

Medical Policy:

Gazyva® (obinutuzumab) injection

POLICY NUMBER	LAST REVIEW	ORIGIN DATE
MG.MM.PH.147	March 1, 2024	

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EmblemHealth established the clinical review criteria based upon a review of currently available clinical information (including clinical outcome studies in the peer reviewed published medical literature, regulatory status of the technology, evidence-based guidelines of public health and health research agencies, evidence-based guidelines and positions of leading national health professional organizations, views of physicians practicing in relevant clinical areas, and other relevant factors). EmblemHealth expressly reserves the right to revise these conclusions as clinical information changes and welcomes further relevant information. Each benefit program defines which services are covered. The conclusion that a particular service or supply is medically necessary does not constitute a representation or warranty that this service or supply is covered and/or paid for by EmblemHealth, as some programs exclude coverage for services or supplies that EmblemHealth considers medically necessary.

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Definitions

Gazyva (obinutuzumab): is a recombinant fully humanized monoclonal antibody (IgG1 subclass) that targets the CD20 antigen on the surface of pre B- and mature B-lymphocytes and after binding causes B-cell lysis. B- cell depletion in peripheral blood is not directly correlated with depletion in solid organs or in malignant deposits and has not been shown to directly correlate to clinical response. It has been shown that obinutuzumab activates polymorphoneclear neutrophils (PMNs), produces radical oxygen, and mediates phagocytosis by binding to CD16A and CD16B NA1 and NA2 isoforms.

Length of Authorization

Coverage will be provided for 6 months and may be renewed.

Dosing Limits [Medical Benefit]

Max Units (per dose and over time):

1. Chronic Lymphocytic Leukemia/Small Lymphocytic Lymphoma (CLL/SLL) Loading Dose:

• 10 billable units day 1, 90 billable units day 2, 100 billable units day 3, 200 billable units days 8 and 15 of Cycle 1 (21 days)

Maintenance Dose:

200 billable units every 21 days

2. All other indications

Loading dose:

100 billable units days 1, 8, 15 of cycle 1 (21 days)

Maintenance Dose:

• 100 billable units every 21 days for 8 cycles; then every 2 months for 2 years

Guideline

I. Initial Approval Criteria

<u>Gazyva</u> may be considered medically necessary if one of the below conditions are met **AND** use is consistent with the medical necessity criteria that follows:

1. Chronic Lymphocytic Leukemia (CLL)/Small Lymphocytic Lymphoma (SLL)

- A. Used as first-line therapy; AND
 - i. Used in combination with chlorambucil; OR
 - ii. Used in combination with acalabrutinib; OR
 - iii. Used in combination with venetoclax; OR
 - iv. Used as a single agent (excluding use in patients without del(17p)/TP53 mutation who are <65 years of age without significant comorbidities); **OR**
 - v. Used in combination with bendamustine for disease without del(17p)/TP53 mutation (excluding use in frail patients); **OR**
- B. Used as subsequent therapy; **AND**
 - i. Used for disease without del(17p)/TP53 mutation; AND
 - a. Used as a single agent (if not given as first-line therapy); AND
 - 1) Used for relapsed or refractory disease after prior BTK inhibitor (e.g., ibrutinib, calabrutinib, etc.)- and venetoclax-based regimens; **OR**
 - b. Used in combination with venetoclax (if previously used as first-line therapy); AND
 - 1) Used as retreatment for relapsed disease after a period of remission

2. Follicular Lymphoma

- A. Patient is > 18 years of age; AND
- B. Gazyva is used in **ONE** of the following situations (i, ii, or iii):
 - i. In combination with chemotherapy; **OR**Note: Examples include CHOP (cyclophosphamide, doxorubicin, vincristine, and prednisone), CVP (cyclophosphamide, vincristine, and prednisone), or bendamustine.
 - ii. For maintenance treatment following Gazyva in combination with chemotherapy; OR
 - iii. Patient experienced an adverse event or intolerance to a rituximab product

 Note: Examples of adverse events or intolerance includes paraneoplastic pemphigus, Stevens Johnson syndrome, lichenoid dermatitis, vesiculobullous dermatitis, toxic epidermal necrolysis. Examples of rituximab products include Rituxan and biosimilars.

