

ARIKAYCE[®] Prescription and Arikares[®] Support Program Enrollment Form



Fax: 1-800-604-6027 or E-mail: enrollment@arikares.com

Please complete all fields on pages 1 and 3 to prevent any delays and include scanned copies of both sides of the patient's insurance card (fields marked with an asterisk [*] are mandatory/required).



Questions?

Phone: 1-833-ARIKARE (1-833-274-5273)
Alternate Phone: 1-973-437-2376

PATIENT INFORMATION

*Patient First Name: _____ *Patient Last Name: _____ *MI: _____
*DOB: _____ *Gender: Male Female Last 4 of SSN: _____
*Physical Address: _____
*City: _____ *State: _____ *ZIP: _____
*Mailing Address: _____ Same as Physical Address
*City: _____ *State: _____ *ZIP: _____
*Primary Phone: _____ Secondary Phone: _____ E-mail: _____
Preferred Contact Method(s): (check all that apply) Phone E-mail Text
Preferred Time to Contact: Morning Afternoon Evening
Preferred Contact Language: English Spanish Other: _____
Authorized Alternate Contact: _____
Alternate Contact Phone: _____ Relationship to Patient: _____

Prescription Insurance Information (Please Send a Copy of Insurance Card)

*Prescription Coverage Plan Name: _____
Beneficiary/Cardholder: _____ Relationship to Cardholder: _____
*Primary Rx Insurance ID #: _____ *Group #: _____
*BIN: _____ *PCN: _____ *Phone: _____
*Primary Rx Plan Type: Private/Commercial Medicare Part D Medicaid TRICARE Other
Secondary Rx Plan Name: _____
Beneficiary/Cardholder: _____ Relationship to Cardholder: _____
Secondary Rx Insurance ID #: _____ Group #: _____
BIN: _____ PCN: _____ Phone: _____
Secondary Rx Plan Type: Private/Commercial Medicare Part D Medicaid TRICARE Other

Patient Does Not Have Insurance

Patient Authorization Signature

Information Disclosure—I have read and understand the Patient Authorization on page 2, and I agree to allow My Information (as defined in the Patient Authorization) to be used and shared as described in the Authorization.

*Patient Signature: _____ *Date: _____

Program Enrollment—By signing below, I agree to enroll in the Arikares Support Program and verify that the information in the "Patient Information" section of this form is accurate and complete.

*Patient Signature: _____ *Date: _____

Please see Indication and Important Safety Information for ARIKAYCE, including Boxed Warning, on page 4. Please see accompanying full Prescribing Information.

PATIENT AUTHORIZATION

By signing the Patient Authorization block on page 1, I authorize my healthcare providers, including the pharmacies I use, and my health insurance plan(s) to disclose my personal information, including information about me (eg, my name, address) and my health, including my finances, insurance, prescriptions, pharmacy fills/claims, and medical condition (“My Information”) to Insmmed (the manufacturer of ARIKAYCE® [amikacin liposome inhalation suspension]) and its affiliates, agents, and contractors, including the administrators of the Insmmed *Arikares*® *Support Program*, the dispensing pharmacies of Insmmed products, and any other person or entity assisting Insmmed in the administration of the *Arikares* Program (collectively, the “Insmmed *Arikares* Team”), for the purposes listed below:

1. To investigate, verify, and determine my insurance coverage for ARIKAYCE
2. To provide financial assistance, and support to facilitate access to ARIKAYCE and the Lamira® System as prescribed by my treating physician
3. To facilitate a voluntary training session educating on device use and successful treatment initiation
4. To determine my initial and continuing eligibility for other assistance programs
5. To contact me by phone, mail, e-mail (if my e-mail address was provided), cell phone, or text message (if my cell phone was provided) to request further information, discuss the application process, administer the Program, evaluate treatment progress and/or the effectiveness of the Program, and to conduct market research
6. For Insmmed’s internal business purposes of continuous improvement, including ongoing quality control
7. To help ensure the accuracy and completeness of my applications
8. To send me marketing information, offers, and educational materials related to MAC (*Mycobacterium avium* complex) lung disease and/or ARIKAYCE

I understand that my pharmacy provider may receive remuneration from Insmmed in exchange for the health information provided and/or for any therapy support services provided to me. I also understand that once My Information has been disclosed under this Authorization, federal privacy laws may no longer protect it and that it may be subject to further disclosure. I specifically authorize the Insmmed *Arikares* Team to use and disclose My Information for the purposes listed above. I further understand that if I decline to sign this Authorization, that will not affect my eligibility for health plan benefits and treatment by my healthcare providers, but I will not have access to the education and services available through the *Arikares Support Program*. I understand that I may revoke this Authorization at any time by calling **1-833-274-5273** (alternate phone 1-973-437-2376) or writing to Insmmed Incorporated, Attn: Arikares, 700 US Highway 202/206, Bridgewater, NJ 08807. If I do revoke this Authorization, the Insmmed *Arikares* Team will stop accessing, using, and disclosing My Information thereafter, but the uses and disclosures previously made in reliance on the Authorization will not be deemed invalid. This Authorization expires ten (10) years from the date of my Program Enrollment signature on page 1, unless specified or mandated to be shorter by applicable state law. I understand that I am entitled to a copy of this Authorization once signed.

