A GUIDE FOR **RADICAVA ORS® PATIENTS**

The importance of managing ALS and how RADICAVA ORS may help

INDICATION

RADICAVA ORS® (edaravone) is indicated for the treatment of amyotrophic lateral sclerosis (ALS).

IMPORTANT SAFETY INFORMATION

Do not receive RADICAVA ORS® (edaravone) if you are allergic to edaravone or any of the ingredients in RADICAVA ORS.

Please see the full Prescribing Information and Patient Information, also available at www.radicavaors.com.

Radicava ORS® (edaravone) Oral Suspension 105mg/5mL

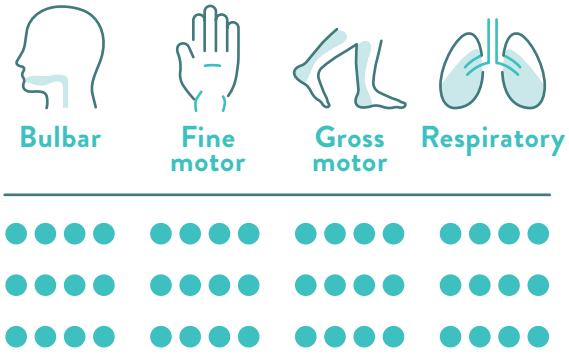
The importance of managing ALS

Although there are no known treatments that cure or reverse amyotrophic lateral sclerosis (ALS), RADICAVA ORS® (edaravone) can make a difference in helping to slow the loss of physical function.

ALS continues to progress, which can make it hard to know if treatment is helping. The ALS Functional Rating Scale–Revised (ALSFRS-R) measures how you are doing now and will help track disease progression. It consists of 4 categories that add up to a total of 48 points. In addition to treatment with RADICAVA ORS, talk to your doctor about other ways to help slow your progression or adapt to functional losses over time.

Points represent physical function

Decline on the ALSFRS-R may vary widely among those with ALS. Some can lose 0 to 1 point per month while others may lose up to 6 points per month.



Losing or keeping a single point on the ALSFRS-R can have a significant impact on those living with ALS.

Even though you might not feel a change, being on RADICAVA ORS is an important step that may help slow the loss of physical function over time.

IMPORTANT SAFETY INFORMATION

Before you take RADICAVA ORS, tell your healthcare provider about all of your medical conditions, including if you:

- have asthma.
- are allergic to other medicines.
- are pregnant or plan to become pregnant. It is not known if RADICAVA ORS will harm your unborn baby.
- are breastfeeding or plan to breastfeed. It is not known if RADICAVA ORS passes into your breastmilk. You and your healthcare provider should decide if you will receive RADICAVA ORS or breastfeed.

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

Please see the full Prescribing Information and Patient Information, also available at www.radicavaors.com.

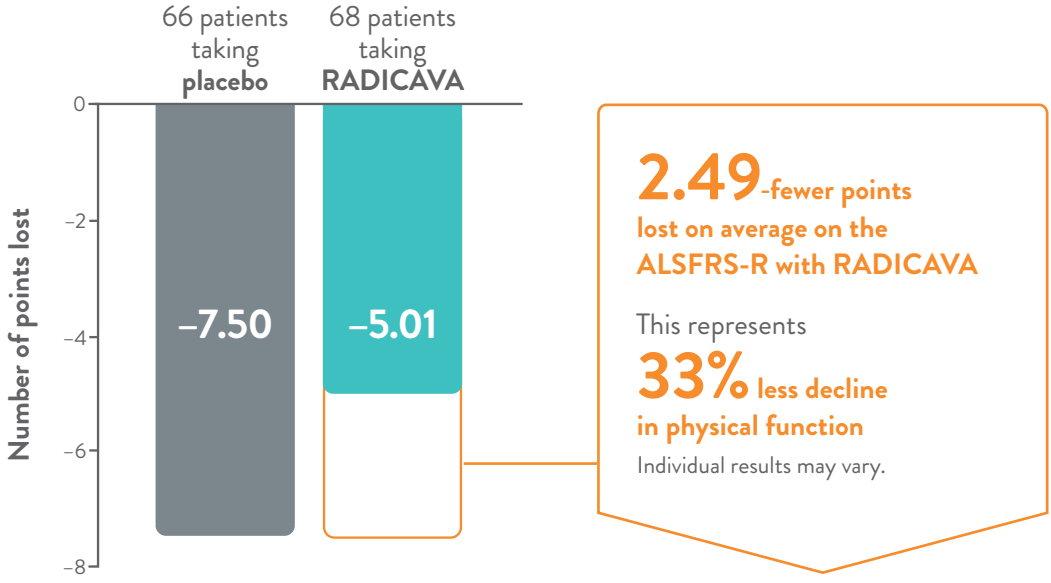
Radicava ORS® (edaravone) Oral Suspension 105mg/5mL