Limitations/Exclusions

Gazyva is not considered medically necessary for when any of the following selection criteria is met:

- 1. The member has an active infection requiring systemic treatment.
- 2. Disease progression while taking Gazyva (obinutuzumab).
- 3. Dosing exceeds single dose limit of Gazyva (obinutuzumab) 1000 mg.
- 4. Indications not supported by CMS recognized compendia or acceptable peer reviewed literature may be deemed as not approvable and therefore not reimbursable.

II. Renewal Criteria

Authorizations for CLL/SLL may NOT be renewed

Authorizations for other indications may be renewed based on the following criteria:

- 1. Patient continues to meet criteria in INITIAL APPROVAL CRITERIA.
- 2. Disease response with treatment as defined by stabilization of disease or decrease in size of tumor or tumor spread; **AND**
- 3. Absence of unacceptable toxicity from the drug including: progressive multifocal leukoencephalopathy (PML), hepatitis B reactivation, severe neutropenia/febrile neutropenia, severe thrombocytopenia, severe infusion reactions, hypersensitivity reactions, tumor lysis syndrome, etc.; **AND**
- 4. Length of therapy does not exceed 2 years.

Dosage/Administration

Indication	Dose	
CLL/SLL	Combination therapy: 100 mg day 1, 900 mg day 2, then 1000 mg days 8 and 15 of Cycle 1 (loading do 1000 mg on Day 1 of Cycles 2-6 (28-day cycle) Monotherapy:	
	Cycle 1 (21-day cycle): 100 mg day 1, 900 mg day 2, then 1000 mg days 8 and 15	
	Cycles 2-8 (21-day cycle): 1000 mg on day 1 -OR-	
	Cycle 1 (21-day cycle): 100mg day 1, 900 mg day 2, 1000 mg day 3, 2000 mg days 8 and 15	
	Cycles 2-8 (21-day cycle): 2000 mg on day 1	
All other indications	1000 mg days 1, 8, & 15 of cycle 1 (loading doses); given in combination with chemotherapy Combination chemotherapy: 1000 mg day 1 of cycles 2-6 (28-day cycle) in combination with bendamustine 1000 mg day 1 of cycles 2-6 (21-day cycle) in combination with CHOP, followed by 2 additional 21-day cycles of Gazyva 1000 mg day 1 of cycles 2-8 (21-day cycle) with CVP 1000 mg every 2 months for two years as monotherapy	

