

CAMPATH- alemtuzumab injection

Genzyme Corporation

HIGHLIGHTS OF PRESCRIBING INFORMATION

These highlights do not include all the information needed to use CAMPATH safely and effectively. See full prescribing information for CAMPATH.

CAMPATH® (alemtuzumab) injection, for intravenous use
Initial U.S. Approval: 2001

WARNING: CYTOPENIAS, INFUSION-RELATED REACTIONS, AND INFECTIONS

See full prescribing information for complete boxed warning.

Serious, including fatal, cytopenias, infusion-related reactions, and infections can occur (5.1-5.3).

- Limit doses to 30 mg (single) and 90 mg (cumulative weekly); higher doses increase risk of pancytopenia. (2.1)
- Escalate dose gradually and monitor patients during infusion. Withhold therapy for Grade 3 or 4 infusion-related reactions. (5.2)
- Administer prophylaxis against *Pneumocystis jirovecii* pneumonia (PCP) and herpes virus infections. (2.2, 5.3)

----- INDICATIONS AND USAGE -----

CAMPATH is a CD52-directed cytolytic antibody indicated as a single agent for the treatment of B-cell chronic lymphocytic leukemia (B-CLL). (1)

----- DOSAGE AND ADMINISTRATION -----

- Administer as an intravenous infusion over 2 hours. (2.1)
- Escalate to recommended dose of 30 mg/day three times per week for 12 weeks. (2.1)
- Premedicate with oral antihistamine and acetaminophen. (2.2)

----- DOSAGE FORMS AND STRENGTHS -----

Injection: 30 mg/1 mL single-dose vial (3)

----- CONTRAINDICATIONS -----

None (4)

----- WARNINGS AND PRECAUTIONS -----

- Cytopenias: Obtain complete blood counts (CBC) and platelet counts at weekly intervals during therapy and CD4 counts after therapy until recovery to ≥ 200 cells/ μ L. Withhold for severe cytopenias. Discontinue for autoimmune or severe hematologic adverse reactions. (2.2, 5.1)
- Immunosuppression/Infections: CAMPATH induces severe and prolonged lymphopenia and increases risk of infection. If a serious infection occurs, withhold treatment until infection resolves. (5.3)
- Immunization: Do not administer live viral vaccines to patients who have recently received CAMPATH. (5.4)

----- ADVERSE REACTIONS -----

Most common adverse reactions ($\geq 10\%$): cytopenias, infusion-related reactions, cytomegalovirus (CMV) and other infections, nausea, emesis, diarrhea, and insomnia. (6)

To report SUSPECTED ADVERSE REACTIONS, contact Genzyme Corporation at 1-877-4-CAMPATH (1-877-422-6728) or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch.

----- USE IN SPECIFIC POPULATIONS -----

- Pregnancy: May cause fetal harm. (8.1)
- Lactation: Advise not to breastfeed. (8.2)

See 17 for PATIENT COUNSELING INFORMATION.

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WARNING: CYTOPENIAS, INFUSION-RELATED REACTIONS, AND INFECTIONS

Cytopenias: Serious, including fatal, pancytopenia/marrow hypoplasia, autoimmune idiopathic thrombocytopenia, and autoimmune hemolytic anemia can occur in patients receiving CAMPATH. Single doses of CAMPATH greater than 30 mg or cumulative doses greater than 90 mg per week increase the incidence of pancytopenia [see *Warnings and Precautions (5.1)*].

Infusion-Related Reactions: CAMPATH administration can result in serious, including fatal, infusion-related reactions. Carefully monitor patients during infusions and withhold CAMPATH for Grade 3 or 4 infusion-related reactions. Gradually escalate CAMPATH to the recommended dose at the initiation of therapy and after interruption of therapy for 7 or more days [see *Dosage and Administration (2.1)* and *Warnings and Precautions (5.2)*].

Immunosuppression/Infections: Serious, including fatal, bacterial, viral, fungal, and protozoan infections can occur in patients receiving CAMPATH. Administer prophylaxis against *Pneumocystis jirovecii* pneumonia (PCP) and herpes virus infections [see *Dosage and Administration (2.2)* and *Warnings and Precautions (5.3)*].

1 INDICATIONS AND USAGE

CAMPATH is indicated as a single agent for the treatment of B-cell chronic lymphocytic leukemia (B-CLL).

2 DOSAGE AND ADMINISTRATION

2.1 Dosing Schedule and Administration

- Administer as an intravenous infusion over 2 hours. **Do not administer as intravenous push or bolus.**
- Recommended Dosing Regimen
 - Gradually escalate to the maximum recommended single dose of 30 mg. Escalation is required at initiation of dosing or if dosing is held ≥ 7 days during treatment. Escalation to 30 mg ordinarily can be accomplished in 3 to 7 days.
 - Escalation Strategy:
 - Administer 3 mg daily until infusion-related reactions are \leq Grade 2 [see *Adverse Reactions (6.1)*].
 - Then administer 10 mg daily until infusion-related reactions are \leq Grade 2.
 - Then administer 30 mg/day three times per week on alternate days (e.g., Mon-Wed-Fri). The total duration of therapy, including dose escalation, is 12 weeks.
- **Single doses of greater than 30 mg or cumulative doses greater than 90 mg per week increase the incidence of pancytopenia.**

2.2 Recommended Concomitant Medications

- Premedicate with diphenhydramine (50 mg) and acetaminophen (500–1000 mg) 30 minutes prior to first infusion and each dose escalation. Institute appropriate medical management (e.g., glucocorticoids, epinephrine, meperidine) for infusion-related reactions as needed [see *Warnings and Precautions (5.2)* and *Adverse Reactions (6.1)*].
- Administer trimethoprim/sulfamethoxazole double strength (DS) twice daily 3 times per week (or equivalent) as *Pneumocystis jirovecii* pneumonia (PCP) prophylaxis.
- Administer famciclovir 250 mg BID or equivalent as herpetic prophylaxis.

Continue PCP and herpes viral prophylaxis for a minimum of 2 months after completion of CAMPATH or until the CD4+ count is ≥ 200 cells/ μL , whichever occurs later [see *Warnings and Precautions (5.3)*].

