

WHAT I CAN DO NEXT RELIES ON WHAT I DO NOW

I started RADICAVA ORS early to help slow the loss of function for as long as possible

Actor portrayals. Individual results may vary.

Proven to help slow the loss of physical function in patients with amyotrophic lateral sclerosis (ALS).

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.

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.

Please see additional Important Safety Information throughout and full Prescribing Information and Patient Information.

Why You Should Be Thinking About Treatment Now

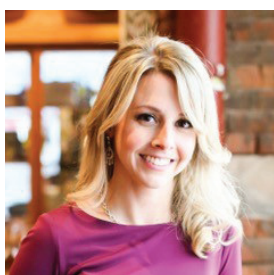
What you should know

Nonfamilial (sporadic) ALS is a form of ALS that does not have a known family history or genetic predisposition. It accounts for approximately 90% of all ALS cases in the United States.^a

As of 2025, there are only 2 available treatments in the US for nonfamilial ALS: riluzole and edaravone.

Your doctor may prescribe both RADICAVA ORS[®] (edaravone) and riluzole.

Talk to your doctor about the right treatment option for you.



“One thing I really like about RADICAVA ORS is how it fits into my daily routine.”

— Stephanie, living with ALS

Hear from real patients about their experience navigating an ALS diagnosis and treatment with RADICAVA ORS.

^aNational Institute of Neurological Disorders and Stroke. Amyotrophic lateral sclerosis (ALS). <https://www.ninds.nih.gov/health-information/disorders/amyotrophic-lateral-sclerosis-als>.

IMPORTANT SAFETY INFORMATION

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

- 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.

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

Recognized by the FDA as a Major Contribution to Patient Care*



Maintenance Dose.
Not actual size.

***RADICAVA ORS[®] can be taken orally (by mouth), which is a less burdensome option than intravenous (IV) administration.^a**



5 mL oral dose may take only a few minutes to administer on treatment days^b



Can be stored at room temperature so you can take it at home or on the go^c

RADICAVA ORS should be taken in the morning on an empty stomach after overnight fasting. Food should not be consumed for 1 hour after administration except water.

RADICAVA ORS has been used as a treatment for people living with ALS since 2022.

Prescribed by
1200+ doctors^d

Over **11,600** people with ALS have
chosen treatment with RADICAVA ORS
for **1,181,300+** days of therapy^{d,e}

82% of patients have
stayed on treatment
for at least **3 months**^f

^aUS Food and Drug Administration. Clinical superiority findings. <https://www.fda.gov/industry/designating-orphan-product-drugs-and-biological-products/clinical-superiority-findings>

^bUse the provided 5 mL syringe that comes with the product. Do not use a household teaspoon to measure the medication.

^cStore RADICAVA ORS upright at room temperature between 68°F and 77°F. Protect from light.

^dBased on RADICAVA ORS prescriptions submitted in the US as of December 2024. Not independently verified.

^eDays of therapy based on number of Starter Kit and Maintenance Kit cartons sold for RADICAVA ORS as of December 2024. Each Starter Kit includes 14 days of therapy, and each Maintenance Kit includes 10 days of therapy.

^fBased on patients who have started therapy between June 20, 2022 and October 16, 2024, with at least 1 shipment.

FDA=Food and Drug Administration.

IMPORTANT SAFETY INFORMATION

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Monitoring Progression With the ALSFRS-R Scale

ALS is a neurological disease than can progress more quickly or more slowly for different people. To help keep track of your body’s ability to function over time, your doctor may use a tool called the ALS Functional Rating Scale–Revised (ALSFRS-R). This scale can help track how ALS progresses for you.

The ALSFRS-R consists of 12 items about your body’s abilities across 4 categories, with each task rated on a scale from 0 (unable to perform) to 4 (normal function) for a total of 48 points^a

	Bulbar	<ul style="list-style-type: none">• Speech	<ul style="list-style-type: none">• Salivation	<ul style="list-style-type: none">• Swallowing
	Fine Motor	<ul style="list-style-type: none">• Handwriting	<ul style="list-style-type: none">• Cutting food	<ul style="list-style-type: none">• Dressing and hygiene
	Gross Motor	<ul style="list-style-type: none">• Turning in bed	<ul style="list-style-type: none">• Walking	<ul style="list-style-type: none">• Climbing stairs
	Respiratory	<ul style="list-style-type: none">• Difficulty breathing or shortness of breath (dyspnea)	<ul style="list-style-type: none">• Trouble breathing when lying down (orthopnea)	<ul style="list-style-type: none">• Respiratory insufficiency

Higher scores mean more physical function.

