**Infectious Disease Agents: Antibiotics – Cephalosporins**

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| Criteria 1 | NP- Cephalexin 750mg, Cefpodoxime, Cefixime Cap |
| Criteria 2 | NP with AR- Cefixime Susp, Suprax Chewable Tab |
| Criteria 3 | P with AR- Cefaclor Susp, Cefprozil Susp |

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| **Criteria Title** | Infectious Disease Agents: Antibiotics – Cephalosporins | | |
| **Criteria Subtitle** | Non-Preferred Products | | |
| **Approval Level** | GCNSeqNo | | |
| **Products**   |  |  | | --- | --- | | Preferred |  | | Non-Preferred | X | | Brand |  | | Generic |  | | Other |  | | Drug Name | Corresponding Code (s) | Type of Code (GCNSeqNo, HICL, NDC) |
| CEPHALEXIN750 mg | 060783 | GCNSeqNo |
| CEFPODOXIME | 016929 | GCNSeqNo |
| CEFPODOXIME | 016930 | GCNSeqNo |
| CEFPODOXIME | 016931 | GCNSeqNo |
| CEFPODOXIME | 016932 | GCNSeqNo |
| CEFIXIME CAP | 028142 | GCNSeqNo |

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| **Sequence Number** | **Question ID** | **Default Next Question ID** | **Question Type** | **Question Text** | **Choice Text** | **Next Question ID** |
| 1 | 0996 |  | Select and Free Text | Does the patient have an infection that is caused by an organism resistant to **ALL** preferred antibiotics?    If yes, please provide documentation of the diagnosis and any culture and sensitivity reports. | Y | END (Approve x 30 days) |
| N | 0997 |
| 2 | 0997 |  | Select | Is the patient completing a course of therapy that was started in the hospital or other similar location or was started before Medicaid eligibility?  Please note: only the remaining course will be authorized. | Y | END (Approve x 30 days) |
| N | 0998 |
| 3 | 0998 |  | Select | Is the patient new to therapy (initial authorization request) or continuing therapy (re-authorization request)? | New Start (initial authorization request) | 0999 |
| Continuation (re-authorization request) | 1234 |
| 4 | 0999 |  | Select | Is this request being prescribed in accordance with Food and Drug Administration (FDA) approved labeling? | Y | 1000 |
| N | 1235 |
| 5 | 1000 |  | Select and Free Text | Has the patient had an inadequate clinical response of at least 3 days with at least one preferred antibiotic?  The preferred alternatives may include the following: Cefadroxil, Cephalexin 250, 500mg, Cefaclor, Cefprozil, Cefuroxime, Cefdinir.  If yes, please submit the medication trials and dates. | Y | 1002 |
| N | 1001 |
| 6 | 1001 |  | Select and Free Text | Has the provider submitted documentation of medical necessity beyond convenience for why the patient cannot be changed to a preferred drug (i.e., allergies, drug-drug interactions, contraindications, or intolerances)?  If yes, please submit the medication name and reason for inability to use. | Y | 1002 |
| N | 1236 |
| 7 | 1002 |  | Select | Is the request for any of the following:  1) a nonsolid oral dosage formulation  2) a non-preferred extended release formulation  3) a non-preferred brand name that has a preferred generic product | Y | 1003 |
| N | END (Approve x 30 days) |
| 8 | 1003 |  | Select and Free Text | Has the provider submitted documentation of medical necessity for the requested product (i.e. medical reasons for why the patient cannot be changed to a solid oral dosage formulation, inadequate clinical response with a product’s immediate release formulation, or inadequate clinical response or allergy of two or more generic labelers)? | Y | END (Approve x 30 days) |
| N | 1235 |
| 9 | 1234 |  | Select and Free Text | Has the provider submitted documentation of the patient’s clinical response to treatment, ongoing safety monitoring, AND medical necessity for continued use? | Y | END (Approve x 30 days) |
| N | 1235 |
| 10 | 1235 |  | Free Text | Please provide the rationale for the medication being requested. | END (Pending Manual Review) | |
| 11 | 1236 |  | Free Text | Please explain the reason(s) why the patient is unable to use medications not requiring prior approval. | END (Pending Manual Review) | |

LENGTH OF AUTHORIZATIONS: Based on indication

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| **Last Approved** | 5/5/2023 |
| **Other** |  |

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| **Criteria Title** | Infectious Disease Agents: Antibiotics – Cephalosporins | | |
| **Criteria Subtitle** | Cefixime Susp, Suprax Chewable Tab | | |
| **Approval Level** | GCNSeqNo | | |
| **Products**   |  |  | | --- | --- | | Preferred |  | | Non-Preferred | X | | Brand |  | | Generic |  | | Other |  | | Drug Name | Corresponding Code (s) | Type of Code (GCNSeqNo, HICL, NDC) |
| CEFIXIME SUSP | 009182 | GCNSeqNo |
| CEFIXIME SUSP | 044428 | GCNSeqNo |
| SUPRAX CHEWABLE TAB | 070122 | GCNSeqNo |
| SUPRAX CHEWABLE TAB | 070123 | GCNSeqNo |

