| Field Name | Field Description |
|-------------------------|--|
| Prior Authorization | Radicava |
| Group Description | |
| Drugs | Radicava, Radivaca ORS (edaravone) |
| | and any other newly marketed agent |
| | |
| | *** riluzole (Rilutek) is Preferred and does not require prior |
| Covered Uses | authorization*** Medically accepted indications are defined using the following |
| Covered Uses | sources: the Food and Drug Administration (FDA), Micromedex, the |
| | Drug Package Insert, and/or per the standard of care guidelines |
| Exclusion Criteria | N/A |
| Required Medical | |
| Information | See "other criteria" |
| Age Restrictions | N/A |
| Prescriber Restrictions | Prescriber must be a neurologist |
| Coverage Duration | If the criteria are met, requests will be approved for up to 6 month |
| | duration |
| Other Criteria | Initial Authorization: |
| | • Member must have a diagnosis of ALS |
| | • Member must have a documented baseline evaluation of |
| | functionality using the revised ALS functional rating scale |
| | $(ALSFRS-R)$ score ≥ 2 |
| | • Member's disease duration is 2 years or less |
| | • Member has a baseline forced vital capacity (FVC) of $\geq 80\%$ |
| | • Member has been on riluzole (Rilutek), is beginning |
| | therapy as an adjunct to treatment with Radicava, or |
| | provider has provided a medical reason why patient is unable to use riluzole |
| | |
| | • Dose is within FDA approved limits |
| | Reauthorization: |
| | Member is not ventilator-dependent |
| | • Provider documents clinical stabilization in symptoms (e.g. |
| | stabilization of ALSFRS-R score) |
| | • Dose is within FDA approved limits |
| Revision/Review Date | Madical Divertary/alinical variance and the second second second |
| 4/2024 | Medical Director/clinical reviewer must override criteria when, |
| | in his/her professional judgement, the requested item is medically |
| | necessary. |