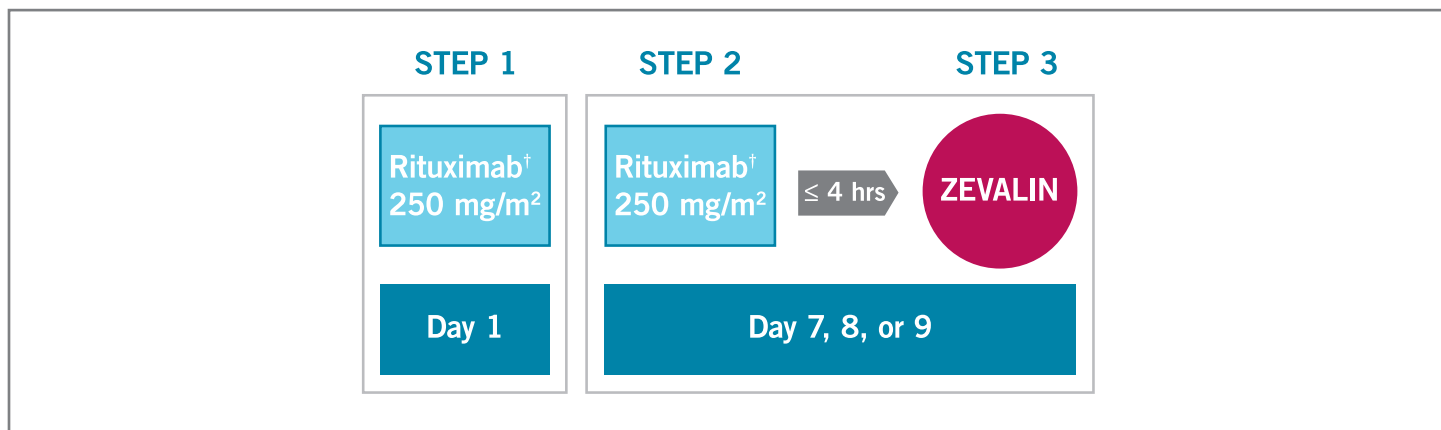


Important Ordering Information for ZEVALIN Administration Facilities

ZEVALIN[®]
ibritumomab tiuxetan
INJECTION FOR INTRAVENOUS USE

The FDA has recently removed the requirement to complete an Indium-111 bioscan prior to ZEVALIN administration.

ZEVALIN treatment has been simplified*



Discontinue rituximab and ZEVALIN infusions in patients who develop severe infusion reactions or severe cutaneous or mucocutaneous reactions.

The Ordering Process for ZEVALIN has NOT changed

- Place orders for ZEVALIN through your radiopharmacy
 - » Orders for ZEVALIN must be placed at least 8 business days prior to ZEVALIN administration
 - » At time of order, you will be asked to provide:
 - ZEVALIN administration date
 - Purchase Order made out to Spectrum Pharmaceuticals, Inc.
- Notify your radiopharmacy of any cancellations or schedule changes immediately

For more information, please contact:

ZEVALIN Support Services

TEL: 866-298-8433

FAX: 877-264-8483

zevalinsupport@sppirx.com

Monday-Friday

8:30 am–8:00 pm EST

ZEVALIN provides flexible treatment scheduling for your practice and patients

Please see additional Important Safety Information on the reverse side.

Please see enclosed full Prescribing Information, including **BOXED WARNINGS**, for ZEVALIN.

* Although no longer required, administration sites may choose to continue to conduct the biodistribution scan by injecting In-111 on Day 1, followed by a scan on Day 3 or 4. If so, imaging date must be provided at time of order.

[†] Pretreatment with acetaminophen 650 mg and diphenhydramine 50 mg orally

© 2011 Spectrum Pharmaceuticals, Inc. All Rights Reserved. May not be reproduced, altered, or distributed without express permission. SPECTRUM PHARMACEUTICALS, INC.[®] and ZEVALIN[®] are registered trademarks of Spectrum Pharmaceuticals, Inc. and its subsidiaries. The Spectrum Pharmaceuticals logo and ZEVALIN logo are trademarks owned by Spectrum Pharmaceuticals, Inc. and its subsidiaries.

Single ZEVALIN Treatment. It's about time.



www.ZEVALIN.com

Indications and Usage

ZEVALIN® (ibritumomab tiuxetan) is a CD20-directed radiotherapeutic antibody administered as part of the ZEVALIN therapeutic regimen indicated for the treatment of patients with:

- Relapsed or refractory, low-grade or follicular B-cell non-Hodgkin's lymphoma (NHL)
- Previously untreated follicular NHL who achieve a partial or complete response to first-line chemotherapy

Important Safety Information

WARNING: SERIOUS INFUSION REACTIONS, PROLONGED AND SEVERE CYTOPENIAS, and SEVERE CUTANEOUS AND MUCOCUTANEOUS REACTIONS

Serious Infusion Reactions: Deaths have occurred within 24 hours of rituximab infusion, an essential component of the ZEVALIN therapeutic regimen. These fatalities were associated with hypoxia, pulmonary infiltrates, acute respiratory distress syndrome, myocardial infarction, ventricular fibrillation, or cardiogenic shock. Most (80%) fatalities occurred with the first rituximab infusion. Discontinue rituximab and Y-90 ZEVALIN infusions in patients who develop severe infusion reactions.