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| **Sequence Number** | **Question ID** | **Default Next Question ID** | **Question Type** | **Question Text** | **Choice Text** | **Next Question ID** |
| 1 | 0996 |  | Select and Free Text | Does the patient have an infection that is caused by an organism resistant to **ALL** preferred antibiotics?    If yes, please provide documentation of the diagnosis and any culture and sensitivity reports. | Y | END (Approve x 30 days) |
| N | 0997 |
| 2 | 0997 |  | Select | Is the patient completing a course of therapy that was started in the hospital or other similar location or was started before Medicaid eligibility?  Please note: only the remaining course will be authorized. | Y | END (Approve x 30 days) |
| N | 0998 |
| 3 | 0998 |  | Select | Is the patient new to therapy (initial authorization request) or continuing therapy (re-authorization request)? | New Start (initial authorization request) | 0999 |
| Continuation (re-authorization request) | 1233 |
| 4 | 0999 |  | Select | Is this request being prescribed in accordance with Food and Drug Administration (FDA) approved labeling? | Y | 1000 |
| N | 1235 |
| 5 | 1000 |  | Select and Free Text | Has the patient had an inadequate clinical response of at least 3 days with at least one preferred antibiotic?  The preferred alternatives may include the following: Cefadroxil, Cephalexin 250, 500mg, Cefaclor, Cefprozil, Cefuroxime, Cefdinir.  If yes, please submit the medication trials and dates. | Y | 1002 |
| N | 1001 |
| 6 | 1001 |  | Select and Free Text | Has the provider submitted documentation of medical necessity beyond convenience for why the patient cannot be changed to a preferred drug (i.e., allergies, drug-drug interactions, contraindications, or intolerances)?  If yes, please submit the medication name and reason for inability to use. | Y | 1002 |
| N | 1236 |
| 7 | 1002 |  | Select | Is the request for any of the following:  1) a nonsolid oral dosage formulation  2) a non-preferred extended release formulation  3) a non-preferred brand name that has a preferred generic product | Y | 1003 |
| N | 1004 |
| 8 | 1003 |  | Select and Free Text | Has the provider submitted documentation of medical necessity for the requested product (i.e. medical reasons for why the patient cannot be changed to a solid oral dosage formulation, inadequate clinical response with a product’s immediate release formulation, or inadequate clinical response or allergy of two or more generic labelers)? | Y | 1004 |
| N | 1235 |
| 9 | 1004 |  | Select | Is the patient 12 years of age and older? | Y | 1235 |
| N | END (Approve x 30 days) |
| 10 | 1233 |  | Select and Free Text | Has the provider submitted documentation of the patient’s clinical response to treatment, ongoing safety monitoring, AND medical necessity for continued use? | Y | 1234 |
| N | 1235 |
| 11 | 1234 |  | Select | Is the patient 12 years of age and older? | Y | 1235 |
| N | END (Approve x 30 days) |
| 12 | 1235 |  | Free Text | Please provide the rationale for the medication being requested. | END (Pending Manual Review) | |
| 13 | 1236 |  | Free Text | Please explain the reason(s) why the patient is unable to use medications not requiring prior approval. | END (Pending Manual Review) | |

LENGTH OF AUTHORIZATIONS: Based on indication

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| **Last Approved** | 5/5/2023 |
| **Other** |  |

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| **Criteria Title** | Infectious Disease Agents: Antibiotics – Cephalosporins | | |
| **Criteria Subtitle** | Cefaclor Susp, Cefprozil Susp | | |
| **Approval Level** | GCNSeqNo | | |
| **Products**   |  |  | | --- | --- | | Preferred | X | | Non-Preferred |  | | Brand |  | | Generic |  | | Other |  | | Drug Name | Corresponding Code (s) | Type of Code (GCNSeqNo, HICL, NDC) |
| CEFACLOR SUSP | 009106 | GCNSeqNo |
| CEFACLOR SUSP | 009108 | GCNSeqNo |
| CEFACLOR SUSP | 009109 | GCNSeqNo |
| CEFPROZIL SUSP | 016582 | GCNSeqNo |
| CEFPROZIL SUSP | 016583 | GCNSeqNo |

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| **Sequence Number** | **Question ID** | **Default Next Question ID** | **Question Type** | **Question Text** | **Choice Text** | **Next Question ID** |
| 1 | 1233 |  | Select | Is the patient 12 years and older?  Please note: a PA is only required for patients 12 years and older. | Y | 1234 |
| N | 1236 |
| 2 | 1234 |  | Select and Free Text | Is the patient able to swallow a standard tablet and/or capsule formulation?  If no, please submit documentation. | Y | 1235 |
| N | END (Pending Manual Review) |
| 3 | 1235 |  | Free Text | Please provide the rationale for the medication being requested. | END (Pending Manual Review) | |
| 4 | 1236 |  | Free Text | A PA is not required for those younger than 12 years of age. | END (Pending Manual Review) | |

LENGTH OF AUTHORIZATIONS: Based on indication

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| **Last Approved** | 5/5/2023 |
| **Other** |  |