

**2025 Blue Cross and Blue Shield Service Benefit Plan - Standard and Basic Option**  
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**Accidental injury**

An injury caused by an external force or element such as a blow or fall that requires immediate medical attention, including animal bites and poisonings. Note: Injuries to the teeth while eating are **not** considered accidental injuries. Dental care for accidental injury is limited to dental treatment necessary to repair sound natural teeth.

**Admission**

The period from entry (admission) as an inpatient into a hospital (or other covered facility) until discharge. In counting days of inpatient care, the date of entry and the date of discharge count as the same day.

**Advanced care planning**

Receiving information on the types of life-sustaining treatments that are available, completing advance directives and other standard forms, and/or if you are diagnosed with a terminal illness and making decisions about the care you would want to receive if you become unable to speak for yourself.

**Agents**

Medications and other substances or products given by mouth, inhaled, placed on you, or injected in you to diagnose, evaluate, and/or treat your condition. Agents include medications and other substances or products necessary to perform tests such as bone scans, cardiac stress tests, CT scans, MRIs, PET scans, lung scans, and X-rays, as well as those injected into the joint.

**Assignment**

An authorization by the enrollee or spouse for us to issue payment of benefits directly to the provider. We reserve the right to pay you, the enrollee, directly for all covered services. Benefits provided under the contract are not assignable by the member to any person without express written approval of the Carrier, and in the absence of such approval, any such assignment shall be void. Your specific written consent for a designated authorized representative to act on your behalf to request reconsideration of a claim decision (or, for an urgent care claim, for a representative to act on your behalf without designation) does not constitute an Assignment. OPM's contract with us, based on federal statute and regulation, gives you a right to seek judicial review of OPM's final action on the denial of a health benefits claim but it does not provide you with authority to assign your right to file such a lawsuit to any other person or entity. Any agreement you enter into with another person or entity (such as a provider, or other individual or entity) authorizing that person or entity to bring a lawsuit against OPM, whether or not acting on your behalf, does not constitute an Assignment, is not a valid authorization under this contract, and is void.

Please visit [www.fepblue.org](http://www.fepblue.org) to obtain a valid authorization form.

**Assisted reproductive technology (ART)**

Reproductive services, testing, and treatments involving manipulation of eggs, sperm, and embryos to achieve pregnancy. In general, assisted reproductive technology (ART) procedures are used to retrieve eggs from an ovulating individual, combine them with sperm in the laboratory, and then implant the embryos or donate them to an individual capable of pregnancy.

**Biologic drug**

A complex drug or product that is manufactured in a living organism, or its components, that is used as a diagnostic, preventive or therapeutic agent.

**Biosimilar drug**

A U.S. FDA-approved biologic drug, which is considered highly similar to an original brand-name biologic drug, with no clinically meaningful differences from the original biologic drug in terms of safety, purity and potency.

**Biosimilar, interchangeable drug**

A U.S. FDA-approved biosimilar drug that may be automatically substituted for the original brand-name biologic drug.

**Calendar year**

January 1 through December 31 of the same year. For new enrollees, the calendar year begins on the effective date of their enrollment and ends on December 31 of the same year.

**Carrier**

The Blue Cross and Blue Shield Association, on behalf of the local Blue Cross and Blue Shield Plans.

**Clinical trials**

An approved clinical trial includes a phase I, phase II, phase III, or phase IV clinical trial that is conducted in relation to the prevention, detection, or treatment of cancer or other life-threatening disease or condition, and is either Federally funded; conducted under an investigational new drug application reviewed by the U.S. Food and Drug Administration (U.S. FDA); or is a drug trial that is exempt from the requirement of an investigational new drug application.

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