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Subject: Daratumumab (Darzalex[®]) Infusion and Daratumumab-Hyaluronidase-fihj (Darzalex Faspro[™]) Injection

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Dosage/ Administration	Position Statement	Billing/Coding	Reimbursement	Program Exceptions	Definitions
Related Guidelines	Other	References	Updates		

DESCRIPTION:

Daratumumab (Darzalex) is an immunoglobulin G1 kappa (IgG1κ) human monoclonal antibody that binds to CD38, a transmembrane glycoprotein expressed on the surface of many hematopoietic cells and causes lysis of multiple myeloma cells. It was first approved by the FDA in November 2015 as monotherapy for the treatment of patients with multiple myeloma (MM) who have received at least three prior lines of therapy including a proteasome inhibitor (PI) and an immunomodulatory agent, or who are double refractory to a PI and an immunomodulatory agent. Daratumumab was previously granted orphan drug designation for the treatment of MM in May 2013. The indication for use was expanded in November 2016 for use in combination with lenalidomide and dexamethasone, or bortezomib and dexamethasone, for the treatment of patients with MM who have received at least one prior therapy. In June 2017, the FDA-approved indication was expanded again to include use, in combination with pomalidomide (Pomalyst) and dexamethasone, for the treatment of patients with MM who have received at least two prior therapies including lenalidomide and a proteasome inhibitor. In May 2018, the FDA approved the use of daratumumab in combination with bortezomib, melphalan and prednisone for the treatment of patients with newly diagnosed MM who are ineligible for autologous stem cell transplant. This was the first daratumumab treatment regimen FDA-approved for first line use in previously untreated patients. In June 2019, the FDA approved daratumumab in combination with lenalidomide and dexamethasone for patients with newly diagnosed multiple myeloma who are ineligible for autologous stem cell transplant. In September 2019, the FDA approved daratumumab in combination with bortezomib, thalidomide, and dexamethasone for patients with newly diagnosed multiple myeloma who are eligible for autologous stem cell transplant. In August 2020, the FDA

approved daratumumab in combination with carfilzomib and dexamethasone for patients with relapsed or refractory multiple myeloma who have received one to three prior lines of therapy.

In May 2020 a new subcutaneously administered version of daratumumab, daratumumab-hyaluronidase-fihj (Darzalex Faspro) was approved by the FDA for five of the eight MM indications approved for intravenous Darzalex. Since the initial approval, Darzalex Faspro has gained the three additional indications and now has the same eight MM indications as IV Darzalex [i.e., “in combination with bortezomib, thalidomide, and dexamethasone for patients with newly diagnosed MM who are eligible for autologous stem cell transplant” (January 2021), “in combination with pomalidomide and dexamethasone for adult patients with multiple myeloma who have received at least one prior line of therapy including lenalidomide and a proteasome inhibitor” (July 2021), and “in combination with carfilzomib and dexamethasone in patients with relapsed or refractory multiple myeloma who have received one to three prior lines of therapy” (December 2021)]. Also, in January 2021, a new indication for the treatment of adult patients with newly diagnosed light chain (AL) amyloidosis in combination with bortezomib, cyclophosphamide and dexamethasone was approved. The advantages of Darzalex Faspro (vs. Darzalex) are a shorter administration duration (a 3 to 5-min SQ injection vs. a 3 to 4-hour IV infusion), fewer systemic administration-related reactions (13% vs. 34%) and simplified flat dosing (1,800 mg SQ) vs. weight-based dosing (16 mg/kg IV).

The safety and efficacy of IV daratumumab, leading to the initial FDA-approval, were assessed in an open-label, multicenter, phase 2, single-arm study (n=106). An overall response rate (ORR) of 29.2% (n=31) was achieved with daratumumab monotherapy in patients with relapsed or refractory MM. Patients had received at least three prior therapies including a PI and an immunomodulatory agent or were double refractory to a PI and an immunomodulatory agent. A stringent complete response was reported in 2.8% (n=3), complete response (CR) in 0%, very good partial response (VGPR) in 9.4% (n=10), and partial response in 17% (n=18). The median time to response was 1 month, and median duration of response was 7.4 months. In this study, patients had received a median of five prior therapies, 97% of patients had refractory disease to the last therapy, and 80% of patients had received an autologous stem cell transplantation (ASCT). Prior therapies included bortezomib (99%), lenalidomide (99%), pomalidomide (63%) and carfilzomib (50%).

The National Comprehensive Cancer Network (NCCN) Guidelines for MM (Version 5.2022 – March 9, 2022) list four daratumumab triplet drug therapies [daratumumab + bortezomib + dexamethasone, daratumumab + carfilzomib (Kyprolis) + dexamethasone, daratumumab + lenalidomide (Revlimid) + dexamethasone, and daratumumab + pomalidomide (Pomalyst) + dexamethasone] under “Preferred Regimens for Early Relapses (1–3 prior therapies)” as category 1 recommendations for the treatment of previously treated MM. There are numerous other category 1 preferred regimens listed by NCCN. The pomalidomide regimen is listed under the subsection of “After two prior therapies including lenalidomide and a PI”. The triplet regimen with bortezomib is supported by an open-label, phase III trial (n=498) in which subjects were randomized to receive daratumumab + bortezomib + dexamethasone triplet therapy, or bortezomib + dexamethasone doublet therapy. Subjects were excluded if there was previous evidence of refractoriness or intolerance to bortezomib or another PI, like ixazomib and carfilzomib. The ORR was 82.9% with triplet therapy vs. 63.2% with doublet therapy (p<0.001). The rates of VGPR and CR were doubled with triplet vs. doublet therapy (59.2% vs. 29.1%, p<0.001, and 19.2% vs. 9%, p<0.001, respectively). Adverse effects were more common with triplet therapy with grade 3 or 4 thrombocytopenia and neutropenia occurring more frequently (45.3% vs. 32.9%, and 12.8% vs. 4.2%,

respectively). The triplet regimen with lenalidomide is supported by a phase III trial (n=569) in which subjects were randomized to receive daratumumab + lenalidomide + dexamethasone triplet therapy, or lenalidomide + dexamethasone doublet therapy. Subjects were excluded if there was previous evidence of refractoriness or intolerance to lenalidomide. The ORR was 92.9% with triplet therapy vs. 76.4% with doublet therapy (p<0.001). The CR was more than doubled with triplet vs. doublet therapy (43.1% vs. 19.2%, p<0.001). Grade 3 or 4 neutropenia occurred more frequently with triplet therapy (51.9% vs. 37%) and daratumumab-associated infusion-related reactions occurred in 47.7% of patients. The quadruplet regimen of daratumumab + cyclophosphamide + bortezomib + dexamethasone is listed under “Other Recommended Regimens for Early Relapses (1–3 prior therapies)” as a category 2A recommendation for the treatment of previously treated MM. Daratumumab monotherapy and the triplet regimen of daratumumab + selinexor (Xpovio) + dexamethasone are listed under “Useful In Certain Circumstances for Early Relapses (1–3 prior therapies)” as category 2A recommendations. The monotherapy regimen is listed under the subsection of “After at least three prior therapies, including a proteasome inhibitor and an immunomodulatory agent, or are double refractory to a proteasome inhibitor and an immunomodulatory agent”. The therapies for previously treated MM include a general footnote of “Regimens included under 1-3 prior therapies can also be used later in the disease course. Attempt should be made to use drugs/drug classes the patients have not been exposed to or exposed to >1 line prior”.

