

Phone: 1-800-809-1265 ext 105 Fax: 1-888-417-3658

MRN:	 	 	
DOB:			

Patient Name:	Height (inches):Weight (lbs):
2. Allergies:	
3. Diagnosis:   G12.21 Amyotrophic	: Lateral Sclerosis
Other ICD-10 Code:diagnosis descript	tion:
*Product information does re-medication:	not suggest premedication prior to infusion
	Property of Palmetto Infusion / CONFIDENTIAL Property of Palmetto Infusion
. Dose:	
, ,	Iministered per (2 consecutive 30 mg/100 ml IV bags)
over a	total of 60 minutes
•	itive days, followed by cessation for 14 days
	days, followed by cessation for 14 days formonths
pecial orders:	
	ilize the ADVERSE DRUG REACTION GUIDELINES
If adverse drug reaction occurs, uti	
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8. Fax updated supporting clinical MD notes with each order renewal or change in orders

Infusion order forms and Adverse Drug Reaction Guidelines are available at <u>www.palmettoinfusion.com</u>



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## **Guidelines for Prescribing Radicava® (edaravone)**

(Required documentation with all initial referrals)

Patier	ent Name: Referra	al Date:
<u> </u>	Include signed and completed <b>Plan of Treatment</b> . (MD must com (Infusion order forms & Standard Adverse Reactions orders are available at www.palr	•
	Include patient demographic information and insurance informati	On. (Copy of insurance cards if available)
<del>1 - 1</del>	Supporting clinical MD notes to include any past tried and/or fair outcomes or contraindications to conventional therapy. Include support diagnosis: EMG results, Nerve conduction studies, MRI Muscle biopsy results as available.  • Radicava® is indicated for the treatment of Amyotrophic Landscape in the streatment of Amyotrophic Landscape.	e any lab results and/or tests to results, Lumbar puncture or
	Other as requested:	

\*\* Warnings/Precautions: Hypersensitivity Reactions: Do not receive RADICAVA® if you are allergic to edaravone or any of the Ingredients. Redness, wheals, erythema multiforme and cases of anaphylaxis (urticaria, decreased blood pressure, and dyspnea) have been reported in spontaneous post marketing reports. Sulfite Allergic Reactions: RADICAVA® 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 allergic reactions, for example, asthma episodes, in certain people. Sulfite sensitivity can happen more often in people who have asthma than in people who do not have asthma. The most common side effects of RADICAVA include bruising (contusion), problems walking (gait disturbance), and headache. These are not all the possible side effects of RADICAVA®. See full prescribing information.

You may report side effects to MT Pharma America, Inc. at 1-888-292-0058 or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch.

Dosing and Administration: https://www.radicava.com/hcp/dosing-and-administration/#administering-radicava

Palmetto Infusion Services will complete insurance verification and submit all required clinical documentation to the patient's insurance company for eligibility. Our office will notify you if any further information is required. We will review financial responsibility with the patient and refer them to any available Co-pay assistance as required. Thank you for the referral.

Please fax all information to 1-888-417-3658 or call 1-800-809-1265 for assistance.