

Prior Authorization Criteria: Weight Loss

Drug Name: Contrave (naltrexone HCl/bupropion HCl), Belviq (locaserin), Belviq XR (locaserin extended-release), Saxenda (liraglutide), Qsymia (Phentermine/Topiramate)

Date: Original 7-2018

| Drug Name: | Contrave (naltrexone HCl/bupropion HCl), Belviq (locaserin), Belviq XR (locaserin extended-release), Saxenda (liraglutide), Qsymia (Phentermine/Topiramate) |
|-------------------------------------|---|
| Prescriber | n/a |
| Restrictions: | |
| Age Restrictions: | Patient is at least 18 years of age. |
| Exclusion Criteria: | n/a |
| Required Medical Information: | Patient has tried and failed, or has contraindications to, <u>both</u> phentermine and Alli; and Patient will use Contrave/Belviq/Saxenda/Qysmia in combination with a nutritional reduced calorie diet and weight reduction program (documentation of both nutritional and weight reduction programs required) in addition to at least one of the following: Patient has BMI equal to, or above, 30 kg/m²; or Patient has BMI greater than or equal to 27 kg/m² and has at least one additional risk factor (e.g. Coronary heart disease, dyslipidemia, hypertension, type II diabetes mellitus, sleep apnea). |
| Renewal Criteria: | Patient has completed at least 3 months (12 weeks) of Belviq/Saxenda therapy, with demonstrated adherence, and lost at least 5 percent of baseline body weight. OR Patient has completed at least 4 months (16 weeks) of Contrave therapy, with demonstrated adherence, and lost at least 5 percent of baseline body weight. OR Patient has completed at least 3 months (12 weeks) of Qysmia 15mg/92mg therapy, with demonstrated adherence, and lost at least 5 percent of baseline body weight; OR Patient has completed at least 3 months (12 weeks) of Qysmia 7.5mg/46mg therapy, with demonstrated adherence, and lost at least 3 percent of baseline body weight; |
| Coverage Duration: | Initial: 3 months (12 weeks) if Belviq/Saxenda/Qysmia, 4 months if Contrave Continuation of therapy: 12 months |