

MEDICAL POLICY

POLICY TITLE	DRUG INFUSION SITE OF SERVICE
POLICY NUMBER	MP 3.016

CLINICAL BENEFIT	<input type="checkbox"/> MINIMIZE SAFETY RISK OR CONCERN. <input type="checkbox"/> MINIMIZE HARMFUL OR INEFFECTIVE INTERVENTIONS. <input type="checkbox"/> ASSURE APPROPRIATE LEVEL OF CARE. <input type="checkbox"/> ASSURE APPROPRIATE DURATION OF SERVICE FOR INTERVENTIONS. <input type="checkbox"/> ASSURE THAT RECOMMENDED MEDICAL PREREQUISITES HAVE BEEN MET. <input checked="" type="checkbox"/> ASSURE APPROPRIATE SITE OF TREATMENT OR SERVICE.
Effective Date:	12/1/2025

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I. POLICY

Note: An injectable medication must meet applicable medical necessity criteria for coverage. When coverage criteria are met for the injectable medication, this policy is used to determine the medical necessity of the requested site of care.

This policy applies to the following infusions administered by health care professionals:

- Abatacept; **and**
- Agalsidase beta; **and**
- Alemtuzumab; **and**
- Alglucosidase alfa; **and**
- Avalglucosidase alfa-ngpt; **and**
- Belimumab; **and**
- Benralizumab; **and**
- Burosumab-twza; **and**
- C1 esterase inhibitor (see coding section for applicable codes); **and**
- Certolizumab pegol; **and**
- Crizanlizumab-tmca; **and**
- Crovalimab-akkz (subcutaneous); **and**
- Denosumab (see coding section for applicable codes); **and**
- Eculizumab; **and**
- Efgartigimod alfa-fcab; **and**
- Efgartigimod alfa and hyaluronidase-qvfc; **and**
- Elosulfase alfa; **and**
- Eptinezumab-jjmr; **and**
- Evinacumab-dgnb; **and**
- Galsulfase; **and**
- Givosiran; **and**
- Golimumab; **and**
- Idursulfase; **and**

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- Imiglucerase; **and**
- Immune globulin intravenous (IVIG) (see coding section for applicable codes); **and**
- Immune globulin intravenous human-ifas; **and**
- Immune globulin intravenous human-slra; **and**
- Immune globulin intravenous, human-stwk; **and**
- Immune globulin subcutaneous (see coding section for applicable codes); **and**
- Immune globulin subcutaneous human-hipp; **and**
- Immune globulin subcutaneous human-klhw
- Inclisiran; **and**
- Inebilizumab-cdon; **and**
- Infliximab; **and**
- Infliximab-dyyb; **and**
- Infliximab-abda; **and**
- Infliximab-axxq; **and**
- Laronidase; **and**
- Lumasiran; **and**
- Mepolizumab; **and**
- Natalizumab; **and**
- Ocrelizumab; **and**
- Ocrelizumab and hyaluronidase-ocsq; **and**
- Olipudase alfa-rpcp; **and**
- Omalizumab; **and**
- Patisiran; **and**
- Pegunigalsidase alfa-iwxj; **and**
- Ravulizumab-cwvz; **and**
- Reslizumab; **and**
- Romosozumab-aqqg; **and**
- Rozanolixizumab-noli; **and**
- Sebelipase alfa; **and**
- Sutimlimab-jome; **and**
- Taliglucerase alfa; **and**
- Teprotumumab-trbw; **and**
- Tocilizumab; **and**
- Tocilizumab-bavi; **and**
- Ublituximab-xiyy; **and**
- Ustekinumab (subcutaneous); **and**
- Ustekinumab-auub (subcutaneous); **and**
- Ustekinumab-ttwe (subcutaneous); **and**
- Ustekinumab-aekn (subcutaneous); **and**
- Vedolizumab; **and**
- Velaglucerase alfa; **and**
- Velmanase alfa-tycv; **and**
- Vestronidase alfa-vjbk; **and**

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- Vutrisiran.