Applicable Procedure Codes

Code	Description	
J9301	Injection, obinutuzumab, 10 mg, 1 billable unit = 10 mg	

Applicable NDCs

Code	Description
50242-0070-xx Gazyva 1000 mg/ 40 ml single dose vial	

ICD-10 Diagnoses

Code	Description
C82.00	Follicular lymphoma grade I unspecified site
C82.01	Follicular lymphoma grade I lymph nodes of head, face, and neck
C82.02	Follicular lymphoma grade I intrathoracic lymph nodes
C82.03	Follicular lymphoma grade I intra-abdominal lymph nodes
C82.04	Follicular lymphoma grade I lymph nodes of axilla and upper limb
C82.05	Follicular lymphoma grade I lymph nodes of inguinal region and lower limb
C82.06	Follicular lymphoma grade I intrapelvic lymph nodes
C82.07	Follicular lymphoma grade I spleen
C82.08	Follicular lymphoma grade I lymph nodes of multiple sites
C82.09	Follicular lymphoma grade I extranodal and solid organ sites
C82.10	Follicular lymphoma grade II unspecified site
C82.11	Follicular lymphoma grade II lymph nodes of head, face, and neck
C82.12	Follicular lymphoma grade II intrathoracic lymph nodes
C82.13	Follicular lymphoma grade II intra-abdominal lymph nodes
C82.14	Follicular lymphoma grade II lymph nodes of axilla and upper limb
C82.15	Follicular lymphoma grade II lymph nodes of inguinal region and lower limb
C82.16	Follicular lymphoma grade II intrapelvic lymph nodes
C82.17	Follicular lymphoma grade II spleen
C82.18	Follicular lymphoma grade II lymph nodes of multiple sites
C82.19	Follicular lymphoma grade II extranodal and solid organ sites
C82.20	Follicular lymphoma grade III unspecified site
C82.21	Follicular lymphoma grade III lymph nodes of head, face, and neck
C82.22	Follicular lymphoma grade III intrathoracic lymph nodes
C82.23	Follicular lymphoma grade III intra-abdominal lymph nodes
C82.24	Follicular lymphoma grade III lymph nodes of axilla and upper limb
C82.25	Follicular lymphoma grade III lymph nodes of inguinal region and lower limb
C82.26	Follicular lymphoma grade III intrapelvic lymph nodes
C82.27	Follicular lymphoma grade III spleen
C82.28	Follicular lymphoma grade III lymph nodes of multiple sites
C82.29	Follicular lymphoma grade III extranodal and solid organ sites
C82.30	Follicular lymphoma grade IIIa unspecified site
C82.31	Follicular lymphoma grade IIIa lymph nodes of head, face, and neck
C82.32	Follicular lymphoma grade IIIa intrathoracic lymph nodes
C82.33	Follicular lymphoma grade IIIa intra-abdominal lymph nodes
C82.34	Follicular lymphoma grade IIIa lymph nodes of axilla and upper limb
C82.35	Follicular lymphoma grade IIIa lymph nodes of inguinal region and lower limb
C82.36	Follicular lymphoma grade IIIa intrapelvic lymph nodes
C82.37	Follicular lymphoma grade IIIa spleen
C82.38	Follicular lymphoma grade IIIa lymph nodes of multiple sites
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C82.87 Other types of follicular lymphoma spleen lymph nodes of multiple sites C82.88 Other types of follicular lymphoma lymph nodes of multiple sites C82.89 Other types of follicular lymphoma extranodal and solid organ sites C82.90 Follicular lymphoma, unspecified site C82.91 Follicular lymphoma, unspecified lymph nodes of head, face, and neck C82.92 Follicular lymphoma, unspecified intrathoracic lymph nodes C82.93 Follicular lymphoma, unspecified intra-abdominal lymph nodes C82.94 Follicular lymphoma, unspecified lymph nodes of axilla and upper limb	C82.85	Other types of follicular lymphoma lymph nodes of inguinal region and lower limb
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C82.93 Follicular lymphoma, unspecified intra-abdominal lymph nodes C82.94 Follicular lymphoma, unspecified lymph nodes of axilla and upper limb	C82.91	Follicular lymphoma, unspecified lymph nodes of head, face, and neck
C82.94 Follicular lymphoma, unspecified lymph nodes of axilla and upper limb	C82.92	
	C82.93	Follicular lymphoma, unspecified intra-abdominal lymph nodes
C82.95 Follicular lymphoma, unspecified lymph nodes of inguinal region and lower limb	C82.94	
	C82.95	Follicular lymphoma, unspecified lymph nodes of inguinal region and lower limb

C82.97 Foll C82.98 Foll C82.99 Foll C83.00 Sma C83.01 Sma C83.02 Sma C83.03 Sma C83.04 Sma C83.05 Sma C83.06 Sma C83.07 Sma	licular lymphoma, unspecified spleen licular lymphoma, unspecified spleen licular lymphoma, unspecified lymph nodes of multiple sites licular lymphoma, unspecified extranodal and solid organ sites all cell B-cell lymphoma unspecified site all cell B-cell lymphoma lymph nodes of head, face, and neck all cell B-cell lymphoma intrathoracic lymph nodes all cell B-cell lymphoma intra-abdominal lymph nodes all cell B-cell lymphoma lymph nodes of axilla and upper limb all cell B-cell lymphoma lymph nodes of inguinal region and lower limb all cell B-cell lymphoma intrapelvic lymph nodes all cell B-cell lymphoma spleen
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C83.06 Sma C83.07 Sma	all cell B-cell lymphoma intrapelvic lymph nodes
C83.07 Sma	
	all cell B-cell lymphoma spleen
C83.08 Sma	all cell B-cell lymphoma lymph nodes of multiple sites
C83.09 Sma	all cell B-cell lymphoma extranodal and solid organ sites
C83.80 Oth	ner non-follicular lymphoma unspecified site
C83.81 Oth	ner non-follicular lymphoma lymph nodes of head, face, and neck
C83.82 Oth	ner non-follicular lymphoma intrathoracic lymph nodes
C83.83 Oth	ner non-follicular lymphoma intra-abdominal lymph nodes
	ner non-follicular lymphoma lymph nodes of axilla and upper limb
C83.85 Oth	ner non-follicular lymphoma lymph nodes of inguinal region and lower limb
C83.86 Oth	ner non-follicular lymphoma intrapelvic lymph nodes
	ner non-follicular lymphoma spleen
C83.88 Oth	ner non-follicular lymphoma lymph nodes of multiple sites
	ner non-follicular lymphoma extranodal and solid organ sites
C85.80 Oth	ner specified types of non-Hodgkin lymphoma, unspecified site
C85.81 Oth	ner specified types of non-Hodgkin lymphoma, lymph nodes of head, face, and neck
	ner specified types of non-Hodgkin lymphoma, intrathoracic lymph nodes
	ner specified types of non-Hodgkin lymphoma, intra-abdominal lymph nodes
C85.84 Oth	ner specified types of non-Hodgkin lymphoma, lymph nodes of axilla and upper limb
C85.85 Oth	ner specified types of non-Hodgkin lymphoma, lymph nodes of inguinal region and lower limb
C85.86 Oth	ner specified types of non-Hodgkin lymphoma, intrapelvic lymph nodes
C85.87 Oth	ner specified types of non-Hodgkin lymphoma, spleen
C85.88 Oth	ner specified types of non-Hodgkin lymphoma, lymph nodes of multiple sites
C85.89 Oth	ner specified types of non-Hodgkin lymphoma, extranodal and solid organ sites
C88.4 Extr	ranodal marginal zone B-cell lymphoma of mucosa-associated lymphoid tissue [MALT-lymphoma]
	ronic lymphocytic leukemia of B-cell type not having achieved remission
C91.12 Chr	ronic lymphocytic leukemia of B-cell type in relapse

