

Policy References: [Local Coverage Determination Respiratory Assist Device \(RAD\) \(L33800\) and Policy Article \(A52512\)](#)

Documentation References: [Standard Documentation Requirements Policy Article \(PA\) A55426](#)

Central Sleep Apnea (CSA) or Complex Sleep Apnea (CompSA)

For a Respiratory Assist Device (RAD) to be covered, the treating clinician must complete the following items;

- [Standard Written Order \(SWO\)](#)
- Medical record documentation requirements (see below)
The treating clinician must fully document in the patient's medical record symptoms characteristic of sleep-associated hypoventilation, such as daytime hyper-somnolence, excessive fatigue, morning headache, cognitive dysfunction, dyspnea, etc.
- Prior to initiating therapy, a complete facility-based, attended polysomnogram (PSG) must be performed documenting both of the following:
 - The diagnosis of CSA or CompSA
 - Significant improvement of the sleep-associated hypoventilation with the use of a RAD device on the settings that will be prescribed for initial use at home, while breathing the beneficiary's prescribed FIO2
- [Medical record requirements after the first 90-days of RAD therapy](#)