

**OLIVE VIEW-UCLA MEDICAL CENTER
RESPIRATORY CARE SERVICES – SLEEP MEDICINE
POLICY & PROCEDURE**

NUMBER: 11429

VERSION: 1

SUBJECT/TITLE: PAP ASSESSMENT

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

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.

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** Each patient prescribed CPAP will be offered a patient education session called CPAP Bootcamp, that will include a follow up visit or phone call within 1 month. The subjective response of the patient will be obtained through a questionnaire.
 - 2.1** Subjective response will be documented in the patient medical record
- 3.0** Device download information will be remotely downloaded at the time of the follow-up visit or phone call and documented in the patient medical record.
- 4.0** If the patient does not arrive for the follow-up visit or respond to the telephone appointment, facility staff will attempt to contact the patient again by telephone and letter to complete a telephone or in-person questionnaire assessment.

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5.0 Attempts to follow-up with the patient will be documented in the patient medical record.

6.0 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.

7.0 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:

7.1 Assessment of use of the device

7.2 Assessment of intolerance or non-acceptance of the device

7.3 Review the device download

7.4 Review telephone questionnaire response/previous subjective response

7.5 Review device patient interface

7.6 Address other underlying causes of inadequate response

References:	
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