

# FULLY FUNDED CLINICAL TRIAL FOR CANINE B-CELL LYMPHOMA

## Study Goal

To assess an advanced new therapy specifically designed to treat canine lymphoma. The primary goal of the study is to measure the extension of survival time of cancer dogs treated with the novel therapy when compared to untreated historical controls.

## Background

Lymphoma is one of the most common blood cancers affecting dogs of different type of breeds and ages. Lymphoma is a devastating disease that, when untreated, has a survival of 4-6 weeks. New treatment options/additions are needed.

The clinical trial treatment is not a chemotherapeutic agent. It is a product designed specifically for canine lymphoma and has been shown in preliminary trials to be safe in dogs.

This type of treatment has been used for the treatment of lymphoma in humans. It improves survival in people when used in conjunction with chemotherapy and was a groundbreaking advancement.

## Eligibility

- Newly diagnosed case of lymphoma
- Adult dog of any breed preferably less than <40kg (80 lbs)
- Owner must give written signed and dated consent to participate

## Sequence of Events

- Candidates are evaluated and confirmed to meet the inclusion criteria by undergoing a complete physical examination, lymph node assessment, body weight, blood sample for lymphocyte level, CBC/Chemistry/Urinalysis, and fine needle aspiration.
- Patients will then begin to receive intravenous injections of the new product for 8 weeks.
- Patients will be closely monitored throughout the treatment process.

With the exception of first exam and consultation required to establish eligibility, this is a fully funded study.



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