Infusion of a medication initiated in the hospital outpatient setting is subject to a one-time 30-day approval period to facilitate transition to a medically necessary alternative less intensive site of care. Alternative less intensive site of care facilities include:

- Non-hospital affiliated outpatient infusion (e.g., ambulatory infusion center or physician office); **or**
- Home infusion

Infusion of one of the listed medications administered in an alternative less intensive site of care facility (see definition above) when criteria for coverage of the medication are met is considered **medically necessary** unless both of the following criteria are met:

- There is not a non-hospital affiliated, outpatient infusion center within an acceptable distance to the patients' home. Refer to link below:
 - [PA Code 9.679 Access requirements in service areas](#); **or**
- There is not a non-hospital affiliated, outpatient infusion center within **20 miles** of the patient's home **and** the patient does not live in the state of Pennsylvania; **and**
- The member's home is not eligible for home infusion services (such as home is not within the service area or is deemed unsuitable for care by the home infusion provider).

Or ONE of the following;

Infusion of one of the listed medications in a hospital outpatient setting or at a hospital-affiliated infusion suite is considered **medically necessary** for an individual when there is clinical documentation of **ANY** one of the following:

- The patient is under the age of 18; **or**
- The prescribed medication has a site of care restriction for administration per the Food and Drug Administration (FDA) approved label; **or**
- Clinical documentation of a severe or potentially life-threatening adverse event during or following infusion of the prescribed drug, and the adverse event cannot be managed through pre-medication in the home or office setting; **or**
- There is clinical documentation of a significant comorbidity (e.g., cardiopulmonary disorder) or concerns regarding fluid overload status that precludes treatment at an alternative less intensive site of care; **or**
- Clinical documentation of unstable vascular access; **or**
- Clinical documentation of physical or cognitive impairments such that home infusion would present an unnecessary health risk; **or**
- Patients current condition requires monitoring that cannot be provided in a less intensive site of care; **or**
- Patient is concurrently being treated with another medication that must be administered in a hospital setting; **or**
- Initiating a new therapy; **or**
- Reinitiating therapy after being off therapy for at least six months.

When the criteria above are not met, infusion of one of the listed medications in a hospital outpatient setting or hospital-affiliated infusion suite is considered **not medically necessary**.

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Note: A hospital outpatient setting or a hospital-affiliated infusion suite is expected to have immediate access to specific services of a medical center/hospital setting, including having emergency resuscitation equipment and personnel (Advanced Cardiac Life Support (ACLS) protocol), emergency services, and inpatient admission or intensive care, if necessary.

Cross-References:

- For medical necessity criteria, refer to the specific medical injectable policy **MP 2.176 Self Administered Medications**

II. PRODUCT VARIATIONS

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This policy is only applicable to certain programs and products administered by Capital Blue Cross and subject to benefit variations as discussed in Section VI. Please see additional information below.

FEP PPO - Refer to FEP Medical Policy Manual. The FEP Medical Policy manual can be found at: <https://www.fepblue.org/benefit-plans/medical-policies-and-utilization-management-guidelines/medical-policies>.

III. DESCRIPTION/BACKGROUND

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This policy outlines the site of care for medication infusions. It provides the criteria used to determine the medical necessity of the site of care for delivery of infused medications.

IV. RATIONALE

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N/A

V. DEFINITIONS

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N/A

VI. DISCLAIMER

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Capital Blue Cross' medical policies are used to determine coverage for specific medical technologies, procedures, equipment, and services. These medical policies do not constitute medical advice and are subject to change as required by law or applicable clinical evidence from independent treatment guidelines. Treating providers are solely responsible for medical advice and treatment of members. These policies are not a guarantee of coverage or payment. Payment of claims is subject to a determination regarding the member's benefit program and eligibility on the date of service, and a determination that the services are medically necessary and appropriate. Final processing of a claim is based upon the terms of contract that applies to the members' benefit program, including benefit limitations and exclusions. If a provider or a member has a question concerning this medical policy, please contact Capital Blue Cross' Provider Services or Member Services.

