

CHOLESTYRAMINE POWDER

Pre - PA Allowance

None

Prior-Approval Requirements

Diagnoses

Patient must have **ONE** of the following:

1. Primary hypercholesterolemia (elevated LDL cholesterol)
 - a. Inadequate treatment response to **ALL** of the following:
 - i. Diet and exercise
 - ii. High intensity HMG-CoA reductase
 - iii. Zetia
2. Pruritus associated with partial biliary obstruction
 - a. Inadequate treatment response to **ALL** of the following:
 - i. Colestipol
 - ii. Rifampin
 - iii. Opioid antagonist
 - iv. Sertraline

AND ALL of the following:

- a. Inadequate treatment response to the commercially available product
- b. The concentration of the final product doesn't exceed the maximum recommended daily dose of 24 grams of anhydrous cholestyramine resin
- c. **NO** history of complete biliary obstruction

Prior - Approval Limits

Duration 6 months

* PA is only applicable to cholestyramine bulk powder. All other formulations are excluded from this policy

Prior – Approval *Renewal* Requirements

Same as above

Prior – Approval *Renewal* Limits

Same as above