

05/01/2017

Dear SoonerCare Provider,

The purpose of this fax is to provide information regarding criteria recently established for the reimbursement of various **phosphate binder products**. You are receiving this fax because you recently prescribed or dispensed a phosphate binder product for SoonerCare member(s). **Effective 05/08/2017, Velphoro® chewable tablets, Auryxia™ tablets, and Fosrenol® 1,000mg chewable tablets, 750mg oral powder, and 1,000mg oral powder will require prior authorization. The following medications do not require prior authorization:** generic calcium acetate containing products, Phoslyra®, Renvela®, Renagel®, and Fosrenol® 500mg and 750mg tablets. The authorization criteria for reimbursement for the prior authorized medications is as follows:

**Velphoro® (Sucroferric Oxyhydroxide) and Auryxia™ (Ferric Citrate) Approval Criteria:**

1. A diagnosis of hyperphosphatemia in patients with chronic kidney disease (CKD) on dialysis; and
2. Documented trials of inadequate response to at least two of the phosphate binders available without a prior authorization or a patient-specific, clinically significant reason why the member cannot use a phosphate binder available without a prior authorization.
3. For Auryxia™, a quantity limit of 12 tablets per day will apply.

**Fosrenol® (Lanthanum Carbonate) 1,000mg Chewable Tablets, 750mg Oral Powder, and 1,000mg Oral Powder Approval Criteria:**

1. A diagnosis of hyperphosphatemia in patients with end stage renal disease (ESRD); and
2. Documented trials of inadequate response to at least two of the phosphate binders available without a prior authorization or a patient-specific, clinically significant reason why the member cannot use a phosphate binder available without a prior authorization; and
3. For the approval of Fosrenol® oral powder, a patient-specific, clinically significant reason why a special formulation is needed over a phosphate binder available without a prior authorization, such as Fosrenol® 500mg or 750mg chewable tablets which can be crushed, must be provided; and
4. For the approval of Fosrenol® 1,000mg chewable tablets, a patient-specific, clinically significant reason why the member cannot use a phosphate binder available without a prior authorization, such as Fosrenol® 500mg or 750mg chewable tablets, must be provided.

Please note all members receiving Velphoro®, Auryxia™, and Fosrenol® 1,000mg tablets, 750mg powder, and 1,000mg powder will require that a manual prior authorization be submitted by their prescriber. No grandfathering will be allowed. If a member requires use of one of these non-preferred products, prior authorization requests can be submitted for consideration to SoonerCare Pharmacy Services, including patient-specific, clinically significant supporting information for use of the requested medication in place of the preferred products.

Updated versions of prior authorization criteria for phosphate binder medications can be downloaded from [www.okhca.org/rx-pa](http://www.okhca.org/rx-pa), then clicking "Chelating/Binding Agents". Prior authorization request forms can be found online at [www.okhca.org/forms](http://www.okhca.org/forms) (PHARM-04).

Thank you for the services you provide to Oklahomans insured by SoonerCare!