

## PHARMACY PRE-AUTHORIZATION CRITERIA



<b>DRUG (S)</b>	Relistor (methylnaltrexone tablets)
<b>POLICY #</b>	14138
<b>INDICATIONS</b>	Relistor tablets are indicated for the treatment of opioid-induced constipation in patients in adults with chronic non-cancer pain. Use of Relistor beyond four months has not been studied.
<b>CRITERIA</b>	ConnectiCare requires Relistor to be medically necessary in patient who met all of the following criteria: <ul style="list-style-type: none"><li>• Patient is on a stable opioid regimen</li><li>• Patient has documented opioid-induced constipation</li><li>• Patient has an intolerance to, or treatment failure of at least three laxative therapies (i.e. senna, bisacodyl, polyethelene glycol, lactulose, phosphasoda enema)</li></ul>
<b>LIMITATIONS</b>	<ul style="list-style-type: none"><li>• If the above criteria are met authorization will be granted for 2 months. Subsequent authorization (an additional 2 months) will be based on physician documentation of efficacy.</li><li>• Authorization will be limited to a maximum of 4 months therapy- use of Relistor beyond four months has not been studied.</li></ul>
<b>REFERENCES</b>	<ol style="list-style-type: none"><li>1. Facts &amp; Comparisons Online</li><li>2. Relistor manufacturer's insert, Salix Pharmaceuticals, Bridgewater, NJ 08807</li></ol>
<b>P&amp;T REVIEW HISTORY</b>	6/08, 9/09, 9/10, 12/11, 10/12, 10/13, 10/14, 11/15, 11/16, 11/17, 11/18
<b>REVISION RECORD</b>	10/14, 11/16, 11/19 (removed injection, adopted EH Relistor medical policy)