## **Potassium Binders**

Point-of-Sale (POS) edits are safety limitations that are automatically verified through computer programming at the time that a prescription claim is submitted at the pharmacy. These edits can be applied to *any* medication, whether or not it is listed in the Preferred Drug List / Non-Preferred Drug List (PDL/NPDL). The first section of this document is organized to follow the order of the therapeutic classes in the PDL/NPDL and explains the POS edits for those medications.

## **POS Abbreviations**

<b>AL</b> – Age Limit	<b>DD</b> – Drug-Drug Interaction	MD – Maximum Dose Limit	TD - Therapeutic Duplication
<b>BH</b> – Behavioral Health Clinical Authorization for Children Younger than 7 Years of Age	<b>DS</b> Maximum Days' Supply Allowed	<b>PR</b> – Enrollment in a Physician- Supervised Program Required	UN – Drug Use Not Warranted
<b>BY</b> – Diagnosis Codes Bypass Some Requirements	<b>DT</b> – Duration of Therapy Limit	PU – Prior Use of Other Medication is Required	X – Prescriber Must Have 'X' DEA Number
CL – Additional Clinical Information is Required	<b>DX</b> – Diagnosis Code Requirement	QL – Quantity Limit	<b>YQ</b> – Yearly Quantity Limit
CU – Concurrent Use with Other Medication is Restricted	ER – Early Refill	RX – Specific Prescription Requirement	

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## **POS Edits**

**CL** – Additional clinical information (prescriber specialty, diagnosis, etc.) is required for patiromer (Veltassa®) and sodium zirconium eyclosilicate (Lokelma®).

**QL** – Patiromer (Veltassa®) has a quantity limit of 30 packets per 30 days.

Revision / Date	Implementation Date	
Created POS Document	July 2021	
Removed clinical requirement for Lokelma / May 2023	<u>July 2023</u>	