The NCCN Guidelines list the triplet regimen of daratumumab + lenalidomide + dexamethasone under “Preferred Regimens” as a category 1 recommended initial primary therapy for non-transplant candidates. The triplet regimen of bortezomib + lenalidomide + dexamethasone is the only other category 1 listed “Preferred Regimen” for initial primary therapy for non-transplant candidates. The triplet regimen is supported by an open-label phase III trial (MAIA, n=737) in which newly diagnosed MM patients ineligible for autologous stem cell transplant were randomized to receive daratumumab + lenalidomide + dexamethasone, or lenalidomide + dexamethasone. The daratumumab-containing regimen had a better 30-month progression free survival (PFS) rate (70.6% vs. 55.6%), and median PFS (not reached vs 31.9 months; HR, 0.56 [95% CI, 0.43 to 0.73]). The daratumumab-containing regimen also had a better ORR (92.9% vs. 81.3%) including a complete response rate or better (47.6% vs. 24.9%). Grade 3 or 4 adverse events occurring in a higher frequency with daratumumab included neutropenia (50% vs. 35.3%), any infection (32.1% vs. 23.3%), and pneumonia (13.7% vs. 7.9%). The NCCN lists the quadruplet regimen of daratumumab with bortezomib + melphalan + prednisone (VMP) under “Other Recommended Regimens” as a category 1 recommended initial primary therapy for non-transplant candidates. The quadruplet regimen is supported by an open-label phase III trial (ALCYONE, n=769) in which newly diagnosed MM patients ineligible for autologous stem cell transplant were randomized to receive daratumumab + VMP or VMP. The ORR was 90.9% with D-VMP vs. 73.9% with VMP (p<0.0001), and the D-VMP group showed an improvement in PFS (not reached vs. 18.1 months; HR=0.5, 95% CI: 0.38 to 0.65; p<0.0001). The most frequent adverse reactions (≥20% with at least 5% greater frequency in the D-VMP arm) were infusion reactions (28% vs. 0%), upper respiratory tract infection (48% vs. 28%), and peripheral edema (21% vs. 14%). The NCCN also lists the quadruplet regimen of daratumumab with cyclophosphamide + bortezomib + dexamethasone under “Other Recommended Regimens” as a category 2A recommended initial primary therapy for non-transplant candidates.

For transplant candidates, the NCCN lists the quadruplet regimen of daratumumab with bortezomib + lenalidomide + dexamethasone under “Other Recommended Regimens” as a category 2A recommended

initial primary therapy. Three quadruplet regimens of daratumumab with bortezomib + thalidomide + dexamethasone, daratumumab with carfilzomib + lenalidomide + dexamethasone, and daratumumab with cyclophosphamide + bortezomib + dexamethasone are listed under “Useful in Certain Circumstances” as category 2A recommendations for initial primary therapy for transplant candidates. The carfilzomib regimen includes a footnote stating, “Generally reserved for the treatment of aggressive MM”. The triplet regimen of bortezomib + lenalidomide + dexamethasone is the only category 1 recommended regimen listed under “Preferred Regimens” for transplant candidates.

Regarding the use of subcutaneous daratumumab-hyaluronidase (Darzalex Faspro), the NCCN Guidelines for MM include a footnote on all daratumumab-containing regimens of “Includes both daratumumab for intravenous infusion and daratumumab and hyaluronidase-fihj for subcutaneous injection. Daratumumab and hyaluronidase-fihj for subcutaneous injection has different dosing and administration instructions compared to daratumumab for intravenous infusion.” The NCCN Guidelines for Systemic Light Chain Amyloidosis (Version 1.2022 – June 29, 2021) list the quadruplet regimen of daratumumab-hyaluronidase + bortezomib + cyclophosphamide+ dexamethasone under “Preferred Regimen” as a category 1 recommendation for initial primary therapy for both transplant and non-transplant candidates. It is the only preferred regimen and the only category 1 recommendation for primary therapy. Daratumumab and daratumumab-hyaluronidase monotherapy are listed as category 2A recommended options for previously treated disease.

POSITION STATEMENT:

Initiation of daratumumab (Darzalex) or daratumumab-hyaluronidase (Darzalex Faspro) **meets the definition of medical necessity** when **ANY** of the following criteria is met (“1”, “2”, or “3”):

1. The member has a diagnosis of symptomatic multiple myeloma (MM), and **BOTH** of the following (“a” and “b”):
 - a. **ANY** of the following (“i”, “ii”, or “iii”)
 - i. Member has newly diagnosed MM, **AND** is eligible for an autologous stem cell transplant as determined by the treating physician, and meets **BOTH** of the following (“1” and “2”):
 1. Member has not previously received treatment for their MM
 2. Daratumumab or daratumumab-hyaluronidase will be used in combination with any of the following:
 - Quadruplet therapy with bortezomib, lenalidomide (Revlimid), and dexamethasone
 - Quadruplet therapy with bortezomib, thalidomide, and dexamethasone
 - Quadruplet therapy with bortezomib, cyclophosphamide, and dexamethasone
 - Quadruplet therapy with carfilzomib (Kyprolis), lenalidomide, and dexamethasone*
 - ii. Member has newly diagnosed MM, **AND** is ineligible for an autologous stem cell transplant as determined by the treating physician (i.e., a non-transplant candidate), and meets **BOTH** of the following (“1” and “2”):
 1. Member has not previously received treatment for their MM

**This regimen is reserved for the treatment of aggressive MM only*

2. Daratumumab or daratumumab-hyaluronidase will be used in **ANY** of the following regimens:
 - Triplet therapy with both lenalidomide and dexamethasone
 - Quadruplet therapy with bortezomib, melphalan and prednisone
 - Quadruplet therapy with bortezomib, cyclophosphamide and dexamethasone
- iii. Member has relapsed or refractory MM, and meets **BOTH** of the following (“1” and “2”):
 1. Member has previously received at least **ONE** appropriate prior line of therapy of adequate duration for the treatment of their MM
 2. Daratumumab or daratumumab-hyaluronidase will be used in **ANY** of the following regimens:
 - Monotherapy
 - Triplet therapy with both bortezomib and dexamethasone
 - Triplet therapy with both carfilzomib and dexamethasone
 - Triplet therapy with both lenalidomide and dexamethasone
 - Triplet therapy with both pomalidomide (Pomalyst) and dexamethasone, **AND** the member has received at least two appropriate prior therapies of adequate duration for the treatment of their MM that include an immunomodulatory agent (e.g., lenalidomide, thalidomide) and a proteasome inhibitor (e.g., bortezomib, carfilzomib, ixazomib)
 - Triplet therapy with both selinexor (Xpovio) and dexamethasone
 - Quadruplet therapy with bortezomib, cyclophosphamide, and dexamethasone
- b. The dosage of IV daratumumab does not exceed 16 mg/kg (based on actual body weight) or the dosage of SQ daratumumab-hyaluronidase does not exceed 1,800 mg daratumumab and 30,000 units hyaluronidase given more often than the following:
 - i. If used as monotherapy (one cycle is 28 days or 4 weeks):
 - Cycles 1 and 2 (weeks 1 to 8): Weekly (i.e., day 1, 8, 15, and 22 of each cycle) for 8 doses*
 - Cycles 3 to 6 (weeks 9 to 24): Every two weeks (i.e., day 1 and 15 of each cycle) for 8 doses
 - Cycle 7 and beyond (week 25 and beyond): Every four weeks (i.e., day 1 of each cycle)
 - ii. If used as quadruplet therapy in combination with bortezomib, lenalidomide, and dexamethasone (one cycle is 21 days or 3 weeks):
 - Induction therapy:
 - Cycle 1 and 4 (weeks 1 to 12): Weekly (i.e., day 1, 8, and 15 of each cycle) for 12 doses*
 - After cycle 4 (week 12): Stop for high-dose chemotherapy and autologous stem cell transplant