2.3 Dosage Modification

- Withhold CAMPATH during serious infection or other serious adverse reactions until resolution.
- Discontinue CAMPATH for autoimmune anemia or autoimmune thrombocytopenia.
- There are no dose modifications recommended for lymphopenia.

Dose Modification for Neutropenia or Thrombocytopenia [see *Warnings and Precautions (5.1)*]

Hematologic Values	Dosage Modification*
ANC $< 250/\mu\text{L}$ and/or platelet count $\leq 25,000/\mu\text{L}$	
For first occurrence:	Withhold CAMPATH therapy. Resume CAMPATH at 30 mg when ANC $\geq 500/\mu\text{L}$ and platelet count $\geq 50,000/\mu\text{L}$.
For second occurrence:	Withhold CAMPATH therapy. Resume CAMPATH at 10 mg when ANC $\geq 500/\mu\text{L}$ and platelet count $\geq 50,000/\mu\text{L}$.
For third occurrence:	Discontinue CAMPATH therapy.
$\geq 50\%$ decrease from baseline in patients initiating therapy with a baseline ANC $\leq 250/\mu\text{L}$ and/or a baseline platelet count $\leq 25,000/\mu\text{L}$	
For first occurrence:	Withhold CAMPATH therapy. Resume CAMPATH at 30 mg upon return to baseline value(s).
For second occurrence:	Withhold CAMPATH therapy. Resume CAMPATH at 10 mg upon return to baseline value(s).
For third occurrence:	Discontinue CAMPATH therapy.

* If the delay between dosing is ≥ 7 days, initiate therapy at CAMPATH 3 mg and escalate to 10 mg and then to 30 mg as tolerated [see *Dosage and Administration (2.1)*].

2.4 Preparation and Administration

Parenteral drug products should be inspected visually for particulate matter and discoloration prior to administration. If particulate matter is present or the solution is discolored, discard the vial. **DO NOT SHAKE VIAL.**

Use aseptic technique during the preparation and administration of CAMPATH. Withdraw the necessary amount of CAMPATH from the vial into a syringe.

- To prepare the 3 mg dose, withdraw 0.1 mL into a 1 mL syringe calibrated in increments of 0.01 mL.
- To prepare the 10 mg dose, withdraw 0.33 mL into a 1 mL syringe calibrated in increments of 0.01 mL.
- To prepare the 30 mg dose, withdraw 1 mL in either a 1 mL or 3 mL syringe calibrated in 0.1 mL increments.

Inject syringe contents into 100 mL sterile 0.9% Sodium Chloride USP or 5% Dextrose in Water USP. **Gently invert the bag to mix the solution.** Discard syringe.

The vial contains no preservatives and is intended for single use only. DISCARD VIAL including any unused portion after withdrawal of dose.

Use within 8 hours after dilution. Store diluted CAMPATH at room temperature between 15°C to 30°C (59°F to 86°F) or refrigerated at 2°C to 8°C (36°F to 46°F). Protect from light.

2.5 Incompatibilities

CAMPATH is compatible with polyvinylchloride (PVC) bags and PVC or polyethylene-lined PVC administration sets. Do not add or simultaneously infuse other drug substances through the same intravenous line.

3 DOSAGE FORMS AND STRENGTHS

Injection: 30 mg/1 mL as a clear, colorless solution in single-dose vial

4 CONTRAINDICATIONS

None.

5 WARNINGS AND PRECAUTIONS

5.1 Cytopenias

Severe, including fatal, autoimmune anemia and thrombocytopenia, and prolonged myelosuppression have been reported in patients receiving CAMPATH.

In addition, hemolytic anemia, pure red cell aplasia, bone marrow aplasia, and hypoplasia have been reported after treatment with CAMPATH at the recommended dose. Single doses of CAMPATH greater than 30 mg or cumulative doses greater than 90 mg per week increase the incidence of pancytopenia.

Withhold CAMPATH for severe cytopenias (except lymphopenia). Discontinue for

autoimmune cytopenias or recurrent/persistent severe cytopenias (except lymphopenia) [see *Dosage and Administration (2.3)*]. No data exist on the safety of CAMPATH resumption in patients with autoimmune cytopenias or marrow aplasia [see *Adverse Reactions (6.1)*].

Obtain complete blood counts (CBC) at weekly intervals during CAMPATH therapy and more frequently if worsening anemia, neutropenia, or thrombocytopenia occurs. Assess CD4+ counts after treatment until recovery to ≥ 200 cells/ μL [see *Dosage and Administration (2.3)* and *Adverse Reactions (6)*].

5.2 Infusion-Related Reactions

Adverse reactions occurring during or shortly after CAMPATH infusion include pyrexia, chills/rigors, nausea, hypotension, urticaria, dyspnea, rash, emesis, and bronchospasm [see *Adverse Reactions (6.1)*]. In clinical trials, the frequency of infusion-related reactions was highest in the first week of treatment. Monitor for the signs and symptoms listed above and withhold infusion for Grade 3 or 4 infusion-related reactions.

The following serious, including fatal, infusion-related reactions have been identified in postmarketing reports: syncope, pulmonary infiltrates, acute respiratory distress syndrome (ARDS), respiratory arrest, cardiac arrhythmias, myocardial infarction, acute cardiac insufficiency, cardiac arrest, angioedema, and anaphylactoid shock.

Initiate CAMPATH according to the recommended dose-escalation scheme [see *Dosage and Administration (2.1)*]. Premedicate patients with an antihistamine and acetaminophen prior to each dose. Institute appropriate medical management (e.g., glucocorticoids, epinephrine, meperidine) for infusion-related reactions as needed [see *Dosage and Administration (2.2)*]. If therapy is interrupted for 7 or more days, reinstitute CAMPATH with gradual dose escalation [see *Dosage and Administration (2.1)*].

5.3 Immunosuppression/Infections

CAMPATH treatment results in severe and prolonged lymphopenia with a concomitant increased incidence of opportunistic infections [see *Adverse Reactions (6.1)*]. Administer PCP and herpes viral prophylaxis during treatment with CAMPATH and for a minimum of 2 months after completion of CAMPATH or until the CD4+ count is ≥ 200 cells/ μL , whichever occurs later [see *Dosage and Administration (2.2)*]. Prophylaxis does not eliminate these infections.