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VII. CODING INFORMATION

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Note: This list of codes may not be all-inclusive, and codes are subject to change at any time. The identification of a code in this section does not denote coverage as coverage is determined by the terms of member benefit information. In addition, not all covered services are eligible for separate reimbursement.

Covered when medically necessary:

Procedure Codes
J0129, J0180, J0202, J0217, J0218, J0219, J0221, J0222, J0223, J0224, J0225, J0490, J0517, J0584, J0597, J0598, J0717, J0791, J0897, J1299, J1302, J1303, J1305, J1306, J1307, J1322, J1458, J1602, J1743, J1745, J1786, J1823, J1931, J2182, J2323, J2329, J2350, J2351, J2357, J2508, J2786, J2840, J3032, J3060, J3111, J3241, J3262, J3357, J3380, J3385, J3397, J9332, J9333, J9334, Q5103, Q5104, Q5121, Q5133, Q5137, Q9996, Q9998

Covered when medically necessary for IVIG products:

Procedure Codes
J1459, J1552, J1554, J1556, J1557, J1561, J1566, J1568, J1569, J1572, J1576

Covered when medically necessary for subcutaneous immune globulin products:

Procedure Codes
J1551, J1555, J1558, J1559, J1575

ICD-10-CM Diagnosis Code*	Description
See above	Please reference the medical policy specific to the drug to determine coverage and appropriate diagnosis codes.

VIII. REFERENCES

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2. American Academy of Allergy Asthma and Immunology. *Guidelines for the site of care for administration of IGIV therapy.* 2011 Dec. Polinski JM, Kowal MK, Gagnon M, et al. *Home infusion: safe clinically effective, patient preferred, and cost saving. Healthcare.* 2016.

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IX. POLICY HISTORY

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MP 3.016	12/08/2020 Minor Review. IVIG and infliximab products added. References Reviewed. Coding updated.
	01/31/2021 Minor Review. Multiple drugs added. References updated. Coding updated.
	02/10/2022 Minor Review. Multiple drugs added. References reviewed. Coding updated. Added new code J2356. Cross reference updated. Removal of GamaSTAN from policy.
	07/07/2022 Administrative Update. Removal of J2356
	09/26/2022 Minor Review. Addition of J2182 and J9332
	03/10/2023 Minor Review. Annual review. References updated. Addition of J0225 and J1302.
	08/04/2023 Minor Review. Addition of J0218, J0219, J0896, J1551, J1554, J2998. References updated. Alemtuzumab added to drug list as editorial update. Code has been on policy since adopted 2021.
	10/09/2023 Minor Review Addition of J0256, J0257, J2793, J0490, J2786, J0584, J1951, J0638, J1306, J9316, J3285, J3032
	12/13/2023 Administrative Update. Added J9334 eff 01/01/2024.
	01/04/2023 Administrative Update. Removed J9334 from policy

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03/26/2024 Minor Review. Addition of J1576, J0217, J2329, J2508, J9334, J9333. References updated.
10/17/2024 Administrative Update. Moved Velmanase alfa-tycv to correct place in alphabetical order. Corrected coding table by adding J1951 as Fensolvi is already in the policy statement. No change to intent.
03/13/2025 Administrative Update. Deleted code J1300 and replaced with new code J1299, eff 04/01/2025.
06/13/2025 Minor Review. Addition of J1307, J1552, J2351, Q5133, Q5137, Q9996, and Q9998. Removed codes J0256, J0257, J0596, J0638, J1951, J0896, J9316, J2998, J2793, J3245, J3285, J1599, 90281, 90283, 90284. Removed erroneous code J1591. Added specification around IVIG and SCIG products in drug list and coding tables.
09/10/2025 Administrative Update. Removed Benefit Variations Section and updated Disclaimer.

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