- Consolidation therapy: weekly for a total of 6 doses (weeks 1 to 6) upon re-initiation of treatment following autologous stem cell transplant
 - Maintenance therapy (daratumumab + lenalidomide): every 4 weeks following completion of consolidation therapy for up to 26 doses (2 years)
- iii. If used as quadruplet therapy in combination with bortezomib, thalidomide, and dexamethasone (one cycle is 28 days or 4 weeks):
- Induction therapy:
 - Cycle 1 and 2 (weeks 1 to 8): Weekly (i.e., day 1, 8, 15, and 22 of each cycle) for 8 doses*
 - Cycles 3 to 4 (weeks 9 to 16): Every 2 weeks (i.e., days 1 and 15 of each cycle) for 4 doses
 - After cycle 4 (week 16): Stop for high-dose chemotherapy and autologous stem cell transplant
 - Consolidation therapy: every 2 weeks for a total of 4 doses (weeks 1 to 8) upon re-initiation of treatment following autologous stem cell transplant
- iv. If used as quadruplet therapy in combination with carfilzomib, lenalidomide, and dexamethasone (one cycle is 28 days or 4 weeks):
- Induction therapy:
 - Cycle 1 and 2 (weeks 1 to 8): Weekly (i.e., day 1, 8, 15, and 22 of each cycle) for 8 doses*
 - Cycles 3 to 6 (weeks 9 to 24): Every 2 weeks (i.e., day 1 and 15 of each cycle) for 8 doses
 - Cycle 7 and 8 (week 25 to 32): Every 4 weeks (i.e., day 1 of each cycle) for 2 doses
 - After cycle 8 (week 32): Stop for high-dose chemotherapy and autologous stem cell transplant
- v. If used as quadruplet therapy in combination with cyclophosphamide, bortezomib, and dexamethasone for newly diagnosed, transplant-eligible members (one cycle is 28 days or 4 weeks):
- Induction therapy:
 - Cycle 1 and 2 (weeks 1 to 8): Weekly (i.e., day 1, 8, 15, and 22 of each cycle) for 8 doses*
 - Cycles 3 to 6 (weeks 9 to 24): Every 2 weeks (i.e., day 1 and 15 of each cycle) for 8 doses
 - Cycle 7 and 8 (week 25 to 32): Every 4 weeks (i.e., day 1 of each cycle) for 2 doses
 - After cycle 8 (week 32): Stop for high-dose chemotherapy and autologous stem cell transplant

- Maintenance therapy (as daratumumab monotherapy): every 4 weeks for up to 12 doses (weeks 1 to 48) upon re-initiation of treatment following autologous stem cell transplant
- vi. If used as quadruplet therapy in combination with cyclophosphamide, bortezomib, and dexamethasone for either newly diagnosed, transplant-ineligible members or members with relapsed or refractory disease (one cycle is 28 days or 4 weeks):
 - Cycles 1 to 2 (weeks 1 to 8): Weekly (i.e., day 1, 8, 15, and 22 of each cycle) for 8 doses*
 - Cycles 3 to 6 (weeks 9 to 24): Every 2 weeks (i.e., day 1 and 15 of each cycle) for 8 doses
 - Cycle 7 and beyond (week 25 and beyond): Every 4 weeks (i.e., day 1 of each cycle)
 - vii. If used as quadruplet therapy in combination with bortezomib, melphalan, and prednisone (one cycle is 42 days or 6 weeks):
 - Cycle 1 (weeks 1 to 6): Weekly for 6 doses*
 - Cycles 2 to 9 (weeks 7 to 54): Every 3 weekly (i.e., days 1 and 21 of each cycle) for 16 doses
 - After cycle 9 (week 55): Every 4 weeks
 - viii. If used as triplet therapy with bortezomib and dexamethasone (one cycle is 21 days or 3 weeks):
 - Cycles 1 to 3 (weeks 1 to 9): Weekly (i.e., days 1, 8, and 15 of each cycle) for 9 doses*
 - Cycles 4 to 9 (weeks 10 to 24): Every 3 weeks (i.e., day 1 of each cycle) for 5 doses
 - After cycle 9 (week 25): Every 4 weeks
 - ix. If used as triplet therapy with either carfilzomib, lenalidomide, pomalidomide, or selinexor, and dexamethasone (one cycle is 28 days or 4 weeks):
 - Cycles 1 to 2 (weeks 1 to 8): Weekly (i.e., day 1, 8, 15, and 22 of each cycle) for 8 doses*
 - Cycles 3 to 6 (weeks 9 to 24): Every 2 weeks (i.e., day 1 and 15 of each cycle) for 8 doses
 - Cycle 7 and beyond (week 25 and beyond): Every 4 weeks (i.e., day 1 of each cycle)

**If given IV, the first prescribed 16 mg/kg dose at Week 1 may be split over two consecutive days (i.e., 8 mg/kg IV on Day 1 and 8 mg/kg IV on Day 2)*

2. The member has a diagnosis of systemic light chain amyloidosis, and **BOTH** of the following (“a” and “b”):
 - a. **EITHER** of the following (“i” or “ii”):
 - i. **BOTH** of the following:
 - Daratumumab or daratumumab-hyaluronidase is being used in combination with bortezomib, cyclophosphamide, and dexamethasone

- Member has newly diagnosed disease, **OR** member has relapsed/refractory disease and was previously treated with the above four-drug regimen and was relapse-free at least two years
- ii. Member has relapsed/refractory disease, **AND** daratumumab or daratumumab-hyaluronidase is being used as monotherapy
- b. The member's dosage of IV daratumumab does not exceed 16 mg/kg (based on actual body weight) or the dosage of SQ daratumumab-hyaluronidase does not exceed 1,800 mg daratumumab and 30,000 units hyaluronidase given more often than the following:
 - Cycles 1 to 2 (weeks 1 to 8): Weekly (i.e., day 1, 8, 15, and 22 of each cycle) for 8 doses*
 - Cycles 3 to 6 (weeks 9 to 24): Every 2 weeks (i.e., day 1 and 15 of each cycle) for 8 doses
 - Cycle 7 and beyond (week 25 and beyond): Every four weeks (i.e., day 1 of each cycle)

**If given IV, the first prescribed 16 mg/kg dose at Week 1 may be split over two consecutive days (i.e., 8 mg/kg IV on Day 1 and 8 mg/kg IV on Day 2)*
3. Member has another FDA-approved or NCCN-supported diagnosis (not previously listed above), and **BOTH** of the following are met ("a" and "b"):
 - a. **EITHER** of the following ("i" or "ii"):
 - i. Member is diagnosed with a condition that is consistent with an indication listed in the product's FDA-approved prescribing information (or package insert) **AND** member meets any additional requirements listed in the "Indications and Usage" section of the FDA-approved prescribing information (or package insert)
 - ii. Indication **AND** usage is recognized in NCCN Drugs and Biologics Compendium as a Category 1 or 2A recommendation
 - b. The dosage of daratumumab or daratumumab-hyaluronidase does not exceed the maximum recommended in the FDA-approved prescribing information or the maximum recommended by the applicable NCCN guidelines for the diagnosis