Routinely monitor patients for CMV infection during treatment with CAMPATH and for at least 2 months following completion of CAMPATH. Withhold CAMPATH for serious infections and during antiviral treatment for CMV infection or confirmed CMV viremia (defined as polymerase chain reaction [PCR] positive CMV in ≥ 2 consecutive samples obtained 1 week apart). Initiate therapeutic ganciclovir (or equivalent) for CMV infection or confirmed CMV viremia.

Epstein-Barr virus (EBV) infection, including severe and fatal EBV-associated hepatitis, has been reported in patients who received CAMPATH.

Monitor for sign and symptoms of EBV infections. Withhold CAMPATH for EBV reactivation or severe infection.

Administer only irradiated blood products to avoid transfusion associated Graft versus Host Disease (TAGVHD), unless emergent circumstances dictate immediate transfusion.

In patients who received CAMPATH as initial therapy, recovery of CD4+ counts to ≥ 200 cells/ μL occurred by 6 months following completion of CAMPATH; however, at 2 months post treatment, the median was 183 cells/ μL . In previously treated patients who received CAMPATH, the median time to recovery of CD4+ counts to ≥ 200 cells/ μL was 2 months; however, full recovery (to baseline) of CD4+ and CD8+ counts may take more than 12 months [see *Adverse Reactions (6)*].

5.4 Immunization

The safety of immunization with live viral vaccines following CAMPATH therapy has not been studied. Do not administer live viral vaccines to patients or infants born to patients receiving CAMPATH. The ability to generate an immune response to any vaccine following CAMPATH therapy has not been studied.

6 ADVERSE REACTIONS

The following clinically significant adverse reactions are discussed in greater detail in other sections of the label:

- Cytopenias [see *Warnings and Precautions (5.1)*]
- Infusion-Related Reactions [see *Warnings and Precautions (5.2)*]
- Immunosuppression/Infections [see *Warnings and Precautions (5.3)*]

6.1 Clinical Trials Experience

Because clinical trials are conducted under widely varying conditions, adverse reaction rates observed in the clinical trials of a drug cannot be directly compared to rates in the clinical trials of another drug and may not reflect the rates observed in practice.

The data below reflect exposure to CAMPATH in 296 patients with CLL of whom 147 were previously untreated and 149 received at least 2 prior chemotherapy regimens. The median duration of exposure was 11.7 weeks for previously untreated patients and 8 weeks for previously treated patients.

The most common adverse reactions with CAMPATH are: infusion-related reactions (pyrexia, chills, hypotension, urticaria, nausea, rash, tachycardia, dyspnea), cytopenias (neutropenia, lymphopenia, thrombocytopenia, anemia), infections (CMV viremia, CMV infection, other infections), gastrointestinal symptoms (nausea, emesis, abdominal pain), and neurological symptoms (insomnia, anxiety). The most common serious adverse reactions are cytopenias, infusion-related reactions, and immunosuppression/infections.

Lymphopenia

Severe lymphopenia and a rapid and sustained decrease in lymphocyte subsets occurred in previously untreated and previously treated patients following administration of CAMPATH. In previously untreated patients, the median CD4+ was 0 cells/ μL at one month after treatment and 238 cells/ μL [25%–75% interquartile range 115 to 418 cells/ μL at 6 months post treatment [see *Warnings and Precautions (5.3)*].

Neutropenia

In previously untreated patients, the incidence of Grade 3 or 4 neutropenia was 42% with a median time to onset of 31 days and a median duration of 37 days. In previously treated patients, the incidence of Grade 3 or 4 neutropenia was 64% with a median

duration of 28 days. Ten percent of previously untreated patients and 17% of previously treated patients received granulocyte colony stimulating factors.

Anemia

In previously untreated patients, the incidence of Grade 3 or 4 anemia was 12% with a median time to onset of 31 days and a median duration of 8 days. In previously treated patients, the incidence of Grade 3 or 4 anemia was 38%. Seventeen percent of previously untreated patients and 66% of previously treated patients received either erythropoiesis stimulating agents, transfusions or both.

Thrombocytopenia

In previously untreated patients, the incidence of Grade 3 or 4 thrombocytopenia was 14% with a median time to onset of 9 days and a median duration of 14 days. In previously treated patients, the incidence of Grade 3 or 4 thrombocytopenia was 52% with a median duration of 21 days. Autoimmune thrombocytopenia was reported in 2% of previously treated patients with one fatality.

Infusion-Related Reactions

Infusion-related reactions, which included pyrexia, chills, hypotension, urticaria, and dyspnea, were common. Grade 3 and 4 pyrexia and/or chills occurred in approximately 10% of previously untreated patients and in approximately 35% of previously treated patients. The occurrence of infusion-related reactions was greatest during the initial week of treatment and decreased with subsequent doses of CAMPATH. All patients were pretreated with antipyretics and antihistamines; additionally, 43% of previously untreated patients received glucocorticoid pre-treatment.

Infections

In the study of previously untreated patients, patients were tested weekly for CMV using a PCR assay from initiation through completion of therapy, and every 2 weeks for the first 2 months following therapy. CMV infection occurred in 16% (23/147) of previously untreated patients; approximately one-third of these infections were serious or life threatening. In studies of previously treated patients in which routine CMV surveillance was not required, CMV infection was documented in 6% (9/149) of patients; nearly all of these infections were serious or life threatening.

Other infections were reported in approximately 50% of patients across all studies. Grade 3 to 5 sepsis ranged from 3% to 10% across studies and was higher in previously treated patients. Grade 3 to 4 febrile neutropenia ranged from 5% to 10% across studies and was higher in previously treated patients. Infection-related fatalities occurred in 2% of previously untreated patients and 16% of previously treated patients. There were 198 episodes of other infection in 109 previously untreated patients; 16% were bacterial, 7% were fungal, 4% were other viral, and in 73% the organism was not identified.