RADICAVA was evaluated in a phase 3 clinical study

In ALS, every point matters

On the ALSFRS-R, the more points preserved, the greater the physical function. In the pivotal clinical study, patients taking RADICAVA® (edaravone) lost an average of 2.49-fewer points vs those taking placebo who declined more rapidly at 24 weeks.

RADICAVA slowed the loss of physical function vs placebo at 24 weeks



RADICAVA ORS® (edaravone) was generally well tolerated

- The safety profile of RADICAVA was evaluated in multiple placebo-controlled studies of 184 patients with ALS
- RADICAVA ORS was generally well tolerated in a 6-month clinical study of 185 patients with ALS

The most common side effects for patients taking RADICAVA were bruising (contusion) [15%], problems with walking (gait disturbance) [13%], and headache [10%]. Fatigue was also reported in 7.6% of patients taking RADICAVA ORS.

These are not all of the possible side effects with RADICAVA or RADICAVA ORS.

IMPORTANT SAFETY INFORMATION

What are the possible side effects of RADICAVA ORS?

RADICAVA ORS may cause serious side effects, including hypersensitivity (allergic) reactions and sulfite allergic reactions.

- Hypersensitivity reactions have happened in people taking RADICAVA ORS and can happen after your medicine has been taken.

Please see the full Prescribing Information and Patient Information, also available at www.radicavaors.com.

Radicava ORS® (edaravone) Oral Suspension 105mg/5mL

The impact of RADICAVA ORS on patients

In 2024, RADICAVA ORS® (edaravone) was recognized as a **major contribution to patient care.**

This is due to its oral suspension route of administration that provides a less burdensome option vs IV administration.



Not actual size.

FDA=Food and Drug Administration; IV=intravenous.

IMPORTANT SAFETY INFORMATION

What are the possible side effects of RADICAVA ORS? *(continued)*

- RADICAVA ORS contains sodium bisulfite, a sulfite that may cause a type of allergic reaction that can be serious and life-threatening. Sodium bisulfite can also cause less severe asthma episodes in certain people. Sulfite sensitivity can happen more often in people who have asthma than in people who do not have asthma.
- Tell your healthcare provider right away or go to the nearest emergency room if you have any of the following symptoms: hives; swelling of the lips, tongue, or face; fainting; breathing problems; wheezing; trouble swallowing; dizziness; itching; or an asthma attack (in people with asthma).

Your healthcare provider will monitor you during treatment to watch for signs and symptoms of all the serious side effects and allergic reactions.

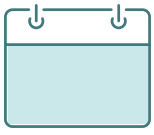
Common side effects reported include bruising (contusion), problems walking (gait disturbance), headache and fatigue. These are not all the possible side effects of RADICAVA ORS.

Please see the full Prescribing Information and Patient Information, also available at www.radicavaors.com.