Approval duration: 6 months (except for the following: quadruplet therapy with daratumumab or daratumumab-hyaluronidase + cyclophosphamide + bortezomib + dexamethasone for newly-diagnosed, transplant-eligible members with MM - approved for 32 weeks; quadruplet therapy with daratumumab or daratumumab-hyaluronidase + carfilzomib + lenalidomide + dexamethasone - approved for 32 weeks; quadruplet therapy with daratumumab or daratumumab-hyaluronidase + bortezomib + lenalidomide + dexamethasone – approved for 12 weeks; quadruplet therapy with daratumumab or daratumumab-hyaluronidase + bortezomib + thalidomide + dexamethasone – approved for 16 weeks)

Continuation of daratumumab (Darzalex) or daratumumab-hyaluronidase (Darzalex Faspro) **meets the definition of medical necessity** when **BOTH** of the following criteria are met ("1" and "2"):

1. An authorization or reauthorization for daratumumab or daratumumab-hyaluronidase has been previously approved by Florida Blue or another health plan in the past 2 years for the treatment of multiple myeloma, systemic light chain amyloidosis, or other FDA-approved or NCCN-supported diagnosis; **OR** the member previously met **ALL** indication-specific initiation criteria
2. **ANY** of the following based on the member's diagnosis ("a", "b", or "c"):

- a. Multiple myeloma, and **ALL** of the following (“i”, “ii”, and “iii”):
- i. Daratumumab or daratumumab-hyaluronidase is being used in **ANY** of the following regimens:
 1. Monotherapy
 2. Triplet therapy with both bortezomib and dexamethasone
 3. Triplet therapy with both carfilzomib and dexamethasone
 4. Triplet therapy with both lenalidomide and dexamethasone
 5. Triplet therapy with both pomalidomide and dexamethasone
 6. Triplet therapy with both selinexor and dexamethasone
 7. Quadruplet therapy with bortezomib, melphalan, and prednisone
 8. Quadruplet therapy with bortezomib, thalidomide, and dexamethasone
 9. Quadruplet therapy with bortezomib, lenalidomide, and dexamethasone
 10. Quadruplet therapy with bortezomib, cyclophosphamide, and dexamethasone
 11. Quadruplet therapy with carfilzomib, lenalidomide, and dexamethasone
 - ii. The member’s dosage of IV daratumumab does not exceed 16 mg/kg (based on actual body weight) or the dosage of SQ daratumumab-hyaluronidase does not exceed 1,800 mg daratumumab and 30,000 units hyaluronidase given more often than the following:
 1. If used as monotherapy (one cycle is 28 days or 4 weeks):
 - Cycles 1 to 2 (weeks 1 to 8): Weekly (i.e., day 1, 8, 15, and 22 of each cycle) for 8 doses
 - Cycles 3 to 6 (weeks 9 to 24): Every 2 weeks (i.e., day 1 and 15 of each cycle) for 8 doses
 - Cycle 7 and beyond (week 25 and beyond): Every four weeks (i.e., day 1 of each cycle)
 2. If used as quadruplet therapy in combination with bortezomib, lenalidomide, and dexamethasone (one cycle is 21 days or 3 weeks):
 - Induction therapy:
 - Cycle 1 and 4 (weeks 1 to 12): Weekly (i.e., day 1, 8, and 15 of each cycle) for 12 doses*
 - After cycle 4 (week 12): Stop for high-dose chemotherapy and autologous stem cell transplant
 - Consolidation therapy: weekly for a total of 6 doses (weeks 1 to 6) upon re-initiation of treatment following autologous stem cell transplant
 - Maintenance therapy (daratumumab + lenalidomide): every 4 weeks following completion of consolidation therapy for up to 26 doses (2 years)
 3. If used as quadruplet therapy in combination with bortezomib, melphalan and prednisone (one cycle is 42 days or 6 weeks):

- Cycle 1 (weeks 1 to 6): Weekly for 6 doses
 - Cycles 2 to 9 (weeks 7 to 54): Every 3 weeks (i.e., days 1 and 21 of each cycle) for 16 doses
 - After cycle 9 (week 55): Every 4 weeks
4. If used as quadruplet therapy in combination with bortezomib, thalidomide, and dexamethasone (one cycle is 28 days or 4 weeks):
- Induction therapy:
 - Cycle 1 and 2 (weeks 1 to 8): Weekly (i.e., day 1, 8, 15, and 22 of each cycle) for 8 doses*
 - Cycles 3 to 4 (weeks 9 to 16): Every 2 weeks (i.e., days 1 and 15 of each cycle) for 4 doses
 - After cycle 4 (week 16): Stop for high-dose chemotherapy and autologous stem cell transplant
 - Consolidation therapy: every 2 weeks for a total of 4 doses (weeks 1 to 8) upon re-initiation of treatment following autologous stem cell transplant
5. If used as quadruplet therapy in combination with carfilzomib, lenalidomide, and dexamethasone (one cycle is 28 days or 4 weeks):
- Induction therapy:
 - Cycle 1 and 2 (weeks 1 to 8): Weekly (i.e., day 1, 8, 15, and 22 of each cycle) for 8 doses*
 - Cycles 3 to 6 (weeks 9 to 24): Every 2 weeks (i.e., day 1 and 15 of each cycle) for 8 doses
 - Cycle 7 and 8 (week 25 to 32): Every 4 weeks (i.e., day 1 of each cycle) for 2 doses
 - After cycle 8 (week 32): Stop for high-dose chemotherapy and autologous stem cell transplant
6. If used as quadruplet therapy in combination with cyclophosphamide, bortezomib, and dexamethasone for newly diagnosed, transplant-eligible members (one cycle is 28 days or 4 weeks):
- Induction therapy:
 - Cycle 1 and 2 (weeks 1 to 8): Weekly (i.e., day 1, 8, 15, and 22 of each cycle) for 8 doses*
 - Cycles 3 to 6 (weeks 9 to 24): Every 2 weeks (i.e., day 1 and 15 of each cycle) for 8 doses
 - Cycle 7 and 8 (week 25 to 32): Every 4 weeks (i.e., day 1 of each cycle) for 2 doses
 - After cycle 8 (week 32): Stop for high-dose chemotherapy and autologous stem cell transplant