^aCedarbaum JM, Stambler N, Malta E, et al; BDNF ALS Study Group (Phase III). The ALSFRS-R: a revised ALS functional rating scale that incorporates assessments of respiratory function. *J Neurol Sci.* 1999;169(1-2):13-21.

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 additional Important Safety Information throughout and full [Prescribing Information](#) and [Patient Information](#).



Every Point Matters

Losing a single point on the ALSFRS-R may have a big impact on physical function.



Actor
portrayal.

On average, patients lose 1 point out of 48 total points per month on the ALSFRS-R^a



Bulbar

1 point of functional loss in the ability to swallow could mean going from being able to eat on your own to needing to use a feeding tube.



Fine Motor

1 point of functional loss in the hands could mean going from being able to feed yourself without assistance to needing help.



Gross Motor

A 1-point loss in the legs could mean going from being able to walk with an aid (for example, using a cane) to not being able to walk.



Respiratory

A 1-point loss in breathing ability could mean going from breathing on your own to occasionally needing to use a ventilator (or machine) to help with breathing.

The sooner you start treatment, the sooner you may begin slowing the loss of physical function.

^aCedarbaum JM, Stambler N, Malta E, et al; BDNF ALS Study Group (Phase III). The ALSFRS-R: a revised ALS functional rating scale that incorporates assessments of respiratory function. *J Neurol Sci.* 1999;169(1-2):13-21.

IMPORTANT SAFETY INFORMATION

What are the possible side effects of RADICAVA ORS? (cont.)

- 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.

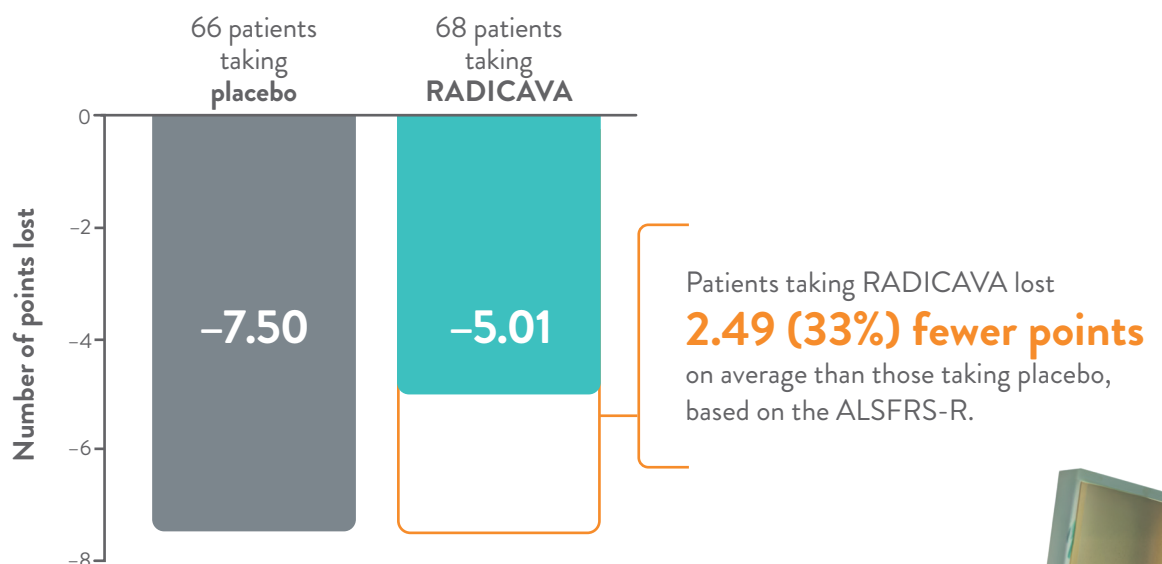
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Radicava ORS[®]
(edaravone) Oral Suspension
105mg/5mL

RADICAVA ORS[®]: Proven to Help Slow the Loss of Physical Function

RADICAVA ORS is an oral form of RADICAVA[®] (edaravone)

In the clinical study, RADICAVA slowed the loss of physical function, as measured by the ALSFRS-R.