Cardiac

Cardiac dysrhythmias occurred in approximately 14% of previously untreated patients. The majority were tachycardias and were temporally associated with infusion; dysrhythmias were Grade 3 or 4 in 1% of patients.

Previously Untreated Patients

Table 1 contains selected adverse reactions observed in 294 patients randomized (1:1) to receive CAMPATH or chlorambucil as first line therapy for B-CLL. CAMPATH was administered at a dose of 30 mg intravenously three times weekly for up to 12 weeks. The median duration of therapy was 11.7 weeks with a median weekly dose of 82 mg (25%-75% interquartile range: 69-90 mg).

Table 1: Per Patient Incidence of Selected* Adverse Reactions in Treatment Naive B-CLL Patients

		CAMPATH (n=147)		Chlorambucil (n=147)	
		All Grades† %	Grades 3-4 %	All Grades %	Grades 3-4 %
Blood and Lymphatic System Disorders	Lymphopenia	97	97	9	1
	Neutropenia	77	42	51	26
	Anemia	76	13	54	18
	Thrombocytopenia	71	13	70	14
General Disorders and Administration Site Conditions	Pyrexia	69	10	11	1
	Chills	53	3	1	0
Infections and Infestations	CMV viremia‡	55	4	8	0
	CMV infection	16	5	0	0
	Other infections	74	21	65	10
Skin and Subcutaneous Tissue Disorders	Urticaria	16	2	1	0
	Rash	13	1	4	0
	Erythema	4	0	1	0
Vascular Disorders	Hypotension	16	1	0	0
	Hypertension	14	5	2	1
Nervous System Disorders	Headache	14	1	8	0
	Tremor	3	0	1	0
Respiratory, Thoracic and Mediastinal Disorders	Dyspnea	14	4	7	3
Gastrointestinal Disorders	Diarrhea	10	1	4	0
Psychiatric Disorders	Insomnia	10	0	3	0
	Anxiety	8	0	1	0
Cardiac Disorders	Tachycardia	10	0	1	0

* Adverse reactions occurring at a higher relative frequency in the CAMPATH arm

† NCI CTC version 2.0 for adverse reactions; NCI CTCAE version 3.0 for laboratory values

‡ CMV viremia (without evidence of symptoms) includes both cases of single

PCR positive test results and of confirmed CMV viremia (≥ 2 occasions in consecutive samples 1 week apart). For the latter, ganciclovir (or equivalent) was initiated per protocol.

Previously Treated Patients

Additional safety information was obtained from 3 single arm studies of 149 previously treated patients with CLL administered 30 mg CAMPATH intravenously three times weekly for 4 to 12 weeks (median cumulative dose 673 mg [range 2–1106 mg]; median duration of therapy 8.0 weeks). Adverse reactions in these studies not listed in Table 1 that occurred at an incidence rate of $>5\%$ were fatigue, nausea, emesis, musculoskeletal pain, anorexia, dysesthesia, mucositis, and bronchospasm.

6.2 Immunogenicity

As with all therapeutic proteins, there is potential for immunogenicity. The incidence of antibody formation is highly dependent on the sensitivity and specificity of the assay. Additionally, the observed incidence of antibody (including neutralizing antibody) positivity in an assay may be influenced by several factors including assay methodology, sample handling, timing of sample collection, concomitant medications, and underlying disease. For these reasons, comparison of the incidence of antibodies with the incidence of antibodies to other alemtuzumab products may be misleading.

Using an ELISA assay, anti-human antibodies (HAHA) were detected in 11 of 133 (8.3%) previously untreated patients. In addition, two patients were weakly positive for neutralizing activity. Limited data suggest that the anti-CAMPATH antibodies did not adversely affect tumor response. Four of 211 (1.9%) previously treated patients were found to have antibodies to CAMPATH following treatment.

6.3 Postmarketing Experience

CAMPATH

The following adverse reactions have been identified during postapproval use of CAMPATH. Because these reactions are reported voluntarily from a population of uncertain size, it is not always possible to reliably estimate their frequency or establish a causal relationship to drug exposure.

General Disorders and Administration Site Conditions: Fatal infusion-related reactions.

Cardiovascular Disorders: Congestive heart failure, cardiomyopathy, decreased ejection fraction (some patients had been previously treated with cardiotoxic agents).

Cerebrovascular Disorders: Cervicocephalic arterial dissection, stroke, including hemorrhagic and ischemic stroke.

Gastrointestinal Disorders: Acute acalculous cholecystitis.

Immune System Disorders: Goodpasture's syndrome, Graves' disease, aplastic anemia, Guillain-Barré syndrome, chronic inflammatory demyelinating polyradiculoneuropathy, serum sickness, fatal transfusion associated graft versus host disease, hemophagocytic lymphohistiocytosis (HLH).

Infections: Epstein-Barr virus (EBV) infection, progressive multifocal leukoencephalopathy (PML), reactivation of latent viruses.

Metabolism and Nutrition Disorders: Tumor lysis syndrome.

Neoplasms: EBV-associated lymphoproliferative disorder.

Nervous System Disorders: Optic neuropathy.

Renal and Urinary Disorders: Glomerular nephropathies.

Other Alemtuzumab Products

The following adverse reactions have been identified during postapproval use of another alemtuzumab product. Because these reactions are reported voluntarily from a population of uncertain size, it is not always possible to reliably estimate their frequency or establish a causal relationship to drug exposure.

Endocrine Disorders: Hypothyroidism, hyperthyroidism, and thyroiditis.

Nervous System Disorders: Autoimmune encephalitis.

7 DRUG INTERACTIONS

No formal drug interaction studies have been performed with CAMPATH.

8 USE IN SPECIFIC POPULATIONS

8.1 Pregnancy

Risk Summary

Based on findings from animal studies, CAMPATH may cause fetal harm when administered to a pregnant woman. Available data from published cohort studies in pregnant women are insufficient to establish a CAMPATH-associated risk of major birth defects, miscarriage or adverse maternal or fetal outcomes. Alemtuzumab was embryolethal in pregnant huCD52 transgenic mice when administered during organogenesis (*see Data*). Human IgG antibodies are known to cross the placental barrier; therefore, CAMPATH may be transmitted from the mother to the developing fetus. Advise women of the potential risk to the fetus. Infants born to pregnant women treated with CAMPATH may be at increased risk of infection (*see Clinical Considerations*).

