you and your patient, along with careful review of the patient education tool (*What You Need to Know About LEMTRADA Treatment: A Patient Guide*) with your patient, will help patients to:

- > Comply with periodic tests
- > Identify and report symptoms early
- > Receive prompt and appropriate treatment if needed

To enhance your understanding of the duration of the effects of treatment and the length of required follow-up, please refer to the diagrams below titled **Overview of LEMTRADA Treatment** and **Overview of LEMTRADA Monitoring**.

Overview of LEMTRADA Treatment



Overview of LEMTRADA Monitoring

Condition	Activity	Timing	
Immune Thrombocytopenia (ITP)	Complete blood count with differential	Prior to initiating LEMTRADA treatment	Monthly until 48 months after last infusion
Glomerular Nephropathies, including anti-GBM disease	Serum creatinine	Prior to initiating LEMTRADA treatment	Monthly until 48 months after last infusion
Glomerular Nephropathies, including anti-GBM disease	Urinalysis with microscopy	Prior to initiating LEMTRADA treatment	Monthly until 48 months after last infusion
Thyroid Disorders	Thyroid function tests (such as TSH)	Prior to initiating LEMTRADA treatment	Every 3 months until 48 months after last infusion

Infusion Reactions

Most patients treated with LEMTRADA in controlled clinical trials in MS experienced infusion reactions during or after LEMTRADA administration. In some patients, infusion reactions were reported more than 24 hours after LEMTRADA infusion. Serious reactions occurred in 3% of patients, including cases of anaphylaxis in 2 patients (including anaphylactic shock), angioedema, bronchospasm, hypotension, chest pain, bradycardia, tachycardia (including atrial fibrillation), transient neurologic symptoms, hypertension, headache, pyrexia, and rash. Other infusion reactions included nausea, urticaria, pruritus, insomnia, chills, flushing, fatigue, dyspnea, pulmonary infiltrates, dysgeusia, dyspepsia, dizziness and pain. In clinical studies, 0.6% of patients with infusion reactions received epinephrine or atropine.

Premedicate with high-dose corticosteroids (1000 mg of methylprednisolone or equivalent) immediately prior to LEMTRADA infusion and for the first 3 days of each treatment course. Consider pretreatment with antihistamines and/or antipyretics prior to LEMTRADA administration. Infusion reactions may occur in patients despite pretreatment.

LEMTRADA can only be administered in certified healthcare settings that have on-site access to equipment and personnel trained to manage infusion reactions (including anaphylaxis and cardiac and respiratory emergencies).

Patients must be observed for infusion reactions during and for at least 2 hours after each LEMTRADA infusion. Consider longer periods of observation if clinically indicated. Vital signs should be monitored before the infusion and periodically during the infusion. If an infusion reaction occurs, appropriate symptomatic treatment should be provided as needed. The duration of the infusion may be extended if clinically indicated. If severe infusion reactions occur, immediate discontinuation of the infusion should be considered.



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Malignancies

LEMTRADA is an immunomodulatory therapy, and caution should be exercised in initiating LEMTRADA in patients with pre-existing or ongoing malignancies.

Thyroid Cancer

LEMTRADA may increase the risk of thyroid cancer. In controlled clinical studies, 0.3% of LEMTRADA-treated patients developed thyroid cancer, compared to none in the interferon beta-1a-treated group. However, screening for thyroid cancer was performed more frequently in the LEMTRADA-treated group, because of the higher incidence of autoimmune thyroid disorders in those patients. Two additional cases of thyroid cancer in LEMTRADA-treated patients occurred in uncontrolled studies.

Melanoma

LEMTRADA may increase the risk of melanoma. In uncontrolled studies, 0.3% of LEMTRADA-treated patients developed melanoma or melanoma in situ. One of those patients had evidence of locally advanced disease.

Lymphoproliferative Disorders and Lymphoma

Cases of lymphoproliferative disorders and lymphoma have occurred in LEMTRADA-treated patients with MS, including a MALT lymphoma, Castleman's Disease, and a fatality following treatment of non-Epstein Barr Virus-associated Burkitt's lymphoma. There are postmarketing reports of Epstein Barr Virus-associated lymphoproliferative disorders in non-MS patients.

Monitoring for Malignancies

Patients and healthcare providers should monitor for symptoms of thyroid cancer, including a new lump or swelling in the neck, pain in the front of the neck, persistent hoarseness or other voice changes, trouble swallowing or breathing, or a constant cough not due to an upper respiratory tract infection.

Perform baseline and yearly skin examinations to monitor for melanoma in patients receiving LEMTRADA.

Patient Enrollment, Counseling, and Management

To enroll your patient in the LEMTRADA REMS Program, you must:

- > Complete the LEMTRADA REMS Patient Enrollment Form for each patient and provide a completed copy to the patient. The completed form should be submitted to the LEMTRADA REMS Program and a copy stored in the patient's records.
- > Enrollment forms can be obtained online (www.LemtradaREMS.com) or by phone (1-855-676-6326).
- > Completed forms should be faxed to 1-855-557-2478.
- > Genzyme will provide confirmation of patient enrollment.