RADICAVA ORS has a foundation of clinical and real-world experience

RADICAVA ORS® (edaravone) is an oral formulation of RADICAVA® (edaravone)—a proven therapy evaluated in over 2 decades of clinical research, including 4 phase 3 clinical trials.

Since 2017



9+

years of RADICAVA experience since FDA approval



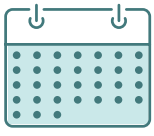
~600

people have taken part in multiple clinical studies that have demonstrated the safety and tolerability of RADICAVA



17,300+

people have been treated with RADICAVA ORS[a]



1,900,000+

days of therapy[b]

For more about RADICAVA ORS, visit radicavaors.com

[a]Based on RADICAVA ORS prescriptions submitted in the US as of May 2026. Not independently verified.
[b]Days of therapy based on number of Starter Kit and Maintenance Kit cartons sold for RADICAVA ORS as of May 2026. Each Starter Kit includes 14 days of therapy and each Maintenance Kit includes 10 days of therapy.

IMPORTANT SAFETY INFORMATION

Call your doctor for medical advice about side effects. You may report side effects to FDA at 1-800-FDA-1088. You may also report side effects to www.fda.gov/medwatch or Shionogi Inc. at 1-888-292-0058.

Please see the full Prescribing Information and Patient Information, also available at www.radicavaors.com.

Radicava ORS® (edaravone) Oral Suspension 105mg/5mL

JourneyMate SUPPORT PROGRAM™

Dedicated Team. Patient-Focused Approach.

A patient-focused approach to ALS and treatment support

No matter where you are in your ALS and RADICAVA ORS® (edaravone) journey, the ***JourneyMate Support Program***™ gives you the understanding, personalized answers, and resources to help you move forward. Experienced program team members are trained to address your educational needs as you continue treatment and provide you with personalized answers and resources for living with ALS. Call the ***JourneyMate Support Program***™ or visit radicavaors.com.[a] This program is here to supplement the resources that your doctor provides.

Have questions about ALS or your RADICAVA ORS treatment?

Call a JourneyMate Resource Specialist—your go-to resource in the ***JourneyMate Support Program***™—at 1-855-457-6968 from 9 AM to 9 PM ET, Monday through Friday.

Some patients with commercial health insurance may be eligible for an Out-of-Pocket Savings Program to help save on out-of-pocket costs for your RADICAVA ORS prescription.[b]

Annual re-enrollment, available upon re-verification of commercial insurance benefits will confirm your continued eligibility for the Out-of-Pocket Savings Program.

A JourneyMate Resource Specialist discusses basic information about Shionogi Inc. products and does not take the place of a patient's doctor. Patients should be sure to talk to their doctor about all treatment-related questions, as their doctor is the best person to help a patient decide if treatment is right for them. If a patient has a medical emergency, they should call 911. Adverse events or product complaints should be reported by calling 1-888-292-0058.

[a]Read the Instructions for Use before you take RADICAVA ORS.

[b]This is not insurance. This offer is for eligible patients who have private, commercial health insurance with prescription coverage for RADICAVA ORS. Patients enrolled in commercial prescription drug insurance and Medicare Part A and/or Part B are eligible for assistance so long as they meet all other eligibility criteria and are not enrolled in or become enrolled in Medicare Parts C or D. Support is not valid for patients covered, in whole or in part, by Medicaid, Department of Veterans Affairs (VA), Department of Defense (DoD), or any other federal or state health insurance program. Annual maximum benefit per patient. Additional terms and conditions apply. See full Eligibility Requirements & Terms and Conditions at RadicavaOOP.com.

SHIONOGI

© 2026 Shionogi Inc. Florham Park, NJ 07932. All Rights Reserved.
RADICAVA, RADICAVA ORS, and the RADICAVA ORS logo are registered trademarks of Shionogi Inc.
JourneyMate Support Program is a trademark of Shionogi Inc.
CP-OE-US-1893 08/26

Radicava ORS® (edaravone) Oral Suspension 105mg/5mL