The background risk of major birth defects and miscarriage for the indicated population is unknown. All pregnancies have a background risk of birth defect, loss, or other adverse outcomes. In the U.S. general population, the estimated background risk of major birth defects and miscarriage in clinically recognized pregnancies is 2% to 4% and 15% to 20%, respectively.

Clinical Considerations

Fetal/Neonatal adverse reactions

Monoclonal antibodies are transported across the placenta as pregnancy progresses, with the largest amount transferred during the third trimester. Consider the risks and benefits of administering live or live-attenuated vaccines to infants exposed to CAMPATH *in utero* [*see Warnings and Precautions (5.3, 5.4)*].

Data

Animal data

When alemtuzumab was administered to pregnant huCD52 transgenic mice during organogenesis (gestation days [GD] 6–10 or GD 11–15) at intravenous doses of 3 or 10 mg/kg, no teratogenic effects were observed. However, there was an increase in embryoletality (increased postimplantation loss and the number of dams with all fetuses dead or resorbed) in pregnant animals dosed during GD 11–15. In a separate study in pregnant huCD52 transgenic mice, administration of alemtuzumab during organogenesis (GD 6–10 or GD 11–15) at intravenous doses of 3 or 10 mg/kg, decreases in B-lymphocyte and T-lymphocyte populations were observed in the offspring at both doses tested.

In pregnant huCD52 transgenic mice administered alemtuzumab at intravenous doses of 3 or 10 mg/kg/day throughout gestation and lactation, there was an increase in pup deaths during the lactation period at 10 mg/kg. Decreases in T-lymphocyte and B-lymphocyte populations and in antibody response were observed in offspring at both doses tested.

8.2 Lactation

Risk Summary

There are no data on the presence of alemtuzumab in human milk, effects on milk production, or the breastfed child. The effects of local gastrointestinal exposure and limited systemic exposure in the breastfed infant to alemtuzumab are unknown. Alemtuzumab was detected in the milk of lactating huCD52 transgenic mice administered alemtuzumab (*see Data*). Maternal IgG is known to be present in human milk and when a drug is present in animal milk, it is likely that the drug will be present in human milk.

Because of the potential for serious adverse reactions from CAMPATH in a breastfed child, including reduced lymphocyte counts, advise lactating women not to breastfeed during treatment with CAMPATH and for at least 3 months following the last dose.

Data

Alemtuzumab was detected in the milk of lactating huCD52 transgenic mice following intravenous administration of alemtuzumab at a dose of 10 mg/kg on postpartum days 8–12. Serum levels of alemtuzumab were similar in lactating mice and offspring on postpartum day 13 and were associated with evidence of pharmacological activity (decrease in lymphocyte counts) in the offspring.

8.3 Females and Males of Reproductive Potential

CAMPATH may cause embryo-fetal harm when administered to pregnant women [*see Use in Specific Populations (8.1)*].

Pregnancy Testing

Pregnancy testing is recommended for females of reproductive potential prior to initiating CAMPATH therapy.

Contraception

Females

Advise female patients of reproductive potential to use effective contraception during

treatment with CAMPATH and for at least 3 months after the last dose.

Infertility

Based on findings from animal studies, alemtuzumab may impair fertility in females and males of reproductive potential [see *Nonclinical Toxicology (13.1)*]. The reversibility of the effect on fertility is unknown.

8.4 Pediatric Use

Safety and effectiveness of CAMPATH have not been established in pediatric patients.

8.5 Geriatric Use

Of 147 previously untreated B-CLL patients treated with CAMPATH, 35% were \geq age 65 and 4% were \geq age 75. Of 149 previously treated patients with B-CLL, 44% were \geq 65 years of age and 10% were \geq 75 years of age. Clinical studies of CAMPATH did not include sufficient number of subjects age 65 and over to determine whether they respond differently than younger subjects. Other reported clinical experience has not identified differences in responses between the elderly and younger patients.

10 OVERDOSAGE

Across all clinical experience, the reported maximum single dose received was 90 mg. Bone marrow aplasia, infections, or severe infusion-related reactions occurred in patients who received a dose higher than recommended.

One patient who received an 80 mg dose intravenously experienced acute bronchospasm, cough, and dyspnea, followed by anuria and death. Another patient received two 90 mg doses intravenously one day apart during the second week of treatment and experienced a rapid onset of bone marrow aplasia.

There is no known specific antidote for CAMPATH overdose. Discontinue CAMPATH and provide supportive therapy.

11 DESCRIPTION

Alemtuzumab, a CD52-directed cytolytic antibody, is a recombinant DNA-derived humanized monoclonal antibody (CAMPATH-1H). CAMPATH-1H is an IgG1 kappa antibody with human variable framework and constant regions, and complementarity-determining regions from a murine (rat) monoclonal antibody (CAMPATH-1G). The CAMPATH-1H antibody has an approximate molecular weight of 150 kD. CAMPATH is produced in mammalian cell (Chinese hamster ovary) suspension culture in a medium containing neomycin. Neomycin is not detectable in the final product.

CAMPATH (alemtuzumab) injection is a sterile, clear, colorless, isotonic solution (pH 6.8–7.4) in a single-dose vial for intravenous use. Each single-dose vial of CAMPATH contains 30 mg alemtuzumab, 8.0 mg sodium chloride, 1.44 mg dibasic sodium phosphate, 0.2 mg potassium chloride, 0.2 mg monobasic potassium phosphate, 0.1 mg polysorbate 80, and 0.0187 mg disodium edetate dihydrate. No preservatives are added.

12 CLINICAL PHARMACOLOGY

12.1 Mechanism of Action

CAMPATH binds to CD52, an antigen present on the surface of B and T lymphocytes, a majority of monocytes, macrophages, NK cells, and a subpopulation of granulocytes. A proportion of bone marrow cells, including some CD34+ cells, express variable levels of CD52. The proposed mechanism of action is antibody-dependent cellular-mediated lysis following cell surface binding of CAMPATH to the leukemic cells.