Prolonged and Severe Cytopenias: Y-90 ZEVALIN administration results in severe and prolonged cytopenias in most patients. Do not administer Y-90 ZEVALIN to patients with $\geq 25\%$ lymphoma marrow involvement and/or impaired bone marrow reserve.

Severe Cutaneous and Mucocutaneous Reactions: Severe cutaneous and mucocutaneous reactions, some fatal, can occur with the ZEVALIN therapeutic regimen. Discontinue rituximab and Y-90 ZEVALIN infusions in patients experiencing severe cutaneous or mucocutaneous reactions.

Dosing: The dose of Y-90 ZEVALIN should not exceed 32.0 mCi (1184 MBq).

Leukemia and Myelodysplastic Syndrome: Among 204 patients receiving Y-90-ZEVALIN following first-line chemotherapy, two patients (1%) were diagnosed with AML within 3 years of receiving ZEVALIN.

Myelodysplastic syndrome (MDS) and/or acute myelogenous leukemia (AML) were reported in 5.2% (11/211) of patients with relapsed or refractory NHL enrolled in clinical studies and 1.5% (8/535) of patients included in the expanded-access trial, with median follow-up of 6.5 and 4.4 years, respectively. Among the 19 reported cases, the median time to diagnosis of MDS or AML was 1.9 years following treatment with the ZEVALIN therapeutic regimen; however, the cumulative incidence continues to increase.

Embryo-fetal Toxicity: May cause fetal harm if given during pregnancy.

Extravasation: Monitor for extravasation and terminate infusion if it occurs. Resume infusion in another limb.

Immunization: Do not administer live viral vaccines to patients who recently received ZEVALIN.

Laboratory Monitoring: Obtain complete blood counts (CBC) and platelet counts at least weekly.

Radionuclide Precautions: During and after radiolabeling ZEVALIN with Y-90, minimize radiation exposure to patients and to medical personnel, consistent with institutional good radiation safety practices and patient management procedures.

Creutzfeldt-Jakob Disease (CJD): The ZEVALIN therapeutic regimen contains albumin, a derivative of human blood. Based on effective donor screening and product manufacturing processes, ZEVALIN carries an extremely remote risk for transmission of viral diseases. A theoretical risk for transmission of CJD also is considered extremely remote. No cases of transmission of viral diseases or CJD have ever been identified for albumin.

Impairment of Fertility: There is a potential risk that the ZEVALIN therapeutic regimen could cause toxic effects on the male and female gonads. Effective contraceptive methods should be used during treatment and for up to 12 months following the ZEVALIN therapeutic regimen.

Nursing Mothers: Patients should be advised to discontinue nursing during and after ZEVALIN treatment.

Adverse Reactions: The most common adverse reactions of ZEVALIN are cytopenias, fatigue, abdominal pain, nausea, nasopharyngitis, asthenia, diarrhea, cough, and pyrexia. Common adverse reactions ($\geq 40\%$) in clinical trials were: neutropenia, leukopenia, thrombocytopenia, anemia, infection, asthenia, musculoskeletal symptoms, and gastrointestinal symptoms. The most serious adverse reactions of ZEVALIN are prolonged and severe cytopenias (thrombocytopenia, anemia, lymphopenia, neutropenia) and secondary malignancies.

When administered following first-line chemotherapy, grade 3/4 adverse reactions of ZEVALIN include prolonged and severe cytopenias (thrombocytopenia [51%], neutropenia [41%], leukopenia [36%], lymphopenia [18%], and anemia [5%]) and secondary malignancies. Cytopenias were more severe and more prolonged among eleven (5%) patients who received ZEVALIN after first-line fludarabine or a fludarabine-containing chemotherapy regimen as compared to patients receiving non-fludarabine-containing regimens. Grade 3/4 infections occurred in 8% of ZEVALIN-treated patients and in 2% of controls and included neutropenic sepsis (1%), bronchitis, catheter sepsis, diverticulitis, herpes zoster, influenza, lower respiratory tract infection, sinusitis, and upper respiratory tract infection.

Grade 3/4 adverse reactions of ZEVALIN in relapsed or refractory NHL patients include prolonged and severe cytopenias (thrombocytopenia [63%], neutropenia [60%], anemia [17%], and ecchymosis [$<1\%$]) and secondary malignancies. Serious infections occurred in 3% of patients (urinary tract infection, febrile neutropenia, sepsis, pneumonia, cellulitis, colitis, diarrhea, osteomyelitis, and upper respiratory tract infection). Life-threatening infections were reported in 2% of patients (sepsis, empyema, pneumonia, febrile neutropenia, fever, and biliary stent-associated cholangitis).

Please see enclosed full Prescribing Information, including BOXED WARNINGS, for ZEVALIN. Because the ZEVALIN therapeutic regimen includes the use of rituximab, please also consult Prescribing Information for rituximab.