- Maintenance therapy (as daratumumab monotherapy): every 4 weeks for up to 12 doses (weeks 1 to 48) upon re-initiation of treatment following autologous stem cell transplant
7. If used as quadruplet therapy in combination with cyclophosphamide, bortezomib, and dexamethasone for either newly diagnosed, transplant-ineligible members or members with relapsed or refractory disease (one cycle is 28 days or 4 weeks):
 - Cycles 1 to 2 (weeks 1 to 8): Weekly (i.e., day 1, 8, 15, and 22 of each cycle) for 8 doses*
 - Cycles 3 to 6 (weeks 9 to 24): Every 2 weeks (i.e., day 1 and 15 of each cycle) for 8 doses
 - Cycle 7 and beyond (week 25 and beyond): Every 4 weeks (i.e., day 1 of each cycle)
 8. If used as triplet therapy with bortezomib and dexamethasone (one cycle is 21 days or 3 weeks):
 - Cycles 1 to 3 (weeks 1 to 9): Weekly (i.e., days 1, 8, and 15 of each cycle) for 9 doses
 - Cycles 4 to 9 (weeks 10 to 24): Every 3 weeks (i.e., day 1 of each cycle) for 5 doses
 - After cycle 9 (week 25): Every 4 weeks
 9. If used as triplet therapy with either carfilzomib, lenalidomide, pomalidomide, or selinexor, and dexamethasone (one cycle is 28 days or 4 weeks)
 - Cycles 1 to 2 (weeks 1 to 8): Weekly (i.e., day 1, 8, 15, and 22 of each cycle) for 8 doses
 - Cycles 3 to 6 (weeks 9 to 24): Every 2 weeks (i.e., day 1 and 15 of each cycle) for 8 doses
 - Cycle 7 and beyond (week 25): Every 4 weeks (i.e., day 1 of each cycle)
- iii. Member has not had disease progression during daratumumab or daratumumab-hyaluronidase treatment
- b. Systemic light chain amyloidosis (SLCA) and **ALL** of the following (“i”, “ii”, and “iii”):
- i. Daratumumab or daratumumab-hyaluronidase is being used as **EITHER** monotherapy **OR** in combination with bortezomib, cyclophosphamide and dexamethasone
 - ii. The member’s dosage of IV daratumumab does not exceed 16 mg/kg (based on actual body weight) or the dosage of SQ daratumumab-hyaluronidase does not exceed 1,800 mg daratumumab and 30,000 units hyaluronidase given more often than the following:
 - Cycles 1 to 2 (weeks 1 to 8): Weekly (i.e., day 1, 8, 15, and 22 of each cycle) for 8 doses
 - Cycles 3 to 6 (weeks 9 to 24): Every 2 weeks (i.e., day 1 and 15 of each cycle) for 8 doses
 - Cycle 7 and beyond (week 25 and beyond): Every four weeks (i.e., day 1 of each cycle)
 - If used for the treatment for newly diagnosed disease, the total duration of therapy does **NOT** exceed a maximum of 2 years
 - iii. Member has not had disease progression during daratumumab or daratumumab-hyaluronidase treatment

- c. Other FDA-approved or NCCN-supported diagnosis (not listed above or as an orphan indication), and **ALL** of the following (“i”, “ii”, and “iii”):
- i. The dosage of daratumumab or daratumumab-hyaluronidase does not exceed the maximum recommended in the FDA-approved prescribing information or the maximum recommended by the applicable NCCN guideline for the specific diagnosis
 - ii. Daratumumab or daratumumab-hyaluronidase is used in a treatment regimen in accordance with the FDA-approved prescribing information or applicable NCCN guideline recommendation for the diagnosis
 - iii. The member has had a beneficial response to treatment with daratumumab or daratumumab-hyaluronidase

Approval duration: 1 year (except for quadruplet therapy with daratumumab or daratumumab-hyaluronidase + bortezomib + thalidomide + dexamethasone; approve for 3 months to allow up to 4 total doses to be given in the consolidation therapy phase; quadruplet therapy with daratumumab or daratumumab-hyaluronidase + carfilzomib + lenalidomide + dexamethasone – allow up to 8 cycles of induction treatment before transplant only). For newly diagnosed light chain amyloidosis, use will not be approved beyond 2 years of total treatment.

DOSAGE/ADMINISTRATION:

THIS INFORMATION IS PROVIDED FOR INFORMATIONAL PURPOSES ONLY AND SHOULD NOT BE USED AS A SOURCE FOR MAKING PRESCRIBING OR OTHER MEDICAL DETERMINATIONS. PROVIDERS SHOULD REFER TO THE MANUFACTURER’S FULL PRESCRIBING INFORMATION FOR DOSAGE GUIDELINES AND OTHER INFORMATION RELATED TO THIS MEDICATION BEFORE MAKING ANY CLINICAL DECISIONS REGARDING ITS USAGE.

FDA-Approved

Darzalex

- In combination with lenalidomide and dexamethasone for the treatment of patients with newly diagnosed multiple myeloma who are ineligible for autologous stem cell transplant, and in patients with relapsed or refractory multiple myeloma who have received at least one prior therapy.
- In combination with bortezomib, melphalan and prednisone for the treatment of patients with newly diagnosed multiple myeloma who are ineligible for autologous stem cell transplant
- In combination with bortezomib, thalidomide, and dexamethasone for the treatment of patients with newly diagnosed multiple myeloma who are eligible for autologous stem cell transplant
- In combination with bortezomib and dexamethasone for the treatment of patients with multiple myeloma who have received at least one prior therapy
- In combination with carfilzomib and dexamethasone for the treatment of patients with relapsed or refractory multiple myeloma who have received one to three prior lines of therapy
- In combination with pomalidomide and dexamethasone for the treatment of patients with multiple myeloma who have received at least two prior therapies including lenalidomide and a proteasome inhibitor
- As monotherapy, for the treatment of patients with multiple myeloma who have received at least three prior lines of therapy including a proteasome inhibitor (PI) and an immunomodulatory agent or who are double refractory to a PI and an immunomodulatory agent

- Combination therapy with bortezomib, melphalan and prednisone (6-week cycle regimen)
 - The recommended dose is 16 mg/kg actual body weight administered as an IV infusion (after dilution) according to the following dosing schedule:
 - Weeks 1 to 6: Weekly (total of 6 doses)
 - Weeks 7 to 54: Every three weeks (total of 16 doses)
 - Week 55 onwards until disease progression: Every four weeks
- Monotherapy, and combination therapy with lenalidomide or pomalidomide and low-dose dexamethasone (4-week cycle regimen)
 - The recommended dose is 16 mg/kg actual body weight administered as an IV infusion (after dilution) according to the following dosing schedule:
 - Weeks 1 to 8: Weekly (total of 8 doses)
 - Weeks 9 to 24: Every two weeks (total of 8 doses)
 - Week 25 onwards until disease progression: Every four weeks
- Combination therapy with bortezomib, thalidomide, and dexamethasone (4-week cycle regimen)
 - The recommended dose is 16 mg/kg actual body weight administered as an IV infusion (after dilution) according to the following dosing schedule:
 - Induction treatment phase:
 - Weeks 1 to 8: Weekly (total of 8 doses)
 - Weeks 9 to 16: Every two weeks (total of 4 doses)
 - Stop for high-dose chemotherapy and autologous stem cell transplant
 - Consolidation treatment phase:
 - Weeks 1 to 8: Every two weeks (total of 4 doses) upon re-initiation of treatment following autologous stem cell transplant
- Combination therapy with bortezomib and dexamethasone (3-week cycle regimen)
 - The recommended dose is 16 mg/kg actual body weight administered as an IV infusion (after dilution) according to the following dosing schedule:
 - Weeks 1 to 9: Weekly (total of 9 doses)
 - Weeks 10 to 24: Every three weeks (total of 5 doses)
 - Week 25 onwards until disease progression: Every four weeks
- Combination therapy with carfilzomib and dexamethasone (4-week cycle regimen)
 - Week 1: 8 mg/kg on days 1 and 2 (total 2 doses)
 - Weeks 2 to 8: 16 mg/kg weekly (total of 7 doses)
 - Weeks 9 to 24: 16 mg/kg every two weeks (total of 8 doses)
 - Week 25 onwards until disease progression: 16 mg/kg every four weeks
- To facilitate administration, the first prescribed 16 mg/kg dose at Week 1 may be split over two consecutive days i.e. 8 mg/kg on Day 1 and Day 2 respectively.
- The initial infusion rate is 50 mL/hr with a maximum rate of 200 mL/hr. Only increase the rate in the absence of infusion reactions with the previous infusion. See the package insert for more detailed recommendations.