This chart shows that in the clinical study, the decline in physical function for patients who received RADICAVA was **33% less** than for those who received placebo at 24 weeks.



IMPORTANT SAFETY INFORMATION

What are the possible side effects of RADICAVA ORS? (cont.)

- 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).

Please see additional Important Safety Information throughout and full [Prescribing Information](#) and [Patient Information](#).

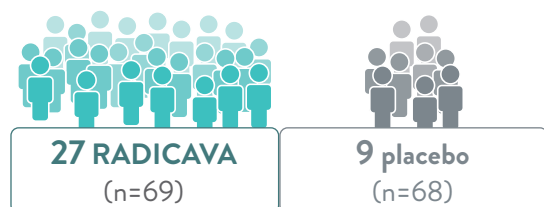
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RADICAVA ORS®: Proven to Help Slow the Loss of Physical Function (cont.)

Patients on RADICAVA® lost fewer points on the ALSFRS-R at 6 months. Fewer points lost means more function retained

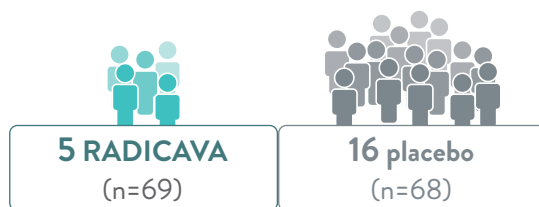
3X more patients
given RADICAVA lost **2 or less points**

(39.1% RADICAVA vs 13.2% placebo)



3X more patients
given placebo lost **11 or more points**

(23.5% placebo vs 7.2% RADICAVA)



A similar number of patients in both treatment arms lost 3 to 10 points
(37/69 with RADICAVA and 43/68 with placebo)

Over 90% of patients in each group were also being treated with riluzole

Your doctor may prescribe RADICAVA ORS in addition to riluzole—the only approved treatments for nonfamilial ALS.

“There are 3 reasons why I take RADICAVA ORS. One, my neurologist and I feel the data is compelling. Two, it’s accessible. And three, the administration fits into my lifestyle so well.”

— Gwen
Patient Advocate & Paid Consultant

IMPORTANT SAFETY INFORMATION

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

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RADICAVA ORS[®] Was Generally Well Tolerated

Side effect profile



The safety profile of RADICAVA was evaluated in multiple placebo-controlled studies in 184 patients with ALS



RADICAVA ORS was generally well tolerated in a 6-month clinical study of 185 patients with ALS

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.

Fewer than 6% (11/185) of patients discontinued RADICAVA ORS because of side effects in the 6-month safety study in 185 patients with ALS.

- Approximately 1% (2/185) of patients discontinued RADICAVA ORS due to gastrointestinal side effects (diarrhea and trouble swallowing)
 - Other reasons for discontinuation in ~1% (2/185) of patients included respiratory failure and muscular weakness

Be sure to tell your doctor if you are allergic to other medications and share all of the medications you are taking.

The safety profiles of RADICAVA and RADICAVA ORS were established in 500+ patients with ALS in multiple clinical studies.

IMPORTANT SAFETY INFORMATION

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

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Taking RADICAVA ORS[®]

RADICAVA ORS is an oral treatment that can fit into your routine



Can be taken orally (by mouth) or via feeding tube



5 mL oral dose may take only a few minutes to administer on treatment days^a



Can be stored at room temperature so you can take it at home or on the go^b

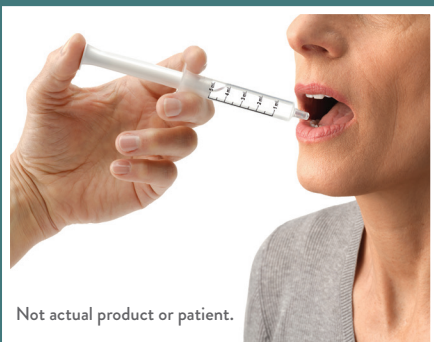


Requires shaking but does not require water or additional mixing steps

RADICAVA ORS should be taken in the morning on an empty stomach after overnight fasting. Food should not be consumed for 1 hour after administration except water.

^aUse the provided 5 mL syringe that comes with the product. Do not use a household teaspoon to measure the medication.