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Phone: **1-833-ARIKARE** (1-833-274-5273)

Alternate Phone: 1-973-437-2376

HEALTHCARE PROFESSIONAL & PRESCRIPTION INFORMATION

*Prescriber First Name: _____ *Prescriber Last Name: _____

*Practice Name: _____ Specialty: _____

*Address: _____ *City: _____ *State: _____ *ZIP: _____

*Phone: _____ *Fax: _____ *NPI #: _____

Office Contact Name: _____ Office Contact Phone: _____

Office Contact E-mail: _____

If Applicable, Check Appropriate Box for Specialty Pharmacy Preference:

No Preference

Maxor Specialty Pharmacy

PANTHERx Specialty Pharmacy



Official Prescription Information

*Patient First Name: _____ *Patient Last Name: _____ *DOB: _____

*Product: ARIKAYCE[®] (amikacin liposome inhalation suspension) *Quantity: 28-Day Supply: 28-Vial Pack (28 Vials of Medication, 4 Aerosol Heads, and 1 Handset)
(First Shipment Includes Lamira[®] System)

*Dosing Info: Once-Daily 590 mg/8.4 mL

*# of Refills: _____

New York prescribers, please submit prescription on an original NY State prescription blank. The prescriber is to comply with his or her state-specific form, fax language, etc. Non-compliance with state-specific requirements could result in outreach to the prescriber.

***Substitution Permitted?** Yes No

Prescriber Certification

I certify that the above therapy is medically necessary and that the information provided is accurate to the best of my knowledge. I certify that I am the prescriber who has prescribed ARIKAYCE to the previously identified patient and that I provided the patient with a description of the Arikares Support Program. I authorize the Arikares Support Program to act on my behalf for the purposes of transmitting this prescription to the appropriate pharmacy.

***Prescriber Signature:** _____ ***Date:** _____

No stamped signatures accepted

Special Instructions:

Please see Indication and Important Safety Information for ARIKAYCE, including Boxed Warning, on page 4. Please see accompanying full Prescribing Information.

INDICATION

LIMITED POPULATION: ARIKAYCE® is indicated in adults, who have limited or no alternative treatment options, for the treatment of *Mycobacterium avium* complex (MAC) lung disease as part of a combination antibacterial drug regimen in patients who do not achieve negative sputum cultures after a minimum of 6 consecutive months of a multidrug background regimen therapy. As only limited clinical safety and effectiveness data for ARIKAYCE are currently available, reserve ARIKAYCE for use in adults who have limited or no alternative treatment options. This drug is indicated for use in a limited and specific population of patients.

This indication is approved under accelerated approval based on achieving sputum culture conversion (defined as 3 consecutive negative monthly sputum cultures) by Month 6. Clinical benefit has not yet been established. Continued approval for this indication may be contingent upon verification and description of clinical benefit in confirmatory trials.

Limitation of Use: ARIKAYCE has only been studied in patients with refractory MAC lung disease defined as patients who did not achieve negative sputum cultures after a minimum of 6 consecutive months of a multidrug background regimen therapy. The use of ARIKAYCE is not recommended for patients with non-refractory MAC lung disease.

IMPORTANT SAFETY INFORMATION

WARNING: RISK OF INCREASED RESPIRATORY ADVERSE REACTIONS

ARIKAYCE has been associated with an increased risk of respiratory adverse reactions, including hypersensitivity pneumonitis, hemoptysis, bronchospasm, and exacerbation of underlying pulmonary disease that have led to hospitalizations in some cases.

Hypersensitivity Pneumonitis has been reported with the use of ARIKAYCE in the clinical trials. Hypersensitivity pneumonitis (reported as allergic alveolitis, pneumonitis, interstitial lung disease, allergic reaction to ARIKAYCE) was reported at a higher frequency in patients treated with ARIKAYCE plus background regimen (3.1%) compared to patients treated with background regimen alone (0%). Most patients with hypersensitivity pneumonitis discontinued treatment with ARIKAYCE and received treatment with corticosteroids. If hypersensitivity pneumonitis occurs, discontinue ARIKAYCE and manage patients as medically appropriate.

Hemoptysis has been reported with the use of ARIKAYCE in the clinical trials. Hemoptysis was reported at a higher frequency in patients treated with ARIKAYCE plus background regimen (18.4%) compared to patients treated with background regimen alone (13.4%). If hemoptysis occurs, manage patients as medically appropriate.