- Administer with an in-line, sterile, non-pyrogenic, low protein-binding polyethersulfone (PES) filter (pore size 0.22 or 0.2 micrometer).
- Pre-medicate patients with a corticosteroid, oral acetaminophen, and oral or IV antihistamine one to three hours prior to every infusion. See the full prescribing information for specific recommendations.
- For monotherapy, post-medicate patients with an oral corticosteroid on the first and second day after all infusions. For combination therapy, consider administering low-dose oral methylprednisolone (≤ 20 mg) or equivalent, the day after the infusion. However, if a background regimen-specific corticosteroid (e.g. dexamethasone) is administered the day after the infusion, additional post-infusion medications may not be needed.
- Initiate antiviral prophylaxis to prevent herpes zoster reactivation within 1 week of starting daratumumab and continue for 3 months following treatment.

Darzalex Faspro

- In combination with bortezomib, melphalan and prednisone for the treatment of patients with newly diagnosed multiple myeloma who are ineligible for autologous stem cell transplant
- In combination with lenalidomide and dexamethasone for the treatment of patients with newly diagnosed multiple myeloma who are ineligible for autologous stem cell transplant, and in patients with relapsed or refractory multiple myeloma who have received at least one prior therapy
- In combination with bortezomib, thalidomide, and dexamethasone for the treatment of patients with newly diagnosed multiple myeloma who are eligible for autologous stem cell transplant
- In combination with bortezomib and dexamethasone, for the treatment of patients with multiple myeloma who have received at least one prior therapy
- In combination with pomalidomide and dexamethasone in patients who have received at least one prior line of therapy including lenalidomide and a proteasome inhibitor
- In combination with carfilzomib and dexamethasone in patients with relapsed or refractory multiple myeloma who have received one to three prior lines of therapy
- As monotherapy, for the treatment of patients with multiple myeloma who have received at least three prior lines of therapy including a proteasome inhibitor (PI) and an immunomodulatory agent or who are double refractory to a PI and an immunomodulatory agent
- In combination with bortezomib, cyclophosphamide and dexamethasone for the treatment of patients with newly diagnosed light chain (AL) amyloidosis
- Combination therapy with bortezomib, melphalan and prednisone (6-week cycle regimen)
 - The recommended dose is 1,800 mg/30,000 units (1,800 mg daratumumab and 30,000 units hyaluronidase) administered subcutaneously over approximately 3 to 5 minutes according to the following dosing schedule:
 - Weeks 1 to 6: Weekly (total of 6 doses)
 - Weeks 7 to 54: Every three weeks (total of 16 doses)
 - Week 55 onwards until disease progression: Every four weeks
- Monotherapy, or combination therapy with carfilzomib and dexamethasone, lenalidomide and dexamethasone, or pomalidomide and dexamethasone (4-week cycle regimen)
 - The recommended dose is 1,800 mg/30,000 units (1,800 mg daratumumab and 30,000 units hyaluronidase) administered subcutaneously over approximately 3 to 5 minutes according to the following dosing schedule:
 - Weeks 1 to 8: Weekly (total of 8 doses)

- Weeks 9 to 24: Every two weeks (total of 8 doses)
 - Week 25 onwards until disease progression: Every four weeks
- Combination therapy with bortezomib, thalidomide, and dexamethasone (4-week cycle regimen)
 - The recommended dose is 1,800 mg/30,000 units (1,800 mg daratumumab and 30,000 units hyaluronidase) administered subcutaneously over approximately 3 to 5 minutes according to the following dosing schedule:
 - Induction treatment phase:
 - Weeks 1 to 8: Weekly (total of 8 doses)
 - Weeks 9 to 16: Every two weeks (total of 4 doses)
 - Stop for high-dose chemotherapy and autologous stem cell transplant
 - Consolidation treatment phase:
 - Weeks 1 to 8: Every two weeks (total of 4 doses) upon re-initiation of treatment following autologous stem cell transplant
- Combination therapy with bortezomib and dexamethasone (3-week cycle regimen)
 - The recommended dose is 1,800 mg/30,000 units (1,800 mg daratumumab and 30,000 units hyaluronidase) administered subcutaneously over approximately 3 to 5 minutes according to the following dosing schedule:
 - Weeks 1 to 9: Weekly (total of 9 doses)
 - Weeks 10 to 24: Every three weeks (total of 5 doses)
 - Week 25 onwards until disease progression: Every four weeks
- Combination therapy with bortezomib, cyclophosphamide and dexamethasone (4-week cycle regimen for light chain amyloidosis)
 - The recommended dose is 1,800 mg/30,000 units (1,800 mg daratumumab and 30,000 units hyaluronidase) administered subcutaneously over approximately 3 to 5 minutes according to the following dosing schedule:
 - Weeks 1 to 8: Weekly (total of 8 doses)
 - Weeks 9 to 24: Every two weeks (total of 8 doses)
 - Week 25 onwards until disease progression: Every four weeks
- Pre-medicate patients with a corticosteroid, oral acetaminophen, and oral or IV antihistamine one to three hours prior to every dose. See the full prescribing information for specific recommendations.
- For monotherapy, post-medicate patients with an oral corticosteroid on the first and second day after all infusions. For combination therapy, consider administering low-dose oral methylprednisolone (≤ 20 mg) or equivalent, the day after the infusion. However, if a background regimen-specific corticosteroid (e.g. dexamethasone) is administered the day after the infusion, additional post-infusion medications may not be needed. If the patient does not experience a major systemic administration-related reaction after the first 3 doses, consider discontinuing the administration of corticosteroids (excluding any background regimen-specific corticosteroid).
- Initiate antiviral prophylaxis to prevent herpes zoster reactivation within 1 week of starting daratumumab and continue for 3 months following treatment.

Dose Adjustments

- Dosage adjustments are not recommended for adverse reactions. Adverse reactions are addressed with treatment interruption and slower infusion rates.
- Renal impairment: no dosage adjustment is necessary for patients with pre-existing renal impairment.
- Hepatic impairment: no clinically significant differences in the pharmacokinetics of daratumumab as monotherapy or as combination therapy were observed for mild [total bilirubin 1 to 1.5 times upper limit of normal (ULN) or aspartate aminotransaminase (AST)>ULN] and moderate (total bilirubin 1.5 to 3 times ULN and any AST) hepatic impairment. The effect of severe (total bilirubin >3 times ULN and any AST) hepatic impairment on daratumumab pharmacokinetics is unknown.

Drug Availability

Darzalex

- 100 mg/5 mL and 400 mg/20 mL solution for injection in single-dose vials
- Store in a refrigerator at 2°C to 8°C (36°F to 46°F). Do not freeze or shake. Protect from light. This product contains no preservative.