^bStore RADICAVA ORS upright at room temperature between 68°F and 77°F. Protect from light.



Not actual product or patient.

Get your questions answered

Talk with a JourneyMate Resource Specialist who helps you and your loved ones learn about RADICAVA ORS as a treatment option.

Call toll-free 1-855-457-6968 between 9 AM and 9 PM ET, Monday through Friday.^c

Read the Instructions for Use before you take RADICAVA ORS.

^cA JourneyMate Resource Specialist discusses basic information about Tanabe Pharma America, 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.

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 Tanabe Pharma America, Inc. at 1-888-292-0058.

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RADICAVA ORS® Dosing Schedule and Guidance

Mark the calendar to stay on track with treatment

The initial treatment cycle starts with daily dosing of RADICAVA ORS for 14 consecutive days, followed by a 14-day drug-free period. Subsequent treatment cycles include dosing for 10 out of 14 days, followed by a 14-day drug-free period.

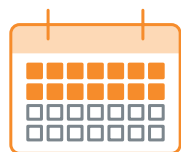


Not actual size.



Not actual size.

Starter treatment cycle (first cycle only)



14 days on
14 days off

RADICAVA ORS is taken once daily for 14 days, followed by 14 days off in a row.

Maintenance treatment cycle (all other cycles)



10 out of 14 days on
14 days off

RADICAVA ORS is taken on 10 of 14 days, followed by 14 days off in a row each month.

Taking RADICAVA ORS daily can occur on **any 10 days** in the 14-day treatment period.

IMPORTANT SAFETY INFORMATION

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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.

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RADICAVA ORS[®] Dosing Schedule and Guidance (cont.)

Fasting guidance for RADICAVA ORS

RADICAVA ORS should be taken in the morning on an empty stomach after overnight fasting.

- No food or drink should be consumed (except water) for 1 hour after administration
- Your doctor can help guide you on any dosing questions or concerns

Type of meal/supplement consumed	Fasting times before and after administration	
	Before	After
High-fat meal (800-1000 calories, 50% fat)	8 hours	1 hour
Low-fat meal (400-500 calories, 25% fat)	4 hours	1 hour
Caloric supplement (250 calories, eg, protein drink)	2 hours	1 hour

Seek immediate medical care if you experience signs or symptoms of an allergic reaction.

[Visit **radicavaors.com/dosingadminguide**](https://radicavaors.com/dosingadminguide) to download the Dosing and Administration Guide for more information.

IMPORTANT SAFETY INFORMATION

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

- 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.

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JourneyMate

SUPPORT PROGRAM™

Dedicated Team. Patient-Focused Approach.

No matter where you are in your ALS journey—from diagnosis to treatment—the **JourneyMate Support Program™** gives you the understanding, answers, and resources to help you move forward. Experienced program team members are trained to address your educational needs and provide you with personalized answers and resources for living with ALS. This program is here to supplement the resources that your doctor provides.



Resource Specialist

A go-to resource in the **JourneyMate Support Program™** for general information about ALS and RADICAVA ORS® for people with ALS and their care partners.



Insurance & Access Specialist

This specialist can help answer questions people with ALS and their care partners have about financial support options and help them understand how to access RADICAVA ORS medication.



ALS Clinical Educator

Once you're prescribed RADICAVA ORS, an ALS Clinical Educator can provide personalized education to you and your family about RADICAVA ORS and will also provide resources to support your treatment.

To learn more, call 1-855-457-6968 between 9 AM and 9 PM ET, Monday through Friday, or visit [radicavaors.com](https://www.radicavaors.com).

The **JourneyMate Support Program™** offers educational support and resources for patients who are considering or have already been prescribed a Tanabe Pharma America, Inc. (TPA) product. An ALS Clinical Educator is an educational resource for patients who have been prescribed an TPA product. An ALS Clinical Educator is provided by TPA and Momentum Life Sciences and is not affiliated with or provided by a doctor. An ALS Clinical Educator does not provide medical advice. The program does not provide medical advice and does not take the place of a patient's doctor. All questions about a condition, diagnosis, or treatment should be referred to the patient's doctor. 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.