Revision History

Company(ies)	DATE	REVISION
EmblemHealth & ConnectiCare	3/1/2024	Annual Review: Updated dosing limits for other indications, updated dosing chart. No criteria changes/
EmblemHealth & ConnectiCare	6/28/2023	Annual Review: <u>Dosing Limits:</u> Chronic Lymphocytic Leukemia/Small Lymphocytic Lymphoma (CLL/SLL) Loading Dose: Removed

		"10 billable units day 1, 90 billable units day 2, 100 billable units days 8, 15 of cycle 1 (28 days)" Added "10 billable units day 1, 90 billable units day 2, 100 billable units day 3, 200 billable units days 8 and 15 of Cycle 1 (21 days)" Maintenance Dose: Removed "100 billable units every 28 days" Added "200 billable units every 21 days " Chronic Lymphocytic Leukemia (CLL)/Small Lymphocytic Lymphoma (SLL): Initial Criteria: Removed "a. The member has stage II-IV CD20 positive CLL/SLL, or if Stage 0-I disease, member must have bulky adenopathy, splenomegaly, OR systemic symptoms; AND b. The member had no prior CLL therapy; AND c. In combination with chlorambucil for disease without del(17p)/TP53 mutation in members age ≥65 years, in younger members with significant comorbidities, or in frail patients unable to tolerate purine analogs." Added "a. Used as first-line therapy; AND i. Used in combination with chlorambucil †; OR ii. Used in combination with chlorambucil †; OR iii. Used in combination with venetoclax; OR iv. Used as a single agent (excluding use in patients without del(17p)/TP53 mutation who are <65 years of age without significant comorbidities); OR v. Used in combination with bendamustine for disease without del(17p)/TP53 mutation who are <65 years of age without significant comorbidities); OR v. Used as subsequent therapy; AND i. Used as subsequent therapy; AND i. Used for disease without del(17p)/TP53 mutation; OR b. Used as a single agent (if not given as first-line therapy); AND 1) Used for relapsed or refractory disease after prior BTK inhibitor (e.g., ibrutinib, calabrutinib, etc.)- and venetoclax-based regimens; OR b) Used in combination with venetoclax (if previously used as first-line therapy); AND 1) Used as retreatment for relapsed disease after a period of remission" Removed Non-Hodgkin Lymphoma Criteria Limitations and Exclusions: removed: "Treatment with Gazyva (obinutuzumab) exceeds the total duration limit of 6 cycles."
EmblemHealth & ConnectiCare	6/17/20022	Transferred policy to new template
EmblemHealth & ConnectiCare	7/15/2019	Annual review

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- 5. AHFS Drug Information. American Society of Health-Systems Pharmacist or Wolters Kluwer Lexi-Drugs. Bethesda, MD. 2017.