Darzalex Faspro

- 1,800 mg of daratumumab and 30,000 units of hyaluronidase per 15 mL in a single-dose vial
- Store in a refrigerator at 2°C to 8°C (36°F to 46°F). Do not freeze or shake. Protect from light. This product contains no preservative.

PRECAUTIONS:

Boxed Warnings

Darzalex

- None

Darzalex Faspro

- None

Contraindications

Darzalex

- History of severe hypersensitivity to daratumumab or any of the components of the formulation

Darzalex Faspro

- History of severe hypersensitivity to daratumumab, hyaluronidase or any of the components of the formulation

Precautions/Warnings

Darzalex

- **Infusion Reactions:** Daratumumab can cause severe infusion reactions including anaphylactic reactions (e.g., bronchospasm, hypoxia, dyspnea, hypertension, tachycardia, headache, laryngeal edema and pulmonary edema). These reactions can be life-threatening and fatal outcomes have been reported. Signs and symptoms may include respiratory symptoms, such as nasal congestion, cough, throat irritation, as well as chills, vomiting and nausea. Less common symptoms were wheezing, allergic rhinitis, pyrexia, chest discomfort, pruritus, and hypotension. Approximately half of all patients experienced a reaction, most during the first infusion. Pre-medicate patients with antihistamines, antipyretics and corticosteroids. Frequently monitor patients during the entire infusion. Interrupt the infusion for infusion reactions of ANY severity. Permanently discontinue the infusion in case of life-threatening infusion reactions (Grade 4). For patients with Grade 1, 2, or 3 reactions, reduce the infusion rate when re-starting the infusion. Ocular adverse reactions, including acute myopia and narrowing of the anterior chamber angle due to ciliochoroidal effusions with potential for increased intraocular pressure or glaucoma, have occurred. If ocular symptoms occur, interrupt infusion and seek immediate ophthalmologic evaluation prior to restarting.
- **Interference with Serological Testing:** Daratumumab binds to CD38 on red blood cells (RBCs) and results in a positive Indirect Antiglobulin Test (Coombs test). Daratumumab-mediated positive indirect antiglobulin test may persist for up to 6 months after the last daratumumab infusion. Type and screen patients prior to starting treatment. Inform blood banks that a patient has received daratumumab.
- **Neutropenia:** Daratumumab may increase neutropenia induced by background therapy. Monitor complete blood cell counts periodically during treatment according to manufacturer's prescribing information for background therapies. Monitor patients with neutropenia for signs of infection. A dose delay may be required to allow recovery of neutrophils. No dose reduction is recommended. Consider supportive care with growth factors
- **Thrombocytopenia:** Daratumumab may increase thrombocytopenia induced by background therapy. Monitor complete blood cell counts periodically during treatment according to manufacturer's prescribing information for background therapies. A dose delay may be required to allow recovery of platelets. No dose reduction of DARZALEX is recommended. Consider supportive care with transfusions.
- **Interference with Determination of Complete Response:** Daratumumab is a human IgG kappa monoclonal antibody that can be detected on both, the serum protein electrophoresis (SPE) and immunofixation (IFE) assays used for the clinical monitoring of endogenous M-protein. This interference can impact the determination of complete response and of disease progression in some patients with IgG kappa myeloma protein.
- **Pregnancy:** Based on the mechanism of action, daratumumab can cause fetal harm when administered to a pregnant woman. Daratumumab may cause depletion of fetal immune cells and decreased bone density. Advise pregnant women of the potential risk to a fetus.

Darzalex Faspro

- **Hypersensitivity and Other Administration Reactions:** Both systemic administration-related reactions, including severe or life-threatening reactions, and local injection-site reactions can occur. Fatal reactions have been reported with daratumumab-containing products, including Darzalex Faspro. Permanently discontinue for life-threatening reactions
- **Cardiac Toxicity in Patients with Light Chain (AL) Amyloidosis:** Serious or fatal cardiac adverse reactions occurred in patients with light chain (AL) amyloidosis who received Darzalex Faspro in combination with bortezomib, cyclophosphamide and dexamethasone. Serious cardiac disorders occurred in 16% and fatal cardiac disorders occurred in 10% of patients. Patients with NYHA Class IIIA or Mayo Stage IIIA disease may be at greater risk. Patients with NYHA Class IIIB or IV disease were not studied. Monitor patients with cardiac involvement of light chain (AL) amyloidosis more frequently for cardiac adverse reactions and administer supportive care as appropriate.

- **Hypersensitivity and Other Administration Reactions:** Both systemic administration-related reactions, including severe or life-threatening reactions, and local injection-site reactions can occur. Fatal reactions have been reported with daratumumab-containing products, including Darzalex Faspro. In a pooled safety population, 9% of patients experienced a systemic administration-related reaction and 8% had an injection-site reaction. Pre-medicate patients with histamine-1 receptor antagonist, acetaminophen and corticosteroids. Monitor patients for systemic administration-related reactions, especially following the first and second injections. For anaphylactic reaction or life-threatening (Grade 4) administration-related reactions, immediately and permanently discontinue treatment. Consider administering corticosteroids and other medications after the administration of daratumumab depending on dosing regimen and medical history to minimize the risk of delayed (defined as occurring the day after administration) systemic administration-related reactions.
- **Interference with Serological Testing:** Daratumumab binds to CD38 on red blood cells (RBCs) and results in a positive Indirect Antiglobulin Test (Coombs test). Daratumumab-mediated positive indirect antiglobulin test may persist for up to 6 months after the last daratumumab infusion. Type and screen patients prior to starting treatment. Inform blood banks that a patient has received daratumumab.
- **Neutropenia:** Daratumumab may increase neutropenia induced by background therapy. Monitor complete blood cell counts periodically during treatment according to manufacturer's prescribing information for background therapies. Monitor patients with neutropenia for signs of infection. Consider withholding treatment until recovery of neutrophils. In lower body weight patients, higher rates of Grade 3-4 neutropenia were observed.
- **Thrombocytopenia:** Daratumumab may increase thrombocytopenia induced by background therapy. Monitor complete blood cell counts periodically during treatment according to manufacturer's prescribing information for background therapies. Consider withholding treatment until recovery of platelets.
- **Interference with Determination of Complete Response:** Daratumumab is a human IgG kappa monoclonal antibody that can be detected on both, the serum protein electrophoresis (SPE) and immunofixation (IFE) assays used for the clinical monitoring of endogenous M-protein. This interference can impact the determination of complete response and of disease progression in some patients with IgG kappa myeloma protein.
- **Pregnancy:** Based on the mechanism of action, daratumumab can cause fetal harm when administered to a pregnant woman. Daratumumab may cause depletion of fetal immune cells and decreased bone density. Advise pregnant women of the potential risk to a fetus. Advise females with reproductive potential to use effective contraception during treatment and for 3 months after the last dose.