IMPORTANT SAFETY INFORMATION

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

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Support for Veterans With ALS

VA (Veterans Affairs) benefits

The VA recognizes ALS as a service-connected disease and provides medical and financial support, including access to treatment, to Veterans with at least 90 continuous days of military service.^a



RADICAVA ORS[®] is on the VA National Formulary subject to prior authorization criteria.

To learn more about assistance and benefits for Veterans with ALS, visit [ALSVeterans.com](https://www.alsveterans.com).

^aA service connection to ALS for any Veteran who develops the disease at any time after separation from service.

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.

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RADICAVA ORS[®]: Cost and Coverage

RADICAVA ORS is widely available



of all Medicare plans cover RADICAVA ORS (with medical exceptions).^a



of all commercial health insurance plans cover RADICAVA ORS.^a

The RADICAVA ORS Out-of-Pocket Assistance Program



of eligible patients with commercial health insurance pay as little as **\$0** per prescription of RADICAVA ORS^b with the Out-of-Pocket Assistance Program. Terms and conditions apply. See page 17 for more details.



How can the *JourneyMate Support Program*[™] help reduce your out-of-pocket costs?

Once your doctor has prescribed RADICAVA ORS and submitted a **Benefit Investigation and Enrollment Form (BIF)** to check how your health insurance covers RADICAVA ORS, an Insurance & Access Specialist will reach out to help you understand the insurance process.

A *JourneyMate Support Program*[™] Insurance & Access Specialist is provided by UBC on behalf of Tanabe Pharma America, Inc. (TPA). A *JourneyMate Support Program*[™] Insurance & Access Specialist may provide information obtained from outside sources about a patient's insurance coverage, financial support options, and whether treatment is covered by their health plan. This information does not require a patient or their doctor to use any TPA product. Because the information provided comes from outside sources, a *JourneyMate Support Program*[™] Insurance & Access Specialist cannot guarantee the information will be accurate or complete.

^aData on file, Tanabe Pharma America, as of February, 2025.

^bSee [Eligibility Requirements & Terms and Conditions](#) on page 17.

IMPORTANT SAFETY INFORMATION

What are the possible side effects of RADICAVA ORS? (cont.)

- 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.

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- 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.

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 Tanabe Pharma America, Inc. at 1-888-292-0058.

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



Actor
portrayals.

**WHAT I CAN DO NEXT
RELIES ON WHAT
I DO NOW**

**Starting RADICAVA ORS[®] now can help slow the loss of
physical function for as long as possible**



Proven to help **slow the loss
of physical function by 33%**
at 24 weeks



An oral treatment
that **can fit into
your routine**



Widely available
and covered by
most insurance plans

IMPORTANT SAFETY INFORMATION

What are the possible side effects of RADICAVA ORS? (cont.)

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Eligibility Requirements & Terms and Conditions for the Out-of-Pocket Assistance Program for RADICAVA ORS®

- Patients who meet all eligibility criteria and are enrolled in the Out-of-Pocket Assistance Program may pay as little as \$0 per RADICAVA ORS prescription dispense, subject to a maximum annual benefit limit per calendar year
- You currently have private, commercial health insurance with prescription coverage for RADICAVA ORS, and your insurance does not cover the entire cost of the medication. Offer is not valid for cash-paying patients
- You are not eligible for RADICAVA ORS assistance if you are enrolled in or become enrolled in Medicare Part C (Medicare Advantage), Medicare Part D (prescription drug benefit), Medicaid, VA, DoD, or any other federal or state insurance program
- If you are enrolled in commercial prescription insurance, Medicare Part A and Medicare Part B, you are eligible for assistance so long as you meet all other eligibility criteria
- You may not seek reimbursement or compensation, in whole or in part, from any government health insurance
- You agree that this Out-of-Pocket Assistance Program is intended solely for the benefit of you as the patient and is not intended for patients in accumulator or maximizer programs. Except where prohibited by applicable state law, Tanabe Pharma America, Inc. reserves the right to modify or discontinue assistance at any time for patients found to be subject to an accumulator adjustment or co-pay maximizer program
- Not valid where prohibited by law, taxed, or otherwise restricted
- This Out-of-Pocket Assistance Program is not health insurance
- Tanabe Pharma America, Inc. has the right to modify, alter, or cancel the Out-of-Pocket Assistance Program at any time without prior notification
- Additional program terms and restrictions apply. Visit [radicava.com](https://www.radicava.com) for full details