

## Spinal Muscular Atrophy

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>DS</b> – Maximum Days’ Supply Allowed	<b>PU</b> – Prior Use of Other Medication is Required
<b>BH</b> – Behavioral Health Clinical Authorization for Children Younger than 7 Years of Age	<b>DT</b> – Duration of Therapy Limit	<b>QL</b> – Quantity Limit
<b>BY</b> – Diagnosis Codes Bypass Some Requirements	<b>DX</b> – Diagnosis Code Requirement	<b>RX</b> – Specific Prescription Requirement
<b>CL</b> – Additional Clinical Information is Required	<b>ER</b> – Early Refill	<b>TD</b> – Therapeutic Duplication
<b>CU</b> – Concurrent Use with Other Medication is Restricted	<b>MD</b> – Maximum Dose Limit	<b>YQ</b> – Yearly Quantity Limit
<b>DD</b> – Drug-Drug Interaction	<b>MME</b> – Maximum Morphine Milligram Equivalent is Restricted	

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POS Edits		
CL – Additional clinical information (diagnosis, etc.) is required for all spinal muscular atrophy agents.		
<del>QL</del> <b>QL</b> – These agents are limited to a maximum quantity listed in the chart to the right. <del>Risdiplam (Evrysdi™) is limited to a maximum quantity of 160ml (2-80ml bottles) every 24 days.</del>	<u>Generic (Brand Example)</u>	<u>Quantity Limit</u>
	<u>Risdiplam (Evrysdi™) oral solution</u>	<u>160ml (2-80ml bottles) every 24 days</u>
	<u>Risdiplam (Evrysdi™) tablet</u>	<u>1 tablet per day</u>

Revision / Date	Implementation Date
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## Spinal Muscular Atrophy

Created POS Document / May 2021 (Evrysdi®)	January 2021
Formatting changes / August 2023 (Evrysdi®)	October 2023
Combined all spinal muscular atrophy agents / October 2023	January 2024
<u>Added quantity limit for Evrysdi® tablet formulation / March 2025</u>	<u>August 2025</u>