BILLING/CODING INFORMATION:

The following codes may be used to describe:

HCPGS Coding

J9144	Injection, daratumumab, 10 mg and hyaluronidase-fihj
J9145	Injection, daratumumab, 10 mg

ICD-10 Diagnosis Codes That Support Medical Necessity

C90.00	Multiple myeloma not having achieved remission
C90.02	Multiple myeloma in relapse

C90.10	Plasma cell leukemia not having achieved remission
C90.12	Plasma cell leukemia in relapse
C90.20	Extramedullary plasmacytoma not having achieved remission
C90.22	Extramedullary plasmacytoma in relapse
C90.30	Solitary plasmacytoma not having achieved remission
C90.32	Solitary plasmacytoma in relapse
E85.81	Light chain (AL) amyloidosis
E85.89	Other amyloidosis
E85.9	Amyloidosis, unspecified

REIMBURSEMENT INFORMATION:

Refer to section entitled [POSITION STATEMENT](#).

PROGRAM EXCEPTIONS:

Federal Employee Program (FEP): Follow FEP guidelines.

State Account Organization (SAO): Follow SAO guidelines.

Medicare Part D: BCBSF has delegated to Prime Therapeutics authority to make coverage determinations for the Medicare Part D services referenced in this guideline.

Medicare Advantage: No National Coverage Determination (NCD) and/or Local Coverage Determination (LCD) were found at the time of the last guideline review date.

DEFINITIONS:

Heavy chain – the larger component of an immunoglobulin. There are five types: IgG, IgA, IgM, IgD, and IgE.

Immunoglobulins (a.k.a., antibodies) – proteins made by normal plasma cells that have an important role in fighting infection as part of the humoral immune response. Antibodies are composed of two heavy chains and two light chains that form a larger complex. Each plasma cell produces only one type of heavy chain and one type of light chain.

Light chain – the smaller component of an immunoglobulin. There are two types: kappa and lambda.

Minimal response (MR) (according to the Revised Uniform Response Criteria by the International Myeloma Working Group) – ALL of the following:

- ≥25% but ≤49% reduction of serum M-protein
- Reduction in 24-hour urine M-protein by 50% to 89%
- In addition, if present at baseline, 25% to 49% reduction in size of soft tissue plasmacytomas is also required

- No increase in size or number of lytic bone lesions (development of compression fractures does not exclude response).

Myeloma Protein (M-Protein) – a nonfunctional immunoglobulin protein or protein fragment produced by malignant plasma cells (or myeloma cells). Since myeloma cells are monoclonal, the M-proteins for a given patient are structurally identical. Both portions of an immunoglobulin (the heavy chain and light chain) can be found in the serum, while only light chains can be found in the urine.

Plasma cell - a fully differentiated B lymphocyte (a type of white blood cell) that is specialized for immunoglobulin production and is found primarily in bone marrow.

Primary refractory MM - patients who never achieve at least a MR to initial induction therapy and progress while on therapy

Progressive MM - at least a 25% increase from nadir in the serum M-protein (absolute increase must be ≥ 0.5 g/dL) or urine M-protein (absolute increase must be ≥ 200 mg/24 hours), or in the difference between involved and uninvolved serum-free light-chain (FLC) levels (with an abnormal FLC ratio and FLC difference >100 mg/L).

Relapsed and refractory MM - patients who never achieve at least a MR or who progress within 60 days of their last therapy

Serum free light chain assay (SFLCA) – a test that detects the amount of unbound or free light chains (i.e., not attached to a heavy chain) in the serum. Normal values - kappa: 3.3 to 19.4 mg/L, lambda: 5.71 to 26.3 mg/L, kappa/lambda ratio: 0.26–1.65 (0.37–3.1 if renal impairment).

Serum Protein Electrophoresis (SPEP) – a test that detects and quantifies the amount of M-protein in the serum. An serum M-protein of greater than 30 g/L is consistent with a diagnosis of MM.

RELATED GUIDELINES:

[Allogeneic Bone Marrow and Stem Cell Transplantation, 02-38240-01](#)

[Belantamab Mafodotin-blmf \(Blenrep\), 09-J3000-80](#)

[Bortezomib Injection, 09-J0000-92](#)

[Carfilzomib \(Kyprolis\) IV, 09-J1000-81](#)

[Chimeric Antigen Receptor \(CAR\) T-Cell Therapies, 09-J3000-94](#)

[Doxorubicin HCl Liposome \(Doxil\) IV, 09-J0000-91](#)

[Elotuzumab \(Empliciti\) IV, 09-J2000-50](#)

[Isatuximab \(Sarclisa\) Injection, 09-J3000-67](#)

[Ixazomib \(Ninlaro\), 09-J2000-51](#)

[Oral Oncology Medications, 09-J3000-65](#)

[Thalidomide \(Thalomid\) Capsules, 09-J1000-56](#)

OTHER:

None

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COMMITTEE APPROVAL:

This Medical Coverage Guideline (MCG) was approved by the Florida Blue Pharmacy Policy Committee on 06/08/22.

GUIDELINE UPDATE INFORMATION:

03/15/16	New Medical Coverage Guideline.
04/15/16	Revision to guideline consisting of clarifying the position statement.
07/01/16	Revision to guideline consisting of HCPCS code updates.
11/15/16	Revision to guideline consisting of updating the position statement, description section, and references based on new NCCN recommendations.
12/15/16	Revision to guideline consisting of updating the position statement with a new triplet regimen and updating the lab documentation requirements.
01/01/17	Revision to guideline consisting of updating the description section, dosage/administration, and references based on new FDA-approved indications. Added HCPCS code J9145.
02/16/17	Revision: Update to Position Statement.
07/15/17	Review and revision to guidelines consisting of updating the description section, position statement, and references.
09/15/17	Revision to guidelines consisting of updating the description section, position statement, dosage/administration section, and references based on new FDA-approved indication for use in combination with pomalidomide and dexamethasone.
07/15/18	Review and revision to guidelines consisting of updating the description section, position statement, dosage/administration, precautions, billing/coding information, and references.
07/15/19	Review and revision to guidelines consisting of updating the description section, position statement, dosage/administration, precautions, billing/coding information, and references.
08/15/19	Revision to guidelines consisting of updating the description section, position statement, dosage/administration section, and references based on new FDA-approved indication for first-line use in combination with lenalidomide and dexamethasone.
11/15/19	Revision to guidelines consisting of updating the description section, position statement, dosage/administration section, and references based on NCCN guideline updates and a new FDA-approved indication for first-line use in combination with bortezomib, thalidomide, and dexamethasone.

07/15/20	Review and revision to guidelines consisting of updating the description section, position statement, dosage/administration, precautions, billing/coding information, related guidelines, and references.
10/01/20	Revision: Added HCPCS code C9062 (for Darzalex Faspro).
10/15/20	Updated description, position statement, dosage/administration, and references based on new FDA-approved combination regimen with carfilzomib (Kyprolis) and dexamethasone and several new NCCN recommended combination regimens.
01/01/21	Revision: Added HCPCS code J9144 and deleted codes C9062 and J9999.
03/15/21	Updated description section, position statement, dosage/administration, precautions, and references based on an FDA-approved expanded MM indication and new indication for light chain amyloidosis.
07/15/21	Review and revision to guidelines consisting of updating the description section, position statement, related guidelines, and references.
09/15/21	Revision to guidelines consisting of updating the description section, dosage/administration section, precautions, and references based on a newly approved MM regimen for Darzalex Faspro.
01/15/22	Revision to guidelines consisting of updating the description section, dosage/administration section, and references based on a newly approved MM regimen for Darzalex Faspro.
07/15/22	Review and revision to guidelines consisting of updating the description section, position statement, precautions, billing/coding, and references.