



MO HealthNet PA Criteria Proposal

Medical Procedure Class:	DME RESPIRATORY ASSIST DEVICE (RAD) WITH BACK- UP RATE - E0471 (KJRR)	
Implementation Date:	5/1/08	
Prepared for:	MO HealthNet	
Prepared by:	ACS-Heritage Information Systems, Inc.	
New Criteria	Revision of Existing Criteria	
Executive Summary		
Purpose:	To allow a more consistent and streamlined process for authorization of RAD with back-up rate.	
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Why was this Issue Selected:	Senate Bill 577 passed by the 94 th General Assembly directs MO HealthNet to utilize an electronic web-based system to authorize Durable Medical Equipment using best medical evidence and care and treatment guidelines, consistent with national standards to verify medical need.	
Procedures subject to Pre- Certification	E0471: Respiratory assist device, bi-level pressure capability with backup rate feature, used with noninvasive interface, e.g. nasal mask or facial mask	
Setting & Population:	All MO HealthNet fee-for-service participants	
Data Sources:		⊠ MHN Consultants

Setting & Population

- Procedure Group for review: E0471 (KJRR)
- Age range: All MO HealthNet fee-for-service participants

Approval Criteria

- Patient has been re-evaluated no sooner than 61 days after initiation of RAD with back-up rate therapy.
- Medical record documents progress of symptoms noted during re-evaluation and patient benefits from use of the device.
- Medical record documents patient use of RAD with back-up rate an average of 4 hours per 24 hour period by time of re-evaluation.
- Provider record contains patient-signed and dated statement that patient is compliantly using device for at least 4 hours per 24 hour period and has used the device for at least 2 months.

Denial Criteria

- Patient is being re-evaluated less than 61 days after initiation of RAD with back-up rate therapy.
- Medical record does not document progress of symptoms noted during reevaluation, or patient does not benefit from use of the device.
- Medical record does not document patient use of RAD with back-up rate an average of 4 hours per 24 hour period by time of re-evaluation.
- Provider record does not contain patient-signed and dated statement that patient is compliantly using device for at least 4 hours per 24 hour period and has used the device for at least 2 months.

Approval Period

E0471 ST (KJRR): Months 4 – 12 (12 Month Approval)