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| --- | --- | --- | --- |
| **Criteria Title** | GH Receptor Antagonist | | |
| **Criteria Subtitle** | Somavert (pegvisomant) | | |
| **Approval Level** | GCNSeqNo | | |
| **Products**   |  |  | | --- | --- | | Preferred |  | | Non-Preferred | X | | Brand |  | | Generic |  | | Other |  | | Drug Name | Corresponding Code(s) | Type of Code (GCNSeqNo, HICL, NDC) |
| SOMAVERT | 051916 | GCNSeqNo |
| SOMAVERT | 051917 | GCNSeqNo |
| SOMAVERT | 051918 | GCNSeqNo |
| SOMAVERT | 072769 | GCNSeqNo |
| SOMAVERT | 072770 | GCNSeqNo |

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| --- | --- | --- | --- | --- | --- | --- |
| **Sequence Number** | **Question ID** | **Default Next Question ID** | **Question Type** | **Question Text** | **Choice Text** | **Next Question ID** |
| 1 | 0999 |  | Select | Is the patient new to therapy (initial authorization request) or continuing therapy (re-authorization request)? | New Start (initial authorization request) | 1000 |
| Continuation (re-authorization request) | 2000 |
| 2 | 1000 |  | Select | Is this request being prescribed in accordance with Food and Drug Administration (FDA) approved labeling? | Y | 1001 |
| N | 1235 |
| 3 | 1001 |  | Select and Free Text | Has the patient had an inadequate clinical response or contraindication to other therapies?  If yes, please submit documentation. | Y | 1002 |
| N | 1235 |
| 4 | 1002 |  | Select and Free Text | Has the provider submitted documentation of the patient’s  baseline liver function tests (LFTs)?  If yes, please submit documentation. | Y | END (Pending Manual Review) |
| N | 1235 |
| 5 | 2000 |  | Select and Free Text | Has the provider submitted documentation of the patient’s liver function tests (LFTs)?    If yes, please submit documentation. | Y | END (Pending Manual Review) |
| N | 1235 |
| 6 | 1235 |  | Free Text | Please provide the rationale for the medication being requested. | END (Pending Manual Review) | |

LENGTH OF AUTHORIZATIONS: Initial authorizations will be for 180 days; subsequent authorizations will be for 180 days and require documentation of LFTs

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| **Last Approved** | 4/10/2023 |
| **Other** |  |