As part of patient management and counseling, you must:

- > Inform your patient about the risks associated with LEMTRADA, including the risks of autoimmune conditions, infusion reactions, and malignancies, and the need for baseline and periodic monitoring. A patient-directed educational guide has been developed for you to use in counseling your patients on the risks associated with LEMTRADA (*What You Need to Know About LEMTRADA Treatment: A Patient Guide*). You should review this guide with your patient on an ongoing basis. You must provide each patient with a copy of this guide and a LEMTRADA Patient Safety Information Card.
- > Perform the baseline and periodic monitoring described above and in the Prescribing Information for LEMTRADA.
- > Complete the LEMTRADA REMS Patient Status Form 6 months after the patient's first infusion with LEMTRADA, and every 6 months thereafter, until 48 months after the completion of the patient's last infusion of LEMTRADA, and submit the completed form to the LEMTRADA REMS Program.
- > Notify Genzyme if an enrolled patient who has received LEMTRADA within the last 48 months is no longer under your care.

Ordering LEMTRADA

To order LEMTRADA, you must submit a LEMTRADA REMS Prescription Ordering Form for each LEMTRADA prescription to the LEMTRADA REMS Program. The ordering form can be obtained online (www.LemtradaREMS.com) or by phone (1-855-676-6326).

Completed forms should be faxed to 1-855-557-2478.



Administering LEMTRADA

As part of the LEMTRADA REMS Program, a healthcare facility must be enrolled in the LEMTRADA REMS Program to be able to dispense/administer LEMTRADA. It is important that you select a healthcare facility that is enrolled and active in the LEMTRADA REMS Program for your patient's infusion. A database of certified healthcare facilities is available by phone at 1-855-676-6326.

Prior to your patient's infusion, you must submit a LEMTRADA REMS Patient Authorization and Baseline Lab Form to the LEMTRADA REMS Program indicating completion of each patient's baseline labs within 30 days of the infusion date.

PRIOR TO EACH TREATMENT COURSE OF LEMTRADA

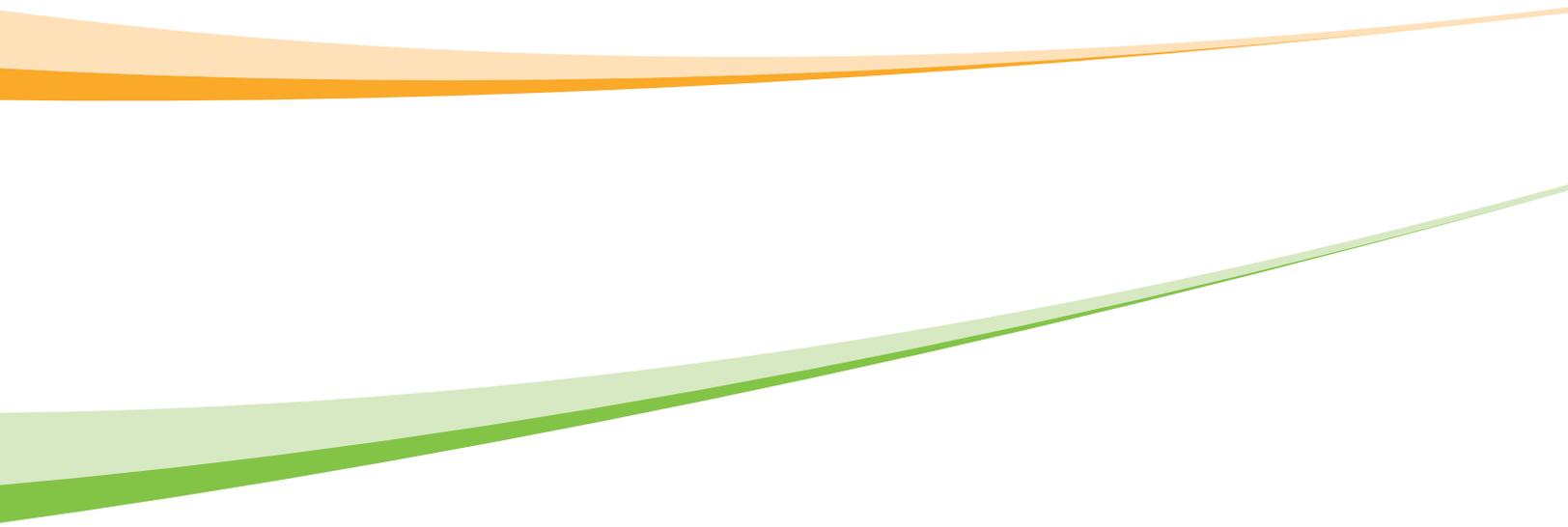
- > Administer corticosteroids (1000 mg methylprednisolone or equivalent) immediately prior to LEMTRADA administration for the first 3 days of any treatment course
- > Administer anti-viral prophylaxis for herpetic viral infection starting on the first day of each treatment course and continuing for a minimum of 2 months following treatment with LEMTRADA or until the CD4+ lymphocyte count is ≥ 200 cells per microliter, whichever occurs later
- > Consider pretreating patients with antihistamines and/or antipyretics prior to LEMTRADA administration as needed

Adverse Event Reporting

Report suspected adverse events to Genzyme Medical Information at 1-800-745-4447 (option 2) or to FDA at 1-800-FDA-1088 or www.FDA.gov/medwatch



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