12.2 Pharmacodynamics

Cardiac Electrophysiology

The effect of multiple doses of alemtuzumab (12 mg/day for 5 days) on the QTc interval was evaluated in a single-arm study in 53 patients without malignancy. No large changes in the mean QTc interval (i.e., >20 ms) were detected in the study. A mean increase in heart rate of 22 to 26 beats/min was observed for at least 2 hours following the initial infusion of alemtuzumab. This increase in heart rate was not observed with subsequent doses.

12.3 Pharmacokinetics

CAMPATH pharmacokinetics were characterized in a study of 30 previously treated B-CLL patients in whom CAMPATH was administered at the recommended dose and schedule. After 12 weeks of dosing, patients exhibited a 7-fold increase in mean AUC.

Distribution

After the last 30 mg dose, the mean volume of distribution at steady-state was 0.18 L/kg (range 0.1 to 0.4 L/kg).

Elimination

CAMPATH pharmacokinetics displayed nonlinear elimination kinetics. Systemic clearance decreased with repeated administration due to decreased receptor-mediated clearance (i.e., loss of CD52 receptors in the periphery). Mean half-life was 11 hours (range 2 to 32 hours) after the first 30 mg dose and was 6 days (range 1 to 14 days) after the last 30 mg dose.

Specific Populations

The effects of renal or hepatic impairment on the pharmacokinetics of CAMPATH have not been studied.

13 NONCLINICAL TOXICOLOGY

13.1 Carcinogenesis, Mutagenesis, Impairment of Fertility

Studies to assess the carcinogenic or genotoxic potential of CAMPATH have not been conducted.

In fertility studies, alemtuzumab (3 or 10 mg/kg intravenously) was administered to huCD52 transgenic male mice on 5 consecutive days prior to cohabitation with untreated wild-type females. No effect on fertility or reproductive performance was observed. However, adverse effects on sperm parameters (including abnormal

morphology [detached/no head] and reduced total count and motility) were observed at both doses tested.

When alemtuzumab (3 or 10 mg/kg intravenously) was administered to huCD52 transgenic female mice for 5 consecutive days prior to cohabitation with untreated wild-type males, there was a decrease in the average number of corpora lutea and implantation sites and an increase in postimplantation loss, resulting in fewer viable embryos at the higher dose tested.

14 CLINICAL STUDIES

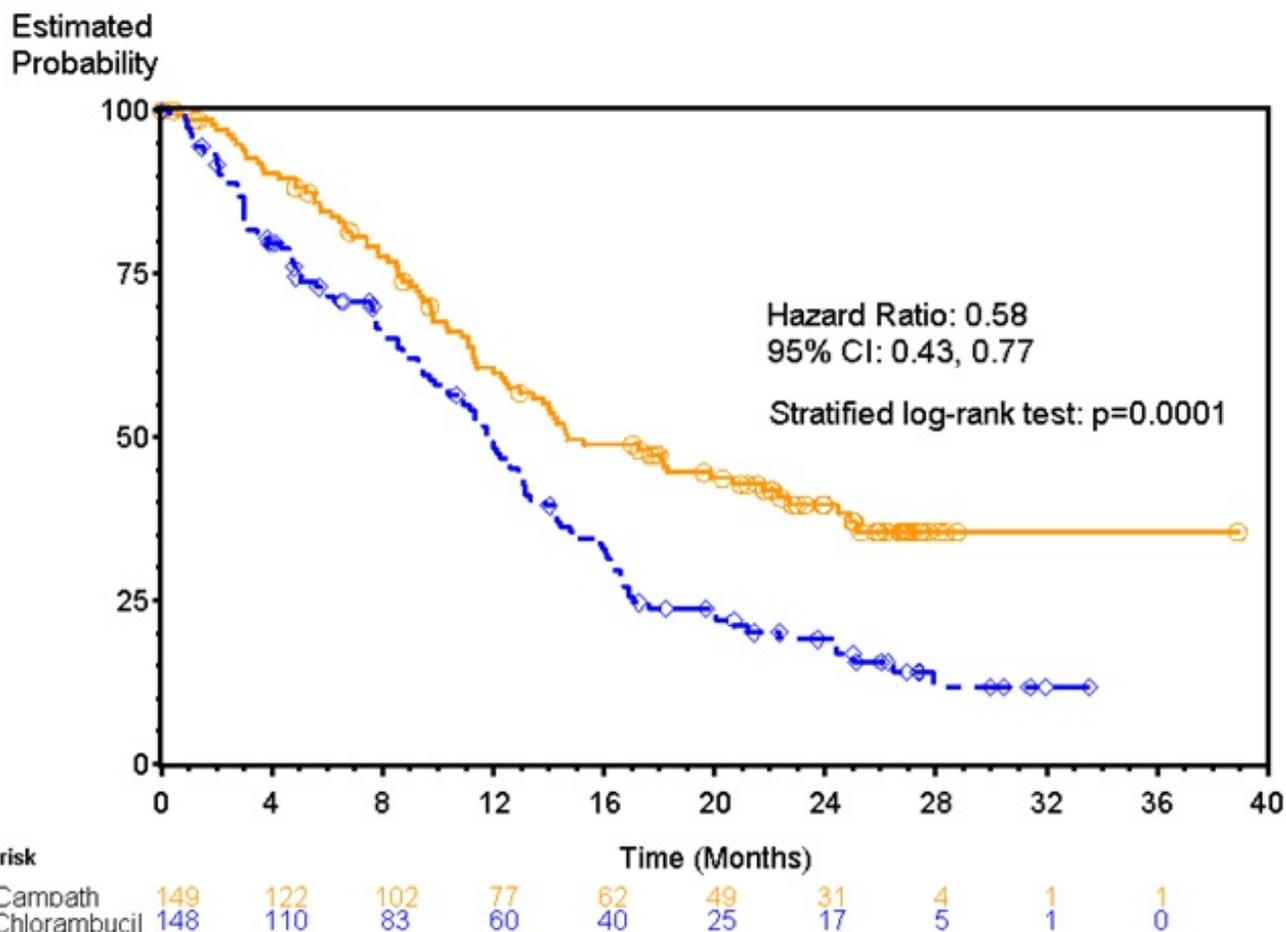
14.1 Previously Untreated B-CLL Patients

CAMPATH was evaluated in an open-label, randomized (1:1) active-controlled study in previously untreated patients with B-CLL, Rai Stage I-IV, with evidence of progressive disease requiring therapy. Patients received either CAMPATH 30 mg intravenously 3 times per week for a maximum of 12 weeks or chlorambucil 40 mg/m² orally once every 28 days for a maximum of 12 cycles.