Bronchospasm has been reported with the use of ARIKAYCE in the clinical trials. Bronchospasm (reported as asthma, bronchial hyperreactivity, bronchospasm, dyspnea, dyspnea exertional, prolonged expiration, throat tightness, wheezing) was reported at a higher frequency in patients treated with ARIKAYCE plus background regimen (28.7%) compared to patients treated with background regimen alone (10.7%). If bronchospasm occurs during the use of ARIKAYCE, treat patients as medically appropriate.

Exacerbations of underlying pulmonary disease have been reported with the use of ARIKAYCE in the clinical trials. Exacerbations of underlying pulmonary disease (reported as chronic obstructive pulmonary disease (COPD), infective exacerbation of COPD, infective exacerbation of bronchiectasis) have been reported at a higher frequency in patients treated with ARIKAYCE plus background regimen (15.2%) compared to patients treated with background regimen alone (9.8%). If exacerbations of underlying pulmonary disease occur during the use of ARIKAYCE, treat patients as medically appropriate.

Anaphylaxis and Hypersensitivity Reactions: Serious and potentially life-threatening hypersensitivity reactions, including anaphylaxis, have been reported in patients taking ARIKAYCE. Signs and symptoms include acute onset of skin and mucosal tissue hypersensitivity reactions (hives, itching, flushing, swollen lips/tongue/uvula), respiratory difficulty (shortness of breath, wheezing, stridor, cough), gastrointestinal symptoms (nausea, vomiting, diarrhea, crampy abdominal pain), and cardiovascular signs and symptoms of anaphylaxis (tachycardia, low blood pressure, syncope, incontinence, dizziness). Before therapy with ARIKAYCE is instituted, evaluate for previous hypersensitivity reactions to aminoglycosides. If anaphylaxis or a hypersensitivity reaction occurs, discontinue ARIKAYCE and institute appropriate supportive measures.

Ototoxicity has been reported with the use of ARIKAYCE in the clinical trials. Ototoxicity (including deafness, dizziness, presyncope, tinnitus, and vertigo) were reported with a higher frequency in patients treated with ARIKAYCE plus background regimen (17%) compared to patients treated with background regimen alone (9.8%). This was primarily driven by tinnitus (8.1% in ARIKAYCE plus background regimen vs 0.9% in the background regimen alone arm) and dizziness (6.3% in ARIKAYCE plus background regimen vs 2.7% in the background regimen alone arm). Closely monitor patients with known or suspected auditory or vestibular dysfunction during treatment with ARIKAYCE. If ototoxicity occurs, manage patients as medically appropriate, including potentially discontinuing ARIKAYCE.

Nephrotoxicity was observed during the clinical trials of ARIKAYCE in patients with MAC lung disease but not at a higher frequency than background regimen alone. Nephrotoxicity has been associated with the aminoglycosides. Close monitoring of patients with known or suspected renal dysfunction may be needed when prescribing ARIKAYCE.

Neuromuscular Blockade: Patients with neuromuscular disorders were not enrolled in ARIKAYCE clinical trials. Aminoglycosides may aggravate muscle weakness by blocking the release of acetylcholine at neuromuscular junctions. Closely monitor patients with known or suspected neuromuscular disorders, such as myasthenia gravis. If neuromuscular blockade occurs, it may be reversed by the administration of calcium salts but mechanical respiratory assistance may be necessary.

Embryo-Fetal Toxicity: Aminoglycosides can cause fetal harm when administered to a pregnant woman. Aminoglycosides, including ARIKAYCE, may be associated with total, irreversible, bilateral congenital deafness in pediatric patients exposed *in utero*. Patients who use ARIKAYCE during pregnancy, or become pregnant while taking ARIKAYCE should be apprised of the potential hazard to the fetus.

Contraindications: ARIKAYCE is contraindicated in patients with known hypersensitivity to any aminoglycoside.

Most Common Adverse Reactions: The most common adverse reactions in Trial 1 at an incidence $\geq 5\%$ for patients using ARIKAYCE plus background regimen compared to patients treated with background regimen alone were dysphonia (48% vs 2%), cough (40% vs 17%), bronchospasm (29% vs 11%), hemoptysis (18% vs 13%), musculoskeletal pain (18% vs 9%), upper airway irritation (18% vs 2%), ototoxicity (17% vs 10%), fatigue and asthenia (16% vs 10%), exacerbation of underlying pulmonary disease (15% vs 10%), diarrhea (13% vs 5%), nausea (12% vs 4%), headache (10% vs 5%), pneumonia (9% vs 9%), pyrexia (8% vs 5%), decreased weight (7% vs 1%), vomiting (7% vs 4%), rash (6% vs 1%), change in sputum (6% vs 1%), and chest discomfort (5% vs 3%).

