

(formerly WellFirst Health)

Coverage of any drug intervention discussed in a Medica prior authorization guideline is subject to the limitations and exclusions outlined in the member's benefit certificate or policy and applicable state and/or federal laws.

| 🛛 Commercial (Small & Large Group) | 🖾 ASO | 🛛 Exchange/ACA | | |
|------------------------------------|-------|----------------|--|--|
| Medicare Advantage (MAPD) | | | | |

| Site of Service | MB2206 |
|----------------------------------|--|
| Covered Service: | Yes |
| Prior Authorization Required: | Yes |
| Additional Information: | None |
| Medicare Policy: | Prior authorization is not required for Medicare Cost products (Dean Care Gold) and Medicare Supplement (Select) when this drug is provided by participating providers. Prior authorization is required if a member has Medicare primary and the plan secondary coverage. This policy is not applicable to our Medicare Replacement products. |
| Wisconsin Medicaid Policy | Coverage of prescription drug benefits is administered by the Wisconsin Medicaid program. Coverage of medical drug benefits is administered by the Wisconsin Medicaid fee-for-service program. Medical drugs not paid on a fee-for-service basis by the Wisconsin Medicaid program are covered by the plan with no PA required. |

Program requirements:

- 1.0 The Site of Service program requirements will be administered as part of the existing prior authorization program.
 - 1.1 All drugs in the Site of Service program require prior authorization.
- 2.0 Requests for select specialty drugs as listed in the list in section 'Drugs in Scope' to be administered in a hospital outpatient setting may be directed to a preferred alternative site of care, such as home infusion provider or a physician office.
- 3.0 To prevent delay in care and allow adequate transition time for the Plan's members to an alternate infusion site, Site of Service program requirements will be waived for 60-



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90 days, depending upon the specific drug, after prior authorization approval so that members can transition to a different infusion site.

Drugs in Scope:

- 1.0 Select infused specialty medications included in the Site of Service program are subject to change.
- 2.0 Changes to the Drugs in Scope
 - 2.1 If currently available infused specialty medications are added to the Site of Service program medication list, prescribers will receive advanced notification per the terms of the provider contract with the Plan.

| HCPCS | Brand Name | Generic Name |
|-------|---------------|------------------------------------|
| J3262 | ACTEMRA | tocilizumab |
| J1931 | ALDURAZYME | laronidase |
| Q5121 | AVSOLA | infliximab-axxq |
| J0490 | BENLYSTA | belimumab |
| J1556 | BIVIGAM | Intravenous immune globulin (IVIG) |
| Q5139 | BKEMV | eculizumab-aeeb |
| J2329 | BRIUMVI | ublituximab-xiiy |
| J1786 | CEREZYME | imiglucerase |
| J2786 | CINQAIR | reslizumab |
| J7318 | CRYSVITA | burosumab |
| J1743 | ELAPRASE | idursulfase |
| J3060 | ELELYSO | taliglucerase alfa |
| J3380 | ENTYVIO | vedolizumab |
| Q5151 | EPYSQLI | eculizumab-aagh |
| J0180 | FABRAZYME | agalsidase beta |
| J1744 | FIRAZYR | icatibant |
| J1572 | FLEBOGAMMA | IVIG |
| J1569 | GAMMAGARD | IVIG |
| J1566 | GAMMAGARD S/D | IVIG |
| J1561 | GAMMAKED | IVIG |
| J1557 | GAMMAPLEX | IVIG |



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| J1561 | GAMUNEX | IVIG |
|-------|--------------------|------------------------------------|
| J1599 | IMMUNE GLOBULIN IV | IVIG |
| Q5103 | INFLECTRA | infliximab-dyyb |
| J1290 | KALBITOR | ecallantide |
| J2840 | KANUMA | sebelipase alfa |
| J2507 | KRYSTEXXA | pegloticase |
| J0202 | LEMTRADA | alemtuzumab |
| J0221 | LUMIZYME | alglucosidase alfa |
| J3397 | MEPSEVII | vestronidase alfa-vjbk |
| J1458 | NAGLAZYME | galsufase |
| J2182 | NUCALA | mepolizumab |
| J2350 | OCREVUS | ocrelizumab |
| J2351 | OCREVUS ZUNOVO | ocrelizumab and hyaluronidase-ocsq |
| J1568 | OCTAGAM | IVIG |
| J0129 | ORENCIA | abatacept |
| J1459 | PRIVIGEN | IVIG |
| J1301 | RADICAVA | edaravone |
| J1745 | REMICADE | infliximab |
| Q5104 | RENFLEXIS | infliximab-abda |
| J1602 | SIMPONI ARIA | golimumab |
| J1300 | SOLIRIS | eculizumab |
| J1746 | TROGARZO | ibalizumab-uiyk |
| J2323 | TYSABRI | natalizumab |
| J1303 | ULTOMIRIS | ravulizumab |
| J1322 | VIMIZIM | elosulfase alfa |
| J3385 | VPRIV | velaglucerase alfa |
| J2357 | XOLAIR | omalizumab |



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Exceptions:

- 1.0 Exceptions to the Site of Service program requirements are reviewed through the prior authorization process and may be granted on a case-by-case basis on medical necessity.
- 2.0 The administration of the infusion and injectable therapy referenced in this policy in a hospital outpatient setting is not considered medically necessary unless the below criteria are met:
 - 2.1 Hospital outpatient administration of infusion or injectable therapy is considered medically necessary for up to a 60 to 90 day period for members beginning a new treatment OR initial review of continuation of therapy
 - 2.2 An outpatient infusion or injectable therapy service in a hospital outpatient setting is considered medically necessary for the applicable validity period when any of the following are present:
 - 2.2.1 Potential changes in the member's clinical condition are such that immediate access to specific services of a hospital setting, having emergency resuscitation equipment and personnel, and inpatient admission or intensive care is necessary. For example, the member is at significant risk of sudden life-threatening changes in medical status based on clinical conditions including but not limited to:
 - 2.2.1.1 Intolerable fluid overload, including impaired or unstable renal function, or
 - 2.2.1.2 History of anaphylaxis to prior infusion therapy with a related pharmacologic or biologic age despite standard premedication, or
 - 2.2.1.3 Acute mental status/cognitive changes or physical impairment AND no home caregiver available; or
 - 2.2.1.4 Vascular access not stable; or
 - 2.2.1.5 Documented clinical history of cardiopulmonary conditions that may cause an increased risk of severe adverse react ions (including but not limited to thromboembolism, hypotension, seizures, aseptic meningitis syndrome, anaphylaxis, acute respiratory distress, pulmonary edema, apnea and transfusion associated lung disease); or
 - 2.2.1.6 The member does not have access to home infusion AND the nearest office based provider who can provide that service exceeds the travel distance to the currently-servicing hospital outpatient center.
 - 2.2.2 Home deemed unsafe environment for infusion (e.g. too many pets, esp. birds, aggressive dogs, etc.); or



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- 2.2.3 Member reasoning (e.g. often members don't want children exposed to the medication, etc.); or
- 2.2.4 Financial impact to member is high in a setting other than hospital outpatient center. These are reviewed on a case-by-case basis only.

| | Committee/Source | Date(s) |
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