Of the 297 patients randomized, the median age was 60 years, 72% were male, 99% were Caucasian, 96% had a WHO performance status 0-1, 23% had maximum lymph node diameter \geq 5 cm, 34% were Rai Stage III/IV, and 8% were treated in the U.S.

Patients randomized to receive CAMPATH experienced longer progression free survival (PFS) compared to those randomized to receive chlorambucil (median PFS 14.6 months vs. 11.7 months, respectively). The overall response rates were 83% and 55% ($p < 0.0001$) and the complete response rates were 24% and 2% ($p < 0.0001$) for CAMPATH and chlorambucil arms, respectively. The Kaplan-Meier curve for PFS is shown in Figure 1.

Figure 1: Progression Free Survival in Previously Untreated B-CLL Patients*



* Log-rank test adjusted for Rai Stage (I-II vs. III-IV).

14.2 Previously Treated B-CLL Patients

CAMPATH was evaluated in three multicenter, open-label, single-arm studies of 149 patients with B-CLL previously treated with alkylating agents, fludarabine, or other chemotherapies. Patients were treated with the recommended dose of CAMPATH 30 mg intravenously 3 times per week for up to 12 weeks. Partial response rates of 21% to 31% and complete response rates of 0% to 2% were observed.

16 HOW SUPPLIED/STORAGE AND HANDLING

CAMPATH (alemtuzumab) is supplied in clear glass single-dose vial containing 30 mg of alemtuzumab in 1 mL of solution. Each carton contains three CAMPATH vials (NDC 58468-0357-3) or one CAMPATH vial (NDC 58468-0357-1).

Store CAMPATH at 2°C to 8°C (36°F to 46°F). Do not freeze. If accidentally frozen, thaw at 2°C to 8°C before administration. Protect from direct sunlight.

17 PATIENT COUNSELING INFORMATION

Cytopenias

Advise patients to report any signs or symptoms such as bleeding, easy bruising, petechiae or purpura, pallor, weakness or fatigue [see *Warnings and Precautions (5.1)* and *Adverse Reactions (6.1)*].

Infusion-Related Reactions

Advise patients of the signs and symptoms of infusion-related reactions and of the need to take premedications as prescribed [see *Warnings and Precautions (5.2) and Adverse Reactions (6.1)*].

Immunosuppression/Infections

Advise patients to immediately report symptoms of infection (e.g., pyrexia) and to take prophylactic anti-infectives for PCP (trimethoprim/sulfamethoxazole DS or equivalent) and for herpes virus (famciclovir or equivalent) as prescribed [see *Warnings and Precautions (5.3) and Adverse Reactions (6.1)*].

Advise patients that irradiation of blood products is required [see *Warnings and Precautions (5.3)*].

Immunization

Advise patients that they should not be immunized with live viral vaccines if they have recently been treated with CAMPATH. Advise females with infants exposed to CAMPATH in utero to inform the pediatrician of the exposure [see *Warnings and Precautions (5.4)*].

Embryo-Fetal Toxicity

Advise pregnant women and females of reproductive potential of the potential risk to a fetus. Advise females to inform their healthcare provider of a known or suspected pregnancy [see *Use in Specific Populations (8.1, 8.3)*].

Advise female patients of reproductive potential to use effective contraception during treatment with CAMPATH and for 3 months after the final dose [see *Use in Specific Populations (8.1, 8.3)*].

Lactation

Advise females not to breastfeed during treatment with CAMPATH and for 3 months after the final dose [see *Use in Specific Populations (8.2)*].

Infertility

Advise females and males of reproductive potential that CAMPATH may impair fertility [see *Use in Specific Populations (8.3) and Nonclinical Toxicology (13.1)*].

Glomerular Nephropathies

Advise patients on signs and symptoms of glomerular nephropathies that may occur months to years after receiving CAMPATH [see *Adverse Reactions (6.2)*].

Manufactured and distributed by:
Genzyme Corporation, Cambridge, MA 02141
A SANOFI COMPANY
US License Number: 1596

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PRINCIPAL DISPLAY PANEL - 1 Vial Carton

NDC 58468-0357-1

Rx only

Campath[®]

alemtuzumab

Injection

30 mg/mL

For Intravenous Infusion After Dilution

Sterile - No U.S. Standard of Potency

One single-dose vial

Discard unused portion

sanofi

FPO

For Intravenous Infusion After Dilution

30 mg/mL

Campath[®]
alemtuzumab
Injection

NDC 58468-0357-1 Rx only

Information. See full prescribing
information.
Store refrigerated at 2°C to 8°C (36°F to 46°F).
Do not freeze. If accidentally frozen, thaw at 2°C to 8°C.
Protect from direct sunlight.
DO NOT SHAKE THE VIAL.