Drug Interactions: Avoid concomitant use of ARIKAYCE with medications associated with neurotoxicity, nephrotoxicity, and ototoxicity. Some diuretics can enhance aminoglycoside toxicity by altering aminoglycoside concentrations in serum and tissue. Avoid concomitant use of ARIKAYCE with ethacrynic acid, furosemide, urea, or intravenous mannitol.

Overdosage: Adverse reactions specifically associated with overdose of ARIKAYCE have not been identified. Acute toxicity should be treated with immediate withdrawal of ARIKAYCE, and baseline tests of renal function should be undertaken. Hemodialysis may be helpful in removing amikacin from the body. In all cases of suspected overdosage, physicians should contact the Regional Poison Control Center for information about effective treatment.



Receiving your ARIKAYCE treatment

Box 1: 28-Day ARIKAYCE Kit



- 1 4 weekly boxes
- 2 1 ARIKAYCE Quick Start Guide
- 3 1 Instructions for Use insert
- 4 1 Full Prescribing Information and Medication Guide
- 5 Lamira Nebulizer Handset
- 6 28 once-daily vials of ARIKAYCE
- 7 Lamira Aerosol Head
- 8 Cooler return form

Box 2: The Lamira® Nebulizer System



Getting started box



- i 1 carrying case
- ii 1 connection cord
- iii 1 A/C power supply
- iv 4 AA batteries
- v 1 eBase® controller
- vi Getting started box
- vii Lint-free towel
- viii 2 oz dish soap sample
- ix Lint-free drying mat
- x Timer

IMPORTANT SAFETY INFORMATION (cont'd)

- **serious allergic reactions.** Serious allergic reactions that may lead to death have happened to people who take ARIKAYCE. Stop taking ARIKAYCE right away and get emergency medical help if you have any of the following symptoms of a serious allergic reaction: hives, itching, redness or blushing of the skin (flushing), swollen lips, tongue or throat, trouble breathing or wheezing, shortness of breath, noisy high-pitched breathing (stridor), cough, nausea, vomiting, diarrhea, feel cramps in your stomach area, fast heart rate, feeling light headed, feeling faint, loss of control of the bowels or bladder (incontinence), and dizziness

While using ARIKAYCE, these side effects may become serious enough that treatment in a hospital is needed. Call your healthcare provider or get medical help right away if you have any of these serious side effects while taking ARIKAYCE. Your healthcare provider may ask you to stop using ARIKAYCE for a short period of time or completely stop using ARIKAYCE.

Do not use ARIKAYCE if you are allergic to any aminoglycoside, or any of the ingredients in ARIKAYCE.

Before using ARIKAYCE, tell your healthcare provider about all medical conditions, including if you:

- have asthma, COPD, shortness of breath, or wheezing (bronchospasm)
- have been told you have poor lung function
- have hearing problems, such as ringing in your ears or hearing loss
- have dizziness or a sense of the room spinning
- have kidney problems
- have neuromuscular disease, such as myasthenia gravis
- are pregnant or plan to become pregnant. It is not known if ARIKAYCE can harm your unborn baby. ARIKAYCE is in a class of medicines that may be connected with complete deafness in babies at birth. The deafness affects both ears and cannot be changed
- are breastfeeding or plan to breastfeed. It is not known if the medicine in ARIKAYCE passes into your breast milk and if it can harm your baby. Talk to your healthcare provider about the best way to feed your baby during treatment with ARIKAYCE

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

ARIKAYCE may cause serious side effects, including:

- **hearing loss or ringing in the ears (ototoxicity).** Ototoxicity is a serious and common side effect of ARIKAYCE. Tell your healthcare provider right away if you have hearing loss or you hear noises in your ears, such as ringing or hissing. Tell your healthcare provider if you start having problems with balance or dizziness (vertigo)
- **worsening kidney problems (nephrotoxicity).** ARIKAYCE is in a class of medicines which may cause worsening kidney problems. Your healthcare provider may do a blood test to check how well your kidneys are working during your treatment with ARIKAYCE
- **worsening muscle weakness (neuromuscular blockade).** ARIKAYCE is in a class of medicines which can cause muscle weakness to get worse in people who already have problems with muscle weakness (myasthenia gravis)

The most common side effects of ARIKAYCE include: changes in voice and hoarseness (dysphonia), cough during or after a dose of ARIKAYCE, especially in the first month after starting treatment, muscle pain, sore throat, tiredness (fatigue), diarrhea, nausea, headache, fever, decreased weight, vomiting, rash, increased sputum, or chest discomfort. These are not all of the possible side effects of ARIKAYCE. **Call your doctor or pharmacist for medical advice about side effects. You may report side effects to FDA at 1-800-FDA-1088 or www.fda.gov/medwatch.**

Please see additional Important Safety Information and full Prescribing Information, including Boxed Warning.

