University of Texas Medical Branch Center for Sleep Disorders Policy: PAP Assessment Effective Date: June 30, 2017 Revised Date: January 18, 2019 Campus: Galveston and Webster

PAP ASSESSMENT

PURPOSE

Completing a PAP assessment for patients prescribed PAP treatment by sleep facility staff will provide a measure to determine if patients have an adequate response to the prescribed treatment.

POLICY

It is the policy of the facility that all patients' prescribed PAP treatment by the sleep facility medical staff will be offered PAP assessment follow-up within 12 weeks of treatment initiation.

PROCEDURE

- **1.0** Each patient prescribed PAP treatment by facility medical staff will be offered a follow-up PAP assessment within 12 weeks of treatment initiation.
- **2.0** Patients will be contacted by telephone eight weeks after treatment initiation. Technical staff will gain the subjective response of the patient through a telephone questionnaire.
 - 2.1 Subjective response will be documented in the patient medical record
- **3.0** Device download information will be remote or manually downloaded after 8 weeks of treatment and documented in the patient medical record.
- **4.0** If the patient does not respond to the telephone inquiry, facility staff will attempt to contact the patient again by telephone and letter to complete a telephone questionnaire assessment.
 - 4.1 If the telephone questionnaire is not completed, staff will attempt to schedule an inperson appointment with the patient.
 - 4.2 Attempts to follow-up with the patient will be documented in the patient medical record.
- **5.0** If the patient reports a positive response to therapy and the device download confirms use and response to therapy, this will be documented in the patient record.
- **6.0** If the patient reports a negative response regarding therapy and/or the device download shows an inadequate response, facility staff will schedule an in-person appointment to discuss the response to therapy, which will include but not be limited to:
 - 6.1 Assessment of use of the device
 - 6.2 Assessment of intolerance or non-acceptance of the device
 - 6.3 Review the device download
 - 6.4 Review telephone questionnaire response/previous subjective response
 - 6.5 Review device patient interface
 - 6.6 Address other underlying causes of inadequate response