NDC 58468-0357-1

Rx only

Campath[®]
alemtuzumab
Injection

30 mg/mL

For Intravenous Infusion After Dilution

Sterile – No U.S. Standard of Potency

One single-dose vial
Discard unused portion

sanofi

6220

FPO

XXXXXX

Area reserved for
serialization data/LOT/EXP.
EXP date format: MMM YYYY

MMM is alpha
YYYY is numeric

Each single-dose vial contains 30 mg alemtuzumab,
8.0 mg sodium chloride, 1.44 mg dibasic sodium
phosphate, 0.2 mg potassium chloride, 0.2 mg
monobasic potassium phosphate, 0.1 mg polysorbate 80,
and 0.0187 mg disodium edetate dihydrate.
Contains no preservatives.

sanofi

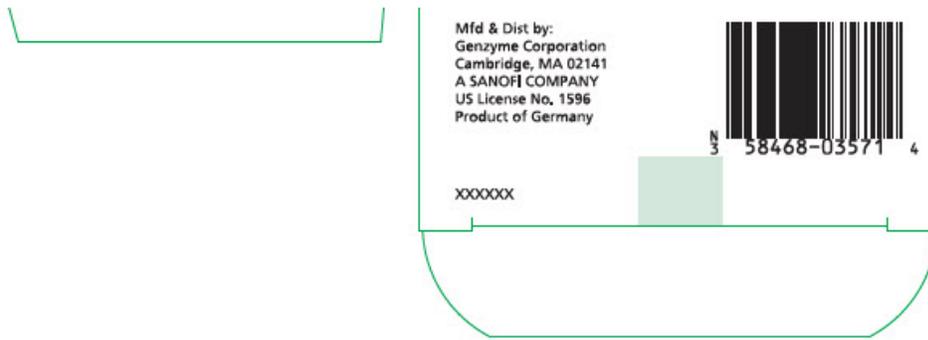
NDC 58468-0357-1

Rx only

Campath[®]
alemtuzumab
Injection

30 mg/mL

For Intravenous Infusion After Dilution



PRINCIPAL DISPLAY PANEL - 3 Vial Carton

NDC 58468-0357-3

Rx only

Campath®
alemtuzumab
Injection

30 mg/mL

For Intravenous Infusion After Dilution

Sterile - No U.S. Standard of Potency

Three single-dose vials
Discard unused portion

sanofi

FPO

For Intravenous Infusion After Dilution

30 mg/mL

Injection
alemtuzumab
Campath[®]

NDC 58468-0357-3 Rx only

Dose and Administration: See full prescribing information.
Store refrigerated at 2°C to 8°C (36°F to 46°F). Do not freeze. If accidentally frozen, thaw at 2°C to 8°C. Protect from direct sunlight. DO NOT SHAKE THE VIAL.

NDC 58468-0357-3 Rx only

Campath[®]
alemtuzumab
Injection

30 mg/mL

For Intravenous Infusion After Dilution

Sterile – No U.S. Standard of Potency

Three single-dose vials
Discard unused portion

sanofi

6220

FPO

XXXXXX

Area reserved for serialization data/LOT/EXP.
EXP date format: MMM YYYY

MMM is alpha
YYYY is numeric

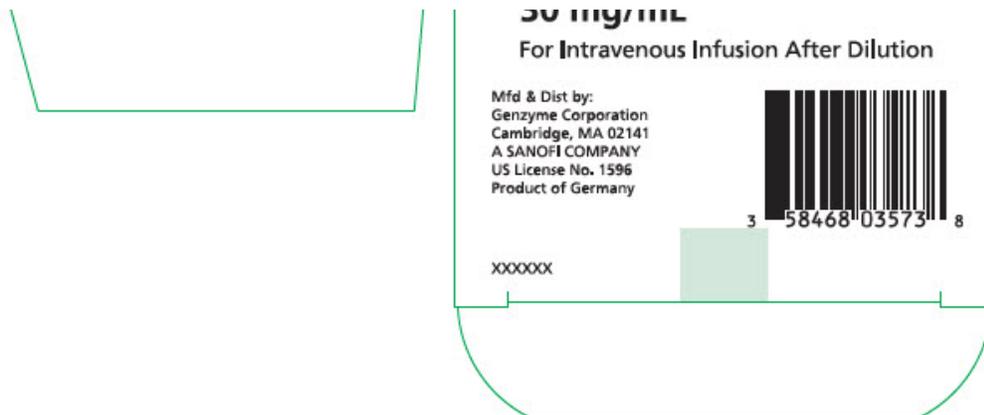
Each single-dose vial contains 30 mg alemtuzumab, 8.0 mg sodium chloride, 1.44 mg dibasic sodium phosphate, 0.2 mg potassium chloride, 0.2 mg monobasic potassium phosphate, 0.1 mg polysorbate 80, and 0.0187 mg disodium edetate dihydrate. Contains no preservatives.

sanofi

NDC 58468-0357-3 Rx only

Campath[®]
alemtuzumab
Injection

30 mg/mL



CAMPATH

alemtuzumab injection

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:58468-0357
Route of Administration	INTRAVENOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALEMTUZUMAB (UNII: 3A189DH42V) (ALEMTUZUMAB - UNII:3A189DH42V)	ALEMTUZUMAB	30 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM CHLORIDE (UNII: 451W47IQ8X)	8 mg in 1 mL
SODIUM PHOSPHATE, DIBASIC, UNSPECIFIED FORM (UNII: GR686LBA74)	1.44 mg in 1 mL
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	0.2 mg in 1 mL
MONOBASIC POTASSIUM PHOSPHATE (UNII: 4J9FJ0HL51)	0.2 mg in 1 mL
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	0.1 mg in 1 mL
EDETATE DISODIUM (UNII: 7FLD91C86K)	0.0187 mg in 1 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:58468-0357-1	1 in 1 CARTON	11/30/2009	
1		1 mL in 1 VIAL, SINGLE-USE; Type 0: Not a Combination Product		
2	NDC:58468-0357-3	3 in 1 CARTON	11/30/2009	
2		1 mL in 1 VIAL, SINGLE-USE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA103948	11/30/2009	

Labeler - Genzyme Corporation (025322157)

Establishment

Name	Address	ID/FEI	Business Operations
Boehringer Ingelheim Pharma GmbH & Co. KG		340700520	ANALYSIS(58468-0357) , API MANUFACTURE(58468-0357) , MANUFACTURE(58468-0357)

Establishment

Name	Address	ID/FEI	Business Operations
Sanofi-Aventis Deutschland GmbH		313218430	ANALYSIS(58468-0357)

Establishment

Name	Address	ID/FEI	Business Operations
EUROAPI UK LIMITED		229522842	PACK(58468-0357) , LABEL(58468-0357)

Revised: 5/2023

Genzyme Corporation